

# **VA/DoD CLINICAL PRACTICE GUIDELINE FOR MANAGEMENT OF POST-TRAUMATIC STRESS**

**Department of Veterans Affairs**

**Department of Defense**

Prepared by:

**The Management of Post-Traumatic Stress Working Group**

With support from:

**The Office of Quality and Performance, VA, Washington, DC**

**&**

**Quality Management Division, United States Army MEDCOM**

## **QUALIFYING STATEMENTS**

The Department of Veterans Affairs (VA) and The Department of Defense (DoD) guidelines are based on the best information available at the time of publication. They are designed to provide information and assist in decision-making. They are not intended to define a standard of care and should not be construed as one. Also, they should not be interpreted as prescribing an exclusive course of management.

Variations in practice will inevitably and appropriately occur when providers take into account the needs of individual patients, available resources, and limitations unique to an institution or type of practice. Every healthcare professional making use of these guidelines is responsible for evaluating the appropriateness of applying them in any particular clinical situation.

**Version 2.0 – 2010**

## Table of Contents

|  | <i>Page</i> |
|--|-------------|
| <b>Introduction</b>  | <b>3</b>    |
| <b>Guideline Update Working Group</b>  | <b>11</b>   |
| <b>ALGORITHMS AND ANNOTATIONS</b>  |             |
| <b>CORE Module: Post-traumatic Stress, Screening</b>   | <b>14</b>   |
| <b>Module A: Management of Acute Stress Reaction,<br/>            and Prevention of PTSD</b> | <b>29</b>   |
| <b>Module B: Management of Post-Traumatic Stress<br/>            Disorder (PTSD)</b>         | <b>60</b>   |
| <b>TREATMENT INTERVENTIONS</b>   |             |
| <b>Module I: Treatment Interventions for<br/>            Post-Traumatic Stress</b>           | <b>106</b>  |
| I1 – Early Intervention to prevent PTSD  |             |
| I2 – Treatment of PTSD   |             |
| I3 – Management of Specific Symptoms   |             |
| <b>APPENDICES</b>  |             |
| <b>Appendix A. Guideline Development Process</b>   | <b>210</b>  |
| <b>Appendix B. Acronym List</b>  | <b>217</b>  |
| <b>Appendix C. PTSD Screening Tools</b>  | <b>220</b>  |
| <b>Appendix D. Participant List</b>  | <b>224</b>  |
| <b>Appendix E. Bibliography</b>  | <b>230</b>  |

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2  
3  
4  
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## INTRODUCTION

This update of the Clinical Practice Guideline for the Management of Post-Traumatic Stress was developed under the auspices of the Veterans Health Administration (VHA) and the Department of Defense (DoD) pursuant to directives from the Department of Veterans Affairs (VA). VHA and DoD define clinical practice guidelines as:

*“Recommendations for the performance or exclusion of specific procedures or services derived through a rigorous methodological approach that includes:*

*• Determination of appropriate criteria such as effectiveness, efficacy, population benefit, or patient satisfaction; and, Literature review to determine the strength of the evidence in relation to these criteria.”*

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This 2010 VA/DoD Post-Traumatic Stress guideline update builds on the 2004 VA/DoD *Clinical Practice Guideline for the Management of Post-Traumatic Stress*. The 2004 Post-Traumatic Stress Guideline was the first effort to bring evidence-based practice to clinicians providing care to trauma survivors and patients with stress disorders in VA and DoD. The development of the Guideline originated with recognition of the need to diagnose and treat Post-Traumatic Stress among the military and veteran population. The Guideline presented evidence-based recommendations that were thoroughly evaluated by practicing clinicians and reviewed by clinical experts from the VHA and DoD.

### Algorithms:

The VA/DoD also utilized an algorithmic approach for the 2004 Guideline for the Management Post-Traumatic Stress. This guideline update has also been developed using an algorithmic approach to guide the clinician in determining care and the sequencing of the interventions on a patient specific basis. The clinical algorithm incorporates the information presented in the guideline in a format which maximally facilitates clinical decision-making. The use of the algorithmic format was chosen because such a format improves data collection, facilitates diagnostic and therapeutic decision-making, and changes in patterns of resource use. However, *this should not prevent providers from using their own clinical expertise in the care of an individual patient*. Guideline recommendations are intended to support clinical decision-making and should never replace sound clinical judgment.

During the past 6 years a number of well-designed randomized controlled trials of pharmacological and psychotherapeutic interventions for post-traumatic stress have been conducted. Therefore, the goal of this update is to integrate the results of this research and update the recommendations of the original guideline to reflect the current knowledge of effective treatment intervention. As in the original guideline, this update will explore the most important research areas of intervention to prevent the development of PTSD in persons who have developed stress reaction symptoms after exposure to trauma.

### Target Population:

This guideline applies to adult patients with traumatic stress reaction treated in any VA or DoD clinical setting.

### Audiences:

The guideline is relevant to all healthcare professionals providing or directing treatment services to patients with traumatic stress at any VA/DoD healthcare setting.

### Post-Traumatic Stress:

Post-traumatic stress consists of a spectrum of traumatic stress disorders, hence this Clinical Practice Guideline for the Management of Post-traumatic Stress. These disorders can be arranged along a temporal

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40 axis from Acute Stress Reaction, to Acute Stress Disorder, Acute PTSD, and Chronic PTSD. Each of these  
41 may be associated with serious mental and physical comorbidities. Some survivors will experience only a  
42 part of this spectrum while others will progress through the entire range.

43 **Acute stress reaction (ASR)**, which in military settings is often labeled as a **Combat and Operational**  
44 **Stress Reaction (COSR)**, reflects acute reactions to a high stress or combat related event. ASR can present  
45 with a broad group of physical, mental and emotional symptoms and signs (e.g., depression, fatigue,  
46 anxiety, decreased concentration/memory, hyperarousal and others) that have not resolved within 4 days  
47 after the event, and after ruling out other disorders.

48 Acute stress disorder (ASD) occurs when the individual has experienced trauma(s) as described above, has  
49 symptoms lasting more than 2 days, but less than one month after exposure to the trauma (may progress to  
50 PTSD if symptoms last >1 month) and exhibits at least three out of five dissociative symptoms.

51 Post-traumatic stress disorder (PTSD) is a clinically significant condition with symptoms continuing more  
52 than 1 month after exposure to a trauma that has caused significant distress or impairment in social,  
53 occupational, or other important areas of functioning. Patients with PTSD may exhibit persistent re-  
54 experiencing of the traumatic event(s), persistent avoidance of stimuli associated with the trauma, numbing  
55 of general responsiveness (not present before the trauma), and persistent symptoms of increased arousal  
56 (not present before the trauma). PTSD can also have a delayed onset which is described as a clinically  
57 significant presentation of symptoms (causing significant distress or impairment in social, occupational, or  
58 other important areas of functioning) at least 6 months after exposure to trauma.

59 PTSD is further sub-divided into Acute PTSD (symptoms lasting more than one month, but less than three  
60 months after exposure to trauma) and Chronic PTSD (symptoms lasting more than three months after  
61 exposure to trauma). PTSD can appear as *Simple* (presenting with common symptoms of PTSD), or  
62 *Complicated by other co-occurring conditions* (persistent difficulties in interpersonal relations, mood,  
63 chronic pain, sleep disturbances, somatization and profound identity problems), or psychiatric disorders  
64 (meeting DSM criteria for another disorder such as substance abuse, depression, or anxiety disorder).

65 OEF/OIF veterans and service members who have sustained a concussion (mild-TBI) in the combat  
66 environment are at significantly greater risk of PTSD. Moreover, the diagnosis of either condition may be  
67 complicated by the fact that PTSD is associated with generalized health symptoms including  
68 neurocognitive impairment and the other symptoms considered in the persistent postconcussion syndrome  
69 definition.

70 Evidence-based practices to prevent and treat PTSD include screening, cognitive behavioral therapies, and  
71 medications. There are many new strategies involving enhancement of cognitive fitness and psychological  
72 resilience to reduce the detrimental impact of trauma. In terms of screening; evidence suggests that  
73 identifying PTSD early, and quickly referring people to treatment, can shorten their suffering and lessen the  
74 severity of their functional impairment. Several types of cognitive behavioral therapies, counseling, and  
75 medications have been shown to be effective in treating PTSD.

76 The VA and DoD Health care systems have undergone significant changes in the past 10-15 years that is  
77 transforming the two into an integrated system that provides high quality care. In response to the increased  
78 demands for services to treat OEF/OIF veterans with PTSD, these systems have invested resources in  
79 expanding outreach activities enhancing the availability and timeliness of specialized PTSD services.  
80 Nevertheless, access to care is still unacceptably variable and service members continue to face barriers to  
81 seeking care such as stigma and access difficulties (e.g., difficulty scheduling or getting to appointments).

## 82 **Post-Traumatic Stress in VA population:**

83 The numbers of veterans seeking and receiving treatment for post-traumatic stress in general, and PTSD in  
84 particular, continues to increase. In a follow-up to a study by Dohrenwend et al. (2006) 9.1% of Vietnam  
85 veterans sampled still suffered from symptoms of PTSD in 1990. During a five year span (2004-2008) the  
86 number of unique veterans seeking help for PTSD in the VA system increased from 274,000 to 442,000.  
87 Also, according to a review of several studies investigating the prevalence of PTSD in U.S Veterans of the  
88 first Persian Gulf War, the Board on Population Health and Public Health Practice at the National  
89 Academies of Science (2008) reported that PCL based prevalence of PTSD in a sample of 11,441 veterans

90 was 12.1%. This review also cited evidence that ten years after the 1990 Gulf War 6.2% of a sample of  
91 veterans still suffered from PTSD.

92 The number of Iraq and Afghanistan veterans separated from service and seen in U.S. Veteran's  
93 Administration health care facilities and diagnosed with PTSD was reported by Seal and colleagues (2007).  
94 Of the 103,788 Veterans included in this record review, the overall prevalence of PTSD was 13%, higher  
95 than any other mental health diagnostic category reported by these authors'. The VA's Uniform Services  
96 Handbook sets standards for mental health care across VA facilities and is intended to both, improve  
97 quality of care, and also facilitate implementation of evidence-based practice. The combination of  
98 improved diagnostic and treatment techniques for all stress related disorders, the needs of veterans from  
99 past wars as far back as Vietnam, the co-morbid conditions many veterans experience in addition to PTSD  
100 (chronic medical conditions, SUD), and the on-going nature of the current wars in Iraq and Afghanistan  
101 with improved survival rates and increased rates of PTSD and comorbid conditions have the obvious effect  
102 of exponentially increasing the necessity of services/programs throughout the VA system to effectively  
103 address the needs of veterans.

#### 104 **Post-Traumatic Stress in DoD population:**

105 A number of studies have been conducted to estimate the prevalence and incidence of PTSD in military  
106 personnel during the Iraq and Afghanistan wars. These studies have shown high consistency in rates, when  
107 grouped according to study type (e.g., studies involving Army or Marine combat infantry units versus  
108 studies involving samples of the deploying population at large, including personnel from support units or  
109 services not involved in direct combat).

110 One of the first and most cited epidemiological surveys to provide estimates of PTSD prevalence in  
111 military personnel who served in Afghanistan or Iraq was published by Hoge et al. (2004). The prevalence  
112 of PTSD 3 months post-deployment among infantry soldiers and Marines who returned from high intensity  
113 combat in Iraq was 12.9% and 12.2% respectively (n = 894 soldiers, 815 Marines) based on a stringent  
114 definition for PTSD supported in a study by Terhakopian et al. (2008) (PCL score of at least 50 combined  
115 with DSM criteria). By comparison the rate among soldiers who had deployed to Afghanistan, where there  
116 was very low intensity combat at that time, was 6.2%, and the baseline rate in a group of soldiers before  
117 deployment was 5%. This study also highlighted the impact that stigma and barriers to care have on  
118 willingness to receive help. Less than half of soldiers in need of mental health services received care, and  
119 many reported concerns that they would be treated differently by peers or leaders if they sought care.

120 In a subsequent survey involving active and National Guard brigade combat teams (infantry), rates of 15%  
121 were documented 3 months post-deployment, and rose to 17-25% 12 months post-deployment using the  
122 same definitions as in the 2004 article (Thomas J, et al., 2010 Arch Gen Psychiatry).

123 In-theater assessments of personnel in ground combat units have been conducted on nearly an annual basis  
124 in Iraq and several times in Afghanistan since the start of the wars (Army Mental Health Advisory Team  
125 Assessments –MHATs). These studies have found rates of acute stress or PTSD (based on the PCL  $\geq$ 50  
126 points) of 10-20%, with a strong correlation to combat frequency and intensity. Rates in units exposed to  
127 minimal combat are similar to baseline rates in the population (5%) and there is a linear increase up to 25%  
128 in units involved in the highest intensity combat. The Afghanistan theater showed lower rates earlier in the  
129 war (7% in 2005), but this increased to levels comparable with Iraq in 2007 and subsequently due to  
130 increased combat intensity.

131 In addition to studies based on infantry samples, there have been a number of studies based on post-  
132 deployment health assessments, health care utilization records, and random samples of military or veteran  
133 populations, including those not engaged in direct combat (Hoge, Auchterlonie, Milliken, 2006; Milliken,  
134 Auchterlonie, Hoge 2007; Rand Corporation 2008; Smith, Ryan, Wingard, Slymen, Sallis, & Kritz-  
135 Silverstein, 2008; Fear NT, et al., 2010; and others). General population samples that do not focus  
136 specifically on combat units have resulted in lower rates than reported in infantry samples, but estimates  
137 approach infantry samples when analyses are restricted to Army or Marine personnel with combat  
138 experience. While most studies have focused on point prevalence of PTSD, one study has looked at the  
139 three-year incidence in a large representative population sample (Smith, Ryan, Wingard, et al., 2008). The  
140 cumulative incidence was 9% in Army personnel who had experienced combat, which would equate to a  
141 prevalence of approximately 12%, including those excluded for PTSD at baseline. Overall, baseline pre-

142 deployment rates in military samples have ranged from 3-6%, comparable to civilian rates reported in the  
143 National Comorbidity study (Kessler et al., 2009); and post-deployment rates have ranged from 6-20%.  
144 The strongest predictor of increased prevalence post-deployment has been combat frequency and intensity.  
145 There are also many other types of traumatic experience that service members encounter, both in their  
146 professional military occupations and in their pre-military or off-duty time, including exposure to accidents,  
147 assault, rape, natural disasters, and other experiences.

#### 148 **Outcome Measures:**

149 The Working Group (WG) agreed on the following health related outcomes for management of post-  
150 traumatic stress:

- 151 • Improvement in quality of life (social and occupational functioning)
  - 152 • Reduce morbidity/mortality
  - 153 • Improvement over long term
  - 154 • Patient Satisfaction
  - 155 • Comorbidity
  - 156 • Improvement of symptoms
- 157

#### 158 **Guideline Goals:**

159 The most important goal of the VA/DoD Clinical Practice Guideline for the Management of Post-  
160 Traumatic Stress is to provide a scientific evidence-base for practice evaluations and interventions. The  
161 guideline was developed to assist facilities to implement processes of care that are evidence-based and  
162 designed to achieve maximum functionality and independence as well as improve patient and family  
163 quality of life. Related specifics include;

- 164 • To identify the critical decision points in management of patients with post traumatic stress disorder
- 165 • To allow flexibility so that local policies or procedures, such as those regarding referrals to, or  
166 consultation with, specialty care (mental health care), can be accommodated
- 167 • To decrease the development of complications and comorbidity
- 168 • To improve patient outcomes, i.e., reduce symptoms, decrease comorbidity, increase functional status  
169 and enhance the quality of life

#### 170 **Development Process:**

171 The development process of this guideline follows a systematic approach described in “Guideline-for-  
172 Guidelines,” an internal working document of the VA/DoD Evidence-Based Practice Working Group that  
173 requires an ongoing review of the work in progress. [Appendix A](#) clearly describes the guideline  
174 development process followed for this guideline.

175 The Offices of Quality and Performance and Patient Care Service of the VA, in collaboration with the  
176 network Clinical Managers, and the Medical Center Command of the DoD identified clinical leaders to  
177 champion the guideline development process. During a preplanning conference call, the clinical leaders  
178 defined the scope of the guideline and identified a group of clinical experts from the VA and DoD that  
179 formed the Guideline Development Working Group.

180 At the start of the update process, the clinical leaders, guideline panel members, outside experts, and  
181 experts in the field of guideline and algorithm development were consulted to determine which aspects of  
182 the 2004 guideline required updating. These consultations resulted in the following recommendations that  
183 guided the update efforts: (1) update any recommendations from the original guideline likely to be effected  
184 by new research findings; (2) provide information and recommendations on health systems changes  
185 relevant to management of post-traumatic stress (3) address content areas and models of treatment for  
186 which little data existed during the development of the original guideline; and (4) review the performance  
187 and lessons learned since the implementation of the original guideline

#### 188 **Review of Literature and Evidence:**

189 Recommendations for the performance or inclusion of specific procedures or services in this guideline were  
190 derived through a rigorous methodological approach that included the following:

- 191 • Determining appropriate criteria such as effectiveness, efficacy, population benefit, or patient  
192 satisfaction
- 193 • Performing a comprehensive literature search and selection of relevant studies from January 2002 to  
194 August, 2009 to identify the best available evidence and ensure maximum coverage of studies at the  
195 top of the hierarchy of study types
- 196 • Reviewing the selected studies to determine the strength of the evidence in relation to these criteria
- 197 • Formulating the recommendations and grading the level of evidence supporting each recommendation

198 This 2010 update builds on the 2004 version of the guideline and incorporates information from the  
199 following existing evidence-based guidelines/reports identified by the Working Group as appropriate seed  
200 documents:

- 201 • ISTSS (2009), Effective Treatments for PTSD: Practice Guidelines from the International Society for  
202 Traumatic Stress Studies. Foa EB, Keane TM, Friedman MJ, Cohen J (Eds) 2009. New York: Guilford  
203 Press.
- 204 • IOM (2007), Institute of Medicine (IOM). 2008. Treatment of posttraumatic stress disorder: An  
205 assessment of the evidence. Washington, DC: The National Academies Press.
- 206 • APA (2009), Practice Guideline for the Treatment of Patients with Acute Stress Disorder and  
207 Posttraumatic Stress Disorder: Guideline Watch, March 2009

208 Literature searches were conducted covering the period from January 2002 through August, 2009 that  
209 combined terms for post-traumatic stress, acute stress reaction (ASR), acute stress disorder (ASD), acute  
210 post-traumatic stress disorder, and chronic post-traumatic stress disorder. Electronic searches were  
211 supplemented by reference lists and additional citations suggested by experts. The identified and selected  
212 studies on those issues were critically analyzed, and evidence was graded using a standardized format. The  
213 evidence rating system for this document is based on the system used by the U.S. Preventive Services Task  
214 Force (USPSTF, 2007).

215 If evidence exists, the discussion following the recommendations for each annotation includes an evidence  
216 table identifying the studies that have been considered, the quality of the evidence, and the rating of the  
217 strength of the recommendation [SR]. The Strength of Recommendation, based on the level of the evidence  
218 and graded using the USPSTF rating system (see Table: Evidence Rating System), is presented in brackets  
219 following each guideline recommendation.

## 220 Evidence Rating System

| SR |   |
|----|---|
| A  | A strong recommendation that clinicians provide the intervention to eligible patients.<br>Good evidence was found that the intervention improves important health outcomes and concludes that benefits substantially outweigh harm.   |
| B  | A recommendation that clinicians provide (the service) to eligible patients.<br>At least fair evidence was found that the intervention improves health outcomes and concludes that benefits outweigh harm.  |
| C  | No recommendation for or against the routine provision of the intervention is made.<br>At least fair evidence was found that the intervention can improve health outcomes, but concludes that the balance of benefits and harms is too close to justify a general recommendation. |
| D  | Recommendation is made against routinely providing the intervention to asymptomatic patients.<br>At least fair evidence was found that the intervention is ineffective or that harms outweigh benefits.   |
| I  | The conclusion is that the evidence is insufficient to recommend for or against routinely providing the intervention.<br>Evidence that the intervention is effective is lacking, of poor quality, or conflicting, and the balance of benefits and harms cannot be determined.     |

221 *SR = Strength of recommendation*

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223 Where existing literature was ambiguous or conflicting, or where scientific data was lacking on an issue,  
224 recommendations are based on the clinical experience of the Working Group. Although several of the  
225 recommendations in this guideline are based on weak evidence, some of these recommendations are  
226 strongly recommended based on the experience and consensus of the clinical experts and researchers of the  
227 Working Group. Recommendations that are based on consensus of the Working Group include a  
228 discussion of the expert opinion on the given topic. No [SR] is presented for these recommendations. A  
229 complete bibliography of the references in this guideline can be found in [Appendix E](#)

230 This Guideline is the product of many months of diligent effort and consensus building among  
231 knowledgeable individuals from the VA, DoD, and a guideline facilitator from the private sector. An  
232 experienced moderator facilitated the multidisciplinary Working Group. The draft document was discussed  
233 in [two](#) face-to-face group meetings. The content and validity of each section was thoroughly reviewed in a  
234 series of conference calls. The final document is the product of those discussions and has been approved  
235 by all members of the Working Group.

236 The list of participants is included in [Appendix D](#) to the guideline.

### 237 **Implementation:**

238 The guideline and algorithms are designed to be adapted by individual facilities in consideration of local  
239 needs and resources. The algorithms serve as a guide that providers can use to determine best interventions  
240 and timing of care for their patients in order to optimize quality of care and clinical outcomes.

241 Although this guideline represents the state of the art practice on the date of its publication, medical  
242 practice is evolving and this evolution requires continuous updating of published information. New  
243 technology and more research will improve patient care in the future. The clinical practice guideline can  
244 assist in identifying priority areas for research and optimal allocation of resources. Future studies  
245 examining the results of clinical practice guidelines such as these may lead to the development of new  
246 practice-based evidence.

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### 248 **KEY POINTS ADDRESSED BY THIS GUIDELINE**

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1. Triage and management of acute traumatic stress
  2. Routine primary care screening for trauma and related symptoms
  3. Diagnosis of trauma syndromes and comorbidities
  4. Evidence-based management of trauma related symptoms and functioning
  5. Collaborative patient/provider decision making, education, and goal setting
  6. Coordinated and sustained follow-up
  7. Identification of major gaps in current knowledge
  8. Outline for psychological care in ongoing military operations
  9. Proactive strategies to promote resilience and prevent trauma related stress disorders
  10. Standardized longitudinal care (DoD/VA, Primary Care/Mental Health)



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**OVERVIEW OF GUIDELINE UPDATE**

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This clinical practice guideline updates the 2004 version of the VA/DoD Guideline on Management of Post-traumatic Stress.

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The Working Group (WG) developed a revised comprehensive clinical algorithm. The objective of the VA/DoD Working Group in developing this revision was to incorporate the accumulating experience in the field and information from the original guideline recommendations into a format that would maximally facilitate clinical decision-making. Randomized controlled trials and systematic reviews published in the past 5 years were identified and have been carefully appraised and included in the analysis of the evidence for this update. Promoting evidence-based treatment ultimately enhances and optimizes treatment outcomes, thus contributing to optimal care across institutional boundaries and promoting a smooth transition of care between the DOD and the VA health care systems.

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The current revision incorporates the four Modules of the 2004 guideline into two modules focusing on acute stress reactions/acute stress disorder and PTSD. Where evidence suggests differences in the management of Acute Stress Reactions (ASR), Acute Stress Disorder (ASD), and Posttraumatic Stress Disorder (PTSD), specific treatment intervention recommendations are provided.

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This effort drew heavily from the International Society for Traumatic Stress Studies clinical practice guideline (Foa et al., 2009). The VA/DoD Working Group reviewed this guideline and made the decision to adopt several of their evidence-based recommendations. Identified randomized controlled trials and systematic reviews published in the past 7 years have been carefully appraised and included in the analysis of the evidence for this update.

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The first Module incorporates the assessment, diagnosis, and management of symptoms of **Acute Stress Reaction (ASR)** in the immediate period after exposure to trauma, the management of **Acute Stress Disorder (ASD)**, and the effective early interventions to prevent progression of stress reactions to full PTSD.

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Additional recommendations were added addressing specific issues that the WG considered to be of importance to patients in the healthcare systems of the VA and DoD. Hence, this VA/DoD updated version of the Post Traumatic Stress guideline includes evidence-based recommendations for routine primary care in non-combat environments and additional recommendations suggesting specific actions for assessment and management of **Combat and Operational Stress Reactions (COSR)**. These specific recommendations may better serve providers caring for service persons with symptoms among the active duty population.

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Key recommendations for acute and early intervention include:

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- Trials assessing the use of beta-blocker, propranolol, anxiolytic, anticonvulsant, or gabapentin showed no significant benefit over placebo on depressive or posttraumatic stress symptoms. Thus, the recommendations from the original guideline suggesting pharmacotherapy interventions to prevent PTSD have been removed.

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- Brief early cognitive behavior therapy that includes exposure and cognitive processing components is strongly recommended, for symptomatic trauma survivors, as it has been shown to accelerate recovery and prevent progression to PTSD.

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- Single session psychological debriefing, in individual or group formats, is not recommended.

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Consensus based recommendations are added to the 2010 revision of the CPG regarding specific adjunct treatment intervention that target specific symptoms frequently seen in patient with acute stress reactions (beyond the core symptoms of ASD/PTSD). These include sleep disturbance, pain, and hyper-agitation. These consensus-based recommendations are aimed to help the primary care practitioners and others to provide brief symptom-focused treatment to help patients adhere to the core treatment. If there is no improvement in these symptoms, patients should be referred to specialty care for evaluation and treatment.

306 The second Module addresses the diagnosis and management of patients with **Post Traumatic Stress**  
307 **Disorder (PTSD)**. The WG revised the algorithm for this module in a patient-centered approach that  
308 emphasizes the decisions and interventions shown to be effective in treating PTSD, regardless of the  
309 treatment setting. This approach should allow the use of the guideline as a starting point for innovative  
310 plans that improve collaborative efforts and focus on key aspects of care. The recommendations outlined in  
311 this guideline should serve as a framework for the care that is provided in specialty mental health care  
312 settings and primary care. The optimal setting of care for the individual patient will depend on patient  
313 preferences, the level of expertise of the provider, and available resources.

314 Key recommendations for treatment of PTSD include;

- 315 • Patient and provider preferences should drive the selection of evidence-based psychotherapy  
316 and/or evidence-based pharmacotherapy as the first-line treatment
- 317 • Education provided to patients and their families should be one of the first steps of PTSD  
318 treatment and should continue throughout treatment
- 319 • First line psychotherapeutic treatments that have been shown to be comparatively effective for  
320 patients with PTSD include Exposure-based Therapy [ET],; Cognitive-based Therapy [CT],;  
321 Stress Inoculation Training [SIT], and Eye Movement Desensitization and Reprocessing [EMDR]
- 322 • First line pharmacotherapy includes the use of antidepressants, particularly serotonergic reuptake  
323 inhibitors that have been found to be effective in treating PTSD (i.e., SSRIs, for which fluoxetine,  
324 paroxetine or sertraline have the strongest support, or SNRIs, for which venlafaxine has the  
325 strongest support)
- 326 • There is evidence that treatment effect sizes and adherence to cognitive behavioral therapy (CBT)  
327 is similar whether the treatment is delivered via Tele-psychiatry or by traditional means.
- 328 • The revised guideline includes a section describing alternative modalities of adjunctive treatment  
329 for PTSD, in which the following recommendations are made:
  - 330 - Acupuncture may be considered as an efficacious and acceptable non-exposure treatment  
331 option for PTSD
  - 332 - Complementary Alternative Medicine (CAM) can be effective in eliciting a relaxation  
333 response and reducing physiological hyperarousal, but should not be considered evidence-  
334 based PTSD treatments. They may be useful as adjunctive modalities and may help foster  
335 engagement approaches have not been demonstrated to be effective and cannot be  
336 recommended. When patients decline standard mental health interventions, alternative  
337 approaches may foster engagement

338 The WG recognizes that PTSD is often accompanied by other psychiatric conditions. Such comorbidities  
339 require clinical attention at the point of diagnosis and throughout the process of treatment. Disorders of  
340 particular concern are **substance-use disorder**, **major depression**, and post-concussive symptoms  
341 attributed to **mild TBI**. The WG also recognizes the fact that most clinical trials excluded patients with  
342 comorbid psychiatric illnesses and no trials have been published that can provide guidance on how to  
343 manage PTSD complicated by comorbid illness. The revised guideline includes recommendations based on  
344 the experience and opinion of the experts providing suggestion for the approach to treatment of PTSD in  
345 the presence of comorbid psychiatric conditions.

346 Finally, clinicians following these updated guidelines should not limit themselves only to the approaches  
347 and techniques addressed in the guideline. All current treatments have limitations; either not all patients  
348 respond to them, patients drop out of treatment, or providers are not comfortable using a particular  
349 intervention. To promote the development of improved treatments, creative integration of combined  
350 treatments that are driven by sound evidence-based principles are encouraged in the field.

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**PTS GUIDELINE WORKING GROUP**

| <b>VHA</b>   | <b>DoD</b>   |
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| 1  | <b>TABLE OF CONTENTS</b>  |    |
| 2  | <b>CORE MODULE: ALGORITHM</b>   | 14 |
| 3  | <b>CORE MODULE: ANNOTATIONS</b>   | 15 |
| 4  | <b>1. PRIMARY PREVENTION</b> .....                                      | 15 |
| 5  | A. Education and Training Fostering Resilience                          | 15 |
| 6  | <b>2. POPULATIONS AT-RISK FOR DEVELOPING PTSD</b> .....                 | 17 |
| 7  | B. Person Exposed to Trauma   | 17 |
| 8  | <b>3. SECONDARY PREVENTION</b> .....                                    | 19 |
| 9  | C. Screen for PTSD Symptoms   | 19 |
| 10 | D. Are Trauma Related Symptoms Present?                                 | 22 |
| 11 | E. Educate about additional care if needed, provide contact information | 26 |
| 12 | <b>Module A: ACUTE STRESS REACTION (ASR)</b>                            |    |
| 13 | <b>and Prevention of Post-Traumatic Stress Disorder (PTSD)</b>          | 29 |
| 14 | <b>ALGORITHM</b>  | 31 |
| 15 | <b>ANNOTATIONS</b>  | 32 |
| 16 | <b>4. ASSESSMENT &amp; TRIAGE</b> .....                                 | 32 |
| 17 | A. Trauma Exposure (within the past 30 days)                            | 32 |
| 18 | B. Assess briefly based on general appearance and behavior              | 33 |
| 19 | C. Unstable, Dangerous to Self or Others                                | 35 |
| 20 | D. Ensure Basic Physical Needs Are Met                                  | 36 |
| 21 | E. Person has Trauma Related Symptoms, Significant                      |    |
| 22 | Impaired Function, or Diagnosis of ASD                                  | 41 |
| 23 | F. Assess Medical and Functional Status                                 | 44 |
| 24 | G. Assess Pre-Existing Psychiatric and Medical Conditions               | 45 |
| 25 | H. Assess Risk Factors for Developing ASD/PTSD                          | 45 |
| 26 | Provide Education and Normalization / Expectancy of Recovery            | 46 |
| 27 | <b>5. TREATMENT</b> .....   | 48 |
| 28 | J. Initiate Brief Intervention  | 48 |
| 29 | K. Acute Symptom Management   | 51 |
| 30 | L. Facilitate Social and Spiritual Support                              | 52 |
| 31 | <b>6. RE-ASSESSMENT</b> .....   | 55 |
| 32 | M. Reassess Symptoms and Function                                       | 55 |
| 33 | <b>7. FOLLOW-UP</b> .....   | 56 |
| 34 | N. Persistent (>1 Month) or Worsening Symptoms, Significant Functional  |    |
| 35 | Impairment or High Risk for Development of PTSD.                        | 56 |
| 36 | O. Monitor and Follow-Up  | 58 |

|    |   |           |
|----|---|-----------|
| 37 | <b>MODULE B: Management of Post-traumatic Disorder</b>                  |           |
| 38 | <b>ALGORITHM</b>  | 60        |
| 39 | <b>ANNOTATIONS</b>  | 62        |
| 40 | <b>1. Assessment</b> .....  | <b>62</b> |
| 41 | A. Assessment of Trauma Exposure-Related Symptoms                       | 62        |
| 42 | B. Assessment of Trauma Exposure  | 64        |
| 43 | C. Assessment of Dangerousness to Self or Others                        | 65        |
| 44 | D. Obtain Medical History, Physical Examination/Laboratory              |           |
| 45 | Tests, and Psychosocial Assessment                                      | 69        |
| 46 | E. Assessment of Functioning/Fitness for Duty                           | 73        |
| 47 | F. Assessment of Risk/Protective Factors                                | 74        |
| 48 | <b>2. Triage</b> .....  | <b>81</b> |
| 49 | G. Are There Clinically Significant Symptoms Suggestive Of PTSD?        | 81        |
| 50 | H. Assess for Co-Occurring Disorders                                    | 83        |
| 51 | I. Educate Patient and Family   | 86        |
| 52 | J. Determine Optimal Setting for Management of PTSD                     |           |
| 53 | and Co-Occurring Disorders  | 88        |
| 54 | J1. Management of PTSD with Comorbidity                                 | 88        |
| 55 | J2. Concurrent PTSD and Substance Abuse                                 | 90        |
| 56 | J3. The Role of the Primary Care Practitioner                           | 92        |
| 57 | K. Facilitate Social and Spiritual Support [See Module A: annotation L] | 94        |
| 58 | L. Assess Duty/Work Responsibilities and Patient’s Fitness              | 94        |
| 59 | <b>3. Treatment</b> .....   | <b>96</b> |
| 60 | M. Initiate Treatment Using Effective Interventions for PTSD            | 96        |
| 61 | <b>4. Re-ASSESSMENT</b> .....   | <b>99</b> |
| 62 | O. Assess response to treatment   | 99        |
| 63 | P. Follow-Up  | 100       |
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**CORE MODULE: ALGORITHM**

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**CORE MODULE: ANNOTATIONS**

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**1. PRIMARY PREVENTION**

7

**A. Education and Training Fostering Resilience**

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**OBJECTIVE**

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Prepare individuals and groups for exposure to traumatic experiences in ways that minimize the likelihood of development of PTSD and other trauma-related problems.

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**BACKGROUND**

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Because exposure to traumatic stressors is part of the expected work experience of some occupations (e.g., military personnel and emergency services workers), it is sensible to make efforts to prepare individuals in these professions for their encounters with traumatic events. This preparation is not explicitly undertaken in most workplaces, with some exceptions (e.g., some military training environments). To date, research has not examined our capacity to prepare individuals or communities for trauma exposure. However, general principles of preparation can be outlined that are consistent with theoretical models of the development of PTSD, research on risk factors for development of PTSD, and emerging concepts of resilience and hardiness.

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**RECOMMENDATIONS**

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1. In high-risk occupations for which probability of trauma exposure is moderate or high, efforts should be undertaken to increase psychological resilience of workers to the negative effects of trauma exposure.

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**DISCUSSION**

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Although little is directly known about our capacity to prepare individuals or communities for trauma exposure, it is possible to identify principles of preparation that are consistent with empirical research on risk and resilience factors and with current theories of PTSD development. Such pre-trauma preparation can include attention to both the ability to cope during the trauma itself and to shaping the post-trauma environment so that it will foster post-trauma adaptation.

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Some influential theories of PTSD posit that a process of classical fear conditioning can lead to development of chronic PTSD symptomatology. In this process, stimuli associated with the traumatic experience can elicit responses similar to those experienced during the trauma itself (e.g., intense anxiety). Other theories suggest that individuals who develop negative trauma-related beliefs (e.g., about personal guilt) will be more likely to experience continuing trauma reactions because such beliefs will maintain a sense of threat and personal incompetence. Research on risk factors for PTSD indicates that post-trauma social support and life stress affect the likelihood of development of the disorder. Protective factors have also been identified that mitigate the negative effects of stress. Research is beginning to delineate the psychological processes that moderate an individual's response to stress, and to explore training programs for increasing resilience to stress. Hardiness (Kobasa, Maddi, & Kahn, 1982) is one personality factor that has been demonstrated to buffer against traumatic stress and PTSD in military veterans (King et al., 1998; Bartone, 2000). Hardiness is characterized by three key attributes: ability to perceive *control* over life's events; ability to make strong *commitment* to

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49 tasks; and ability to see stressful experiences as a *challenge* to be overcome.  
50 Training programs, personnel policies, and leadership strategies that promote  
51 hardiness may thereby increase an individual's ability to resist the negative effects of  
52 traumatic stress.

53 Such findings and theories are consistent with the following principles of preparation:

- 54 1. *Provide realistic training* that includes vicarious, simulated, or actual exposure to  
55 traumatic stimuli that may be encountered. Examples of application of this  
56 principle in military training include exposure to live weapons fire, survival  
57 training, or, for subgroups of military personnel, mock captivity training. This  
58 principle can be applied to many work roles; for example, those likely to be  
59 involved in body handling might be trained in mortuary environments. It is  
60 consistent with classical conditioning theories in that this can help reduce arousal  
61 or anxiety associated with particular traumatic stimuli.
- 62 2. *Strengthen perceived ability to cope* during the trauma and with the aftermath.  
63 Realistic training contributes to this goal. Instruction and practice in the use of a  
64 variety of coping skills (e.g., stress inoculation training, problem-solving,  
65 assertion, and cognitive restructuring) may be helpful in enabling workers to  
66 tolerate stressful work environments. In addition, individuals can be trained to  
67 cope with acute stress reactions that are common following trauma exposure.  
68 Such training experiences help to maximize expectations of mastery of traumatic  
69 situations and their physical and emotional sequelae. The training must include  
70 specific, practical actions to do to change the threatening or horrifying situation  
71 for the better. Without such positive action learning, "simulated" terrifying or  
72 horrifying situations and stimuli can induce feelings of helplessness that make the  
73 training itself traumatizing.
- 74 3. *Create supportive interpersonal work environments* that are likely to provide  
75 significant social support during and after traumatic events. Efforts to build  
76 teams and establish group cohesion among work group members are important  
77 in this regard. Identification and training of peer stress management resource  
78 persons, and training and practice in the provision of peer social support may  
79 also be useful. Families are crucial in post-trauma support and can be given  
80 information about, and training in, ways of providing social support. Finally,  
81 competent, ethical leadership at all levels of the organization helps protect  
82 against traumatization.
- 83 4. *Develop and maintain adaptive beliefs* about the work role and traumatic  
84 experiences that may be encountered within it. Key beliefs will be related to  
85 realistic expectancies about the work environment, confidence in leadership,  
86 confidence in the meaningfulness or value of the work role, positive but realistic  
87 appraisals of own coping ability, and knowledge about the commonness and  
88 transitory nature of most acute stress reactions. It may be useful to identify and  
89 discuss negative beliefs that sometimes arise in the specific work environment, in  
90 order to "inoculate" against such beliefs.
- 91 5. *Develop workplace-specific comprehensive traumatic stress management*  
92 *programs*. Such programs can be a significant source of post-trauma support  
93 (e.g., via mental health professionals) that can minimize trauma-related  
94 problems among workers. It is important to take steps to increase awareness of  
95 such services and to de-stigmatize and reduce potential negative consequences  
96 of their use. For example, employees should be helped to understand that  
97 seeking help in confronting symptoms and problems early in their development is



98 likely to be more effective than avoiding them or keeping them secret from  
99 others, but that even long hidden or persisting PTSD can be treated.

100 Comprehensive preparation programs that target and integrate these principles and  
101 that are, themselves, integrated into existing unit/community support systems may  
102 be expected to be most helpful (Gist & Lubin, 1999).

### 103 EVIDENCE

|   | Evidence   | Sources                 | LE  | QE   | SR |
|---|--|-------------------------|-----|------|----|
| 1 | Foster resilience by preparing workers in high-risk occupations for trauma exposure. | Working Group Consensus | III | Poor | I  |

104 *LE=Level of Evidence; QE = Quality of Evidence; SR =Strength of Recommendation (see Appendix A)*

## 105 2. POPULATIONS AT-RISK FOR DEVELOPING PTSD

### 106 B. Person Exposed to Trauma

#### 107 OBJECTIVE

108 Identify persons at risk for developing a traumatic stress disorder (PTSD) after  
109 trauma exposure.

#### 110 BACKGROUND

111 Although exposure to trauma is common, several risk factors for the development of  
112 PTSD have been identified. Risk factors for developing PTSD can be grouped as  
113 characteristics related to the trauma itself, pre-trauma factors and post-trauma  
114 factors.

- 115 • *Trauma*-related risks include the nature, severity, and duration of the trauma  
116 exposure. For example, life-threatening traumas such as physical injury or rape  
117 pose a high risk of PTSD (Kilpatrick, 1989). A prior history of trauma exposure  
118 conveys a greater risk of PTSD from subsequent trauma (Breslau et al., 1999).
- 119 • *Pre-trauma* risk factors include adverse childhood, younger age, female gender  
120 (not in military cohorts), minority race, and low socioeconomic or educational  
121 status.
- 122 • *Post-trauma* risks include poor social support and life stress (Brewin et al., 2000).  
123 A greater risk for developing PTSD may be conveyed by post-trauma factors (e.g.,  
124 lack of social support and additional life stress) than pre-trauma factors.

#### 125 RECOMMENDATIONS

- 126 1. Persons exposed to trauma should be assessed for known risk factors for  
127 developing PTSD – both pre-trauma risks and post trauma risks.
- 128 2. The trauma type, nature, and severity should be assessed.
- 129 3. Assessment of existing social supports and ongoing stressors is important.
- 130 4. Patients with Acute Stress Disorder (ASD) warrant careful clinical attention, as  
131 they are at high-risk for developing PTSD.
- 132 5. Patients with persisting dissociative symptoms may also warrant careful clinical  
133 attention.

134

**DISCUSSION**

135 A meta-analysis of risk factors for PTSD of assessed studies of trauma-exposed  
136 adults reported that 14 different risk factors in the literature have a modest  
137 association with PTSD development (Brewin et al., 2000). Overall, factors such as  
138 gender, age at trauma, and race predicted PTSD in some populations, but not in  
139 others. Further, factors such as education, prior trauma, and childhood adversity  
140 predicted PTSD more consistently (Harvey & Bryant, 2000; Harvey & Bryant, 1998).  
141 However, this varies with the population and study methods. Prior psychiatric  
142 history, childhood abuse, and family psychiatric history have more consistent  
143 predictive effects. Factors operating during or after the trauma (e.g., trauma  
144 severity, lack of social support, and additional life stress) have somewhat stronger  
145 effects than pre-trauma factors. This finding is consistent with other studies that  
146 suggest poor social supports and ongoing life stress to be predictors of PTSD  
147 development. This may have clinical implications as early interventions that increase  
148 social support after trauma exposure may reduce the likelihood of PTSD (Litz et al.,  
149 2002).

150 The development of Acute Stress Disorder (ASD) at the time of the trauma is also a  
151 risk for developing PTSD (Classen et al., 1998). Numerous prospective cohort studies  
152 with various types of trauma exposure (e.g., violent assault and accidents) support  
153 that ASD is a predictor of later PTSD (Brewin et al., 1999; Bryant et al., 2000;  
154 Harvey & Bryant, 1998; Mellman et al., 2001). In these studies among persons with  
155 ASD, 40 to 80 percent did develop PTSD. Finally, most studies suggest an increased  
156 risk of PTSD development among individuals with peritraumatic dissociation (Birmes  
157 et al., 2001; Murray et al., 2002). Subsequent research indicates that it is the  
158 posttraumatic duration of dissociation, rather than peritraumatic occurrence of  
159 dissociation (Panasetis & Bryant, 2003), that predicts the development of PTSD.

160

**EVIDENCE**

|   | Evidence  | Sources   | LE | QE   | SR |
|---|---|---|----|------|----|
| 1 | Assessment of persons exposed to trauma for risk factors for developing PTSD (pre-trauma and post-trauma risks).      | Brewin et al., 2000   | II | Good | B  |
| 2 | Assessment of trauma type, nature and severity.   | Brewin et al., 1999<br>Bryant et al., 2000<br>Harvey & Bryant, 2000<br>Mellman et al., 2001   | II | Good | B  |
| 3 | Assessment of existing social supports and ongoing stressors.   | Litz et al., 2002   | II | Good | B  |
| 4 | Patients with dissociative symptoms or ASD warrant careful clinical attention due to a high risk for developing PTSD. | Birmes et al., 2001<br>Brewin et al., 1999<br>Bryant et al., 2000<br>Harvey & Bryant, 2000<br>Mellman et al., 2001<br>Murray et al., 2002 | II | Good | B  |
| 5 | Patients with dissociative symptoms warrant careful clinical attention  | Brewin et al., 1999<br>Murray et al., 2002  | II | Fair | C  |

161

LE – Level of Evidence QE = Quality of Evidence; SR = Strength of Recommendation (see Appendix A)

162 **3. SECONDARY PREVENTION**163 **C. Screen for PTSD Symptoms**

---

164 **OBJECTIVE**

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165 Identify possible cases of post-traumatic stress

166 **BACKGROUND**

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167 Patients don't often self-identify as suffering with PTSD, and patients with  
168 unrecognized PTSD are often difficult to treat because of poor patient/provider  
169 rapport, anger and distrust, a focus on somatic symptoms, and other trauma-related  
170 problems. Research supports the utility of brief screening tools for identifying  
171 undiagnosed cases of PTSD. Identification of PTSD may help facilitate development  
172 of rapport, suggest treatment options, and potentially improve outcomes for these  
173 patients.

174 **RECOMMENDATIONS**

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- 175 1. All new patients should be screened for symptoms of PTSD initially and then on  
176 an annual basis or more frequently if clinically indicated due to clinical suspicion,  
177 recent trauma exposure (e.g., major disaster), or history of PTSD.
- 178 2. Patients should be screened for symptoms of PTSD using paper and pencil or  
179 computer-based screening tools.
- 180 3. There is insufficient evidence to recommend one PTSD screening tool versus  
181 another. However, the following screening tools have been validated and should  
182 be considered for use. For example: (See Appendix C)
  - 183 - Primary Care PTSD Screen (PC-PTSD)
  - 184 - PTSD Brief Screen
  - 185 - Short Screening Scale for DSM IV PTSD
  - 186 - PTSD Checklist (PCL).
- 187 4. There is insufficient evidence to recommend special screening for members of  
188 any cultural or racial groups.

189 **DISCUSSION**

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190 The benefit of screening is well established for diseases with high prevalence. In one  
191 study (Taubman et al., 2001) 23 percent of patients presenting in the primary care  
192 setting reported exposure to traumatic events and 39 percent of those met criteria  
193 for PTSD. Screening strategies should, however, balance efficacy with practical  
194 concerns (e.g., staffing, time constraints, and current clinical practices). Brevity,  
195 simplicity, and ease of implementation should encourage compliance with  
196 recommended screening. Care should be exercised in implementing screening in  
197 ways that avoid social stigmatization and adverse occupational effects of positive  
198 screens.

199 Brewin (2005) reviewed published screening instruments for civilian PTSD consisting  
200 of 30 items or fewer and that validated against structured clinical interviews.  
201 Thirteen instruments were identified meeting these criteria, all consisting of  
202 symptoms of traumatic stress. The review concluded that the performance of some  
203 currently available instruments is near to their maximal potential effectiveness, and  
204 that instruments with fewer items, simpler response scales, and simpler scoring  
205 methods perform as well as if not better than longer and more complex measures.

206  
207**208 Screening Tools: (See Appendix C)**

209 *Primary Care PTSD Screen (PC-PTSD):* This is a 4-item screen that was designed for  
210 use in primary care and other medical settings and is currently used to screen for  
211 PTSD in veterans at the VA. The screen includes an introductory sentence to cue  
212 respondents to traumatic events. The authors suggest that in most circumstances  
213 the results of the PC-PTSD should be considered "positive" if a patient answers "yes"  
214 to any 3 items. Those screening positive should then be assessed with a structured  
215 interview for PTSD. The screen does not include a list of potentially traumatic events  
216 (Prins et al., 2002, 2003). Internal consistency (KR2- = .79) and test-retest reliability  
217 (r. -.84) of the PC-PTSD was found as good (Prins et al., 1999). The operating  
218 characteristics of the screen suggest that the overall efficiency (i.e., optimal  
219 sensitivity and specificity = .87) is best when any two items are endorsed. The PC-  
220 PTSD screen has been validated in a military population (Bliese, et al.,) and has been  
221 used extensive in post-deployment screening efforts (Hoge, 2006)

222 *PTSD Brief Screen (Leskin et al., 1999):* The PTSD Brief Screen was developed using  
223 the rationally derived approach based on data from the National Comorbidity Survey.  
224 Construct validity has generally been adequate. The overall efficiency of this screen  
225 was good (.78), whereas the correlations were significantly lower or negative for  
226 other mental disorders indicating good construct validity.

227 *PTSD Checklist (PCL):* The PCL has been used extensively in military and civilian  
228 populations and there are numerous validation studies, including studies in military  
229 populations (Terhakopian et al., 2008).

**230 Screening for Sexual Trauma:**

231 Within the Veterans Health Administration, every new woman veteran must be  
232 screened for history of sexual trauma according to public law (see screening for  
233 sexual trauma at <http://www.va.gov/publ/direc/health/direct/12000008.html>).

234 The passage of Public Law (Pub. L.) 102-585, in 1992, authorized the Department of  
235 Veterans Affairs (VA) to include outreach and counseling services for women  
236 veterans who experienced incidents of sexual trauma while they served on active  
237 duty in the military. The law defines sexual trauma as sexual harassment, sexual  
238 assault, rape and other acts of violence. It further defines sexual harassment as  
239 repeated unsolicited verbal or physical contact of a sexual nature, which is  
240 threatening in nature. NOTE: This law was later amended by Pub. L. 103-452, which  
241 authorizes the VA to provide counseling to men as well as women. The Veterans  
242 Millennium Health Care Act, signed on November 30, 1999, has significant  
243 implications under Section 115, Counseling and Treatment for Veterans Who Have  
244 Experienced Sexual Trauma. Provisions of Pub. L. 106-117. Section 115 are to: (1)  
245 Expand the focus on sexual trauma beyond counseling and treatment, (2) Mandate  
246 that counseling and appropriate care and services will be provided, (3) Extend the  
247 period of the program to December 31, 2004, and (4) Require a formal mechanism  
248 be implemented to report on outreach activities.

**249 Special Screening of any Cultural or Racial Groups:**

250 Research has centered on three broadly defined groups, Hispanics, Blacks/African-  
251 Americans, and Whites/Caucasians in the attempt to answer two questions: first, are  
252 members of one or more groups more susceptible to developing PTSD? And second,  
253 are the symptoms shown by members of any group more severe or otherwise  
254 different from symptoms shown by other veterans with PTSD?

255 There are data to suggest that Blacks/African-Americans and Hispanics experience  
256 higher rates of PTSD than do Whites/Caucasians (Frueh et al., 1998; Ortega &  
257 Rosenheck, 2000). But, as Frueh and his colleagues note in a systematic review,  
258 "secondary analyses within the existing epidemiological studies suggest that  
259 differential rates of PTSD between racial groups may be a function of differential  
260 rates of traumatic stressors and other pre-existing conditions. This finding, in  
261 combination with the general paucity of empirical data and certain methodological  
262 limitations, significantly moderates the conclusions that should be reached from this  
263 body of literature." Studies in military samples have generally shown no or minimal  
264 race/ethnic differences in PTSD prevalence.

265 In terms of symptom severity and clinical course, the evidence is also mixed. Among  
266 the studies reviewed here, the following conclusions were reached:

- 267 • Two studies found Black/African-American veterans to be more severely affected  
268 than Hispanics or Whites/Caucasians (Frueh et al., 1996; Penk et al., 1989)
- 269 • The National Vietnam Veterans Readjustment Study (NVVRS) found higher PTSD  
270 prevalence among Hispanic veterans than among Whites or Blacks after  
271 controlling for combat exposure. (Kulka et al., 1993)
- 272 • One study found Hispanics to be more severely affected than Whites/Caucasians,  
273 but not to suffer from higher functional impairment levels than Whites (Ortega  
274 and Rosenheck, 2000)
- 275 • Three studies found no significant clinical differences between Black/African-  
276 American veterans and White/Caucasian veterans (Frueh et al., 1997; Rosenheck  
277 and Fontana, 1996; Trent et al., 2000)
- 278 • One review found no clinical differences among Hispanics, Blacks/African-  
279 Americans, and Whites/Caucasians (Frueh et al., 1998)
- 280 • One study found American-of-Japanese-Ancestry Vietnam Veterans had lower  
281 PTSD prevalence than Caucasians (Friedman et al.,(2004)

282 These results support Frueh et al. (1998) in their conclusion that: "despite the  
283 prevailing zeitgeist and clinical lore, the limited extant empirical evidence suggests  
284 that veterans of different races are more similar to each other than they are different  
285 when it comes to the clinical manifestation and response to treatment of combat-  
286 related PTSD and associated features."

#### 287 EVIDENCE

|   | Recommendation  | Sources   | LE   | QE   | SR |
|---|---|---|------|------|----|
| 1 | Screening all patients for PTSD symptoms.   | Breslau et al., 1999a<br>Leskin & Westrup, 1999<br>Prins et al., 1999<br>Taubman et al., 2001   | II-2 | Fair | B  |
| 2 | Frequency of PTSD screening   | Working Group Consensus   | III  | Poor | I  |
| 3 | Primary Care PTSD Screen<br>PTSD Brief Screen<br>Short Screening Scale for DSM IV<br>PTSD Checklist (PCL) | Breslau et al., 1999a<br>Leskin & Westrup, 1999<br>Prins et al., 1999<br>Terhakopian, et al 2008  | II-2 | Fair | B  |
| 4 | Special screening for members of any cultural or racial groups.   | Frueh et al., 1996, 1997, 1998<br>Ortega & Rosenheck, 2000<br>Penk et al., 1989<br>Rosenheck & Fontana, 1996<br>Trent et al., 2000<br>Friedman 2004 | III  | Poor | I  |

288 LE – Level of Evidence; QE = Quality of Evidence; SR = Strength of Recommendation (see Appendix A)

## 289 **D. Are Trauma Related Symptoms Present?**

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### 290 OBJECTIVE

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291 Identify people exposed to trauma who are at risk for developing acute stress  
292 reactions (ASR), acute stress disorder (ASD), or Post-Traumatic Stress Disorder  
293 (PTSD).

### 294 BACKGROUND

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#### 295 *Warning Signs of Trauma Related Stress (APA)*

296 Individuals who have experienced a traumatic event often experience psychological  
297 stress reactions related to the incident. In most instances, these are common  
298 normal reactions to extreme situations. Individuals who feel they are unable to  
299 regain control of their lives or who experience the following symptoms for more than  
300 a month should consider seeking outside professional mental health assistance.  
301 Some symptoms to watch out for include:

- 302 • Recurring thoughts, mental images, or nightmares about the event
- 303 • Having trouble sleeping
- 304 • Changes in appetite
- 305 • Experiencing anxiety and fear, especially when exposed to events or situations
- 306 • Feeling on edge, being easily startled or becoming overly alert
- 307 • Feeling depressed, sad and having low energy
- 308 • Experiencing memory problems including difficulty in remembering aspects of the
- 309 trauma
- 310 • Feeling "scattered" and unable to focus on work or daily activities
- 311 • Having difficulty making decisions
- 312 • Feeling irritable, easily agitated, or angry and resentful
- 313 • Feeling emotionally "numb," withdrawn, disconnected, or different from others
- 314 • Spontaneously crying, feeling a sense of despair and hopelessness
- 315 • Feeling extremely protective of, or fearful for, the safety of loved ones
- 316 • Not being able to face certain aspects of the trauma, and avoiding activities,
- 317 places, or even people that remind you of the event
- 318

### 319 RECOMMENDATIONS

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- 320 1. Individuals who are presumed to have symptoms of PTSD or who are positive for  
321 PTSD on the initial screening should receive a more detailed assessment of their  
322 symptoms.
- 323 2. Useful symptom-related information may include details such as time of onset,  
324 frequency, course, severity, level of distress, and degree of functional  
325 impairment.
- 326 3. The elapsed time since the exposure to trauma should be considered when  
327 assessing the risk of developing PTSD and determining the diagnosis appropriate  
328 intervention. The following definitions will help providers select the appropriate  
329 treatment algorithm:

## 330 **Stress Related Disorders & Syndromes Definitions**

### 331 **Trauma**

332 An extreme traumatic stressor involving direct personal experience of an event that  
333 involves actual or threatened death or serious injury, or other threat to one's  
334 physical integrity; or witnessing an event that involves death, injury, or a threat to  
335 the physical integrity of another person; or learning about unexpected or violent  
336 death, serious harm, or threat of death or injury experienced by a family member or  
337 other close associate. According to DSM-IV-TM criteria, the person's response to the  
338 event must involve intense fear, helplessness, or horror. However, there is evidence  
339 that military personnel do not respond in the same way as civilian victims of  
340 traumatic events, and the A2 criteria for "fear, helpless, or horror" is being  
341 reconsidered in the proposed new DSM criteria (Adler, 2008).

### 342 **Acute Stress Reactions (ASR)**

343 Acute stress reaction is a transient condition that develops in response to a traumatic  
344 event. Onset of at least some signs and symptoms may be simultaneous with the  
345 trauma itself or within minutes of the traumatic event and may follow the trauma  
346 after an interval of hours or days. In most cases symptoms will disappear within  
347 days (even hours). Symptoms include a varying mixture of:

348 Acute stress reactions include broad group of physical, mental and emotional  
349 signs and symptoms that result from heavy mental and emotional work during  
350 exposure to difficult potentially traumatic conditions

351 Symptoms include depression, fatigue, anxiety, decreased  
352 concentration/memory, hyperarousal or any of the four categories of reactions  
353 (See Table CORE - 1) that have not resolved during 4 days after the event, after  
354 a rule-out of other disorders.

355 The traumatic events that can lead to an acute stress reaction are of similar severity  
356 to those involved in post-traumatic stress disorder.

### 357 **Combat and Operational Stress Reactions (COSR) during an Ongoing Military Operation**

358 COSR is the term used to describe an acute stress reaction in the combat  
359 environment and can include virtually any symptom and sign, including physical and  
360 neurological symptoms, resulting from exposure to extremely stressful events or  
361 combat experiences. It may result from exhaustion from cumulative effects of  
362 chronic sleep deprivation and extreme physical stress, or occur in response to  
363 specific combat traumatic experiences.

### 364 **Acute Stress Disorder (ASD)**

365 ASD refers to clinically significant (causing significant distress or impairment in  
366 social, occupational, or other important areas of functioning) symptoms >2 days, but  
367 <1 month after exposure to a trauma as defined above (may progress to PTSD if  
368 symptoms last >1 month). Criteria for diagnosis include:

- 369 • Exposure to trauma as defined above
- 370 • Either while experiencing or after experiencing the distressing event, the  
371 individual has at least three of the following dissociative symptoms:
- 372 • A subjective sense of numbing, detachment, and/or absence of emotional  
373 responsiveness.
- 374 • A reduction in awareness of his/her surroundings (e.g., "being in a daze").

- 375 • Derealization (the feeling that familiar surroundings or people are unreal or have  
376 become strange).
- 377 • Depersonalization (the feeling in an individual that (s)he is no longer him/herself.  
378 His/Her personality, body, external events, the whole world may be no longer  
379 appear real).
- 380 • Dissociative amnesia (i.e., the inability to recall an important aspect of the  
381 trauma).
- 382 • The traumatic event is persistently re-experienced in at least one of the following  
383 ways: recurrent images, thoughts, dreams, illusions, flashback episodes, or a  
384 sense of reliving the experience; or distress on exposure to reminders of the  
385 traumatic event.
- 386 • Marked avoidance of stimuli that arouse recollections of the trauma (e.g.,  
387 thoughts, feelings, conversations, activities, places, people, sounds, smells, or  
388 others)
- 389 • Marked symptoms of anxiety or increased arousal (e.g., difficulty sleeping,  
390 irritability, poor concentration, hypervigilance, exaggerated startle response, and  
391 motor restlessness).

### 392 ***Post-Traumatic Stress Disorder (PTSD)***

393 Clinically significant symptoms that are causing significant distress or impairment in  
394 social, occupational, or other important areas of functioning, and occur more than 1  
395 month after exposure to a trauma. Symptoms may include:

396 The traumatic event is **persistently re-experienced** in one (or more) of the  
397 following ways:

- 398 • Recurrent and intrusive recollections of the event, including images, thoughts, or  
399 perceptions.
- 400 • Recurrent distressing dreams of the event.
- 401 • Acting or feeling as if the traumatic event were recurring (includes a sense of  
402 reliving the experience, illusions, hallucinations, and dissociative flashback  
403 episodes, including those that occur on awakening or when intoxicated).
- 404 • Intense psychological distress at exposure to internal or external cues that  
405 symbolize or resemble an aspect of the traumatic event.
- 406 • Physiologic reactivity on exposure to internal or external cues that symbolize or  
407 resemble an aspect of the traumatic event.

408 Persistent **avoidance** of stimuli associated with the trauma and numbing of general  
409 responsiveness (not present before the trauma), as indicated by three or more of the  
410 following:

- 411 • Efforts to avoid thoughts, feeling, or conversations associated with the trauma.
- 412 • Efforts to avoid activities, places, or people that arouse recollections of the  
413 trauma.
- 414 • Inability to recall an important aspect of the trauma.
- 415 • Markedly diminished interest or participation in significant activities.
- 416 • Feeling of detachment or estrangement from others.
- 417 • Restricted range of affect (e.g., unable to have loving feelings).



- 418 • Sense of foreshortened future (e.g., does not expect to have a career, marriage,  
419 children, or a normal life span).

420 Persistent symptoms of increased **arousal** (not present before the trauma) as  
421 indicated by at least two of the following:

- 422 • Difficulty falling or staying asleep  
423 • Irritability or outbursts of anger  
424 • Difficulty concentrating  
425 • Hypervigilance  
426 • Exaggerated startle response.

### 427 ***Acute PTSD***

428 The above clinically significant symptoms continue to cause significant distress or  
429 impairment in social, occupational, or other important areas of functioning lasting >1  
430 month, but <3 months after exposure to trauma.

### 431 ***Chronic PTSD***

432 The above clinically significant symptoms cause significant distress or impairment in  
433 social, occupational, or other important areas of functioning lasting >3 months after  
434 exposure to trauma. Chronic PTSD is unlikely to improve without effective treatment.

- 435 • Some PTSD patients may exhibit persistent difficulties in interpersonal relations,  
436 mood, somatization and profound identity problems. Such presentation may be  
437 often associated with sustained or repeated trauma during childhood or  
438 adolescence (such as longstanding incest or physical abuse), but it may also be  
439 associated with sustained trauma in later life or may appear as a late  
440 consequence of chronic PTSD, even if the original traumatic stressor was a single  
441 event.
- 442 • Comorbid – also meeting DSM criteria for another disorder such as substance use  
443 disorder, major depression disorder, other anxiety disorder, and mTBI among  
444 military personal.

### 445 ***PTSD with Delayed Onset***

446 Onset of the above clinically significant symptoms causing significant distress or  
447 impairment in social, occupational, or other important areas of functioning at least 6  
448 months after exposure to trauma.

449 Figure A1. Stress Reaction Timeline.

450

451

451

452

**Table CORE - 1 Common Signs Following Exposure to Trauma**

| Physical  | Cognitive/Mental   | Emotional   | Behavioral  |
|---|--|---|---|
| <ul style="list-style-type: none"> <li>• Chills</li> <li>• Difficulty breathing</li> <li>• Dizziness</li> <li>• Elevated blood pressure</li> <li>• Fainting</li> <li>• Fatigue</li> <li>• Grinding teeth</li> <li>• Headaches</li> <li>• Muscle tremors</li> <li>• Nausea</li> <li>• Pain</li> <li>• Profuse sweating</li> <li>• Rapid heart rate</li> <li>• Shock symptoms</li> <li>• Thirst</li> <li>• Twitches</li> <li>• Visual difficulties</li> <li>• Vomiting</li> <li>• Weakness</li> </ul> | <ul style="list-style-type: none"> <li>• Blaming someone</li> <li>• Change in alertness</li> <li>• Confusion</li> <li>• Difficulty identifying familiar objects or people</li> <li>• Hyper-vigilance</li> <li>• Increased or decreased awareness of surroundings</li> <li>• Intrusive images</li> <li>• Loss of orientation to time, place, person</li> <li>• Memory problems</li> <li>• Nightmares</li> <li>• Poor abstract thinking</li> <li>• Poor attention</li> <li>• Poor concentration</li> <li>• Poor decision-making</li> <li>• Poor problem-solving</li> </ul> | <ul style="list-style-type: none"> <li>• Agitation</li> <li>• Anxiety</li> <li>• Apprehension</li> <li>• Denial</li> <li>• Depression</li> <li>• Emotional shock</li> <li>• Fear</li> <li>• Feeling overwhelmed</li> <li>• Grief</li> <li>• Guilt</li> <li>• Inappropriate emotional response</li> <li>• Irritability</li> <li>• Loss of emotional control</li> </ul> | <ul style="list-style-type: none"> <li>• Increased alcohol consumption</li> <li>• Antisocial acts</li> <li>• Change in activity</li> <li>• Change in communication</li> <li>• Change in sexual functioning</li> <li>• Change in speech pattern</li> <li>• Emotional outbursts</li> <li>• Erratic movements</li> <li>• Inability to rest</li> <li>• Change in appetite</li> <li>• Pacing</li> <li>• Startle reflex intensified</li> <li>• Suspiciousness</li> <li>• Social withdrawal</li> </ul> |

453

**E. Educate about additional care if needed, provide contact information****OBJECTIVE**

Provide normalization for survivors and responders whose reactions are not clinically significant

**BACKGROUND**

Trauma survivors and responders who are NOT experiencing signs or symptoms or who are experiencing few symptoms should receive education. It should emphasize that the observed reactions in the symptomatic survivors are common in the aftermath of trauma and do not signify personal inadequacy, health problems, mental illness, or other enduring negative consequences.

Contemporary approaches to early intervention following trauma exposure emphasize the importance of "normalization" of acute stress reactions. Survivors or responders who show distressing symptoms or disturbed behavior should be educated to understand that their reactions are common, normal responses to the extreme events. Such an approach follows from the common clinical observation that individuals experiencing acute stress reactions often interpret their reactions as signs of "personal weakness" or evidence that they are "going crazy," which increases their demoralization and distress. Normalization is undermined if survivors or responders who are not experiencing disruptive distress show a derogatory or punitive attitude to others who are.

Also, the persons with distress who most strongly deny or dissociate from their distress may be at increased risk for developing acute stress disorder (ASD) and subsequent PTSD. The education and normalization may therefore help them recognize how to protect themselves better, and to seek care early if symptoms do

478 interfere with their “self-control”. Even those who go on to develop PTSD may  
479 benefit from an understanding that their symptoms do not represent “personal  
480 weakness” and that although their symptoms may be severe and painful, they are  
481 not losing control of their minds.

#### 482 RECOMMENDATIONS

- 483 1. Pre- and post-trauma education should include helping the asymptomatic trauma  
484 survivor or responder understand that the acute stress reactions of other people  
485 are common and probably transient, and do not indicate personal failure or  
486 weakness, mental illness, or health problems.
- 487 2. Education should include sufficient review of the many ways that post-traumatic  
488 problems can present, including symptoms in the ASD/PTSD spectrum,  
489 behavioral problems with family and friends, occupational problems, and alcohol  
490 or other substance misuse/abuse.
- 491 3. Education should also include positive messages by identifying and encouraging  
492 positive ways of coping and describing simple strategies to resolve or cope with  
493 developing symptoms and problems, setting expectations for mastery and/or  
494 recovery.
- 495 4. Provide contact information should post traumatic symptoms emerge later.
- 496 5. Early psychotherapy intervention, or psychological debriefing is not beneficial for  
497 asymptomatic individuals and may be harmful [D]

#### 498 DISCUSSION

499 Individuals who do not exhibit symptoms may have family members or close friends  
500 who are symptomatic. The clinician should educate them about their role in  
501 supporting their loved ones and emphasize that normalization is a concept that can  
502 incorporate helping asymptomatic survivors to:

- 503 • View other people’s (and their own possible future) stress reactions as normal,  
504 common, and expectable responses to trauma
- 505 • Recognize that sometimes peoples’ inadequate attempts to cope with their  
506 reactions are also within the range of “normal” for the strange situation
- 507 • See that it is natural for them to wonder how they’re doing and to be surprised or  
508 upset by the intensity, duration, or uncontrollability of their reactions.

509 The evidence base for the utility of normalization is weak. Few studies have  
510 attempted to assess the degree of normalization of survivor attitudes and establish a  
511 relationship with PTSD and other outcomes. Also unstudied is whether reassurance  
512 of normality and likely recovery provided by co-survivor peers or helpers actually  
513 serves to promote normalization. Nonetheless, the concept of normalization is  
514 consistent with theories of the development and maintenance of PTSD and with  
515 research showing a relationship between negative reactions to symptoms and PTSD  
516 (Steil & Ehlers, 2000).

517 Recent literature in the area of trauma has highlighted the potential for interventions  
518 to exacerbate trauma reactions. Asymptomatic survivors should not be offered  
519 services that extend beyond delivery of Psychological First Aid and education.  
520 Psychotherapy intervention may actually cause harm in persons not experiencing  
521 symptoms of acute stress (Roberts et al., 2009). The general rule of “do no harm”  
522 should apply not only to professionals but volunteers alike.

523

524 **EVIDENCE**

|   | Recommendation  | Sources                             | LE  | QE   | SR |
|---|---|-------------------------------------|-----|------|----|
| 1 | Providing pre- and post-trauma education can help individuals to understand and cope with exposure experiences. | Working Group Consensus             | III | Poor | I  |
| 2 | Early psychological interventions for asymptomatic trauma survivors are NOT effective and may be harmful        | Roberts 2009b                       | I   | Good | D  |
|   | Psychological debriefings are not effective   | See module I-1: Early Interventions |     |      |    |

525 *LE =Level of Evidence; QE = Quality of Evidence; SR = Strength of Recommendation (see Appendix A)*

526

1

1  
2 **MODULE A: ACUTE STRESS REACTION (ASR)**  
3 **and**  
4 **PREVENTION OF POST TRAUMATIC STRESS DISORDER (PTSD)**

5 **KEY POINTS**

- 6 1. Provide for basic survival needs and comfort (e.g., liquids, food, shelter,  
7 clothing, heat/cooling).  
8 2. Help survivors achieve restful and restorative sleep.  
9 3. Preserve an interpersonal safety zone protecting basic personal space (e.g.,  
10 privacy, quiet, personal effects).  
11 4. Provide nonintrusive ordinary social contact (e.g., a "sounding board,"  
12 judicious uses of humor, small talk about current events, silent  
13 companionship).  
14 5. Address immediate physical health problems or exacerbations of prior  
15 illnesses.  
16 6. Assist in locating and verifying the personal safety of separated loved  
17 ones/friends.  
18 7. Reconnect survivors with loved ones, friends, trusted other persons (e.g.,  
19 work mentors, health care, clergy).  
20 8. Help survivors take practical steps to resume ordinary day-to-day life (e.g.,  
21 daily routines or rituals).  
22 9. Help survivors take practical steps to resolve pressing immediate problems  
23 caused by the disaster (e.g., loss of a functional vehicle, finance, housing).  
24 10. Facilitate resumption of normal family, community, school, and work roles.  
25 11. Provide opportunities for grieving for losses.  
26 12. Help survivors reduce problematic tension or anxiety to manageable levels.  
27 13. Support survivors' helpers through consultation and training about common  
28 stress reactions and stress management techniques.

29 *Military Considerations*

- 30 14. Preserve the fighting force.  
31 15. Return the service member (SM) to functional status.  
32 16. Maintain and enhance unit capabilities and readiness.

33 *Military Clinical Objectives:*

- 34 17. Prevent exacerbation of symptoms / mitigate symptoms of acute stress.  
35 18. Prevent development of traumatic stress sequelae (e.g., ASD, PTSD,  
36 depressive disorders, anxiety disorders, and substance use disorders).  
37 19. Keep SM with his/her unit and prevent unnecessary medical evacuation.  
38

39 **DISCUSSION**

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40 Screening and needs assessments for individuals, groups and populations are  
41 important for the provision of informed early intervention following a major incident

42 or traumatic event. Initial reactions following trauma are varied, complex, and  
43 unstable.

44 Although acute stress reaction (ASR) is not defined in the DSM-IV, there has long  
45 been recognition among mental health professionals that individuals who experience  
46 a traumatic event react in certain predictable ways. A key point in the World Health  
47 Organization definition (WHO, 1992) of ASR is the assertion that "the symptoms  
48 usually appear within minutes of the impact of the stressful stimulus or event, and  
49 disappear within 2-3 days (often within hours)." This view is echoed in a Guideline  
50 for Evidence-Based Early Psychological Intervention for Victims/Survivors of Mass  
51 Violence, released in 2002 by a collaborative group of Federal Departments and the  
52 American Red Cross: "a sensible working principle in the immediate post-incident  
53 phase is to expect normal recovery" (NIMH, 2002).

54 The authors of this guideline have formulated the recommendations discussed below  
55 for the management of persons with acute stress reactions (ASR) following a  
56 traumatic event. Most of the recommendations in this module are based on group  
57 consensus. When available, the evidence and supporting research are presented in  
58 evidence tables.

59

60 The approach to triage in the immediate response to traumatic exposure for service  
61 members with symptoms during Ongoing Military Operations may vary from the  
62 management of civilians exposed to traumatic events. Combat and Operational  
63 Stress Reactions (COSR) management is targeted to preserve the fighting force and  
64 return the service member (SM) to functional status. The annotations for this  
65 Module include, when appropriate, specific recommendations addressing the service  
66 members with COSR.

67

**MODULE A: ALGORITHM**

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70

**MODULE A: ANNOTATIONS**

71

**1. ASSESSMENT & TRIAGE**

72

**A. Trauma Exposure (within the past 30 days)**

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73

**Acute Stress Reactions** (ASR) are transient symptoms or problems that often develop in response to a traumatic event. Traumatic events are events that cause a person to fear that he/she may die or be seriously injured or harmed. These events also can be traumatic when the person witnesses them happening to others. Such events often create feelings of intense fear, helplessness, or horror for those who experience them. The traumatic events that can lead to an acute stress reaction are of similar severity to those involved in post-traumatic stress disorder (PTSD).

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**Combat or Operational Stress Reaction** (COSR) is an acute stress reaction of service members during Ongoing Military Operations. COSR specifically refers to negative adaptation to high stress events and potentially traumatic event exposure. This reaction is not attributed to an identified medical/surgical condition requiring other urgent treatment (a service member can have COSR concurrent with minor wounds/illnesses).

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Among the common types of traumatic events are:

86

- Combat in a war zone

87

- Ongoing military operations (including HA/DR]

88

- Rape, sexual or other physical assault

89

- Natural disaster (e.g., hurricanes, floods or fires)

90

- Child physical and/or sexual abuse

91

- Domestic violence (battering)

92

- Motor vehicle accidents (MVAs)

93

- Exposure to the sudden or unexpected death of others

94

- Sudden life-threatening physical illness (e.g., heart attack or cancer)

95

- Continuous or reoccurring exposure to traumatic event(s)

96

*Events specific to COSR:*

97

- Intense emotional demands (e.g., rescue personnel and caregivers searching for possibly dying survivors, or interacting with bereaved family members)

98

99

- Extreme fatigue, weather exposure, hunger, sleep deprivation

100

- Extended exposure to danger, loss, emotional/physical strain

101

- Exposure to environment hazards, such as toxic contamination (e.g., gas or fumes, chemicals, radioactivity)

102

103

- While COSR can result from a specific traumatic event, it generally emerges from cumulative exposure to multiple stressors.

104

105

106

Onset of at least some signs and symptoms may be simultaneous with the trauma itself or may follow the trauma after an interval of hours or days. Symptoms may include depression, fatigue, anxiety, decreased concentration/memory, irritability, agitation, and exaggerated startle response.

107

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111 **B. Assess briefly based on general appearance and behavior**

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112 **OBJECTIVE**

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113 Identify individuals who may be at risk for endangering themselves or others due to  
114 emotional distress or functional incapacity.

115 **BACKGROUND**

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116 Acute stress reactions (ASR) are transient symptoms or problems that often develop  
117 in response to a traumatic event. The symptoms of acute stress reaction begin  
118 within minutes of the traumatic event and disappear within days (even hours).  
119 Symptoms vary greatly, but can include a mixture of:

- 120 • Anxiety symptoms (e.g., sweating, increased heart rate, and flushing)
- 121 • An initial state of 'daze' - narrowing of attention
- 122 • Reduced levels of consciousness - disorientation
- 123 • Agitation or overactivity
- 124 • Depression
- 125 • Withdrawal

126 There are a number of possible reactions to a traumatic situation, which are  
127 considered within the "norm" for persons experiencing traumatic stress. These  
128 reactions are considered 'normal' in the sense of affecting most survivors, being  
129 socially acceptable, psychologically effective, and self-limited. In the early stage  
130 (the first 4 days after the trauma exposure), it is important not to classify these  
131 reactions as "symptoms" in the sense of being indicative of a mental disorder.

132 **RECOMMENDATIONS**

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- 133 1. Identification of a patient with ASR symptoms is based on observation of  
134 behavior and function; there is insufficient evidence to recommend a specific  
135 screening tool.
- 136 2. Individuals exhibiting the following responses to trauma should be screened for  
137 ASR:
  - 138 a. Physical: exhaustion, hyperarousal, somatic complaints (GI, GU, MS, CV,  
139 Resp, NS), or symptoms of conversion disorder
  - 140 b. Emotional: anxiety, depression, guilt/hopelessness
  - 141 c. Cognitive/mental: amnesic or dissociative symptoms, hypervigilance,  
142 paranoia, intrusive re-experiencing
  - 143 d. Behavioral: avoidance, problematic substance use.
- 144 3. Individuals who experience ASR should receive a comprehensive assessment of  
145 their symptoms to include details about the time of onset, frequency, course,  
146 severity, level of distress, functional impairment, and other relevant information.
- 147 4. Assess for capability to perform routine functions.

148 *Assessment specific to COSR:*

- 149 5. Assess service member's function, to include:
  - 150 a. Any changes in productivity
  - 151 b. Co-worker or supervisor reports of recent changes in appearance, quality  
152 of work, or relationships

- 153 c. Any tardiness/unreliability, loss of motivation, or loss of interest
- 154 d. Forgetful or easily distracted
- 155 e. Screening for substance use
- 156 6. Document symptoms of COSR and obtain collateral information from unit leaders,  
157 coworkers or peers about stressors, function, medical history, and absence or  
158 impairment in operation or mission.
- 159 7. Consider the service member's role and functional capabilities and the complexity  
160 and importance of his/her job.

## 161 DISCUSSION

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162 An acute stress reaction (ASR/COSR) may appear concurrent with other wounds or  
163 illnesses. Providers should confirm that the symptoms are not due to identified  
164 medical/surgical conditions requiring other urgent treatment. ASR does not require  
165 a specific traumatic event, and may result from cumulative exposure to multiple  
166 stressors.

167 In the aftermath of any extreme stressful event, most of those suffering from acute  
168 traumatic stress reactions will be easy to spot. Those who have been injured will be  
169 obvious. Among the uninjured there will also be many who look stunned, appear  
170 pale and faint, or can be seen to be shaking. Some of those who appear to be  
171 suffering from trauma may not even be the actual victims of the disaster, but  
172 witnesses or rescuers who may be deeply affected by what they are seeing. Some  
173 may not be immediately identifiable as traumatized because they may be highly  
174 active - looking for others or looking after others, organizing help and rescue. A  
175 percentage of these may, in the next days or weeks, develop post-traumatic stress  
176 disorder (PTSD).

177 Practitioners who are managing service members suffering from stress reactions or  
178 COSR should consider a variety of factors when deciding when a service member is  
179 ready to return to duty, considering type of job and level of responsibility:

- 180 • Service members in lower skilled jobs, such as truck drivers, food service  
181 personnel, and basic supply functions, can be expected to function effectively  
182 despite continuing anxiety. The cost of functional degradation of individuals  
183 in these roles is likely limited.
- 184 • Service members in higher risk or skilled jobs (e.g., artillery forward  
185 observers, combat controllers, physicians, and pilots) should not be returned  
186 to full duty unless or until the service member has resumed effective  
187 functioning.

188 Individuals in leadership positions should be required to demonstrate a higher level  
189 of reconstitution.

190

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**C. Unstable, Dangerous to Self or Others, or Need for Urgent Medical Attention**

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**OBJECTIVE**

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Protect individuals who may be at risk for endangering themselves or others due to emotional distress or functional incapacity.

**BACKGROUND**

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Emergency treatment, administered to an injured person before professional medical care is available, can be applied to stress reactions of the mind as well as to physical injuries of the body. Acute interventions can be envisioned as the mental health correlate of physical first aid, with the goal being to “stop the psychological bleeding”. The first most important measure should be to eliminate (if possible) the source of the trauma, or to remove the victim from the traumatic, stressful environment. Once the patient is in a safe situation, the provider should attempt to reassure the patient, encourage a professional healing relationship, encourage a feeling of safety, and identify existing social supports. Establishing safety and assurance may enable people to get back on track, and maintain their pre-trauma stable condition.

**RECOMMENDATIONS**

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1. Address acute medical/behavioral issues to preserve life and avoid further harm by:
  - a. Providing appropriate medical/surgical care or referring to stabilize
  - b. Avoiding use of prescribed medications, mood or mind-altering substances
  - c. Preventing possible biological or chemical agent exposure
  - d. Managing substance intoxication or withdrawal
  - e. Stopping self-injury or mutilation
  - f. Addressing inability to care for oneself
2. Arrange a safe, private, and comfortable environment for continuation of the evaluation:
  - a. Assess danger to self or others (e.g., suicidal, homicidal behavior)
  - b. Establish a working treatment alliance with the patient
  - c. Maintain a supportive, non-blaming non-judgmental stance throughout the evaluation
  - d. Assist with the removal of any ongoing exposure to stimuli associated with the traumatic event
  - e. Minimize further traumas that may arise from the initial traumatic event
  - f. Assess and optimize social supports
  - g. Secure any weapons and explosives.
3. Legal mandates should be followed:
  - a. Reporting of violence, assault
  - b. Confidentiality for the patient
  - c. Mandatory testing
  - d. Attending to chain of evidence in criminal cases (e.g., rape, evaluation)

- 232 e. Involuntary Commitment procedures if needed
- 233 4. Carefully consider the following potential interventions to secure safety:
- 234 a. Find safe accommodation and protect against further trauma
- 235 b. Voluntary admission if suicidal
- 236 c. Restraint/seclusion only if less restrictive measures are ineffective
- 237 d. Provide medications managing specific symptoms as needed (e.g.,
- 238 sleep, pain)
- 239 5. Educate and “normalize” observed psychological reactions to the chain of
- 240 command.
- 241 6. Evacuate to next level of care if unmanageable, if existing resources are
- 242 unavailable, or if reaction is outside of the scope of expertise of the care
- 243 provider.

#### 244 DISCUSSION

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245 Foa et al. (2000) rank “suicidality” among factors that will affect treatment decisions  
246 for PTSD. This factor must also be considered in the immediate post-trauma period:  
247 “self-destructive and impulsive behaviors, while not part of the core PTSD symptom  
248 complex, are recognized as associated features of this disorder that may profoundly  
249 affect clinical management. Therefore, the routine assessment of all patients  
250 presenting with acute stress symptoms after exposure to a traumatic stressor should  
251 include a careful evaluation of current suicidal ideation and past history of suicidal  
252 attempts. Risk factors for suicide should also be assessed, such as current  
253 depression and substance abuse. If significant suicidality is present, it must be  
254 addressed before any other treatment is initiated.” Likewise, the patient must be  
255 assessed for any signs of violence toward others, or threat of violence in the home  
256 environment (e.g., ongoing battering) and any risk of violence should be an  
257 indication for immediate treatment.

258 While there is little research on these issues for acute stress reaction per se, the  
259 literature suggests some general trends for persons with PTSD that may inform  
260 clinical treatment of ASR. For example, individuals with subthreshold PTSD are at  
261 high risk for suicidal ideation (Marshall et al., 2001) and, for women, suicide  
262 attempts (Breslau, 2001; Ferrada-Noli et al., 1998; Kaslow et al., 2000; Prigerson et  
263 al., 1999). For young adults, aggressive symptoms may be predictive of suicidality  
264 in men and elevated symptoms of PTSD and/or depression may be more predictive  
265 in women (Prigerson et al., 1999). Some individuals with stress reactions could be  
266 at risk for violence toward others. This can be manifested through explosivity and  
267 anger problems and may predict risk for violent behavior.

268 Optimizing existing social supports is helpful in settings of acute stress and may  
269 decrease risk of suicidality in PTSD (Kotler et al., 2001). For example, higher social  
270 support for women who have experienced domestic violence may reduce risk of  
271 PTSD and other mental disorders (Coker et al., 2002).

#### 272 D. Ensure Basic Physical Needs Are Met

---

##### 273 OBJECTIVE

---

274 Ensure that trauma-exposed persons with acute stress symptoms have their basic  
275 needs met.

276

**BACKGROUND**

277 Trauma victims often have significant disruption to their routines for sleep, nutrition,  
278 exercise, access to finances, and health care. Their normal shelter, clothing, and  
279 other basic resources may be destroyed or inaccessible. These disruptions can be  
280 additionally traumatizing.

281 (See specific protocols for DoD specific services)

282 Early interventions should typically seek to address the needs of the individual  
283 person, with the aim of promoting normal recovery, resiliency, and personal growth  
284 and avoid additional harm. (See Table A - 1)

285 Individual persons who were exposed to trauma as members of a group/unit that  
286 existed prior to the trauma event (e.g., police units, firefighters or military units)  
287 may also benefit from interventions addressing the collective outcomes, such as  
288 social order and community or unit cohesion. Some of the acute interventions, such  
289 as psychoeducation, may be provided in a group format to maintain unit integrity  
290 and promote continuity with established relationship.

291

292 **Table A - 1 Early Intervention after Exposure to Trauma (<4 days)**

| SR | Significant Benefit | Some Benefit  | Unknown | No Benefit Potential Harm |
|----|---------------------|---|---------|---------------------------|
| C  | --                  | Psychological First Aid   | --      | --                        |
| I  | --                  | Provision of basic needs<br>Psychoeducation,<br>normalization<br>Social, and spiritual<br>support | --      | --                        |
| D  | --                  | --  | --      | Psychological debriefing  |

293 *SR = Strength of recommendation (see Appendix A)*

294

**RECOMMENDATIONS**

296

1. Acute intervention should ensure that the following needs are met:

297

a. Safety/security/survival

298

b. Food, hydration, clothing and shelter

299

c. Sleep

300

d. Medications (i.e., replace medications destroyed/lost)

301

e. Education as to current status

302

f. Communication with family, friends and community

303

g. Protection from ongoing threats/toxins/harm. If indicated reduce use  
of alcohol, tobacco, caffeine and illicit psychoactive substances

304

305

2. Provide Psychological First Aid to include:

306

a. Protect survivors from further harm

307

b. Reduce physiological arousal

- 308 c. Mobilize support for those who are most distressed  
 309 d. Keep families together and facilitate reunion with loved ones  
 310 e. Provide information, foster communication and education  
 311 f. Use effective risk communication techniques.

312 *Interventions Specific for Members of Pre-existing Group (e.g., COSR)*

- 313 3. Treat according to member's prior role and not as a "patient"  
 314 4. Assure or provide the following, as needed:  
 315 a. Reunion or ongoing contact with group/unit  
 316 b. Promote continuity with established relationship (e.g., primary group)  
 317 c. Respite from intense stress  
 318 d. Comfortable environment (e.g., thermal comfort)  
 319 e. Consider psychoeducation and discussion in a group format  
 320 f. Assign job tasks and recreational activities that will restore focus and  
 321 confidence and reinforce teamwork (limited duty)

322 **DISCUSSION**

---

323 Psychological first aid should be envisioned as the mental health correlate of physical  
 324 first aid, with the goal being to "stop the bleeding." The patient should be removed  
 325 from the traumatic situation. When the patient is in a safe situation the clinician  
 326 should attempt to reassure the patient and encourage a feeling of safety.

327 In their Disaster Mental Health Response Handbook (Raphael, 2000) a group of PTSD  
 328 experts propose three stages of care:

329 ***Protect:***

330 Find ways to protect survivors from further harm and from further exposure to  
 331 traumatic stimuli. If possible, create a "shelter" or safe haven for them, even if it is  
 332 symbolic. The fewer traumatic stimuli people see, hear, smell, taste, or feel, the  
 333 better off they will be.

334 ***Direct:***

335 Kind and firm direction is needed and appreciated. Survivors may be stunned, in  
 336 shock, or experiencing some degree of dissociation. When possible, direct  
 337 ambulatory survivors:

- 338 1. Away from the site of destruction  
 339 2. Away from severely injured survivors  
 340 3. Away from continuing danger

341 ***Connect:***

342 Survivors who are encountered will usually have lost connection to the world that  
 343 was familiar to them. A supportive, compassionate, and nonjudgmental verbal or  
 344 nonverbal exchange between you and survivors may help to give the experience of  
 345 connection to the shared societal values of altruism and goodness. Help survivors  
 346 connect:

- 347 o To loved ones  
 348 o To accurate information and appropriate resources  
 349 o To locations where they will be able to receive additional support

- 350                   ○ To unit comrades and mission, fostering vertical and horizontal cohesion

351                   **Triage:**

352                   The majority of survivors experience normal stress reactions. However, some may  
353                   require immediate crisis intervention to help manage intense feelings of panic or  
354                   grief. Signs of panic are trembling, agitation, rambling speech, erratic behavior.  
355                   Signs of intense grief may be loud wailing, rage, or catatonia. In such cases,  
356                   attempt to quickly establish therapeutic rapport, ensure the survivor's safety,  
357                   acknowledge and validate the survivor's experience, and offer empathy. Medication  
358                   may be appropriate and necessary, if available.

359                   **Psychological First Aid**

360                   Psychological first aid was first coined in Raphael's book 'When Disaster Strikes: how  
361                   individual and communities cope with catastrophe' (1986). It is included as part of  
362                   the *Fundamental Criteria for First Aid* knowledge and skills that soldiers should be  
363                   trained in order to save themselves or other soldiers in casualty situation. The FM  
364                   21-11 First Aid for Soldiers document (1991) includes the following:

365                    "The Psychological first aid is most needed at the first sign that a soldier cannot  
366                   perform the mission because of emotional distress. Stress is inevitable in  
367                   combat, in hostage and terrorist situations, and in civilian disasters, such as  
368                   floods, hurricanes, tornadoes, and industrial and aircraft catastrophes. Most  
369                   emotional reactions to such situations are temporary, and the person can still  
370                   carry on with encouragement. Painful or disruptive symptoms may last for  
371                   minutes, hours, or a few days. However, if the stress symptoms are seriously  
372                   disabling, they may be psychologically contagious and endanger not only the  
373                   emotionally upset individual but also the entire unit. Even when there is no  
374                   immediate danger of physical injury, psychological harm may occur.

375                   Psychological first aid really means assisting people with emotional distress whether  
376                   it results from physical injury, disease, or excessive traumatic stress. Emotional  
377                   distress is not always as visible as a wound, a broken leg, or a reaction to pain from  
378                   physical damage. However, overexcitement, severe fear, excessive worry, deep  
379                   depression, misdirected aggression, or irritability and anger are signs that stress has  
380                   reached the point of interfering with effective coping.

381                   Psychological first aid should go hand in hand with physical first aid. The discovery  
382                   of a physical injury or cause for an inability to function does not rule out the  
383                   possibility of a psychological injury (or vice versa). A physical injury and the  
384                   circumstances surrounding it may actually cause an emotional injury that is  
385                   potentially more serious than the physical injury; both injuries need treatment. The  
386                   person suffering from pain, shock, fear of serious damage to his body, or fear of  
387                   death does not respond well to joking, indifference, or fearful-tearful attention. Fear  
388                   and anxiety may take as high a toll of the soldier's strength as does the loss of  
389                   blood." (The Department of the Army; Washington, DC, 4 December 1991)

390                   For extended discussion of dangerousness to self or others, [see Module B –](#)  
391                   [Annotation C – Assessment of Dangerousness.](#)

392

393

**Table A - 2 Key Elements of Psychological First Aid (PFA)**

1. *Contact and Engagement* - Respond to contacts initiated by affected persons, or initiate contacts in a non-intrusive, compassionate, and helpful manner
2. *Safety and Comfort* - Enhance immediate and ongoing safety, and provide physical and emotional comfort
3. *Stabilization (if needed)* - Calm and orient emotionally overwhelmed or distraught survivors
4. *Information Gathering - Current Needs and Concerns* - Identify immediate needs and concerns, gather additional information, and tailor PFA interventions
5. *Practical Assistance* - Offer practical help to the survivor in addressing immediate needs and concerns
6. *Connection with Social Supports* - Help establish opportunities for brief or ongoing contacts with primary support persons or other sources of support, including family members, friends, and community helping resources
7. *Information on Coping* - Provide information (about stress reactions and coping) to reduce distress and promote adaptive functioning
8. *Linkage to Collaborative Services* - Link survivors with needed services and inform them about available services that may be needed in the future.

These core goals of PFA constitute the basic objectives of providing early assistance (e.g., within days or weeks following an event). The amount of time spent on each goal will vary from person to person, and with different circumstances according to need.

Complete document describing PFA components can be found at:  
<http://www.vdh.state.va.us/EPR/pdf/PFA9-6-05Final.pdf>

394

395



395

396

**Specific Interventions for COSR:**

397

Combat Operation Stress Control (COSC) utilizes the management principles of brevity, immediacy, contact, expectancy, proximity, and simplicity (BICEPS). These principles apply to all COSC interventions or activities throughout the theater, and are followed by COSC personnel in all BH/COSC elements. These principles may be applied differently based on a particular level of care and other factors pertaining to mission, enemy, terrain and weather, troops and support available, time available, and civil considerations (METT-TC).

404

The actions used for COSC (commonly referred to as the 6 R's) involve the following actions:

405

406

**Reassure** of normality (normalize the reaction)

407

**Rest** (respite from combat or break from work)

408

**Replenish** bodily needs (such as thermal comfort, water, food, hygiene, and sleep).

409

410

**Restore** confidence with purposeful activities and talk

411

**Retain** contact with fellow Soldiers and unit

412

**Remind** recognize emotion of reaction (specifically potentially life-threatening thoughts and behaviors).

413

414

**E Person has Trauma Related Symptoms, Significant Impaired Function, or Diagnosis of ASD**

415

Identify patients who have excessive post-traumatic stress symptoms or significant distress impaired function, or are diagnosed with ASD.

416

417

**BACKGROUND**

418

Since people who develop ASD are at greater risk of developing PTSD, they should be identified and offered treatment as soon as possible. Although ASD does not occur in all people who later develop PTSD, treatment should be considered for all acutely traumatized people with ASD, those with severe PTSD symptoms but do not meet ASD diagnostic criteria, as well as those with functional impairment because of acute physiological symptoms (e.g., hyper-arousal).

419

420

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424

Some patients with an acute stress reaction may benefit from augmentation of the acute intervention and additional follow-up. Because people vary in their reaction and in the rate they recover from traumatic stress, some individuals may require more time or an adjustment of the treatment prior to improvement. Some want and feel a need to discuss the event, and some have no such need. Respect individual and cultural preferences in the attempt to meet their needs as much as possible. Allow for normal recovery, and monitor.

425

426

427

428

429

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431

**RECOMMENDATIONS**

432

1. Acutely traumatized people who meet criteria for diagnosis of ASD, and those with significant levels of post-trauma symptoms after at least two weeks post-trauma, as well as those who are incapacitated by acute psychological or physical symptoms should receive further assessment and early intervention to prevent PTSD.

433

434

435

436

- 437 2. Trauma survivors, who present with symptoms that do not meet the diagnostic  
438 threshold for ASD, or those who recovered from the trauma and currently show  
439 no symptoms, should be monitored and may benefit from follow-up and provision  
440 of ongoing counseling or symptomatic treatment.
- 441 3. Service members with COSR who do not respond to initial supportive  
442 interventions may warrant referral or evacuation.
- 443 (See Module I-1: Early Intervention for prevention of PTSD).

#### 444 DISCUSSION

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445 Stress reactions produce biologic, psychological, and behavioral changes. Biologic  
446 alterations include disruptions in neurochemicals, sleep patterns, hyper-arousal, and  
447 somatic symptoms (e.g., pain, gastrointestinal symptoms). Psychological changes  
448 include: mood disturbances (e.g., emotional lability, irritability, blunting, numbing)  
449 anxiety (e.g., increased worry, ruminations) and cognitive disturbances (e.g.,  
450 memory impairment, confusion, and impaired task completion).

451 Not all individuals who are exposed to trauma or who have a COSR require a mental  
452 health referral. However, those service members who are deteriorating or are not  
453 responding to acute supportive interventions, need to be identified and evacuated to  
454 a more definitive level of care. Also, patients who have a high potential for  
455 dangerousness, or the development of symptoms suggestive of a stress related  
456 disorder (i.e., ASD) also need to be identified and referred to a facility that may  
457 provide appropriate mental health care.

458 Patients who do not respond to first line supportive interventions may warrant  
459 treatment augmentation or a mental health referral. Clear indications for a mental  
460 health referral include; a worsening of stress related symptoms, new onset of  
461 dangerousness or maladaptive coping to stress, exacerbation of comorbid psychiatric  
462 conditions, or deterioration in function. Because patients with new onset stressors,  
463 poor social supports, or inadequate coping skills may be at heightened risk to  
464 develop PTSD, a mental health referral is also indicated.

#### 465 ***Acute Stress Disorder (ASD)***

466 Different types of trauma can lead to ASD, from interpersonal assaultive violence, to  
467 accidents, to combat related trauma. As many as ninety percent of individuals who  
468 experience sexual assault, will have acute stress symptoms but not ASD (Breslau,  
469 1996). Additionally, surveys from the OIF/OEF combat theaters indicate that ~10%-  
470 18% of deployed US combat forces experience trauma related stress symptoms (as  
471 measured with PCL cutoff score of 50+). Beliese (2009) states that to facilitate  
472 provision of physical needs, normalization, and psycho-education it may be prudent,  
473 when possible, to wait 24 to 48 hours before beginning medications.

474 Prior to DSM-IV (American Psychiatric Association, 1994), severe distress occurring  
475 in the month after a traumatic event was not regarded as a diagnosable clinical  
476 problem. Although this prevented the pathologizing of transient reactions, it  
477 hampered the identification of more severely traumatized individuals who might  
478 benefit from early interventions. To address this issue, DSM-IV introduced the  
479 diagnosis of acute stress disorder (ASD) to describe those acute reactions associated  
480 with an increased likelihood of developing chronic PTSD (See Table A - 3). A  
481 diagnosis of ASD is given when an individual experiences significantly distressing  
482 symptoms of reexperiencing, avoidance, and increased arousal within 4 weeks of the  
483 trauma. These symptoms must be present for at least two days before the diagnosis  
484 of ASD can be made. The DSM-IV diagnosis of ASD requires that the victim report at

485 least three of the following five symptoms labeled as indicators of dissociation:  
 486 numbing, reduced awareness of surroundings, derealization, depersonalization, and  
 487 dissociative amnesia. These requirements are based on evidence found in previous  
 488 studies that dissociative symptoms at the time of (or shortly after) the traumatic  
 489 event are predictive of the subsequent development of chronic PTSD (Bremner et al.,  
 490 1992; Marmar et al., 1994; Koopman et al., 1994). Thus the fundamental  
 491 differences between PTSD and ASD involve time elapsed since the trauma and the  
 492 relative emphasis on dissociative symptoms in the ASD diagnosis.

493  
 494

**Table A - 3 Diagnostic criteria for 308.3 Acute Stress Disorder (DSM-IV)**

- |   |
|---|
| <p>A. The person has been exposed to a traumatic event in which both of the following were present:</p> <ul style="list-style-type: none"> <li>• the person experienced, witnessed, or was confronted with an event or events that involved actual or threatened death or serious injury, or a threat to the physical integrity of self or others</li> <li>• the person's response involved intense fear, helplessness, or horror</li> </ul> <p>B. Either while experiencing or after experiencing the distressing event, the individual has three (or more) of the following dissociative symptoms:</p> <ul style="list-style-type: none"> <li>• a subjective sense of numbing, detachment, or absence of emotional responsiveness</li> <li>• a reduction in awareness of his or her surroundings (e.g., "being in a daze")</li> <li>• derealization</li> <li>• depersonalization</li> <li>• dissociative amnesia (i.e., inability to recall an important aspect of the trauma)</li> </ul> <p>C. The traumatic event is persistently reexperienced in at least one of the following ways: recurrent images, thoughts, dreams, illusions, flashback episodes, or a sense of reliving the experience; or distress on exposure to reminders of the traumatic event.</p> <p>D. Marked avoidance of stimuli that arouse recollections of the trauma (e.g., thoughts, feelings, conversations, activities, places, and people).</p> <p>E. Marked symptoms of anxiety or increased arousal (e.g., difficulty sleeping, irritability, poor concentration, hypervigilance, exaggerated startle response, motor restlessness).</p> <p>F. The disturbance causes clinically significant distress or impairment in social, occupational, or other important areas of functioning or impairs the individual's ability to pursue some necessary task, such as obtaining necessary assistance or mobilizing personal resources by telling family members about the traumatic experience.</p> <p>G. The disturbance lasts for a minimum of 2 days and a maximum of 4 weeks and occurs within 4 weeks of the traumatic event.</p> <p>H. The disturbance is not due to the direct physiological effects of a substance (e.g., a drug of abuse, a medication) or a general medical condition, is not better accounted for by Brief Psychotic Disorder, and is not merely an exacerbation of a preexisting Axis I or Axis II disorder.</p> |
|---|

495

495 **2. TREATMENT and FOLLOW-UP**496 **F. Assess Medical and Functional Status**

---

497 **OBJECTIVE**

---

498 Obtain complete history, physical examination, relevant laboratory tests, and  
499 assessment of functioning to determine course of treatment.

500 **BACKGROUND**

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501 One of the key goals of ASR supportive care is to address immediate physical health  
502 problems and to assist the individual in beginning to return to a normal level of  
503 function. In order to do this, the clinician or caregiver must assess the individual's  
504 current state of health and functioning.

505 **RECOMMENDATIONS**

---

- 506 1. Medical status should be obtained for all persons presenting with symptoms to  
507 include:
- 508 g. History, physical examination and a neurological examination
  - 509 h. Use of prescribed medications, mood or mind-altering substances, and  
510 possible biological or chemical agent exposure
  - 511 i. A minimal mental status examination (MMSE) to assess cognitive  
512 function
- 513 2. The history and physical examination findings should lead the provider to other  
514 assessments as clinically indicated. Assessment may include:
- 515 j. Screen for toxicology if the symptom presentation indicates
  - 516 k. Radiological assessment of patients with focal neurological findings or  
517 possible head injury
  - 518 l. Appropriate laboratory studies to rule out medical disorders that may  
519 cause symptoms of acute stress reactions (e.g., complete blood count  
520 [CBC], chemistry profile, thyroid studies, HCG, EKG, EEG.)
- 521 3. A focused psychosocial assessment should be performed to include assessment of  
522 active stressors, losses, current social supports, and basic needs (e.g., housing,  
523 food, and financial resources).
- 524 4. A brief assessment of function should be completed to evaluate: 1) objectively  
525 impaired function based on general appearance and behavior; 2) subjectively  
526 impaired function; 3) baseline level of function (LOF) vs. current LOF; and 4)  
527 family and relationship functioning.

528 **DISCUSSION**

---

529 Whenever possible, providers should include assessment of any physical injuries,  
530 review of targeted H&P and laboratory results (if available), assessment of the  
531 individual's level of functioning and level of family and relationship functioning.  
532 Ideally, the current clinical picture should be compared to the individual's pre-trauma  
533 state, but often this may not be possible in the immediate aftermath of a traumatic  
534 event. Evaluation of the patient's level of function is warranted because evidence

535 has shown that functional impairment after trauma is a predictor for later  
536 development of PTSD (Norris et al., 2002).

## 537 **G. Assess Pre-Existing Psychiatric and Medical Conditions**

---

### 538 **BACKGROUND**

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539 Circumstances brought about by a traumatic event may complicate any existing  
540 psychiatric conditions or may exacerbate pre-existing pathology.

### 541 **RECOMMENDATIONS**

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- 542 1. Assess patients for pre-existing psychiatric conditions to identify high risk  
543 individuals and groups.
- 544 2. Assure access and adherence to medications the patient is currently taking.
- 545 3. Refer patients with pre-existing psychiatric conditions to mental health specialty  
546 when indicated, or emergency hospitalization if needed.

### 547 **DISCUSSION**

---

548 The NIMH (2002) guideline addresses the need to manage pre-existing psychiatric  
549 and medical conditions. The authors point to the "special needs of those who have  
550 experienced enduring mental health problems, those who are disabled and other  
551 high-risk groups who may be vulnerable and less able to cope with unfolding  
552 situations." They also call for additional attention to be paid to members of these  
553 groups in the immediate post-trauma period. However, they also emphasize "the  
554 presumption of clinically significant disorders in the early post-incident phase is  
555 inappropriate, except for individuals with preexisting conditions."

## 556 **H Assess Risk Factors for Developing ASD/PTSD**

---

### 557 **BACKGROUND**

---

558 Not all trauma survivors develop permanent stress disorders. Early identification of  
559 those at-risk for negative outcomes following trauma can facilitate prevention,  
560 referral, and treatment. Screening for those at greatest risk should address past and  
561 current psychiatric and substance use problems and treatment, prior trauma  
562 exposure, pre-injury psychosocial stressors, and existing social support.

### 563 **RECOMMENDATIONS**

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- 564 1. Trauma survivors who exhibit symptoms or functional impairment should be  
565 screened for the following risk factors for developing ASD/PTSD:

#### 566 **Pre-traumatic factors**

- 567 1. Ongoing life stress
- 568 2. Lack of social support
- 569 3. Age at time of trauma (school age youth, 40-60 years of age)
- 570 4. Pre-existing psychiatric disorders, or substance misuse
- 571 5. History of traumatic events (i.e. MVA)
- 572 6. History of post-traumatic stress disorder (PTSD)
- 573 7. Other pre-traumatic factors including: female gender, low socioeconomic  
574 status, lower level of education, lower level of intelligence, race (Hispanic,  
575 Japanese, other ethnic minority), reported abuse in childhood, report of other

576 previous traumatization, report of other adverse childhood factors, family  
577 history of psychiatric disorders, poor training or preparation for the traumatic  
578 event.

#### 579 Peri-traumatic or trauma related factors

- 580 1. Severe trauma
- 581 2. Physical injury to self or other
- 582 3. Type of trauma (interpersonal traumas such as torture, rape or assault  
583 convey high risk of PTSD)
- 584 4. High perceived threat to life of self or others
- 585 5. Age at trauma (school age youth, 40-60 years of age)
- 586 6. Community (mass) trauma
- 587 7. Other peri-traumatic factors including: history of peri-traumatic dissociation  
588 and interpersonal trauma.

#### 589 Post-traumatic factors

- 590 1. Ongoing life stress
- 591 2. Lack of positive social support
- 592 3. Bereavement or traumatic grief
- 593 4. Major loss of resources
- 594 5. Negative social support (shaming or blaming environment)
- 595 6. Other post-traumatic factors including: children at home and a distressed  
596 spouse.

#### 597 DISCUSSION

---

##### 598 **Risk Factors for ASD**

599 When evaluating risk factors for ASD, the clinician should keep in mind that ASD is  
600 no longer diagnosed later than four weeks after a traumatic event. Thus, not enough  
601 time will have passed following the trauma for many post-trauma factors to have had  
602 full impact on the course of symptoms.

##### 603 **Risk Factors for PTSD**

604 When evaluating risk factors for developing PTSD, the clinician should keep in mind  
605 that PTSD is defined as occurring only after four weeks have elapsed following a  
606 traumatic event. PTSD symptoms, however, may not appear until a considerable  
607 time has passed, sometimes surfacing years later.

608 For further discussion of risk factors for PTSD - See [Module B: Annotation F](#)

#### 609 **Provide Education and Normalization / Expectancy of Recovery**

---

##### 610 BACKGROUND:

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611 Education for trauma survivors and their families may help normalize common  
612 reactions to trauma, improve coping, enhance self-care, facilitate recognition of  
613 significant problems, and increase knowledge of, and access to, services. Individuals  
614 should be reassured about common reactions to traumatic experiences and advised  
615 regarding positive and problematic forms of coping with them.

616 Information about social support and stress management is particularly important.  
617 Opportunities to discuss emotional concerns in individual, family, or group meetings  
618 can enable survivors to reflect on what has happened. Education regarding  
619 indicators that initial acute reactions are failing to resolve will be important. Signs  
620 and symptoms of PTSD; anxiety, depression, substance use disorders, and other  
621 difficulties should be explained. Survivors will need information about financial,  
622 mental health, rehabilitation, legal, and other services available to them, as well as  
623 education about common obstacles to pursuing needed services.

624 **RECOMMENDATION:**

---

- 625 1. All survivors should be given educational information to help normalize common  
626 reactions to trauma, improve coping, enhance self-care, facilitate recognition of  
627 significant problems, and increase knowledge of and access to services. Such  
628 information can be delivered in many ways, including public media, community  
629 education activities, and written materials.

630 **DISCUSSION:**

---

631 Immediate post-trauma distress will remit naturally for many patients (Blanchard et  
632 al., 1995) and provision of mental health services may be unnecessary.  
633 Hypothetically, it is even possible that too much focus on mental health issues may  
634 be iatrogenic for some survivors, centering their attention on symptoms and  
635 problems and making attention and caring contingent on needing such help.

636

636

637 **2. TREATMENT**

638 **J. Initiate Brief Intervention**

639 **OBJECTIVE**

640 To lessen the physical, psychological, and behavioral morbidity associated with acute  
 641 stress reaction (ASR), hasten the return to full function (duty), and reduce risk for  
 642 development of ASD or PTSD following traumatic event.

643 **BACKGROUND**

644 It is likely that not all patients will require intervention immediately following a  
 645 traumatic occurrence. Depending on the intensity and duration of the trauma, there  
 646 will be people who will make it through unharmed. Often, if a person appears to be  
 647 coping well and denies symptoms of ASD or PTSD, they may not need specialized  
 648 care.

649 For people who show symptoms of ASD or PTSD (including symptoms of intrusive  
 650 recollections, avoidance, numbing, and physiological hyperarousal when confronted  
 651 with reminders of the trauma) brief acute intervention may be indicated.

652 Early intervention may need to assist the individual with anticipating problems in  
 653 using their support system. This may be particularly important in light of the fact  
 654 that the psychological aftermath of trauma may significantly disrupt a person’s  
 655 capacity to use others to cope with and manage post-traumatic symptoms and daily  
 656 demands. Table A-4 summarizes the interventions and their potential benefit in the  
 657 first month after exposure to the trauma.

658 **Table A - 4 Early Intervention after Exposure to Trauma (4 to 30 days after exposure)**

| SR | Significant Benefit                               | Some Benefit   | Unknown Benefit  | No Benefit Potential Harm   |
|----|---|--|--|---|
| A  | Brief Cognitive Behavioral Therapy (4-5 sessions) |  |  |   |
| B  |   | Prazosin for nightmares                                  | Imipramine   |   |
| C  |   | Social, and spiritual support                            | Propranolol  |   |
| I  |   | Provide basic Needs<br>Psychoeducation,<br>normalization | Benzodiazpines<br>Sympatholytics<br>Antidepressants<br>Anticonvulsants<br>Atypical Antipsychotics<br>Chloral hydrate |   |
| D  |   |  | Psychological First Aid  | Psychological debriefing<br>Typical Antipsychotics<br>Formal psychological interventions for asymptomatic survivors |

659 SR = Strength of recommendation

660



661 **RECOMMENDATIONS**

- 662 1. The following treatment recommendations should apply for all acutely  
663 traumatized people who meet criteria for diagnosis of ASD, and for those with  
664 significant levels of post-trauma symptoms after at least two weeks post-trauma,  
665 as well as those who are incapacitated by acute psychological or physical  
666 symptoms.
- 667 2. Treatment should be provided after education, normalization, and Psychological  
668 First Aid has been provided, and basic needs following the trauma were made  
669 available.
- 670 3. There is insufficient evidence to recommend for or against the use of  
671 Psychological First Aid to address symptoms beyond 4 days following trauma. [I]
- 672 4. Recommend monitoring for development of PTSD using validated symptom  
673 measures (e.g., PTSD Checklist, other screening tools for ASD/PTSD).
- 674 5. Survivors who present symptoms that do not meet the diagnostic threshold of  
675 ASD or PTSD should be monitored and may benefit from follow-up and provision  
676 of ongoing counseling or symptomatic treatment.

677 **6. Psychotherapy:**

- 678 m. Consider early brief intervention (4 to 5 sessions) of cognitive based  
679 therapy (CBT) that includes exposure based therapy, alone or  
680 combined with a component of cognitive re-structuring therapy for  
681 patients with significant early symptom levels, especially those  
682 meeting diagnostic criteria for ASD.[A]
- 683 n. Early psychotherapy intervention for *asymptomatic* individuals is not  
684 beneficial and may be harmful. [D]
- 685 o. Recommend against individual Psychological Debriefing as a viable  
686 means of reducing acute stress disorder (ASD) or progression to post-  
687 traumatic stress disorder (PTSD). [D]
- 688 p. Recommend against group Psychological Debriefing as a viable means  
689 of reducing acute stress disorder (ASD) or progression to post-  
690 traumatic stress disorder. (Note: this is not a recommendation against  
691 Operational Debriefing) [D]
- 692 q. Groups may be effective vehicles for providing trauma related  
693 education, training in coping skills, and increasing social support  
694 especially in the context of multiple group sessions.
- 695 r. Group participation should be voluntary.

696 **7. Pharmacotherapy:**

- 697 s. There is insufficient evidence to support a recommendation for use of a  
698 pharmacological agent to prevent the development of ASD or PTSD.  
699 [I]

700 **DISCUSSION**

701 For discussion of the supporting evidence and grading of the recommendations see  
702 Module I-1: [Early Interventions to Prevention of PTSD](#)

703 **ASD Treatment**

704 The relationship between ASD and PTSD was examined in three prospective studies.  
705 Classen and colleagues (1998) studied the acute stress reactions of bystanders to a  
706 mass shooting in an office building. They assessed 36 employees (bystanders) 8  
707 days after the shooting. Between 7 and 10 months later, they reassessed 32  
708 employees for post-traumatic stress symptoms and found that 33 percent of them  
709 met criteria for ASD and that meeting criteria for ASD was a strong predictor of PTSD  
710 (accounting for 19 percent of the variance) as well as intrusion (accounting for 53  
711 percent of the variance) and avoidance (accounting for 45 percent of the variance).

712 In another prospective study, Harvey and Bryant (1998a) examined the relationship  
713 between ASD and PTSD in 92 motor vehicle accident survivors. From the twelve  
714 participants (13 percent) who met criteria for ASD within 2 to 26 days of the  
715 accident, 78 percent met criteria for PTSD 6 months later. Nineteen participants (21  
716 percent) met some but not all of the criteria for ASD; of the 15 individuals available  
717 for follow-up, 9 (60 percent) met criteria for PTSD. From the 61 participants who did  
718 not meet the criteria for ASD; only 2 met criteria for PTSD. This study provides  
719 strong evidence of ASD being a predictor of PTSD. Nevertheless, Harvey and Bryant  
720 concluded that the current criteria for ASD might be too stringent for ASD to be used  
721 to predict the risk for PTSD. Bryant and Harvey (1998b) also examined the  
722 relationship between ASD and PTSD for a subset (n=79) of the motor vehicle  
723 accident survivors who suffered mild traumatic brain injury as a result of the  
724 accident. They were particularly interested in the utility of ASD as a predictor of  
725 PTSD in individuals with postconcussive symptoms that could overlap with ASD  
726 symptoms. Their results were similar to previously reported findings: 14 percent  
727 met criteria for ASD; 6 months after the event, 82 percent of those with ASD also  
728 met criteria for PTSD.

729 In another prospective study, Brewin and colleagues (1999) evaluated the use of  
730 ASD to predict PTSD in 157 survivors of violent assault. Participants were assessed  
731 for several ASD symptoms using items from the Post-traumatic Stress Disorder  
732 Symptoms Scale (Foa et al. 1993); additional items were generated to determine  
733 whether the event met the ASD criterion. Nineteen percent of participants met  
734 criteria for ASD and 20 percent met criteria for PTSD at 6-month follow-up. They  
735 found that meeting full criteria for ASD was a better predictor of PTSD than any of  
736 the symptom clusters. Eighty three percent of participants who met criteria for ASD  
737 were diagnosed with PTSD six months later.

738 Research suggests that relatively brief but specialized interventions may effectively  
739 prevent PTSD in some subgroups of trauma patients. Several controlled trials have  
740 suggested that brief (i.e., 4 to 5 sessions) cognitive-behavioral treatments,  
741 comprised of education, breathing training/relaxation, imaginal and *in vivo* exposure,  
742 and cognitive restructuring, delivered within weeks of the traumatic event, can often  
743 prevent PTSD in survivors of sexual and non-sexual assault (Foa et al., 1995) and  
744 MVAs and industrial accidents (Bryant et al., 1998a, 1999). Brief intervention with  
745 patients hospitalized for injury has been found to reduce alcohol consumption in  
746 those with existing alcohol problems (Gentilello et al., 1999). Controlled trials of  
747 brief early intervention services targeted at other important trauma sequelae (e.g.,  
748 problems returning to work, depression, family problems, trauma recidivism, and  
749 bereavement-related problems) remain to be conducted, but it is likely that targeted  
750 interventions may be effective in these arenas for at least some survivors.

751 Two well-designed studies offer evidence that brief treatment interventions utilizing a  
752 combination of cognitive behavioral techniques may be effective in preventing PTSD

753 in a significant percentage of subjects. In a study of a brief treatment program for  
754 recent sexual and nonsexual assault victims who all met criteria for PTSD, Foa et al.  
755 (1995) compared repeated assessments vs. a Brief Prevention Program (BPP) (four  
756 sessions of trauma education, relaxation training, imaginal exposure, *in vivo*  
757 exposure, and cognitive restructuring). Two months posttrauma, only 10 percent of  
758 the BPP group met criteria for PTSD, whereas 70 percent of the repeated  
759 assessments group met criteria for PTSD. In a study of motor vehicle and industrial  
760 accident victims who met criteria for ASD, Bryant et al. (1998a) compared five  
761 sessions of nondirective supportive counseling (support, education, and problem-  
762 solving skills) vs. a brief cognitive-behavioral treatment (trauma education,  
763 progressive muscle relaxation, imaginal exposure, cognitive restructuring, and  
764 graded *in vivo* exposure to avoided situations). Immediately post-treatment, 8  
765 percent in the CBT group met criteria for PTSD, versus 83 percent in the supportive  
766 counseling group. Six Months Post-Trauma, 17 percent in CBT met criteria for PTSD,  
767 versus 67 percent in supportive counseling. One important caveat to these  
768 interventions is that dropout rate was high, and the authors concluded that *those*  
769 *with more severe symptoms may need supportive counseling prior to more intensive*  
770 *cognitive behavioral interventions.*

771 In addition to targeted brief interventions, some trauma survivors may benefit from  
772 follow-up provision of ongoing counseling or treatment. Candidates for such  
773 treatment would include survivors with a history of previous traumatization (e.g.,  
774 survivors of the current trauma who have a history of childhood physical or sexual  
775 abuse) or preexisting mental health problems.

776

#### EVIDENCE

|   | Recommendation   | Sources  | LE | QE   | SR |
|---|--|--|----|------|----|
| 1 | Monitor patient with ASD for development of PTSD. (ASD predictor of PTSD). | Brewin et al. 1999<br>Bryant et al, 1998a & 1998 b                           | I  | Good | A  |
| 2 | Brief intervention of CBT (4 to 5 sessions).                               | Bryant et al., 1998a<br>Foe et al, 1995<br>[See Module I-1: Brief early CBT] | I  | Good | A  |

777 *LE = Level of Evidence; QE = Quality of Evidence; SR = Strength of Recommendation (see Appendix A)*

#### 778 K. Acute Symptom Management

779

##### BACKGROUND

780 Survivors of trauma may not complain directly of ASD symptoms such as re-  
781 experiencing or avoidance. Instead, they may complain of sleeping problem, pain or  
782 other somatic concerns. After addressing immediate needs and providing education  
783 and intervention, alleviating these symptoms will make it easier for survivors to cope  
784 and recover from their traumatic experience.

785

##### RECOMMENDATIONS

- 786 1. Specific symptoms treatment should be provided after education, normalization  
787 and basic needs are met.
- 788 2. Consider a short course of medication (less than 6 days) targeted for specific  
789 symptoms in patients post trauma
- 790 a. Sleep disturbance/insomnia (e.g., hypnotics such as ambien,  
791 trazodone, benzodiazepines, and seroquel)

- 792                   b. Management of pain (e.g., analgesics)
- 793                   c. Irritation/excessive arousal/anger (e.g., benzodiazepines)
- 794           3. Provide non-pharmacologic intervention to address specific symptoms (e.g.,  
795           relaxation, breathing techniques, avoiding caffeine) to address both general  
796           recovery and specific symptoms (sleep disturbance, pain, hyper-arousal or  
797           agitation).

798           For discussion of the supporting evidence of the recommendations  
799           see Module I-3: [Management of Specific Symptoms](#)

## 800 **L. Facilitate Social and Spiritual Support**

### 801 **BACKGROUND**

802           Social support will be critical for helping the individual cope after a trauma has  
803           occurred. It may be necessary to identify potential sources of support and facilitate  
804           support from others (e.g., partners, family, friends, work colleagues, and work  
805           supervisors).

806           The terms “religious” and “spiritual” are both used in the clinical literature to refer to  
807           beliefs and practices to which individuals may turn for support following a traumatic  
808           event.

809           Research on the relationship between religion and health has been hampered by the  
810           lack of consistent and adequate multidimensional measures of religiousness and  
811           spirituality (Idler et al., 2003). Examinations of research findings must take care to  
812           look at the particular dimensions of religiousness / spirituality being measured.  
813           Because the terms are so closely related, and because researchers in this area have  
814           not consistently differentiated between the two concepts, the reader should assume  
815           that in the discussion below religion/spirituality are referred to in the general sense  
816           and not in any specific terms.

### 817 **RECOMMENDATIONS**

- 818           1. Preserve an interpersonal safety zone protecting basic personal space (e.g.,  
819           privacy, quiet, personal effects).
- 820           2. Reconnect with previously supportive relationships (family, friends, unit  
821           members)
- 822           3. Provide nonintrusive ordinary social contact (e.g., a "sounding board," judicious  
823           use of humor, small talk about current events, silent companionship). Provide  
824           opportunities for grieving for losses. (Providing space and opportunities for  
825           prayers, mantras, rites and rituals and end-of-life care as determined important  
826           by the patient).
- 827           4. Consider assessing for spiritual needs, and providing direct spiritual care or  
828           ensuring patient access to spiritual care when sought. [C]

### 829 **DISCUSSION**

830           Religion may provide a framework by which many survivors of trauma construct a  
831           meaningful account of their experience, and may be a useful focus for intervention  
832           with trauma survivors.

**833 Religion seeking is an observed post-traumatic phenomenon:**

834 There is a large body of anecdotal literature documenting the propensity of  
835 individuals to seek religious/spiritual comfort following a traumatic event. The  
836 terrorist attacks of September 11, 2001 provide a recent instance of this  
837 phenomenon. Meisenhelder (2002) notes "the events of September 11, 2001  
838 triggered a widespread national response that was two-fold: a post-traumatic stress  
839 reaction and an increase in attendance in religious services and practices  
840 immediately following the tragic events." Schuster and his colleagues performed a  
841 nationwide phone survey of 569 adults within a week of the event (2001), and found  
842 that "forty-four percent of the adults reported one or more substantial stress  
843 symptoms; 91 percent had one or more symptoms to at least some degree.  
844 Respondents throughout the country reported stress syndromes. They coped by  
845 talking with others (98 percent), turning to religion (90 percent)."

**846 Demonstrated benefits of the practice: religious/spiritual care:**

847 Some longitudinal studies have found support for religious /spiritual practices  
848 extending longevity (Hummer et al., 1999; Strawbridge et al., 1997), and reducing  
849 the duration of depression (Koenig et al., 1998). However, others examining the  
850 overall literature have found it to be "weak and inconsistent" (Sloan, et al., 1999).  
851 Strawbridge and his colleagues (1998) used a large public health survey to  
852 investigate "associations between two forms of religiosity and depression as well as  
853 the extent to which religiosity buffers relationships between stressors and  
854 depression". The authors conclude that "religiosity may help those experiencing  
855 non-family stressors, but may worsen matters for those facing family crises."

856 Several possible pathways for positive physical / mental health benefits from  
857 religious/spiritual practice have been identified. These include; (1) reduction of  
858 behavioral risks through healthy religious lifestyles (e.g. less drinking or smoking)  
859 (2) expanded social support through involvement in spiritual communities, (3)  
860 enhancement of coping skills and helpful cognitive appraisals resulting in meaning-  
861 making, and (4) physiological mechanisms such as activation of the "relaxation  
862 response" through prayer or meditation.

863 Recent research on cognitive processes following traumatic events indicates that  
864 challenges to an individual's basic life assumptions may occur. These assumptions  
865 involve the security and meaningfulness of the world and the individual's sense of  
866 self-worth in relation to perception of the environment (Janoff-Bulman, 1979).  
867 Specifically, these assumptions are; (1) that one's environment is physically and  
868 psychologically safe, (2) that events are predictable, meaningful and fair, (3) that  
869 one's own sense of self-worth is positive in relation to experiences with other people  
870 and events (Hunter, 1996). The style and intensity of religious / spiritual appraisals  
871 and coping behaviors made in the aftermath of trauma may both reflect and  
872 influence the recovery trajectory and ultimate outcomes of traumatic events.

873 A recent meta-analytic study (Ano & Vasconcelles, 2005) of religious / spiritual  
874 coping with stressful events examined results from 49 studies meeting inclusion  
875 criteria from the 109 studies initially identified. In this study, Pargament's (1997)  
876 model of religious coping was used to distinguish between positive and negative uses  
877 of religion / spirituality to deal with life crises. Bivariate correlations between  
878 religious coping (positive and negative) measures and psychological adjustment  
879 (positive and negative) were averaged to obtain effect size estimates for findings for  
880 four key relationships; 1) positive religious coping and positive adjustment, 2)  
881 positive coping and negative adjustment, 3) negative coping and positive  
882 adjustment, and 4) negative coping and negative adjustment. The strongest

883 average effect size estimate (.33) was obtained for the 29 studies examining the  
884 relationship between positive coping and positive adjustment. The 22 studies of  
885 negative coping and negative adjustment produced a significant moderate effect size  
886 estimate of .22. Finally, from 38 studies examining correlations between positive  
887 coping and negative adjustment, a small, yet significant, effect size estimate of -.12  
888 was obtained. Taken together, findings from the Ano & Vasconcelles (2005) meta-  
889 analysis show that both forms of religious coping are related to psychological  
890 adjustment to stress. Positive religious coping is related to both positive and  
891 negative forms of adjustment in the directions expected; whereas negative coping is  
892 only significantly related to negative adjustment.

893 A study of help-seeking military veterans found significant associations between  
894 negative religious coping, lack of forgiveness and worse mental health outcomes  
895 (PTSD and Depression) (Witvliet et al., 2004). Similarly, loss of religious faith was  
896 found to be associated with worse mental health outcomes (i.e. greater utilization of  
897 mental health services) among military veterans in treatment for PTSD (Fontana &  
898 Rosenheck, 2004). Most recently, in a study of religiously active trauma survivors  
899 positive relationships were found between a measure of positive religious coping,  
900 seeking spiritual support, and posttraumatic growth. In the same study a negative  
901 religious coping indicator, religious strain, was significantly related to posttraumatic  
902 symptoms (Harris, Erbes, Engdahl, Olsen, Winskowski & McMahill, 2008). Thus,  
903 there is some consistency in findings of a complex relationship between religious  
904 coping and psychological adjustment following stress.

905 An integrative review of the literature on spirituality and bereavement by Wortman &  
906 Park (2008) suggests that religion/spirituality has a mixed but generally positive  
907 relationship with adjustment to bereavement depending on the dimension measured.  
908 Making-meaning appeared to be an important pathway associating  
909 religion/spirituality with adjustment for several dimensions, including general  
910 religiousness, intrinsic religiousness, religious coping, and belief in the helpfulness of  
911 religion.

### 912 ***The role of the Chaplain:***

913 Recent studies of healthcare systems indicate strong association between the degree  
914 to which patient emotional / spiritual needs are met and overall patient treatment  
915 satisfaction. Healthcare accrediting bodies are recommending that  
916 Chaplains/pastoral care teams work in close collaboration with medical care teams to  
917 provide patients with an in-depth spiritual care experience that results in emotional  
918 comfort and improved satisfaction (Clark et al., 2003).

919 The Chaplain may play an important role in helping individuals regain a sense that  
920 their basic life assumptions are true. Chaplains receive training in a wide variety of  
921 supportive techniques, and they stand ready to assist all individuals, including those  
922 who do not subscribe to an organized religion. Chaplains may provide assistance in  
923 one or more of the following ways:

- 924 ○ Organizing and mobilizing community action by;
- 925 ○ Providing education and consultation with advice for leaders
- 926 ○ Assisting in the mobilization of action plans and recovery processes
- 927 ○ Offering facilitation of adaptation and mastery in social change
- 928 ○ Assisting in the development of community networks
- 929 ○ Supporting the development of a positive recovery organization
- 930 ○ Serving as a source of communication

- 931                   ○ Fostering Community Theater and art geared to encouraging working  
932                   through and recovery from the trauma.
- 933           Providing space and opportunities for prayers, mantras, rites and rituals and end-of-  
934           life care as determined important by the patient is another significant contribution of  
935           the Chaplain. (Canda and Phaobtong, 1992; Lee, 1997).

936 **EVIDENCE**

|   | Recommendation   | Sources   | LE  | QE   | SR |
|---|--|---|-----|------|----|
| 1 | Consider referral for religious/spiritual counseling as indicated for patient symptoms, consistent with available resources, and resonant with patient beliefs | Baldacchino and Draper, 2001<br>Bell Meisenhelder, 2002<br>Calhoun et al., 2000<br>Humphreys et al., 2001<br>Nixon et al., 1999<br>Strawbridge et al., 1998 | III | Poor | I  |
| 2 | Consider providing direct spiritual care or ensuring patient access to spiritual care  | Bogia and Preston, 1985<br>Everly, 2000b  | II  | Fair | C  |

937 *LE = Level of Evidence; QE = Quality of Evidence; SR = Strength of Recommendation (see Appendix A)*  
938

939 **3. RE-ASSESSMENT**940 **M. Reassess Symptoms and Function**941 **OBJECTIVE**

942           Identify patients with persistent traumatic stress symptoms, related dysfunction, or  
943           additional treatment needs.

944 **BACKGROUND**

945           Clinical reassessment of response to the acute intervention is indicated to determine  
946           if there are persistent symptoms and, if necessary, to develop a follow-up plan.

947           Especially important are acute levels of traumatic stress symptoms, which predict  
948           chronic problems; for example, more than three-quarters of MVA patients diagnosed  
949           with ASD will have chronic PTSD at 6 months post-trauma (Bryant and Harvey,  
950           2000).

951           In follow-up appointments, it will be important to screen for PTSD and other anxiety  
952           disorders, depression, alcohol and substance abuse, problems with return to work  
953           and other productive roles, adherence with medication regimens and other  
954           appointments, and potential for re-traumatization.

955 **RECOMMENDATIONS**

- 956           1. Assessment of the response to the acute intervention should include an  
957           evaluation for the following risk factors:
- 958                   a. Persistent or worsening traumatic stress symptoms (e.g., dissociation,  
959                   panic, autonomic arousal, cognitive impairment)
  - 960                   b. Significant functional impairments (e.g., role/work, relationships)
  - 961                   c. Dangerousness (suicidal or violent ideation, plan and/or intent)
  - 962                   d. Severe psychiatric comorbidity (e.g., psychotic spectrum disorder,  
963                   substance use disorder or abuse)

- 964 e. Maladaptive coping strategies (e.g., pattern of impulsivity, social  
965 withdrawal, or other reactions under stress)
- 966 f. New or evolving psychosocial stressors
- 967 g. Poor social supports.
- 968 2. Follow-up after acute intervention to determine patient status should include:
- 969 a. Patient does not improve or status worsens – continue management  
970 of PTSD (See Module B) in consultation or referral to PTSD specialty  
971 care or mental health provider. Recommend involvement of the  
972 primary care provider in the treatment. Patients with multiple  
973 problems may benefit from a multi-disciplinary approach to include  
974 occupational therapy, spiritual counseling, recreation therapy, social  
975 work, psychology and/or psychiatry
- 976 b. Patient demonstrates partial improvement (e.g., less arousal, but no  
977 improvement in sleep) – consider augmentation or adjustment of the  
978 acute intervention and follow up within 2 weeks
- 979 c. Patient recovers from acute symptoms – provide education about  
980 acute stress reaction and contact information with instructions for  
981 available follow-up if needed

## 982 DISCUSSION

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983 After initiating an acute intervention, it is crucial for providers to follow-up and  
984 assess for treatment response and for any new or additional risk factors. Studies of  
985 trauma exposed populations show that poor social supports and severe stress after  
986 the trauma may increase the risk of developing PTSD. Persons with stress reactions  
987 may respond with maladaptive coping styles or health risk behaviors, so an  
988 assessment of coping styles and health risk behaviors is warranted. Those patients  
989 who respond well to acute interventions can then be offered contact information for  
990 follow-up should they later become symptomatic.

## 991 4. FOLLOW-UP

### 992 N. Persistent (>1 Month) or Worsening Symptoms, Significant Functional Impairment or High 993 Risk for Development of PTSD.

---

#### 994 OBJECTIVE

---

995 Identify patients with PTSD or high risk for developing PTSD who may benefit from  
996 PTSD treatment.

#### 997 BACKGROUND

---

998 A crucial goal of follow-up activities is referral, as necessary, for appropriate mental  
999 health services. In fact, referral, and subsequent delivery of more intensive  
1000 interventions, will depend upon adequate implementation of screening. Screening,  
1001 whether conducted in formal or informal ways, can best help determine who is in  
1002 need of referral. But even if those who might benefit from mental health services are  
1003 adequately identified, factors such as embarrassment, fear of stigmatization,  
1004 practical barriers (e.g., distance from services), and cultural norms that do not  
1005 support help-seeking may all limit motivation to seek help or pursue a referral. Those  
1006 making referrals can directly discuss these attitudes about seeking help and attempt  
1007 to preempt avoidance of needed services. Motivational interviewing techniques



1008 (Rollnick et al., 1992) may help increase rates of referral acceptance.

#### 1009 RECOMMENDATIONS

---

- 1010 1. Individuals who fail to respond to early interventions should be referred for PTSD  
1011 treatment when they have:
  - 1012 a. Worsening of stress-related symptoms
  - 1013 b. High potential or new onset potential for dangerousness
  - 1014 c. Development of ASD/PTSD
  - 1015 d. Maladaptive coping with stress (e.g., social withdrawal, alcohol use)
  - 1016 e. Exacerbation of pre-existing psychiatric conditions
  - 1017 f. Deterioration in function
  - 1018 g. New onset stressors
  - 1019 h. Poor social supports
- 1020 2. Primary Care provider should consider initiating therapy pending referral or if the  
1021 patient is reluctant or unable to obtain specialty services.
- 1022 3. Primary Care provider should continue evaluating and treating comorbid physical  
1023 illnesses, addressing any other health concerns, as well as educating and  
1024 validating the patient regarding his/her illness.

#### 1025 DISCUSSION

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1026 Not all individuals who are exposed to trauma or who have an Acute Stress Reaction  
1027 (ASR) following trauma require a mental health referral. However, patients who are  
1028 deteriorating or not responding to acute supportive interventions need to be  
1029 identified and referred to mental health. Also, those patients who have a high  
1030 potential for dangerousness or potential for the development of PTSD also need to be  
1031 identified and referred to specialty care.

1032 Some patients with an acute stress reaction who show partial improvement may  
1033 benefit from augmentation of the acute intervention and additional follow-up.  
1034 Because people recover from traumatic stress-related problems at different rates,  
1035 some individuals may require more time, or an adjustment of the treatment, prior to  
1036 improvement. For example, early in treatment, medications may be adjusted to  
1037 target prominent symptoms.

1038 Patients with partial PTSD exhibit clinically meaningful levels of functional  
1039 impairment in association with their symptoms (Stein 1997). Functional impairment,  
1040 rates of co-morbid disorders, and rates of suicidal ideation were shown to increase  
1041 linearly with an increasing number of PTSD symptoms, and individuals with sub-  
1042 threshold PTSD had increased suicidal ideation even after controlling for the presence  
1043 of co-morbid major depressive disorder (Marshall, 2001).

1044 Patients who do not respond to first line interventions may warrant treatment  
1045 augmentation or a mental health referral. Clear indications for a mental health  
1046 referral include: a worsening of stress related symptoms, new onset of  
1047 dangerousness or maladaptive coping to stress, exacerbation of comorbid psychiatric  
1048 conditions, or deterioration in function. Because patients with new onset stressors,  
1049 poor social supports, or inadequate coping skills may be at heightened risk to  
1050 develop PTSD, mental health referral is also indicated.

1051 Primary Care providers who identify patients with possible PTSD should consider  
1052 referral to a Mental Health, or PTSD clinic. This referral should be made in  
1053 consultation with the patient, and with consideration of the patient’s severity of  
1054 problems and preferences.

1055 Several treatment modalities can be initiated and monitored in the primary care  
1056 setting (e.g. Pharmacotherapy, Supportive Counseling). Therefore, the Primary Care  
1057 practitioner should consider initiating therapy pending referral. However, if the  
1058 patient is reluctant or unable to obtain specialty services (see Module B) the Primary  
1059 Care provider should continue evaluating and treating comorbid somatic illnesses,  
1060 and addressing any other health concerns, as well as educating and validating the  
1061 patient regarding his/her illness. If patients are referred to specialty care, it is vital  
1062 that the Primary Care team (including the Health Care Integrator) stay actively  
1063 involved in coordination with the Specialist, in the care of patients with PTSD.

1064 Additional Points:

- 1065 ○ Don’t suggest or insinuate that physical or cognitive symptoms co-existing  
1066 with ASD/PTSD are related to a “stress”, “emotional”, or “psychological”  
1067 problem. Educate patients about the physiological dysregulation  
1068 associated with PTSD and how this can impact physical and cognitive  
1069 functioning.
- 1070 ○ Encourage referral to mental health via collaborative discussion, if  
1071 indicated
- 1072 ○ Educate patients about the physiological basis also of emotional and  
1073 psychological symptoms in ASD/PTSD.
- 1074 ○ Primary care providers should not hesitate to ask questions about trauma-  
1075 related symptoms, but not delve into details of the traumatic experience  
1076 itself unless there is the time and skill to manage resultant distress and  
1077 adequately support the patient. Providers should be aware that narration  
1078 of traumatic experiences may be associated with increased distress  
1079 temporarily.
- 1080 ○ Primary care providers should follow-up with patients about issues related  
1081 to trauma in an ongoing way. Patients with initial sub-threshold  
1082 presentation are at increased risk of developing PTSD.

## 1083 **O. Monitor and Follow-Up**

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### 1084 **BACKGROUND**

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1085 Many trauma survivors experience some symptoms in the immediate aftermath of a  
1086 traumatic event. In most instances these symptoms will eventually remit and do not  
1087 require long-term follow-up. Those exposed to traumatic events and who manifest  
1088 no or few symptoms after a period of time (approximately two months) do not  
1089 require routine follow-up, but follow-up should be provided if requested.

### 1090 **RECOMMENDATIONS**

---

- 1091 1. Follow-up should be offered to individuals who request it, or those at high risk of  
1092 developing adjustment difficulties following exposure to major incidents and  
1093 disasters, including individuals who:
  - 1094 a. Have acute stress disorder or other clinically-significant symptoms  
1095 stemming from the trauma
  - 1096 b. Are bereaved

- 1097 c. Have a pre-existing psychiatric disorder
- 1098 d. Require medical or surgical attention
- 1099 e. Were exposed to a major incident or disaster that was particularly
- 1100 intense and of long duration.

1101 **DISCUSSION**

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1102 For many types of trauma, experience indicates that relatively few survivors make  
1103 use of available mental health services. This may be due to a lack of awareness of  
1104 the availability of such services, low perceived need for them, lack of confidence in  
1105 their utility or negative attitudes toward mental health care. Therefore, those  
1106 planning follow-up and outreach services for survivors must consider how to reach  
1107 trauma survivors to educate them about sources of help and market their services to  
1108 the intended recipients (Excerpted from Raphael, 2000).

1109 In the chaos of some kinds of traumatic events (e.g., natural disaster), it is  
1110 important that workers systematically obtain detailed contact information to facilitate  
1111 later follow-up and outreach. In addition, it is important that those providing  
1112 outreach and follow-up efforts be opportunistic in accessing settings where survivors  
1113 are congregating. Each contact with the system of formal and informal services  
1114 available to survivors affords an opportunity to screen for risk and impairment and  
1115 intervene appropriately. Settings providing opportunities for contact with survivors  
1116 are diverse (e.g., remembrance ceremonies, self-help group activities, settings  
1117 where legal and financial services are delivered, interactions with insurance  
1118 companies). For survivors injured or made ill during the traumatic event, follow-up  
1119 medical appointments represent opportunities for reassessment, referral, and  
1120 treatment.

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**MODULE B: ALGORITHM**

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## ANNOTATIONS

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## 1. ASSESSMENT

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A. Assessment of Trauma Exposure-Related Symptoms

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BACKGROUND

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Post-traumatic stress disorder (PTSD) is the development of characteristic and persistent symptoms along with difficulty functioning after exposure to a life-threatening experience or to an event that either involves a threat to life or serious injury. Symptoms of PTSD may diminish with the passage of time, or they may persist for many years. PTSD often occurs together with, or precedes other, psychiatric illnesses. Patients are most likely to present to primary care with unexplained somatic and/or psychological symptoms (e.g., sleep disturbance, night sweats, fatigue, and difficulty with memory or concentration). The common symptoms after exposure to trauma are included in [Table B-1](#).

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The symptoms required for the diagnosis of PTSD may be divided into 3 clusters and should be present for at least 1 month.

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- **Intrusion or re-experiencing** - memories of the trauma or "flashbacks" that occur unexpectedly; these may include nightmares, intrusive mental images or extreme emotional distress and/or physiological reactivity on exposure to reminders of the traumatic event.

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- **Avoidance** - avoiding people, places, thoughts, or activities that bring back memories of the trauma; this may involve feeling numb or emotionless, withdrawing from family and friends, or "self-medicating" by abusing alcohol or other drugs.

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- **Hyperarousal** - feeling "on guard" or irritable, having sleep problems, having difficulty concentrating, feeling overly alert and being easily startled, having sudden outbursts of anger.

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PTSD is frequently under-recognized and therefore, often goes untreated. In a general survey in Israel, 9 percent of patients in a primary care setting were found to have PTSD. Only 2 percent of the sample was recognized as having the disorder. Despite this lack of recognition, more than 80 percent of men and 92 percent of women with PTSD in this survey reported significant distress from the disorder. Even individuals with "subthreshold" symptoms who do not meet full diagnostic criteria for the disorder suffer from significant impairments, including increased suicidal ideation.

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In some cases, providers may initially consider PTSD and use this guideline first, whereas in others it may be useful to follow the algorithms and recommendation of the DoD/VA guideline for Post Deployment, the VA/DoD guideline for medically unexplained symptoms or the VA/DoD guideline for Major Depressive Disorder (MDD). All these guidelines provide a link to this module when appropriate.

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RECOMMENDATIONS

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1. Patients who are presumed to have symptoms of PTSD or who are positive for PTSD on the initial screening should receive a thorough assessment of their symptoms to guide accurate diagnosis and appropriate clinical decision making.

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2. Consider using well validated self-administered checklists to ensure systematic, standardized, and efficient review of the patient's symptoms and history of

48

- 49 trauma exposure. Routine ongoing use of these checklists may allow assessment  
 50 of treatment response and patient progress. (see Appendix C: PCL-C)
- 51 3. Useful information may include details such as time of onset, frequency, course,  
 52 severity, level of distress, functional impairment, and other relevant information.
- 53 4. Behavioral health clinicians should obtain a comprehensive diagnostic assessment  
 54 that includes, but is not limited to, the symptoms that characterize PTSD.  
 55 Structured psychiatric interviews, such as the Clinician-Administered PTSD scale  
 56 (CAPS), may be considered.
- 57 5. The assessment should also include review of other salient symptoms (guilt,  
 58 dissociation, derealization, depersonalization, reduction in awareness of  
 59 surroundings) that may affect treatment planning and decision-making.

60 **Table B - 1 Common Symptoms following Exposure to Trauma**

| Physical                | Cognitive/Mental                                  | Emotional                        | Behavioral                    |
|-------------------------|---|----------------------------------|-------------------------------|
| Chills                  | Blaming someone                                   | Agitation                        | Increased alcohol consumption |
| Difficulty breathing    | Change in alertness                               | Anxiety                          | Antisocial acts               |
| Dizziness               | Confusion   | Apprehension                     | Change in activity            |
| Elevated blood pressure | Difficulty identifying familiar objects or people | Denial                           | Change in communication       |
| Fainting                | Hyper-vigilance                                   | Depression                       | Change in sexual functioning  |
| Fatigue                 | Increased or decreased awareness of surroundings  | Emotional shock                  | Change in speech pattern      |
| Grinding teeth          | Intrusive images                                  | Fear                             | Emotional outbursts           |
| Headaches               | Loss of orientation to time, place, person        | Feeling overwhelmed              | Erratic movements             |
| Muscle tremors          | Intrusive images                                  | Grief                            | Inability to rest             |
| Nausea                  | Loss of orientation to time, place, person        | Guilt                            | Change in appetite            |
| Pain                    | Memory problems                                   | Inappropriate emotional response | Pacing                        |
| Profuse sweating        | Nightmares  | Irritability                     | Startle reflex intensified    |
| Rapid heart rate        | Poor abstract thinking                            | Loss of emotional control        | Suspiciousness                |
| Shock symptoms          | Poor attention                                    |                                  | Social withdrawal             |
| Thirst                  | Poor concentration                                |                                  |                               |
| Twitches                | Poor decision-making                              |                                  |                               |
| Visual difficulties     | Poor problem-solving                              |                                  |                               |
| Vomiting                |   |                                  |                               |
| Weakness                |   |                                  |                               |

61  
 62 DISCUSSION

63 Initial screening is discussed in the CORE module (See [Core Module annotation C](#);  
 64 and [Appendix C: Screening Tools](#)).

65 The DSM-IV (1994) describes three-symptom clusters characteristic of PTSD  
 66 (reexperiencing, avoidance, and arousal) as follows:

67 The traumatic event is persistently **reexperienced** in one (or more) of the following  
 68 ways:

- 69 • Recurrent and intrusive distressing recollections of the event, including images,  
 70 thoughts, or perceptions
- 71 • Recurrent distressing dreams of the event
- 72 • Acting or feeling as if the traumatic event were recurring (includes a sense of  
 73 reliving the experience, illusions, hallucinations, and dissociative flashback  
 74 episodes, including those that occur upon awakening or when intoxicated)

- 75 • Intense psychological distress at exposure to internal or external cues that  
76 symbolize or resemble an aspect of the traumatic event
- 77 • Physiological reactivity on exposure to internal or external cues that symbolize or  
78 resemble an aspect of the traumatic event

79 Persistent **avoidance** of stimuli associated with the trauma and numbing of general  
80 responsiveness (not present before the trauma), as indicated by three (or more) of  
81 the following:

- 82 • Efforts to avoid thoughts, feelings, or conversations associated with the trauma
- 83 • Efforts to avoid activities, places, or people that arouse recollections of the  
84 trauma
- 85 • Inability to recall an important aspect of the trauma
- 86 • Markedly diminished interest or participation in significant activities
- 87 • Feeling of detachment or estrangement from others
- 88 • Restricted range of affect (e.g., unable to have loving feelings)
- 89 • Sense of a foreshortened future (e.g., does not expect to have a career,  
90 marriage, children, or a normal life span)

91 Persistent symptoms of increased **arousal** (not present before the trauma), as  
92 indicated by two (or more) of the following:

- 93 • Difficulty falling or staying asleep
- 94 • Irritability or outbursts of anger
- 95 • Difficulty concentrating
- 96 • Hypervigilance
- 97 • Exaggerated startle response

98 In the case that this syndrome originates in war experiences, the presumed cause  
99 presents itself as an exceptional event overcoming the individual's resources. The  
100 notion of war traumatization has been extended to other events such as  
101 catastrophes, physical attacks, rapes, child and wife battering, and sexual abuses.  
102 However, the events which cause PTSD are significantly more numerous. For  
103 example, it can be seen that medical events such as giving birth, miscarriage, heart  
104 attack, cancer, or hospitalization following resuscitation may give rise to PTSD.  
105 Further, people experiencing prolonged periods of distress may equally develop a  
106 post-traumatic syndrome without any one particular event having occurred to  
107 surpass their defenses.

## 108 EVIDENCE

|     | Recommendation                               | Sources  | LE  | QE   | SR |
|-----|--|--|-----|------|----|
| 109 | 1 Obtain thorough assessment of the symptoms | Lagomasino et al., 1999<br>Williams & Shepherd, 2000 | III | Poor | I  |

LE=Level of Evidence; QE = Quality of Evidence; SR =Strength of Recommendation (see Appendix A)

## 110 B. Assessment of Trauma Exposure

### 111 BACKGROUND

112 Assessment should include a careful examination of the traumatic experience itself,  
113 including the nature of the event and the patient's involvement in it, the patient's  
114 emotional, physical, and behavior responses at time of traumatization, thoughts and  
115 feelings about those responses (e.g., what he or she did or did not do).



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**RECOMMENDATIONS**

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- 117 1. Assessment of the trauma exposure experience should include:
  - 118 a. History of exposure to traumatic event(s)
  - 119 b. Nature of the trauma
  - 120 c. Severity of the trauma
  - 121 d. Duration and frequency of the trauma
  - 122 e. Age at time of trauma
  - 123 f. Patient's reactions during and immediately following trauma exposure
  - 124 (e.g., helplessness, horror, and fear)
  - 125 g. Existence of multiple traumas
- 126 6. When assessing trauma exposure, the clinician must consider the patient's ability
- 127 to tolerate the recounting of traumatic material, since it may increase distress
- 128 and/or exacerbate PTSD symptoms.
- 129 7. The assessment should be performed cautiously, especially in situations where
- 130 the trauma source is still present and the patient perceives himself or herself to
- 131 be in danger.

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**DISCUSSION**

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133 The history also should include an assessment of prior stressful life events; coping  
134 skills; ego resources and self capacities; environmental and social resources;  
135 cognitive functioning; psychiatric history; medical, family, social, and occupational  
136 history; and cultural and religious background. This background is necessary to  
137 establish an appropriate treatment plan specific to the individual patient. For  
138 example, if the individual does not feel safe in his or her current living situation,  
139 issues concerning safety need to be addressed first. Or, if the individual has a  
140 history of childhood abuse and has learned to use dissociation to protect the self,  
141 treatment will need to focus on helping the trauma victim manage his or her  
142 tendency to dissociate under stress. The repeatedly traumatized individual may also  
143 need to work through earlier childhood traumas as well as the more recent traumatic  
144 event.

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**C. Assessment of Dangerousness to Self or Others**

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**BACKGROUND**

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147 It is crucial to assess for safety and dangerousness in persons with PTSD, including  
148 current risk to self or others, as well as historical patterns of risk. Assessment of  
149 dangerousness needs to take place in a safe and secure environment and should  
150 begin with the building of rapport. If the patient has thoughts of self-harm, assess  
151 whether they have current intent and have had previous suicidal ideation, intent, or  
152 history of a suicide attempt.

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**RECOMMENDATION**

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- 154 1. All patients with PTSD should be assessed for safety and dangerousness including  
155 current risk to self or others, as well as historical patterns of risk:
  - 156 a. Suicidal or homicidal ideation, intent (plan), means (e.g., weapon,  
157 excess medications), history (e.g. violence or suicide attempts),

- 158 behaviors (e.g., aggression, impulsivity), comorbidities (substance  
159 abuse, medical conditions)
- 160 b. Family and social environment – including domestic or family  
161 violence, risks to the family
- 162 c. Ongoing health risks or risk-taking behavior
- 163 d. Medical/psychiatric comorbidities or unstable medical conditions
- 164 e. Potential to jeopardize mission in an operational environment

## 165 DISCUSSION

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166 Any history of suicidal attempts or a family history of a completed or attempted  
167 suicide should be taken seriously. Pay careful attention to patients with behaviors  
168 that may signal dangerousness (e.g., agitation, threatening, intimidation, paranoia).  
169 Access to weapons or other means of harm should also be taken seriously. Assess  
170 for domestic or family violence, because these are elevated in those with PTSD.  
171 Assessment of medical, psychiatric, and social/environmental risks is also warranted.

172 Assessment of dangerousness can include questions such as:

- 173 ○ You sound like you've had a very difficult time recently. Has life ever  
174 seemed like its not worth living?
- 175 ○ Have you ever thought about acting on those feelings? Have you thought  
176 of how you would do this?
- 177 ○ Sometimes, when people get really upset or angry, they feel like doing  
178 harm to other people. Have you had any thoughts recently about harming  
179 others?
- 180 ○ How do you express your feelings?
- 181 ○ Are there times you are afraid to go home?

### 182 ***Dangerousness to Self***

183 *Suicidality* - Persons with PTSD, including sub-threshold PTSD, are at high-risk for  
184 suicidal ideation (Marshall et al., 2001) and, for women, suicide attempts (Breslau,  
185 2000; Ferrada-Noli et al., 1998; Kaslow et al., 2000; Prigerson & Slimack, 1999).

186 Suicidal behavior is best assessed with the following criteria: presence of active  
187 depression or psychosis, presence of substance abuse, past history of suicidal acts,  
188 formulation of plan, a stated intent to carry out the plan, feeling that the world would  
189 be better off if the patient were dead, availability of means for suicide (e.g., firearms  
190 and pills), disruption of an important personal relationship, and failure at an  
191 important personal endeavor (Simon, 1992). The presence of these factors often  
192 constitutes a psychiatric emergency and must always be taken seriously. Among  
193 young adults, aggressive symptoms may be predictive of suicidality in men and  
194 elevated symptoms of PTSD and/or depression may be more predictive in women  
195 (Prigerson & Slimack, 1999). Other predictors of completed suicide in general  
196 include history of suicide attempts, family history of suicide, and access to weapons,  
197 male gender and Caucasian race. Rates of suicidal ideation in treatment-seeking  
198 Vietnam veterans have been 70 to 80 percent (Kramer et al., 1994). Additionally,  
199 Vietnam veterans with diagnosed PTSD have an increased risk of death due to  
200 suicide as compared to those without PTSD (Bullman & Kang, 1994). Among  
201 veterans with PTSD, intensive combat-related guilt has been linked to suicidality  
202 (Hendin & Haas, 1991). These findings point to the need for greater clinical

203 attention to the role of guilt in the evaluation and treatment of suicidal veterans with  
204 PTSD.

205 Individuals with severe childhood trauma (e.g., sexual abuse) may present with  
206 complex PTSD symptoms and parasuicidal behaviors, (e.g., self mutilation,  
207 medication overdoses) (Roth et al., 1997). Further, limited cognitive coping styles in  
208 PTSD have been linked to a heightened suicide risk (Amir et al., 1999). Fostering  
209 competence and social support may reduce this risk (Kotler et al., 2001). Comorbid  
210 substance use disorders may increase the risk of suicidality. Additionally, persons  
211 with PTSD may also be at personal risk of danger through ongoing or future  
212 victimization in relationships (e.g. domestic violence/battering, or rape).

- 213 • Many war veterans suffer from post-traumatic stress disorder (PTSD), depression  
214 or both disorders (Tanielian 2008 RAND). The majority of US soldiers in Iraq were  
215 exposed to some kind of traumatic, combat-related situations, such as being  
216 attacked or ambushed (92%), seeing dead bodies (94.5%), being shot at (95%)  
217 and/or knowing someone who was seriously injured or killed (86.5%) (Hoge,  
218 2004).
- 219 • In a nationally representative sample ( $N = 5877$ ; age, 15-54) that compared the  
220 relationship between anxiety disorders and suicidal ideation or suicide attempts,  
221 PTSD was significantly associated with suicidal ideation (adjusted odds ratio =  
222 2.79;  $p < 0.01$ ) and suicide attempts (adjusted odds ratio = 2.67;  $p < 0.01$ ).  
223 None of the other anxiety disorders were significantly associated with suicidal  
224 ideation or attempts (Sareen, 2005).
- 225 • Unlike the general population, older and younger veterans are more prone to  
226 suicide than are middle-aged veterans. (Zivin, 2007). Veterans with PTSD have  
227 been reported to have high levels of suicidal ideation and behaviors (Oquendo,  
228 2005).
- 229 • Jakupcak (2009) found PTSD to be a risk factor for suicidal ideation in Iraq and  
230 Afghanistan War veteran. Veterans from OEF/OIF who screened positive for PTSD  
231 were more than 4 times as likely to endorse suicidal ideation relative to non-  
232 PTSD veterans. Among veterans who screened positive for PTSD ( $n = 202$ ), the  
233 risk for suicidal ideation was 5.7 times greater in veterans who screened positive  
234 for two or more comorbid disorders relative to veterans with PTSD only.
- 235 • Patients with co-occurring disorders, such as depression and alcohol abuse or  
236 depression and posttraumatic stress disorder (PTSD), have been reported to be  
237 at much higher risk for suicide than patients with only 1 of these disorders.
- 238 • Male veterans with schizophrenia or schizoaffective disorder and comorbid PTSD  
239 were reported to have higher rates of suicidal ideation and suicidal behaviors  
240 compared to those without comorbid PTSD (Straus, 2006).
- 241 • In a large, nationally representative, longitudinal data set of depressed veterans  
242 whose causes of death have been definitively identified using linked National  
243 Death Index data, veterans who received a PTSD diagnosis had a lower rate of  
244 suicide than did veterans without PTSD (68.16 vs 90.66, respectively). The  
245 suicide rate was higher in the South than in the Northeast (88.93 vs 73.55,  
246 respectively) or central regions (88.93 vs 83.09, respectively, but slightly lower  
247 than rates in the West (88.93 vs 90.04, respectively). Veterans with a service-  
248 connected disability had a lower rate of suicide than those without a service-  
249 connected disability (70.06 vs 92.20) (Zivin, 2007).

**250      *Dangerousness to Others***

251      Some individuals with PTSD may be at risk for violence toward others (Swanson et  
252      al., 2002). Explosivity, anger problems, and past history of violence are associated  
253      with an increased risk for violent behavior. Violence often emerges as a response to  
254      a perceived threat or marked frustration by the patient stemming from his or her  
255      inability to meet goals by nonviolent means. The specific factors that contribute to  
256      violent behavior may include psychiatric, medical, environmental and  
257      situational/social engagements. Often, it is a combination of these factors that  
258      precipitates and aggravates the potential for violence, which may quickly escalate to  
259      agitation or the carrying out of violent impulses. Whatever the cause, the following  
260      situations may serve as warning signs pointing towards a very real threat of  
261      violence:

- 262      • Ideation and/or intent to harm others
- 263      • Past history of violent behaviors
- 264      • Severely agitated, aggressive, threatening or hostile behaviors
- 265      • Actively psychotic presentation

266      Special attention to the risk of domestic violence is warranted. Careful attention to  
267      the home environment and relationships is essential. If there are children, an  
268      assessment of parenting skills, anger management, caregiver burden, and discipline  
269      style is crucial. Advising high-risk patients and their families on gun removal and  
270      safe storage practices has been recommended to decrease the risk of violence  
271      (Seng, 2002). PTSD is a predictor of violence in persons with severe mental illness  
272      (Swanson et al., 2002). Also, substance use disorders are highly comorbid in PTSD  
273      and can also predict violence. Immediate attention and intervention may be  
274      required in order to ward off the potential for escalation of agitation or violent  
275      impulses.

**276      *Health Risks***

277      Persons with PTSD may have high rates of health risk behaviors, health problems,  
278      and medical conditions. Thus, an assessment of health and behavioral risks in  
279      individuals with PTSD is warranted. In addition to alcohol and drug use, persons  
280      with PTSD are at high-risk for cigarette smoking (Acierno et al., 1996). PTSD is a  
281      predictor of several HIV-risk behaviors as well as a risk factor for related blood-borne  
282      infections, such as hepatitis B and C (Hutton et al., 2001). Other potentially  
283      dangerous comorbid medical conditions are intoxication or withdrawal syndromes  
284      requiring medical detoxification (e.g., alcohol, benzodiazepine, barbiturates, and  
285      possibly opiates). Medical conditions that can present a danger to others include the  
286      risk of transmission of blood-borne pathogens such as HIV and HCV/HBV, thus risk  
287      assessment and serotesting is warranted.

**288      *Medical Conditions***

289      *Urgent conditions* - Any condition immediately threatening to life, limb, eyesight,  
290      or requiring emergency medical care requires immediate attention.

291      *Chronic diseases* - PTSD has also been linked to cardiovascular disease, anemia,  
292      arthritis, asthma, back pain, diabetes, eczema, kidney disease, lung disease,  
293      ulcers, chronic pain, work absenteeism, and other generalized health problems  
294      (Weisberg et al., 2002; Hoge et al, 2007). One explanation for these problems  
295      may relate to the association of PTSD with dysregulation of neuroendocrine, and  
296      autonomic nervous system functions. Patients who have PTSD and other chronic  
297      medical diseases may find that PTSD worsens their medical conditions. Some

298 medical conditions, which can be acutely dangerous in the presence of PTSD,  
299 include bronchial asthma, peptic ulcer disease, GI bleed and malignant  
300 hypertension, (Davidson et al., 1991).

### 301 **Psychiatric Conditions**

302 *Delirium* - (also known as organic brain syndrome, organic psychosis, acute  
303 confusional state, acute brain syndrome and various other names) is a disorder  
304 of cognition and consciousness with abrupt onset that is frequently overlooked.  
305 This is common in the elderly and medically ill (Farrell & Ganzini, 1995).

306 *Acute or marked psychosis* - "Psychosis," in and of itself, is not a psychiatric  
307 disorder. Rather, psychosis is a symptom, which may present in a variety of  
308 conditions. Psychotic patients have an impaired sense of reality, which may  
309 manifest in several forms (hallucinations, delusions, mental confusion or  
310 disorganization). Acute psychosis represents a medical emergency.

311 *Severe debilitating depression* (e.g., catatonia, malnourishment, severe  
312 disability) - While many mild to moderate illnesses may not necessarily present  
313 situations mandating immediate attention, the presence of severe depressive  
314 symptoms may represent a medical emergency, even in the absence of suicidal  
315 ideation.

#### 316 **EVIDENCE:**

|   | Recommendation  | Sources  | LE   | QE   | R |
|---|---|--|--|------|---|
| 1 | Assess for dangerousness including suicidal or homicidal ideation, intent, means, history, behaviors, and comorbidities | Breslau, 2000<br>Bullman & Kang, 1994<br>Ferrada-Noli et al., 1998<br>Kaslow et al., 2000<br>Marshall et al., 2001<br>Prigerson & Slimack, 1999<br>Swanson et al., 2002<br>Zivin, 2007 | III<br>II-2<br>III<br>II-2<br>II<br>II<br>II<br>II-2 | Good | B |
| 2 | Assess family and social environment – including risks for family   | Seng, 2002<br>Swanson, 2002  | III<br>II  | Good | B |
| 3 | Assess ongoing health risks or risk-taking behaviors  | Acierno et al., 1996<br>Hutton et al., 2001  | II-2<br>II   | Good | B |
| 4 | Assess medical or psychiatric comorbidities or unstable medical condition   | Davidson et al., 1991<br>Farrell et al., 1995<br>Weisberg et al., 2002   | II<br>III<br>III                                     | Good | B |
| 5 | In operational environment, consider the potential to jeopardize the mission  | Working Group Consensus  | III  | Poor | I |

317 *LE = Level of Evidence; QE = Quality of Evidence; SR= Recommendation ;(see Appendix A)*

## 318 **D. Obtain Medical History, Physical Examination/Laboratory Tests, and Psychosocial Assessment**

### 319 **OBJECTIVE**

320 Obtain comprehensive patient data in order to reach a working diagnosis.

### 321 **BACKGROUND**

322 A wide range of medical conditions and treatments may result in abnormal behavior,  
323 and many medical disorders may produce or exacerbate psychiatric symptoms in

324 patients with pre-existing mental illness. Multiple studies indicate high rates of  
325 medical disease (24 to 50 percent) in patients presenting with psychiatric symptoms  
326 (Williams & Shepherd, 2000). Failure to detect and diagnose underlying medical  
327 disorders may result in significant and unnecessary morbidity and mortality  
328 (Lagomasino et al., 1999). The converse problem is far greater in primary care:  
329 patients present with somatic symptoms and have psychiatric disorders that have  
330 not been properly diagnosed or treated. In one study, 5 of 6 patients with a  
331 psychiatric diagnosis had a somatic presentation, and the primary care physician  
332 made the diagnosis only half the time, whereas for the 16% with a psychological  
333 complaint, the correct diagnosis was made 94% of the time. (Bridges et al., 1985) A  
334 standardized approach to medical evaluation including a thorough history, physical  
335 examination, laboratory evaluation, and occasionally other ancillary testing prevents  
336 the omission of important aspects of the evaluation (Williams & Shepherd, 2000).

### 337 RECOMMENDATIONS

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- 338 1. All patients should have a thorough assessment of medical and psychiatric  
339 history, with particular attention paid to the following:
  - 340 a. Baseline functional status
  - 341 b. Baseline mental status
  - 342 c. Past medical history
  - 343 d. Medications: to include herbal & over-the-counter (OTC) drugs
  - 344 e. Past psychiatric history: to include prior treatment for mental health  
345 and substance use disorder, and past hospitalization for depression or  
346 suicidality
  - 347 f. Current life stressors
- 348 2. If trauma exposure is recent (<1 month) particular attention should be given to  
349 the following:
  - 350 a. Exposure to/Environment of trauma
  - 351 b. Ongoing traumatic event exposure
  - 352 c. Exposure, perhaps ongoing, to environmental toxins
  - 353 d. Ongoing perceived threat
- 354 3. All patients should have a thorough physical examination. On physical  
355 examination, particular attention should be paid to the neurological exam and  
356 stigmata of physical/sexual abuse, self-mutilation, or medical illness. Note  
357 distress caused by, or avoidance of, diagnostic tests/examination procedures.
- 358 4. All patients, particularly the elderly, should have a Mental Status Examination  
359 (MSE) to include assessment of the following:
  - 360 a. Appearance and behavior
  - 361 b. Language/speech
  - 362 c. Thought process (loose associations, ruminations, obsessions) and  
363 content (delusions, illusions and hallucinations)
  - 364 d. Mood (subjective)
  - 365 e. Affect (to include intensity, range, and appropriateness to situation  
366 and ideation)

- 367 f. Level of Consciousness (LOC)
- 368 g. Cognitive function
- 369 h. All patients should have routine laboratory tests as clinically indicated,  
370 such as TSH, Complete Metabolic Panel, Hepatitis, HIV, and HCG (for  
371 females). Also consider CBC, UA, Tox/EtoH panel and other tests.
- 372 i. Other assessments may be considered (radiology studies, ECG, and  
373 EEG), as clinically indicated.
- 374 j. All patients should have a narrative summary of psychosocial  
375 assessments to include work/school, family, relationships, housing,  
376 legal, financial, unit/community involvement, and recreation, as  
377 clinically appropriate.

## 378 DISCUSSION

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379 Differential diagnosis is important given the many co-morbidities associated with  
380 PTSD including dementia, depression, substance abuse/withdrawal, bereavement,  
381 psychosis, bipolar disorder, seizure disorder, persistent post concussion syndrome,  
382 thyroid disease, neoplasm, somatoform-spectrum disorders (including irritable  
383 bowel, chronic fatigue, headaches, and non-cardiac chest pain), anxiety disorders,  
384 toxicosis, rheumatoid-collagen vascular disease, hypoxia, sleep apnea, closed head  
385 injury, CHF, and delirium.

### 386 **Medical and Psychiatric History**

387 The medical history may be obtained from the patient, family, friends, or coworkers  
388 or from official accounts of a traumatic event, and needs to include:

- 389 • Baseline functional/mental status
- 390 • Past medical history
- 391 • Exposure to/environment of trauma (severity, duration, ongoing risk, and  
392 individual vs. community exposure)
- 393 • Medications: to include herbal & OTC drugs
- 394 • Past psychiatric history to include: prior treatment, past hospitalization for  
395 depression or suicidality, and/or inability to function in usual life roles
- 396 • Substance use and misuse can cause, be caused by, and/or exacerbate PTSD.  
397 Use of screening tools (such as the AUDIT, MAST or DAST can improve detection  
398 of substance use disorders (see the VA/DoD Guideline for Substance Use  
399 Disorder)
- 400 • Family history of PTSD or psychiatric disorder
- 401 • Sleep/eating patterns
- 402 • Current life stressors
- 403 • Risk factors suggesting the need for a higher than usual index of suspicion -  
404 certain physiological and psychological conditions or life events may contribute to  
405 the development or exacerbation of PTSD symptoms (see Annotation F).

### 406 **Physical Examination**

407 A brief, screening physical examination may uncover endocrine, cardiac,  
408 cerebrovascular, or neurologic disease that may be exacerbating or causing  
409 symptoms. Particular attention should be given to a neurologic examination and  
410 stigmata of physical/sexual abuse, self-mutilation, or medical illness. Special note  
411 should also be made of distress caused by or avoidance of diagnostic tests or

412 examination procedures, since these reactions may be suggestive of prior physical or  
413 sexual abuse. Careful attention should also be given to complying with legal  
414 mandates for documentation, reporting, and collection of evidence.

#### 415 ***Mental Status Examination (MSE)***

416 Particularly in the elderly patient, a full Mental Status Examination (MSE) includes a  
417 cognitive screening assessment. The assessment may consist of using a  
418 standardized instrument such as the Folstein Mini-Mental State Examination (MMSE)  
419 (Crum RM, et al., 1993; Cummings JL, 1993; Folstein et al., 1975). If screening is  
420 suggestive of cognitive impairment and the patient is not delirious, then a laboratory  
421 evaluation to assess for reversible causes of dementia is appropriate. However, the  
422 PTSD assessment should be continued. If delirium is present, consider it an  
423 emergency and stabilize the patient before continuing with the PTSD assessment.

424 Level of Consciousness (LOC) should be assessed to rule out delirium. Abnormal tics  
425 or movements should be noted as well as dysarthria, dysprosody, aphasia, agraphia,  
426 and alexia, which may suggest underlying neurological disease. Sensory illusions  
427 may be seen in neurologic syndromes and intoxications (Lagomasino et al., 1999).

428 Consider seeking further evaluation and consultation from neuropsychology specialty  
429 in cases of suspected cognitive disorders.

#### 430 ***Laboratory Evaluation***

431 The history and physical examination findings should be used to direct a conservative  
432 laboratory evaluation. There is no test for PTSD, but PTSD is frequently co-morbid  
433 with substance use disorders, depression, and high-risk behaviors. Therefore,  
434 testing is directed toward detection of associated medical conditions and to rule out  
435 any contraindications to medical therapy. Appropriate laboratory studies include:  
436 TSH, Complete Metabolic Panel, Hepatitis, HIV, and HCG (for females). Also consider  
437 CBC, UA, Tox/EtoH Panel and other tests, as clinically indicated.

#### 438 ***Other Evaluation***

- 439 • Diagnostic imaging and neuropsychological testing are not a part of the standard  
440 evaluation for PTSD. Proceed with management while awaiting the completion of  
441 the laboratory evaluation.
- 442 • MRI/CT of the head may be indicated to rule out mass lesions, intracranial  
443 bleeding, hydrocephalus, or subdural hematomas (Lagomasino et al., 1999).
- 444 • ECG may rule out underlying cardiac abnormalities that preclude the use of  
445 medications, such as tricyclic antidepressants (Lagomasino et al., 1999).
- 446 • Consider EEG or other diagnostic testing, as indicated by history and physical  
447 exam.

#### 448 ***Psychosocial Assessment***

- 449 • Past psychiatric illness, treatment, or admission
- 450 • Past/ongoing problems with anxiety, impulsivity, mood changes, intense/unstable  
451 interpersonal relationships, suicidality, and hallucinations
- 452 • Recreational use of drugs/alcohol/tobacco/caffeine
- 453 • Social supports (family, friends, homelessness/housing, community, and financial  
454 status)
- 455 • Losses (bereavement, friend/family member injuries/death, occupation, and  
456 moral injury/betrayal)
- 457 • Occupational/educational/military history



- 458 • Legal issues  
 459 • Religious/spiritual history  
 460 Consider use of checklists to determine if psychosocial rehabilitation services are  
 461 indicated in PTSD treatment (Foa et al., 2000). (See Module I-2: D. Psychosocial  
 462 Rehabilitation Intervention)

463 **EVIDENCE**

|   | Recommendation                               | Sources  | QE  | QE   | R |
|---|--|--|-----|------|---|
| 1 | Obtain thorough history and physical         | Lagomasino et al., 1999<br>Williams & Shepherd, 2000 | III | Poor | I |
| 2 | Obtain appropriate laboratory evaluations    | Lagomasino et al., 1999<br>Williams & Shepherd, 2000 | III | Poor | I |
| 3 | Perform radiological assessment as indicated | Lagomasino et al., 1999                              | III | Poor | I |
| 4 | Obtain thorough psychosocial assessment      | Lagomasino et al., 1999<br>Penk, 2009                | III | Poor | I |

464 *LE = Level of Evidence; QE = Quality of Evidence; SR= Recommendation (see Appendix A)*

465 **E. Assessment of Functioning/Fitness for Duty**466 **BACKGROUND**

467 One of the key goals of care is to assist the individual in beginning to return to a  
 468 normal level of functioning. The clinician must assess the individual's current level of  
 469 family, relationship, work/school, and social functioning.

470 **RECOMMENDATION**

- 471 1. Assessment of global function should be obtained through use of standardized,  
 472 targeted, and validated instruments designed to assess family/relationship,  
 473 work/school, and/or social functioning.

474 **DISCUSSION**475 ***Global Functional Assessment***

476 Consider using instruments such as the GAF (American Psychiatric Association, 1994)  
 477 or the SF-36 (McHorney, 1994) to assess global function. Such measures are useful  
 478 for directing therapeutic interventions and monitoring response to treatment.

479 ***Narrative Functional Assessment***

480 Functional assessment must be considered from the patient's point of view as well as  
 481 from the clinician's point of view. A narrative account provides a more complete  
 482 picture of the patient and his/her response to trauma. It allows for targeted social  
 483 and behavioral interventions. Components of functional assessment should include:  
 484 work/school, relationships, housing, legal, financial, unit/community involvement,  
 485 and recreation.

486 **Table B - 2 Components of Functional Assessment**

|             |   |
|-------------|---|
| <b>Work</b> | <ul style="list-style-type: none"> <li>• Is the person unemployed or seeking employment?</li> <li>• If employed, any changes in productivity?</li> <li>• Have co-workers or supervisors commented on any recent changes in appearance, quality of work, or relationships?</li> <li>• Tardiness, loss of motivation, loss of interest?</li> <li>• Been more forgetful, easily distracted?</li> </ul> |
|-------------|---|

|                                     |   |
|-------------------------------------|---|
| <b>School</b>                       | <ul style="list-style-type: none"> <li>• Changes in grades?</li> <li>• Changes in relationships with friends?</li> <li>• Recent onset or increase in acting out behaviors?</li> <li>• Recent increase in disciplinary actions?</li> <li>• Increased social withdrawal?</li> <li>• Difficulties with concentration and short-term memory?</li> </ul>   |
| <b>Family Relationships</b>         | <ul style="list-style-type: none"> <li>• Negative changes in relationship with significant others?</li> <li>• Irritable or easily angered by family members?</li> <li>• Withdrawal of interest in or time spent with family?</li> <li>• Any violence within the family?</li> <li>• Parenting difficulties?</li> </ul>   |
| <b>Recreation</b>                   | <ul style="list-style-type: none"> <li>• Changes in recreational interests?</li> <li>• Decreased activity level?</li> <li>• Poor motivation to care for self?</li> <li>• Sudden decrease in physical activity?</li> <li>• Anhedonia?</li> </ul>   |
| <b>Housing</b>                      | <ul style="list-style-type: none"> <li>• Does the person have adequate housing?</li> <li>• Are there appropriate utilities and services (electricity, plumbing, other necessities of daily life)?</li> <li>• Is the housing situation stable?</li> </ul>  |
| <b>Legal</b>                        | <ul style="list-style-type: none"> <li>• Are there outstanding warrants, restraining orders, or disciplinary actions?</li> <li>• Is the person regularly engaging in, or at risk to be involved in, illegal activity?</li> <li>• Is patient on probation or parole?</li> <li>• Is there family advocacy/ Dept. of Social Services (DSS) involvement?</li> </ul>   |
| <b>Financial</b>                    | <ul style="list-style-type: none"> <li>• Does the patient have the funds for current necessities, including food, clothing, and shelter?</li> <li>• Is there a stable source of income?</li> <li>• Are there significant outstanding or past-due debts, alimony, child support?</li> <li>• Has the patient filed for bankruptcy?</li> <li>• Does the patient have access to healthcare and/or insurance?</li> </ul> |
| <b>Unit / Community Involvement</b> | <ul style="list-style-type: none"> <li>• Does the patient need to be put on profile, MEB, or limited duty?</li> <li>• Is patient functional and contributing in the unit environment?</li> <li>• Is there active/satisfying involvement in a community group or organization?</li> </ul>  |

487

488 **EVIDENCE**

|   | Recommendation                          | Sources of Evidence     | QE  | QE   | SR |
|---|---|-------------------------|-----|------|----|
| 1 | Conduct Global Functional Assessment    | Working Group Consensus | III | Poor | I  |
| 2 | Conduct Narrative Functional Assessment | Working Group Consensus | III | Poor | I  |

489 *LE = Level of Evidence; QE = Quality of Evidence; SR= Recommendation (see Appendix A)*

490 **F. Assessment of Risk/Protective Factors**

491 **BACKGROUND**

492 Following a traumatic event, a majority of those exposed may experience post-  
493 traumatic stress responses. Of the population of persons who experience a traumatic  
494 event, only a subset will ultimately develop the disorder. After 9 to 12 months, 15 to  
495 25 percent continue to be disturbed by these symptoms. This group with persistent  
496 symptoms may have a distinct combination of characteristics that determine the  
497 presence of these ongoing problems. The presence of, and interplay among three  
498 groups of risk factors; biological factors (including genetics), the nature of the  
499 trauma, and the recovery environment (psychological and social support) work  
500 together to contribute to an individual's vulnerability or resilience to PTSD.

501 **RECOMMENDATIONS**

- 502 1. Patients should be assessed for risk factors for developing PTSD. Special  
503 attention should be given to post-traumatic factors (i.e., social support, ongoing  
504 stressors, and functional incapacity) that may be modified by intervention.
- 505 2. When evaluating risk factors for PTSD, the clinician should keep in mind that  
506 PTSD is defined as occurring only after four weeks have elapsed following a  
507 traumatic event. PTSD symptoms, however, may not appear until a considerable  
508 time has passed, sometimes surfacing years later.

509 **DISCUSSION**510 **Risk Factors for PTSD**

511 Two major systematic reviews of predictors of PTSD have been published (Brewin *et al*,  
512 2000; Ozer *et al*, 2003). The main outcome measure considered in the reviews  
513 was effect size calculated for the different factors. Effect sizes give an indication of  
514 the magnitude of the associations found.

515 Brewin and colleagues (2000) found that gender, psychiatric history, history of child  
516 abuse, and prior adversity play a role in the development of PTSD. Individually, the  
517 effect size of all the risk factors is modest, but factors operating during or after the  
518 trauma, such as level of trauma severity, lack of social support, and additional life  
519 stress have somewhat stronger effects than pre-traumatic factors (Brewin *et al.*,  
520 2000).

521 The review by Ozer *et al* (2003) focused on personal characteristics salient for  
522 psychological processing and functioning, and aspects of the traumatic event or its  
523 sequelae. Dissociation during the trauma, perceived support and perceived life threat were  
524 strongly associated with PTSD. Prior trauma and prior (in early childhood or in adult life)  
525 adjustment factors were identified among the pre-trauma factors. Prior trauma was more  
526 strongly related to PTSD when the traumatic experience involved non-combat interpersonal  
527 violence than when the traumatic experience resulted from combat or an accident. Perceived  
528 life threat was more associated when assessment was further away from the traumatic event  
529 and in non-combat interpersonal violence than in accidents. Perceived social support was also  
530 more significant in studies that assessed individuals further away from the time of the traumatic  
531 event. Family history of psychiatric disorders was more significant among survivors of non-  
532 combat interpersonal violence than when the traumatic experience was combat exposure.

533 The following characteristics have been reported in studies to be risk factors for the  
534 development of PTSD:

535 **Pre-traumatic factors**

- 536 • Ongoing life stress or demographics
- 537 • Lack of social support
- 538 • Age at time of trauma (school age youth, 40-60 years of age at highest risk)
- 539 • Pre-existing psychiatric disorder
- 540 • Female gender
- 541 • Low socioeconomic status, lower level of education, lower level of intelligence,
- 542 race (Hispanic, Japanese, other Ethnic minority)
- 543 • Prior trauma exposure ( reported abuse in childhood, report of other previous
- 544 traumatization, report of other adverse childhood factors
- 545 • Family history of psychiatric disorders (Genetics)

546 **Peri-traumatic or trauma-related factors**

- 547 • Severe trauma
- 548 • Type of trauma (interpersonal traumas such as torture, rape or assault convey a
- 549 high risk of PTSD)
- 550 • High perceived threat to life
- 551 • Age at trauma (School age youth, and 40-60 years of age)
- 552 • Community (mass) trauma
- 553 • Peri-traumatic dissociation

554 **Post-traumatic factors**

- 555 • Ongoing life stress
- 556 • Lack of positive social support
- 557 • Negative social support (e.g., negative reactions from others)
- 558 • Bereavement
- 559 • Major loss of resources
- 560 • Other post-traumatic factors including: children at home and distressed spouse

561

562 ***Pre-Traumatic Factors***

563 *Prior exposure* to traumatic events is a risk indicator for chronic PTSD (Brewin et al.,  
564 2000; Ozer et al., 2003). In particular, a history of exposure to interpersonal  
565 violence, in childhood or adulthood, substantially increases the risk for chronic PTSD  
566 following exposure to any type of traumatic event (Breslau, 2002<sup>a</sup>; Brewin et al.,  
567 2000; Ozer et al., 2003). Green et al. (2000) surveyed 1,909 college-aged women  
568 and found that those who had experienced interpersonal trauma, and those who had  
569 experienced multiple traumas, exhibited elevated symptoms. Dougall et al. (2000)  
570 hypothesized that prior trauma history sensitizes victims to the new stressor, thus  
571 potentiating its impact. They argued that evaluating trauma history is essential for  
572 improving early intervention efforts.

573 Epidemiological studies have yielded higher rates of PTSD *in women* than in men in  
574 general populations, and there are also a number of gender differences in clinical  
575 presentation after trauma. Seedat and Stein (2000) studied a series of patients  
576 presenting with physical trauma after interpersonal violence and found that “women  
577 were more likely than men to have been previously assaulted, or to have sustained

578 injury by a relative or someone known to them, but less likely to have used  
579 substances at the time of the assault or to require emergency surgery". Although  
580 there is considerable evidence suggesting a gender difference in PTSD prevalence, it  
581 is unclear whether this difference may be related to a higher risk of traumas that  
582 result in increased risk (e.g. rape) or greater willingness to seek mental health care  
583 for PTSD among women. One analysis in military personnel suggested that women  
584 and men who are working in support units with similar level of combat exposure  
585 appear to have an equivalent risk of developing PTSD (Hoge et al., 2007), and  
586 further research is needed.

587 *Pre-existing psychiatric problems* are associated with more adverse responses to  
588 trauma (Norris et al., 2002; Breslau, 2002a), in a review of recent epidemiological  
589 studies, found that preexisting psychiatric disorder was one of 3 factors that had a  
590 predictable effect on the development of PTSD. Two recent meta-analyses of risk or  
591 predictive factors for PTSD have identified prior psychiatric history as a risk factor for  
592 the development of PTSD (Brewin et al., 2000; Ozer et al., 2003). A *family history* of  
593 psychiatric disorders may also contribute to a person's vulnerability to PTSD. Brewin  
594 and colleagues (2000) found that "factors such as psychiatric history, reported  
595 *childhood abuse*, and family psychiatric history ... had more uniform predictive  
596 effects" than did other risk factors such as gender or age at trauma.

597 *Genetics* – Family history of any psychiatric disorders or possible genetic differences  
598 in regulating pre-synaptic uptake of serotonin (or other neurobiological mechanism)  
599 can increase risk. Genetic research has shown that, of the two variants of the gene  
600 regulating pre-synaptic uptake of serotonin, the long form appears to be associated  
601 with resilience and the short form with the vulnerability to stress events. Individuals  
602 who inherited the short form and were exposed to four of more stressful life events  
603 were much more likely to develop PTSD and depression and to attempt suicide  
604 (Koenen et al., 2009). Other genes that may confer vulnerability or resilience are  
605 currently under investigation. Twin studies have also indicated that there is a genetic  
606 vulnerability to PTSD. Twin research to date suggests that exposure to assaultive  
607 trauma is moderately heritable whereas exposure to non-assaultive trauma is not.  
608 PTSD symptoms are moderately heritable, and comorbidity of PTSD with other  
609 disorders may be partly due to shared genetic and environmental influences  
610 (Koenen, 2008, Afifi, 2010)

### 611 ***Peri-Traumatic Factors***

612 Foy et al (1984) published one of the first formal studies to look at risk factors for  
613 PTSD and reported characteristics of trauma exposure to be of central importance.  
614 Numerous studies have since observed a dose-response relationship between trauma  
615 severity and PTSD. The more severe the trauma, the more likely the person  
616 experiencing it will develop PTSD. Armenian and colleagues (2000) found this to be  
617 true among disaster victims. Feehan et al. (2001) found higher PTSD rates among  
618 more severely-traumatized members of a general cohort.

619 With regards to type of trauma, interpersonal violence (rape, torture, physical  
620 assault) was found to be more likely to produce PTSD than more impersonal events  
621 (such as accidents or group trauma) Holbrook et al. (2001).

622 Situations where the trauma is potentially life-threatening also carry a high risk of  
623 PTSD: in a meta-analysis of 68 PTSD studies, Ozer et al. (2003) found "perceived life  
624 threat" to have a high risk value, and in Woods' study of abused women, the  
625 perceived threat of homicide played a role in the later development of PTSD.  
626 Holbrook et al. (2001) diagnosed 261 (32%) of 824 individuals as having PTSD 6  
627 months after major physical trauma. Patients who were totally incapacitated,

628 experienced physical injury, or suffered major losses were also at higher risk for  
629 developing PTSD. Factors associated with a PTSD diagnosis included perceived threat  
630 to life, female gender, younger age and lower income.

631 Ozer et al. (2003) also found that dissociation at the time of the trauma is predictive  
632 of later development of PTSD. Demographic factors may also be predictive.  
633 Finnsdottir & Elklit (2002) found higher rates of PTSD among disaster victims who  
634 were young at the time of the trauma, and in a general group of psychiatric patients.  
635 Neria et al. (2002) found young age at trauma to be a risk factor for PTSD. Finally,  
636 biological factors may also be relevant to predicting PTSD. Shalev et al. (1998)  
637 measured the heart rate and blood pressure of eighty-six trauma survivors at the  
638 time of their presentation at a hospital emergency room, and concluded that  
639 "Elevated heart rate shortly after trauma is associated with the later development of  
640 PTSD." In a meta-analysis, Yehuda et al. (1998a) reported that studies  
641 "demonstrated increased heart rate and lower cortisol levels at the time of the  
642 traumatic event in those who have PTSD at a follow-up time compared to those who  
643 do not."

#### 644 ***Post-Traumatic Factors***

645 The post-trauma environment has been shown to be an important predictor of  
646 chronicity (Berwin, 2000). The experience of traumatization may have life-altering  
647 consequences in terms of social status, employment, and health, and continuing  
648 difficulties in these areas may contribute to the likelihood that a person will develop  
649 PTSD. Feehan et al. (2001) in interviews with 374 trauma survivors, found  
650 unemployment to be a risk factor. Likewise, in the meta-analysis performed by  
651 Norris et al. (2002) "resource loss" was cited as a risk for PTSD.

652 Impaired social support is a not-infrequent outcome of a traumatic experience.  
653 Armenian et al. (2000), Brewin et al. (2000), Gregurek et al. (2001), and Ozer et al.  
654 (2003) all reported that the loss of support from significant others can pose a risk for  
655 development of PTSD.

656 And finally, general ongoing life stress may also play a role. Brewin et al. (2000)  
657 reported "life stress" to be more predictive of PTSD development than pre-traumatic  
658 factors such as gender or age at trauma. Norris et al. (2002) found that, in disaster  
659 victims, "secondary stressors" increased the likelihood of adverse outcomes.

660 Some have suggested that secondary gain related to compensation may predict  
661 treatment outcome. Laffaye et al. (2008) found that initial levels of perceived  
662 support and stressors did not predict the course of chronic PTSD symptoms.  
663 Furthermore, the literature indicates that veterans who are seeking or have been  
664 awarded compensation participate in treatment at similar or higher rates than do  
665 their non-compensation-seeking counterparts. Veteran treatment outcome studies  
666 produced either null or mixed findings, with no consistent evidence that  
667 compensation-seeking predicts worse outcomes. Studies of motor vehicle accident  
668 survivors found no association between compensation status and course of recovery  
669 (Laffaye, 2007).

#### 670 **Risk Factors for PTSD in the military veterans**

671 Friedman et al. (1994) concluded that, "the likelihood of developing chronic PTSD  
672 depends on premilitary and postmilitary factors in addition to features of the trauma  
673 itself. Premilitary factors include negative environmental factors in childhood,  
674 economic deprivation, family psychiatric history, age of entry into the military,  
675 premilitary educational attainment, and personality characteristics. Postmilitary

676 factors include social support and the veteran's coping skills. Among military  
677 personnel, there are three populations at risk for unique problems that may amplify  
678 the psychological impact of war-zone stress. They are women whose war-zone  
679 experiences may be complicated by sexual assault and harassment; . . . and those  
680 with war-related physical disabilities, whose PTSD and medical problems often  
681 exacerbate each other."

682 Among military service members, combat exposures are reported as the strongest  
683 predictors of subsequent PTSD ( Berwin, 2000; Clancy 2006; Foy, 1987; Baker  
684 1997; Smith 2008). The frequency and intensity of direct combat appears to be one  
685 of the strongest predictors of PTSD. A number of studies have found a strong dose  
686 response relationship of combat frequency and intensity to PTSD prevalence (e.g.  
687 Hoge, et Al., 2004; Dohrenwend Science 2006; MHAT6 report, Smith Millen Cohort  
688 Study 2008).

689 Studies of veterans have reported gender differences in PTSD risks: war-zone  
690 stressors appear preeminent for PTSD in men, post-trauma resilience-recovery  
691 variables more important for women (King et al., 1999).

692 Clancy et al., 2006 examined the effect of exposure before, during or after military  
693 service. Findings indicated that nonmilitary-related trauma was prevalent among the  
694 veterans sample (90%). Regression analyses for PTSD symptom severity revealed  
695 that age, greater combat exposure, and a history of physical assault after military  
696 service were significantly associated with more severe PTSD symptoms. Childhood  
697 physical abuse, adult sexual trauma, and a history of being physically assaulted  
698 during military service were also significantly associated with PTSD symptom  
699 severity.

700 Injury severity was a significant predictor of any mental health diagnosis and PTSD  
701 diagnosis. Gunshot wounds and diastolic blood pressure were significant predictors of  
702 any mental health diagnosis, but not PTSD. A study of a sample of 1,968 men (831  
703 battle injuries and 1,137 nonbattle injuries) injured during Operation Iraqi Freedom  
704 (OIF) found, those that battle injuries compared with nonbattle injuries had a greater  
705 risk of PTSD and other mental health diagnosis, and there was a positive association  
706 with injury severity (MacGregor et al., 2009). Aggressive pain control after injury  
707 has shown, in one study, to reduce the incidence of PTSD. The study (Holbrook,  
708 2010) found that the use of morphine during trauma care may reduce the risk of  
709 subsequent development of PTSD after serious injury.

710 One semi-prospective study, (Zohar, 2009) examined risk factors for the  
711 development of posttraumatic stress disorder following combat trauma by comparing  
712 a large sample of war veterans (Israeli Defense Force) who developed PTSD with a  
713 matched control group of veterans who did not. Neither behavioral assessment nor  
714 training was found to predict PTSD. The predictive factors that were found were  
715 essentially nonspecific, such as cognitive functioning, education, rank, and position  
716 during the trauma, with little effect from training. The author concluded that "... an  
717 armed force that uses universal recruitment, carefully structured predrafting  
718 psychological assessment of social and individual qualifications (including motivation)  
719 failed to identify increased risk factors for PTSD. However, nonspecific factors were  
720 found to be associated with an increased risk for PTSD. This study suggests that the  
721 focus of future research on risk factors for PTSD should incorporate other domains  
722 rather than behavioral assessment alone" (Zohar et al., 2009).

723 Phillips et al. (2010) identified risk factors for PTSD among military service members  
724 as related to their combat exposure. The threat of death, serious injury, and  
725 witnessing of injury or death are significant risk factors for screening positive for

726 postdeployment PTSD among male Marines as well as violence exposures prior to  
727 entering the Marine Corps, which are independent of future combat exposures. Prior  
728 assault was also found to increase vulnerability for, rather than resilience against,  
729 PTSD symptoms among military professionals in the Millennium Cohort Study of US  
730 military cohort deployed in the wars in Iraq and Afghanistan (Smith, 2008).

731 PTSD symptoms among service members deployed to Iraq or Afghanistan have been  
732 associated with lower rank, being unmarried, less formal education, and a history of  
733 childhood adversity (Smith 2008; Iversen 2008).

734 The intrapersonal characteristic of hardiness as well as postwar social support may  
735 be protective against developing PTSD. In contrast, negative life events in the  
736 postwar or trauma period are linked to PTSD (King et al., 1998).

737 There is evidence that a strong social support network, indicated by unit cohesion, is  
738 protective. A large social support network may diminish the association between  
739 stressful life events and PTSD symptoms (Schnurr et al., 2004; Benotsch et al., 2000  
740 Brailey et al., 2007).

741

742



742 **2. TRIAGE**743 **G. Are There Clinically Significant Symptoms Suggestive Of PTSD?**744 **BACKGROUND**

745 In the primary care setting, providers often do not have time or resources to  
746 accomplish a detailed mental health intake evaluation, so it is important for them to  
747 be comfortable with the initial evaluation and management of stress-related  
748 disorders without having to be concerned with the fine details of DSM-IV and making  
749 a definite diagnosis. Providers who perform the initial evaluation of patient with  
750 suspected PTSD should recognize that a detailed recounting of the traumatic  
751 experience may cause further distress to the patient.

752 Please refer to [Annotation A](#) for a discussion of post-traumatic symptoms.

753 **RECOMMENDATION**

- 754 1. Providers should formulate a presumptive diagnosis of stress-related disorder  
755 consistent with the DSM IV criteria for PTSD.
- 756 2. Primary care providers should consider initiating treatment or referral based on a  
757 working diagnosis of stress-related disorder.
- 758 3. Patients with difficult or complicated presentation of the psychiatric component  
759 should be referred to PTSD specialty care for diagnosis and treatment.

760 **DISCUSSION**

761 Approximately 90 percent of patients with a mental health diagnosis are seen in  
762 primary care (Gebhart, 1996).

763 Many options are available to primary care providers to treat stress-related disorders  
764 and to relieve the burden of suffering for PTSD patients, including pharmacotherapy,  
765 supportive counseling, and referral. Because these interventions can be helpful in a  
766 variety of psychiatric disorders, it is not essential that a detailed diagnostic  
767 assessment be completed prior to treating the patient.

768 In addition, a detailed recounting of the traumatic experience may cause further  
769 distress to the patient and is not advisable unless a provider has been trained and is  
770 able to support the patient through this experience.

771

772 **Table B - 3 Diagnostic criteria for Post-traumatic Stress Disorder (DSM-IV)**

|  |
|--|
| <p>A. The person has been <b>exposed to a traumatic event</b> in which both of the following were present:</p> <ol style="list-style-type: none"> <li>1. The person experienced, witnessed, or was confronted with an event or events that involved actual or threatened death or serious injury, or a threat to the physical integrity of self or others.</li> <li>2. The person's response involved intense fear, helplessness, or horror.<br/>Note: In children, this may be expressed instead by disorganized or agitated behavior.</li> </ol> |
|--|

- B. The traumatic event is persistently **re-experienced** in one (or more) of the following ways:
1. Recurrent and intrusive distressing recollections of the event, including images, thoughts, or perceptions. Note: In young children, repetitive play may occur in which themes or aspects of the trauma are expressed
  2. Recurrent distressing dreams of the event. Note: In children, there may be frightening dreams without recognizable content
  3. Acting or feeling as if the traumatic event were recurring (includes a sense of reliving the experience, illusions, hallucinations, and dissociative flashback episodes, including those that occur on awakening or when intoxicated). Note: In young children, trauma-specific reenactment may occur
  4. Intense psychological distress at exposure to internal or external cues that symbolize or resemble an aspect of the traumatic event
  5. Physiological reactivity on exposure to internal or external cues that symbolize or resemble an aspect of the traumatic event.
- C. Persistent **avoidance** of stimuli associated with the trauma and numbing of general responsiveness (not present before the trauma), as indicated by three (or more) of the following:
1. Efforts to avoid thoughts, feelings, or conversations associated with the trauma
  2. Efforts to avoid activities, places, or people that arouse recollections of the trauma
  3. Inability to recall an important aspect of the trauma
  4. Markedly diminished interest or participation in significant activities
  5. Feeling of detachment or estrangement from others
  6. Restricted range of affect (e.g., unable to have loving feelings)
  7. Sense of a foreshortened future (e.g., does not expect to have a career, marriage, children, or a normal life span).
- D. Persistent symptoms of increased **arousal** (not present before the trauma), as indicated by two (or more) of the following:
1. Difficulty falling or staying asleep
  2. Irritability or outbursts of anger
  3. Difficulty concentrating
  4. Hypervigilance
  5. Exaggerated startle response.
- E. Duration of the disturbance (symptoms in Criteria B, C, and D) is **more than 1 month**.
- F. The disturbance causes clinically significant distress or impairment in social, occupational, or other important areas of functioning.

Specify if:

**Acute:** if duration of symptoms is less than 3 months

**Chronic:** if duration of symptoms is 3 months or more

**With Delayed Onset:** if onset of symptoms is at least 6 months after the stressor

## 774 DSM-IV &amp; DSM-IV-TR Cautionary Statement

- 775 • The specified diagnostic criteria for each mental disorder are offered as  
776 guidelines for making diagnoses, because it has been demonstrated that the  
777 use of such criteria enhances agreement among clinicians and investigators.  
778 The proper use of these criteria requires specialized clinical training that  
779 provides both a body of knowledge and clinical skills.
- 780 • These diagnostic criteria and the DSM-IV Classification of mental disorders  
781 reflect a consensus of current formulations of evolving knowledge in our field.  
782 They do not encompass, however, all the conditions for which people may be  
783 treated or that may be appropriate topics for research efforts.
- 784 • The purpose of DSM-IV is to provide clear descriptions of diagnostic  
785 categories in order to enable clinicians and investigators to diagnose,  
786 communicate about, study, and treat people with various mental disorders. It  
787 is to be understood that inclusion here, for clinical and research purposes, of  
788 a diagnostic category such as Pathological Gambling or Pedophilia does not  
789 imply that the condition meets legal or other nonmedical criteria for what  
790 constitutes mental disease, mental disorder, or mental disability. The clinical  
791 and scientific considerations involved in categorization of these conditions as  
792 mental disorders may not be wholly relevant to legal judgments, for example,  
793 that take into account such issues as individual responsibility, disability  
794 determination, and competency.

795 **H. Assess for Co-Occurring Disorders**796 **OBJECTIVE**

797 Improve management of PTSD symptoms when they are complicated by the  
798 presence of a medical or psychiatric comorbidity.

799 **BACKGROUND**

800 Comorbid medical and psychiatric conditions are important to recognize, because  
801 they can modify clinical determinations of prognosis, patient or provider treatment  
802 priorities, selection of interventions, and the setting where PTSD care will be  
803 provided. Patients with PTSD have been found to frequently report physical  
804 symptoms, cognitive health concerns, and to utilize high levels of medical care  
805 services. Providers should also expect that 50 to 80 percent of patients with PTSD  
806 will have one or more coexisting mental health disorders. PTSD is strongly  
807 associated, among veterans from recent deployment (OEF/OIF), with generalized  
808 physical and cognitive health symptoms attributed to concussion/mild traumatic  
809 brain injury – mTBI.)

810 Because of the many potential etiologies of these symptoms, it is generally best to  
811 develop a collaborative care treatment strategy based in primary care, and address  
812 these health concerns simultaneously with PTSD symptoms. (See VA/DoD Clinical  
813 Practice Guideline for Post-Deployment Health). Management should focus on  
814 identifying and treating the symptoms that are causing the most impairment,  
815 regardless of the cause or diagnosis.

816 Some comorbid medical or psychiatric conditions may require early specialist  
817 consultation, in order to assist in determining treatment priorities. In some cases,  
818 these disorders may require stabilization before (or in concert with) initiation of PTSD  
819 treatment.

820

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**RECOMMENDATIONS**

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- 821 1. Providers should recognize that medical disorders/symptoms, mental health  
822 disorders, and psychosocial problems commonly coexist with PTSD and should  
823 screen for them during the evaluation and treatment of PTSD.
- 824 2. Because of the high prevalence of psychiatric comorbidities in the PTSD  
825 population, screening for depression and other psychiatric disorders is warranted  
826 (see also VA/DoD Clinical Practice Guidelines for the Management of Major  
827 Depressive Disorder [MDD] and for Bipolar Disorder).
- 828 3. Patterns of current and past use of substance by persons with trauma histories or  
829 PTSD should be routinely assessed to identify substance misuse or dependency  
830 (alcohol, nicotine, prescribed drugs, and illicit drugs) (see also VA/DoD Clinical  
831 Practice Guideline for Substance Use Disorders).
- 832 4. Pain (acute and chronic) and sleep disturbances should be assessed in all patients  
833 with PTSD.
- 834 5. Generalized physical and cognitive health symptoms - also attributed to  
835 concussion/mild traumatic brain injury (mTBI) ) and many other causes - should  
836 be assessed and managed in patients with PTSD and co-occurring diagnosis of  
837 mTBI. (See also VA/DoD CPG for mild TBI and the CPG for Post-Deployment  
838 Health)
- 839 6. Associated high-risk behaviors (e.g., smoking, alcohol/drug abuse, unsafe  
840 weapon storage, dangerous driving, HIV and hepatitis risks) should be assessed  
841 in patients with PTSD.
- 842 7. Providers should consider the existence of comorbid conditions when deciding  
843 whether to treat patients in the primary care setting or refer them for specialty  
844 mental health care. (See Annotation J)
- 845 8. Patients with complicated co-morbidity may be referred to mental health or PTSD  
846 specialty care for evaluation and diagnosis. (see Annotation J)

847

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**DISCUSSION**

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848

**Co-Occurring Conditions**

849 Comorbid conditions and psychosocial problems of significant importance to  
850 treatment planning include:

851 **Medical Conditions:** PTSD is associated with elevated rates of generalized physical  
852 and cognitive health concerns, which are thought to be mediated in part by  
853 neuroendocrine dysregulation and autonomic nervous system reactivity (Hoge et al.,  
854 2006, ). These health conditions can include chronic headaches, chronic  
855 musculoskeletal pain, memory and attention problems, fatigue, dizziness,  
856 gastrointestinal symptoms, sleep dysfunction, hypertension, rapid heart rate  
857 (sometimes in association with panic symptoms), cardiovascular disease, impulsivity,  
858 anger, sexual problems, and a variety of other health complaints. These health  
859 concerns can sometimes cluster together, and may present as multisystem problems  
860 in the same manner as somatoform-spectrum or medically unexplained physical  
861 symptom (MUPS) conditions. These symptoms have been commonly described after  
862 all wars, overlap with numerous conditions, and often have more than one potential  
863 etiology (see DoD/VA Post-Deployment Health CPG). For example, service members  
864 or veterans who present to primary care with headaches, cognitive problems,  
865 fatigue, dizziness, and / or irritability may be experiencing these symptoms as a

866 result of chronic sleep deprivation, neuroendocrine/autonomic nervous system  
867 dysregulation associated with PTSD, residual effects of injuries during deployment  
868 (including concussions/mTBIs), chronic pain, medication side effects, depression,  
869 substance misuse, or other causes. For veterans of combat, their experiences may  
870 have involved the extremes of physiological stress contributing to long-term  
871 dysregulation of neuroendocrine and autonomic nervous systems.

872 It is important for clinicians to be aware of the high medical co-morbidity of PTSD,  
873 and the fact that physical health concerns (e.g. chronic pain, headaches) may make  
874 it more difficult to treat PTSD symptoms. Because of the many potential etiologies of  
875 these symptoms, it is generally best to develop a collaborative care treatment  
876 strategy based in primary care, and address these health concerns simultaneously  
877 with PTSD symptoms. (See also the VA/ DoD Guideline for Post-Deployment Health)

878 Some medical disorders may restrict PTSD treatment options (e.g., dementia limits  
879 psychotherapeutic options; cardiac conduction problems may limit some  
880 pharmacotherapeutic options; and disorders that restrict mobility may limit ability to  
881 attend weekly treatment sessions). It is generally best to maximize medical  
882 management of these conditions first and then focus on PTSD treatment.

883 **mTBI** – Providers should have specific awareness of traumatic brain injury (TBI),  
884 particularly concussion/mTBI, in the post deployment population because of the high  
885 prevalence (occurring in 10-20% of combat veterans) and high co-morbidity with  
886 PTSD. Concussions/mTBIs are common from sports injuries, motor vehicle accidents,  
887 military training (e.g. hand-to-hand combatives), and combat. Concussion/mTBI is  
888 distinct from moderate and severe TBI clinically and epidemiologically within a  
889 neurophysiological continuum, and the approach to care for these conditions is  
890 completely different (Hoge, et al., 2009). Persistent PCS include all of the same  
891 symptoms that veterans report after combat service, and overlap with the physical  
892 and cognitive health problems associated with PTSD. It is often difficult to precisely  
893 attribute symptoms to concussive events that occurred months or years earlier.  
894 Combat-related concussions (particularly those associated with loss of  
895 consciousness) are associated with an increased risk of PTSD, presumably because of  
896 the life-threatening context of the concussion (distinct from concussions occurring in  
897 non-life threatening situations, such as sports accidents, which are not associated  
898 with PTSD).

899 **Substance Use Disorders:** Patients with PTSD frequently use alcohol and other  
900 substances in maladaptive ways to cope with their symptoms. (Approximately 40 to  
901 50 percent of PTSD patients treated in the VA have current substance use problems.  
902 Effective PTSD treatment is extremely difficult in the face of active substance use  
903 problems, unless substance use disorders are also treated. Most often, attempts to  
904 address substance problems should proceed concurrently with the direct  
905 management of PTSD. However, in cases when the substance use is severe,  
906 substance use may require initial treatment and stabilization before progressing to  
907 PTSD care (e.g., patient requires detoxification from opiates) (see Annotation J2 -  
908 Concurrent PTSD and Substance Abuse).

909 **Psychiatric Disorders:** In addition to substance use disorders, other commonly  
910 occurring mental disorders that co-exist with PTSD include: major depression,  
911 dysthymia, panic disorder, obsessive-compulsive disorder, and generalized anxiety  
912 disorder. Treatment of these disorders often occurs concurrently with therapy for  
913 PTSD, but on occasion they will take precedence. These disorders have evidence-  
914 based therapies that may pose additional effective treatment options. Comorbid  
915 disorders that are less common with PTSD, but not rare, include psychotic disorders

916 and bipolar disorder. Practitioners should be alert to comorbid eating disorders, such  
 917 as bulimia, particularly in women.

918 **Personality Disorders:** Personality disorders are long-term problems of coping that  
 919 begin in childhood or adolescence and are often associated with past abuse or  
 920 neglect and recurrent relationship problems. These patterns often result in poor  
 921 adherence to prescribed PTSD management and the primary care provider may  
 922 require early assistance and advice from the mental health care provider.

923 **Psychosocial Problems:** Associated behavior problems and psychosocial deficits  
 924 commonly present in patients with chronic PTSD include:

- 925 • Homelessness
- 926 • Suicidality
- 927 • Domestic violence or abuse
- 928 • Aggression, rage
- 929

|   | Recommendation  | Sources                 | QE  | QE   | R |
|---|---|-------------------------|-----|------|---|
| 1 | Screen for medical disorders/symptoms, mental disorders, and psychosocial problems that commonly coexist with PTSD during the evaluation and treatment of PTSD. | Working Group Consensus | III | Poor | I |
| 2 | Stabilize acute coexisting medical and/or psychiatric disorders prior to initiating PTSD treatment.   | Working Group Consensus | III | Poor | I |

930

931 **I. Educate Patient and Family**

932 **OBJECTIVE**

933 Help trauma survivors cope with ASD/PTSD by providing information that may help  
 934 them manage their symptoms and benefit from treatment.

935 **BACKGROUND**

936 Education of the trauma survivor is a core component of all PTSD treatment.  
 937 Survivors need to better understand what they are experiencing, how to cope with  
 938 reactions or symptoms, and what happens in treatment. It is also helpful to provide  
 939 this information to family members or to the patient’s significant others so that they  
 940 can more effectively support the patient’s recovery.

941 **RECOMMENDATIONS**

- 942 1. Trauma survivors and their families should be educated about PTSD symptoms,  
 943 other potential consequences of exposure to traumatic stress, practical ways of  
 944 coping with traumatic stress symptoms, co-morbidity with other medical health  
 945 concerns, and processes of recovery from PTSD, and the nature of treatments.  
 946 [C]

947 **DISCUSSION**

948 PTSD education involves teaching the survivor to label, recognize, and understand  
 949 PTSD symptoms (and other trauma-related problems) that he or she is experiencing.  
 950 Education should include discussion of the adaptive nature of many of the symptoms,

951 which have to do with survival and the body's normal responses to threat. This is  
952 particularly important if PTSD occurred after exposure encountered in an  
953 occupational context, where the person was trained to respond to critical incidents  
954 (e.g., military, firefighter, police, and other first responders). Education should also  
955 provide simple advice regarding coping (such as sleep hygiene instruction), explain  
956 what can be done to facilitate recovery, and describe treatment options. Education  
957 can help make symptoms more understandable and predictable, decrease fear of  
958 symptoms, increase social support and lessen feelings of isolation, increase  
959 awareness of coping options and reduce maladaptive coping, and help survivors  
960 decide whether to seek treatment or learn how to better participate in treatment.

961 Education should be one of the first steps of PTSD treatment. It can help establish  
962 the credibility of the treatment provider, make treatment seem immediately helpful  
963 to the patient, and help prepare the patient for next steps in treatment. In fact,  
964 education should continue throughout PTSD treatment, sometimes in brief  
965 discussions when the patient has questions and sometimes more systematically as a  
966 formal activity. It can be delivered to individuals or to groups. Because patients  
967 with PTSD often have difficulties with concentration and memory, repetition of  
968 educational information and provision of written information are important.

969 The content of PTSD-related education can include the following topics:

970 *Nature of PTSD symptoms:* It is useful to help the survivor identify and label the  
971 reactions that he or she may be experiencing, recognize that emotional and  
972 physical reactions are expected after trauma, understand how the body's  
973 response to trauma includes many of the symptoms of PTSD, and understand  
974 that anxiety and distress are often "triggered" by reminders of the traumatic  
975 experience that can include sights, sounds, or smells associated with the trauma,  
976 physical sensations (e.g., heart pounding), or behaviors of other people.

977 *Practical steps to cope with trauma-related problems:* Survivors can also be  
978 educated about ways of coping with their PTSD symptoms in order to minimize  
979 their impact on functioning and quality of life. While education about coping is  
980 not a substitute for more systematic coping skills training, information on specific  
981 topics can be useful. Survivors can be helped to distinguish between positive and  
982 negative coping actions. Positive coping includes actions that help to reduce  
983 anxiety, lessen other distressing reactions and improve the situation; they  
984 include relaxation methods, physical exercise in moderation, talking to another  
985 person for support, positive distracting activities, and active participation in  
986 treatment. Negative coping methods may help to perpetuate problems and can  
987 include continual avoidance of thinking about the trauma, use of alcohol or drugs,  
988 social isolation, and aggressive or violent actions.

989 *Nature of the recovery process and PTSD treatment:* Survivors will sometimes  
990 have unrealistic or inaccurate expectations of recovery, and may benefit from  
991 understanding that recovery is an ongoing daily gradual process (i.e., it doesn't  
992 happen through sudden insight or "cure"), and that healing doesn't mean  
993 forgetting about the trauma or having no emotional pain when thinking about it.  
994 Education about what happens in treatment is also important. Treatment  
995 providers should explain and encourage discussion of treatment options,  
996 including evidence-based treatments. This can help build motivation to  
997 participate or persist in treatment.

998

999 Despite the fact that education is a component in all PTSD treatment, and the strong  
1000 clinical consensus that exists as to the importance of education, there is little  
1001 empirical evidence that it reduces PTSD symptoms. Education is a component of  
1002 empirically-supported treatments, but has not been evaluated as a “stand alone”  
1003 treatment (nor is it intended to be delivered in the absence of other treatment  
1004 elements).

#### 1005 EVIDENCE

|   | Recommendation  | Sources                 | QE  | QE   | R |
|---|---|-------------------------|-----|------|---|
| 1 | Educate patients and family members regarding the trauma, its effects, ways of coping, and the treatment process. | Working Group Consensus | III | Poor | C |

1006 *LE = Level of Evidence; QE = Quality of Evidence; SR= Recommendation (see Appendix A)*

1007

## 1008 J. Determine Optimal Setting for Management of PTSD and Co-Occurring Disorders

### 1009 J1. Management of PTSD with Comorbidity

#### 1010 BACKGROUND

1011 When PTSD has been determined to be the primary target of intervention because it  
1012 is significantly impairing a patient’s functioning or causing a high level of distress,  
1013 the patient’s preferences and motivation, the co-occurrence of other conditions, and  
1014 the capacity to provide the necessary services should be considered in determining  
1015 the optimal setting for treatment and long-term management. The referral to  
1016 specialty care may be considered in this context.

1017 When there are co-occurring medical or psychiatric conditions, the clinician will need  
1018 to determine the best strategy for prioritizing and treating multiple disorders. In  
1019 general, these disorders should be treated concurrently with PTSD treatment,  
1020 although there are exceptions, such as severe substance dependence that requires  
1021 medical detoxification prior to other forms of treatment. One important decision  
1022 point is whether PTSD and its psychiatric comorbidities should be treated in the  
1023 primary care setting or referred to specialty mental health care.

#### 1024 RECOMMENDATIONS

##### 1025 *Consultation / Referral*

- 1026 1. PTSD and comorbid mental health conditions should be treated concurrently for  
1027 all conditions through an integrated treatment approach, which considers patient  
1028 preferences, provider experience, severity of the conditions, and the availability  
1029 of resources.
- 1030 2. Patients with PTSD and severe comorbid mental health conditions should be  
1031 treated either through referral or in consultation with a provider experienced in  
1032 treating the co-morbid conditions.
- 1033 3. Because of the profound social impairment of PTSD (caused, for example, by the  
1034 patient’s anger and avoidance symptoms), close friends and family members in  
1035 the patient’s immediate daily environment (e.g., parents, spouse, or children)  
1036 should be provided education and advised to consider assistance from specialty  
1037 care, both for individual treatment and couples/family treatment.



- 1038 4. Factors to consider when determining the optimal setting for treatment include:
- 1039 a. Local availability of service options (specialized PTSD programs,
- 1040 evidence-based treatments, behavioral health specialty care, primary
- 1041 care, integrated care for co-occurring disorders, Vet Centers, other)
- 1042 b. Level of provider comfort and experience in treating psychiatric
- 1043 comorbidities
- 1044 c. Patient preferences
- 1045 d. The need to maintain a coordinated continuum of care for chronic
- 1046 comorbidities
- 1047 e. Availability of resources and time to offer treatment

#### 1048 DISCUSSION

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1049 A number of guiding principles should be considered in making treatment decisions  
1050 with these patients:

- 1051 • Integrated care models, in which the physical and mental health needs of  
1052 patients are addressed in a single setting by a multidisciplinary provider team,  
1053 have potential to reduce perceived stigma associated with help-seeking.
- 1054 • In systems where integrated care models do not exist, consultation and  
1055 comprehensive assessment by a mental health provider are recommended.
- 1056 • In general, referral to specialty mental health is indicated if a patient with PTSD  
1057 has comorbid mental disorders that are severe or unstable. Examples include:  
1058 patients whose depression is accompanied by suicidality, patients with substance  
1059 dependence disorder, and patients with concurrent psychotic or bipolar disorder.  
1060 If the patient is referred to mental health for treatment of PTSD, then it is usually  
1061 best for the mental health provider to provide comprehensive treatment for all  
1062 mental disorders.
- 1063 • For patients referred to specialty mental health care, it is important to preserve  
1064 the continuity of care by ensuring ongoing communication with the primary care  
1065 provider and to ensure coordination of care when multiple providers are involved.

#### 1066 **Considerations Related to Possible Referral**

1067 *Complicated severe PTSD:* Some patients with PTSD have complicated, challenging  
1068 presentations. These patients warrant referral to specialty PTSD care that includes  
1069 access to cognitive-behavioral evidence-based treatments. (See [Module I-2:](#)  
1070 [Treatment for PTSD](#))

1071 *Co-occurring major depressive disorder (MDD)* in the absence of significant  
1072 suicidality, panic, or generalized anxiety often shows reduction in intensity when the  
1073 PTSD is treated. Depression of mild severity may not require referral to specialty  
1074 care or additional treatments outside those targeting PTSD. Patients should be  
1075 carefully monitored for change in symptoms. A reduction of PTSD symptoms that is  
1076 not accompanied by reduction of symptoms in depression, or anxiety would justify a  
1077 more formally targeted treatment. (Refer to the VA/DoD guideline for MDD)

1078 *Co-occurring mild to moderate disorders such as substance use, pain disorders, and*  
1079 *sleep problems* can frequently be effectively treated in the context of PTSD  
1080 treatment and do not require a referral to specialty care. Consultation, to integrate  
1081 adjunctive interventions, may be considered. (See the respective VA/DoD CPGs)

1082 *Co-occurring severe psychiatric disorders*, while not precluding concurrent PTSD  
1083 treatment, typically justify referral to specialty care for evaluation and treatment.

1084 These disorders may include: *Severe Major Depression or Major Depression with*  
 1085 *suicidality, Unstable Bipolar disorder, Severe Personality Disorders, Psychotic*  
 1086 *disorders, Significant TBI, and Severe substance use disorder (SUD), or substance*  
 1087 *abuse* of such intensity that PTSD treatment components are likely to be difficult to  
 1088 implement.

1089 *Persistent Post Concussion Symptoms* in patients who present with PTSD and a  
 1090 history of concussion/mTBI may be best managed within the primary care setting  
 1091 without specialty referral, provided that the provider(s) has a reasonable level of  
 1092 comfort with this topic. Providers should recognize that mTBI/concussion is one of  
 1093 numerous possible etiologies of co-morbid post-deployment symptoms occurring in  
 1094 veterans and service members with PTSD, and it is often difficult to precisely  
 1095 attribute symptoms to concussive events that occurred months or years earlier.  
 1096 From a treatment standpoint, physical or cognitive symptoms, such as headaches or  
 1097 memory problems, can be treated symptomatically whether or not their underlying  
 1098 cause is PTSD, concussion/mTBI, or another condition. Primary care clinicians  
 1099 should not get caught up in debating causation, but maintain focus on identifying  
 1100 and treating the symptoms that are causing the most impairment, regardless of the  
 1101 cause.

## 1102 **J2. Concurrent PTSD and Substance Abuse**

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### 1103 **OBJECTIVE**

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1104 Improve management of PTSD symptoms when they are complicated by a  
 1105 concurrent substance abuse problem.

### 1106 **BACKGROUND**

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1107 Research has documented a strong relationship between PTSD and substance abuse  
 1108 problems in civilian and military populations of both genders (e.g., Brady, 2005;  
 1109 Kessler et al., 1995). For example, among male veterans seeking treatment for  
 1110 combat-related PTSD, high rates of lifetime alcohol use disorders and drug  
 1111 abuse/dependence have been documented (Roszell et al., 1991). Similarly, an  
 1112 extensive literature has documented high rates of PTSD among male veterans  
 1113 seeking substance abuse treatment. For example, Triffleman et al. (1995) found  
 1114 that 40 percent of substance abuse inpatient veterans had a lifetime history of  
 1115 combat-related PTSD, 58 percent had a lifetime history of PTSD due to combat or  
 1116 other traumatic exposure, and 38 percent had current PTSD.

1117 A prospective and retrospective study (Breslau et al., 2003) reported an increased  
 1118 risk for the onset of nicotine dependence and drug abuse or dependence in persons  
 1119 with PTSD, but no increased risk or a significantly ( $P = .004$ ) lower risk (for nicotine  
 1120 dependence, in the prospective data) in persons exposed to trauma in the absence of  
 1121 PTSD, compared with unexposed persons. Exposure to trauma in either the presence  
 1122 or the absence of PTSD did not predict alcohol abuse or dependence.

1123 Clinicians should note that substance abuse may mask or suppress PTSD symptoms,  
 1124 causing an individual to apparently fail to meet criteria for PTSD diagnosis.

### 1125 **RECOMMENDATIONS**

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- 1126 1. Substance use patterns of patients with trauma histories or PTSD should be  
 1127 routinely assessed (see the VA/DoD Clinical Practice Guideline for the  
 1128 Management of Substance Use Disorders).
- 1129 2. Substance abusers should be routinely screened for trauma exposure and PTSD.

- 1130 3. Patients with SUD and PTSD should be educated about the relationships between  
1131 PTSD and substance abuse, referred for concurrent PTSD and SUD treatment, or  
1132 provided with integrated PTSD/Substance Abuse treatment.
- 1133 4. There is insufficient evidence to recommend for or against any specific approach  
1134 to addressing PTSD that is comorbid with substance abuse/dependence. [I]
- 1135 5. Substance Abuse-PTSD patients should receive follow-up care that includes a  
1136 continued focus on PTSD issues.

#### 1137 DISCUSSION

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1138 Substance abusers with PTSD experience higher levels of subjective distress and  
1139 other problems than substance abusers without PTSD. For instance, compared with  
1140 women who abuse substances but do not meet diagnostic criteria for PTSD, female  
1141 PTSD-substance abusers report greater psychopathology, substance abuse problems,  
1142 dissociation, and behaviors associated with borderline personality disorder (Ouimette  
1143 et al., 1996). Patients with concurrent PTSD and substance abuse may benefit less  
1144 from conventional substance abuse treatment than those with substance abuse only  
1145 (Ouimette et al., 1998a), and PTSD is frequently under-diagnosed among individuals  
1146 receiving treatment for substance abuse (Dansky et al., 1997). These considerations  
1147 have led some authorities to develop specialized treatments that integrate treatment  
1148 for PTSD and substance abuse (Najavits, 2002; Triffleman et al., 1999).

1149 Because substance abusers with PTSD may be at higher risk for relapse and their  
1150 relapses may be “triggered” in part by trauma reminders and cues, clinicians should  
1151 adapt relapse prevention methods to help substance abuse patients identify their  
1152 trauma-related relapse cues and prepare them to cope with those triggers without  
1153 drinking or using.

1154 Because withdrawal symptoms experienced during early abstinence may be  
1155 associated with a resurgence of traumatic memories, worsening PTSD symptoms,  
1156 and, possibly, increased risk for suicidal thoughts or attempts (Kosten & Krystal,  
1157 1988), the client should be supported closely through this period, prepared for  
1158 possible short-term worsening of PTSD symptoms, and helped to develop strategies  
1159 for managing symptoms and urges to drink or use.

1160 12-step programs can play an important role in the treatment of PTSD/substance  
1161 abuse. In PTSD/substance abuse veterans hospitalized for substance abuse  
1162 disorder, greater 12-step involvement was associated with a number of positive  
1163 changes during treatment (Ouimette et al., 1998b) and with remission from  
1164 substance abuse/dependence over a two-year period. Involvement in 12-Step  
1165 groups may be especially helpful for patients who are socially isolated, lack positive  
1166 social activities and support, or lack a social group supportive of abstinence. It is  
1167 possible that PTSD patients may have special difficulties in affiliating with the groups  
1168 (e.g., social anxiety, social skills deficits, difficulties with intimacy and trust, feeling  
1169 unsafe in groups of people) and it may be appropriate in some circumstances for  
1170 clinicians to target affiliation as a treatment goal. With regard to women,  
1171 consideration should be given to the fact that exposure to 12-step environments  
1172 comprised largely of men may present a real problem for those with a history of  
1173 male-perpetrated sexual assault; use of women’s meetings may be preferable,  
1174 especially early in recovery.

1175

1176

1177 **EVIDENCE**

|   | Recommendation   | Sources   | QE   | QE   | R |
|---|--|---|------|------|---|
| 1 | Routinely assess substance use patterns of clients with trauma histories or PTSD.                      | Working Group Consensus   | III  | Poor | I |
| 2 | Integrate PTSD-Substance Abuse treatment when possible.  | Working Group Consensus   | III  | Poor | I |
| 3 | Screen substance abusing patients for trauma exposure and PTSD.  | Dansky et al., 1997   | III  | Poor | I |
| 4 | Educate substance-abusing patients with PTSD about the relationships between PTSD and substance abuse. | Working Group Consensus   | III  | Poor | I |
| 5 | Consider concurrent PTSD treatment, or provision of integrated PTSD/substance abuse treatment.         | Najavits, 2002<br>Ouimette et al., 1998<br><a href="#">Zatzick 2004</a> | II-2 | Mod  | C |
| 6 | Follow-up care for substance abuse-PTSD patients should include a continued focus on PTSD issues.      | Ouimette et al., 2000   | II-3 | Fair | I |

1178 *LE = Level of Evidence; QE = Quality of Evidence; SR= Recommendation (see Appendix A)*

1179 **J3. The Role of the Primary Care Practitioner**1180 **BACKGROUND**

1181 Primary care clinicians may decide to refer for specialized psychiatric care at any  
1182 point, depending on their level of comfortable, experience, and experience in treating  
1183 PTSD, the particular needs and preferences of the patient, and the availability of  
1184 other services.

1185 **RECOMMENDATIONS**

- 1186 1. Primary care providers should routinely provide the following services for all  
1187 patients with trauma-related disorders, especially those who are reluctant to seek  
1188 specialty mental health care:
- 1189 • Education about the disorder and importance of not letting stigma  
1190 and barriers to care interfere with specialty treatment if needed
  - 1191 • Supportive counseling
  - 1192 • PTSD-related education
  - 1193 • Regular follow-up and monitoring of symptoms
  - 1194 • Regular follow-up and monitoring of co-morbid health concerns
- 1195 2. Primary care providers should consider consultation with mental health providers  
1196 for patients with PTSD who warrant a mental health referral but refuse it or seem  
1197 reluctant to talk to a mental health provider.
- 1198 3. Primary care providers should take leadership in providing a collaborative multi-  
1199 disciplinary treatment approach. Team members may include the primary care  
1200 providers, mental health specialists, other medical specialists (e.g., neurology,  
1201 pain management), chaplains, pastors, social workers, occupational or  
1202 recreational therapists, Vet Center staff members, staff of family support centers,  
1203 exceptional family member programs, VA benefits counselors, vocational  
1204 rehabilitation specialists, peer counselors, and others.

- 1205 4. Primary care providers should continue to be involved in the treatment of  
 1206 patients with acute or chronic stress disorders. All patients with PTSD should  
 1207 have a specific primary care provider assigned to coordinate their overall  
 1208 healthcare.

1209 **DISCUSSION**

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1210 Because it is difficult for PCPs to be able to provide psychotherapy, it is  
 1211 recommended that primary care providers who identify patients with possible PTSD  
 1212 consider referral to a mental health or PTSD specialist early in the treatment process.  
 1213 This referral should be made in consultation with the patient and with consideration  
 1214 of the patient’s preferences.

1215 It is also recommended that the primary care practitioner consider initiating therapy  
 1216 (e.g., pharmacotherapy, supportive counseling) pending referral or if the patient is  
 1217 reluctant or unable to obtain mental health services. The primary care provider also  
 1218 has a vital role to play in addressing the health of patients with PTSD by evaluating  
 1219 and treating comorbid physical illnesses, by mobilizing community resources (e.g.,  
 1220 OT, Family Support, Command and Unit supports, Family Members, and Chaplains),  
 1221 and by educating and validating the patient regarding his/her illness. It is vital that  
 1222 the primary care provider and the primary care team stay actively involved, in  
 1223 coordination with the behavioral health specialist, in the care of patients with PTSD.

1224 Veterans initially seen in the primary care setting who are diagnosed with PTSD and  
 1225 are in need of a referral for treatment can be referred to the Vet Center for  
 1226 treatment. This is especially helpful if the veteran is not service-connected, cannot  
 1227 afford treatment, or feels uncomfortable in a hospital setting or sitting in large  
 1228 waiting areas with many other people. If the veteran needs a referral to community  
 1229 services, the Vet Center is also available. The staff at the Vet Center is available for  
 1230 crisis intervention and works closely with VAMC staff during the assessment phase  
 1231 and in partnership during the course of treatment.

1232 **Additional Points for Primary Care:**

- 1233 • Be careful not to suggest or insinuate that physical or cognitive symptoms co-  
 1234 existing with PTSD are related to a “stress”, “emotional”, or “psychological”  
 1235 problem. Educate patients about the physiological dysregulation associated with  
 1236 PTSD and how this can impact physical and cognitive functioning.
- 1237 • Encourage referral to mental health via collaborative discussion.
- 1238 • Educate patients about the physiological basis also of emotional and  
 1239 psychological symptoms in PTSD.
- 1240 • Do not hesitate to ask questions about trauma-related symptoms, but not delve  
 1241 into details of the traumatic experience itself unless there is the time and skill to  
 1242 manage resultant distress and adequately support the patient. Providers should  
 1243 be aware that narration of traumatic experiences, although an important  
 1244 component of treatment, may be associated with increased distress temporarily.
- 1245 • Follow-up with patients about issues related to trauma in an ongoing way.  
 1246 Patients with initial subthreshold PTSD are at increased risk of developing  
 1247 delayed PTSD.

1248  
 1249 **EVIDENCE**

---

| Recommendation | Sources | QE | QE | R |
|----------------|---------|----|----|---|
|----------------|---------|----|----|---|

|   |   |                         |     |      |   |
|---|---|-------------------------|-----|------|---|
| 3 | Consider the existence of comorbid conditions when deciding whether to refer to specialty mental health care. | Working Group Consensus | III | Poor | I |
|---|---|-------------------------|-----|------|---|

1250 LE = Level of Evidence; QE = Quality of Evidence; SR= Recommendation (see Appendix A)

1251 **K. Facilitate Social and Spiritual Support [See Module A: annotation L]**

1252 Family, religious organizations and community leaders can be helpful when dealing  
 1253 with an unfamiliar culture and/or religion. It may also be appropriate to consult a  
 1254 local cultural adviser. But particular attention should be paid to the individual’s own  
 1255 beliefs and values, and confidentiality always must be maintained when getting input  
 1256 from other sources. Patient’s beliefs should be seen in the context of their social,  
 1257 religious and cultural environment and if need be, a trusted member of the person’s  
 1258 faith or cultural group should be consulted.

1259 **RECOMMENDATIONS**

- 1260 1. Provide nonintrusive ordinary social contact (e.g., a "sounding board," judicious  
 1261 use of humor, small talk about current events, silent companionship). Provide  
 1262 opportunities for grieving for losses. (Providing space and opportunities for  
 1263 prayers, mantras, rites and rituals and end-of-life care as determined important  
 1264 by the patient).
- 1265 2. Consider providing direct spiritual care or ensuring patient access to spiritual care  
 1266 when sought.

1267 **L. Assess Duty/Work Responsibilities and Patient’s Fitness (In Relation To Military Operations)**

1268 **BACKGROUND**

1269 Ideally, service members who become ineffective as a result of PTSD will be returned  
 1270 to duty at the earliest possible time. For most military specialties, the time required  
 1271 to enlist and train the soldier to minimal operational readiness often exceeds a year.  
 1272 Consequently, service members who become ineffective due to stress-related  
 1273 conditions constitute a significant source of trained personnel who potentially have  
 1274 much to offer despite their disability. Assessment of fitness for duty may also have  
 1275 implications for medical boards and vocational rehabilitation.

1276 **RECOMMENDATION**

- 1277 1. The determination of when to return to work/duty should take into consideration  
 1278 the complexity, and importance of the patient’s job role, and functional  
 1279 capabilities.
- 1280 2. The continuing presence of symptoms of PTSD should not be considered in itself  
 1281 as sufficient justification for preventing a return to work/duty.

1282  
 1283 **DISCUSSION**

1284 Practitioners who are managing patients suffering from stress reactions or PTSD  
 1285 should consider a variety of factors when deciding if and when the individual is ready  
 1286 to return to work or military duty.

1287 First, what is the patient’s job and level of responsibility? Many patients in lower skill  
 1288 jobs (e.g., truck drivers, food service personnel, and basic supply functions) can be  
 1289 expected to function effectively despite continuing symptoms. In addition, the cost

1290 of functional failure of individuals in these roles is likely limited. In contrast, patients  
1291 in higher skill jobs or those that involve greater risks (e.g., artillery forward  
1292 observers, combat controllers, physicians, and pilots) should not be returned to duty  
1293 unless there appears to be a high probability that they have resumed effective  
1294 functioning. Individuals in leadership positions should be required to demonstrate a  
1295 higher level of reconstitution as errors on the part of these individuals can potentially  
1296 lead to much greater consequences.

1

2 **3. TREATMENT**3 **M. Initiate Treatment Using Effective Interventions for PTSD**

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4 **BACKGROUND**

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5 Many treatment strategies are available to treat stress-related disorders and to  
6 relieve the burden of suffering for PTSD patients. Options include pharmacotherapy,  
7 psychotherapy, and somatic and alternative medicine interventions. Treatment may  
8 be provided by primary care providers, specialty mental health providers, or some  
9 combination of these.

10 Primary care is an ideal setting in which to educate patients and their families about  
11 treatment options for post-traumatic stress. Patients should learn that  
12 psychotherapy for PTSD requires special training and should be referred to a  
13 therapist who has experience providing CBT. Such educational efforts must include  
14 informing patients that even if they respond to medication therapy, treatment for a  
15 longer period may be needed. The patient's preferences along with provider  
16 recommendations should drive the selection of treatment interventions in a shared  
17 decision-making process.

18 Discussion of the evidence supporting the recommendations for treatment  
19 intervention is included in Module I-2: Treatment for PTSD.

20 **RECOMMENDATIONS**

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- 21 1. A supportive and collaborative treatment relationship or therapeutic alliance  
22 should be developed and maintained with patients with PTSD.
- 23 2. Providers should explain to all patients with PTSD the range of available and  
24 effective options for PTSD treatment.
- 25 3. Patient education is recommended as an element of treatment of PTSD for all  
26 PTSD patients and their family members. [C]
- 27 4. Patient preferences along with provider recommendations should drive the  
28 selection of treatment interventions in a shared decision-making process.
- 29 5. Evidence-based psychotherapy and/or evidence-based pharmacotherapy are  
30 recommended as first-line treatment options.
- 31 6. Psychotherapies should be provided by practitioners who have been trained in the  
32 particular method of treatment, whenever possible. [Expert Consensus]
- 33 7. Patients with PTSD who are experiencing clinically significant symptoms,  
34 including chronic pain, insomnia, anxiety, or depression, should receive specific  
35 symptom management interventions.
- 36 8. A stepped care approach to treatment delivery may be considered. [Expert  
37 Consensus]
- 38 9. Specialized PTSD psychotherapies may be augmented by additional problem-  
39 specific methods/services and pharmacotherapy.
- 40 10. Consider referral for alternative care modalities (Complementary Alternative  
41 Medicine) for patient symptoms, consistent with available resources, and  
42 resonant with patient belief systems.



43 11. Management of PTSD or related symptoms may be initiated based on a  
44 presumptive diagnosis of PTSD. Long-term pharmacotherapy will be coordinated  
45 with other intervention.

46 **SPECIFIC TREATMENT MODALITIES:**

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47 **Pharmacotherapy:**

48 See Recommendations for Pharmacotherapy - Module I-2: Interventions for PTSD

49 **Psychotherapy:**

50 See Recommendations for Psychotherapy - Module I-2: Interventions for PTSD

51 **Adjunctive Treatment:**

52 See Adjunctive Treatment and other modalities, including Complementary  
53 Alternative Medicine – Module I-2: Interventions for PTSD

54 **DISCUSSION**

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55 ***Establishing Therapeutic Alliance***

56 Many people with PTSD find that their relationships with others have changed as a  
57 result of exposure to trauma. They often report that they have difficulty trusting  
58 others, are suspicious of authority, dislike even minor annoyances, and generally  
59 want to be left alone. Since the clinician-patient relationship draws heavily on trust,  
60 respect, and openness, and since the relationship often has to be formed in a busy  
61 clinical or bureaucratic setting, the provider may find the PTSD patient to seem to be  
62 withholding, negativistic, or even hostile at the initial meeting. The patient may  
63 seem to have “an attitude” or an “Axis II” comorbidity. As a result, many combat  
64 veterans have been misunderstood and misdiagnosed by otherwise competent  
65 professionals, and ultimately the patient suffers through feeling betrayed and  
66 misunderstood by the mental health professional. If a therapeutic relationship is to  
67 have any opportunity to develop, the treatment provider must adopt a stance of  
68 caring and concerned involvement that takes what the patient says at face value,  
69 doesn’t judge or label this type of behavior, and doesn’t withdraw into an “objective”  
70 “professional” role. In short, the clinician who can relate honestly and openly is  
71 more likely to have a patient who is willing to relate to him/her as a fellow human  
72 being and an effective partner in treatment.

73 A general understanding of what has happened to the veteran is critical in this  
74 process of developing a therapeutic relationship. Every provider working with  
75 combat veterans should be advised to read some basic material on the experience of  
76 combat and watch documentaries of the same. The provider must develop an  
77 understanding that wartime and military service involve some of the most intense  
78 human experiences, and those feelings of profound rage, fear, and grief can be an  
79 expected part of these experiences. These feelings will be present in the interview  
80 setting and must be met with respect and compassion. It is also helpful for the  
81 professional to be careful not to assume that they have any understanding of the  
82 military experience based if they have not themselves served in the military, and  
83 should not be afraid to ask questions when they don’t understand something about  
84 the military that the patient is referring to.

85 ***PTSD Treatment***

86 Refer to the evidence-based treatment strategies for PTSD, summarized in the  
87 section on Pharmacotherapy and Psychotherapy Intervention of this guideline. The

88 section also includes medication tables that summarize indications/benefits,  
89 contraindications/adverse effects, and usual dosages. (See [Module I-2](#))

90 Supportive counseling for PTSD has received little study to date and cannot be  
91 endorsed as an evidence-based psychotherapeutic strategy. However, it has been  
92 shown to be effective compared with no treatment and may be the sole  
93 psychotherapeutic option available for the patient with PTSD who is reluctant to seek  
94 specialty mental health care. It may be a useful engagement strategy to provide  
95 temporary support with the ultimate goal to convince patient to accept evidence-  
96 based treatment.

97 **EVIDENCE**

|   | Recommendation   | Sources                 | QE  | QE   | R |
|---|--|-------------------------|-----|------|---|
| 1 | Take steps to create a supportive, therapeutic alliance. | Working Group Consensus | III | Poor | I |

98 *LE = Level of Evidence; QE = Quality of Evidence; SR= Recommendation (see Appendix A)*

99

99 **4. RE-ASSESSMENT**100 **0. Assess response to treatment**

---

101 **OBJECTIVE**

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102 Re-assess patient status following therapeutic intervention to determine response to  
103 treatment, inform treatment decisions, and identify need for additional services. Re-  
104 assessment should address PTSD symptoms, diagnostic status, functional status,  
105 quality of life, additional treatment needs, and patient preferences.

106 **RECOMMENDATIONS**

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- 107 1. At a minimum, providers should perform a brief PTSD symptom assessment at  
108 each treatment visit. The use of a validated PTSD symptom measure, such as  
109 the PTSD Checklist, should be considered (see Appendix C).
- 110 2. More comprehensive re-assessment and evaluation of treatment progress should  
111 be conducted at least every 90 days, and perhaps with greater frequency for  
112 those in active treatment, and should include a measure of PTSD  
113 symptomatology (PCL) and strongly consider a measure of Depression  
114 symptomatology (PHQ9).
- 115 3. Other specific areas of treatment focus (e.g., substance abuse), should also be  
116 reevaluated and measured by standardized measures of outcome.
- 117 4. Assessment of functional impairment should also be made, at minimum, by  
118 asking patients to rate to what extent their symptoms make it difficult to engage  
119 in vocational, parental, spousal, familial, or other roles.
- 120 5. Consider continued assessment of:
  - 121 • Patient preferences
  - 122 • Treatment adherence
  - 123 • Adverse treatment effects

124

125 **DISCUSSION**

---

126 Patients should be assessed at least every three months after initiating treatment for  
127 PTSD, in order to monitor changes in clinical status and revise the intervention plan  
128 accordingly. The interval of three months is suggested because many controlled  
129 trials of first line therapies for PTSD demonstrate clinically-significant changes during  
130 this time frame. Assessment of the following domains is advised: (a) symptom  
131 severity and diagnostic status of PTSD, comorbid mental disorders, and comorbid  
132 medical conditions; (b) functional status and quality of life in major areas of  
133 adjustment (e.g., occupation, social and family relations, activities of daily living and  
134 capacity for self-care, physical health needs, and spiritual fulfillment); (c)  
135 psychosocial needs (e.g., financial and housing deficits); (d) patient satisfaction with  
136 treatment received and preferences for type and amount of continued treatment, (e)  
137 compliance or adherence with treatments provided; and (e) adverse side effects of  
138 pharmacological or psychosocial treatments administered.

139 A number of interview and questionnaire methods are recommended for assessing  
140 the diagnostic status and clinical severity of patients (see Annotation E). These  
141 measures may be used to identify the presence/absence of major mental disorders,

142 including PTSD, as well as the degree of symptom severity. Much of this information  
143 is important to share with patients in assessing progress of treatment and making  
144 collaborative decisions about future directions of care. The DSM-IV criteria for PTSD  
145 domains (b – e) can be routinely measured using standard clinical interview  
146 methods.

### 147 ***Regular Follow-Up and Monitoring***

148 The use of pencil-and-paper measures of PTSD symptom severity such as the PTSD  
149 Checklist (PCL; see appendix C) should be considered. Scores on the PCL may be  
150 plotted serially over time to create a longitudinal record of symptom severity and  
151 may be helpful for recognizing environmental (e.g., renewed proximity to a  
152 previously abusive parent) or seasonal (e.g., anniversary of a traumatic war event)  
153 precipitants of PTSD symptoms.

### 154 ***Early Recognition of a Psychosocial Crisis and Referral to Specialists***

155 Primary care providers may be the first to recognize that a patient with PTSD is  
156 entering a psychosocial crisis. Depending on the severity and disability associated  
157 with the crisis and the potential for harm to the patient or others, the primary care  
158 provider may be obliged to obtain specialty mental health services, even if that  
159 patient is reluctant to seek those services.

### 160 ***Coordination of General Health Care***

161 The traditional role of the primary care provider as the coordinator of various  
162 disciplines and consultants involved in the treatment of any single patient is  
163 especially relevant for the patient with PTSD. Particularly in patients with chronic  
164 PTSD, medically unexplained symptoms or problems with substance use (including  
165 smoking) may lead to the need for a wide range of specialists. Coordination of these  
166 services is important to avoid confusion and unnecessary health care use.

167

## 168 **P. Follow-Up**

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### 169 **BACKGROUND**

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170 Prevalence of partial or sub-threshold PTSD is substantial. Because of risk of relapse  
171 following discontinuation of therapy in patients with chronic PTSD, long-term  
172 treatment is often needed. Most patients with chronic PTSD (defined by the DSM-IV  
173 as full-criterion symptoms lasting 3 months or more), should be continued for at  
174 least 1 year, with regularly scheduled follow-up in order to prevent relapse. In other  
175 cases, CBT may be effective for symptoms that do not respond to SSRIs.

176 The continued importance of psychoeducation and reinforcement of health-promoting  
177 behaviors by the primary care physician is important but generally neglected area of  
178 public health.

### 179 **RECOMMENDATIONS**

---

- 180 1. If patient does not improve or status worsens, consider one of the following  
181 treatment modification options:
  - 182 a. Continue application of the same modality at intensified dose and/or  
183 frequency
  - 184 b. Change to a different treatment modality
  - 185 c. Apply adjunctive therapies

- 186 d. Consider a referral to adjunctive services for treatment of comorbid  
187 disorders or behavioral abnormalities (e.g., homelessness, domestic  
188 violence, or aggressive behavior)
- 189 e. For patient with severe symptoms or coexisting psychiatric problems  
190 consider referrals to:
- 191 • Specialized PTSD programs
  - 192 • Specialized programs for coexisting problems and conditions
  - 193 • Partial psychiatric hospitalization or “day treatment” programs
  - 194 • Inpatient psychiatric hospitalization
- 195 2. If patient demonstrates partial (insufficient) remission, consider one of the  
196 following treatment modification options:
- 197 a. Continue the present treatment modality to allow sufficient time for full  
198 response
  - 199 b. Continue application of the same modality at intensified dose and/or  
200 frequency
  - 201 c. Change to a different treatment modality
  - 202 d. Apply adjunctive therapies
  - 203 e. Increase level of care (e.g., referral facility, partial hospitalization, inpatient  
204 hospitalization, residential care)
  - 205 f. Consider a referral to adjunctive services for treatment of comorbid  
206 disorders or behavioral abnormalities (e.g., homelessness or domestic  
207 violence)
- 208 3. If patient demonstrates improved symptoms and functioning but requires  
209 maintenance treatment:
- 210 a. Continue current course of treatment
  - 211 b. Consider stepping down the type, frequency, or dose of therapy
  - 212 c. Consider:
    - 213 • Transition from intensive psychotherapy to case management  
214 contacts
    - 215 • Transition from individual to group treatment modalities
    - 216 • Transition to as needed treatment
  - 217 d. Discuss patient status and need for monitoring with the primary care  
218 provider
  - 219 e. Consider a referral to adjunctive services for treatment of comorbid  
220 disorders or behavioral abnormalities (e.g., homelessness or domestic  
221 violence)
- 222 4. If patient demonstrates remission from symptoms and there are no indications  
223 for further therapy:
- 224 a. Discontinue treatment
  - 225 b. Educate the patient about indications for and route of future care access

- 226 c. Monitor by primary care for relapse/exacerbation
- 227 5. Evaluate psychosocial function and refer for psychosocial rehabilitation, as
- 228 indicated. Available resources include, but are not limited to: chaplains, pastors,
- 229 Family Support Centers, Exceptional Family Member Programs, VA benefits
- 230 counselors, occupational or recreational therapists, Vet Centers, and peer-
- 231 support groups. (See [Module I-2 D: Psychosocial Rehabilitation](#))
- 232 6. Provide case management, as indicated, to address high utilization of medical
- 233 resources.

234 **DISCUSSION**

---

235 ***Patient Does Not Improve or Status Worsens:***

236 Re-assessment of patients' clinical status may occasionally show that symptoms

237 and/or functional status are failing to improve or are deteriorating in a sustained

238 way. It is important to determine that this static or deteriorated state is not simply

239 the result of a major life crisis unrelated to the therapy being administered.

240 The clinician must next determine if a patient's unimproved clinical status reflects a

241 temporary exacerbation of symptoms expected to occur in the course of treatment

242 that will ultimately prove to be effective. For example, it is common for patients in a

243 range of trauma-focused therapies to experience some brief distress or symptom

244 exacerbation during initial phases of treatment where they focus on emotions

245 associated with traumatic memories. In this case, it is important to reassure the

246 patient about the natural course of recovery through treatment, assist him/her in

247 coping with symptoms, and enlist him/her in the decision to continue with the

248 current method of treatment. Increasing session contacts and or increasing the dose

249 of medications may provide needed support.

250 If the clinician and patient agree that the current treatment regimen is ineffective,

251 then a collaborative decision can be made to switch to a different modality. Another

252 approach is to hold the course of a current therapy, which may appear ineffective,

253 but apply adjunctive treatments (see Module I-2: PTSD Interventions) There is no

254 empirical evidence that supports the effectiveness of combination treatments for

255 PTSD. However, there is clinical consensus that some treatments can act

256 synergistically (e.g., combining coping skills and symptom management approaches

257 with exposure-based treatments).

258 Clinicians should consider changing the treatment plan by increasing the level of care

259 offered to patients. Levels of care for PTSD vary in intensity, including infrequent

260 visits administered in outpatient clinics, partial hospital programs, specialized

261 inpatient PTSD programs, PTSD residential care programs and domiciliaries, and

262 acute inpatient hospitalization. Patients who fail to progress in outpatient treatment

263 may benefit from a temporary transition to a higher level of care, followed by return

264 to outpatient management after greater stabilization of symptoms has been

265 achieved.

266 Often, progress in PTSD treatment may be compromised by a concurrent behavioral

267 disorder (e.g., domestic violence), life crisis (e.g., homelessness), or uncontrolled

268 substance use disorder. Referral to ancillary clinical services should be considered

269 for patients for whom these problems emerge during the course of treatment, as

270 identified upon re-assessment.

271 **Patient Demonstrates Improved Symptoms and Functioning but Requires Maintenance**  
272 **Treatment:**

273 Treatment may also lead to slight or moderate improvement that nonetheless leave  
274 the patient with significant distress and impairment in functioning. If the patient  
275 demonstrates partial (insufficient) remission, consider one of the following treatment  
276 modification options:

- 277 1. Continue the present treatment approach to allow sufficient time for full  
278 response. This option might be worth considering when a treatment involves  
279 acquisition of skills (e.g., cognitive restructuring or anxiety management). In  
280 such a case, it is possible that the patient may be in the process of learning  
281 the skill, with the full impact of therapy dependent on increased practice and  
282 skill mastery. Or, treatment may not have yet yielded its maximum potential  
283 effect because of limited patient compliance; steps taken to increase  
284 adherence to treatment prescriptions may accelerate responsiveness to the  
285 intervention.
- 286 1. If the moderate level of improvement obtained is less than would be  
287 expected, given what is known about the patient and the treatment modality,  
288 a change to a different treatment approach may be indicated.
- 289 2. In certain circumstances, a move to an increased level of care may be  
290 warranted. For example, if current functioning remains poor despite some  
291 symptom improvement or the patient stands to experience major  
292 consequences for failure to improve more rapidly (e.g., marital separation), it  
293 may be desirable to move from outpatient care to a higher level of care (e.g.,  
294 residential care).
- 295 3. Improvement in PTSD symptoms may be inhibited by the presence of  
296 untreated additional problems, such as substance abuse or exposure to  
297 domestic violence. In such situations, it is important to initiate services for  
298 these problems in order to improve the capacity of the PTSD treatment to  
299 effect change.
- 300 4. Patients with partial PTSD may exhibit clinically meaningful levels of  
301 functional impairment in association with their symptoms. Functional  
302 impairment, rates of co-morbid disorders, and rates of suicidal ideation were  
303 shown to increase linearly with increasing number of PTSD symptoms, in one  
304 study, and individuals with sub-threshold PTSD had increased suicidal ideation  
305 even after controlling for the presence of co-morbid major depressive disorder  
306 (Marshall, 2001).
- 307 5. Studies in which the prevalence of partial, or sub-threshold PTSD was  
308 examined found it to be substantial. . In one study of infantry soldiers  
309 returning from Iraq, the prevalence of PTSD was estimated to be 12% when a  
310 stringent PCL definition of PTSD was utilized, but rose to 18-20% when a  
311 more liberal DSM symptom-based definition was applied (Hoge, 2004). A  
312 large Canadian epidemiological study assessing for current PTSD found the  
313 incidence to be 5.0% (women) and 1.7% (men), but the incidence of partial  
314 PTSD to be even higher at 5.7% and 2.2% for women and men respectively.  
315 Individuals with sub-threshold PTSD showed similar levels of social and  
316 occupational impairment as those meeting full criteria (Stein, 1997; marshall,  
317 2001).

318

319 ***When Symptoms and Other Trauma-Related Problems show Significant Improvement,***  
320 ***the Options include the Following:***

- 321 • Discontinue treatment.
- 322 • Continue the course of treatment as is.
- 323 • “Step down” to a treatment requiring less intensive resources.

324 Clinician judgment, based on discussion with the patient, will be the basis of such a  
325 decision.

326 When therapy has resulted in clinically-significant improvement, but the  
327 improvement in functioning is recent and of limited duration, a continuation of the  
328 existing type and intensity of treatment may be indicated if the clinician judges that  
329 time is required for the patient to continue practicing new skills or to otherwise  
330 consolidate treatment gains. This will be especially true if the clinician judges that a  
331 reduction in level of therapeutic support would threaten treatment gains.

332 If treatment has produced clear benefit, but the patient is continuing to show  
333 treatment gains week-by-week, it may also be helpful to maintain the treatment as  
334 is, in hopes of continued improvement. For many patients, some level of continuing  
335 care may be indicated after more intensive help has produced improvements. A  
336 step-down to less resource-intensive help can often be accomplished by changing  
337 treatment type (e.g., from individual psychotherapy to periodic group support),  
338 reducing frequency of contacts (e.g., from once-per-week to twice-per-month  
339 contact), or reducing treatment dose (e.g., medication).

340 If treatment has resulted in significant reductions in PTSD, but related problems  
341 (e.g., anger, social isolation, guilt) have shown little change, it will be important to  
342 consider adding treatment components to address those problems or referring the  
343 patient for additional services.

344 ***Patient Demonstrates Remission from Symptoms:***

345 When the patient demonstrates remission from symptoms and there are no  
346 indications for further therapy, it is time to discontinue treatment. Discontinuation of  
347 treatment may be anxiety-provoking for some patients, who have come to depend  
348 on the therapist. If this is the case, it may be helpful to discontinue treatment by  
349 using the step-down approach noted above, and gradually moving toward  
350 termination. Whether treatment is ended gradually, or more quickly, it is important  
351 to educate the patient about expected levels of continuing symptoms, indicators of  
352 relapse or need for future care, and ways of accessing care should the need arise.  
353 The patient can be encouraged to talk with his or her primary care provider about  
354 the treatment experience and enlist help in monitoring improvement.

355 ***Psychosocial Rehabilitation for All Patients with PTSD***

356 Patients with persistent mental health symptoms and needs may benefit from a  
357 range of assistance strategies provided by a range of disciplines. In addition to the  
358 usual general health and mental health specialists, available resources include, but  
359 are not limited to, case-management, chaplains, pastors, Family Support Centers,  
360 Exceptional Family Member Programs, VA Benefits Counselors, vocational counselors,  
361 occupational or recreational therapy, Vet Centers, and peer-support groups.

362 In the primary care setting, appropriate encouragement of patients to obtain a  
363 mental health referral is important, even if patients are initially hesitant or reluctant  
364 to seek it. Mental health referral options include outpatient psychology, social work,  
365 or psychiatry clinics, depending on local resources and policies.



366 In the specialty mental health settings, patients may be referred to specialized PTSD  
 367 programs or programs that focus treatment on important coexisting problems, such  
 368 as substance use disorder programs or programs for domestic violence or sexual  
 369 assault/abuse. Depending on the level of associated disability, complexity of  
 370 medication regimen, and level of threat to self or others, patients with persistent  
 371 PTSD symptoms and needs may require inpatient or partial psychiatric  
 372 hospitalization.

373 Providers referring from either the primary or specialty mental health setting should  
 374 consider the need for case-management to ensure that the range of patient needs is  
 375 addressed and that follow-up contact is maintained.

376 **EVIDENCE**

|   | Recommendations   | Sources                 | QE  | QE   | R |
|---|---|-------------------------|-----|------|---|
| 1 | The primary care provider should remain involved in treatment and patient monitoring. | Working Group Consensus | III | Poor | I |
| 2 | Multidisciplinary team approach   | Working Group Consensus | III | Poor | I |

377

|    |   |            |
|----|---|------------|
| 1  | <b>Module I: TREATMENT INTERVENTIONS</b>                              |            |
| 2  | <b>I-1. EARLY INTERVENTION TO PREVENT PTSD .....</b>                  | <b>108</b> |
| 3  | A. PSYCHOTHERAPY  | 110        |
| 4  | A1. Psychological Debriefing  | 110        |
| 5  | A2. Brief Early Cognitive-Behavioral Intervention                     | 114        |
| 6  | A3. Other Early Interventions   | 115        |
| 7  | B. Early Pharmacotherapy Interventions to Prevent Development of PTSD | 116        |
| 8  |   |            |
| 9  | <b>I-2. TREATMENT FOR POST-TRAUMATIC STRESS DISORDER (PTSD) ....</b>  | <b>120</b> |
| 10 | A. Selection of Therapy for PTSD                                      | 120        |
| 11 | B. PSYCHOTHERAPY INTERVENTIONS FOR PTSD                               | 121        |
| 12 | B1. Cognitive Therapy (CT)  | 125        |
| 13 | B2. Exposure Therapy (ET)   | 129        |
| 14 | B3. Stress Inoculation Training (SIT)                                 | 132        |
| 15 | B4. Eye Movement Desensitization and Reprocessing (EMDR)              | 133        |
| 16 | B5. Imagery Rehearsal Therapy (IRT)                                   | 136        |
| 17 | B6. Psychodynamic Therapy   | 137        |
| 18 | B7. Patient Education   | 139        |
| 19 | B8. Group Therapy   | 140        |
| 20 | B9. Dialectical Behavior Therapy                                      | 148        |
| 21 | B10. Hypnosis   | 150        |
| 22 | B11. Behavioral Couples Therapy                                       | 152        |
| 23 | B12. Telemedicine and Web-based Interventions                         | 153        |
| 24 | <b>C. PHARMACOTHERAPY FOR PTSD .....</b>                              | <b>156</b> |
| 25 | <b>D. ADJUNCTIVE SERVICES .....</b>                                   | <b>175</b> |
| 26 | D1. Psychosocial Rehabilitation                                       | 175        |
| 27 | D2. Spiritual Support   | 180        |
| 28 | <b>E. SOMATIC TREATMENT.....</b>                                      | <b>183</b> |
| 29 | E1. Biomedical Somatic Therapies                                      | 183        |
| 30 | E2. Acupuncture   | 185        |
| 31 | <b>F. COMPLEMENTARY AND ALTERNATIVE MEDICINE.....</b>                 | <b>186</b> |
| 32 | F1. Natural Products (Biologically Based Practices)                   | 188        |
| 33 | F2. Mind-Body Medicine  | 188        |
| 34 | F3. Manipulation and Body Based Practices (Exercise and Movement)     | 188        |
| 35 | F4. Energy medicine   | 189        |
| 36 | F5. Whole Medial Systems  | 189        |
| 37 | F6. Other Approaches  | 189        |
| 38 |   |            |

|    |   |            |
|----|---|------------|
| 39 | <b>I-3. MANAGEMENT OF SPECIFIC SYMPTOMS .....</b> | <b>190</b> |
| 40 | A. Sleep Disturbances                             | 190        |
| 41 | B. Pain   | 196        |
| 42 | C. Irritability, Severe Agitation, or Anger       | 205        |
| 43 |   |            |

**I-1. EARLY INTERVENTION TO PREVENT PTSD**

2 **BACKGROUND**

3 Several studies have examined the effectiveness of treatment interventions and  
 4 acute symptom management early (within one month) following a traumatic event in  
 5 preventing PTSD. This includes the use of various medications for the prevention of  
 6 PTSD and brief multiple sessions of psychotherapy.

7 This section summarizes the evidence supporting the recommendations for early  
 8 intervention. Table I-1 summarizes the recommendations for interventions and  
 9 their potential benefit discussed in Module A, Annotation J: Brief Intervention.

10

11 **Table I - 1 Early Intervention after Exposure to Trauma (>4 days, < 30 days)**

| SR | Significant Benefit                               | Some Benefit   | Unknown  | No Benefit Potential Harm   |
|----|---|--|--|---|
| A  | Brief Cognitive Behavioral Therapy (4-5 sessions) |  |  |   |
| B  |   | Prazosin for nightmares                                  | Imipramine   |   |
| C  |   | Social, and spiritual support                            | Propranolol  |   |
| I  |   | Provide basic Needs<br>Psychoeducation,<br>normalization | Sympatholytics<br>Antidepressants<br>Anticonvulsants<br>Atypical Antipsychotics<br>Chloral hydrate | Benzodiazpines **   |
| D  |   |  | Psychological First Aid  | Psychological debriefing<br>Typical Antipsychotics<br>Formal psychological interventions for asymptomatic survivors |

12 *SR = Strength of recommendation (see Appendix A); \*\* Potential harm*

13

13

14 **RECOMMENDATIONS**

- 15 **1.** The following treatment recommendations should apply for all acutely  
16 traumatized people who meet criteria for diagnosis of ASD, and for those with  
17 significant levels of post-trauma symptoms after at least two weeks post-trauma,  
18 as well as those who are incapacitated by acute psychological or physical  
19 symptoms.
- 20 **2.** Treatment should be provided after education, normalization, and Psychological  
21 First Aid has been provided, and basic needs following the trauma were made  
22 available.
- 23 **3.** There is insufficient evidence to recommend for or against the use of  
24 Psychological First Aid to address symptoms beyond 4 days following trauma. [I]
- 25 **4.** Recommend monitoring for development of PTSD using validated symptom  
26 measures (e.g., PTSD Checklist, other screening tools for ASD/PTSD).
- 27 **5.** Survivors who present symptoms that do not meet the diagnostic threshold  
28 should be monitored and may benefit from follow-up and provision of ongoing  
29 counseling or symptomatic treatment.
- 30 **6. Psychotherapy:**
  - 31 a. Consider early brief intervention (4 to 5 sessions) of cognitive  
32 behavioral therapy (CBT) that includes exposure based therapy, alone  
33 or combined with a component of cognitive re-structuring therapy for  
34 patients with significant early symptom levels, especially those  
35 meeting diagnostic criteria for ASD.[A]
  - 36 b. Early psychotherapy intervention for *asymptomatic* individuals is not  
37 beneficial and may be harmful. [D]
  - 38 c. Recommend against individual Psychological Debriefing as a viable  
39 means of reducing acute stress disorder (ASD) or progression to post-  
40 traumatic stress disorder (PTSD). [D]
  - 41 d. Recommend against group Psychological Debriefing as a viable means  
42 of reducing acute stress disorder (ASD) or progression to post-  
43 traumatic stress disorder. (Note: this is not a recommendation against  
44 Operational Debriefing) [D]
  - 45 e. Groups may be effective vehicles for providing trauma related  
46 education, training in coping skills, and increasing social support  
47 especially in the context of multiple group sessions.
  - 48 f. Group participation should be voluntary.
- 49 **7. Pharmacotherapy:**
  - 50 a. There is insufficient evidence to support a recommendation for use of a  
51 pharmacological agent to prevent the development of ASD or PTSD.  
52 [I]

53 For discussion of use of medication for specific symptom management  
54 during the early phase after trauma see [Module I-3: Symptom](#)  
55 [Management](#)

56

57

57

58 **DISCUSSION**

---

59 **A. PSYCHOTHERAPY**60 **A1. Psychological Debriefing**

---

61 Psychological debriefing grew out of practices and experiences involving the military  
62 of the United States and other western nations. For soldiers exhibiting signs of acute  
63 stress reactions (ASR) following combat-related traumatic events, the practice of  
64 conducting early debriefings as part of a larger restoration approach, appeared to  
65 have significant impact on reducing more permanent disability.

66 The use of debriefings soon after exposure to traumatic events became part of  
67 military doctrine in the United States and elsewhere, as well as part of standards for  
68 early response to catastrophe for organizations such as the Red Cross.

69 Unfortunately, the technique appears to be of little help, and potentially harmful if  
70 used in individual victims of trauma as prophylaxis for PTSD.

71 In considering the use of debriefing procedures as part of early interventions  
72 following trauma exposure, a distinction between the general approaches of  
73 psychological versus operational debriefings is in order. Moreover, distinction should  
74 be made between debriefing procedures that are targeted at all exposed individuals,  
75 irrespective of symptom level, and by contrast, briefer versions of empirically  
76 supported brief psychotherapy interventions that are targeted at symptomatic  
77 individuals over a few sessions (see Annotation [A2: Brief CBT](#)).

78 **DEFINITIONS**

---

79 **Psychological Debriefing** is a broad umbrella term used to describe a variety of  
80 one-time individual and/or group procedures that involve review of a traumatic  
81 event, by survivors or other impacted persons, for the purpose of actively  
82 encouraging individuals to: (a) talk about their experiences during the event, (b)  
83 recognize and verbalize their thoughts, emotions and physical reactions during and  
84 since the event, and (c) learn about coping methods. Specially trained debriefers  
85 lead psychological debriefings following several protocols. Some protocols emphasize  
86 normalization of symptoms and group support. Some include psycho-education and  
87 information about resources.

88 The term "Psychological Debriefing" does not include purely informational briefings or  
89 debriefings used in professional military or other workgroups (e.g. psychological  
90 education lectures or stress management briefings such as Battlemind Training,  
91 Battlemind Debriefing, or operational debriefings) discussed below.

92 **Operational Debriefing** is a routine individual or team review of the details of an  
93 event from a factual perspective, for the purpose of: (a) learning what actually  
94 happened for the historical record or planning purposes; (b) improving results in  
95 similar future situations or missions; and (c) increasing the readiness of those being  
96 debriefed for further action. Operational debriefings are conducted by leaders or  
97 specialized debriefers according to the organization's standing operational procedure.

98 Although operational debriefings achieve important short-term objectives of the  
99 organization, there is insufficient evidence that they can also reduce subsequent  
100 PTSD or other long-term negative outcomes. Organizations that use operational

101 debriefings should train their debriefers to avoid causing unintentional psychological  
102 harm and to identify individuals who need behavioral health follow-up.

103 **Critical Incident Stress Debriefing (CISD)** is a formalized structured review  
104 method in a group format of the stressful experience of a disaster that includes  
105 psychological debriefing. In fact, CISD was developed to assist first responders such  
106 as fire and police personnel, not the victims/survivors of a disaster or their relatives.  
107 CISD was never intended as a substitute for therapy, was designed to be delivered in  
108 a group format with professional workgroups, and is meant to be incorporated into a  
109 larger, multi-component crisis intervention system labeled Critical Incident Stress  
110 Management (CISM).

111 **Critical Incident Stress Management (CISM)** incorporates several components,  
112 including pre-crisis intervention, disaster or large-scale incident demobilization and  
113 informational briefings, "town meetings," staff advisement, defusing, CISD, one-on-  
114 one crisis counseling or support, family crisis intervention, organizational  
115 consultation, and follow-up and referral mechanisms for assessment and treatment,  
116 if necessary.

117 **Battlemind Debriefing** is a recently developed intervention aimed specifically at  
118 professional military teams/workgroups (like CISD) and designed to reduce any  
119 potential iatrogenic effects of psychological debriefing noted in some studies;  
120 specifically, less emphasis is given to personal disclosure and review of index events  
121 (there is no requirement for individual disclosure, and the focus of the debriefing is  
122 more broadly on the transition from the entire deployment, rather than a single  
123 critical incident) and more emphasis is given to enhancement of peer support.  
124 Battlemind debriefing can be delivered in either small group or large group lecture  
125 formats.

### 126 ***Individual Debriefing***

127 Reviews and meta-analyses of studies of psychological debriefing as an early  
128 intervention to reduce or prevent PTSD symptoms in individuals have concluded that  
129 this technique is ineffective or potentially harmful (Rose, Bisson, & Wessely, 2002).  
130 Of note, two well-controlled studies with longer-term follow-up of individual patients  
131 have suggested that this intervention may be related to a poorer outcome compared  
132 to controls (Bisson, 1997; Mayou et al., 2000 which is a follow-up on Hobbs, 1996).  
133 Bisson et al. (2008) in a summary of the evidence in the ISTSS guideline (2009) also  
134 found no evidence to support the preventive value of individual debriefing delivered  
135 in a single session. Of the 10 studies that compared psychological debriefing with no  
136 interventions, 2 were positive, 5 were neutral and 3 had negative results. A meta-  
137 analysis conducted by Van Emmerick et al. (2002) included 7 studies and found that  
138 psychological debriefing interventions (non-CISD) and no intervention improved  
139 symptoms of post-traumatic stress disorder, but psychological debriefing did not  
140 improve symptoms. Cuijpers et al. (2005) assessed the results of studies examining  
141 the effect of prevention and found that the risk of posttraumatic stress disorder was  
142 somewhat increased after debriefing, but not significantly (RR=1.33), indicating a  
143 possible adverse effect. The RCTs to date cover only a limited variety of traumatic  
144 stressors, subject populations, and debriefing protocols. Most controlled studies  
145 have been of individually administered, one-time individual debriefings of victims of  
146 motor vehicle accidents or crimes such as rape. However, findings have been  
147 consistent across trials.

**Group Debriefing**

148  
149 The recommendation against conducting structured group debriefing is based  
150 primarily on the lack of effectiveness in studies; there does not appear to be any  
151 evidence of harm. In a partially randomized trial, Deahl et al. (2000) found no  
152 benefit of debriefing over assessment only in terms of PTSD symptoms; however,  
153 the group receiving debriefing evidenced lower alcohol misuse scores. The non-  
154 random assignment to groups weakens conclusions of this study (commanders blind  
155 to condition separated approximately 100 soldiers into two groups based on  
156 schedules and responsibilities; the groups were then randomly designated  
157 'debriefing' or 'control', thus outcomes are confounded by whatever factors were  
158 used for separating soldiers into groups by commanders). In another study by  
159 Campfield and Hills (2001), robbery victims were randomly assigned to immediate  
160 (less than 10 hours) or delayed (greater than 48 hours) CISD groups. Immediate  
161 CISD produced more pronounced reduction in symptoms, but no control group was  
162 employed and thus no conclusions regarding efficacy relative to no treatment can be  
163 made. This is particularly necessary with this intervention given that most people will  
164 recover spontaneously without any intervention and because of the potentially  
165 iatrogenic effects found in some studies of CISD with individuals. Other studies of  
166 group debriefing that have been conducted were of poor design, either, or both, in  
167 terms of low sample size and non-random assignment to group, and preclude  
168 conclusions regarding efficacy (Eid et al., 2001; Richards, 2001).

169 Three more recent RCT's with group debriefing have been reported. In an analogue  
170 study with students, Devilly et al. (2008) found no advantage of debriefing following  
171 a distressing video relative to a post-video snack. Adler et al. (2008) conducted a  
172 randomized trial of Critical Incident Stress Debriefing (CISD) of groups of soldiers  
173 deployed to a Kosovo peacekeeping mission. This trial randomised 1,050 soldiers  
174 from 19 platoons into 62 groups receiving one of three conditions: Debriefing (23  
175 groups), Stress Education (20 groups) and No Intervention (19 groups). The authors  
176 reported no differences between groups on all behavioural outcomes. In a second  
177 RCT by Adler et al. (2009) with returning Iraq soldiers, results indicated that  
178 compared to a Stress Education control condition, the Battlemind Debriefing had no  
179 overall effect on PTSD; moreover, within the subgroup of high combat-exposed  
180 soldiers, Battlemind Debriefing was no more effective than the Battlemind Training  
181 lecture (given in both small group and large group formats), with both treatments  
182 producing extremely minimal improvements in PTSD Check List (PCL) scores. A  
183 small but significant reduction in PTSD symptoms, depression symptoms, and sleep  
184 problems was observed for Battlemind Debriefing compared with standard stress  
185 education, although similar benefits were observed for the two other Battlemind  
186 training interventions. Thus, given the similar efficacy of the Battlemind Training  
187 lecture program, and the very small effect sizes observed, there is no reason to  
188 recommend Battlemind Debriefing or the Battlemind lecture program.

189 It remains possible that group interventions with pre-existing workgroups (teams,  
190 units, EMTs, co-workers) may assist with non-PTSD areas of improvement, such as  
191 group cohesion, morale, and other important variables, but this has not been  
192 demonstrated empirically. Similarly, group interventions may be useful for screening,  
193 education, and support. Trained personnel should lead these group interventions and  
194 if group approaches are used, group participation should be voluntary. Operational  
195 debriefings after traumatic events during on-going military operations also share  
196 these considerations, but they have other objectives that may override individual  
197 mental health protection. All operational debriefings should select protocols and  
198 train the debriefers to minimize psychological harm to the participants.



199 In conclusion, routine use of individual debriefing or the use of group psychological  
200 debriefing in the aftermath of trauma cannot be recommended in either military or  
201 civilian life. Of importance is the fact that other early treatment interventions have  
202 been found to prevent PTSD in symptomatic individuals (see Annotation B2). It  
203 appears appropriate to continue to focus resources on identifying and treating those  
204 with recognizable psychiatric disorders arising after trauma. The emphasis should be  
205 placed on the early detection of those at risk of developing psychopathology and those  
206 early interventions that have been found effective should be aimed at this group.

207 **EVIDENCE TABLE**

|   | Evidence  | Sources  | LE      | QE   | NET            | SR |
|---|---|--|---------|------|----------------|----|
| 1 | Individual psychological debriefing (or critical incident stress debriefing) is ineffective in preventing PTSD and may have adverse long-term effects | Rose, 2002 (Cochrane SR)<br>Van Emmerick et al., 2002<br>Cuijpers et al., 2005<br>Bisson et al., 2009 (ISTSS)<br>Sijbrandji et al., 2006<br>Hobbs et al., 1996<br>Mayou et al., 2000<br>Bisson, 1997 | I       | Good | Zero,<br>Small | D  |
| 3 | Group debriefing is ineffective in preventing PTSD  | Deahl et al., 2000<br>Campfield and Hills, 2001<br>Eid et al., 2001<br>Richards, 2001<br>Devilly et al., 2008<br>Adler et al., 2008<br>Adler et al., 2009  | I, II-1 | Fair | Zero,<br>Small | D  |

208 *LE =Level of Evidence; QE = Quality of Evidence; NET=Net benefit; SR = Strength of*  
209 *Recommendation (see Appendix A)*  
210

210 **A2. Brief Early Cognitive-Behavioral Intervention**

211 Research suggests that relatively brief but specialized interventions may effectively  
 212 prevent PTSD in some subgroups of trauma patients. Several controlled trials have  
 213 suggested that brief (i.e., 4 to 5 sessions) cognitive-behavioral treatments,  
 214 comprised of education, breathing training/relaxation, imaginal and in vivo exposure,  
 215 and cognitive restructuring, delivered within weeks of the traumatic event, can often  
 216 prevent PTSD in survivors of sexual and non-sexual assault (Foa et al., 1995) and  
 217 MVAs and industrial accidents (Bryant et al., 1998a , 1999). Brief intervention with  
 218 patients hospitalized for injury has been found to reduce alcohol consumption in  
 219 those with existing alcohol problems (Gentilello et al., 1999). Controlled trials of brief  
 220 early intervention services targeted at other important trauma sequelae (e.g.,  
 221 problems returning to work, depression, family problems, trauma recidivism, and  
 222 bereavement-related problems) remain to be conducted, but it is likely that targeted  
 223 interventions may be effective in these arenas for at least some survivors.

224 At present it is unknown how much time should elapse after a traumatic experience  
 225 before cognitive-behavioral intervention is initiated (Litz & Bryant: in Foa 2009  
 226 [ISTSS]). If provided too early, individuals who may not need therapy will consume  
 227 helping resources. For this reason, trials have not commenced before 2 weeks after  
 228 the trauma (Bryant, 1998, 1999, 2003).

229 **Target Population for Brief CBT**

230 Studies that have targeted all trauma survivors, regardless of levels of stress  
 231 reactions, have been ineffective in preventing PTSD (Roberts et al., 2009b).  
 232 Trauma-focused CBT has been found to be effective in reducing and preventing post-  
 233 traumatic stress symptoms in individuals who were symptomatic, especially those  
 234 meeting criteria for ASD (Roberts et al., 2009a; Stapleton, 2006). These  
 235 interventions have focused on the traumatic experience via exposure to memories  
 236 and trauma reminders, sometimes combined with cognitive therapy or other  
 237 behavioral interventions. One study has indicated that combined imaginal and in vivo  
 238 exposure is significantly more effective than pure cognitive restructuring in reducing  
 239 subsequent PTSD among individuals diagnosed with ASD (Bryant, et al., 2008). This  
 240 is an important finding that requires replication.

241 Cognitive behavioral therapy was more effective in reducing symptoms than a self-  
 242 help booklet or repeated assessment. The combination of an elevated initial  
 243 symptom score and failure to improve with self-monitoring was effective in  
 244 identifying a group of patients with early PTSD symptoms who were unlikely to  
 245 recover without intervention. (Ehlers, 2003)

246 **Evidence Table**

|   | Evidence   | Sources  | LE | QE   | SR |
|---|--|--|----|------|----|
| 1 | Brief cognitive-behavioral intervention (4 to 5 sessions) may prevent PTSD in those reporting clinically-significant symptoms of acute post-traumatic stress | Roberts 2009a (§)<br>Kornor, 2008<br>Bryant et al., 1998a, 1999<br>Bryant et al., 2003, 2008 | I  | Good | A  |
| 2 | Multisession early psychological interventions for asymptomatic trauma survivors are not effective and may be harmful.                                       | Roberts 2009b (§)  | I  | Good | D  |

247 *LE = Level of Evidence; QE = Quality of Evidence; SR = Strength of Recommendation; §-Systematic*  
 248 *Review (see Appendix A)*

249 **A3. Other Early Interventions**

250 Efficacious early interventions have largely been structured as brief versions of  
 251 effective PTSD treatments. This suggests that other interventions may be effective in  
 252 preventing PTSD but more research is needed to investigate other intervention  
 253 methods. Some non-CBT interventions have received research attention. For  
 254 example, brief structured writing has been found ineffective in preventing PTSD in  
 255 two studies (van Emmerik, et al., 2008; Bugg, et al., 2009). A memory restructuring  
 256 intervention failed to show preventive impact relative to a control condition (Gidron  
 257 et al., 2007). Likewise, providing self-help information, as a preventive  
 258 psychoeducation strategy to prevent PTSD has not been found to be efficacious  
 259 (Scholes et al., 2007; Turpin et al., 2005).

260  
261**Table I – 2 Early brief Psychotherapy Studies to Prevent the Development of PTSD**

| Author, Tear                          | Results  | n   | Trauma                   | LE | QE   | NB    |
|---------------------------------------|--|-----|--------------------------|----|------|-------|
| <b>Brief CBT</b>                      |  |     |                          |    |      |       |
| Bryant 1998a                          | Brief (5sessions) CBT w/I 2 weeks > Supportive counseling in preventing PTSD         | 24  | Civilian                 | I  | Fair | Mod   |
| Bryant 1999                           | Brief (5 session) PE or PE + Anxiety mgmt > Supportive counseling in preventing PTSD | 45  | Civilian with ASD        | I  | Good | Sub   |
| Bryant 2003                           | Brief (5sessions) CBT > Supportive Counseling in preventing PTSD                     | 24  | ASD after mTBI           | I  | Good | Sub   |
| Bryant 2008                           | Brief (5 sessions) ET > CT > No Tx in preventing PTSD                                | 90  | ASD civilians            | I  | Good | Sub   |
| Resnick 2007                          | Video intervention reduces PTSD vs. standard care                                    | 140 | Sexual assault           | I  | Good | Mod   |
| Rothbaum                              | 1 session ET   | 10  |                          |    |      | EXC   |
| <b>Self Help (SH)</b>                 |  |     |                          |    |      |       |
| Scholes 2007                          | no group differences between SH and no Tx  | 227 | Emergency Room           | I  | Good | Zero  |
| Turpin 2005                           | no group differences in PTSD between SH and no Tx                                    | 141 | N/R                      | I  | Fair | Zero  |
| <b>Structured Writing Therapy</b>     |  |     |                          |    |      |       |
| van Emmerik, 2008                     | Efficacy of SWT was comparable to CBT  | 125 | ASD and PTSD pts         | I  | Good | Mod   |
| Bugg, 2009                            | No differences between writing and self help (information only) groups               | 67  | Emergency room           | I  | Mod  | Small |
| <b>Memory Structured Intervention</b> |  |     |                          |    |      |       |
| Gidron 2007                           | No differences between MSI and supportive listening.                                 | 34  | traffic accident victims | I  | Fair | Zero  |

262

263

264

---

**B. Early Pharmacotherapy Interventions to Prevent Development of PTSD**

---

**Prevention of PTSD**

Few studies have examined the effectiveness of pharmacological treatment for acute symptom management and PTSD prevention immediately following a traumatic event. This includes the use of various agents for the prevention of PTSD (propranolol, hydrocortisone, and gabapentin) and the use of D-cycloserine, an N-methyl D-aspartate partial agonist, in combination with psychotherapy. Although of interest, none of these approaches is yet advocated in standard treatment guidelines for PTSD (Stein 2009 [SR]). There is insufficient evidence to draw concrete conclusions or make specific recommendations regarding the use of pharmacological agents for prevention of PTSD. While prevention of ASD is ideal, there are currently no evidence-based pharmacologic treatment modalities to arrest symptom formation and prevent progression to ASD during the first days and weeks following the traumatic exposure.

Once potential medical causes of neuropsychiatric impairment are ruled out and other immediate needs are met (e.g., physical needs, practical needs for assistance, normalization, and psychoeducation), then both medications and non-pharmacologic interventions may be considered. The selection and effectiveness of specific interventions administered acutely are not well supported in the literature. Although there are no evidence-based pharmacologic treatments for ASD, there may be a role for pharmacotherapy to aid in the management of specific symptoms (e.g., insomnia, pain, hyper arousal).

**Use of Benzodiazepines**

Historically, benzodiazepines were the primary agent in PTSD treatment, particularly alprazolam, and clonazepam. However, based on the limited data that are available, benzodiazepine administration should be used with caution (or discouraged) both in acute stress disorder (ASD) and post-traumatic stress disorder (PTSD), due to lack of evidence for effectiveness and risks that may outweigh potential benefits. Although benzodiazepines have been frequently used “as needed” and continuously for anxiety disorders, including to augment evidence-based treatment modalities in PTSD, there is theoretical, animal, and human evidence to suggest that benzodiazepines may actually *potentiate* the acquisition of fear responses and worsen recovery from trauma. Benzodiazepines use should be used especially cautiously in combat veterans with PTSD, because of the very high co-morbidity of combat-related PTSD with alcohol misuse and substance use disorders (upwards of 50% of comorbidity) and potential problems with tolerance and dependence. Once initiated in combat veterans, benzodiazepines can be very difficult, if not impossible, to discontinue, due to significant withdrawal symptoms compounded by the underlying PTSD symptoms.

Braun, et al. 1990, in a randomized double blind cross-over study, of alprazolam showed no significant benefit in alleviating PTSD symptoms compared with placebo. A slight reduction in anxiety symptoms was offset by withdrawal effects documented after only five weeks of treatment.

Gelpin, et al. (1996) in an open labeled study, treated 13 patients who had recently experienced trauma (within the past 18 days) and were experiencing excessive distress (panic, agitation, or persistent insomnia) for up to 6 months with alprazolam or clonazepam. These 13 patients were compared with a control group of recently traumatized individuals matched for demographics and symptoms (using the Impact of Events Scale). On follow-up, PTSD occurred at a significantly higher rate in the

312 benzodiazepine treated group (9/13, 69%) than in the control group (2/13, 15%).  
313 Although the strength of the evidence is low (open labeled study), the study  
314 suggested that benzodiazepines may worsen outcomes in the acute period following  
315 trauma, and the authors referenced animal data consistent with the hypothesis that  
316 benzodiazepines may potentiate the acquisition of fear responses.

317 Melman, et Al. 2002. In this small double-blind randomized controlled study during  
318 the acute period after trauma (mean 2 weeks after trauma), the short-term (7 day)  
319 evening use of temazepam in patients with significant ASD/PTSD symptoms was  
320 compared with placebo (11 patients in each group). The study showed no benefits in  
321 preventing PTSD, and the trend was similar to the Gelpin study, with 6 of 11 (55%)  
322 patients who received tamazepam developing PTSD, compared with 3/11 (27%) who  
323 received placebo.

324 Benzodiazepines can be effective against anxiety, insomnia and irritability, but they  
325 should be used with caution in patients with ASD and PTSD because of the high  
326 frequency of co-occurring substance abuse and dependence in patients with PTSD.  
327 The balance between benefit and potential risks, including the risks of dependency  
328 and of withdrawal after discontinuation, should be evaluated when considering  
329 benzodiazepines in patients with acute stress reaction.

### 330 ***Sleep Disturbance***

331 One of the most difficult symptoms to address in the immediate aftermath of  
332 exposure to a traumatic event is sleep disturbance. Theoretically, the more sleep  
333 impairment and trauma-related nightmares an individual continues to experience,  
334 the more likely he or she is to continue to experience the symptoms of ASD and/or  
335 subsequently develop PTSD. There is little evidence for the effectiveness of any  
336 sleep aids in the immediate aftermath of trauma.

337 For Recommendations and discussion of the evidence for sleep disturbance  
338 see [Module I-3: A. Sleep Disturbance](#)

### 339 ***Ineffectiveness of Propranolol***

340 Several studies have examined the use of propranolol, hydrocortisone, and  
341 gabapentin for the prevention of PTSD (Pittman, 2002; Stein, 2007).

342 Four small and brief clinical trials were identified in the peer-reviewed medical  
343 literature that evaluated the use of pharmacologic treatments to prevent the  
344 development of posttraumatic stress disorder (PTSD) symptoms in traumatized  
345 subjects (Pitman, 2002; Stein, 2007; Reist, 2001; Vaiva, 2003). All studies involved  
346 immediate posttraumatic administration of propranolol and 1 study also included a  
347 trial of gabapentin. Two of the studies (Reist, 2001; Vaiva, 2003) were excluded due  
348 to poor quality. Pitman (2002) reported a pilot study of 41 patients who were  
349 randomized to begin, within 6 hours of the event, a 10-day course of double-blind  
350 propranolol (n = 18) versus placebo (n = 23), 40 mg four times daily. Significant  
351 improvement of symptoms was noted in the treatment group. Stein (2007)  
352 conducted a double blind, randomized controlled trial of 14 days of the beta-blocker  
353 propranolol (n = 17), the anxiolytic anticonvulsant gabapentin (n = 14), or placebo  
354 (n = 17), administered within 48 hours of injury to patients admitted to a surgical  
355 trauma center. Of 569 accessible, potentially eligible subjects, 48 (8%) participated.  
356 Although well tolerated, neither study drug showed a significant benefit over placebo  
357 on depressive or posttraumatic stress symptoms.

358 McGhee et al., (2008) examined the relationship between PTSD prevalence and  
359 propranolol administration in 603 soldiers injured in OIF/OEF, of which 226

360 completed the PTSD Checklist-Military. Thirty-one soldiers received propranolol and  
361 34 matched soldiers did not. In propranolol patients, the prevalence of PTSD was  
362 32.3% vs 26.5% in those not receiving propranolol (P = .785). These data suggest  
363 propranolol does not decrease PTSD development in burned soldiers.

364 Although some positive results were noted, the size and weak study designs of the  
365 investigations do not allow for definitive conclusions regarding the value of these  
366 medications in preventing the development of PTSD symptoms after traumatic  
367 events.

### 368 **Other Medications**

369 One study that involved administration of cortisol at the time of cardiac bypass  
370 surgery (Schelling, 2004) suggested that patients who received stress doses of  
371 cortisol had lower PTSD symptom scores than a comparison group (that did not  
372 receive cortisol) when questioned six months after surgery.

373 A crossover trial of 1 month of low-dose cortisol therapy evaluated 3 patients  
374 diagnosed with PTSD (Aerni, 2004). The authors reported that each patient  
375 demonstrated improvement on at least 1 self-reported PTSD measure. The study  
376 was excluded from analysis for this guideline due to small numbers.

377 A single clinical investigation of the effect of the antibiotic D-cycloserine (Heresco-L,  
378 2002) enrolled 11 patients in a crossover trial. While patients reported some  
379 improvement on self-reported measures of PTSD symptoms, similar improvements  
380 were seen in placebo-treated patients.

### 381 **Conclusions:**

382 There is a small amount of evidence that suggests that administration of cortisol at  
383 the time of, or immediately after, a traumatic event may have a preventive effect on  
384 the subsequent development of PTSD symptoms. Little evidence exists suggesting  
385 that gabapentin or propranolol are of value in preventing the development of PTSD  
386 after trauma.

387 Due to the limited support of evidence, the use of medications in the early period  
388 post trauma to prevent PTSD cannot be recommended. Pharmacotherapy may be  
389 considered to aid in the management of specific symptoms (e.g., addressing sleep  
390 disturbance, irritability or control of pain).

### 391 **EVIDENCE TABLE**

|   | Evidence  | Sources   | LE   | QE   | Net Effect | SR |
|---|---|---|------|------|------------|----|
| 1 | Pharmacotherapy prophylaxis for PTSD  | Stein, 2006 [§]   | I    | Poor | -          | I  |
| 2 | Propranolol to reduce hyperarousal, excessive arousal, or panic attacks         | Pittman et al., 2002<br>Stein et al., 2007<br>Schelling., 2004  | I    | Fair | Small      | C  |
| 3 | Benzodiazepines for insomnia, hyperarousal, excessive arousal, or panic attacks | Braun et al., 1990<br>Cates et al., 2004<br>Gelpin et al., 1996 | II-2 | Fair | Small/Neg  | D  |

392 *LE =Level of Evidence; QE = Quality of Evidence; SR = Strength of Recommendation §=Systematic*  
393 *Review (see Appendix A)*  
394  
395

396

**Table I - 3 Pharmacologic Studies to Prevent the Development of PTSD**

| Author, Year         | Results  | n  | Trauma                 | LE | QE   | NB    |
|----------------------|--|----|------------------------|----|------|-------|
| <b>Propranolol</b>   |  |    |                        |    |      |       |
| Pitman, 2002         | Significant improvement post acute stress  | 41 | Any                    | I  | Good | Small |
| Stein, 2007          | No difference from placebo (gabapentin or propranolol)                                   | 48 | Severe physical injury | I  | Good | Zero  |
| Reist, 2001          | Recall of arousing story was reduced   | 38 | N/R                    | I  | Poor | EXC   |
| Vaiva, 2003          | PTSD rate and symptoms lower in the propranolol group                                    | 19 | MVA, assault           | I  | Poor | EXC   |
| <b>Cortisol</b>      |  |    |                        |    |      |       |
| Schelling, 2004      | Hydrocortisone administered during cardiac surgery reduced chronic stress symptom scores | 91 | Bypass surgery         |    | Fair | Mod   |
| Aerni, 2004          | Low dose cortisol for 1 month reduces the cardinal symptoms of PTSD                      | 3  |                        |    | Poor | EXC   |
| <b>D-cycloserine</b> |  |    |                        |    |      |       |
| Heresco-Levy, 2002   | Improvements in numbing, avoidance, and anxiety symptoms                                 | 11 | N/R                    |    | Fair | Zero  |

397

*N/R = no reported; EXC - Excluded; Mod = Moderate*

398

399

**I-2. TREATMENT FOR POST-TRAUMATIC STRESS DISORDER (PTSD)****A. Selection of Therapy for PTSD**

In clinical practice, providers and patients alike are often faced with important decisions relating to type, number, frequency, and dose of various psychotherapies and pharmacologic interventions. Therapies may be broadly divided into (1) evidence-based psychotherapies (particularly cognitive behavioral therapies), (2) evidence-based pharmacotherapies (particularly SSRIs and SNRIs), and (3) key adjunctive or supplemental treatment modalities.

Providers should explain to all patients with PTSD the range of therapeutic options that are available and effective for PTSD. This discussion should include general advantages and disadvantages (including side-effects/risks) associated with each therapeutic option. Contrary to clinical intuition, among the different evidence-based cognitive behavioral psychotherapy treatments, the evidence suggests that they are equivalent in their effectiveness (Rothbaum, 2001). In general, PTSD therapy research has provided insufficient evidence to favor medication or evidence-based psychotherapy as a first-line treatment. There is also insufficient evidence to suggest for or against combined medication and psychotherapy over only one of the two approaches. Patient preferences and the particular evidence-based treatments that the provider has the most training/expertise in will often drive the initial therapeutic approach.

It may be helpful to add therapies using a stepped care approach, even though supporting evidence is lacking. The use of stepped care approaches, which have been best studied in the context of collaborative care management based in primary care, has been advocated for many chronic conditions including hypertension, chronic fatigue, low back pain, depression, and post-war health concerns. In collaborative/stepped care, the intensity of care is augmented for patients who do not achieve an acceptable outcome with lower levels of care. In this approach there are care management techniques to facilitate coordination of care, monitoring, and follow-up; and there is judicious use of referrals supported through collaborative consultations. Stepped care is based on several assumptions: different people require different levels of care; finding the right level of care often depends on monitoring outcomes; moving from lower to higher levels of care based on patient outcomes often offers efficient increases in overall effectiveness; conditions often co-exist and involve more than one organ system; iatrogenic effects can occur from uncoordinated specialty care; unnecessary diagnostic tests, or use on non-evidence-based treatments; and patients often drop out or fail to follow-up with treatment as it becomes more complex.

The level or intensity of care is guided by illness trajectory (degree of chronicity and current illness severity), observed outcomes, and previously attempted therapies. Active follow-up is used to determine the level of care each patient requires over time. In PTSD for example, the patient and provider may determine that the first-line therapy will be psychotherapy. If, after a period of treatment, the patient is not responding adequately, the patient may be "stepped up" in therapeutic intensity by adding a medication, such as a selective serotonin reuptake inhibitor (SSRI) to the regimen of ongoing psychotherapy, and reassessing whether additional measures need to be taken to address co-morbid conditions.



47 **RECOMMENDATIONS**

- 48 1. Providers should explain to all patients with PTSD the range of available and  
49 effective therapeutic options for PTSD.
- 50 2. Patient education is recommended as an element of treatment of PTSD for all  
51 patients and the family members. [C]
- 52 3. Patient and provider preferences should drive the selection of evidence-based  
53 psychotherapy and/or evidence-based pharmacotherapy as the first line  
54 treatment. [Expert Consensus]
- 55 4. Psychotherapies should be provided by practitioners who have been trained in the  
56 particular method of treatment, whenever possible. [Expert Consensus]
- 57 5. A collaborative/stepped care approach to therapy administration may be  
58 considered, though supportive evidence is lacking. [Expert Consensus]
- 59 6. Specialized PTSD psychotherapies may be augmented by additional problem  
60 specific methods /services, and pharmacotherapy. [Expert Consensus]

62 **B. PSYCHOTHERAPY INTERVENTIONS FOR PTSD**

63 **Table I - 4 Intervention for Treatment of PTSD**

| SR | Significant Benefit  | Some Benefit  | Unknown   | No benefit or Harm |
|----|--|---|---|--------------------|
| A  | Cognitive Therapy [CT]<br>Exposure Therapy [ET]<br>Stress Inoculation Training [SIT]<br>Eye Movement Desensitization and Reprocessing [EMDR] |   |   |                    |
| B  |  | Imagery Rehearsal Therapy [IRT]<br>Psychodynamic Therapy<br>Hypnosis<br>Group therapy | WEB-Based CBT<br>Dialectical Behavioral Therapy [DBT] |                    |
| C  |  | Patient Education   |   |                    |
| I  |  | Family Therapy  |   |                    |

64 *SR\_ = level of recommendation*

65  
66 Significant difficulties reside in defining different CBTs and establishing their  
67 parameters. There are a number of reasons for this difficulty, including the diversity  
68 of treatments available and a lack of a common “language” to describe the same  
69 treatment components. The evidence-based psychotherapeutic interventions for  
70 PTSD that are supported by RCTs can be considered broadly as trauma-focused  
71 cognitive-behavioral therapy (CBT) interventions. CBT for PTSD refer to a range of  
72 psychological interventions based primarily on learning theory, cognitive theory,  
73 emotional processing theory, and conditioning models. Psychoeducation is another  
74 important component of all interventions. Non-trauma focused psychological  
75 treatments have not been shown to reduce PTSD symptoms as significantly as these  
76 four evidence-based CBT modalities. (ISTSS 2009, NICE 2005, Bisson 2009)

77 Trauma-focused CBT interventions include explicit review of traumatic experiences as  
78 part of treatment in some form. These include exposure techniques that involve  
79 repetitive review of traumatic memories and trauma-related situations and cognitive  
80 techniques that focus on identification and modification of trauma-related beliefs and  
81 meanings. Although stress inoculation training does not focus as explicitly on the  
82 exploration of traumatic memories or meanings, it is still considered a trauma-  
83 focused CBT because of its focus on assisting individuals with PTSD in reducing  
84 trauma-related avoidance and anxiety through relaxation and other techniques and  
85 because of how extensively it has been tested specifically for PTSD. Other CBT  
86 interventions that are not trauma focused are less effective.

87 In formulating the recommendations,, the working group evaluated the empirical  
88 evidence-base, considering randomized trials as the highest level of the evidence-  
89 based hierarchy. Therapy provided in clinical trial settings differs from therapy that  
90 is practiced in day-to-day care, and the recommendations represent the techniques  
91 and protocols as they were studied and reported in the RCTs.

92 Components of efficacious CBT interventions for PTSD have been packaged in  
93 research trials in various ways. Very few studies have dismantled these individual  
94 components to assess the relative efficacy of each independently. The majority of  
95 interventions included in RCTs have been grouped into four main categories based on  
96 the components most emphasized or the specific names used in the published  
97 literature:

- 98 • **Exposure-based techniques** (for example, prolonged exposure) emphasize in-  
99 vivo, imaginal, and narrative exposure, but also include elements of cognitive  
100 restructuring (e.g. evaluating the accuracy of beliefs about danger) as well as  
101 relaxation techniques and self-monitoring of anxiety.
- 102 • **Cognitive-based therapies** (for example, cognitive processing therapy),  
103 emphasize cognitive restructuring (challenging automatic or acquired beliefs  
104 connected to the traumatic event, such as beliefs about safety or trust) but also  
105 include relaxation techniques and discussion/narration of the traumatic event  
106 either orally and/or through writing.
- 107 • **Stress Inoculation Training** (the specific anxiety management package that  
108 has been most extensively studied in the literature), places more emphasize on  
109 breathing retraining and muscle relaxation, but also includes cognitive elements  
110 (self-dialogue, thought stopping, role playing) and, sometimes, exposure  
111 techniques (in-vivo exposure, narration of traumatic event).
- 112 • **EMDR** (extensively studied in a large number of RCTs as its own modality)  
113 resembles other CBT modalities in that there is an exposure component (e.g.  
114 talking about the traumatic event as well as holding distressing traumatic  
115 memories in mind, without verbalizing them) combined with a cognitive  
116 component (e.g., identifying a negative cognition, an alternative positive  
117 cognition, and assessing the validity of the cognition), and relaxation/self-  
118 monitoring techniques (e.g., breathing, "body scan"). Although alternating eye-  
119 movements are part of the classic EMDR technique (and the name of this type of  
120 treatment), comparable effect sizes have been achieved with or without eye  
121 movements or other forms of distraction or kinesthetic stimulation.  
122 Consequently, the mechanisms of effectiveness are likely to be similar to the  
123 other trauma-focused CBTs.

124 **RECOMMENDATIONS**

- 125 1. Psychotherapy interventions are aimed at reduction of symptoms severity and  
126 improvement of global functioning. However, the clinical relevance and  
127 importance of other outcome indicators (e.g., improvement of quality of life,  
128 physical & mental health) are not currently well known.
- 129 2. Providers should explain to all patients with PTSD the range of available and  
130 effective therapeutic options for PTSD.

131 **Treatment Options:**

- 132 3. Strongly recommend that patients who are diagnosed with PTSD should be  
133 offered one of the following evidence-based psychotherapeutic interventions  
134 (Cognitive Behavioral Therapy (CBT)). Selection will be based on patient  
135 preference, severity of the symptoms, the clinician expertise in one or more  
136 of these treatment methods, and available resources:
- 137 - Cognitive based Therapy [CT], [A]
  - 138 - Exposure based Therapy [ET] [A]
  - 139 - Stress Inoculation Training [SIT] [A]
  - 140 - Eye Movement Desensitization and Reprocessing [EMDR] [A]
- 141 4. Brief **Psychodynamic Therapy** may be considered for patients with PTSD.  
142 [B]
- 143 5. Imagery **Rehearsal Therapy [IRT]** can be considered for treatment of  
144 nightmares and sleep disruption. [B]
- 145 6. Dialectical **Behavioral Therapy (DBT)** can be considered for patients with a  
146 borderline personality disorder typified by parasuicidal behaviors. [B]
- 147 7. Hypnotic **Techniques** can be considered especially for symptoms associated  
148 with PTSD, such as pain, anxiety, dissociation and nightmares, for which  
149 hypnosis has been successfully used. [B]
- 150 8. There is insufficient evidence to recommend for or against **Family or**  
151 **Couples interventions** as first line treatment for PTSD; Family or Couples  
152 therapy may be considered in managing PTSD-related family disruption or  
153 conflict, increasing support, or improving communication.[I]
- 154 9. **Relaxation Therapy** (including biofeedback) is not recommended as single  
155 strategy for the treatment of PTSD. [D]
- 156 10. Relaxation **Techniques** should be considered as a component of CBT  
157 treatment approaches for ASD or PTSD in alleviating symptoms associated  
158 with physiological hyperreactivity. [C]
- 159 11. Group Therapy is recommended as useful treatment for patients with PTSD  
160 and may be considered to improve symptoms of PTSD [B]
- 161 • There is insufficient evidence to favor any particular type of group therapy  
162 over other types.
  - 163 • Group therapy is contraindicated for patients with active psychosis,  
164 cognitive deficits or at current suicidal or homicidal risk.
  - 165 • Patients being considered for group therapy should exhibit; acceptance for  
166 the rationale for trauma work, willingness to self-disclose, and no current  
167 life crisis.

- 168                   • Patients entering group therapy should sign an informed consent.
- 169           12. Consider augmenting with other effective evidence-based interventions for  
170           patients who do not respond to a single approach.
- 171           13. Supportive psychotherapy is not considered to be effective for the treatment  
172           of PTSD. However, it is more helpful than no treatment and may be helpful in  
173           preventing relapse in patients who have reasonable control over their  
174           symptoms and are not in severe and acute distress. For the patient with co-  
175           morbid disorders, supportive therapy may be all they can tolerate without  
176           causing additional harm.

177           Notes:

178           Psychotherapy interventions are aimed at reduction of symptoms severity and  
179           improvement of global functioning. However, the clinical relevance and  
180           importance of other outcome indicators (e.g., improvement of quality of life,  
181           physical & mental health) are not currently well known.

182           Psychodynamic, interpersonal, experiential (e.g., Gestalt therapy), and many  
183           other approaches may also be beneficial parts of an effectively integrated  
184           approach. Most experienced therapists integrate diverse therapies, which are not  
185           mutually exclusive in a fashion that is designed to be especially beneficial to a  
186           given patient.

187           **Delivery of care:**

- 188           14. Telemedicine interventions that have been shown efficacious may be  
189           considered as an alternative to standard mental health treatment for PTSD.
- 190           a. Telemedicine interventions are recommended when face-to-face  
191           interventions are not feasible due to geographic distance between patient  
192           and provider or other barriers to patient access (e.g. agoraphobia,  
193           physical disability); when the patient would benefit from more frequent  
194           contact than is feasible with face-to-face sessions; or when the patient  
195           declines more traditional mental health interventions.
- 196           b. Providers using telemedicine interventions should endeavor to maintain  
197           and strengthen the therapeutic relationship, build patient rapport, stress  
198           practice and assignment completion, and ensure adequacy of safety  
199           protocols using similar techniques as they do in a face-to-face session.
- 200           c. Providers using technology-assisted interventions should take steps to  
201           ensure that their work complies with the regulations and procedures of the  
202           organization in which they are employed, legal standards, and the ethical  
203           standards of their professions. Patient confidentiality and safety should be  
204           monitored closely.
- 205           15. Web-based interventions, that have been shown to be efficacious may be  
206           considered as a standalone intervention, or as an alternative to standard  
207           mental health treatment for PTSD
- 208           a. Clinicians should carefully review the content of any web-based materials  
209           to ensure their accuracy and ethical application before recommending use  
210           to patients.
- 211           b. Web-based interventions are recommended where face-to-face  
212           interventions are not feasible (e.g., geography limits access to other forms  
213           of treatment) or when patients decline more traditional mental health  
214           interventions.

- 215 c. Providers using web-based interventions should regularly encourage  
216 patients to complete the web-based intervention and endeavor to  
217 maintain and strengthen the therapeutic relationship, build patient  
218 rapport, stress practice and assignment completion, and ensure adequacy  
219 of safety protocols.
- 220 d. Providers using technology-assisted interventions should take steps to  
221 ensure that their work complies with the regulations and procedures of the  
222 organization in which they are employed, legal standards, and the ethical  
223 standards of their professions. Patient confidentiality and safety should be  
224 monitored closely.

## 225 DISCUSSION

### 226 **B1. Cognitive Therapy (CT)**

---

227 Cognitive therapy (CT) emerged principally from the work of Albert Ellis (1962) and  
228 Aaron Beck (1964). Initially manualized for the treatment of depression (Beck, Rush,  
229 Shaw, & Emery, 1979), CT has been successfully adapted to the treatment of a  
230 diverse set of psychiatric disorders, including PTSD (Freeman & Datillo, 1992;  
231 Freeman et al., 1989; Scott et al., 1989). The most studied protocols include CPT  
232 and CT as manualized by Bryant (ref).

233 Several randomized controlled trials (RCTs) demonstrate the efficacy of CT for a wide  
234 range of patients with PTSD, demonstrating its use in treating veterans with combat-  
235 related trauma, motor vehicle accident survivors, sexual or physical assault victims,  
236 and victims of natural disasters. Most RCTs have examined CT as delivered in an  
237 individual therapy format, though some studies have investigated group-delivered  
238 CT.

239 For purposes of this guideline, CT is defined as a treatment whose primary goal is to  
240 improve mood and behavior through a deliberate and explicit focus on modifying  
241 dysfunctional thoughts, beliefs, and expectations. In theory, while behavioral change  
242 is a desirable outcome of CT, the treatment components themselves do not explicitly  
243 or directly target behavioral patterns per se (however, it appears that even cognitive  
244 interventions may involve behavioral components such as exposure in that, for  
245 example, discussing the meaning of a traumatic event inevitably involves exposing  
246 oneself to the memories of that event). Likewise, while exposure-based interventions  
247 may result in altered cognitions, exposure therapies, per se, do not involve an  
248 explicit focus on cognitive restructuring procedures seen in CT. Nonetheless, in  
249 practice it is virtually impossible to conduct cognitive trauma-focused therapy  
250 without also involving behavioral or exposure based components, as it is similarly  
251 virtually impossible to conduct behavioral or exposure-based therapy without  
252 involving cognitive therapy components.

253 CT is accomplished through a systematic and prescriptive process of (a) identifying  
254 dysfunctional beliefs, (b) challenging and disputing these beliefs by examining the  
255 evidence for or against them, and (c) restructuring or replacing these beliefs with  
256 those that are more functional, logical, and reality-based. According to theories on  
257 which CT is based, traumatic events may lead to distorted beliefs regarding personal  
258 safety, self-efficacy, relative danger, future consequences of actions, and availability  
259 of support. Over time, these maladaptive beliefs lead to or maintain symptoms of  
260 PTSD and impair global functioning. The goal of CT for PTSD is to correct these  
261 beliefs, which causes a decrease in symptoms and improves functioning.

262 The CT treatment protocol for PTSD typically begins with an introduction of how  
263 thoughts affect emotions and behavior. The cognitive model of change is introduced  
264 and the patient is given a detailed rationale and expectations for participation in  
265 therapy are established. Treatment interventions are focused on identifying and  
266 clarifying patterns of thinking. Several active techniques are used, such as capturing  
267 and recording thoughts about significant events, weighing the evidence in support of  
268 these thoughts, challenging distressing trauma-related thoughts, and replacing  
269 dysfunctional thoughts with more adaptive ones. Through systematic assignments  
270 both during and between therapy sessions, dysfunctional thoughts are examined,  
271 challenged, and replaced. As thoughts become more logical and reality-based,  
272 symptoms decrease and global functioning improves. CT also emphasizes the  
273 identification and modification of distorted core beliefs about self, others, and the  
274 larger world. CT teaches that improved accuracy of thoughts and beliefs about self,  
275 others, and the world leads to improved mood and functioning.

## 276 DISCUSSION

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277 Randomized controlled trials (RCTs) have shown that CT alone is an effective  
278 intervention for patients with PTSD (Lovell et al., 1998; Marks et al., 2001). It is  
279 useful for identifying and modifying the many negative beliefs related to a traumatic  
280 experience. CT can be used effectively to reduce distressing trauma-related  
281 thoughts (e.g., about survival guilt, self-blame for causing the trauma, feelings of  
282 personal inadequacy, or worries about the future). Modifying thoughts about these  
283 and other trauma-related issues can reduce PTSD symptoms and improve mood and  
284 functioning. Numerous other trials support CT as a key component of combination  
285 treatments.

286 CT techniques are often delivered as part of treatment “packages” that can include  
287 exposure therapy, trauma-related education, and anxiety management. For  
288 example, Cognitive Processing Therapy, which has been manualized and validated  
289 for use with female sexual assault-related PTSD in women (Resick et al., 2002) and  
290 in veterans ( ), combines aspects of CT and exposure therapy. CT can also be  
291 delivered in conjunction with a range of other psychological therapies (e.g., EMDR  
292 and psychodynamic therapy). CT techniques may be an especially helpful treatment  
293 component when co-morbid depressive and/or anxiety disorders are present.

294 Significant difficulties reside in defining different CTs and establishing their  
295 parameters. There are a number of reasons for this difficulty, including the diversity  
296 of treatments available, a lack of a common “language” to describe the same  
297 treatment components, and the fact that a number of CTs have Behavioral Therapy  
298 components, such as imaginal and in-vivo exposure, embedded in their approach.  
299 This occurs for both explicit and implicit reasons. Explicitly, a number of treatments  
300 combine cognitive and exposure-based components and have patients engage in  
301 activities that have the dual goals of modifying maladaptive beliefs and facilitating  
302 behavioral habituation to the feared stimulus through repeated exposure to the  
303 trauma memory and reminders of the trauma memory. In effect, these treatments  
304 are combination treatments rather than purely cognitive therapies. Other treatments  
305 integrate exposure-based components in a more implicit, or accidental, manner. For  
306 example, having patients write extensively about their trauma, talk about it, or  
307 discuss it with others may have the result of habituation through repeated behavioral  
308 steps while the treatment itself claims to be purely cognitive. The effect of this  
309 blurring of the treatment components makes classification difficult and leads to a  
310 number of CTs that are best framed as combination or package treatments  
311 containing both cognitive and exposure therapy (ET) elements. In fact, while  
312 cognitive therapy and exposure therapy may be theoretically distinct forms of

313 treatment, in practice these interventions are often combined and most evidenced-  
314 based PTSD treatments combine elements of CT and ET.

315 Contraindications for CT have not been empirically established, but may include  
316 psychosis, severe brain damage, or severe intellectual impairment.

317 **Summary of Studies:**

318 Twenty-one relevant clinical trials that evaluated the use of CT for PTSD were  
319 analyzed. The trials investigated the effect of CT compared with no-treatment  
320 conditions, such as placement on a waiting list, and compared with other therapies.  
321 Both single-session therapy and long-term therapy were studied, with the longest  
322 therapy lasting 30 weeks, plus additional sessions after the end of formal treatment.  
323 Although therapists trained in standardized CT methods provided some treatments,  
324 the actual content of the therapy was often variable, as was the terminology used to  
325 describe it. In these studies, although the patients in the control group and study  
326 group generally improved over time, there was significantly greater improvement in  
327 most treated groups, compared with controls. The studies that enrolled participants  
328 from the general population of PTSD patients examined a primarily female  
329 population. There was a single clinical trial that enrolled male disaster workers.

330 Follow-up intervals ranged from immediate posttreatment to up to 2 years after  
331 completion of therapy. Patient retention rates were generally similar to those  
332 observed in studies of other types of therapy, but ranged from 52% to 100%. Few  
333 studies were blinded, and most relied on self-reported symptom questionnaires to  
334 provide data for analysis.

335 Nine relevant randomized clinical trials compared the effect of CT with that of a  
336 nonactive treatment, such as waitlist control group, treatment as usual (TAU), or  
337 repeated assessment (Beck, et al., 2009; Classen, Koopman, Nevill-Manning, &  
338 Spiegel, 2001; Difede, et al., 2007; Duffy, Gillespie, & Clark, 2007; Ehlers, et al.,  
339 2005; Foa, Zoellner, & Feeny, 2006; Monson, et al., 2006; Sijbrandij, et al., 2007;  
340 Smyth, Hockemeyer, & Tulloch, 2008). Both group and individual CT appeared  
341 effective in reducing PTSD symptoms. This was seen for brief, limited treatment  
342 models and for treatment programs taking several months to complete. Four studies  
343 compared the effect of CT with that of therapies described as support, supportive  
344 care, or Rogerian support therapy (Blanchard, et al., 2003; McDonagh et al., 2005;  
345 Foa, Zoellner, & Feeny, 2006; Cottraux, et al., 2008). In these trials, CT was  
346 reported to be superior to supportive care in reducing PTSD or in retaining patients  
347 in therapy. Notably, in the Cottraux study, there were more drop-outs from the  
348 Rogerian group due to worsening symptoms. Additionally, the CT group patients in  
349 this study demonstrated sustained improvements in PTSD symptoms at two years  
350 follow-up. Trauma-focused CT and present-focused CT were compared in a single  
351 study of Vietnam veterans (Schnurr, et al., 2003). Approximately 40% of all  
352 participants showed significant change in PTSD symptoms, but neither treatment  
353 was superior to the other.

354 The Trauma-Adaptive Recovery Group Education and Therapy (TARGET) model was  
355 studied in a trial that compared it with CT in the treatment of substance abuse  
356 patients (Frisman et al., 2008). Some improvement in PTSD symptoms was noted in  
357 both groups, but TARGET therapy was reported to produce greater improvement in  
358 sobriety self-efficacy. One clinical trial (van Emmerik, et al., 2008) compared CT with  
359 a structured writing therapy that included three components: (a) writing in the first  
360 person, (b) cognitive self-reappraisal of the writing, (c) farewell and sharing the  
361 writing. The authors reported improvement in both study groups, but detected no  
362 differences in efficacy between them.

363 Recently, researchers have attempted to dismantle treatments to examine their  
364 efficacious components. Bryant et al. (2003) reported that patients who received  
365 both CT and ET demonstrated less avoidance, depression, and catastrophic  
366 cognitions relative to patients who received ET only, while there was no difference in  
367 PTSD symptoms between the groups. In a later four arm study to try to determine  
368 what specific components of CBT were more effective, Bryant et al. (2008) compared  
369 cognitive restructuring (CR) alone, in-vivo exposure alone, in-vivo combined with  
370 imaginal exposure, and the combination of CR, in-vivo exposure, and imaginal  
371 exposure. The combined treatment was most effective; supporting the notion that  
372 effective therapy needs to include a combination of exposure and cognitive  
373 techniques. Resick et al. (2008) found no difference between patients assigned to  
374 receive Cognitive Processing Therapy (CPT) and patients assigned to receive only the  
375 cognitive component of CPT. Interestingly, a third group that received only written  
376 narrative exposure without any of the other CPT techniques performed nearly as well  
377 with no significant difference compared with full CPT or the cognitive component by  
378 the time of the 6 month follow-up. In an attempt to isolate the active ingredients,  
379 McDonagh et al. (2005) compared a treatment combining exposure and cognitive  
380 therapy elements to both a waitlist control group and a group given Present-  
381 Centered Therapy (PCT), a form of problem-solving therapy designed to eliminate  
382 the active ingredients found in CBT. Both treatment groups demonstrated improved  
383 symptoms over the waitlist control group but did not differ between themselves.

384 Bisson (2009) performed a systematic review of the randomized trials of all  
385 psychological treatments (Cochrane Collaboration Report). Treatment were  
386 categorized as Trauma-focused cognitive behavioural therapy/exposure therapy  
387 (TFCBT); stress management (SM); other therapies (supportive therapy, non-  
388 directive counselling, psychodynamic therapy and hypnotherapy); group cognitive  
389 behavioural therapy (group CBT); eye movement desensitization and reprocessing  
390 (EMDR). The reduction of clinician assessed PTSD symptoms measured immediately  
391 after treatment of TFCBT did significantly better than waitlist/usual care, SM and  
392 other therapies. Stress management did significantly better than waitlist/usual care,  
393 and there was no significant difference between other therapies and waitlist/usual  
394 care control. Group TFCBT was significantly better than waitlist/usual care. EMDR did  
395 significantly better than waitlist/usual care and other therapies. There was no  
396 significant difference between EMDR and TFCBT or SM.

397 There was evidence that individual TFCBT, EMDR, stress management and group  
398 TFCBT are effective in the treatment of PTSD. Other non-trauma focused  
399 psychological treatments did not reduce PTSD symptoms as significantly. There was  
400 some evidence that individual TFCBT and EMDR are superior to stress management  
401 in the treatment of PTSD at between 2 and 5 months following treatment, and also  
402 that TFCBT, EMDR and stress management were more effective than other therapies.  
403 There was insufficient evidence to determine whether psychological treatment is  
404 harmful. There was some evidence of greater dropout in active treatment groups.  
405 The considerable unexplained heterogeneity observed in these comparisons, and the  
406 potential impact of publication bias on these data, suggest the need for caution in  
407 interpreting the results of this review.

#### 408 **Conclusions:**

409 There is good evidence that individual CT is effective in reducing PTSD symptoms,  
410 and limited evidence that treatment gains persists for up to 2 years. Additional  
411 research is needed to demonstrate the efficacy of CT delivered in a group format.  
412 Given the contrasting outcomes of available studies comparing combinations and



413 dismantling components, there is insufficient data to conclude that CT is superior to  
414 ET at this time.

415 **EVIDENCE TABLE**

|   | Recommendation   | Sources  | LE  | QE   | SR |
|---|--|--|-----|------|----|
| 1 | CT is effective with civilian men and women exposed to combat and non-combat trauma. | Bryant et al., 2003<br>Bryant et al., 2008<br>Cottraux et al., 2008<br>Difede et al., 2007<br>Duffey et al., 2007<br>Ehlers et al., 2005<br>Foa et al., 2005<br>Lovell, et al., 2001<br>Marks et al., 1998<br>Sijbrandij et al., 2007<br>Smyth, Hockemeyer, & Tulloch, 2008<br>vanEmmerik, Kamphuis, & Emmelkamp, 2008 | I   | Good | A  |
| 3 | CT is effective in treating comorbid substance abuse and PTSD                        | Frisman et al., 2001   | I   | Poor | C  |
| 4 | CT is effective in treating PTSD in motor vehicle accident survivors                 | Blanchard et al., 2003   | I   | Mod  | B  |
| 5 | CT is effective in treating PTSD in a group format                                   | Beck et al., 2009  | III | Poor | I  |
| 2 | CT is effective with military and veterans with combat- and non-combat-related PTSD. | Monson et al., 2006  | III | Poor | I  |
| 3 | CT is effective for women with PTSD associated with sexual assault.                  | Chard, 2005<br>Classen et al., 2003<br>Foa et al., 2004<br>Foa, Zoellner, & Feeny, 2006<br>McDonagh et al., 2005<br>Resick et al., 2002<br>Resick et al., 2002, 2008   | I   | Good | A  |

416 *QE = Quality of Evidence; SR = Strength of Recommendation (see Appendix A)*

417 **B2. Exposure Therapy (ET)**

418 Exposure therapy protocols have a high level of evidence for treatment of PTSD, and  
419 generally include the components of psychoeducation, imaginal or narrative  
420 exposure, in-vivo exposure, and processing of thoughts and emotions. The most  
421 commonly used protocol is Prolonged Exposure (PE), although various other  
422 exposure protocols have been used. Protocols that provide only a portion of these  
423 components (e.g. imagery rehearsal therapy, in-vivo exposure or imaginal exposure  
424 alone), show less robust effect sizes. Imaginal exposure involves encouraging the  
425 patient to revisit the experience in imagination, and recalling the experience through  
426 verbally describing the physical and emotional details of the trauma. In vivo  
427 exposure involves asking the patient to physically confront realistically safe but still  
428 feared stimuli (e.g. driving a car after having been in a serious motor vehicle  
429 accident). In vivo exposure is typically arranged in a hierarchical order based on  
430 perceived difficulty of confronting each stimulus. In addition, each item on the

431 hierarchy may be titrated to make it more or less difficult depending on the patient's  
432 progress in treatment. In the preceding example the patient might first sit in a car  
433 in the passenger seat, and then in the driver's seat, and then start the car, etc. The  
434 patient repeats each situation until a reduction in the intensity of emotional and  
435 physiological response is achieved, at which point they move on to the next item in  
436 their hierarchy.

#### 437 DISCUSSION

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438 RCTs have shown that Exposure Therapy (ET) helps men and women with PTSD  
439 symptoms. RCTs of ET have demonstrated its efficacy in female victims of sexual  
440 and non-sexual assault, motor vehicle accidents, male combat-related trauma, war  
441 refugees, and mixed trauma populations. In randomized trials comparing ET with  
442 other cognitive behavioral treatments, ET has performed as well or better than any  
443 other cognitive behavioral therapy (CBT) approach. Findings regarding efficacy in  
444 (mostly Vietnam) combat veterans in VA clinical settings are less consistent and the  
445 degree of improvement in PTSD symptoms may be less pronounced though  
446 preliminary data suggest it is efficacious (Rauch et al., 2009).

447 The mechanism of ET is thought to be related to reduction in negative emotions  
448 (fear, anxiety, sadness, guilt) associated with their experience through repetitive,  
449 therapist-guided confrontation of feared places, situations, memories, thoughts, and  
450 feelings. ET usually lasts from 8 to 15 sessions depending on the trauma and  
451 treatment protocol. In the most common form of ET, patients are repeatedly  
452 exposed to their own individualized trauma stimuli, until their arousal and emotional  
453 responses are consistently diminished. ET providers can vary the pacing and  
454 intensity of exposing patients to the most difficult details of their trauma based on  
455 the patient's emotional response to the trauma and to the therapy itself.

456 Several studies indicate that results are comparable between exposure with other  
457 forms of cognitive behavioral therapy (e.g., cognitive therapy, EMDR, stress  
458 inoculation training, or combinations). Variations on exposure therapy that have  
459 promising results include written exposure and exposure in the context of a broader  
460 narration of the patient's life. For example, in a three arm dismantling RCT by  
461 Resick, et al. (2008), written exposure was compared directly with the CPT program  
462 (without the written exposure component), and the full CPT program (including  
463 written exposure). Treatment sessions for the written exposure only group consisted  
464 of two one-hour sessions to provide overview of treatment and education, followed  
465 by five two-hour sessions where the patient was asked to write for approximately 60  
466 minutes alone about their worst traumatic event, followed by reading this to the  
467 therapist who provided supportive feedback without any of the cognitive  
468 restructuring techniques. The written exposure group did nearly as well as both of  
469 the CPT treatment arms (which consisted of 12 one-hour sessions), and on the six  
470 month follow-up there was no significant difference between the three groups. This  
471 supports the notion that a systematic writing narrative process can be highly  
472 effective in alleviating symptoms.

473 Oral narrative therapy has also been shown to be highly effective in treating PTSD in  
474 war-ravaged refugee populations. In one study of Rwandan refugees with PTSD and  
475 severe war-related trauma (Neuner, 2008), lay counselors had patients construct a  
476 narration of their life from birth to the present while focusing on detailed exploration  
477 of specific traumatic experiences. This resulted in significant improvement in PTSD  
478 symptoms, with effects comparable to any of the most cited CPT or PE studies in  
479 U.S. or European clinical samples. Increasingly, virtual (computer based) exposure  
480 techniques and strategies are being utilized to accomplish exposure therapy.

481 However, to date there are no randomized studies of virtual reality compared with  
482 either wait list or standard exposure techniques that confirm its efficacy.

483 There have, as yet, been no randomized trials comparing ET with pharmacotherapy,  
484 either alone or in conjunction with one another. However, two trials have examined  
485 augmentation strategies. In one trial, the addition of ET following 10 weeks of  
486 sertraline resulted in reduction in relapse and additional symptom reduction in those  
487 patients who either failed to initially respond or partially responded to sertraline  
488 (Rothbaum, et al., 2006). In a second study, augmentation with paroxetine for  
489 patients who partially responded to 6 sessions of ET did not result in additional  
490 benefit (Simon, et al, 2007).

491 As with any treatment, patients need to be screened for their suitability prior to  
492 undergoing ET as it may temporarily increase their level of distress. Patients living  
493 with the threat of domestic violence should not be considered for ET until their  
494 security can be assured. ET has not been studied and providers should therefore use  
495 caution when working with patients with health problems that preclude exposure to  
496 intense physiological arousal, current significant suicide risk, substance dependence,  
497 or current psychosis. Providers should be aware of the possibility of increased  
498 distress as patients confront trauma memories and reminders. As in all PTSD  
499 treatment, providers must take concrete steps to prepare patients for the treatment  
500 (e.g., present clear rationale, explore patient concerns, encourage realistic  
501 expectations, and build commitment to the therapy) in order to reduce the risk of  
502 dropout.

503

**EVIDENCE TABLE**

|   | Evidence   | Sources  | LE | QE   | SR |
|---|--|--|----|------|----|
| 1 | ET is effective in the treatment of PTSD (compared to waiting list, present centered therapy, and other control comparisons) | Basoglu 2005, 2007<br>Cloitre 2002<br>Cooper et al., 1989<br>Feske, 2008<br>Foa et al., 1991 & 1999a<br>Ironson et al., 2002<br>Keane et al., 1989<br>Marks et al., 1998<br>McDonah, 2005<br>Neuner 2004, 2008 (life narration),<br>Schnurr 2007<br>Tarrier et al., 1999   | I  | Good | A  |
| 2 | ET compared to other forms of therapy show equivalent results.   | Bryant, 2003<br>Bryant, 2008<br>Foa et al., 1991 & 1999a<br>Foa, 2005<br>Marks et al., 1998<br>Paunovic & Ost, 2001<br>Power, 2002<br>Resick ,2002<br>Resick & Nishith, 2001<br>Resick, 2008 (written exposure)<br>Rothbaum, 2005<br>Schnurr, 2001<br>Tarrier et al., 1999 | I  | Good | A  |

504

QE = Quality of Evidence; R = Recommendation (see Appendix A)

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**B3. Stress Inoculation Training (SIT)**

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507 Several therapy protocols have been developed that focus on anxiety management  
508 and coping skills training, including Stress Inoculation Training and Relaxation  
509 Training. Stress inoculation training (SIT), is presented as a tool box or set of skills  
510 for managing anxiety and stress (Hembree & Foa, 2000). This treatment was  
511 developed for the management of anxiety symptoms and adapted for treating  
512 women rape trauma survivors. SIT typically consists of education and training of  
513 coping skills, including deep muscle relaxation training, breathing control,  
514 assertiveness, role playing, covert modeling, thought stopping, positive thinking and  
515 self-talk, and in-vivo exposure. The rationale for this treatment is that trauma  
516 related anxiety can generalize to many situations (Rothbaum et al., 2000). The  
517 Expert Consensus Guideline Series: Treatment of Post-traumatic Stress Disorder  
518 notes that anxiety management is among the most useful psychotherapeutic  
519 treatments for patients (Foa et al., 1999b). Relaxation protocols have also  
520 demonstrated encouraging results (Marks, et al., 1998; Taylor, et al., 2003; Vaughn,  
521 et al., 1994).

522 SIT is designed to “inoculate” people with PTSD from heightened stress responses  
523 through teaching anxiety management skills which can include:

- 524 • Relaxation training: teaching patients to control fear and anxiety through the  
525 systematic relaxation of the major muscle groups
- 526 • Breathing retraining: teaching slow, abdominal breathing to help the patient relax  
527 and/or avoid hyperventilation with its unpleasant and often frightening physical  
528 sensations
- 529 • Positive thinking and self-talk: Teaching the person how to replace negative  
530 thoughts (e.g., ‘I’m going to lose control’) with positive thoughts (e.g., ‘I did it  
531 before and I can do it again’) when anticipating or confronting stressors. This is  
532 often combined with in-vivo exposure.
- 533 • Assertiveness training: teaching the person how to express wishes, opinions, and  
534 emotions appropriately and without alienating others
- 535 • Thought stopping: distraction techniques to overcome distressing thoughts by  
536 inwardly ‘shouting stop’ (Foa et al., 1999b).

537 Many SIT protocols also include cognitive restructuring and elements of exposure  
538 therapy.

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**DISCUSSION**

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540 There have been two RCTs that have evaluated SIT and both studies found SIT to be  
541 effective with women who have survived sexual assault. A study by Foa and  
542 colleagues (1991) with 45 female sexual assault victims compared SIT, Prolonged  
543 Exposure (PE) (see Annotation B2), Supportive Counseling (SC) and wait list control.  
544 SIT was found to be the most effective treatment for short term symptom  
545 improvement and both SIT and PE were effective for long term improvement with PE  
546 superior to SIT. Rothbaum, (2001) reports, “results suggested that all conditions  
547 produced improvement on all measures immediately post-treatment and at follow-  
548 up. At follow-up, clients who received PE continued to improve after treatment  
549 termination, whereas clients in the SIT and SC conditions evidenced no change  
550 between post-treatment and follow-up.” Another study with 96 female sexual  
551 assault victims compared SIT, PE, combined SIT and PE, and wait list controls (Foa  
552 et al., 1999a). The study found all treatments were better than wait list control for

553 ameliorating PTSD severity at post-treatment and at 6-month follow-up.  
 554 Interestingly, although all three treatments were effective, the combined treatment  
 555 was not superior to either SIT or PE alone. Although this may be related to the fact  
 556 that clients in the combined treatment group received less PE-or SIT-specific  
 557 techniques than participants in the individual treatments, the most likely explanation  
 558 presented in the paper was an unusually low drop out rate that happened to occur in  
 559 the PE-only group.

560 A study of 15 women by Kilpatrick et al. (1982) found SIT to be effective in reducing  
 561 rape related fear and anxiety.

562 Motor vehicle accident survivors (Hickling & Blanchard, 1997) had a 68 percent  
 563 reduction of PTSD symptoms after involvement in a modified version of Foa et al.'s  
 564 SIT/PE combination program.

565 A controlled study comparing three different forms of relaxation (relaxation,  
 566 relaxation plus deep breathing, and relaxation plus deep breathing plus biofeedback)  
 567 for 90 Vietnam veterans found that all treatments were equally, but only mildly,  
 568 effective in leading to improvement (Watson et al., 1997).

569 Vaughn, et al. (1994) found that relaxation training was superior to waitlist. Taylor  
 570 et al (2003) also found support for reduction of PTSD with a relaxation protocol  
 571 though effects were less than for ET. In a head-to-head comparison study by Marx  
 572 (1998), relaxation training produced nearly the same beneficial effect as PE.

573 **EVIDENCE TABLE**

|   | Evidence   | Sources  | LE | OE   | SR |
|---|--|--|----|------|----|
| 1 | SIT is effective as a treatment for PTSD particularly related to sexual assault. | Foa et al., 1999a<br>Foa et al., 1991<br>Kilpatrick et al, 1982<br>Rothbaum, 2000a | I  | Good | A  |

574 *LE=Level of Evidence QE=Quality of Evidence; SR= Strength of Recommendation (see Appendix A)*

575 **B4. Eye Movement Desensitization and Reprocessing (EMDR)**

576 Eye Movement Desensitization and Reprocessing (EMDR) is a psychological  
 577 treatment designed to alleviate the distress associated with traumatic memories  
 578 (Shapiro, 1989a, and 1989b). The objective of EMDR is to assist patients to  
 579 accessing and process traumatic memories while bringing these to an adaptive  
 580 resolution (Shapiro, 2001).

581 In EMDR, the therapist collaborates with patients to: (1) access a disturbing image  
 582 associated with the traumatic event; (2) solicit the experience of body sensations  
 583 associated with the disturbing image; (3) identify an aversive self-referring cognition  
 584 (in concise words) that expresses what the patient "learned" from the trauma; (4)  
 585 identify an alternative positive self-referring cognition that the patient wishes could  
 586 replace the negative cognition. The patient is then asked to hold the disturbing  
 587 image, sensations, and the negative cognition in mind while tracking the clinician's  
 588 moving finger back and forth in front of his or her visual field for about 20 seconds.  
 589 In successive tracking episodes, the patient concentrates on whatever changes or  
 590 new associations have occurred. Eye movement episodes are repeated until there  
 591 are no new associations. Subsequent tracking episodes attempt to replace the  
 592 negative cognitive self-statement with the alternate positive cognition.

593 Between sessions, the patient is directed to keep a journal of any situations that  
 594 provoke PTSD symptoms and of any new insights or dreams about the trauma. The

595 number of sessions is dependent upon observed improvements and the number of  
596 traumatic events experienced.

597 Within a session, standard self-rating scales document changes in the intensity of the  
598 symptoms and the negative cognition, and the patient's acceptance of the alternative  
599 positive cognition. The patient reports following each set of eye movement episode  
600 to inform the therapist of the strength of both negative and positive cognitions;  
601 changes in cognitions, the images, emotions, or body sensations are also reported.

602 EMDR protocols allow for substitution of left-right alternating tone or touch as  
603 modifications to the use of the eye movements, suggesting that it is not the eye  
604 movements per se, but rather side to side alternating stimulation that is sought.  
605 Studies attempting to ascertain the relative contribution of the eye-movement  
606 component suggest that comparable outcomes are attained with or without eye  
607 movements. These findings are seen as indicating that this aspect (i.e., eye-  
608 movements or alternating stimulation of any type) of the treatment protocol may not  
609 be critical components.

610 Given the success of EMDR and the lack of support for the alternating stimulation  
611 components, many theorists are considering the active ingredients for the observed  
612 treatment gains. Specifically, EMDR is gaining acceptance as a treatment that shares  
613 components with other existing, successful treatments. Derived from desensitization  
614 strategies, EMDR counters avoidance of the traumatic memories and related cues by  
615 repeatedly accessing the aversive traumatic images themselves, promotes emotional  
616 processing by soliciting the emotional responses attendant to the aversive  
617 memories; identifies a novel and alternative view of the traumatic experience in  
618 conjunction with the patient, and then challenges the patient to consider the validity  
619 or accuracy of the alternative perspective. With the focus upon physiological arousal  
620 and reactivity, EMDR as a desensitization treatment also provides a component of  
621 arousal management that is inherent in the treatment. Thus, EMDR at its most basic  
622 level incorporates components of a) exposure to trauma related cues; and b)  
623 processing of emotional responses. Each of these EMDR components involves efforts  
624 to mitigate strategic avoidance reactions theoretically viewed as maintaining current  
625 symptomatology. EMDR also includes: c) elements of corrective and rational  
626 restructuring of the patient's views of the traumatic event; and d) self monitoring of  
627 cognitive and emotional responses that are often viewed as key homework  
628 components of cognitive behavior therapy in general, and e) a focus on heightened  
629 physiological arousal and reactivity.

## 630 DISCUSSION

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631 EMDR possesses efficacy for treating patients with PTSD: this conclusion is based  
632 upon a thorough review of the literature in the treatment guidelines generated by a  
633 task force for the International Society for Traumatic Stress Studies (Spates et al.,  
634 2009) as well as by Division 12 of the American Psychological Association (APA).  
635 The United Kingdom's NICE Guidelines for PTSD (2005) also recommend EMDR as a  
636 treatment supported by multiple efficacy studies. While the results of numerous  
637 controlled published studies found medium to large effect sizes for EMDR, the claims  
638 for EMDR as a modality that is more efficient or rapid than other forms of cognitive  
639 behavioral treatment remains unconfirmed across the studies that specifically  
640 address the issue. Similarly, suggestions that EMDR is more easily tolerated than  
641 other psychological treatments remain unsupported empirically.

642 Results of meta-analytic studies, review articles, and extant practice guidelines  
643 suggest that EMDR successfully treats symptoms of PTSD when compared to no  
644 treatment or delayed treatment conditions. When compared to other treatment

645 modalities, most studies reviewed indicated that EMDR possessed comparable  
646 efficacy to other well accepted cognitive behavioral treatments to include: stress  
647 inoculation training (SIT) and exposure therapies.

648 Maxfield and Hyer (2002) conducted a meta-analysis involving comparisons of EMDR  
649 against wait list controls, cognitive behavior therapy involving exposure, and  
650 treatment modalities described as other than CBT. Results indicated superiority of  
651 EMDR to the wait list control condition. Also, the authors found an overall superiority  
652 of EMDR compared to the other active treatment conditions, though they noted  
653 sufficient variability that they judged the summed results to indicate comparable vs.  
654 superior effectiveness of EMDR over other treatments.

655 Four studies specifically compared EMDR with Exposure Therapy. (Lee et al., 2002;  
656 Power et al., 2002; Rothbaum, et al., 2005; and Taylor et al., 2002). Lee et al.  
657 (2002) and Power et al. (2002) found that EMDR had equivalent or better results  
658 than CBT and was more efficient in that it worked faster. Taylor et al., (2002) didn't  
659 observe differential efficiency in their trial, but they also used therapist-assisted *in*  
660 *vivo* work plus imaginal work. Rothbaum, et al., (2005) found symptom  
661 improvement at post-test to be equivalent between EMDR and prolonged exposure.  
662 She writes in the abstract, "PE and EMDR

663 did not differ significantly for change from baseline to either posttreatment or 6-  
664 month follow-up measurement on any quantitative scale." Although a measure  
665 termed "end-state functioning" was described as favoring PE, this was a composite  
666 variable derived from three validated scales, that seemed to magnify the small non-  
667 significant differences on these individual scales when they were combined.

668 Criticisms of EMDR stem from its theoretical premises to the necessity of its  
669 components to achieve the desired outcome. With respect to componential analyses,  
670 the data are substantive at this time. Spates et al. (2009) reviews aptly the  
671 literature on dismantling studies in EMDR and concludes that "the best provisional  
672 conclusion so far is that the bilateral stimulation component of EMDR does not  
673 incrementally influence treatment outcome". Notwithstanding the lack of necessity  
674 for eye movements, when viewed within the framework of all other trauma-focused  
675 CBTs, EMDR is equivalent.

676 Some support for the inclusion of therapeutic eye movement is provided by a set of  
677 seven studies but most of these are studies with analog populations, or in clinical  
678 populations exposed to a traumatic event, but who didn't necessarily develop full  
679 clinical PTSD (Andrade et al., 1997; Barrowcliff et al., 2004; Christman and Garvey,  
680 2000; Kavanaugh et al., 2001; Kuiken et al., 2001-2002; Sharpley et al., 1996;  
681 Wilson, Silver, Covi, & Foster, 1996; and van den Hout et al., 2001).

682 There may be some basis for or against recommending this treatment depending  
683 upon the type of trauma leading to PTSD. Specifically, studies of EMDR efficacy with  
684 combat veterans have demonstrated considerable variability, with several authors  
685 suggesting that the treatment may be less than optimal for this condition  
686 (Boudewyns et al., 1993; Jensen, 1994). Other studies that are more recent have  
687 suggested the opposite (Carlson et al., 1998; Devilly et al., 1998). It should be  
688 noted that only two of the cited studies had a full course of treatment – all the others  
689 were short duration studies. However, it should be also noted that studies of other  
690 CBT modalities and SSRIs have also shown inconsistent results in combat veterans,  
691 and thus, based on current evidence, there is no reason to believe that EMDR would  
692 not be as effective as other trauma-focused CBTs in this populations.

693 Overall, argument can reasonably be made that there are rigorously controlled  
694 studies to support the conclusion that EMDR is effective in the treatment for PTSD.

695 **EVIDENCE TABLE**

|    | Recommendation   | Sources   | LE | QE   | R        |
|----|--|---|----|------|----------|
| 1  | EMDR is more efficacious for PTSD than wait-list, routine care, and active treatment controls. | Chemtob et al., 2000<br>Davidson & Parker, 2001<br>Maxfield & Hyer, 2002<br>Sheppard et al., 2000<br>Van der Kolk et al. 2007   | I  | Good | A        |
| 2  | Eye movements are <i>not</i> critical to the effects of EMDR.                                  | Davidson & Parker, 2001<br>Spates et al., 2009  | I  | Poor | C        |
| 3. | EMDR compared with ET show consistent comparable results.                                      | Cahill, 2000<br>Davidson & Parker, 2001<br>Foa & Meadows, 1997<br>Ironson et al., 2002<br>Lee et al., 2002<br>Power et al., 2002<br>Rothbaum et al., 2005<br>Servan-Schrieber, 2000<br>Sheppard et al., 2000<br>Taylor et al., 2002<br>Van Etten and Taylor, 1998 | I  | Fair | <b>B</b> |

696 LE=Level of Evidence QE=Quality of Evidence; SR= Strength of Recommendation (see Appendix A)

697 **B5. Imagery Rehearsal Therapy (IRT)**

698 Occurrence of nightmares as a problem is frequent; 4 to 8 percent in the general  
699 population and 60 percent in PTSD. Evidence shows that nightmares are associated  
700 with psychological distress and sleep impairment. A conditioning pattern similar to  
701 classic psychophysiological insomnia is produced in the nightmare disturbed loop,  
702 along with the negative cognition of "fear of going to sleep." Studies using brief CBT  
703 (desensitization and imagery rehearsal) have demonstrated a large reduction in  
704 nightmares. Many studies, including Forbes et al. (2001) suggest that PTSD is  
705 associated with a propensity toward image, particularly where the post-traumatic  
706 symptom picture is characterized by nightmares and flashbacks. IRT incorporates a  
707 system to increase the imagery control.

708 IRT is aimed at changing the content of the patient's nightmares to promote mastery  
709 over the content-threat, thereby altering the meaning, importance, and orientation  
710 to the nightmare. IRT includes elements of 1) psychoeducation about nightmares,  
711 insomnia, and PTSD, 2) positive coping skill building (thought stopping, breathing,  
712 grounding, writing/talking about issues and others), 3) cognitive restructuring, 4)  
713 sleep hygiene, stimulus control, and sleep restriction, 5) and focused use of pleasant  
714 imagery to replace negative imagery in recurrent nightmares. While discussion of  
715 trauma imagery occurs, the model includes a de-emphasis of discussion of this  
716 content in group sessions. The model has been tested primarily in a group format.

717 **DISCUSSION**

718 Several studies have examined IRT with some promising results. While not with a  
719 primary PTSD population, Krakow et al. (1995) studied 58 chronic nightmare  
720 sufferers who were randomly assigned to a treatment group (n = 39) or a wait-list  
721 control group (n = 19). The IRT group demonstrated significant reductions in



722 nightmares and improved sleep quality. Further, reduction in nightmares was a  
723 significant predictor of improvement in sleep. The authors concluded that, for some  
724 chronic sufferers, nightmares may be conceptualized as a primary sleep disorder  
725 which can be effectively and inexpensively treated with CBT.

726 Krakow et al., (2001) randomly assigned 168 female survivors of sexual assault  
727 (95% of the sample met the criteria for PTSD) to receive IRT (n = 88) or wait-list (n  
728 = 80) and found that among completers those women assigned to IRT had a larger  
729 reduction in self reported PTSD severity at 3 month follow up than wait-list. Further,  
730 the impact of nightmares was reduced and sleep quality improved. In a pilot study of  
731 IRT with crime survivors with PTSD, Krakow et al., (2001a) reported significant  
732 reductions in nightmares, improved sleep, and reduced PTSD severity at 3-month  
733 follow-up.

734 Forbes et al. (2001) completed an open trial of group IRT with 12 Vietnam Veterans  
735 with combat-related nightmares and PTSD. Veterans reported significant reduction  
736 in nightmare frequency and intensity for the target nightmare. In addition, self  
737 reported PTSD symptoms were significantly reduced. Follow up data demonstrated  
738 maintenance of gains at 12 months following the conclusion of treatment (Forbes et  
739 al, 2003).

740 While much of the research to date has focused on IRT other versions of nightmare  
741 reduction programs, such as Emotional Relaxation and Rescripting are currently  
742 under empirical examination.

#### 743 EVIDENCE TABLE

|   | Recommendation  | Sources   | LE       | QE   | R |
|---|---|---|----------|------|---|
| 1 | IRT can be considered for treatment of PTSD (nightmare and sleep disruption in particular). | Krakow et al., 1995; 2001a; 2001b<br>Forbes et al, 2001, 2003 | I<br>I-1 | Fair | B |

744 LE=Level of Evidence QE=Quality of Evidence; SR= Strength of Recommendation (see Appendix A)

## 745 B6. Psychodynamic Therapy

### 746 BACKGROUND

747 In 1895, Joseph Breuer and Sigmund Freud based their *Studies on Hysteria* on the  
748 proposition that traumatic life events can cause mental disorder (Breuer & Freud,  
749 1955). This principle, radical for its time, grew in scope and application over the  
750 next century and strongly influenced military psychiatry in World War I (Kardiner,  
751 1941; Rivers, 1918) and World War II (Grinker & Spiegel, 1945). Psychodynamic  
752 principles were later applied to the psychological problems of Holocaust survivors  
753 (Krystal, 1968; De Wind, 1984), Vietnam veterans (Lindy, 1996), rape survivors  
754 (Rose, 1991), adult survivors of childhood sexual trauma (Courtois, 1999; Roth &  
755 Batson, 1997; Shengold, 1989), and survivors of other traumatic events (Horowitz,  
756 1997). Psychodynamic ideas have also helped providers manage the sometimes  
757 complex issues that may surface in the relationship between survivor and  
758 psychotherapist (Pearlman & Saakvitne, 1995; Wilson & Lindy, 1994).  
759 Psychodynamic psychotherapies operate on the assumption that addressing  
760 unconscious mental contents and conflicts (including those that may have been  
761 blocked from consciousness as part of a maladaptive response) can help survivors  
762 cope with the effects of psychological trauma. Psychological meanings of post-  
763 traumatic responses are explored by examination of the fears, fantasies, and  
764 defenses stirred up by the traumatic event.

765

DISCUSSION

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Individual case reports comprise the bulk of the psychodynamic literature on the treatment of psychological trauma, but a small group of empirical investigations are available to support recommending that 2 psychodynamically-informed treatments can be considered as treatment options for PTSD.

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Three RCTs have supported the efficacy of Gersons' Brief Eclectic Psychotherapy for reducing PTSD symptoms in police (Gersons, Carlier, Lamberts, & van der Kolk, 2000) and community patients with PTSD (Lindauer et al., 2005; Lindauer et al., 2007). This 16-week individual psychotherapy includes both CBT (e.g., psychoeducation, imaginal exposure, cognitive restructuring) and psychodynamic elements (focus on shame and guilt, attention to the patient-therapist relationship) and a farewell ritual at the end of treatment. At present, it is unclear which elements of treatment are responsible for the improved outcomes.

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Brom and colleagues (1989) conducted a RCT that compared Horowitz' (1976) Brief Psychodynamic Therapy to hypnotherapy, trauma desensitization, and a wait-list control group in the treatment of PTSD. They found that symptoms of intrusion and avoidance improved significantly in each of the treatment groups but not in the control group; no differences across the three treatments were observed.

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While research evidence and clinical experience suggest that psychodynamic psychotherapy can be effectively combined with other forms of psychotherapy and with psychopharmacological interventions for depression (DiMascio et al., 1979; van Praag, 1989), this approach has not been sufficiently researched in work with PTSD.

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Psychodynamic ideas have, in some instances, been misapplied in clinical work with trauma survivors giving rise to concern about the creation or elaboration of so-called *false memories* (Roth & Friedman, 1997). It may be that trauma survivors are particularly prone to this phenomenon given their tendency towards dissociation. It is important that clinicians be properly trained before undertaking psychodynamic treatment of trauma survivors

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Because of its focus on basic problems in interpersonal relationships, psychodynamic psychotherapy may be useful in working with patients with complex PTSD. Clinical case studies suggest that psychodynamic psychotherapy may be of particular value in work with adult survivors of childhood sexual abuse (Courtois, 1999; Roth & Batson, 1997; Shengold, 1989).

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EVIDENCE TABLE

|   | Recommendation   | Sources   | LE   | QE   | SR |
|---|--|---|------|------|----|
| 1 | Some forms of psychodynamic psychotherapy can be considered for the treatment of PTSD. | Brom et al., 1989<br>Gersons, Carlier, Lamberts, & van der Kolk, 2000<br>Lindauer et al., 2005<br>Lindauer et al., 2007 | I    | Good | B  |
| 2 | Psychodynamic psychotherapy for patients with comorbidity PTSD.                        | Courtois, 1999<br>Roth & Batson, 1997<br>Shengold, 1989   | II-2 | Fair | B  |

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LE=Level of Evidence QE=Quality of Evidence; SR= Strength of Recommendation (see Appendix A)

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803 **B7. Patient Education**

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804 **BACKGROUND**

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805 Education of the trauma survivor is a core part of all PTSD treatment. Survivors need  
806 to better understand what they are experiencing, how to cope with reactions or  
807 symptoms, and what happens in treatment. It is also helpful to provide this  
808 information to family members or to the patient's significant others so that they can  
809 more effectively support the patient's recovery.

810 **DISCUSSION**

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811 PTSD education involves teaching the survivor to label, recognize, and understand  
812 PTSD symptoms (and other trauma-related problems) that he or she is experiencing,  
813 providing simple advice regarding coping, explaining what he or she can do to  
814 facilitate recovery, and describing treatment options. Education can help make  
815 symptoms more understandable and predictable, decrease fear of symptoms,  
816 increase awareness of coping options, and help survivors decide whether to seek  
817 treatment or learn how to better participate in treatment.

818 Education should be one of the first steps of PTSD treatment. It can help establish  
819 the credibility of the treatment provider, make treatment seem immediately helpful  
820 to the patient, and help prepare the patient for next steps in treatment. In fact,  
821 education should continue throughout PTSD treatment, sometimes in brief  
822 discussions when the patient has questions and sometimes more systematically as a  
823 formal helping activity. It can be delivered to individuals or to groups. Because those  
824 with PTSD often have difficulties with concentration and memory, repetition of  
825 educational information and provision of written information are important.

826 The content of PTSD-related education can include the following topics:

- 827 1. *Nature of PTSD symptoms:* It is often useful to help the survivor identify and  
828 label the reactions that he or she may be experiencing, recognize that emotional  
829 and physical reactions are very common (and not dangerous), and understand  
830 that anxiety and distress are often "triggered" by reminders of the traumatic  
831 experience that can include sights, sounds, or smells associated with the trauma,  
832 physical sensations (e.g., heart pounding), or behaviors of other people.  
833 However, it is important to include comment on positive steps that the individual  
834 is taking, if appropriate, rather than providing a long list of possible symptoms  
835 for review. Patients can also benefit in understanding how PTSD symptoms have  
836 their basis in adaptive survival responses to life-threatening events.
- 837 2. *Practical steps to cope with trauma-related problems:* Survivors can also be  
838 educated about ways of coping with their PTSD symptoms in order to minimize  
839 their impact on functioning and quality of life. While education about coping is not  
840 a substitute for more systematic coping skills training, simple information can  
841 also be useful. Survivors can be helped to distinguish between positive and  
842 negative coping actions. Positive coping includes actions that help to reduce  
843 anxiety, lessen other distressing reactions, and improve the situation: relaxation  
844 methods, exercise in moderation, talking to another person for support, positive  
845 distracting activities, and active participation in treatment. Negative coping  
846 methods may help to perpetuate problems and can include continual avoidance of  
847 thinking about the trauma, use of alcohol or drugs, social isolation, and  
848 aggressive or violent actions.

849 3. *Nature of the recovery process and PTSD treatment:* Survivors will sometimes  
850 have unrealistic or inaccurate expectations of recovery, and may benefit from  
851 understanding that recovery is an ongoing daily gradual process (i.e., it doesn't  
852 happen through sudden insight or "cure") and that healing doesn't mean  
853 forgetting about the trauma or having no emotional pain when thinking about it.  
854 Education about what happens in treatment is also important. This can help build  
855 motivation to participate or persist in treatment.

856 Despite the ubiquity of education in PTSD treatment, and a strong clinical consensus  
857 as to the importance of such education, there is little evidence bearing on its impact  
858 on chronic PTSD. Education has usually been a component of empirically supported  
859 treatments, but it has not been carefully evaluated as a "stand alone" treatment (nor  
860 is it intended to be delivered in the absence of other treatment elements).

861 Psychoeducation was one of several components in each study, and the effect of the  
862 psychoeducation component per se can thus not be evaluated. There is, therefore,  
863 insufficient evidence to conclude that psychoeducation alone is an effective  
864 treatment for PTSD.

865 Three studies (Krupnick J 2008); (Wallis 2002); (Weine S 2008) compared group  
866 interventions containing a psychoeducation component with WL. There were 9, 12,  
867 and 16 sessions, and the sample sizes were 48, 83, and 166. Although each  
868 intervention contained a psychoeducational component, the focus and content of the  
869 group sessions differed across studies. In 2 studies, the group intervention  
870 decreased PTSD symptoms compared with WL, while in the third; PTSD symptoms  
871 were only evaluated as a mediator for effects on access to mental health services. No  
872 study included a control condition for the psychoeducation component.

### 873 EVIDENCE

|   | Recommendation   | Sources  | LE          | QE           | R      |
|---|--|--|-------------|--------------|--------|
| 1 | Psychoeducation is recommended as component of PTSD treatment. | Foa et al., 1999<br>Lubin et al 1998<br>(Krupnick J 2008);<br>(Wallis 2002);<br>(Weine S 2008) | III<br>II-2 | Poor<br>Fair | C<br>B |

874 LE=Level of Evidence QE=Quality of Evidence; SR= Strength of Recommendation (see Appendix A)

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## 876 B8. Group Therapy

### 877 BACKGROUND

878 The ISTSS (2009) guideline includes a comprehensive critical review of group therapy  
879 approaches for the treatment of PTSD. (Shea, McDevitt-Murphy, Ready, & Schnurr,  
880 2009). Shea et al.'s discussion builds upon the previous edition's chapter by Foy and  
881 colleagues (2000). The authors briefly review the use of group therapy for PTSD. They  
882 note that it first began to be used as a "front-line treatment" for PTSD in the 1970's, and  
883 that it has continued to be used, and researched, up to the present. There is an intuitive  
884 appeal of providing this form of therapy to patients who, by the nature of their disorder,  
885 have to deal with "isolation, alienation, and diminished feelings" (Foy et al., 2000).  
886 Thus, it is possible that group therapy may foster "survivor helping survivor" feelings in  
887 participants.

888 Foy et al., (2000) and Shea et al., (2009) distinguish group treatment approaches by  
889 their emphasis on reintegration of the traumatic experience as an integral change

890 process. Trauma-focused groups assume integration of the traumatic memory and  
891 modify the meaning of the trauma for the individual, while present-centered supportive  
892 approaches aim to decrease isolation and increase sense of competence. The authors  
893 characterize three overarching group therapy orientations (See table I-5):  
894 Psychodynamic/Interpersonal/Process (present- or trauma-focused), Supportive  
895 (present –focused), and Cognitive Behavioral (predominantly trauma-focused). Most  
896 groups share common strategies designed to provide a sense of safety, trust, and  
897 develop cohesion among members. The three approaches do, however, differ in  
898 significant ways in terms of techniques and strategies used:

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**Table I - 5 Group Therapy in PTSD (Shea et al., 2009)**

| Approach   | Techniques/Strategies   |
|--|---|
| Supportive groups<br>(Present focused)                         | -Aim to enhance daily functioning through provision of safety, trust, acceptance, and normalization of symptoms and experiences<br>-Help individuals develop sense of mastery over problems via group feedback, emotional support, and reinforcement of adaptive behaviors<br>-Focus on current life issues rather than traumatic experiences   |
| Psychodynamic/<br>Interpersonal<br>Process<br>(trauma focused) | -Facilitate insight-based learning and change<br>-When an explicit focus on trauma is present, trauma material arises in a less structured manner or covertly and emphasis is on increasing awareness of unconscious fears and maladaptive patterns<br>-Emphasize understanding the meaning of the -trauma symptoms<br>-Help individuals gain insight and make connections into how current difficulties may be linked to the trauma<br>-The Interpersonal Therapy (IPT) model helps groups members identify their specific relationship difficulties and behavioral patterns that promote poor functioning<br>-“Process” groups maintain emphasis on the immediate present experience of the individual, their feelings and needs, and their interactions with other members |
| Cognitive-behavioral<br>therapy groups<br>(Trauma focused)     | -Include psychoeducation on trauma and skills training to manage anxiety and arousal<br>-Trauma is directly addressed via repeated imaginal exposure techniques in session and having individuals listen to audio-recording of their trauma experiences as homework between sessions<br>-Maladaptive thoughts and beliefs are identified and modified or restructured<br>-In final sessions, relapse prevention strategies are planned and coping skills reviewed   |

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The value and necessity of these factors, however, have not been examined empirically.

Foy et al. (2000) provide a useful guide to selecting candidates for group therapy (although, the value and necessity of these factors have not been examined empirically):

- Flexibility in personal schedule
- Ability to establish interpersonal trust
- Prior group experience, including 12-step groups
- Completion of a preparatory course of individual therapy
- Similar traumatic experiences with other group members
- Compatibility for gender, ethnicity, and sexual orientation
- Willingness to abide by rules of group confidentiality
- Not severely paranoid or sociopathic
- Stable living arrangement

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**918 Contraindications for Group Therapy (Shea et al., 2009)**

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- Active psychosis
- Severe organicity or limited cognitive capacity
- Pending litigation or compensation seeking

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923 Trial participants in the studies reviewed commonly lacked previous individual or group  
924 therapy experience.

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926 Although clinical judgment assumes that positive engagement in and outcomes from  
927 trauma-focused group treatment depend on the ability to tolerate high anxiety or other  
928 emotions, no valid method exists for assessing such a trait and studies of trauma-  
929 focused groups have not examined it as a predictor of effect (Shea et al., 2009).

929

**930 Indications for Trauma Focus versus Supportive Groups (from Foy et al., 2000)**

931

- Individual can tolerate high anxiety arousal or other strong affects
- No active suicidality or homicidality
- Substance abuse or other comorbidities are under control
- Individual accepts rationale for trauma uncovering work
- Willingness to self-disclose personal traumatic experiences
- No current life crises

936

937 Although most studies of group treatment for PTSD do focus on a particular trauma  
938 type, the importance of homogeneity of groups in terms of trauma type is an  
939 unanswered question.

940

941 Vicarious traumatization is a concern within trauma-focused groups, but no published  
942 evidence exists indicating that negative effects occur for some members in trauma-  
943 focused group treatment as a result of vicarious traumatization and systematic  
944 investigation of this possibility has not occurred (Shea et al., 2009).

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**RATIONALE**

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947 The empirical literature on group treatment for PTSD has grown since the publication  
948 of the first edition of the Treatment Guidelines for PTSD although there remain  
949 methodological weaknesses in study designs and there is no empirical evidence to  
950 support a conclusion that group treatment is superior to individual treatment for  
951 trauma.

951

952 None-the-less, it does appear that group-based treatment for individuals diagnosed  
953 with PTSD is associated with improvements in symptoms of PTSD and there is  
954 growing belief that some unique attributes of the group treatment format provide  
955 benefits that are superior to individual treatment for trauma. Identified benefits  
956 include; efficiency in treatment provision, development of support and understanding  
957 between group members that may counteract isolation and alienation.

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**DISCUSSION**

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959 Our search for relevant studies began with a review of studies included in Shea et al.  
960 (2009) and was limited to studies in which the sample participants met DSM criteria  
961 for PTSD and the active treatment was solely or predominantly in group format. Of  
962 the total 22 studies in Shea et al. (2009), 14 studies met the inclusion criteria,  
963 including six randomized and two nonrandomized trials comparing at least one active  
964 treatment group to a comparison or control condition, and six studies reporting pre-  
965 to post-treatment effects of a single group treatment condition. Four additional  
966 studies, not included in the Shea et al. (2009), provide additional information to

966 consider when weighing the effectiveness of group therapy. Two were recently  
967 published and one of these focused on a Veteran sample (Beck, Coffey Foy, Keane, &  
968 Blanchard, 2009; Ready, Thomas, Worley, Backscheider, Harvey, Baltzell, et al.,  
969 2008). Two additional RCTs: Rogers et al., (1999), although dated, was a  
970 randomized trial with a Veterans sample and Spielman et al. (2004) compared  
971 trauma and present focus group therapy in women sexually abused. . Fourteen of  
972 the total 18 studies examined cognitive-behavioral interventions; two examined  
973 interpersonal therapy, and one psychodynamic intervention.

#### 974 **Randomized Studies**

975 Eight of the 18 studies reviewed used randomized designs. Of these eight, six  
976 examined cognitive-behavioral therapy (CBT) approaches. Only three of these six  
977 compared the active treatment to a comparison condition that was not a wait-list.  
978 Schnurr and colleagues (2003) investigated Trauma-Focused Group Therapy (TFGT)  
979 in male Veterans of the Vietnam War in the largest and most rigorous study to date  
980 of group therapy for PTSD (Schnurr, Friedman, Foy, Shea, Hsieh, Lavori, et al.,  
981 2003). TFGT incorporates group-based psychoeducation, coping skills training,  
982 imaginal exposure, cognitive challenging, and relapse prevention, with one-third of  
983 all sessions devoted to individual work (Foy, Ruzek, Glynn, Riney, & Gusman, 1997).  
984 Schnurr et al (2003) did not include individual sessions. TFGT was compared with  
985 present-centered group therapy (PCGT), an approach designed to provide the  
986 “nonspecific” factors of support and interpersonal connection inherent in group  
987 treatment. Both groups experienced significant modest-sized pre- to post-treatment  
988 improvement in PTSD, which were maintained at 12 months. The primary intention-  
989 to-treat (ITT) analyses did not find differences on PTSD or any other outcomes  
990 between the group conditions (Schnurr, 2003). Rogers and colleagues (Rogers et al.,  
991 1999) also focused on Vietnam War Veterans in their randomized pilot trial  
992 comparing a single group session of flooding-based exposure therapy with a single  
993 group session of eye movement desensitization and reprocessing (EMDR) in 12  
994 Veterans who were undergoing inpatient treatment for combat-related PTSD. There  
995 were no differences between groups on PTSD symptoms post-treatment. The authors  
996 reported that both groups showed significant improvements. Lastly, Beck and  
997 colleagues (Beck et al., 2009) randomized 44 individuals with PTSD related to motor  
998 vehicle accidents (MVAs) to either Group Cognitive Behavior Therapy (GCBT) or a  
999 minimal contact comparison condition. The GCBT was a 14-week treatment  
1000 adaptation of individual CBT to a group setting (for details on development and  
1001 adaptations, see Beck & Coffey, 2005). The MCC condition consisted of telephone  
1002 contact by the project coordinator once every 4 weeks during the span of 14 weeks  
1003 where a structured script was used to assess PTSD and suicidality, and provide  
1004 minimal support but no active intervention. At post-treatment, GCBT resulted in  
1005 significantly greater reductions in PTSD symptoms among treatment completers,  
1006 with large between group effect sizes and stability of gains at 3-months. Significantly  
1007 more patients in GCBT (88.3%) versus in MCC (31.3%) no longer met criteria for  
1008 PTSD at post-treatment.

1009 The remaining three randomized studies comparing CBT to a wait-list (WL) control  
1010 involved female populations. In the largest of the three trial (n = 168), ITT analyses  
1011 showed that Imagery Rehearsal Therapy (IRT) resulted in significantly more  
1012 improvement in PTSD, nightmares, and sleep, with large between-group effects on  
1013 the CAPS and PTSD Symptom Scale (PSS) and improvements maintained at 6  
1014 months (Krakow, Hollifield, Johnston, Koss, Schrader, Warner, et al., 2001; Krakow,  
1015 Hollifield, Scharader, Koss, Tandberg, Lauriello, et al., 2000). Large effects were  
1016 also found on PTSD symptoms between an affect management group and WL control

1017 where both groups also received individual therapy and medication (Zlotnick, Shea,  
1018 Rosen, Simpson, Mulrenin, Begin, et al., 1997). Both of these trials examined PTSD  
1019 related to childhood abuse. A trial of women with PTSD related to diverse traumas,  
1020 as well as comorbid panic disorder, indicated that multichannel exposure therapy  
1021 (MCET) was superior to control in PTSD reduction (Falsetti, Resnick, Davis, &  
1022 Gallagher, 2001; Falsetti, Resnick, & Davis, 2005).

1023 Two small-scale randomized trials evaluated non-CBT approaches versus wait-list  
1024 controls in women with PTSD related to sexual abuse, mostly in childhood.  
1025 Comparison of a trauma-focused group and a present-focused group, both based on  
1026 psychodynamic principles, showed no differences for either relative to a wait-list  
1027 control (Spiegel, Classen, Thurston, & Butler, 2004) and even when combined, the  
1028 composite treatment group showed significant more improvement only on non-PTSD  
1029 measures (Classen, Koopman, Nevill-Manning, & Spiegel, 2001). In contrast,  
1030 Krupnick and colleagues (Krupnick, Green, Miranda, & Stockton, 2008) found  
1031 significant effects with ITT analyses for Interpersonal Therapy group on PTSD,  
1032 depression, and interpersonal functioning, with a medium-to-large effect for PTSD.

### 1033 **Nonrandomized Controlled Studies**

1034 \_Two studies compared a group treatment to control condition with a nonrandomized  
1035 design. Cloitre and Koenen (2001) reported significant improvement in PTSD, anger,  
1036 and depression for an interpersonal/process therapy compared to a wait-list control,  
1037 but only for therapy groups that did not have members with borderline personality  
1038 disorder (BPD). The effect on PTSD symptoms was large. Resick and Schnicke  
1039 (1992) observed significant improvements for cognitive reprocessing therapy (CPT)  
1040 but not for wait-list on PTSD, depression, distress, and social adjustment, with  
1041 maintenance of improvement at 6 months.

1042

### 1043 **Summary of Studies without Control or Comparison Conditions:**

1044 All seven studies without a control or comparison condition evaluated CBT-based  
1045 group treatments. With a very large sample of 2,223 Vietnam War Veterans, a  
1046 naturalistic program evaluation study by Creamer and colleagues (Creamer, Elliot,  
1047 Forbes, Biddle, & Hawthorne, 2006) examined the effects of a group-based CBT  
1048 approach with components similar to trauma-focused group therapy for combat-  
1049 related PTSD (TFGT) (Foy et al., 1997), along with 6-12 sessions of individual  
1050 therapy through the 2 weeks of group treatment. Outcomes of the naturalistic trial  
1051 included a very low drop-out rate of 3% and moderate-to-large within-treatment  
1052 effect sizes at 6-months on PTSD, depression, alcohol use, family functioning, and  
1053 anger that continued to increase over a 12-month follow-up period. The design had  
1054 the advantage of reflecting routine clinical practice in a real-world treatment setting.

1055 A similar field test approach was employed by Ready et al. (2008), who evaluated an  
1056 on-going model called Group-Based Exposure Therapy (GBET) similar to TFGT in a  
1057 sample of 102 predominantly Vietnam War Veterans. Significant symptom reductions  
1058 were found in all three PTSD symptoms clusters and on both clinician ratings and  
1059 self-report measures. Large effect sizes were found on the clinician ratings of total  
1060 PTSD symptoms and medium-to-large effect sizes were found on PTSD self-report  
1061 measures on both posttreatment and 6-month follow-up. Medium effect sizes were  
1062 also found on suicidal ideation. In addition, 81% of patients with valid total CAPS  
1063 scores at pretest and posttest showed a clinically significant improvement, defined as  
1064 a reduction of 10 or more points on the total CAPS at posttest. The authors note that



1065 group formats have been favored within the VA system, although differential effects  
1066 of such multi-phased treatments as GBET are unknown.

1067 Donovan and colleagues (Donovan, Padin-Rivera, & Kowalix, 2001) evaluated a 12-  
1068 week partial hospitalization program (Transcend) in a sample of 46 male Vietnam  
1069 War Veterans. The program addresses combat-related PTSD and substance abuse  
1070 utilizing skills training, trauma processing, psychoeducation, relapse prevention, and  
1071 cognitive restructuring, among other techniques, and is supplemented by physical  
1072 exercise and community service activities. Significant medium-size decreases in  
1073 PTSD symptoms and addictive behavior were observed posttreatment and  
1074 maintained through 12-month follow-up (Donovan et al., 2001).

1075

1076 Three studies evaluated Seeking Safety, a present-centered 24-session CBT  
1077 intervention for PTSD complicated by substance abuse or dependence. All three used  
1078 small samples of less than 30. In the earliest trial, 17 of of 27 women completed  
1079 treatment and in the completer analyses significant medium-size declines were found  
1080 pretreatment to 3-month follow-up in alcohol use and in subtle symptoms of long-  
1081 term childhood trauma, such as "fear of men", but not in DSM-based PTSD  
1082 symptoms as assessed by the Modified PTSD Symptom Scale (MPSSR). Zlotnick et  
1083 al. (Zlotnick, Najavits, Rohsenow, & Johnson, 2003) evaluated Seeking Safety as an  
1084 adjunct to group treatment-as-usual based on an abstinence-oriented 12-step model  
1085 in 18 incarcerated women. They found that at posttreatment, 53% no longer met  
1086 criteria for PTSD and there were significant decrease in PTSD symptoms.  
1087 Improvements in drug and alcohol use also occurred. Gains were maintained through  
1088 3-month follow-up (Zlotnick et al., 2003). Most recently, results of an evaluation of  
1089 Seeking Safety in 18 Veterans with combat-related PTSD and substance abuse  
1090 disorders at a VA Medical Center indicated significant improvements in self-reported  
1091 PTSD and quality of life posttreatment (Cook, Wasler, Kane, Ruzek, & Woody, 2006).

1092 Lastly, an evaluation of a trauma-focused group therapy that incorporated  
1093 psychoeducation, exposure, and cognitive-behavioral interventions with 29 multiply  
1094 traumatized women showed significant pre- to posttreatment reductions in PTSD  
1095 symptoms, with clinically meaningful reductions (more than one standard deviation  
1096 blow pretreatment levels, about a 50% reduction) in scores on the Clinician-  
1097 Administered PTSD Scale (CAPS) occurring for 38% of the group. However, although  
1098 the trial abstract indicated that participants were "diagnosed with chronic PTSD", the  
1099 mean values for the CAPS reported for each cluster and the total were very small  
1100 (each < 1.7). How the CAPS was scored was not clear. In addition, self-report  
1101 measures of PTSD showed nonsignificant reductions (Lubin, Loris, Burt, & Johnson,  
1102 1998).

1103

#### 1104 **Caveat regarding analysis of the data**

1105 In examining the effects of these group treatments, a significant and prevalent  
1106 methodological limitation warrants discussion. This limitation is that most studies  
1107 failed to use analytic strategies to account for clustering of observations within  
1108 treatment groups. Participants administered treatment in group format share a  
1109 common therapy environment, which may homogenize response to the treatment.  
1110 As explained by Baldwin and colleagues (Baldwin, Murray, & Shadish, 2005), studies  
1111 that do not take into account the magnitude of the dependency among observations  
1112 taken on members of the same group, or intraclass correlation (ICC), underestimate  
1113 the standard error of the treatment effect by pooling the effect of the group with the  
1114 effect of the treatment, so that even if treatment has no effect, an incorrect analysis

1115 can suggest a treatment effect. Baldwin et al. (2005) demonstrated the significance  
1116 of this limitation by correcting the unit of analysis with an adjustment to the  
1117 denominator degrees of freedom within predominantly pre- to posttreatment  
1118 analyses in 33 studies of treatments on the American Psychological Association's list  
1119 of evidence-based group treatment to reflect the number of groups (i.e., cohorts in  
1120 each condition) in the study. This correction resulted in the loss of statistically  
1121 significant effects for over 30% of the studies. Incorporating the ICC in the  
1122 reanalyses resulted in a loss of significance for additional studies.

1123 Out of the 18 studies reviewed herein, only two studies corrected for unit of analysis  
1124 and group ICC (Beck et al., 2009; Schnurr et al., 2003) and two studies accounted  
1125 for ICC (Creamer et al., 2006; Ready et al., 2008) in analyses, with the remaining  
1126 studies taking a typical approach of treating the individual participants as the unit of  
1127 analyses and not correcting for the ICC. As was the case for the APA studies, it is  
1128 likely that true effects for group treatments for PTSD are more modest than  
1129 published effects. With the assumption that the smaller the sample size, the more  
1130 modest the true detectable effects, we adjusted the denominator degrees of freedom  
1131 in the pre- to post-treatment analysis of the primary PTSD outcome for the three  
1132 studies with the smallest sample size of those reviewed (Resick & Schnicke, 1992;  
1133 Rogers et al., 1999; Falsetti et al., 2001, 2005). The RCT of Rogers and colleagues  
1134 (2009), with a total sample of 12, used only one group per condition. Based on  
1135 Baldwin et al. (2005) method, we treated this study as if it had two groups per  
1136 condition. After adjusting the degrees of freedom, the change in symptoms were no  
1137 longer statistically significant ( $p = .06$ ). Falsetti and colleagues (2001, 2005) used a  
1138 sample of 27 in their nonrandomized controlled trial, also with only 1 group per  
1139 condition. Again assuming 2 groups per condition, correction to the degrees of  
1140 freedom resulted in a nonsignificant finding for PTSD symptom change ( $p = .088$ ).  
1141 Lastly, degrees of freedom adjustments in Resick and Schnicke's (1992) uncontrolled  
1142 trial of 19 individuals with three CPT groups resulted in pre- to post-treatment  
1143 changes on the PTSD subscale of the SCL-90-R and the arousal subscale of the PTSD  
1144 Symptom Scale no longer meeting statistical significance ( $p = .062$  and  $p = .068$ ,  
1145 respectively). In reviewing the articles for this annotation, we noted that many  
1146 authors did not indicate the number of groups/cohorts per condition, the number of  
1147 participants per group, or both. Compounding the failure to account for observation  
1148 clustering was the fact that many studies also suffered from small samples,  
1149 particularly given the number of analyses performed.

1150

## 1151 **Conclusions**

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1153 The empirical literature on group treatment for PTSD has grown since the publication  
1154 of the first edition of the Treatment Guidelines for PTSD. However, most studies  
1155 continue to utilize small sample sizes, use wait-list controls, and fail to account for  
1156 clustering of observations in analyses. Advances in methodological rigor are typified  
1157 within studies by Schnurr and colleagues (2003), as well as Beck and colleagues  
1158 (2009). Field tests of interventions developed as part of clinical practice and  
1159 evaluated on large samples, such as those by Creamer et al. (2006) and Ready et al.  
1160 (2008), offer unique information yet pose numerous questions regarding complex  
1161 multi-phase approaches that require further research.

1162 With these caveats in mind, our review indicates that, overall, the research indicates  
1163 that group-based treatment for individuals diagnosed with PTSD is associated with  
1164 improvements in symptoms of PTSD, although results are inconsistent, quality of  
1165 most trials are poor, and there is insufficient evidence regarding the efficacy of group

1166 therapy compared with individual CBTs. Reported pre- to post-treatment effect  
1167 sizes, based on analyses not adjusted for within-group dependence and unit of  
1168 analysis, range from small to large and likely overestimate the true effect of the  
1169 treatment. The amount of change exceeded that of wait-list controls for most  
1170 studies. The trial of a psychodynamic treatment evidenced the weakest within-group  
1171 effects (Classen et al., 2004). The two studies of interpersonal therapy evidenced  
1172 small to large effects (Cloitre & Koenen, 2001; Krupnick et al., 2008). Significant  
1173 support exists for cognitive-behavioral approaches, for both combat veterans and in  
1174 adults with histories of abuse, with effects ranging from small to very large. The four  
1175 studies that did correct for either unit of analysis and/or group intraclass correlation  
1176 yielded between-group effect sizes of .09 (Pre to 7-mo CAPS, ITT analysis: Schnurr  
1177 et al., 2003) and .84 (Pre to Post CAPS: Beck et al., 2009), and within-group effect  
1178 sizes of .59 (Pre to 6-mo PCL: Creamer et al., 2006), and 1.20 (Pre to 6-mo CAPS,  
1179 ITT analysis, Ready et al., 2008). Thus, variability in methodology and resultant  
1180 effects are evident.

1181 As noted above, few studies have directly compared different forms of group therapy  
1182 and the largest controlled study to date (Schnurr et al., 2003) indicated equal benefit  
1183 from trauma-focused and present-centered supportive therapies in the primary  
1184 analyses. The only other study to compare two active treatments (one trauma-  
1185 focused and the either present-focused) did not find differences, but did suffer from  
1186 small samples (Classen et al., 2004). Shea and colleagues' (2009) examination of  
1187 within-group effect sizes pre to post-treatment found no evidence that groups  
1188 focusing on trauma provide superior outcome than those who do not. Only one trial  
1189 examined a group adaptation of an existing and proven individual therapy protocol  
1190 (Beck et al., 2009). Given that within this trial, reductions in PTSD from GCBT were  
1191 comparable to those obtained in previous studies of individual CBT but GCBT did not  
1192 reduce comorbid anxiety and depression, the authors call for direct comparison of  
1193 individual versus group CBT in order to determine differential efficacy.

1194 We concur with Shea et al.'s (2009) assessment that it remains unknown whether  
1195 improvements found in most studies of cognitive-behavioral or  
1196 interpersonal/process-oriented treatments are due to the strategies employed by the  
1197 different group types. Schnurr et al. (2003) remains the only adequately powered  
1198 comparison of an active treatment and control, and primary analyses yielded null  
1199 findings. Ready and colleagues discuss the role of "dose" of exposure therapy as a  
1200 potential factor in the differences observed between their GBET and some of the  
1201 other group models for treating war-related PTSD, such as Schnurr et al.'s (2003)  
1202 TFGT. Group-based exposure therapy has a minimum of 60 hours of exposure  
1203 therapy compared to TFGT's 40 hours (4 hours of within-group war trauma  
1204 presentations per patient, 16 hours of listening to recordings of one of the patients  
1205 own presentations, and 20 hours of hearing other patients war trauma  
1206 presentations). They also note that it also may be significant that the number of  
1207 exposures to a patients' own traumatic experiences in GBET (22 or more) is more  
1208 similar to the number of exposures typically used in individual exposure therapy (35  
1209 or more; Foa, Hembree, & Dancu, 2002, as cited in Ready et al., 2008) than in TFGT  
1210 (10). Perhaps the higher dose of exposure therapy contributed to the differences in  
1211 treatment outcomes between GBET and Schnurr et al.'s (2003) trial of TFGT. Further  
1212 evidence for such a possibility comes from the fact that when Schnurr et al. (2003)  
1213 included participants with at least 24 sessions of treatment in secondary analyses,  
1214 significant differences suggested that trauma-focused groups may be more effective  
1215 than present-centered groups.

1216 EVIDENCE TABLE

|   | Recommendation  | Sources  | LE      | QE   | SR |
|---|---|--|---------|------|----|
| 1 | Consider group treatment for patients with PTSD                                   | Beck et al., 2009 ①<br>Classen et al., 2001; ①<br>Cloitre & Koenen, 2001<br>Cook et al., 2006<br>Creamer et al., 2006 -<br>Donovan et al., 2001 -<br>Falsetti et al., 2001, 2005 ①<br>Krakow et al., 2000, 2001 ①<br>Krupnick et al., 2008 ①<br>Lublin et al., 1998<br>Najavits et al., 1998<br>Ready et al., 2008 -<br>Resick & Schniske, 1992<br>Rogers et al., 1999 ①<br>Schnurr et al., 2003 v①<br>Spiegel et al., 2004①<br>Zlotnick et al., 1997①<br>Zlotnick et al., 2003<br>Foy et al., 2000 (SR)<br>Shea et al., 2009 (SR) | I<br>II | Fair | B  |
| 2 | Current findings do not favor trauma-focused versus present-focused group therapy | Schnurr et al., 2003<br>Classen et al., 2004<br>Shea et al., 2009  | I       | Good | I  |

1217 LE=Level of Evidence QE=Quality of Evidence; SR= Strength of Recommendation (see Appendix A)

1218

1219 **B9. Dialectical Behavior Therapy**

1220 BACKGROUND

1221 Dialectical behavior therapy (DBT) is a comprehensive cognitive-behavioral  
 1222 treatment for complex, difficult-to-treat mental disorders, specifically designed to  
 1223 treat chronically suicidal individuals, and patients with multi-disorders or borderline  
 1224 personality disorder (BPD).

1225 DBT has since been adapted for other seemingly intractable behavioral disorders  
 1226 involving emotion dysregulation, including substance dependence in individuals with  
 1227 BD and binge eating, to other clinical populations (e.g., depressed, suicidal  
 1228 adolescents), and in a variety of settings (e.g., inpatient, partial hospitalization,  
 1229 forensic).

1230 While considerable evidence supports the use of exposure-based treatment for PTSD,  
 1231 its utilization may pose some problems for patients where the symptoms of PTSD are  
 1232 complicated. High rates of attrition, suicidality, dissociation, destructive impulsivity,  
 1233 and chaotic life problems are reasons cited by clinicians for abandoning empirically  
 1234 supported exposure treatment. Some practitioners have suggested that the  
 1235 approach of DBT, designed to address many of these issues, offers useful strategies  
 1236 for addressing the needs of patients considered poor candidates for exposure  
 1237 therapy.

1238 The DBT approach incorporates what is valuable from other forms of therapy, and is  
1239 based on a clear acknowledgement of the value of a strong relationship between  
1240 therapist and patient. Therapy is structured in stages and at each stage a clear  
1241 hierarchy of targets is defined. The techniques used in DBT are extensive and  
1242 varied, addressing essentially every aspect of therapy. These techniques are  
1243 underpinned by a dialectical philosophy that recommends a balanced, flexible and  
1244 systemic approach to the work of therapy. Patients are helped to understand their  
1245 problem behaviors and then deal with situations more effectively. They are taught  
1246 the necessary skills to enable them to do so and helped to deal with any problems  
1247 that they may have in applying those skills. Advice and support is available between  
1248 sessions. The patient is encouraged and helped to take responsibility for dealing with  
1249 life's challenges.

## 1250 DISCUSSION

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1251 Although DBT is becoming more common as a technique for treating patients with  
1252 BPD, no clinical trials have been reported in the literature for the use of DBT in  
1253 patients with PTSD. The following studies concern patients with BPD who attempt  
1254 some form of self-injury; however, for patients with PTSD and comorbid BPD, these  
1255 studies may be applicable to the treatment decision process.

1256 In a meta-analysis of RCTs of "psychosocial and/or psychopharmacological treatment  
1257 versus standard or less intensive types of aftercare" for patients who had shown self-  
1258 harm behaviors, Hawton et al., (2000) compared DBT vs standard after care and  
1259 found that DBT significantly reduced rates of further self-harm (0.24; 0.06 to 0.93)."  
1260 The authors caution, however, that "there still remains considerable uncertainty  
1261 about which forms of psychosocial and physical treatments of self-harm patients are  
1262 most effective, inclusion of insufficient numbers of patients in trials being the main  
1263 limiting factor."

1264 van den Bosch et al., (2002) and Verheul et al., (2003) reported on the effectiveness  
1265 of DBT in a group of 58 female BPD patients. For these women, DBT therapy  
1266 "resulted in better retention rates and greater reductions of self-mutilating and self-  
1267 damaging impulsive behaviours compared with usual treatment, especially among  
1268 those with a history of frequent self-mutilation" (Verheul et al., 2003). In the same  
1269 study group, van den Bosch et al., (2002) compared the results of therapy in women  
1270 with and without comorbid substance abuse. They found that comorbid substance  
1271 abuse did not dilute the effect of the DBT, but that the DBT therapy had no effect on  
1272 the womens' substance problems. Evans et al., (1999) compared the provision of  
1273 self-help booklets alone to six sessions of cognitive therapy linked to the booklets,  
1274 which contained elements of DBT (MACT) in 34 patients who had attempted self-  
1275 harm. The authors reported that MACT therapy led to a lowering of the number of  
1276 suicidal acts per month, and also improved self-rated depressive symptoms.

1277 Linehan and colleagues, (1993) conducted a RCT of 39 women with BPD, who were  
1278 randomly assigned to DBT or usual care for one year, then followed-up at six and  
1279 twelve months following treatment. The authors reported that DBT patients had  
1280 significantly less parasuicidal behavior, less anger, and better self-reported social  
1281 adjustment during the initial 6 months and significantly fewer psychiatric inpatient  
1282 days and better interviewer-rated social adjustment during the final 6 months;  
1283 overall, DBT subjects had significantly higher Global Assessment Scale scores during  
1284 the follow-up year.

1285 Telch et al., (2001) and Safer et al., (2001) expanded the DBT concept to treatment  
1286 of women with binge eating disorder. In both studies, women were randomly  
1287 assigned to DBT or a wait list (Telch study – 44 women; Safer study – 31 women)

1288 and the authors' results were similar; patients improved significantly in reduction of  
1289 binge/purge behaviors, but did not differ on any secondary measures.

1290 Bohus et al., (2000) treated 24 female chronically suicidal patients with DBT and  
1291 found significant improvements in ratings of depression, dissociation, anxiety and  
1292 global stress and a highly significant decrease in the number of parasuicidal acts.

1293 Gould et al., (2003) and Miller and Glinski, (2000) identify DBT as a promising  
1294 treatment for suicide, however, they acknowledge the need for RCTs. In their  
1295 overview of the use of DBT, Koerner and Linehan (2000) also stress the need for  
1296 longitudinal follow-up studies to determine suicide rates and maintenance of  
1297 treatment gains.

#### 1298 EVIDENCE TABLE

|   | Recommendation   | Sources   | QE | QE   | R |
|---|--|---|----|------|---|
| 1 | Consider DBT for patients with a borderline personality disorder typified by parasuicidal behaviors. | Evans et al., 1999<br>Hawton et al., 2000<br>Linehan et al., 1993<br>Safer et al., 2001<br>Telch et al., 2001<br>van den Bosch et al., 2002<br>Verheul et al., 2003 | I  | Fair | B |

1299 *LE=Level of Evidence QE=Quality of Evidence; SR= Strength of Recommendation (see Appendix A)*

1300

## 1301 B10. Hypnosis

### 1302 BACKGROUND

1303 Hypnosis is not held to be an ASD or PTSD therapy per se but may significantly  
1304 enhance the effectiveness of other therapies in their treatment or in the  
1305 management of a variety of related clinical conditions (Kirsch et al., 1998; Spiegel &  
1306 Spiegel, 1987). Historically, hypnotic treatments have played a role in the  
1307 management of shell shock, battle fatigue and traumatic neuroses.

1308 Hypnosis is defined by the APA as "a procedure during which a health professional or  
1309 researcher suggests that a client, patient, or subject experience changes in  
1310 sensations, perceptions, thought, or behavior. The hypnotic context is generally  
1311 established by an induction procedure (Kirsch, 1994). An induction procedure  
1312 typically entails instructions to disregard extraneous concerns and focus on the  
1313 experiences and behaviors that the therapist suggests or that may arise  
1314 spontaneously.

1315 Hypnosis should only be used by credentialed health care professionals, who are  
1316 properly trained in the clinical use of hypnosis and are working within the areas of  
1317 their professional expertise.

### 1318 DISCUSSION

1319 Most of the case studies that reported that hypnosis was useful in treating  
1320 posttrauma disturbances following a variety of traumas lack methodological rigor,  
1321 and therefore strong conclusions about the efficacy of hypnosis to treat PTSD cannot  
1322 be drawn (Rothbaum, 2001).

1323 Brom and colleagues (1989) in a RCT, showed that hypnosis and desensitization  
1324 significantly decreased intrusive symptoms whereas psychodynamic therapy was

1325 useful for reducing avoidance symptoms in patients with various types of  
1326 posttraumatic symptomatology. In a meta-analysis, Sherman (1998) compared the  
1327 effects of the Brom et al. trial with those of other controlled studies and found that  
1328 the major advantage of using hypnosis may appear at long term follow-up rather  
1329 than at the end of treatment: this is consistent with meta-analyses of hypnosis for  
1330 conditions other than PTSD (Kirsch et al., 1999).

1331 Various studies, including meta-analyses, of the treatment of anxiety, pain,  
1332 repetitive nightmares and other conditions often associated with PTSD imply that  
1333 hypnosis can substantially reduce the severity of these problems (Daly & Wulff,  
1334 1987; Jiranek, 1993; Richmond et al., 1996; Kirsch et al., 1995; Eichelman, 1985;  
1335 Kingsbury, 1993) and enhance the effectiveness of psychodynamic and cognitive  
1336 behavioral therapy (Kirsch, 1996; Kirsch et al., 1999; Smith et al., 1980). Most of  
1337 the literature on the use of hypnosis for PTSD is based on service and case studies.  
1338 Shakibaei, (2008) reported that hypnotherapy helped reduce both pain and re-  
1339 experiencing of traumatic events among burn patients in a randomized control trial  
1340 but it should be noted that patients meeting criteria for any acute psychiatric  
1341 disorder were specifically excluded from this study.

1342 Abramowitz (2008) reports on a RCT in which hypnotherapy was compared to  
1343 zolpidem treatment for insomnia among 32 patients with combat PTSD who were  
1344 also suffering from insomnia. All patients were already taking an SSRI. He found  
1345 significant improvement in PTSD symptoms and sleep quality, number of  
1346 awakenings, ability to concentrate in the morning, and morning sleepiness in the  
1347 hypnotherapy group. Sleep time improved equally in both groups.

1348 There are a number of indications for using hypnosis in the treatment of PTSD:

- 1349 1. Hypnotic techniques may be especially valuable as an adjunctive treatment for  
1350 symptoms often associated with PTSD including dissociation, anxiety, pain,  
1351 nightmares and insomnia.
- 1352 2. PTSD patients who manifest at least moderate hypnotizability may benefit from  
1353 the addition of hypnotic techniques to their treatment.
- 1354 3. Because confronting traumatic memories may be very difficult for some PTSD  
1355 patients, hypnotic techniques may provide them with a means to modulate their  
1356 emotional and cognitive distance from such memories as they are worked  
1357 through therapeutically.

1358

1359 There are a number of contraindications for using traditional hypnotic techniques in the  
1360 treatment of PTSD (Cardena et al., 2008):

- 1361 1. In the rare cases of individuals who are refractory or minimally responsive to  
1362 suggestion, hypnotic techniques may not be the best choice, because there is  
1363 some evidence that hypnotizability is related to treatment outcome efficacy  
1364 (Levitt, 1994; Spiegel et al., 1981 & 1993).
- 1365 2. Some PTSD patients may be resistant to hypnotic treatment because of religious  
1366 concerns or other beliefs. If resistance persists, other suggestive techniques may  
1367 be tried including emotional self-regulation therapy (ESRT), which is done with  
1368 open eyes and uses sensory recall exercises rather than a hypnotic induction  
1369 (Bayot et al., 1997; Kirsch et al., 1999).
- 1370 3. For patients who have low blood pressure or are prone to fall asleep, hypnotic  
1371 procedures such as "alert hand," which emphasize alertness and activity rather  
1372 than relaxation, may be substituted (Cardena et al., 1998).

1373 **EVIDENCE**

|   | Recommendation                                   | Sources                            | LE | QE   | R |
|---|--|------------------------------------|----|------|---|
| 1 | Hypnosis may be used to alleviate PTSD symptoms. | Brom et al., 1989<br>Sherman, 1998 | I  | Fair | B |

1374 *LE=Level of Evidence QE=Quality of Evidence; SR= Strength of Recommendation (see Appendix A)*

1375

1376 **B11. Behavioral Couples Therapy**1377 **BACKGROUND**

1378 Perceived social support has been identified as an important resilience factor in  
1379 PTSD. Families report significant distress during the deployment cycle the  
1380 prevalence of family problems, such as divorce. A number of family and couples  
1381 interventions have been developed, including Behavioral Family Therapy (BFT)  
1382 (Mueser & Glynn, 1995), Cognitive-Behavioral Couples Therapy (CBCT; Monson et  
1383 al., 2004), and Support and Family Education (SAFE; Sherman, 2003). However,  
1384 there is as yet little support for these interventions as a first line treatment for PTSD.

1385 **DISCUSSION**

1386 Glynn et al. (1999) conducted a RCT of couples or family treatment for PTSD utilizing  
1387 either an Exposure condition, Exposure followed by BFT, or a wait list control. While  
1388 both active treatment conditions improved on PTSD symptoms, BFT did not  
1389 significantly improve the PTSD symptoms, compared to the Exposure only condition.  
1390 However, BFT did demonstrate improved problem solving skills relative to the other  
1391 two conditions.

1392 Monson et al. (2004, 2005) conducted a small, uncontrolled pilot study of seven  
1393 couples who received CBCT for PTSD. Significant improvements were found on  
1394 PTSD, depression, and anxiety for both veterans and wives. The improvement in  
1395 relationship satisfaction was more mixed, with no improvement for husbands but  
1396 greater improvement for wives.

1397 Devilly (2002) examined a Lifestyle Management course for male veterans with PTSD  
1398 and their partners, in a weeklong residential treatment. Both veterans and their  
1399 partners experienced significant reductions in anxiety, depression, and stress;  
1400 veterans also experienced significant reductions in PTSD. However, the effect size of  
1401 these changes was small and symptom improvements were considered to be of  
1402 limited clinical importance.

1403 No studies that evaluated behavioral couple's therapy (BCT) for treatment of  
1404 posttraumatic stress disorder (PTSD) were identified. One study (Rotunda et al.,  
1405 2008) evaluated BCT for substance use disorder (SUD) in veterans with comorbid  
1406 PTSD. The study was not designed to evaluate BCT for treatment of PTSD, but did  
1407 assess psychological symptoms as a function of BCT. Although, the effects of BCT on  
1408 PTSD symptoms specifically were not reported, the results suggested that BCT may  
1409 reduce general psychological distress and increase abstinence in male veterans with  
1410 SUD and comorbid PTSD. However, caution should be taken in generalizing these  
1411 findings to a population with PTSD alone, given the body of literature demonstrating  
1412 that those with comorbid SUD and PTSD are different from those with PTSD alone on  
1413 a number of important clinical variables (e.g., symptom severity, chronicity of  
1414 illness, treatment refractory) (Robert, Beckham, and Moore, 2007).



1415 No review or meta-analysis publications that addressed BCT, BFT, or CBCT as a  
1416 treatment for PTSD were identified.

1417 **EVIDENCE TABLE**

|   | Evidence   | Sources of Evidence                         | LE | QE   | SR |
|---|--|---|----|------|----|
| 1 | BFT did not significantly improve the PTSD symptoms, and was inferior to other psychotherapies | Glynn et al. (1999)<br>Monson et al. (2004) | I  | Fair | I  |

1418 LE=Level of Evidence QE=Quality of Evidence; SR= Strength of Recommendation (see Appendix A)

1419

1420 **B12. Telemedicine and Web-based Interventions**

1421 ***B12-1. TELEMEDICINE INTERVENTIONS***

1422 **BACKGROUND**

1423 Increasingly, a range of technologies are being adapted to enhance delivery of  
1424 mental health services. Such technologies include the telephone and  
1425 videoconferencing tools. Some technological applications assist human providers in  
1426 delivering their treatments to patients, as when videoconferences or telephones are  
1427 used to reach those for whom attendance may be difficult, or increasing convenience  
1428 for patients by eliminating travel to face-to-face sessions. Telephone-based services  
1429 – phone-based counseling, automated telephone assessment, and interactive  
1430 telephone applications – provide ways of extending assessment and treatment into  
1431 the natural environment.

1432 **DISCUSSION**

1433 A burgeoning body of rigorous research has demonstrated that psychotherapy for  
1434 treatment of depression and anxiety disorders delivered either via  
1435 videoteleconferencing (VTC) or via telephone is not only effective, but clinically  
1436 equivalent to face-to-face delivery (O’Reilly et al., 2007; Bee et al., 2008).  
1437 However, only one study to date has conducted that level of rigorous examination of  
1438 psychotherapy delivered to a PTSD population via VTC (Morland et al., 2009). This  
1439 study found that Anger Management Group therapy via VTC was *as effective as* face-  
1440 to-face delivery in reducing anger symptoms in PTSD patients, both immediately  
1441 post treatment and in short term follow-up (i.e., 3 months). It also found that there  
1442 were no significant differences between the two modalities in satisfaction with treatment,  
1443 treatment credibility, attendance, homework completion, attrition, or alliance among  
1444 group members (Greene et al., 2010). However, patients receiving treatment via  
1445 VTC had lower therapeutic alliance with the group leader than those who received  
1446 face-to-face delivery.

1447 Additional trials of VTC delivery of PTSD-specific treatments have also demonstrated  
1448 clinical effectiveness that was comparable to face-to-face delivery (e.g. Frueh et al.,  
1449 2007). However, due to the methodologies used (e.g. small sample size, non-  
1450 randomized), they were not able to test if VTC was actually equivalent to face-to-  
1451 face treatment. A non-random cohort study demonstrated that CBT delivered via  
1452 VTC improved PTSD symptoms at a level similar to face-to-face group delivery  
1453 (Germain, Marchand, Bouchard, Drouin, & Guay, 2009). A recent pilot study found  
1454 that Prolonged Exposure Therapy delivered via VTC was highly effective, safe, and  
1455 feasible (Tuerk et al., 2010).

1456 There has been somewhat inconsistent evidence of process outcomes such as patient  
1457 and provider satisfaction, patient treatment preference, comfort talking to their  
1458 therapists, and homework compliance among the different trials comparing VTC and  
1459 face-to-face delivery of PTSD interventions. Although several studies have found no  
1460 significant differences between the two modalities, some have found that in-person  
1461 delivery has generated slightly better process outcomes (Morland, Pierce, & Wong,  
1462 2004; Frueh et al., 2007).

1463 The effectiveness of telephone delivery of case management and support has been  
1464 well proven for a wide variety of behavioral health interventions. However, it is  
1465 much less studied with PTSD patients. A small cohort study demonstrated that  
1466 telephone-based monitoring and support improved patient satisfaction and entry into  
1467 aftercare compared to treatment as usual condition (Rosen et al., 2006).

1468 Mobile phone-based interventions present several advantages and capabilities (e.g.,  
1469 web-browsing, text messaging, software applications, etc.) that could address  
1470 common problems in delivering evidence-based treatments (Boschen, 2010);  
1471 however, the evidence to support these technologies in the PTSD interventions has  
1472 yet to be generated.

1473 In summary, telephone delivery and videoconferencing can be effectively used to  
1474 overcome geographical barriers to mental health care. There is an abundance of  
1475 evidence that the modalities are safe and effective. There is preliminary evidence to  
1476 suggest that psychotherapy delivered via these modalities is as effective as face-to-  
1477 face care. As the field develops, additional research needs to examine how TMH  
1478 modalities affect the therapeutic process and also how mobile phone-based  
1479 interventions can be effectively used for PTSD treatment.

## 1480 **B12-2. WEB-BASED INTERVENTIONS**

### 1481 **BACKGROUND**

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1482 Increasingly, a range of computer and Internet technologies are being adapted to  
1483 enhance delivery of mental health services. Web-based applications can deliver  
1484 elements of treatment (such as psychoeducation or skills training) in the absence of  
1485 provider contact or with reduced contact. And it is possible that access to help via  
1486 technologies may increase engagement in care by reducing stigma associated with  
1487 treatment-seeking and increasing accessibility of care (e.g., for rural populations,  
1488 disabled persons, individuals without easy transportation access). To date, research  
1489 conducted with PTSD patients has been very limited, but services are increasingly  
1490 being delivered via these technologies. Newly developed technologies can present  
1491 significant challenges related to patient confidentiality and safety, and these must be  
1492 addressed carefully, by both the individual providers and the organization delivering  
1493 these interventions.

### 1494 **DISCUSSION**

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1495 Web-based interventions have very limited research for treatment of PTSD, although  
1496 several studies have been done to assess these techniques particularly in  
1497 traumatized individuals with general distress or subclinical PTSD symptoms. Web-  
1498 based interventions may provide an effective delivery modality for CBT techniques  
1499 that can be considered in certain circumstances. However, these interventions raise  
1500 a number of privacy and confidentiality issues and have not been directly compared  
1501 with other evidence-based person-to-person CBT modalities that have been shown to  
1502 be efficacious may be considered as a standalone intervention, or as an alternative  
1503 to standard mental health treatment for PTSD.

1504 The Internet provides a potential resource for delivery of both information  
1505 (psychoeducation) and more complex interventions. At present, while there is much  
1506 traumatic stress-related information available on the Web, the accuracy and  
1507 authoritativeness of the information can be difficult for consumers to determine.  
1508 Bremner, Quinn, Quinn, and Veledar (2006) reviewed the quality of 80 websites  
1509 related to psychological aspects of trauma, and found that 42% of sites had  
1510 inaccurate information, 82% did not provide a source of content, and 41% did not  
1511 use a mental-health professional in the development of the content. The authors  
1512 concluded that, although abundant, websites providing information about traumatic  
1513 stress are often not useful, and can sometimes provide inaccurate and potentially  
1514 harmful information to consumers of medical information.

1515 Despite these concerns, prominent authoritative websites that are grounded in  
1516 research on psychological trauma and PTSD do exist and many public organizations  
1517 and universities have developed online information resources related to post-  
1518 traumatic stress (e.g., National Center for PTSD site: [www.ncptsd.va.gov](http://www.ncptsd.va.gov); Center  
1519 for the Study of Traumatic Stress, <http://www.centerforthestudyoftraumaticstress.org/>;  
1520 International Society for Traumatic Stress Studies, [www.istss.org](http://www.istss.org); National Institute  
1521 of Mental Health, [http://www.nimh.nih.gov/health/topics/post-traumatic-stress-  
1522 disorder-ptsd/index.shtml](http://www.nimh.nih.gov/health/topics/post-traumatic-stress-disorder-ptsd/index.shtml)).

1523 Patients and family members should be warned that information about PTSD that is  
1524 obtained from the Internet should be interpreted with caution. Internet sites from  
1525 established health care agencies or patient advocacy organizations are recommended  
1526 over chat rooms or nonspecialist or commercial sites.

1527 Several randomized controlled trials (RCTs) of web-based intervention treatment of  
1528 PTSD have been conducted. Taken together, they provide preliminary support for the  
1529 use of specific web-based CBT approaches. RCTs of web-based therapist-assisted  
1530 interventions (Knauvelsrud & Maercker, 2007; Lange, et al., 2001; Lange et al.,  
1531 2003; Litz, Engel, Bryant, & Papa, 2007) have demonstrated significant  
1532 improvements in trauma-related symptoms compared to wait-list and supportive  
1533 counseling control conditions, with improvements being maintained over short-term  
1534 (i.e., 3 month) follow-up periods. The studies have focused largely on traumatized  
1535 individuals (clinical and non-clinical, such as university students) with generalized  
1536 distress or subclinical PTSD symptoms. Only one of the RCTs selected patients  
1537 based on a PTSD diagnosis. This study involved service members with PTSD related  
1538 to the Pentagon attack of September 11, 2001 or combat in Iraq or Afghanistan (OEF  
1539 & OIF) (Litz et al., 2007). However, the definition of PTSD was not based on a  
1540 standard structured clinical interview, and no difference was found in the intent-to-  
1541 treat analysis between the internet-based CBT and internet-based supportive  
1542 therapy control. In addition, this was not a pure internet-based intervention, as it  
1543 involved a 2 hour initial face-to-face session in addition to periodic telephone  
1544 contact. A meta-analysis of Internet interventions for anxiety (Reger & Gahm,  
1545 2009) found that the effect Sizes for PTSD symptoms fell in the large range (ES =  
1546 .75; CI = .49 to 1.01), but again, this was not based on studies of patients with  
1547 PTSD per se, but rather persons who have sustained trauma and who have distress,  
1548 subclinical PTSD, or in some cases actual PTSD. Reger and Gahm (2009) also noted  
1549 many methodological problems with current studies and indicated that additional  
1550 research is needed to determine evidence for effectiveness.

1551 In conclusion, there is insufficient evidence to recommend web-based interventions  
1552 for treatment of PTSD. The use of the Internet may have relevance as adjunctive  
1553 modalities in assisting distressed traumatized individuals and complement other  
1554 evidence based treatment interventions.

1555 As with face-to-face treatments, it is important to recognize that existing studies  
1556 have looked at the effectiveness of specific web-based protocols. Thus, it cannot be  
1557 inferred that the studied modalities are generalizable to other web-based treatments.  
1558 Three of the studies cited above relate to one intervention entitled Interapy (Lange,  
1559 et al., 2001; Lange et al., 2003; Knauvelsrud & Maercker, 2007). Interapy and  
1560 DeStress (Litz et al., 2007) share several intervention components, including  
1561 repeated writing about the traumatic experience. These evidence-supported web-  
1562 based protocols are also therapist-assisted, with significant input from the provider.  
1563 For example, Interapy involves a mean per-patient total of 14 hours of therapist  
1564 time. Evidence from research on other mental health problems indicates that rates of  
1565 attrition from web-based interventions are high in the absence of provider contact to  
1566 facilitate completion. With regard to PTSD, there is relatively little evidence at  
1567 present for the effectiveness of Internet interventions that are completely self-  
1568 administered (e.g., Hirai & Clum, 2005).

1569 Regardless of intervention mode, it is important that those involved in technology-  
1570 assisted intervention delivery take steps to ensure that their work complies with the  
1571 regulations and procedures of the organization in which they are employed, with  
1572 evolving legal standards, and with the ethical standards of their professions. Newly  
1573 developed technologies can present significant challenges related to patient  
1574 confidentiality and safety, and these must be addressed carefully by both the  
1575 individual providers and the organization delivering these interventions.

### 1576 C. PHARMACOTHERAPY FOR PTSD

1577 There is growing evidence that PTSD is characterized by specific psychobiological  
1578 dysfunctions, which have contributed to a growing interest in the use of medications  
1579 to treat trauma-related biological effects (see Table I-6).

1580 Studies of medication classes used in therapy for PTSD in individuals exposed to  
1581 trauma that assessed clinical outcomes were included in the review for this guideline  
1582 update. Evidence from randomized controlled trials (RCTs) was considered to be of  
1583 highest quality, followed by observational evidence. Other sources were evaluated  
1584 when randomized controlled trials and observational studies were not available or did  
1585 not provide adequate evidence. Studies were excluded if they did not evaluate  
1586 response to pharmacotherapy and if they did not evaluate individuals exposed to  
1587 trauma. The recommendations and tables address only drugs that have been  
1588 studied and are available in the U.S. Other drugs that have not been reported in  
1589 published studies or were tested in open label trials have not been considered and  
1590 therefore do not appear in the table.

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**Table I - 6. Summary Table – Drug Therapy for PTSD**

| SR       | Significant Benefit | Some Benefit  | Unknown   | No Benefit  |
|----------|---------------------|---|---|---|
| <b>A</b> | SSRIs<br>SNRIs      |   |   |   |
| <b>B</b> |                     | Atypical Antipsychotics<br>(adjunctive)<br>TCAs<br>MAOIs (phenelzine)<br>Mirtazapine<br>Nefazodone ** |   |   |
| <b>C</b> |                     |   | Prazosin  |   |
| <b>I</b> |                     |   | Buspirone<br>Non-benzodiazepine<br>hypnotics<br>Bupropion<br>Trazodone (adjunctive) | Gabapentin<br>Guanfacine<br>Lamotrigine<br>Topiramate<br>Valproate<br>Tiagabine<br>Benzodiazepines ** |
| <b>D</b> |                     |   |   | Conventional<br>Antipsychotics **   |

1594 SR\_ = level of recommendation; \*\* Potential harm

1595 **RECOMMENDATIONS**1596 **General Recommendations:**

- 1597 1. Risks and benefits of long-term pharmacotherapy should be discussed prior to  
1598 starting medication and should be a continued discussion item during  
1599 treatment.
- 1600 2. Monotherapy therapeutic trial should be optimized before proceeding to  
1601 subsequent strategies by monitoring outcomes, maximizing dosage  
1602 (medication or psychotherapy), and allowing sufficient response time (For at  
1603 least 8 weeks). [C]
- 1604 3. if there is some response and patient is tolerating the drug, continue for at  
1605 least another 4 weeks.
- 1606 4. If the drug is not tolerated discontinue the current agent and switch to  
1607 another effective medication.
- 1608 5. If no improvement is observed at 8 weeks consider;
- 1609 6. Increasing the dose of the initial drug to maximum tolerated
- 1610 7. Discontinuing the current agent and switching to another effective medication
- 1611 8. Augmenting with additional agents.
- 1612 9. Recommend assessment of adherence to medication at each visit.
- 1613 10. Recommend assessment of side effects and management to minimize or  
1614 alleviate adverse effects.

- 1615 11. Assess for treatment burden (e.g., medication adverse effects, attending  
1616 appointments) after initiating or changing treatment, when the patient is non-  
1617 adherent to treatment, or when the patient is not responding to treatment.
- 1618 12. Since PTSD is a chronic disorder, responders to pharmacotherapy may need  
1619 to continue medication indefinitely; however it is recommended that  
1620 maintenance treatment should be periodically reassessed.
- 1621 13. Providers should give simple educational messages regarding antidepressant  
1622 use (e.g., take daily, understand gradual nature of benefits, continue even  
1623 when feeling better, medication may cause some transient side effects, along  
1624 with specific instructions on how to address issues or concerns and when to  
1625 contact the provider) in order to increase adherence to treatment in the acute  
1626 phase of treatment. [B]

#### 1627 **Monotherapy:**

- 1628 14. Strongly recommend selective serotonin reuptake inhibitors (SSRIs), for  
1629 which fluoxetine, paroxetine or sertraline have the strongest support, or  
1630 serotonin norepinephrine reuptake inhibitors (SNRIs), for which venlafaxine  
1631 has the strongest support, for the treatment of PTSD. [A]
- 1632 15. Recommend mirtazapine, nefazodone, tricyclic antidepressants (TCAs),  
1633 amitriptyline and imipramine, and monoamine oxidase inhibitors (MAOIs),  
1634 phenelzine, for treatments for PTSD. [B]
- 1635 16. There is insufficient evidence to support the use of prazosin as monotherapy  
1636 in the management of PTSD. [I]
- 1637 17. The existing evidence does not support the use of bupropion, buspirone,  
1638 trazodone, guanfacine, anticonvulsants (tiagabine, lamotrigine, topiramate,  
1639 valproate, gabapentin), or atypical or atypical antipsychotics as monotherapy  
1640 in the management of PTSD. [D]
- 1641 18. There is evidence against the use of benzodiazepines in the management of  
1642 PTSD. [D]

#### 1643 **Augmented Therapy for PTSD:**

- 1644 19. Recommend atypical antipsychotics as adjunctive therapy - (risperidone and  
1645 olanzapine [B] Quetiapine [C]).
- 1646 20. There is insufficient evidence to recommend a sympatholytic, or an  
1647 anticonvulsant as an adjunctive therapy for the treatment of PTSD. [I]

#### 1648 **DISCUSSION**

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#### 1649 **Treatment of PTSD core symptoms**

##### 1650 **Antidepressants**

1651 Antidepressants, particularly serotonergic reuptake inhibitors (SSRIs) have proved  
1652 effective in treating PTSD, and are recommended as first-line agents in treatment  
1653 guidelines (Davidson et al., 2001; Brady et al., 2000; Foa et al., 2000; Foa et al.,  
1654 1999). Over 3000 patients have participated in studies of paroxetine, sertraline and  
1655 fluoxetine. Sertraline and paroxetine have FDA approval for PTSD. SSRIs have a  
1656 broad spectrum of action effectively reducing all three core symptoms of PTSD. As a  
1657 class they are generally well tolerated.

1658 Results with SSRIs are conflicting with respect to wartime related PTSD. Martenyi et  
1659 al. (2002) with combat veterans of recent wars found fluoxetine to be significantly

1660 superior to placebo. Martenyi (2007) reported a negative fixed-dose trial with  
1661 fluoxetine. In addition, Friedman et al. (2007) testing Vietnam vets with chronic  
1662 PTSD in a VA Hospital setting observed no difference between sertraline and placebo.  
1663 One should not extrapolate the findings of Friedman's paper to all veterans as  
1664 veterans with chronic PTSD who remain symptomatic after decades of VA treatment  
1665 comprise a chronic treatment refractory cohort that is not representative of all male  
1666 combat veterans with PTSD.

1667 The SSRIs Citalopram, escitalopram and fluvoxamine have been not been studied  
1668 sufficiently to warrant a recommendation.

1669 Venlafaxine, a SNRI, has been shown to have positive results in two trials of more  
1670 than 800 participants with noncombat-related PTSD (Davidson, 2006a, 2006b).  
1671 Duloxetine and desvenlafaxine have not been studied and cannot be recommended  
1672 at this time. In a 24-week comparison trial, venlafaxine performed as well as  
1673 sertraline in civilian population (Davidson, 2006b).

1674 Other monotherapy recommendations are mirtazapine, nefazodone, TCAs, and  
1675 MAOIs although these have been studied in fewer patients and are considered  
1676 second-line treatment options. Of the TCAs, only amitriptyline and imipramine have  
1677 demonstrated positive outcomes, while data on desipramine and nortriptyline have  
1678 been negative, poor quality studies. Nefazodone has been the subject of several  
1679 small- to mid-sized RCTs and case-control studies (Davis et al., 2000; Garfield et al.,  
1680 2001; Gillin et al., 2001; Hertzberg et al., 1998; Hidalgo et al., 1999; Zisook et al.,  
1681 2000). In all six studies, the drug was helpful in improving CAPS, HAM-D, sleep, and  
1682 anxiety. In a trial of combat and sexual assault origin PTSD, nefazodone was more  
1683 effective than placebo (Davis, 2004). Nefazodone has demonstrated efficacy  
1684 equivalent to sertraline in two fair-quality trials (McRae, 2004; Saygin, 2002). Two  
1685 trials with mirtazepine (Davidson, 2003; Chung, 2004) have demonstrated positive  
1686 findings. However, in the placebo controlled trial (Davidson, 2003) both mirtazepine  
1687 and placebo had large effect sizes. In a trial of military veterans, mirtazepine was as  
1688 efficacious as sertraline but there was no placebo comparison arm (Chung, 2004). Of  
1689 the currently available MOAIs, only phenelzine has been studied. In a placebo  
1690 comparison trial, Vietnam veterans assigned to phenelzine had significant  
1691 improvement in IES compared to placebo (Kosten, 1991).

1692 Although **atypical antipsychotics** are not effective as monotherapy, significant  
1693 efficacy as adjunctive treatment to antidepressants has been shown in trials  
1694 composed primarily of veterans. Response was predominantly in hyperarousal and  
1695 re-experiencing symptom clusters. These trials studied risperidone and olanzapine  
1696 with one small quetiapine trial. Only one trial has addressed their role in co-morbid  
1697 psychosis (Hamner, 2003). A significant improvement in psychotic symptoms  
1698 (change in PANS) was found in risperidone treated Veterans compared to placebo;  
1699 both groups improved significantly in their CAPS scores. In an open-label trial,  
1700 quetiapine improved both PANS and CAPS scores compared to baseline (Hamner  
1701 2003 different from above). Olanzapine as an adjunctive treatment improved CAPS  
1702 scores and sleep quality compared to placebo in a small 8-week trial (Stein M,  
1703 2002). No data are available on the use of typical antipsychotics in the treatment of  
1704 PTSD.

1705 The existing evidence does not support the use of **anticonvulsants** as monotherapy  
1706 in the management of PTSD. Tiagabine has been compared to placebo in two RCTs  
1707 with no difference in response (Connor 2006; Davidson 2007). Valproate, as  
1708 monotherapy, did not differ from placebo in one RCT (Davis, 2008) Anticonvulsants  
1709 are frequently used as adjunctive treatments. Only topiramate has been studied in

1710 this role in Veterans with negative results (Lindley, 2007). Data on other  
1711 anticonvulsants are insufficient to recommend their use in PTSD. A meta analysis  
1712 showed useful use of valproate in PTSD (Adamou, 2007)

1713 The Cochrane Collaboration published a review of the evidence regarding  
1714 pharmacological treatments in PTSD (Stein et al., 2006). They found 35 short-term  
1715 RCTs of PTSD (4597 participants) to review, three of which contained a maintenance  
1716 component; five of those were unpublished. The authors concluded that, while no  
1717 clear evidence exists to show that any particular class of medication is more effective  
1718 or better tolerated than any other, the greatest number of trials showing efficacy to  
1719 date, as well as the largest, have been with the SSRIs. On the basis of the data, the  
1720 review recommends the SSRIs as first line agents in the pharmacotherapy of PTSD,  
1721 and supports their value in long-term treatment.

1722 A meta-analysis of 4 RCTs that compared SSRIs to placebo without regard to  
1723 diagnostic criteria, duration, severity, or co-morbid diagnoses reported that  
1724 treatment favored the drug in all 4 trials; however, only one study (with 183  
1725 subjects) reached statistical significance. Two RCTs maintained treatment with an  
1726 SSRI for 64 weeks and 40 weeks, respectively. One study reported that 50% of  
1727 patients experienced worsening symptoms when placebo was substituted for active  
1728 drug and in the second report patients on placebo were 6.4 times more likely to  
1729 relapse compared to the drug group. Although some patients may respond to an  
1730 antidepressant trial within 3 months, some patients may require more than 12 weeks  
1731 to respond to SSRIs (Martenyi et al., 2002).

### 1732 **Benzodiazepines**

1733 Benzodiazepines are widely used for symptomatic control of insomnia, anxiety, and  
1734 irritability, there is no evidence they reduce the core symptoms (e.g., syndromal  
1735 symptoms) of PTSD, such as avoidance or dissociation (Friedman and Southwick,  
1736 1995; Viola et al., 1997). More recent studies have been scarce, and Kosten et al.  
1737 (2000) presents evidence that does not support the use of benzodiazepines in PTSD.

1738 Benzodiazepine administration should be discouraged both in acute stress disorder  
1739 (ASD) and post-traumatic stress disorder (PTSD), due to lack of evidence for  
1740 effectiveness and risks that outweigh potential benefits. Although benzodiazepines  
1741 have been frequently used "as needed" and continuously for anxiety disorders,  
1742 including to augment evidence-based treatment modalities in PTSD, there is  
1743 evidence to suggest that benzodiazepines may actually potentiate the acquisition of  
1744 fear responses and worsen recovery from trauma. Benzodiazepines use should be  
1745 considered relatively contraindicated in combat veterans with PTSD, because of the  
1746 very high co-morbidity of combat-related PTSD with alcohol misuse and substance  
1747 use disorders (upwards of 50% of comorbidity) and potential problems with  
1748 tolerance and dependence. Once initiated in combat veterans, benzodiazepines can  
1749 be very difficult, if not impossible, to discontinue, due to significant withdrawal  
1750 symptoms compounded by the underlying PTSD symptoms.

1751 Although there have been few clinical trials of benzodiazepines to prevent or treat  
1752 PTSD, they have been consistent in showing negative findings.

### 1753 **Clinical Trials**

- 1754 • Braun, et al. 1990 - In a randomized double blind cross-over study, alprazolam  
1755 showed no significant benefit in alleviating PTSD symptoms compared with  
1756 placebo. A slight reduction in anxiety symptoms was offset by withdrawal effects  
1757 documented after only five weeks of treatment.



- 1758 • Gelpin, et al. 1996 - In this open labeled study, 13 patients who had recently  
1759 experienced trauma (within the past 18 days) and were experiencing excessive  
1760 distress (panic, agitation, or persistent insomnia) were treated for up to 6  
1761 months with alprazolam or clonazepam. These 13 patients were compared with a  
1762 control group of recently traumatized individuals matched for demographics and  
1763 symptoms (using the Impact of Events Scale). On follow-up, PTSD occurred at a  
1764 significantly *higher* rate in the benzodiazepine treated group (9/13, 69%) than in  
1765 the control group (2/13, 15%). Although the strength of the evidence is low  
1766 (open labeled study), the study suggested that benzodiazepines may worsen  
1767 outcomes in the acute period following trauma, and the authors referenced  
1768 animal data (see below) consistent with the hypothesis that benzodiazepines may  
1769 potentiate the acquisition of fear responses.
- 1770 • Melman, et al. 2002 - In this small double-blind randomized controlled study  
1771 during the acute period after trauma (mean 2 weeks after trauma), the short  
1772 term (7 day) evening use of temazepam in patients with significant ASD/PTSD  
1773 symptoms was compared with placebo (11 patients in each group). The study  
1774 showed no benefits in preventing PTSD, and the trend was similar to the Gelpin  
1775 study, with 6 of 11 (55%) patients who received tamazepam developing PTSD,  
1776 compared with 3/11 (27%) who received placebo.
- 1777 • Cates, et al. 2004 - This small single blind cross-over placebo controlled study  
1778 compared clonazepam with placebo for the treatment of sleep dysfunction  
1779 associated with combat-related PTSD. The study showed no significant difference  
1780 between the benzodiazepine and placebo treatment.

### 1781 **Other Relevant Studies**

- 1782 • Viola et al. (1997) - At Tripler Army Medical Center, after having treated 632  
1783 patients, the vast majority of whom suffered from combat-related PTSD, between  
1784 1990 and 1996, the staff began to "explore treatment alternatives" to  
1785 benzodiazepines due to the "risks attendant to benzodiazepine management of  
1786 PTSD, coupled with poor clinical outcome" (Viola et al., 1997).
- 1787 • Risse et al. (1990) - This case series reflects the typical clinical experience when  
1788 benzodiazapines are utilized for treating combat-related PTSD. In this study  
1789 alprazolam was used to augment treatment of anxiety symptoms in 8 combat  
1790 veterans with chronic PTSD and co-morbid conditions (mostly alcohol misuse).  
1791 Although anxiety initially improved with treatment, the improvement was short  
1792 lived and resulted in tolerance to increasing doses and eventual failure of the  
1793 treatment. The key problem was encountered upon attempting to gradually  
1794 withdraw the medication after determining that ongoing treatment was not going  
1795 to be of further benefit. All 8 patients experienced severe reactions, including  
1796 anxiety, sleep disturbance, rage, hyperalertness, increased nightmares, and  
1797 intrusive thoughts; 6 of the 8 veterans developed a level of rage with homicidal  
1798 ideation that they had never encountered previously.

- 1799 • Davydow (2008) - In this literature review of the risk factors for developing PTSD  
1800 after serious trauma (requiring ICU treatment), greater ICU benzodiazepine  
1801 administration was found to be one of the consistent predictors of PTSD.
- 1802 • Randal, et al. (1995); and Coupland, et al. (1997) - Flumazenil, a  
1803 benzodiazepine/GABA receptor antagonist, provokes panic attacks in patients  
1804 with panic disorder, but not in healthy controls. In these two studies (one of  
1805 which involved a group of male Vietnam combat veterans), flumazenil was  
1806 compared with placebo to determine if it provoked anxiety, panic, or PTSD  
1807 symptoms. Both studies showed that there were no significant increases in  
1808 anxiety, panic, or PTSD symptoms in subjects as a result of flumazenil  
1809 administration. This suggests that PTSD is dissimilar to panic disorder in terms  
1810 of benzodiazepine receptor functioning, and helps to explain why benzodiazepine  
1811 treatment has produced no significant benefits in clinical trials.
- 1812 • Several studies involving different animal models of PTSD (for example, Matar, et  
1813 al., 2009 and Hebert, et al., 1996) have shown that benzodiazepine  
1814 administration in the immediate aftermath of stress exposure significantly  
1815 increases vulnerability to develop more severe responses upon subsequent  
1816 exposure to stress.

|   | Evidence  | Source   | LE                  | QE        | NB            | SR |
|---|---|--|---------------------|-----------|---------------|----|
| 1 | Benzodiazepines are not recommended for treatment of PTSD | Braun et al., 1990<br>Cates et al., 2004<br>Gelpin et al., 1996<br>Melman et al., 2002 | I<br>I<br>II-2<br>I | Fair/Poor | Small/<br>Neg | D  |

1817 LE =Level of Evidence; QE = Quality of Evidence; SR = Strength of Recommendation; NB=Net benefit (see  
1818 Appendix A)

### 1819 **Symptom-specific adjunctive treatments**

1820 A number of trials have shown the efficacy of certain agents to target specific PTSD  
1821 symptoms.

1822 **Prazosin** as a global treatment for PTSD has yielded mixed results, it has shown  
1823 consistent efficacy in improving sleep and reducing nightmares. In five relatively  
1824 small studies (Raskind et al., 2002,2003 and 2007; and Taylor 2006,2008), prazosin  
1825 has demonstrated a value in reducing nightmares and in improving CAPS, CGI, and  
1826 CGIC scores. (See discussion in [Module I-3 A. Sleep Disturbance](#))

1827 Other agents that have improved insomnia are trazodone, mirtazepine, and  
1828 olanzapine. There are no trials of non-benzodiazepine hypnotics in the treatment of  
1829 sleep disorders associated with PTSD.

1830 **Sympatholytics** have also been investigated as PTSD therapy.

1831 **Buspirone**, a nonbenzodiazepine antianxiety drug, is reported to have “clinical  
1832 efficacy” in two very small studies (Duffy & Malloy, 1994; Wells et al., 1991).

1833

**Table I - 1 Pharmacologic Studies for Treatment of PTSD**

| Drug         | Source of Evidence          | Result   | n   | LE     | QE | NB   | SR |
|--------------|-----------------------------|--|-----|--------|----|------|----|
| <b>SSRI</b>  |                             |  |     |        |    |      |    |
| Sertraline * | Brady et al., 2000          | Significant improvement, CAPS-2, CGI   | 187 | I      | G  | Sub  | A  |
|              | Davidson et al., 2001a      | Effective for prevention of PTSD relapse   | 96  | I      | G  | Sub  |    |
|              | Davidson et al., 2001b      | Significant responder rate, CAPS-2   | 208 | I      | G  | Sub  |    |
|              | Davidson et al., 2002       | Study of effect on individual symptoms   | ?   | II-2   | F  | Mod  |    |
|              | Davidson et al., 2001c      | Effective for preventing PTSD relapse  | 96  | I      | G  | Sub  |    |
|              | Friedman et al., 2007       | No sig diff between Tx and Pbo. Combat trauma.   | 169 | I      | G  | Zero |    |
|              | Londborg et al., 2001       | Significant response maintained x 36 weeks   | 128 | II-1   | G  | Sub  |    |
|              | Rapaport et al., 2002       | Significant response maintained x 64 weeks   | 359 | I/II-1 | G  | Sub  |    |
|              | Tucker et al., 2003         | Significant improvement of primary outcome   | 58  | I      | F  | Zero |    |
|              | Zohar et al., 2002          | Numeric advantage (only), not sig, Israeli vets  | 42  | I      | G  | Zero |    |
| Paroxetine * | Marshal 2007                | Significant improvement, CAPS-2 & CGI  | 52  | I      | G  | Sub  | A  |
|              | Marshall, et al., 2001      | Significant improvement, CAPS-2 & CGI  | 551 | I      | G  | Sub  |    |
|              | Tucker, et al., 2001        | Significant improvement, CAPS-2  | 307 | I      | G  | Sub  |    |
| Fluoxetine   | Barnett et al., 2002        | Study of tolerability. Well-tolerated  | 65  | I      | G  | Sub  | A  |
|              | Connor et al., 1999         | “Superior” response for civilian patients  | 53  | I      | G  |      |    |
|              | Davidson 2005               | Well-tolerated and effective in prevention of relapse, improve CGI. High rate of relapse | 57  | I      | G  | Sub  |    |
|              | Martenyi 2007               | No significant dif. between Tx and Pbo; TOP-8  | 411 | I      | G  | Zero |    |
|              | Martenyi et al., 2002a      | Effective for prevention of PTSD relapse; 50% subjects - combat related                  | 131 | I      | G  | Sub  |    |
|              | Martenyi et al., 2002b      | Effective: improvement in TOP-8, CGI; 50% subjects -combat related                       | 301 | I      | G  | Sub  |    |
|              | Meltzer-Brody, et al., 2000 | Reduced all symptom clusters of PTSD;  | 53  | II-2   | F  |      |    |
|              | Van der ko et al., 1994     | Fluoxetine > placebo, more in non VA pts   | 64  | I      | G  | Sub  |    |
| Citalopram   | Seedat et al., 2000         | Significant improvement, CAPS-2  | 14  | II-1   | F  |      |    |
|              | Tucker et al, 2003          | Sig improvement (↓ BP)   | 58  | I      | F  | Neg  |    |
| Fluvoxamine  | Escalona et al., 2002       | Appears to improve PTSD symptoms   | 15  | III    | P  | -    | I  |
|              | Neylan et al., 2001         | Improved sleep quality for Vietnam vets  | 21  | III    | P  | -    |    |

| Drug                | Source of Evidence           | Result   | n    | LE   | QE | NB  | SR |
|---------------------|------------------------------|--|------|------|----|-----|----|
| <b>TCA</b>          |                              |  |      |      |    |     |    |
| Amitriptyline       | Davidson et al., 1990        | Effective for core symptoms of PTSD  | 46   | I    | G  | Mod | B  |
|                     | Davidson et al., 1993        | Significant improvement: IES, CGI, HAM-D   | 62   | I    | G  |     |    |
| Desipramine         | Reist et al., 1989           | Did not show efficacy; no statistics   | 27   | III  | P  |     |    |
| Imipramine          | Kosten et al., 1991          | Significant improvement, CAPS-2, IES   | 41   | I    | G  |     |    |
| Nortriptyline       | Zygmunt et al., 1998         | Effective for traumatic grief symptoms   | 22   | II-1 | G  |     |    |
|                     | Dow et al., 1997             | Improvement in CGE for PTSD with MDD   | 72   | II-2 | F  |     |    |
| <b>MAOI/RIMA</b>    |                              |  |      |      |    |     |    |
| Phenelzine          | Kosten et al., 1991          | Significant improvement in IES, better than placebo                                      | 37   | I    | G  | Mod | B  |
| <b>SNRI</b>         |                              |  |      |      |    |     |    |
| Venlafaxine         | Davidson et al, 2006;        | Effective in tx PTSD; Improves resilience  | 329  | I    | G  | Sub | A  |
|                     | Davidson et al., 2006b       | Effective similar to sertraline  | 531  | I    | G  | Sub |    |
| <b>Secondary AD</b> |                              |  |      |      |    |     |    |
| Bupropion           | Becker et al., 2007          | Bupropion SR had no effect on PTSD   | 30   | I    | F  | Neg | I  |
|                     | Canive et al., 1998          | No change in total CAPS score - male veterans  | 17   | II-2 | F  |     |    |
| Nefazodone          | Davis et al, 2004            | Nefazodone is effective and well-tolerated ; Combat sexual                               | 42   | I    | G  | Sub | B  |
|                     | Davis et al., 2000           | Significant improvement in CAPS, HAM-D   | 36   | II-2 | G  |     |    |
|                     | Garfield et al., 2001        | Significant improvement in CAPS, anxiety   | 14   | II-2 | F  |     |    |
|                     | Gillin et al., 2001          | Significant improvement in sleep, CAPS   | 12   | II-2 | F  |     |    |
|                     | Hidalgo et al., 1999         | High response rate; pooled data, 6 studies   | 105  | II-2 | F  |     |    |
|                     | McRae et al., 2004           | Nefazodone is effective as sertraline. High attrition rates                              | 37   | I    | F  | Sub |    |
|                     | Saygin et al., 2002          | Nefazodone is effective as sertraline and well-tolerated<br>Earthquake survivors         | 54   | I    | F  | Sub |    |
| Zisook et at., 2000 | PTSD symptoms lessened, CAPS | 19   | II-2 | F    |    |     |    |
| Trazodone           | Warner et al., 2001          | Reduction in nightmares; 9 reports priapism  | 74   | III  | P  |     | I  |
| Mirtazapine         | Chung et al, 2004            | Mirtazapine is effective as sertraline and well tolerated.<br>Military veterans (Korean) | 51   | II-1 | F  | Mod | B  |
|                     | Davidson et al., 2003        | Significant improvement in the SPRINT, SIP, DTS as compared to placebo                   | 26   | I    | F  | Mod | B  |

| Drug                           | Source of Evidence       | Result  | n   | LE   | QE | NB    | SR |
|--------------------------------|--------------------------|---|-----|------|----|-------|----|
| <b>Anticonvulsants</b>         |                          |   |     |      |    |       |    |
| Gabapentin                     | Hamner et al., 2001      | Effective for insomnia, adjunct treatment   | 30  | II-2 | P  | Zero  | I  |
| Lamotrigine                    | Hertzberg et al., 1999   | Promising results   | 14  | I    | P  | Zero  | I  |
| Topiramate                     | Lindley SE 2007          | No significant effect for topiramate over placebo   | 40  | I    | F  | Zero  | C  |
|                                | Tucker P, 2007           | Not significant difference from placebo (non-combat)  | 38  | I    | G  | Zero  |    |
| Valproate                      | Davis L 2008             | Divalproex monotherapy was not effective in the treatment of chronic PTSD                           | 85  | I    | G  | Zero  | C  |
|                                | Adamou 2007 (SR)         | Valproate was generally effective in reducing hyperarousal, improving irritability and anger        | 63  | I    | F  | Small |    |
| Tiagabine                      | Connor K 2006            | No significant improvement was observed on all outcome measures                                     | 29  | I    | F  | Zero  | ?  |
|                                | Davidson J 2007          | No difference from placebo  | 232 | I    | G  | Zero  |    |
| <b>Atypical Antipsychotics</b> |                          |   |     |      |    |       |    |
| Olanzapine                     | Butterfield et al., 2001 | No beneficial effect. High placebo response   | 15  | I    | G  |       | I  |
|                                | Petty et al., 2001       | Significant improvement in CAPS, CGI  | 48  | II-1 | G  |       |    |
|                                | Stein et al., 2002       | Adjunct to SSRI. Significant improvement in measures, but not in global response                    | 19  | I    | F  | Mod   |    |
| Quetiapine                     | Hamner et al., 2003      | Significant improvement in CAPS   | 20  | II-1 | F  |       | I  |
| Risperidone                    | Bartzokis 2005           | Risperidone>placebo; Vet military   | 48  | I    | G  | Small | C  |
|                                | Hamner et al., 2003      | Adjunct to other meds, comorbid psychoses   | 40  | I    | F  | Neg   |    |
|                                | Monnelly 2003            | Risperidone > placebo; Military combat  | 15  | I    | F  | Mod   |    |
|                                | Padala P 2006            | Sexual assault - risperidone monotherapy was more effective than placebo                            | 20  | I    | P  | Small |    |
|                                | Reich D 2004             | Child abuse - Risperidone more effective than placebo;  | 21  | I    | F  | Mod   |    |
|                                | Rothbaum 2008            | Civilian - Risperidone augmentation was helpful in subjects who did not remit with sertraline alone | 45  | I    | F  | Small |    |

| Drug            | Source of Evidence        | Result  | n   | LE   | QE   | NB    | SR |
|-----------------|---------------------------|---|-----|------|------|-------|----|
| Sympatholytics  |                           |   |     |      |      |       |    |
| Clonidine       | Kinzie & Leung, 1989      | Cambodian refugees improved, dual therapy   | 68  | III  | P    | Small | I  |
| Guanfacine      | Horrigan & Barnhill, 1996 | suppression of PTSD associated nightmares in children                                       | 1   | III  | P    | Small | D  |
|                 | Neylan T, 2006            | No effect on PTSD symptoms (in vet military)  | 63  | I    | Good | Neg   |    |
|                 | Davis I 2008              | No effect on PTSD symptoms  | 65  | I    | Good | Neg   |    |
| Prazosin        | Raskind et al., 2003      | Significant improvement, CAPS, CGI  | 10  | I    | F    | Small | B  |
|                 | Raskind et al., 2002      | Significant improvement in dream scores   | 59  | II-2 | F    | Zero  |    |
|                 | Raskind et al., 2007      | Significant improved sleep quality, reduced nightmares, better overall sense of well being. | 34  | I    | G    | Mod   |    |
|                 | Taylor et al., 2006       | Reduction in global PTSD illness severity   | 11  | II   | P    | Mod   |    |
|                 | Taylor et al., 2008       | Significantly improved CGI-I scores; and changed PDRS scores toward normal dreaming         | 13  | I    | F    | Mod   |    |
| Benzodiazepines |                           |   |     |      |      |       |    |
| Benzodiaz.      | Kosten et al., 2000       | Not associated with adverse outcomes  | 370 | II-2 | F    | Zero  | D  |
| Alprazolam      | Braun et al., 1990        | Did not show efficacy.<br>(concern: rebound anxiety)  | 16  | I    | F    | Zero  | D  |
| Clonazepam      | Fossey & Hamner, 1994     | A source of sexual dysfunction  | 42  | III  | P    | Zero  | D  |
|                 | Gelpin et al., 1996       | No beneficial effect in PTSD, may worsen outcome  | 20  | II-1 | F    | Zero  |    |
|                 | Shalev & Rogel 1992       | No effect on auditory startle   | N/A | III  | F    | Zero  |    |
| Temazepam       | Melman et al., 2002       | No benefits in preventing PTSD, May worsen outcome  | 11  | I    | P    | Zero  | D  |
|                 | Cares et al., 2004        | No difference between the benzodiazepine and placebo  | I   | I    | P    | Zero  | D  |

LE=Level of Evidence QE=Quality of Evidence (F=Fair; G=Good; P=Poor); NB=Net benefit (Sub=Substantial; Mod = Moderate; Neg=Negative)  
SR= Strength of Recommendation (see Appendix A);

\* FDA Approved

\*\* Trials added since the 2004 VA/DoD PTS guideline

**Table I - 2 Symptoms Response by Drug Class and Individual Drug (based on controlled trials)**

|                                |                              | Global Improvement | Re-experiencing (B) | Avoidance/ Numbing (C) | Hyper-arousal (D) |
|--------------------------------|------------------------------|--------------------|---------------------|------------------------|-------------------|
| <b>SSRI</b>                    | Fluoxetine                   | X                  | X                   | X                      | X                 |
|                                | Sertraline                   | X                  | X                   | X                      | X                 |
|                                | Paroxetine                   | X                  | X                   | X                      | X                 |
| <b>SNRI</b>                    | Venlafaxine                  | X                  | X                   | X                      | X                 |
| <b>TCAs</b>                    | Amitriptyline/<br>Imipramine | X                  | X                   |                        | X                 |
| <b>MAOIs</b>                   | Phenelzine                   | X                  | X                   |                        | X                 |
| <b>Sympatholytics</b>          | Prazosin                     | X                  | X                   |                        | X                 |
| <b>Other Antidepressants</b>   | Mirtazapine                  | X                  | X                   |                        | X                 |
|                                | Nefazodone                   | X                  | X                   |                        | X                 |
| <b>Atypical antipsychotics</b> | Risperidone                  |                    | X                   |                        | X                 |
|                                | Olanzapine                   |                    |                     |                        | X                 |

**Table I - 3 Drug Details**

| Agent  | *Oral Dose      | Contraindications  | Adverse Events   | Pregnancy Category  | Remarks   |
|--|-----------------|--|--|---|---|
| <b>Selective Reuptake Serotonin Inhibitors (SSRIs)</b> |                 |  |  |   |   |
| Fluoxetine   | 20 – 60 mg/d    | <b>Contraindications :</b> <ul style="list-style-type: none"> <li>MAO inhibitor within 14 days</li> <li>Concurrent use of pimozone or thioridazine</li> <li>Hypersensitivity</li> </ul>  | <ul style="list-style-type: none"> <li>Nausea , diarrhea</li> <li>Headache</li> <li>Dizziness</li> <li>Sexual dysfunction</li> <li>Hyponatremia/SIADH (Syndrome of Inappropriate Antidiuretic Hormone)</li> <li>Nervousness, anxiety, agitation</li> <li>Serotonin syndrome</li> </ul> | <ul style="list-style-type: none"> <li>All except paroxetine are Category C</li> <li>Paroxetine Category D</li> <li>Women planning to breast feed, consider an antidepressant with the lowest excretion into breast milk: paroxetine, sertraline</li> </ul> | <ul style="list-style-type: none"> <li>Avoid abrupt discontinuation of all except fluoxetine</li> <li>Citalopram, escitalopram and sertraline are less likely to be involved in hepatic enzyme drug interactions involving CYP2D6 or 3A4</li> <li>All except escitalopram are generically available</li> <li>St. Johns Wort may in decrease the concentration of SSRIs metabolized by CYP2D6</li> </ul> |
| Paroxetine   | 20 – 60 mg/d    |  |  |   |   |
| Sertraline   | 50 – 200 mg/d   |  |  |   |   |
| Fluvoxamine  | 50 – 150 mg bid |  |  |   |   |
| Citalopram   | 20 – 60 mg/d    |  |  |   |   |
| Escitalopram   | 10 – 20 mg/d    |  |  |   |   |
| <b>Tricyclic Antidepressants</b>                       |                 |  |  |   |   |
| Imipramine   | 150 – 300 mg/d  | <b>Contraindications :</b> <ul style="list-style-type: none"> <li>Clomipramine – seizure disorder</li> <li>MAOI use within 14 days</li> <li>Acute MI within 3 months</li> <li>Hypersensitivity</li> </ul><br><b>Relative Contraindications:</b> <ul style="list-style-type: none"> <li>Coronary artery disease</li> <li>Prostatic enlargement</li> </ul> | <ul style="list-style-type: none"> <li>Dry mouth</li> <li>Dry eyes</li> <li>Constipation</li> <li>Orthostatic hypotension</li> <li>Increased heart rate</li> <li>Ventricular arrhythmias</li> <li>Weight gain</li> <li>Drowsiness</li> </ul>   | <ul style="list-style-type: none"> <li>Category C</li> <li>Women planning to breast feed, consider an antidepressant with the lowest excretion into breast milk: nortriptyline</li> </ul>   | <ul style="list-style-type: none"> <li>Therapeutic blood concentrations not established for PTSD</li> <li>Desipramine and nortriptyline have lower rate of sedation anticholinergic and hypotensive effects</li> <li>Moderate CYP2D6 inhibition</li> <li>St. Johns Wort may decrease the concentration of SSRIs metabolized by CYP2D6</li> </ul>  |
| Amitriptyline  | 150 – 300 mg/d  |  |  |   |   |
| Desipramine  | 100 – 300 mg/d  |  |  |   |   |
| Nortriptyline  | 50 – 150 mg/d   |  |  |   |   |
| Protriptyline  | 30 – 60 mg/d    |  |  |   |   |
| Clomipramine   | 150 – 250 mg/d  |  |  |   |   |



| Agent                               | *Oral Dose  | Contraindications   | Adverse Events   | Pregnancy Category  | Remarks   |
|-------------------------------------|---|---|--|---|---|
| <b>Monoamine Oxidase Inhibitors</b> |   |   |  |   |   |
| Phenelzine                          | 45-75 mg/d in divided doses   | <b>Contraindications:</b> <ul style="list-style-type: none"> <li>All antidepressants within 14 days of start of a MAOI, except fluoxetine is 5 weeks</li> <li>Concurrent use with CNS stimulants or depressants and decongestants</li> <li>CHF, hepatic or renal disease</li> <li>Pheochromocytoma</li> <li>Foods high in tyramine</li> <li>Hypersensitivity</li> </ul> | <ul style="list-style-type: none"> <li>Hypertensive crisis with drug/tyramine interactions</li> <li>Bradycardia</li> <li>Orthostatic hypotension</li> <li>Insomnia</li> <li>Dry mouth</li> <li>Dry Eyes</li> <li>Constipation</li> </ul> | <ul style="list-style-type: none"> <li>Category C</li> </ul>  | <ul style="list-style-type: none"> <li>Patient must maintain a low tyramine diet and avoid foods rich in tyramine</li> <li>Tranycypromine should be taken early in the day to reduce insomnia</li> <li>MAOIs are to be discontinued 2-weeks prior to starting another antidepressant or serotonergic agent</li> </ul> |
| Tranycypromine                      | 10 – 60 mg/d<br>target 1 mg/kg/d<br>target 0.7 mg/kg/d                                  |   |  |   |   |
| <b>Sympatholytics</b>               |   |   |  |   |   |
| Propranolol                         | 40 - ? mg/d   | <b>Contraindications:</b> <ul style="list-style-type: none"> <li>Sinus bradycardia, uncompensated congestive heart failure, 2<sup>nd</sup> or 3<sup>rd</sup> degree heart block, severe COPD or asthma, hypersensitivity to beta-blockers</li> </ul>  | <ul style="list-style-type: none"> <li>Hypotension, bronchospasm, bradycardia</li> </ul>   | <ul style="list-style-type: none"> <li>Category C &amp; D (2<sup>nd</sup> &amp; 3<sup>rd</sup> trimesters)</li> <li>Breast feeding – not recommended</li> </ul> | <ul style="list-style-type: none"> <li>Has only been used in a single dose for prevention of PTSD</li> </ul>  |
| Prazosin                            | Target 6 – 10 mg/d<br>Start with 1 mg at bedtime and increase as blood pressure allows. | <ul style="list-style-type: none"> <li>Hypersensitivity to quinazolines</li> <li>Concurrent use of phosphodiesterase type-5 inhibitors</li> </ul>   | <ul style="list-style-type: none"> <li>First dose syncope</li> </ul>   | <ul style="list-style-type: none"> <li>Category C</li> <li>Breast feeding – affects unknown</li> </ul>  | <ul style="list-style-type: none"> <li>Primarily used for management of recurrent distressing dreams</li> </ul>   |
| <b>Novel Antidepressants</b>        |   |   |  |   |   |

| Agent       | *Oral Dose     | Contraindications  | Adverse Events  | Pregnancy Category   | Remarks  |
|-------------|----------------|--|---|--|--|
| Bupropion   | 150 – 450 mg/d | <p><b>Contraindications :</b></p> <ul style="list-style-type: none"> <li>MAOI use within 14 days (all)</li> <li>Hypersensitivity</li> </ul> <p>Bupropion<br/>– single doses of regular-release &gt;150 mg/d and total daily dose &gt;450 mg/d.</p> <ul style="list-style-type: none"> <li>History of seizures, anorexia or buliemia</li> <li>Nefazodone</li> <li>Active liver disease or increased liver enzymes</li> <li>Use with carbamazepine, pimozone, cisapride, triazolam and alprazolam</li> </ul> | <ul style="list-style-type: none"> <li>Bupropion: headache, insomnia, dizziness, weight loss, decreased appetite, anxiety, agitation, nervousness, sleep disturbances</li> <li>Nefazodone: hepatotoxicity</li> <li>Trazodone and nefazodone: sedation, rare priapism</li> <li>Venlafaxine: hypertension in patients with pre-existing hypertension, headache, insomnia, somnolence, nervousness, dizziness, anorexia</li> <li>Mirtazapine: weight gain, increase appetite, somnolence, dry mouth</li> </ul> | <ul style="list-style-type: none"> <li>Category C (all)</li> </ul> | <ul style="list-style-type: none"> <li>Need to taper venlafaxine to prevent rebound signs/symptoms</li> <li>The group has a lower rate of sexual dysfunction compared to SSRIs</li> <li>Obtain baseline LFTs when treating with nefazodone</li> <li>Nefazodone is a potent CYP3A4 inhibitor</li> <li>St. Johns Wort may increase mirtazepine's metabolism</li> </ul> |
| Nefazodone  | 300 – 600 mg/d |  |   |  |  |
| Trazodone   | 300 – 600 mg/d |  |   |  |  |
| Venlafaxine | 150 – 375 mg/d |  |   |  |  |
| Mirtazapine | 30 – 60 mg/d   |  |   |  |  |

| Agent                  | *Oral Dose   | Contraindications   | Adverse Events  | Pregnancy Category   | Remarks   |   |
|------------------------|--|---|---|--|---|---|
| <b>Anticonvulsants</b> |  |   |   |  |   |   |
| <b>Carbamazepine</b>   | target 400 – 1600 mg/d   | <b>Contraindications:</b> <ul style="list-style-type: none"> <li>bone marrow suppression, particularly leukopenia</li> <li>hypersensitivity to carbamazepine, pimozide or tricyclic antidepressants</li> <li>MAOI use within 14 days</li> <li>Concurrent use of nefazodone</li> </ul> | <ul style="list-style-type: none"> <li>bone marrow suppression, aplastic anemia, leukopenia, SIADH, drowsiness, ataxia, photosensitivity, serious dermatologic reactions including Stevens-Johnson syndrome, A-V block and bradycardia</li> </ul> | <ul style="list-style-type: none"> <li>Category D</li> <li>Excreted into breast milk in high concentrations; measurable in infant serum.</li> </ul>  | <ul style="list-style-type: none"> <li>Therapeutic blood concentration are not established for PTSD, but monitoring may be useful in cases of suspected toxicity (usual range 4 – 12 mcg/mL)</li> <li>Strong inducer of CYP 1A2, 2B6, 2C8, 2C9, 2C19, and 3A4. Induction can reduce effectiveness of many medication such as oral contraceptives</li> </ul> |   |
| <b>Gabapentin</b>      | target 300 – 3600 mg/d   | <ul style="list-style-type: none"> <li>renal impairment</li> </ul>  | <ul style="list-style-type: none"> <li>sedation, ataxia</li> <li>peripheral edema</li> </ul>  | <ul style="list-style-type: none"> <li>Category C</li> <li>Excreted into breast milk; effects unknown</li> </ul>   |   |   |
| <b>Lamotrigine</b>     | Not taking divalproex or CBZ: 25 mg once a day for 2 weeks, then 50 mg/day for 2 weeks, then 100 mg/day for 1 week | <ul style="list-style-type: none"> <li>increased rash with valproate; max dose of 200 mg</li> </ul>   | <ul style="list-style-type: none"> <li>Stevens- Johnson syndrome</li> <li>fatigue</li> <li>Headache</li> <li>Peripheral edema</li> <li>Rash</li> <li>Vision changes</li> </ul>  | <ul style="list-style-type: none"> <li>Category C</li> <li>Excreted in breast milk with measurable</li> </ul>  |   | Adjust dose base on renal function  |
| <b>Topiramate</b>      | target 200 – 400 mg/d.<br>Start with 25 – 50 mg/d and increase by 15 – 50 mg/week to maximum dose or as tolerated. | <ul style="list-style-type: none"> <li>hepatic impairment</li> </ul>  | <ul style="list-style-type: none"> <li>angle closure glaucoma</li> <li>sedation</li> <li>dizziness</li> <li>ataxia</li> <li>cognitive impairment</li> <li>weight loss</li> <li>paresthesia</li> <li>vision changes</li> </ul>                     | <ul style="list-style-type: none"> <li>Category C</li> <li>Excreted in to breast milk; breast feeding not recommended</li> </ul>   |   | <ul style="list-style-type: none"> <li>Taking divalproex: 25 mg every other day for 2 weeks, then 25 mg/day for 2 weeks, then 50 mg/day for 1 week, then 100 mg/day</li> <li>Taking enzyme inducing drug (eg, CBZ): 50 mg/day for 2 weeks, then 100 mg/day for 2 weeks, then 200 mg/day for 1 week, then 300 mg/day for 1 week</li> </ul> |
| <b>Valproate</b>       | target 10 – 15 mg/kg/d   | <ul style="list-style-type: none"> <li>impaired liver function, thrombocytopenia</li> </ul>   | <ul style="list-style-type: none"> <li>nausea/vomiting</li> <li>sedation</li> <li>ataxia</li> <li>thrombocytopenia</li> <li>Alopecia</li> <li>Weight gain</li> <li>pancreatitis</li> </ul>  | <ul style="list-style-type: none"> <li>Category D</li> <li>Low concentrations in breast milk and infant. Theoretical risk for hepatotoxicity or thrombocytopenia. Monitor for jaundice, liver damage, bleeding.</li> </ul> |   |   |

| Agent   | *Oral Dose  | Contraindications  | Adverse Events  | Pregnancy Category   | Remarks  |
|---|---|--|---|--|--|
| <b>Benzodiazepines</b>  |   |  |   |  |  |
| <b>Clonazepam</b><br><br><b>Lorazepam</b><br><b>Alprazolam</b><br><b>Diazepam</b> | Start - 0.25 mg bid,<br>increase by 0.25 mg<br>every 1-2 days;<br>maximum 20 mg/d<br><br>2 – 4 mg/d 1.5 to 6<br>mg/d 10 - 40 mg/d | <b>Contraindications:</b> <ul style="list-style-type: none"> <li>hypersensitivity</li> <li>significant liver disease (clonazepam)</li> <li>narrow angle glaucoma</li> <li>severe respiratory insufficiency (lorazepam)</li> <li>Caution in elderly patients and patients with impaired liver function.</li> <li>Risk of abuse in patients with history of substance abuse</li> </ul> | <ul style="list-style-type: none"> <li>sedation</li> <li>memory impairment</li> <li>ataxia</li> <li>dependence</li> <li>Confusion</li> <li>hypotension</li> </ul> | <ul style="list-style-type: none"> <li>Category D (all)</li> <li>All enter breast milk; breastfeeding not recommended</li> </ul> | <ul style="list-style-type: none"> <li>If doses sustained &gt; 2 months at therapeutic doses, then drug should be tapered over 4-week period</li> <li>Alprazolam – concern with rebound anxiety</li> </ul> |

| Agent                                   | *Oral Dose                                  | Contraindications   | Adverse Events   | Pregnancy Category   | Remarks   |
|---|---|---|--|--|---|
| <b>Typical antipsychotics</b>           |   |   |  |  |   |
| Chlorpromazine<br>Haloperidol           | 100 – 800 mg/d<br>2 – 20 mg/d               | <b>Contraindication :</b> <ul style="list-style-type: none"> <li>• Parkinson’s disease</li> <li>• QTc prolongation or concurrent use with medications that prolong the QTc interval</li> <li>• Severe CNS depression</li> <li>• Hypersensitivity</li> </ul> | <ul style="list-style-type: none"> <li>• Sedation</li> <li>• Orthostatic hypotension with chlorpromazine, thioridazine</li> <li>• Akathisia</li> <li>• Dystonia</li> <li>• drug-induced parkinsonism</li> <li>• Tardive dyskinesia may occur with all antipsychotics with long-term use.</li> <li>• Neuroleptic malignant syndrome</li> <li>• QTc changes</li> </ul> | <ul style="list-style-type: none"> <li>• Category C (all)</li> <li>• All enter breast milk; not recommended</li> </ul>   | <ul style="list-style-type: none"> <li>• Therapeutic doses not established in the treatment of PTSD</li> <li>• Use should be well justified in medical record because of the risk of tardive dyskinesia.</li> <li>•</li> </ul>  |
| <b>Atypical antipsychotics</b>          |   |   |  |  |   |
| Olanzapine<br>Quetiapine<br>Risperidone | 5 – 20 mg/d<br>300 – 800 mg/d<br>1 – 6 mg/d | <b>Relative contraindication:</b> <ul style="list-style-type: none"> <li>• Parkinson’s disease</li> <li>• Hypersensitivity</li> </ul>   | <ul style="list-style-type: none"> <li>• Sedation</li> <li>• Weight gain</li> <li>• Neuroleptic malignant syndrome</li> <li>• Higher doses may cause akathisia, drug-induced parkinsonism, especially with risperidone doses &gt;6 mg/d</li> </ul>   | Olanzapine: Category B<br>Quetiapine: Category C<br>Risperidone: Category C <ul style="list-style-type: none"> <li>• All excreted into breast milk; not recommended, use with caution</li> </ul> | <ul style="list-style-type: none"> <li>• Therapeutic doses not established for PTSD</li> <li>• Weight gain occurs with all agents; however, olanzapine produces significantly greater gain</li> <li>• The relative risk of tardive dyskinesia compared to typical antipsychotics has not been established for these agents</li> <li>• Monitor for development of diabetes/hyperglycemia , increased cholesterol and triglycerides</li> <li>•</li> </ul> |

| Agent  | *Oral Dose                 | Contraindications  | Adverse Events   | Pregnancy Category  | Remarks  |
|--|----------------------------|--|--|---|--|
| <b>Non-benzodiazepine</b>                    |                            |  |  |   |  |
| <i>hypnotics</i><br>- Zaleplon<br>- Zolpidem | 5 – 10 mg/d<br>5 – 10 mg/d | <b>Contraindications:</b> <ul style="list-style-type: none"> <li>Hypersensitivity</li> </ul> <b>Precautions:</b> <ul style="list-style-type: none"> <li>Caution with alcohol/drug abuse history</li> <li>Caution in elderly and patients with liver dysfunction</li> </ul> | <ul style="list-style-type: none"> <li>Sedation</li> <li>Ataxia</li> <li>Rebound insomnia may occur</li> <li>Dizziness</li> <li>Headache</li> <li>Behavioral changes, bizarre behavior, hazardous activities while asleep</li> </ul> | <ul style="list-style-type: none"> <li>Category C (both)</li> <li>Enters breastmilk; avoid zaleplon; use zolpidem with caution</li> </ul> | <ul style="list-style-type: none"> <li>Abuse has occurred resulting in withdrawal reactions</li> <li>Zolpidem is a CYP3A4 substrate and its metabolism can be decreased by 3A4 inhibitors</li> </ul> |
| <i>anti-anxiety</i><br>- Buspirone           | 20 – 60 mg/d               | <b>Precaution:</b> <ul style="list-style-type: none"> <li>MAOI use within 14 days</li> </ul>   | <ul style="list-style-type: none"> <li>Nausea</li> <li>Headache</li> <li>Dizziness</li> <li>Drowsiness</li> </ul>  | <ul style="list-style-type: none"> <li>Category B</li> <li>Excretion into breast milk unknown; not recommended</li> </ul>                 | <ul style="list-style-type: none"> <li>Buspirone is a CYP3A4 substrate and its metabolism can be decreased by 3A4 inhibitors</li> </ul>  |

\*Dose adjustments may be necessary in renal or hepatic impairment.

## D. ADJUNCTIVE SERVICES

### D1. Psychosocial Rehabilitation

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#### BACKGROUND

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Patients with chronic PTSD may develop a persistent incapacitating mental illness marked by severe and intolerable symptoms; marital, social, and vocational disability; and extensive use of psychiatric and community services. These patients may sometimes benefit more from therapeutic intervention that facilitates generalizing skills for coping with PTSD from clinic to home/work/community such as case management and psychosocial rehabilitation than from psycho-or pharmacotherapy.

Psychosocial Rehabilitation involves clinicians providing family psycho-education, supported employment, supported education, supported housing, or some serving as case managers, or others working with peer counselors. VHA's Uniform Mental Health Services policies (Veterans Health Administration, 2009) now mandate psychosocial rehabilitation, expanding such services from inpatient units to outpatient programs in Primary Care settings, Outpatient clinics, Community-Based Outpatient Clinics (CBOC), Vet Centers, Home-Based Care programs, and in partnerships with agencies and providers in communities.

#### RECOMMENDATIONS

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1. Consider psychosocial rehabilitation techniques once the client and clinician identify the following kinds of problems associated with the diagnosis of PTSD: persistent high-risk behaviors, lack of self care/independent living skills, homelessness, interactions with a family that does not understand PTSD, socially inactive, unemployed, and encounters with barriers to various forms of treatment/rehabilitation services.
16. Client and clinician should determine whether such problems are associated with core symptoms of PTSD and, if so, ensure that rehabilitation techniques are used as a contextual vehicle for alleviating PTSD symptoms.
17. Psychosocial rehabilitation should occur concurrently or shortly after a course of treatment for PTSD, since psychosocial rehabilitation is not trauma-focused.

#### DISCUSSION

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Penk and Flannery (2000) listed seven forms of psychosocial rehabilitation as clinical practice guidelines for Post-traumatic Stress Disorder (PTSD):

1. Patient education services
2. Self-Care and Independent Living Skills Techniques
3. Supported Housing
4. Marital/Family Skills Training
5. Social Skills Training
6. Vocational Rehabilitation
7. Case Management

A decade later, Penk & Ainspan (2009) suggest to add to this list: 8) Physical health and well-being and computer-assisted self-management training in reducing PTSD and other mental disorders such as addictions and depression.

46

**Table I - 7 Adjunctive Problem Focused Method/Services**

| If the client and clinician together conclude that the patient with PTSD: |  | Service/Training  |
|---|--|---|
| 1   | Is not fully informed about aspects of health needs and does not avoid high-risk behaviors (e.g., PTSD, substance) | Provide patient education   |
| 2   | Does not have sufficient self-care and independent living skills   | Refer to self-care/independent living skills training services        |
| 3   | Does not have safe, decent, affordable, stable housing that is consistent with treatment goals                     | Use and/or refer to supported housing services                        |
| 4   | Does not have a family that is actively supportive and/or knowledgeable about treatment for PTSD                   | Implement family skills training                                      |
| 5   | Is not socially active   | Implement social skills training                                      |
| 6   | Does not have a job that provides adequate income and/or fully uses his or her training and skills                 | Implement vocational rehabilitation training                          |
| 7   | Is unable to locate and coordinate access to services such as those listed above                                   | Use case management services  |
| 8   | Does request spiritual support   | Provide access to religious/spiritual advisors and/or other resources |
| OTHER CONDITIONS  |  |   |
| 9   | Does have a borderline personality disorder typified by parasuicidal behaviors                                     | Consider Dialectical Behavioral Therapy                               |
| 10  | Does have concurrent substance abuse problem   | Integrated PTSD substance abuse treatment (e.g., Seeking Safety)      |

47

48 The empirical literature on group treatment for PTSD has grown since the publication  
49 of the first edition of the Treatment Guidelines for PTSD.

50 Evidence-based research from randomized clinical trials now is available to support  
51 recommending psychosocial rehabilitation when treating veterans. (Glynn, Drebing,  
52 & Penk, 2009). Psychosocial Rehabilitations are not limited to veterans with  
53 schizophrenia or other psychoses. Psychosocial rehabilitations are recognized as  
54 efficacious in treating Post-traumatic Stress Disorders (PTSD), Major Depression  
55 Disorders (MDD), and Addictions, especially when mental health practices are  
56 delivered through self-management manuals and the Internet integrated into  
57 supported education and supported employment

58 The psychosocial rehabilitation model may include medication when needed; skills  
59 training designed to assist veterans to live productively in the community, and  
60 various forms of psychotherapy. Integrating trauma focused psychotherapies with  
61 psychosocial rehabilitation is currently under-utilized but new interventions are being  
62 empirically-validated to bring together several forms of treatments and rehabilitation  
63 for PTSD.

64



## 65 **Models of Psychosocial Rehabilitation Services**

### 66 1. Education

- 67 • Family psychoeducation is the process of providing education and coping skills  
68 for veterans and their families about relevant medical and mental disorders.  
69 Examples of such psychosocial rehabilitation are the family interventions for  
70 PTSD developed by Shirley Glynn (1999) at the VA in West Los Angeles and  
71 manualized approaches designed by Sherman, Sautter, Lyons, Manguno-Mire,  
72 Han, and Perry (2005) delivered at VHA medical centers in VISN 16—  
73 Oklahoma City, Jackson, and Houston.
- 74 • Family psychoeducation generally takes place in multi-family groups  
75 (producing the added benefit of augmenting social support) but such  
76 techniques can also be given in single family formats or even by books or  
77 online (e.g., Sherman & Sherman, 2005).
- 78 • Family psychoeducation is noted for fostering social support, challenging a key  
79 symptom in Post-traumatic Stress Disorder (PTSD) which is characterized by  
80 social avoidance and isolation. Precautions are needed in fielding family  
81 psycho-education among many different families, since consent of each  
82 individual is always required when information is shared about a veteran's  
83 illness and/or about families' symptoms and ways of coping. Family  
84 psychoeducation is a treatment modality in which families are a partner in  
85 providing services to each other: Families are not objects in treatment.
- 86 • Family psychoeducation is effective, particularly for PTSD (Cf. Glynn, Drebing,  
87 & Penk 2009), hence well-regarded in the VHA and emphasized in mental  
88 health services. Studies from different countries over the past 20 years show  
89 that family psychoeducation reduces the rates of re-hospitalization by an  
90 average of fifty percent.

### 91 2. Self-Care and Independent Living Skills Techniques

- 92 • While social rehabilitative therapies (i.e., teaching social, coping, and life  
93 function skills) have been proven effective in chronic schizophrenic and other  
94 persistently impaired psychiatric cohorts, they have yet to be formally tested  
95 with PTSD clients. Since they appear to generalize well from clients with one  
96 mental disorder to another, it is reasonable to expect that they will also work  
97 with PTSD clients. There is clinical consensus that appropriate outcomes would  
98 be improvement in self-care, family function, independent living, social skills,  
99 and maintenance of employment.
- 100 • Given the positive impact of independent skills training techniques for mental  
101 disorders in general (Halford et al., 1995), PTSD-centered modules should be  
102 developed and tested for effectiveness.

### 103 3. Supported Housing

- 104 • VHA for decades has offered support for housing through residential care  
105 programs, such as residential care in inpatient units, domiciliaries, affiliations  
106 with state and local housing resources, vouchers for single-room occupancy,  
107 congregate housing in private homes.

- 108 • Forms of housing considered more effective are those in which clinical services  
109 are integrated or efforts are made by treating staff to foster community living  
110 (Goldfinger et al., 1997; Schutt & Garrett, 1992)
- 111 • Existing literature for persons with other forms of mental illness demonstrates  
112 that case management linked to specialized clinical services is more effective  
113 than “single-room occupancy” or “warehousing” in shelters without other  
114 forms of support (Goldfinger et al., 1997).
- 115 • The greatest risk to ending housing arrangement and likelihood of  
116 discontinuing rehabilitation arise from addictions (Goldfinger, Schutt,  
117 Tolomiczenko, Seidman, Penk, Turner, 1999; Rog, 2000; Tsemberis &  
118 Eisenberg, 2000; Culhane, Metraux, & Hadley, 2002). Thus, interventions that  
119 provide housing supports are critical to success in rehabilitation (Mares,  
120 Kaspro, & Rosenheck, 2004).
- 121 • Research on outcomes for compensated work therapy transitional residence  
122 model(CWT/TR) have shown such endeavors indeed are quite successful in  
123 transitioning homeless, unemployed veterans who had been hospitalized on  
124 inpatient units from VA medical centers to independent living in the  
125 community (Schutt, Rosenheck, Penk, Drebing, & Seibyl, 2005). The program  
126 requires that unemployed, homeless veterans work in CWT (and, later, other  
127 jobs) in order to gain access to VHA housing for a limited time before  
128 transitioning to housing on one’s own or in private congregate housing with  
129 other veterans.
- 130 • Outcome studies show that such interventions are successful in promoting  
131 tenure in jobs and in personal living arrangements, promoting healthier styles  
132 of living, as well lowered costs due to reduce recidivism, (Cook, 2001; McKay,  
133 Johnsen, Banks and Stein, 2005; Cowell, Pollio, North, et al, 2003; Pelletier,  
134 Ngyuen, Bradley, et al (2005).

#### 135 4. Marital/Family Skills Training

- 136 • Marital and family treatments for trauma survivors fall into one of two general  
137 categories: systemic approaches designed to treat marital or family disruption,  
138 and supportive approaches designed to help family members offer support for  
139 an individual being treated for PTSD. These treatments are usually provided  
140 as an adjunct to other forms of treatment that are designed to directly address  
141 the PTSD symptoms.
- 142 • A single, low-quality RCT compared the addition of family therapy to individual  
143 therapy for war veterans with PTSD (Glynn et al., 1999). It found no  
144 significant benefit to the addition of behavioral family therapy (BFT), largely  
145 due to a high dropout rate, nor did it add significantly to the treatment of  
146 PTSD with direct therapeutic exposure (DTE) (an individual psychotherapy  
147 technique).
- 148 • There are no research studies on the effectiveness of marital/family therapy  
149 for the treatment of PTSD. However, because of trauma's unique effects on  
150 interpersonal relatedness, clinical wisdom indicates that spouses and families  
151 be included in treatment of those with PTSD. Of note, marriage counseling is

152 typically contraindicated in cases of domestic violence, until the batterer has  
153 been successfully (individually) rehabilitated.

#### 154 5. Social Skills Training

- 155 • Effectiveness of social skills training has been well demonstrated over many  
156 years in many RCTs but not specifically for PTSD (Dilk & Bond, 1996).
- 157 • Effectiveness of social skills training has been demonstrated for reducing social  
158 isolation of persons with severe mental disorders (e.g., schizophrenia); similar  
159 techniques may be promising for PTSD, particularly if adapted to address  
160 antecedent conditions involved in trauma and its consequences (Rothbaum &  
161 Foa, 1996).

#### 162 6. Vocational Rehabilitation

- 163 • Effectiveness of vocational rehabilitation techniques in treating mental  
164 disorders has been demonstrated under controlled experimental conditions  
165 (Bell & Lysaker, 1996; Bell et al., 1996; Bell et al., 1993; Bond et al., 1997)  
166 and controlled, clinical studies (Anthony et al., 1995; Drake, 1996; Lehman,  
167 1995; Lysaker et al., 1993).
- 168 • As a form of psychosocial rehabilitation, Supported Employment (SE) means  
169 that individuals with mental health disorders learn how to find and keep  
170 regular, real world, jobs in the community. In SE, vocational rehabilitation  
171 specialists provide continuous support to assist veterans achieve success at  
172 work. Outcomes for SE have been shown to be much better than for  
173 traditional approaches and this finding has been replicated in several countries  
174 (Bond, Drake and Mueser, 1997; Latimer, Lecomte, Becker, et al., 2006;  
175 Oldman, Thomson, Calsaferrri, et al., 2005; Penk, Drebing, Rosenheck, van  
176 Ormer, & Mueller, 2010, 2009).
- 177 • Strong outcome data exists to support the efficacy of Supported Employment  
178 (SE) for veterans with medical and mental disorders (Glynn, Drebing, & Penk,  
179 2009).
- 180 • SE consists of many different kinds of interventions including the “place-and-  
181 train” model that uses on-the-job training within and outside VA medical  
182 centers (Penk, 2000).
- 183 • Cochrane Review reviewed eighteen randomized controlled trials among  
184 nonveteran and veteran samples, mostly those with serious mental disorders,  
185 and found that SE was superior to programs that offered pre-vocational  
186 training (Crowther, Marshall, Bond, and Huxley, 2001).
- 187 • SE was found to be associated with fewer crises, less chaos, more structure,  
188 and on-going support from vocational rehabilitation specialists because  
189 consumers now focus on developing their lives in the community and  
190 managing their illness more independently (Bond, Becker, and Drake, 2001).
- 191 • Effect sizes for treating PTSD with Supported Employment are sizable (e.g.,  
192 Glynn, Drebing, & Penk; 2009; Drebing, Van Ormer, Rosenheck, Rounsaville,  
193 Herz, & Penk, 2005; Drebing, Van Ormer, Schutt, Krebs, Losardo, Boyd,  
194 Penk, & Rosenheck, 2004; Drebing, Van Ormer, Rosenheck, Rounsaville, Herz,  
195 & Penk, 2005; Rogers, Anthony, Lyass, & Penk, 2006; Drebing, Van Ormer,  
196 Mueller, Hebert, Penk, Petry, Rosenheck, & Rounsaville, 2007).

## 197 7. Case Management

198 Although case management has been shown to be useful for a range of other  
199 psychiatric disorders, there is currently no evidence available from RCTs or from  
200 systematic reviews to support or reject the use of case management for PTSD  
201 patients.

- 202 • Among populations with histories of trauma, the assertive community  
203 treatment models have been empirically validated under controlled (but not  
204 with random assignment) conditions (Mueser et al., 1998).
- 205 • Most of the research that empirically validates case management has been  
206 conducted among persons with severe mental disorders (Mueser et al., 1998),  
207 presumably including persons with co-occurring PTSD and other disorders.
- 208 • Evidence suggests that outcomes are more favorable for intensive case  
209 management (well-trained clinician teaches client psychosocial rehabilitation  
210 skills in the client’s home/community) than for simple case management  
211 (clinician links client to needed services).
- 212 • Case management has been demonstrated to reduce inpatient hospitalizations  
213 and severe symptoms, as well as to stabilize housing for formerly homeless  
214 persons; however, there is little evidence to suggest that case management  
215 improves vocational adjustment/social functioning (Mueser et al., 1998).

216 **D2. Spiritual Support**217 **BACKGROUND**

218 Social support will be critical for helping the individual cope after a trauma has  
219 occurred. It may be necessary to identify potential sources of support and facilitate  
220 support from others (e.g., partners, family, friends, work colleagues, and work  
221 supervisors).

222 The terms “religious” and “spiritual” are both used in the clinical literature to refer to  
223 beliefs and practices to which individuals may turn for support following a traumatic  
224 event.

225 **Spiritual & existential issues:** “Given the complex range of PTSD  
226 symptomatology, a successful treatment program will address not only the emotional  
227 issues that characterize the disorder but also its psychophysiological, cognitive, and  
228 interpersonal processes and existential meanings” (Hunter, 1996).

229 Religion may provide a framework by which many survivors of trauma construct a  
230 meaningful account of their experience, and may be a useful focus for intervention  
231 with trauma survivors.

232 **DISCUSSION**233 **Religion seeking is an observed post-traumatic phenomenon:**

234 There is a large body of anecdotal literature documenting the propensity of  
235 individuals to seek religious/spiritual comfort following a traumatic event. The  
236 terrorist attacks of September 11, 2001 provide a recent instance of this  
237 phenomenon. Meisenhelder (2002) notes “the events of September 11, 2001  
238 triggered a widespread national response that was two-fold: a post-traumatic stress  
239 reaction and an increase in attendance in religious services and practices  
240 immediately following the tragic events.” Schuster and his colleagues performed a  
241 nationwide phone survey of 569 adults within a week of the event (2001), and found

242 that "forty-four percent of the adults reported one or more substantial stress  
243 symptoms; 91 percent had one or more symptoms to at least some degree.  
244 Respondents throughout the country reported stress syndromes. They coped by  
245 talking with others (98 percent), turning to religion (90 percent)."

246 **Demonstrated benefits of the practice: religious/spiritual care:**

247 Some longitudinal studies have found support for religious /spiritual practices  
248 extending longevity (Hummer et al., 1999; Strawbridge et al., 1997), and reducing  
249 the duration of depression (Koenig et al., 1998). However, others examining the  
250 overall literature have found it to be "weak and inconsistent" (Sloan, et al., 1999).  
251 Strawbridge and his colleagues (1998) used a large public health survey to  
252 investigate "associations between two forms of religiosity and depression as well as  
253 the extent to which religiosity buffers relationships between stressors and  
254 depression". The authors conclude "religiosity may help those experiencing non-  
255 family stressors, but may worsen matters for those facing family crises."

256 Several possible pathways for positive physical / mental health benefits from  
257 religious/spiritual practice have been identified. These include; (1) reduction of  
258 behavioral risks through healthy religious lifestyles (e.g. less drinking or smoking)  
259 (2) expanded social support through involvement in spiritual communities, (3)  
260 enhancement of coping skills and helpful cognitive appraisals resulting in meaning-  
261 making, and (4) physiological mechanisms such as activation of the "relaxation  
262 response" through prayer or meditation.

263 Recent research on cognitive processes following traumatic events indicates that  
264 challenges to an individual's basic life assumptions may occur. These assumptions  
265 involve the security and meaningfulness of the world and the individual's sense of  
266 self-worth in relation to perception of the environment (Janoff-Bulman, 1979).  
267 Specifically, these assumptions are; (1) that one's environment is physically and  
268 psychologically safe, (2) that events are predictable, meaningful and fair, (3) that  
269 one's own sense of self-worth is positive in relation to experiences with other people  
270 and events (Hunter, 1996). The style and intensity of religious / spiritual appraisals  
271 and coping behaviors made in the aftermath of trauma may both reflect and  
272 influence the recovery trajectory and ultimate outcomes of traumatic events.

273 A recent meta-analytic study (Ano & Vasconcelles, 2005) of religious / spiritual  
274 coping with stressful events examined results from 49 studies meeting inclusion  
275 criteria from the 109 studies initially identified. In this study, Pargament's (1997)  
276 model of religious coping was used to distinguish between positive and negative uses  
277 of religion / spirituality to deal with life crises. Bivariate correlations between  
278 religious coping (positive and negative) measures and psychological adjustment  
279 (positive and negative) were averaged to obtain effect size estimates for findings for  
280 four key relationships; 1) positive religious coping and positive adjustment, 2)  
281 positive coping and negative adjustment, 3) negative coping and positive  
282 adjustment, and 4) negative coping and negative adjustment. The strongest  
283 average effect size estimate (.33) was obtained for the 29 studies examining the  
284 relationship between positive coping and positive adjustment. The 22 studies of  
285 negative coping and negative adjustment produced a significant moderate effect size  
286 estimate of .22. Finally, from 38 studies examining correlations between positive  
287 coping and negative adjustment, a small, yet significant, effect size estimate of -.12  
288 was obtained. Taken together, findings from the Ano & Vasconcelles (2005) meta-  
289 analysis show that both forms of religious coping are related to psychological  
290 adjustment to stress. Positive religious coping is related to both positive and

291 negative forms of adjustment in the directions expected; whereas negative coping is  
292 only significantly related to negative adjustment.

293 A study of help-seeking military veterans found significant associations between  
294 negative religious coping, lack of forgiveness and worse mental health outcomes  
295 (PTSD and Depression) (Witvliet et al., 2004). Similarly, loss of religious faith was  
296 found to be associated with worse mental health outcomes (i.e. greater utilization of  
297 mental health services) among military veterans in treatment for PTSD (Fontana &  
298 Rosenheck, 2004). Most recently, in a study of religiously active trauma survivors  
299 positive relationships were found between a measure of positive religious coping,  
300 seeking spiritual support, and posttraumatic growth. In the same study a negative  
301 religious coping indicator, religious strain, was significantly related to posttraumatic  
302 symptoms (Harris, Erbes, Engdahl, Olsen, Winskowski & McMahill, 2008). Thus,  
303 there is some consistency in findings of a complex relationship between religious  
304 coping and psychological adjustment following stress.

305 An integrative review of the literature on spirituality and bereavement by Wortman &  
306 Park (2008) suggests that religion/spirituality has a mixed but generally positive  
307 relationship with adjustment to bereavement depending on the dimension measured.  
308 Making-meaning appeared to be an important pathway associating  
309 religion/spirituality with adjustment for several dimensions, including general  
310 religiousness, intrinsic religiousness, religious coping, and belief in the helpfulness of  
311 religion.

312

### 313 **The role of the Chaplain:**

314 Recent studies of healthcare systems indicate strong association between the degree  
315 to which patient emotional / spiritual needs are met and overall patient treatment  
316 satisfaction. Healthcare accrediting bodies are recommending that  
317 Chaplains/pastoral care teams work in close collaboration with medical care teams to  
318 provide patients with an in-depth spiritual care experience that results in emotional  
319 comfort and improved satisfaction (Clark et al., 2003).

320 The Chaplain may play an important role in helping individuals regain a sense that  
321 their basic life assumptions are true. Chaplains receive training in a wide variety of  
322 supportive techniques, and they stand ready to assist all individuals, including those  
323 who do not subscribe to an organized religion. Chaplains may provide assistance in  
324 one or more of the following ways:

- 325 • Organizing and mobilizing community action by;
- 326 • Providing education and consultation with advice for leaders
- 327 • Assisting in the mobilization of action plans and recovery processes
- 328 • Offering facilitation of adaptation and mastery in social change
- 329 • Assisting in the development of community networks
- 330 • Supporting the development of a positive recovery organization
- 331 • Serving as a source of communication
- 332 • Fostering Community Theater and art geared to encouraging working  
333 through and recovery from the trauma.

334 Providing space and opportunities for prayers, mantras, rites and rituals and end-of-  
335 life care as determined important by the patient is another significant contribution of  
336 the Chaplain. (Canda and Phaobtong, 1992; Lee, 1997).

337 **EVIDENCE**

|   | Recommendation   | Sources  | LE  | QE   | SR |
|---|--|--|-----|------|----|
| 1 | Consider referral for religious/spiritual counseling as indicated for patient symptoms, consistent with available resources, and resonant with patient beliefs | Baldacchino and Draper, 2001<br>Meisenhelder, 2002<br>Calhoun et al., 2000<br>Humphreys et al., 2001<br>Nixon et al., 1999<br>Strawbridge et al., 1998 | III | Poor | I  |
| 2 | Consider providing direct spiritual care or ensuring patient access to spiritual care  | Bogia and Preston, 1985<br>Everly, 2000b   | II  | Fair | C  |

338 *LE = Level of Evidence; QE = Quality of Evidence; SR = Strength of Recommendation (see Appendix A)*339 **E. SOMATIC TREATMENT**340 **E1. Biomedical Somatic Therapies**341 **OBJECTIVE**

342 Evaluate the evidence for efficacy of Biomedical Somatic Therapies including  
343 Electroconvulsive Therapy (ECT), Cranial Electrotherapy Stimulation (CES), Vagal  
344 Nerve Stimulation (VNS), Repetitive Transcranial Magnetic Stimulation (rTMS), Deep  
345 Brain Stimulation (DBS)) in the treatment of PTSD.

346 **BACKGROUND**

347 There has been little research studying these modalities in the treatment of PTSD.  
348 ECT has strong research support in the treatment of refractory depression. VNS,  
349 rTMS, and CES have been cleared for marketing by the FDA for the treatment of  
350 depression, and DBS has been given a humanitarian exemption clearance for  
351 marketing for the treatment of Obsessive Compulsive Disorder. None of these  
352 modalities have been approved for the treatment of PTSD.

353 **RECOMMENDATIONS**

354 There is insufficient evidence to recommend the use of any of the Biomedical  
355 Somatic Therapies for first-line treatment of PTSD. [D]

356 18. ECT and rTMS may be considered as an alternative in chronic, severe,  
357 medication and psychotherapy-resistant PTSD. [B]

358 **DISCUSSION**

359 Although there is significant interest in biomedical somatic interventions in PTSD,  
360 there is no evidence for their use as a first line treatment for PTSD. ECT and rTMS  
361 may be beneficial in chronic, treatment-resistant PTSD, however their use has to be  
362 further studied in larger patient populations, and specifically in combat veterans.

363 ***Electroconvulsive Therapy (ECT)***

364 Watts (2007) reports a VA retrospective chart review study of twelve hospitalized  
365 Vietnam veterans with severe refractory depression (including bipolar depression)  
366 with co-morbid PTSD who underwent a course of ECT. Results showed good  
367 response for depressive symptoms but minimal response for PTSD symptoms.

368 Margoob et al (2010) reports on an open ECT trial for 20 patients (17 completers)  
369 with severe, chronic, antidepressant- and CBT-refractory PTSD who were

370 prospectively treated with a fixed course of 6 bilateral ECT treatments on an  
371 outpatient basis. The improvement in PTSD (40%), measured by CAPS was  
372 independent of the improvement in depression (57%), and treatment gains were  
373 maintained at 4-6 months follow-up.

#### 374 ***Repetitive Transcranial Magnetic Stimulation (rTMS)***

375 Rosenberg (2002) added rTMS to standard antidepressant therapy in 12 patients  
376 with PTSD and found that depression responded strongly but PTSD benefits were  
377 minimal.

378 Osuch (2009) studied rTMS as an adjunct to exposure therapy and existing  
379 medications in 9 patients with comorbid major depression and PTSD in a double-  
380 blind crossover study which included a sham arm and found decrease in  
381 hyperarousal symptoms alone.

382 Cohen (2004) reported findings of an RCT which showed significant improvement in  
383 PTSD core symptoms of re-experiencing and avoidance, but only when a 10 Hertz  
384 treatment was delivered (note that Osuch utilized no more than a 5 Hertz strength).

385 Boggio et al. (2009) studied the efficacy of 20 Hz rTMS of either right or left  
386 dorsolateral prefrontal cortex (DLPFC) as compared to sham rTMS in 30 patients with  
387 chronic PTSD, in a double blind, placebo-controlled trial with a sham arm. Both  
388 active conditions-20 Hz rTMS of left and right DLPFC induced a significant decrease in  
389 PTSD symptoms based on the PTSD Checklist and Treatment Outcome PTSD Scale;  
390 however, right rTMS induced a larger effect than left rTMS. Improvements in PTSD  
391 symptoms were still significant at 3-month follow-up. Neuropsychological evaluation  
392 showed that active 20 Hz rTMS was not associated with cognitive worsening in  
393 patients with PTSD.

#### 394 ***Vagal Nerve Stimulation (VNS)***

395 There is one open pilot study of Vagal Nerve Stimulation for treatment-resistant  
396 anxiety disorders (George, 2008) that included two patients with PTSD. This study  
397 does not provide sufficient evidence on which to base a recommendation regarding  
398 the use of VNS in the treatment of PTSD.

399 Although there has been significant interest and widespread utilization of CES in the  
400 treatment of PTSD, there is insufficient evidence for or against its use.

#### 401 **Conclusion:**

402 While intriguing, the findings from these studies are limited by a small number of  
403 patients and comorbid symptomatology, and do not provide adequate support to  
404 recommend any of the biomedical somatic interventions as a first-line treatment for  
405 PTSD. rTMS and ECT have had initial evidence of possible benefits in chronic,  
406 treatment-resistant PTSD, however more studies in larger patient populations are  
407 needed.



408 **EVIDENCE TABLE**

|   | Evidence  | Source  | LE                         | QE                   | SR |
|---|---|---|----------------------------|----------------------|----|
| 1 | ECT – for PTSD co-morbid with severe refractory depression.   | Watts 2007<br>Margoob, 2010   | II-3<br>II-2               | Fair                 | B  |
| 2 | rTMS -<br>Good PTSD outcome at higher frequency.<br><br>Primarily for comorbid depression added to antidepressant. Depression benefit robust. | Burt 2002<br>Cohen 2004<br>Boggio, 2009<br>Rosenberg PB 2002<br>Osuch, 2009 | III<br>I<br>I<br>I<br>II-1 | Poor<br>Good<br>Fair | B  |
| 3 | VNS - vagus nerve stimulation (VNS) for treatment-resistant anxiety disorders   | George, 2008  | I                          | Poor                 | I  |

409 *LE=Level of Evidence QE=Quality of Evidence; SR= Strength of Recommendation (see Appendix A)*

410

411 **E2. Acupuncture**412 **OBJECTIVE**

413 Improve management of PTSD symptoms, particularly when accompanied by  
414 associated symptoms of chronic pain, depression, insomnia, anxiety, or substance  
415 abuse.

416 **BACKGROUND**

417 The practice of needling in acupuncture to mediate pain, one of the well accepted  
418 indications for acupuncture, is thought to occur through the production of  
419 endogenous monoamines and neuropeptides. Besides activating neurohumoral  
420 pathways, acupuncture stimulates neural connections associated with the Autonomic  
421 Nervous System, prefrontal cortex, and limbic system, all structures thought to  
422 regulate the pathophysiology of PTSD. Acupuncture investigation for the treatment  
423 of PTSD has been limited to at best two (English) RCTs. However, symptomatic  
424 relief of disturbances associated with PTSD symptom clusters enhances the  
425 consideration of the use of this modality.

426 **RECOMMENDATIONS**427 **1.** Acupuncture may be considered as treatment for patients with PTSD. [B]428 **DISCUSSION**

429 Research focusing on the efficacy of acupuncture is still relatively limited. The few  
430 available studies are well done and demonstrate significant improvement in both  
431 PTSD and PTSD associated symptomatology. A larger numbers of studies exist  
432 concluding Acupuncture's efficacy in pain management, insomnia, depression, and  
433 substance abuse.

434 Hollifield et al (2007) evaluated the potential efficacy and acceptability of  
435 acupuncture for the treatment of PTSD. Individuals diagnosed with PTSD were  
436 randomized to an acupuncture treatment group (ACU), a cognitive-behavioral  
437 therapy group (CBT), or a wait-list control group (WLC). The primary outcome  
438 measure was self-reported PTSD symptoms at baseline, end treatment, and 3-month  
439 follow-up. Repeated measures MANOVA was used to detect predicted Group X Time

440 effects in both intent-to-treat (ITT) and treatment completion models. Compared  
441 with the WLC condition in the ITT model, acupuncture provided large treatment  
442 effects for PTSD ( $F [1, 46] = 12.60$ ;  $p < 0.01$ ; Cohen's  $d = 1.29$ ), similar in  
443 magnitude to group CBT ( $F [1, 47] = 12.45$ ;  $p < 0.01$ ;  $d = 1.42$ ) (ACU vs. CBT,  $d =$   
444  $0.29$ ). Symptom reductions at end treatment were maintained at 3-month follow-up  
445 for both interventions.

446 A recent unpublished DoD/VA RCT studied 55 active duty members with PTSD  
447 randomized to PTSD treatment as usual (TAU) and PTSD treatment as usual plus  
448 eight 90 minute acupuncture sessions delivered twice weekly for four weeks (TAU +  
449 Acupuncture). Outcome measures included: Clinician-Administered PTSD Scale  
450 (CAPS), PTSD Checklist (PCL), Becks Depression Inventory (BDI I-II), Numeric  
451 Rating Scale for Pain (NRS), and SF-36v2. Follow up was at baseline, 4, 8, and 12  
452 weeks post randomization. Compared to usual PTSD care, a four week course of  
453 twice weekly acupuncture resulted in significantly greater improvement in PTSD  
454 symptoms, (Pre-post ES 1.4-1.6 versus usual care ES 0.12-0.74, significant  
455 improvement in Depression, and significant improvement in pain.

#### 456 EVIDENCE TABLE

|   | Evidence  | Source                 | LE | QE   | SR |
|---|---|------------------------|----|------|----|
| 1 | There is some evidence that acupuncture may be helpful with the management of Post-traumatic Stress Disorder, acute or chronic. | Hollified et al., 2007 | I  | Good | B  |

457 *LE=Level of Evidence QE=Quality of Evidence; SR= Strength of Recommendation (see Appendix A)*

## 458 F. COMPLEMENTARY AND ALTERNATIVE MEDICINE

### 459 OBJECTIVE

460 Identify interventions derived from traditional and nontraditional complementary  
461 approaches that may provide effective first line or adjunctive treatment for PTSD.

### 462 BACKGROUND

463 Complementary and alternative medicine (CAM) is a group of diverse medical  
464 practices, products and systems that are not generally considered part of  
465 conventional medicine. While there is limited evidence to suggest that any of the  
466 CAM therapies listed below are efficacious for PTSD, these interventions may be of  
467 value in dealing with other symptoms (particularly those associated with  
468 hyperarousal) or comorbid conditions. There is little evidence that these  
469 interventions are harmful. Some patients who may be reluctant to accept mental  
470 health labels or interventions may be more accepting of these novel treatment  
471 approaches. CAM interventions are affordable and generally accessible in  
472 communities across the nation. Many CAM therapies are practiced in a group setting  
473 which may have the added benefit of increasing socialization. CAM programs may be  
474 engaged as a family that could increase social support and reduce stress for all  
475 family members. CAM approaches often provide an increased sense of mastery and  
476 control that may promote greater resilience.

477 Since complementary medicine may relate to particular cultural backgrounds or other  
478 belief systems, health professionals should be aware of and sensitive to the needs  
479 and desires of the patient and the family. Health professionals should be willing to  
480 discuss the effectiveness of therapy and different options of care within the context  
481 of the current health care system.

#### 482 **CAM Modalities:**

483 CAM modalities are typically grouped into broad categories reflecting putative  
484 mechanism of action: However, this is more for convenience than any real biologic  
485 underpinnings. In fact, some modalities such as acupuncture span more than one  
486 category. Generally consistent with the schema offered by the National Center for  
487 Complimentary and Alternative Medicine these groups are:

- 488 1. **Natural Products (Biologically Based Practices):** Biologically based therapies  
489 that use natural substances (e.g., herbs, foods, vitamins, dietary supplements) to  
490 promote healing and wellness.
- 491 2. **Mind-Body Medicine:** Approaches that seek to enhance conscious mental  
492 control over other bodily processes. These practices use a variety of techniques  
493 to enhance the mind's ability to impact physical function and promote health.  
494 (e.g., meditation, yoga, prayers, guided imagery, progressive muscle  
495 relaxation/deep breathing, dance therapy, art therapy, music therapy.)
- 496 3. **Manipulation and Body Based Practices (Exercise and Movement):** These  
497 practices are based on movement of one or more parts of the body.  
498 Manipulations that focus primarily on the structure of the body including bones  
499 and joints, soft tissue circulatory and lymphatic system such as chiropractic,  
500 spinal manipulation, massage therapy, reflexology, and acupressure.
- 501 4. **Energy medicine:** Energy moving practices involving the use of energetic fields  
502 that purportedly surround and penetrate the human body. (e.g., Qi gong, Reiki,  
503 Therapeutic touch)
- 504 5. **Whole Medial Systems:** Alternative medicine systems based upon complete  
505 systems of theory and practice such as Ayurvedic Medicine, homeopathy,  
506 naturopathy, and Traditional Chinese Medicine (TCM).

#### 507 **RECOMMENDATIONS**

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- 508 1. There is insufficient evidence to recommend CAM approaches other than  
509 acupuncture as first line treatments for PTSD. [I]
- 510 2. CAM approaches may be recommended as adjunctive treatments for PTSD  
511 symptoms (hyperarousal and when comorbid conditions for which CAM  
512 therapies may be effective (e.g. pain, insomnia) are present.
- 513 3. CAM approaches may be recommended as alternatives when patients decline  
514 standard mental health interventions for the purpose of engagement and  
515 potential increase in sense of self-mastery, self-control and social support.
- 516 4. Providers should discuss the effectiveness of therapy and different options of  
517 care within the context of the current health care system.

518 **DISCUSSION**

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519 Surveys of CAM utilization (meditation, yoga, massage, and deep breathing  
520 exercises) among the general US population indicate significant increases in  
521 acceptance of these practices over the past decade (Barnes et al., 2008). A recent  
522 White House Commission reported on CAM highlighted the need for continued  
523 rigorous research regarding these approaches (White House Commission, 2002).

524 Complementary and Alternative Medicine (CAM) approaches to the treatment of  
525 many medical and mental health diagnoses, including PTSD are in use; the research  
526 base to support their effectiveness is improving but not complete. Numerous CAM  
527 modalities have been used for symptomatic relief related to insomnia, anxiety, and  
528 various somatic presentations associated with PTSD. Revisions of the VA/DoD CPGs  
529 are currently underway to include a comprehensive review of the evidence for all  
530 treatments, including CAM.

531 **F1. Natural Products (Biologically Based Practices)**

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532 **Herbal or dietary supplements** have also been used for the treatment of  
533 PTSD. Omega -3 fatty acid, docosahexaenoic acid (DHA) affects catecholamines and  
534 proinflammatory cytokines have been shown to decrease the perception of stress  
535 (Bradbury, 2004). Although there have been some studies of their effectiveness,  
536 the results of these small RCTs provide insufficient evidence to draw firm conclusions  
537 about their effectiveness for PTSD. In addition, the quality and purity of herbals and  
538 dietary supplements available in the United States varies widely, further complicating  
539 their use.

540 **F2. Mind-Body Medicine**

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541 **Mindfulness Based Stress Reduction (MBSR)** is one such modality used to  
542 emphasize the richness of each moment of our lives. Like many of the tools CAM  
543 practices encourage, MBSR fosters a sense of awareness and discovery issuing from  
544 the deep relaxation a contemplative mind experiences. Transcendental Meditation™  
545 used to treat anxiety and depression in Vietnam  
546 Veterans (Books 1990) is another tool or practice shown to be useful for autonomic  
547 arousal, **Yoga Nidra**, a relaxation and meditative form of yoga, has also been used  
548 as an adjunctive treatment for PTSD. Formal studies demonstrating its effectiveness  
549 for PTSD are currently being conducted, and further research is needed on Yoga  
550 Nidra for PTSD before its effectiveness can be commented on. Yogic breathing, a  
551 pattern of of specific breathing techniques, may also lessen anxiety.

552 **F3. Manipulation and Body Based Practices (Exercise and Movement)**

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553 Exercise has been advocated as an integrative approach in the prevention and  
554 treatment of PTSD and other combat related mental health problems. Taylor (2008)  
555 examined the relationship between baseline physical fitness and the development of  
556 PTSD symptoms (as measured by the Impact of Event Scale) in a group of 31  
557 soldiers undergoing military survival training. He found that higher levels of pre-  
558 study physical fitness were inversely related to both trait anxiety levels and IES  
559 scores. Studies have shown that pre-trauma levels of exercise tend to decline after  
560 developing PTSD (de Assis, et al., 2008). Aside from PTSD, Depression was a  
561 frequent condition for which exercise therapy was applied. The majority of reviewed

562 studies utilized an aerobic exercise regimen, e.g., walking, running, stationary  
563 cycling (Diaz & Motta, 2008; Manger & Motta, 2005). One study emphasized the  
564 importance of participant selection of the specific type of exercises that would  
565 comprise their treatment (Donta, et al, 2003). The studies reviewed here utilized  
566 both group and individual exercise formats. All studies demonstrated either a  
567 reduction in symptoms from baseline PTSD measures or relative to a placebo or  
568 control group but the effects were generally modest and did not always extend to  
569 other mental health disorders such as anxiety and depression. A primary  
570 methodological limitation of the papers reviewed here is that exercise interventions  
571 were rarely conducted in isolation from other psychotherapeutic approaches.

#### 572 **F4. Energy medicine**

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573 Reiki and Johrei

- 574 • *Reiki* and *johrei* are both energy medicine techniques that originated in Japan.  
575 In *reiki*, the practitioner places his hands on or near the person receiving  
576 treatment with the intent to transmit *ki*, believed to be life-force energy.  
577 *Johrei*, a form of energy healing that originated in Japan, involves the  
578 practitioner facing the person receiving the treatment, where “spiritual  
579 energy” is transmitted through the practitioner (Brooks, et al., 2006).
- 580 • There are no current controlled studies examining *Reiki* or *johrei* in patients  
581 with PTSD or Acute Stress Disorder. A small number of low-quality studies  
582 have been conducted showing positive improvement in conditions commonly  
583 comorbid with PTSD, such as depression (Collinge, Wentworth, and Sabo,  
584 2005) or anxiety (Brooks et al., 2006).
- 585 • A recent systematic review of randomized clinical trails of *reiki* noted that the  
586 currently available RCTs are “scarce” and lack independent replication (Lee,  
587 Pittler, and Ernst, 2008). The studies that exist suffer from methodological  
588 flaws related to sample size, inadequate design, and poor reporting.

#### 589 **F5. Whole Medical Systems**

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#### 591 **F6. Other Approaches**

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##### 592 ***Animal-Assisted Therapy (AAT)***

593 AAT is a goal-directed intervention in which an animal meeting specific criteria is an  
594 integral part of the treatment process. AAT is delivered and/or directed by a  
595 health/human service provider working within the scope of his or her profession. AAT  
596 is designed to promote improvement in human physical, social, emotional, and/or  
597 cognitive functioning. AAT is provided in a variety of settings and may be group or  
598 individual in nature. Commonly used animals include dogs and horses.

**I-3. MANAGEMENT OF SPECIFIC SYMPTOMS**

2 This section includes recommendations regarding treatment interventions for a  
3 selected list of physical symptoms that are common in patients presenting with  
4 posttraumatic stress symptoms.

5 Survivors of trauma may not complain directly of PTSD symptoms such as re-  
6 experiencing or avoidance. Instead, they may complain of sleeping problems.  
7 When seeking to identify PTSD, providers should consider asking specific  
8 questions about sleep problems, (including flashbacks and nightmares), pain  
9 (including musculoskeletal, headache), or hyperarousal (including an  
10 exaggerated startle response or sleep disturbance). Many individuals with PTSD  
11 experience sleep disturbances (trouble falling asleep or problems with waking up  
12 frequently after falling asleep). Chronic pain and insomnia often occur  
13 simultaneously, with the vast majority of chronic pain patients complaining of  
14 interrupted or poor quality sleep. When a person with PTSD experiences sleep  
15 disturbances, using alcohol as a way to self-medicate becomes a double-edged  
16 sword. Excessive alcohol use can impair one's ability to sleep restfully and to  
17 cope with trauma memories and stress. The need to improve sleep in these  
18 patients is clear, given increasing evidence that sleep disturbance is associated  
19 with heightened pain sensitivity and elevated disability.

20 Chronic pain is frequently observed in patients with PTSD and is often associated  
21 with a significant level of affective distress and physical disability. Chronic pain  
22 may develop because of an injury sustained in a traumatic event such as a motor  
23 vehicle accident, work-related injury, or injury in military combat. Patients with  
24 chronic pain, particularly headache disorders and fibromyalgia (FM), associated  
25 with psychological traumas need a special management strategy. Diagnosis of  
26 headache disorders and FM in traumatized patients and obtaining the clinical  
27 history of a traumatic event or diagnosing PTSD in chronic pain patients is of  
28 great importance.

**A. Sleep Disturbances****BACKGROUND**

31 Many patients with PTSD have had insomnia for years including broken sleep,  
32 frequent awakenings, and nightmares, all of which contribute to poor sleep  
33 quality. Hyperarousal behaviors, part of PTSD symptoms for many people can be  
34 stronger at night and contribute to insomnia. Sleep problems in traumatized  
35 patients may also reflect co-morbid conditions some of which may be of new  
36 onset (pain may be prominent among these).

37 There is no evidence to suggest that insomnia, as a component of traumatic  
38 stress reactions, should be managed differently than insomnia associated with  
39 other conditions. Clinical experience does, however, show that some  
40 psychologically traumatized patients dread sleep because of intense nightmares.

41 Research demonstrates that non-pharmacologic sleep strategies yield outcomes  
42 equal or superior to those obtained with hypnotics alone or hypnotics combined  
43 with non-pharmacologic strategies. Long-term outcomes are better following  
44 non-pharmacologic interventions. The aim of sleep management is to establish a  
45 regular, normalized sleep-wake pattern.

46 RECOMMENDATIONS (BASED ON CONSENSUS OF CLINICAL EXPERTS)47 **Sleep Disturbance**

- 48 1. Encourage patients to practice good sleep hygiene including:
- 49 • Restricting the night-time sleep period to about eight hours
  - 50 • Waking at a regular time
  - 51 • Arising from bed at a regular time
  - 52 • Avoiding going to bed too early
  - 53 • Avoiding alcohol
  - 54 • Avoiding stimulants, caffeinated beverages, power/energy
  - 55 drinks, nicotine and over the counter medications
  - 56 • Avoiding stimulating activities, light, noise and temperature
  - 57 extremes before bedtime (e.g., exercise, video games, T.V.) or
  - 58 in the sleeping area
  - 59 • Reducing (to less than 30 minutes), or abolishing, daytime naps
  - 60 • Practicing relaxation techniques
  - 61 • Engaging in moderate exercise, but not immediately before
  - 62 bedtime
- 63 2. Offer Cognitive Behavioral Therapy for Insomnia that may include:
- 64 • Educating about proper sleep habits and sleep needs
  - 65 • Correcting false and unrealistic beliefs/concerns about sleep
  - 66 • Identifying and addressing anxious, automatic thoughts which
  - 67 disrupt sleep
- 68 3. Consider adjunctive therapy for nightmares using prazosin. [B]
- 69 4. Any significant change in sleep patterns should trigger clinical
- 70 reassessment in order to rule out worsening or new onset of co-morbid
- 71 conditions

72 **Insomnia**

- 73 5. Monitor symptoms to assess improvement or deterioration and reassess
- 74 accordingly.
- 75 6. Explore cause(s) for insomnia, including co-morbid conditions.
- 76 7. Begin treatment for insomnia with non-pharmacologic treatments including
- 77 sleep hygiene and cognitive behavioral treatment (See recommendation for
- 78 Sleep Disturbances).
- 79 8. The selection of sleep agents for the treatment of insomnia in PTSD
- 80 patients may be impacted by other treatment decisions (e.g., medications
- 81 already prescribed for the treatment of PTSD, depression, TBI, pain, or
- 82 concurrent substance abuse/withdrawal) and social/environmental/logistical
- 83 concerns associated with deployment.
- 84 a. **Trazodone** may be helpful in management of insomnia and may
- 85 also supplement the action of other antidepressants.

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- b. **Hypnotics** are a second line approach to the management of insomnia and should only be used for short periods of time. Should hypnotic therapy be indicated, the newer generation of non-benzodiazepines (e.g. zolpidem, eszopiclone, ramelteon) may have a safety advantage by virtue of their shorter half-life and lower risk of dependency. Patients should be warned of and monitored for the possibility of acute confusional states/bizarre sleep behaviors associated with hypnotic use. Benzodiazepines can be effective in chronic insomnia but may have significant adverse effects (confusion, sedation, intoxication) and significant risk of dependency.
  - c. **Atypical antipsychotics** should be avoided due to potential adverse effects but may be of value when agitation or other symptoms are severe.
  - d. If nightmares remain severe, consider adjunctive treatment with **prazosin [B]**
  - e. If symptoms persist or worsen – refer for evaluation and treatment of insomnia

105 Additional information of management of insomnia can be found in:

106 VHA Pharmacy Benefit Management (PBM) guideline for Insomnia:  
107 <http://www.pbm.va.gov/ClinicalRecommendations.aspx>

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109



110 **Sleep Hygiene Patient Education**

- Avoid or limit caffeinated products, nicotine, and alcohol, especially later in the day
- Avoid drinking excess liquids after supper to avoid having to get up during the night to go to the bathroom
- Avoid or limit daytime naps to 30 minutes in the early afternoon before 3:00 pm
- Go to bed only when sleepy. Sleep only as much as needed to feel refreshed. Staying in bed longer can result in fragmented/shallow sleep on following nights
- Create a dark, quiet, temperature-controlled bedroom (e.g., change the number of blankets you use; use earplugs; close the door if noisy)
- Avoid heavy meals within 2 hours of bedtime; a light snack might help if hungry
- Maintain a regular daily schedule of activities including bedtime and awakening times, 7 days/week. Use an alarm clock if needed
- Exercise regularly during the daytime. Avoid active exercise in the late evening when it is close to bedtime
- Use the bed and bedroom only for sleeping or sexual activity. Do not eat, work or watch television while in bed
  - If you cannot sleep, if possible, get out of bed and go to another room read or engage in other quiet activities; or do other relaxation activities before attempting to sleep again. Return to bed only when sleepy. Repeat if necessary. Do not watch the clock; turn the clock around or cover it up
- Solve problems before retiring. If not possible, write down your worries, plans and strategies during the early evening and not at bedtime
- Correct extrinsic factors such as environmental disruption (e.g., pets or snoring partner)
- Establish a “wind-down” routine going to bed and develop and maintain bedtime “rituals” that make going to sleep a familiar routine; for example:
  - Set time to relax before bed with 20-30 minutes of relaxation (e.g. soft music, meditation, breathing exercises)
  - Take a warm bath
  - Have a light snack, which could include: warm milk, foods high in tryptophan, such as bananas, carbohydrates, which can help induce sleep

Adapted from Petit L, et al. *Age Ageing* 2003; 32: 22. Wilson S. and Nutt D. *Prescriber* 2005; 19: 29-41  
Wolkove N, et al. *CMAJ* 2007; 176: 1449-54.

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112 **DISCUSSION**113 **Use of Benzodiazepines for Sleep Disturbance**

114 In a small, double-blind, placebo-controlled temazepam trial in acute  
115 accident/injury victims at a trauma center (Mellman et al., 2002) temazepam 30  
116 mg was administered for 5 nights, tapered for 2 nights, then discontinued. At 6  
117 week follow-up, 6/11 temazepam subjects and 3/11 placebo subjects met PTSD  
118 symptom criteria. Sleep improvement was noted however for the duration of the  
119 trial. However, in a small randomized, controlled trial, alprazolam did not have  
120 substantial benefit for PTSD or for nightmares, although it did improve anxiety  
121 (Braun, 1990). In another small single-blind controlled study, clonazepam did not  
122 demonstrate significant benefit for sleep difficulties, including nightmares (Cates,  
123 2004).

124 An argument can be made for short-term use of a benzodiazepine for the  
125 purpose of reducing hyperarousal symptoms in the immediate trauma aftermath,  
126 in order to help normalize sleep cycles and minimize anxiety. Longer-term use of  
127 benzodiazepines, however, should be avoided, as the limited data available show  
128 that prolonged use of benzodiazepines (1-6 months duration) is associated with a  
129 higher rate of subsequent PTSD (Gelpin et al., 1996).

130 Benzodiazepines use should be considered relatively contraindicated in combat  
131 veterans with PTSD, because of the very high co-morbidity of combat-related  
132 PTSD with alcohol misuse and substance use disorders (upwards of 50% of  
133 comorbidity) and potential problems with tolerance and dependence. Once  
134 initiated in combat veterans, benzodiazepines can be very difficult, if not  
135 impossible, to discontinue, due to significant withdrawal symptoms compounded  
136 by the underlying PTSD symptoms.

### 137 **Prazosin to Treat Sleep Disturbance**

138 Five publications (Raskind 2000, 2002, 2003; Taylor 2006 & 2008) that examined  
139 the role of antiadrenergic medications, commonly used for treating hypertension,  
140 in the treatment of posttraumatic stress disorder (PTSD) were identified in the  
141 peer-reviewed literature.

142 Although Taylor et al. (2006) and Raskind et al. (2003) were excluded from the  
143 analyses (due to small number of subjects that did not meet inclusion criteria),  
144 both have shown positive results in reducing psychological distress specifically to  
145 trauma cues (Taylor et al., 2006). Patients taking prazosin showed significant  
146 improvement on the Clinician-Administered PTSD Scale (Raskind, 2003).

147 Raskind et al. (2007) evaluated prazosin effects on trauma nightmares, sleep  
148 quality, global clinical status, dream characteristics, and comorbid depression.  
149 Forty veterans (mean age 56 +/- 9) with chronic PTSD and distressing trauma  
150 nightmares and sleep disturbance were randomized to evening prazosin (13.3  
151 +/- 3 mg/day) or placebo for 8 weeks. In the evaluable sample (n = 34) primary  
152 outcome measures demonstrated that prazosin was significantly superior to  
153 placebo for reducing trauma nightmares and improving sleep quality. Prazosin  
154 shifted dream characteristics from those typical of trauma-related nightmares  
155 toward those typical of normal dreams.

156 Taylor et al. (2008) was a double blind, placebo-controlled crossover study of 13  
157 civilians with trauma-related PTSD. Prazosin was rapidly titrated to 3 mg/night  
158 during each 3-week treatment phase. Prazosin, compared with placebo,  
159 significantly increased total sleep time by 94 min (p <0.01); and total rapid eye  
160 movement (REM) sleep and mean REM duration were also longer with prazosin.  
161 Reductions in trauma nightmares, total PTSD symptoms (using the PCL-C) and  
162 CGIC scores were significantly changed compared with placebo.

163 The results of these studies were consistent and positive, suggesting that  
164 prazosin therapy is safe and is associated with reduction of nighttime symptoms  
165 of PTSD. Prazosin is an effective and well-tolerated treatment for trauma  
166 nightmares, sleep disturbance and global clinical status in veterans with chronic  
167 PTSD.

168 [Ruff and colleagues \(2009\) found in an observational study that prazosin](#)  
169 [combined with sleep hygiene](#) counseling was an effective initial treatment for a  
170 group of OIF/OEF veterans (n=74) with headaches associated with histories of  
171 mild TBI from exposure to an explosion in combat and with PTSD. Prazosin was  
172 well tolerated. Nine weeks after providing sleep counseling and initiating an

173 increasing dosage schedule of prazosin at bedtime 65 veterans had reduced  
174 headache intensity and frequency, reduced daytime sleepiness, and improved  
175 cognitive performance. These gains maintained 6 months later.

176 Sleep hygiene counseling is beneficial in terms of improving sleep duration and  
177 reducing the time it takes for a person to fall asleep (Morin et al., 1999). By  
178 blocking nightmares in people with PTSD, prazosin prolongs sleep duration by  
179 preventing sleep interruptions. Thus the two interventions may have synergized,  
180 with sleep hygiene counseling reducing the time it took for veterans to fall asleep  
181 and prazosin prolonging sleep.

182 A systematic review by Dierks et al. (2007) did not find any additional publication  
183 to the above. The authors concluded that despite various limitations, all of the  
184 studies showed significant improvements in the sleep-related symptoms of PTSD  
185 following the addition of prazosin therapy, based on the Clinician Administered  
186 PTSD Scale recurrent distressing dreams item and the Clinical Global Impression  
187 of Change [scale](#).

#### 188 EVIDENCE TABLE

|   | Evidence  | Source  | LE   | QE   | NB            | SR |
|---|---|---|------|------|---------------|----|
| 1 | Prazosin to improved sleep quality, reduced trauma nightmares | Dierks et al. (2007) §<br>Raskind et al., 2002, 2007<br>Taylor et al., 2008 | I    | Good | Mod           | B  |
| 2 | Benzodiazepines for sleep disturbance, insomnia               | Gelpin et al., 1996<br>Mellman et al., 1998                                 | II-2 | Fair | Small/<br>Neg | C  |

189 *LE =Level of Evidence; QE = Quality of Evidence; SR = Strength of Recommendation; NB=Net benefit (see*  
190 *Appendix A)*

191

192 **Table I - 8 Pharmacologic Studies - Prazosin for Sleep Disturbances**

| Author, Tear             | Results   | n  | Trauma                         | LE   | QE   | NB  |
|--------------------------|---|----|--------------------------------|------|------|-----|
| Raskind et al. ,<br>2002 | Significant improvement in dream scores after 8 weeks of prazosin treatment | 59 | Retrospective study - Veterans | II-2 | Poor | Mod |
| Raskind et al. ,<br>2003 | Significant improvement, CAPS, CGI.<br>Prazosin > placebo                   | 10 | Veterans                       | I    | Poor | Mod |
| Raskind et al.,<br>2007  | No difference   | 34 | Veterans                       | I    | Good | Mod |
| Taylor et al., 2006      | Reduction in global PTSD illness severity                                   | 11 | Civilian                       | II   | Poor | Mod |
| Taylor et al., 2008      | Reductions of nighttime, significantly increased total sleep time           | 13 | Civilian                       | I    | Fair | Mod |

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195 **B. Pain**196 **BACKGROUND**

197 There is growing body of research that indicates clearly that PTSD and chronic  
198 pain frequently co-occur. People with both PTSD and chronic pain tend to have  
199 greater distress and impairment compared to those with only one of these  
200 conditions and assessment and treatment are more complicated. When  
201 evaluating a patient with known, or suspected, ASD or PTSD it becomes  
202 important to include a pain assessment (acute or chronic) in the examination and  
203 to consider the extent to which PTSD symptoms may be being influenced by pain  
204 or the extent to which pain symptoms are being exacerbated by PTSD. Likewise,  
205 pain evaluations should include an assessment of possible PTSD symptomatology.  
206 For some patients chronic pain may actually serve as a reminder of the traumatic  
207 event, which will tend to exacerbate the PTSD. For others, re-experiencing the  
208 trauma may exacerbate pain.

209 Certain types of chronic pain are more common in individuals who have  
210 experienced specific traumas. For example, adult survivors of physical,  
211 psychological, or sexual abuse tend to be more at risk for developing certain  
212 types of chronic pain later in their lives. The most common forms of chronic pain  
213 involve: pain in the pelvis, lower back, face, and bladder; fibromyalgia;  
214 interstitial cystitis; and nonremitting whiplash syndromes. Chronic pain is a  
215 common problem among returning soldiers. In service persons from OEF/OIF  
216 head, neck and back pain, shoulder and knee pain have been found most  
217 common (Lew, 2009). While the concept of comorbid physical and emotional  
218 problems is not new, the particular combination of chronic pain, PTSD symptoms  
219 and post concussion syndrome and their apparent prevalence is unique to the  
220 OEF/OIF population and appears to be more common in blast injured patients  
221 and may be more difficult to treat than each condition independently.

222 Understanding the development and maintenance of chronic pain and PTSD and  
223 how they interact is of essential importance as the co-occurrence of these  
224 conditions is often overlooked in practice and, as a result, encumbers treatment  
225 outcomes. Fear-based avoidance is a central theme in both PTSD and chronic  
226 pain. While the underlying basis for the avoidance may differ, avoidance  
227 behaviors may exacerbate or maintain the severity of either or both conditions.  
228 Although pharmacological agents have been examined in the treatment of pain  
229 and PTSD individually, little is known regarding the relationship of medication use  
230 with functioning in patients with comorbid conditions. Pain should be assessed  
231 and aggressively treated in early phases of post trauma and providers across  
232 disciplines need to work together to develop treatments that are complementary,  
233 based on theory, and supported by empirical evidence.

234 **RECOMMENDATIONS (BASED ON CONSENSUS OF CLINICAL EXPERTS)**

- 235 1. Recommend pain assessment using a '0 to 10' scale.
- 236 2. Obtain a thorough biopsychosocial history and assess for other medical and  
237 psychiatric problems, including risk assessment for suicidal and homicidal  
238 ideation; misuse of substances, such as drugs or alcohol; over-the-counter  
239 and prescription drugs or narcotics.

- 240 3. Assessment should include questions about the nature of the pain and likely  
241 etiology (i.e., musculoskeletal and neuropathic), locations, quality, quantity,  
242 triggers, intensity, duration of the pain, as well as aggravating and relieving  
243 factors.
- 244 4. Assessment should include evaluation of the impact of pain on function and  
245 activities, pain-related disability or interference with daily activities.
- 246 5. Assessment should include the identification of avoidance behaviors that  
247 contribute to emotional distress and/or impaired functioning.
- 248 6. Management of pain should be multidisciplinary, addressing the physical,  
249 social, psychological and spiritual components of pain in an individualized  
250 treatment plan that is tailored to the type of pain. [C]
- 251 7. Selection of treatment options should balance the benefits of pain control with  
252 possible adverse effects (especially sedating medications) on the individual's  
253 ability to participate in, and benefit from, PTSD treatment. [I]
- 254 8. Musculoskeletal pain syndromes can respond to correcting the underlying  
255 condition and treatment with non-steroidal anti-inflammatory drugs  
256 (NSAIDs).
- 257 9. When appropriate, recommend use of non-pharmacologic modalities for pain  
258 control such as biofeedback, massage, imaging therapy, physical therapy, and  
259 complimentary alternative modalities (yoga, meditation, acupuncture). [C]
- 260 10. Centrally acting medications should be used in caution in patients with PTSD  
261 as they may cause confusion and deterioration of cognitive performance and  
262 interfere with the recovery process
- 263 a. If required, lower doses of opioid therapy or other centrally acting  
264 analgesics should be used for short duration with transition to the  
265 use of NSAIDs. [C]
- 266 11. Consider offering Cognitive Behavioral Therapy that may include:
- 267 b. Encouraging increasing activity by setting goals
- 268 c. Correcting false and unrealistic beliefs/concerns about pain
- 269 d. Teaching cognitive and behavioral coping skills (e.g., activity  
270 pacing)
- 271 e. Practicing and consolidation of coping skills –and reinforcement of  
272 use.

## 273 DISCUSSION

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### 274 **Prevalence**

275 PTSD and chronic pain disorder are highly comorbid (Sharp, 2001). The literature  
276 indicates a high degree of co-occurrence between pain and PTSD, regardless of  
277 whether the pain is being assessed in patients with PTSD or PTSD is being  
278 assessed in patients with chronic pain. Chronic pain and posttraumatic stress  
279 disorder (PTSD) are frequently observed to be comorbid following traumatic  
280 injury (Bryant et al., 1999; Hickling and Blanchard, 1992). Studies have shown  
281 that PTSD symptoms tend to be elevated in patients with chronic pain and  
282 fibromyalgia (Amir, 1997; Engel, 2000; Sherman, 2000) chronic low back pain,  
283 and other musculoskeletal disorders (Sherman, 2000).

284  
285 Sharp (2004) reported on four studies (Bryant et al., 2001; Beckham et al.,  
286 1997; McFarlane et al., 1987; and Benedikt & Kolb, 1986). The samples were  
287 drawn from MVA victims, combat veterans, fire fighters, and chronic pain clinic  
288 patients. In each instance they found a high prevalence of pain in patients  
289 diagnosed with PTSD or a high prevalence of PTSD in patients diagnosed with  
290 chronic pain.

291 Asmundson et al. (2002) discusses the current definitions of chronic pain and  
292 PTSD and finds some overlaps in the cognitive, behavioral, and physiological  
293 domains. He states that "anxiety and hyper-arousal, avoidance behaviour,  
294 emotional lability and elevated somatic focus are frequently observed in both  
295 conditions. Both PTSD and chronic pain are characterized by hyper-vigilance and  
296 attentional bias for stimuli that are specific to each condition."

297 Norman et al. (2008) found that self-reported pain levels within 48 hours after  
298 serious injury were significantly and strongly associated with the subsequent risk  
299 of PTSD. Similarly, in a study of 2,931 seriously injured patients admitted to  
300 acute care hospitals in the United States, Zatzick and Galea (2007) found that  
301 pain after injury was significantly associated with an increased risk of PTSD one  
302 year after hospitalization. The prevalence of PTSD is particularly high when the  
303 chronic pain results directly from a traumatic event (Hickling & Blanchard, 1992;  
304 Taylor & Koch, 1995; Chibnall, 1994; Asmundson et al. 1998; Otis et al., 2003),  
305 and the presence of both PTSD and chronic pain can increase the symptom  
306 severity of either condition (Otis, 2003).

307 Schwartz et al. (2006) noted that between 10% and 50% of patients treated in  
308 tertiary care settings for chronic pain and related conditions have symptoms that  
309 meet criteria for PTSD.

310  
311 Norman (2007) demonstrated that Peritraumatic pain was associated with an  
312 increased risk of PTSD, even after controlling for a number of other significant  
313 risk factors other than acute stress disorder symptoms. An increase of 0.5 S.D.  
314 from the mean in a 0–10 pain rating scale 24–48 hours after injury was  
315 associated with an increased odds of PTSD at 4 months by more than fivefold,  
316 and at 8 months by almost sevenfold. The author suggests that high levels of  
317 peritraumatic pain could be used to identify individuals at elevated risk for PTSD  
318 following traumatic injury.

319 Persons with comorbid pain and PTSD may experience less symptom  
320 improvement after treatment for these conditions (Asmundson, 2002; Baker et  
321 al., 1997; Clark et al., 2009; Hickling, 1992; McClean, 2005; Muse, 1986). The  
322 co-occurrence of chronic pain and PTSD has implications on the experience of  
323 both conditions. Patients with chronic pain related to trauma or PTSD experience  
324 more intense pain and affective distress (Geisser, 1996), higher levels of life  
325 interference (Turk et al., 1996), and greater disability than pain patients without  
326 trauma or PTSD (Sherman, 2000).

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### 328 ***Comorbidity of Pain and PTSD (and PCS) in OEF/OIF***

329 Because of the nature of injuries and the physical demands of OEF and OIF  
330 deployments, there are data to suggest that a significant majority of returned  
331 warriors report ongoing pain problems (Clark, 2004; Clark et al., 2009a; Gironda  
332 et al., 2006; Kalra et al., 2008).

333 Posttraumatic headaches are a common complaint (Gironda et al., 2009; Clark et  
334 al., 2007; Gironda et al., 2006; Lew et al., 2007; Ruff et al., 2008). Other  
335 commonly reported pain problems are low back pain and joint pain (Clark et al.,  
336 2007; Clark et al., 2009a). The high prevalence of chronic pain (pain that lasts  
337 longer than 3 months) places OEF/OIF soldiers at long-term risk for impaired  
338 functional ability, significant emotional distress, interpersonal conflict, substance  
339 misuse, and vocational limitations. Substance misuse (including opioid  
340 medications) has also been found in OEF/OIF returnees, although at lesser  
341 prevalence than pain, (published rates range from 3 to 28%; Clark et al., 2007;  
342 Kalra et al., 2008; Kang & Hyams, 2007; Shipherd et al., 2007).

343 Pain symptoms are a common complaint among post deployment populations  
344 (back pain and headache). In one study of 1800 OEF/OIF veterans, 46.5%  
345 reported some pain, with 59% of those exceeding the VA clinical threshold of  $\geq 4$   
346 on a 0-10 pain scale (Gironda et al., 2006). Recent literature suggests that  
347 many returning service members have multiple comorbid symptoms of post  
348 concussion syndrome, chronic pain, and PTSD (Clark et al., 2007; Clark et al.,  
349 2009; Lew et al., 2009; Sayer et al., 2008). In a sample of OEF/OIF veterans  
350 pain was the single most common complaint recorded and 42% of the sample  
351 reported concurrent PCS, chronic pain, and PTSD symptoms.

#### 352 Evidence Statements

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- 353 • Benedikt and Kolb (1986) reported that 10 percent of a sample of 225  
354 patients referred to a Veterans Administration pain clinic met criteria for  
355 PTSD
- 356 • Muse (1986) reported that 9.5 percent of a sample of patients attending a  
357 multidisciplinary chronic pain center met criteria for "post-traumatic pain  
358 syndrome"
- 359 • When patients are referred for the assessment of a chronic pain problem  
360 resulting from a traumatic event, the prevalence of PTSD increases
- 361 • Asmundson (1998) -Among injured workers with chronic pain who were  
362 referred to a rehabilitation program, 34.7 percent of the sample reported  
363 symptoms consistent with PTSD.
- 364 • Hickling (1992) - high rates of PTSD have been reported by, and  
365 colleagues for patients referred for, psychological treatment following an  
366 MVA
- 367 • Chibnall (1994) - rates of PTSD in patients for which pain is secondary to  
368 an MVA range from 30 to 50 percent
- 369 • Greenfield (1992); Turk (1996) - Among patients with fibromyalgia, 24 to  
370 47 percent of patients attribute the onset of their symptoms to a physical  
371 injury associated with an MVA
- 372 • Geisser (1996) - patients with accident-related pain and high PTSD  
373 symptoms reported higher levels of pain and affective distress relative to  
374 patients with accident-related pain and without PTSD, or nonaccident-  
375 related pain
- 376 • McFarlane (1994) - the prevalence of chronic pain in individuals with a  
377 primary diagnosis of PTSD is high. Pain was the most common physical  
378 complaint (45% back pain and 34% headaches).
- 379 • Beckham (1997) - 80 percent among combat Vietnam veterans with PSTD  
380 who completed self-report questionnaires assessing PTSD reported the  
381 presence of a chronic pain condition Increased levels of PTSD re-  
382 experiencing symptoms were associated with increased pain level and  
383 pain-related disability

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- Tushima (1990) found that patients with post-traumatic headache reported more frequent pain and had a poorer prognosis than did nontraumatic headache patients
  - Peterlin (2008)- found that a chronic migraine sample reported significantly more PTSD than did an episodic migraine sample

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“The mechanism by which chronic pain, and PTSD (and Post Concussion Syndrome) interact is still unclear. Researchers evaluating comorbid pain and PTSD have presented a variety of models to explain this phenomenon, including a Shared Vulnerability model, a Mutual Maintenance model, and a Triple Vulnerability model (Asmundson et al., 2002; Otis et al., 2003; Sharp & Harvey, 2001). These models propose mechanisms of interaction via the dispositional tendency to be fearful or anxious, the belief that anxiety states cause harmful consequences, and the cognitive distortions and behavioral patterns of PTSD and chronic pain that maintain or exacerbate symptoms of the other syndrome. These models have yet to be fully tested and there are no available outcomes data regarding the success of integrated treatment of comorbid pain and PTSD symptoms (Otis et al., 2003). However, such research is now being conducted (Otis, 2008)” (Walker, 2010)

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#### 404 ***Pain & PTSD Sensitivity Assessment***

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Given the high rates of comorbidity of chronic pain and PTSD, clinicians should assess for both disorders. Several well-validated self-report questionnaires are available to help determine a diagnosis and the severity of symptoms. Self-report measures of pain include the 0 to 10 numerical pain rating scale, the McGill Pain Questionnaire or for a more comprehensive assessment, the West Haven-Yale Multidimensional Pain Inventory, or the Pain Outcomes Questionnaire was developed and validated specifically for veterans and will be integrated into the electronic medical record.

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Asmundson et al. (2002) Recommended that clinicians who conduct diagnostic assessments of patients presenting with PTSD symptoms also screen for the presence of existing pain conditions such as fibromyalgia or chronic musculoskeletal pain. The clinician can include self reported questions or use a structured clinical interview format. They recommend the use of self-report measures like the McGill Pain Questionnaire-Short Form, the Multidimensional Pain Inventory, and where appropriate, the Pain Anxiety Symptoms Scale.

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#### 421 ***Interdisciplinary Approach to Management***

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Only a few studies have reported the results of treatments designed to address co-occurring chronic pain and PTSD. Given the current state of the literature, few recommendations can be made regarding preferred treatment modalities for individuals with comorbid pain and PTSD. Several authors support the use of a multidisciplinary treatment approach for patients with PTSD and chronic pain (Muse, 1986).

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Given the broad range of emotional and physical symptoms characteristic of veterans with Co-occurring PTSD, chronic pain and possible PCS, an integrated treatment approach is required (Walker, 2010). Treatment goals need to be



431 clarified (e.g., reduced symptom severity, increased occupational or interpersonal  
432 functioning, reduced ongoing use of health care services) (Clark, 2008).

433 The focus of the integrated approach should be on education and management of  
434 symptoms and improving daily function. The interventions and treatment  
435 modalities employed follow the current evidence based recommendations for  
436 PCS, chronic pain and PTSD (See VA/DoD guidelines for mTBI/Concussion and  
437 the VHA NPMS, 2003)

438 The VHA National Pain Management Strategy (VHA/NPMS, 2003) advocates the  
439 use of evidence based, comprehensive, multicultural, integrated approach to pain  
440 issues that focuses on reducing pain and suffering, improving function, and  
441 enhancing quality-of-life. Specific practice guidelines for managing acute and  
442 chronic pain associated with certain conditions like low back pain (APS-AAPM,  
443 2005; VHA/DoD, 2007), the use of opioids with chronic pain (VHA/DoD, 2010),  
444 and other resource material are recommended for use by practitioners.

445 For the OEF/OIF population of returning soldiers, treatment should be  
446 individualized on an inpatient or outpatient basis depending upon need within  
447 group and individual treatment formats. Treatment should be goal oriented and  
448 time limited, with increased patient function and independence as major goals  
449 (Clark, 2009).

450 “The foundations of treatment are psychoeducation, coping skills training, and  
451 cognitive– behavioral therapy that involves active self-management techniques  
452 for chronic pain and PTSD. In many cases symptom management will be  
453 facilitated by the judicious use of pharmacological agents; minimizing  
454 polypharmacy and adverse effects that may impact comorbid symptoms. This is  
455 particularly important in that many of these patients present with difficulties in  
456 memory, attention, and concentration. While these cognitive deficits may be  
457 mild, many pharmaceuticals useful in the treatment of pain or emotional distress  
458 can negatively impact cognitive functioning” (Walker, 2010).

#### 459 ***Non-Pharmacologic Treatment***

460 Initial treatment of PTSD focuses on providing psychoeducation about the  
461 disorder. This may include specifically addressing how fears and avoidance of the  
462 trauma may serve to maintain the symptoms and decrease the ability to function.  
463 This may also include discussing how your pain may serve as a trigger or  
464 reminder of the trauma and increase arousal, fear, and avoidance and thereby  
465 increase disability and pain (Sharp, 2004).

466 Nonpharmacological ways to manage chronic pain may include **Relaxation** (e.g.,  
467 relax the locus of the pain problems by relaxing muscle tension). **Increasing**  
468 **Activity and Fitness** (e.g., gradual return to more normal levels of activities  
469 and slowly increase patient’s stamina for physical activities), **Reducing**  
470 **Emotional Over-Reactivity** (e.g., practice specific methods of emotional  
471 reaction to stressful triggers); **External Focusing/Distracting** (e.g., learn to  
472 shift and manipulate the focus of attention in a positive way, which will minimize  
473 your experience of the pain).

474 **Complementary Alternative medicine** - approaches are being increasingly  
475 studied to decrease symptom severity. There are numerous interventions that  
476 are being used to help manage chronic pain, including breathing, muscle  
477 relaxation, visual imagery, music, cold/heat, stretching, massage therapy, stress  
478 management, acupuncture, acupressure, hydrotherapy and other.

479 Tan et al. (2007) examined various CAM therapies for chronic pain. For example  
480 heart rate variability (HRV) biofeedback (using a stress eraser portable  
481 biofeedback device that easily can be used by Veterans at home for the purpose  
482 of increasing HRV) has been shown to be effective for reducing the symptoms of  
483 PTSD (e.g., Tan et al., 2009; Zucker, 2009), and for persistent pain associated  
484 with fibromyalgia (Hassett et al., 2007). Regulating Heart rhythm coherence,  
485 using biofeedback devices that computes the heart rhythm patterns have been  
486 shown to improve symptoms such as depression, anxiety, panic disorder, and  
487 PTSD symptoms (McCraty, Atkinson, Tomasino, & Stuppy, 2001).

#### 488 **Treatment: Pharmacotherapy**

489 There are no studies evaluating the pharmacotherapy for acute dissociation or  
490 traumatic pain associated with ASR.

491 The most common first-line treatments for pain have traditionally been  
492 analgesics, which include opioids, NSAIDS (nonsteroidal anti-inflammatory  
493 drugs), antiepileptic drugs, and tricyclic antidepressants for neuropathic pain, and  
494 antidepressants that target the inhibition of norepinephrine reuptake (SNRIs).  
495 With respect to trauma exposure, some data suggests that pain patients with  
496 comorbid PTSD use analgesic medications at higher rates than their non-PTSD  
497 counterparts (Schwartz et al., 2006).

#### 498 **Opioid Therapy**

499 While controversial, the use of opioid medications in the treatment of chronic,  
500 non-malignant pain has increased significantly over the past three decades  
501 (Caudill-Slosberg et al., 2004). The efficacy of opioids in alleviating acute pain is  
502 well established, but less is known regarding their utility in treating chronic pain  
503 or their relationship with patient functioning over extended periods of use  
504 (Ballantyne and Shin, 2008). Side effect profiles associated with opioid use –  
505 tolerance, physical dependence, cognitive impairment – often are cited as factors  
506 contributing to potential decreases in functioning (Ballantyne and Shin, 2008;  
507 Eriksen et al., 2006). Understanding the interrelationships of chronic pain, PTSD,  
508 and opioid use with patient functioning becomes clinically relevant given that  
509 comorbid psychiatric disorders are known to increase risk of abuse and  
510 dependency among persons with chronic pain (Edlund et al., 2007). Some data  
511 suggests that pain patients with comorbid PTSD use analgesic medications at  
512 higher rates than their non-PTSD counterparts (e.g., Schwartz et al., 2006). It  
513 may suggest that the experience of pain in the present may be affected by  
514 previous emotional trauma and ongoing trauma-related stress disorders. Some  
515 findings suggest the possibility that long term use of opioids may lead to opioid-  
516 induced hyperalgesia (Angst & Clark, 2006).

517 Given inconsistent findings regarding the efficacy of opioids for long term pain  
518 control, potential for reductions in overall functioning, and the increased risk of  
519 abuse and dependency, providers should consider the benefits and potential  
520 harm of extended opioid therapy for patients with chronic pain subsequent to  
521 traumatic injury (Clapp, 2010) (VA/DoD COT CPG, 2010).

522 Selective serotonin reuptake inhibitors (SSRIs) are recommended as the first-line  
523 pharmacological intervention for PTSD (See Intervention for PTSD –Module B).  
524 SSRIs also have been examined for use in the treatment of chronic pain, but  
525 support for their efficacy in this population is limited (see reviews by McCleane,  
526 2008; Dworkin et al., 2007). Sedative and anxiolytic medications are sometimes  
527 prescribed to alleviate symptoms associated with both PTSD and chronic pain but

528 are not recommended due to the addictive properties of many anxiolytic agents  
529 (American Psychiatric Association, 2004; Sanders et al., 2005). The relationship  
530 between these pharmacological agents and functioning among patients with  
531 comorbid pain and PTSD has not been examined.

532

533

### Early pain intervention to prevent PTSD

534 Acute pain caused by physical injury may by itself be a precursor for PTSD.  
535 Injury that is also associated with traumatic exposure increases the risk for  
536 PTSD. When pain is treated early and aggressively, patients may have the best  
537 chance of getting better. Though many fear addiction from opioids, they can be  
538 an important part of halting the pain cycle. Few studies have investigated the  
539 effect of pain reduction in the early stages after injury and the development of  
540 PTSD.

541 Bryant et al. (2008<sup>m</sup>) examined the influence of acute administration of  
542 morphine as protective against development of PTSD in a consecutive sample of  
543 patients admitted to hospital after traumatic injury (n = 155). The patients who  
544 met criteria for PTSD at 3 months (14%) received significantly less morphine  
545 than those who did not develop PTSD; there was no difference in morphine levels  
546 in those who did and did not develop major depressive episode or another  
547 anxiety disorder. The authors suggested that administration of morphine in the  
548 acute post-traumatic stage may limit fear conditioning in the aftermath  
549 of traumatic injury and may serve as a secondary prevention strategy to reduce  
550 PTSD development.

551 Holbrook et al. (2010) analyzed data for 696 military personnel (mostly male,  
552 mean age about 24) who were hurt during OIF but who did not have serious  
553 traumatic brain injury. About one-third (35%) of the injured personnel developed  
554 PTSD. The finding was that those who had been administered morphine shortly  
555 after their injury (60% versus 76%) were less likely to develop PTSD (ORs  
556 ranging from 0.48 to 0.66,  $P < 0.05$  for all). Several factors, including severity and  
557 mechanism of injury, need for amputation, resuscitation, and the presence of  
558 mild traumatic brain injury were adjusted for. Although causality could not be  
559 established, the authors concluded that a reduction in perceived pain levels  
560 through the use of morphine or other opioids, as part of trauma care, may lower  
561 the rate of PTSD onset after major trauma.

562

### Cognitive Behavior Therapy (CBT)

563 Cognitive behavioral therapy is recommended as first line therapy for PTSD (See  
564 Intervention for PTSD Module B). CBT is a skills-based treatment approach that  
565 focuses on teaching patients ways to identify and change maladaptive thoughts,  
566 feelings, and behaviors and to replace them with those that are more balanced  
567 and adaptive, with the ultimate goal of improving patients' overall quality of life  
568 and reducing psychological distress. In addition, cognitive-behavioral treatment  
569 approaches focus on changing certain target behaviors that appear to be  
570 problematic, and teach adaptive ways of coping (Otis, 2009). CBT includes  
571 establishing a safe (controlled) setting for the patient to address trauma cues.  
572 Treatment protocols include a range of techniques from desensitization, to  
573 confront mildly distressing cues to exposure (re-experiencing all at once) of the  
574 traumatic event to allow the patient to confront his or her fears and process the  
575 trauma memory until it is no longer distressing (Calhoun & Resick, 1993;  
576 Blanchard & Hickling, 1997). By also learning stress management techniques,  
577 the patient can learn how to decrease his or her anxiety response and learn how

578 to use these skills to manage PTSD-type symptoms. Many of these tools are also  
579 used for adaptive chronic pain management. "By gradually increasing one's  
580 activity level through the exposure exercises, the patient is also able to decrease  
581 his/her focus on pain and fear and engage in life again" (Townsend, 2010). CBT  
582 for pain is aimed at changing maladaptive thoughts and behaviors that serve to  
583 maintain and exacerbate the experience of pain. CBT for chronic pain involves  
584 teaching patients ways of safely reintroducing enjoyable activities into their lives.

585 Using components of cognitive processing therapy (CPT) for PTSD and cognitive  
586 behavioral therapy (CBT) for chronic pain management, a 12-session integrated  
587 treatment for veterans with comorbid chronic pain and PTSD was developed  
588 (Otis, 2009). The key components of the CBT for chronic pain, include cognitive  
589 restructuring (i.e., teaching patients how to recognize and change maladaptive  
590 thoughts), relaxation training (e.g., diaphragmatic breathing, progressive muscle  
591 relaxation), time-based activity pacing (i.e., teaching patients how to become  
592 more active without overdoing it), and graded homework assignments designed  
593 to decrease patients' avoidance of activity and reintroduce a healthy, more active  
594 lifestyle (Otis, 2007). The therapy includes weekly readings and homework  
595 assignments, pre- and post-treatment evaluations using measures of pain, PTSD,  
596 physical disability, and psychological distress.

597 The result of implementing the program in a pilot study demonstrates the  
598 importance of establishing participant trust, regular therapy attendance, and  
599 addressing participant avoidance. Participants reported that they generally liked  
600 the format of treatment, appreciated learning about the ways that chronic pain  
601 and PTSD share some common symptoms, and ways that the two disorders can  
602 interact with one another. The authors concluded, based on this initial small pilot  
603 study that the participants appeared to benefit from receiving the integrated  
604 treatment for pain and PTSD (Otis, 2009).

605

606 **EVIDENCE TABLE Pain in PTSD**

|   | Evidence                                       | Source  | LE  | QE   | SR |
|---|--|---|-----|------|----|
| 1 | Prevalence of co morbid chronic pain and PTSD. | Schwartz., 2006<br>Girona, 2006<br>Sharp, 2004<br>Asmundson et al., 2002<br>Norman et al., 2008<br>Zatzick & Galea, 2007<br>Bryant et al., 2009 | III | Good | A  |
|   | Comorbidity in OEF/OIF                         |   |     |      |    |
| 2 | Pain & Anxiety Sensitivity Assessment          | Asmundson et al., 2002  |     |      | B  |
| 3 | Multidisciplinary Approach to Management       | Clark, 2008   |     |      | C  |
| 4 | Treatment: Pharmacotherapy                     | Schwartz et al., 2006   |     |      | I  |
|   | Early intervention to control acute pain       | Bryant<br>Holbrook et al., 2009   | I   | Good | B  |

|   |                                  |   |  |  |   |
|---|----------------------------------|---|--|--|---|
| 5 | Non-Pharmacologic Treatment      | Sharp, 2004   |  |  |   |
| 6 | Cognitive Behavior Therapy (CBT) | Otis, 2009<br>Townsend, 2010<br>Foa & Meadows, 1997<br>Foa et al., 1991<br>Keane & Kaloupek, 1996<br>Marks et al., 1998<br>Solomon et al., 1991 |  |  | C |

607 LE =Level of Evidence; QE = Quality of Evidence; SR = Strength of Recommendation; NB=Net  
608 benefit (see Appendix A)

609

## 610 C. Irritability, Severe Agitation, or Anger

### 611 BACKGROUND

612 In the most general sense, anger is a feeling or emotion that ranges from mild  
613 irritation to intense fury and rage. Anger is often a central feature of response to  
614 trauma and can be seen as a core component of the survival response in  
615 humans. Mismanaged or uncontrolled anger and rage can lead to a continued  
616 sense of being out of control and may cause conflicts in personal and professional  
617 relationships. Anger and irritability may be associated with domestic violence and  
618 abuse, road rage, and workplace violence even if there is no intent to cause harm  
619 to others. It is important to distinguish between anger and aggression.  
620 Aggression is behavior that is intended to cause harm to another person or  
621 damage property. This behavior can include verbal abuse, threats, or violent  
622 acts. Anger, on the other hand, is an emotion and does not necessarily lead to  
623 aggression. Therefore, a person can become angry without acting aggressively.

624 Anger becomes a problem when it is felt too intensely, is felt too frequently, or is  
625 expressed inappropriately. Anger management interventions include a range of  
626 methods including teaching individuals to recognize signs of becoming angry,  
627 self-calm, avoid escalating conflicts, and respond to anger-eliciting situations in  
628 more positive ways.

### 629 RECOMMENDATIONS (BASED ON CONSENSUS OF CLINICAL EXPERTS)

- 630 1. Assess the nature of symptoms, severity, and dangerousness. Consider  
631 using standardized Anger Scales, such as Spielberger's State-Trait Anger  
632 Expression Inventory,<sup>1</sup> to quantify.
- 633 2. Explore for cause of symptoms and follow-up to monitor change
- 634 3. Consider referral to specialty care for counseling or for marital or family  
635 counseling as indicated. Offer referral for:
  - 636 a. Anger Management therapy
  - 637 b. Training in exercise and relaxation techniques
- 638 4. Promote participation in enjoyable activities - especially with family/ loved  
639 ones.
- 640 5. Promote sleep and relaxation
- 641 6. Avoid stimulants and other substances (caffeine, alcohol)

- 642           7. Address pain (see pain management)
- 643           8. Avoid benzodiazepines
- 644           9. Consider SSRI/SNRI
- 645                 a. If not responding to SSRI/SNRI and other non-pharmacologic
- 646                 interventions, consider low-dose antiadrenergics, or low-dose
- 647                 atypical antipsychotics (risperidone, quetiapine)
- 648                 b. If not responding or worsening, refer to specialty care

649 **DISCUSSION**

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650           In anger management treatments, physical arousal, problem behaviors, and

651           anger-provoking thoughts/beliefs are all addressed in different ways (Chemtob,

652           1997).

653           Cognitive-behavioral treatment such as anxiety management, shows positive

654           results when used to address anger, and applies many techniques to manage

655           these three anger components.

656 **DISCUSSION**

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657 **Prevalence**

658           High levels of anger have been observed in veterans of the Iraq and Afghanistan

659           Wars. (Jakupcak, 2007)

660           A study of sample OEF/OIF veterans found over half of the veterans with PTSD

661           indicated that they had been aggressive in the past 4 months, such as

662           threatening physical violence, destroying property, and having a physical fight

663           with someone. Veterans with subthreshold PTSD syndrome reported just about

664           the same amount of aggressive behavior as the veterans with PTSD. In fact,

665           anger has been shown to be associated with other comorbid conditions to PTSD

666           such as head injury, and alcohol (substance) abuse. Each of these conditions has

667           been associated with elevated anger and hostility in veterans from previous

668           conflicts (Elboquen 2010).

669           In another survey of 2797 US soldiers returning from deployment, overall, 40%

670           of soldiers reported killing or being responsible for killing during their

671           deployment. Even after controlling for combat exposure, killing was a significant

672           predictor of PTSD symptoms, alcohol abuse, anger, and relationship problems

673           Maquan et al., 2010).

674           A study assessing Vietnam combat veterans and comparing them to veterans

675           who did not serve in war found that the combat veterans were not significantly

676           more angry than their veteran peers who did not serve in Southeast Asia.

677           Additionally, various parameters of war zone duty were not highly associated with

678           anger scores. However, combat veterans with PTSD scored significantly higher

679           than veterans without PTSD on measures of anger, arousal, range of anger-

680           eliciting situations, hostile attitudinal outlook, and tendency to hold anger in.

681           These results suggest that PTSD, rather than war zone duty, is associated with

682           various dimensions of angry affect (McFalls et al., 1997).

683           Anger can be a very difficult emotion to deal with and can lead to a number of

684           legal and interpersonal problems, such as domestic violence. In fact, individuals

685           with PTSD are particularly at risk for the perpetration of relationship violence.

686 Research has identified anger as prominent in and an influence on, treatment  
687 outcomes for military veterans with PTSD. To improve treatment effectiveness,  
688 clinicians need to assess veterans' anger, aggression, and alcohol use, as well as  
689 their current fear of anger and elucidate the relationship between these factors  
690 (Forbes, 2008).

691 Chemtob et al., (1997) described three components of posttraumatic anger that  
692 can become maladaptive or interfere with one's ability to adapt to current  
693 situations that do not involve extreme threat:

- 694 • **Arousal:** Anger is marked by the increased activation of the  
695 cardiovascular, glandular, and brain systems associated with emotion and  
696 survival. It is also marked by increased muscle tension. Sometimes with  
697 individuals who have PTSD, this increased internal activation can become  
698 reset as the normal level of arousal and can intensify the actual emotional  
699 and physical experience of anger. This can cause a person to feel  
700 frequently on-edge, keyed-up, or irritable and can cause a person to be  
701 more easily provoked. It is common for traumatized individuals to actually  
702 seek out situations that require them to stay alert and ward off potential  
703 danger. Conversely, they may use alcohol and drugs to reduce overall  
704 internal tension.
- 705 • **Behavior:** Often, the most effective way of dealing with extreme threat is  
706 to act aggressively, in a self-protective way. Additionally, many people  
707 who were traumatized at a relatively young age do not learn different  
708 ways of handling threat and tend to become caught in their ways of  
709 reacting when they feel threatened. This is especially true of people who  
710 tend to be impulsive (who act before they think). Again, as stated above,  
711 while these strategies for dealing with threat can be adaptive in certain  
712 circumstances, individuals with PTSD can become stuck in using only one  
713 strategy when other approaches would be more constructive. Behavioral  
714 aggression may take many forms, including aggression toward others,  
715 passive-aggressive behavior (e.g., complaining, "backstabbing,"  
716 deliberately being late or doing a poor job), or self-aggression (self-  
717 destructive activities, self-blame, being chronically hard on oneself, self-  
718 injury).
- 719 • **Thoughts and Beliefs:** The thoughts or beliefs that people have to help  
720 them understand and make sense of their environment can often  
721 overexaggerate threat. Often the individual is not fully aware of these  
722 thoughts and beliefs, but they cause the person to perceive more hostility,  
723 danger, or threat than others might feel is necessary. For example, a  
724 combat veteran may become angry when others around him (wife,  
725 children, coworkers) don't "follow the rules." The strength of his belief is  
726 actually related to how important it was for him to follow rules during the  
727 war in order to prevent deaths. Often, traumatized persons are not aware  
728 of the ways their beliefs are related to past trauma. For instance, by  
729 acting inflexibly toward others because of their need to control their  
730 environment, they can provoke others into becoming hostile, which  
731 creates a self-fulfilling prophecy. Common thoughts people with PTSD  
732 have include: "You can't trust anyone," "If I got out of control, it would be  
733 horrible/life-threatening/intolerable," "After all I've been through, I  
734 deserve to be treated better than this," and "Others are out to get me, or  
735 won't protect me, in some way."

736 **How can individuals with posttraumatic anger get help?**

737 In anger management treatment, arousal, behavior, and thoughts/beliefs are all  
738 addressed in different ways. Cognitive-behavioral treatment, a commonly utilized  
739 therapy that shows positive results when used to address anger, applies many  
740 techniques to manage these three anger components:

- 741 • For **increased arousal**, the goal of treatment is to help the person learn  
742 skills that will reduce overall arousal. Such skills include relaxation, self-  
743 hypnosis, and physical exercises that discharge tension.
  
- 744 • For **behavior**, the goal of treatment is to review a person's most frequent  
745 ways of behaving under perceived threat or stress and help him or her to  
746 expand the possible responses. More adaptive responses include taking a time  
747 out; writing thoughts down when angry; communicating in more verbal,  
748 assertive ways; and changing the pattern "act first, think later" to "think first,  
749 act later."
  
- 750 • For **thoughts/beliefs**, individuals are given assistance in logging,  
751 monitoring, and becoming more aware of their own thoughts prior to  
752 becoming angry. They are additionally given alternative, more positive  
753 replacement thoughts for their negative ideas (e.g., "Even if I am out of  
754 control, I won't be threatened in this situation," or "Others do not have to be  
755 perfect in order for me to survive/be comfortable"). Individuals often role-  
756 play situations in therapy so they can practice recognizing their anger-  
757 arousing thoughts and applying more positive thoughts.

758 There are many strategies for helping individuals with PTSD deal with the  
759 frequent increase of anger they are likely to experience. Most individuals have a  
760 combination of the three anger components listed above, and treatment aims to  
761 help with all aspects of anger. One important goal of treatment is to improve a  
762 person's sense of flexibility and control so that he or she does not feel re-  
763 traumatized by his or her own explosive or excessive responses to anger triggers.  
764 Treatment is also meant to have a positive impact on personal and work  
765 relationships.



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**APPENDICES**

|  |            |
|--|------------|
| <b>Appendix A. Guideline Development Process</b> | <b>210</b> |
| <b>Appendix B. Acronym List</b>                  | <b>217</b> |
| <b>Appendix C. PTSD Screening Tools</b>          | <b>220</b> |
| <b>Appendix D. Participant List</b>              | <b>224</b> |
| <b>Appendix E. Bibliography</b>                  | <b>230</b> |

14  
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## APPENDIX A: Guideline Development Process

20 The update of the VA/DoD Clinical Practice Guideline for Management of Post-Traumatic Stress was  
21 developed following the steps described in “Guideline for Guidelines,” an internal working document of  
22 the VA/DoD Evidence Based Practice Working Group, that requires an ongoing review of guideline works  
23 in progress.

24  
25 The Offices of Quality Performance and Patient Care Services of the VA, and the Army Medical Command  
26 of the DoD identified clinical leaders to champion the guideline development process. During a  
27 preplanning conference call, the clinical leaders defined the scope of the guideline and identified a group of  
28 clinical experts from the VA and DoD to form the Management of Post-Traumatic Stress Working Group  
29 (WG). For this guideline these WG participants were drawn from the fields of primary care, psychiatry,  
30 psychology, internal medicine, pharmacology, nursing, and social work.

31 The WG participated in 2 face-to-face meetings to reach consensus about the guideline algorithm and  
32 evidence-based recommendations and to prepare a draft update document. The draft continued to be revised  
33 by the Working Group through numerous conference calls and individual contributions to the document.

34 Recommendations for the management of post-traumatic stress were derived through a rigorous  
35 methodological approach that included the following:

- 36 • Determining appropriate criteria such as effectiveness, efficacy, population benefit, or patient  
37 satisfaction
- 38 • Reviewing literature to determine the strength of the evidence in relation to these criteria
- 39 • Formulating the recommendations and grading the level of evidence supporting the  
40 recommendation

41 After orientation to the goals and scope of the guideline update, the WG developed a set of 13 researchable  
42 questions within the focus areas of the guideline and identified associated key terms. For this guideline, two  
43 sets of questions were developed. The First (A) addressed *acute and early intervention aimed at/prevention*  
44 *of PTSD in adults with recent exposure to trauma*. The second set (B) focused on *therapy of adult patients*  
45 *with PTSD to achieve resolution of symptoms and functional outcome*. This approach ensured that the  
46 guideline development work outside of meetings focused on issues that practitioners considered important  
47 and also produced criteria for the literature search and selection of included studies that formed the body of  
48 evidence for this guideline update.

49  
50 All questions specified (adapted from the Evidence-Based Medicine toolbox, Center for Evidence-Based  
51 Medicine, [<http://www.cebm.net> ]):

- 52 • Population – Characteristics of the target patient population
- 53 • Intervention – Exposure, diagnostic, or prognosis
- 54 • Comparison – Intervention, exposure, or control used for comparison
- 55 • Outcome – Outcomes of interest

56 These specifications served as the preliminary criteria for selecting studies. See *PICO Questions to Guide*  
57 *Literature Search* for a complete listing and categorization of the questions (*end of this appendix*).

### 58 Literature Search

59 An initial global literature search yielded 59 systematic reviews/meta-analyses addressing  
60 pharmacotherapy, psychotherapy, combination, enhancement, complementary and other topics. One  
61 hundred and seventy eight (178) RCTs were found on the same subjects. Twenty-four controlled trials (CT)  
62 addressed combination, enhancement, and other areas. Refinement of the review process with input from

63 the WG members resulted in the studies being identified that met the baseline criteria for inclusion,  
64 addressed one or more of the researchable questions, and covered topic areas that had either not been  
65 addressed in the previous version of this guideline or had been included but not fully developed . A more  
66 detailed (full) search was conducted on each question, supplemented by hand searches and cross-  
67 referencing to search for relevant articles. The searches for these questions covered the period since the  
68 publication of the first VA/DoD CPG on management of post-traumatic stress (between January 1, 2002  
69 and August, 2009).

## 70 **Selection of Evidence**

71 The evidence selection process was designed to identify the best available evidence to address each key  
72 question and ensure maximum coverage of studies at the top of the hierarchy of study types. Published,  
73 peer-reviewed RCTs, as well as meta-analyses and systematic reviews that included randomized controlled  
74 studies, were considered to constitute the strongest level of evidence in support of guideline  
75 recommendations. This decision was based on the judgment that RCTs provide the clearest, most  
76 scientifically sound basis for judging comparative efficacy. The WG also recognized the limitations of  
77 RCTs, particularly considerations of generalizability with respect to patient selection and treatment quality.  
78 When available, the search sought out critical appraisals already performed by others that described explicit  
79 criteria for deciding what evidence was selected and how it was determined to be valid. The sources that  
80 have already undergone rigorous critical appraisal include Cochrane Reviews, Best Evidence, Technology  
81 Assessment, AHRQ systematic evidence reports, and other published Evidence-based Clinical Practice  
82 Guidelines.

83 The following databases were searched: Medline/Pubmed, Embase, PsycINFO, OVID, PILOT, and  
84 Cochrane Central Register of Controlled Trials. Limits were set for language (English), and type of  
85 research (RCT, systematic reviews including EPC and HTA reviews and meta-analyses). For prognostic  
86 and diagnostic questions (e.g., does test improve outcome?); cohort or other prospective non-RCT designs  
87 were considered.

88 The following inclusion criteria were used to select the articles identified in the literature search for  
89 possible inclusion:

- 90 • Published in United States, United Kingdom, Europe, Australia, Japan, New Zealand
- 91 • Full articles only published in English
- 92 • Study populations: age limited to adults 18 years of age or older; all races, ethnicities, and cultural  
93 groups
- 94 • Relevant outcomes able to be abstracted from the data presented in the articles
- 95 • Sample sizes appropriate for the study question addressed in the paper. RCTs were included if they  
96 were initiated with 30 or more participants

## 97 **Preparation of Evidence Tables (Reports) and Evidence Rating**

98 The results of the searches were organized in evidence reports, and copies of the original studies were  
99 provided to the WG for further analysis. Each reference was appraised for scientific merit, clinical  
100 relevance, and applicability to the populations served by the VA and DoD health care systems.

## 101 **Recommendation and Quality Rating**

102 Evidence-based practice involves integrating clinical expertise with the best available clinical evidence  
103 derived from systematic research.

104 A group of research analysts read and coded each article that met inclusion criteria. The articles were  
105 assessed for methodological rigor and clinical importance. Clinical experts from the VA and DoD WG  
106 reviewed the results and evaluated the strength of the evidence, considering quality of the body of evidence  
107 (made up of the individual studies) and the significance of the net benefit (potential benefit minus possible  
108 harm) for each intervention.

109 The overall strength of each body of evidence that addresses a particular Key Question was assessed using  
110 methods adapted from the U.S. Preventive Services Task Force (Harris, 2001). To assign an overall quality  
111 [QE] (see [Table A-2](#)) of the evidence (good, fair, or poor), the number, quality, and size of the studies;  
112 consistency of results between studies; and directness of the evidence were considered. Consistent results  
113 from a number of higher-quality studies [LE] (see [Table A-1](#)) across a broad range of populations; supports  
114 with a high degree of certainty that the results of the studies are true and therefore the entire body of

115 evidence would be considered “good” quality. A “fair” quality was assigned to the body of evidence  
116 indicating that the results could be due to true effects or to biases present across some or all of the studies.  
117 For a “poor” quality body of evidence, any conclusion is uncertain due to serious methodological  
118 shortcomings, sparse data, or inconsistent results.

119 The Strength of Recommendation [SR] was then determined based on the Quality of the Evidence [QE],  
120 and the clinical significance of the net benefit [NB] (see [Table A-3](#)) for each intervention, as demonstrated  
121 by the body of evidence. Thus, the grade (i.e., A, B, C, D or I) assigned to guideline recommendations  
122 reflect both variables; the Quality of the evidence and the potential clinical benefit that the intervention  
123 may provide to patients (see [Table A4](#)).  
124

| <b>Table A-1: Level of Evidence (LE)</b> |  |
|--|--|
| <b>I</b>                                 | At least one properly done RCT   |
| <b>II-1</b>                              | Well-designed controlled trial without randomization   |
| <b>II-2</b>                              | Well-designed cohort or case-control analytic study, preferably from more than one source            |
| <b>II-3</b>                              | Multiple time series evidence with/without intervention, dramatic results of uncontrolled experiment |
| <b>III</b>                               | Opinion of respected authorities, descriptive studies, case reports, and expert committees           |

125

| <b>Table A-2: Overall Quality [QE]</b> |  |
|--|--|
| <b>Good</b>                            | High grade evidence (I or II-1) directly linked to health outcome  |
| <b>Fair</b>                            | High grade evidence (I or II-1) linked to intermediate outcome;<br><i>or</i><br>Moderate grade evidence (II-2 or II-3) directly linked to health outcome |
| <b>Poor</b>                            | Level III evidence or no linkage of evidence to health outcome   |

126

| <b>Table A-3: Net Effect of the Intervention [NB]</b> |  |
|---|--|
| Substantial   | More than a small relative impact on a frequent condition with a substantial burden of suffering;<br><i>or</i><br>A large impact on an infrequent condition with a significant impact on the individual patient level.       |
| Moderate  | A small relative impact on a frequent condition with a substantial burden of suffering;<br><i>or</i><br>A moderate impact on an infrequent condition with a significant impact on the individual patient level.              |
| Small   | A negligible relative impact on a frequent condition with a substantial burden of suffering;<br><i>or</i><br>A small impact on an infrequent condition with a significant impact at the individual patient level.            |
| Zero or Negative                                      | Negative impact on patients;<br><i>or</i><br>No relative impact on either a frequent condition with a substantial burden of suffering, or an infrequent condition with a significant impact on the individual patient level. |

127

| <b>Table A-4: Final Grade of Recommendation [SR]</b> |  |          |       |                  |
|--|--|----------|-------|------------------|
|  | <i>The net benefit of the intervention</i> |          |       |                  |
| <b>Quality of Evidence</b>                           | Substantial                                | Moderate | Small | Zero or Negative |
| <b>Good</b>  | A  | B        | C     | D                |
| <b>Fair</b>  | B  | B        | C     | D                |
| <b>Poor</b>  | I  | I        | I     | I                |

128  
129

**Strength of Recommendation Rating [SR]**

|   |  |
|---|--|
| A | A strong recommendation that the clinicians provide the intervention to eligible patients.<br><i>Good evidence was found that the intervention improves important health outcomes and concludes that benefits substantially outweigh harm.</i>   |
| B | A recommendation that clinicians provide (the service) to eligible patients.<br><i>At least fair evidence was found that the intervention improves health outcomes and concludes that benefits outweigh harm.</i>  |
| C | No recommendation for or against the routine provision of the intervention is made.<br><i>At least fair evidence was found that the intervention can improve health outcomes, but concludes that the balance of benefits and harms is too close to justify a general recommendation.</i> |
| D | Recommendation is made against routinely providing the intervention to asymptomatic patients.<br><i>At least fair evidence was found that the intervention is ineffective or that harms outweigh benefits.</i>   |
| I | The conclusion is that the evidence is insufficient to recommend for or against routinely providing the intervention.<br><i>Evidence that the intervention is effective is lacking, of poor quality, or conflicting, and the balance of benefits and harms cannot be determined.</i>     |

130

**Algorithm Format**

131

The clinical algorithm incorporates the information presented in the guideline in a format which maximally facilitates clinical decision-making. The use of the algorithmic format was chosen because of evidence showing that such a format improves data collection, facilitates diagnostic and therapeutic decision-making, and changes patterns of resource use.

132

133

134

135

The algorithmic format allows the provider to follow a linear approach to critical information needed at the major decision points in the clinical process and includes:

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137

- An ordered sequence of steps of care

138

- Recommended observations

139

- Decisions to be considered

140

- Actions to be taken

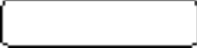



141

A clinical algorithm diagrams a guideline into a step-by-step decision tree. Standardized symbols are used to display each step in the algorithm (Society for Medical Decision-Making Committee, 1992). Arrows connect the numbered boxes indicating the order in which the steps should be followed.

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|   |  |
|---|--|
|  | Rounded rectangles represent a clinical state or condition.  |
|  | Hexagons represent a decision point in the guideline, formulated as a question that can be answered Yes or No. A horizontal arrow points to the next step if the answer is YES. A vertical arrow continues to the next step for a negative answer. |
|  | Rectangles represent an action in the process of care.   |
|  | Ovals represent a link to another section within the guideline.  |

145

146

A letter within a box of an algorithm refers the reader to the corresponding annotation. The annotations elaborate on the recommendations and statements that are found within each box of the algorithm.

147

148 Included in the annotations are brief discussions that provide the underlying rationale and specific evidence  
149 tables. Annotations indicate whether each recommendation is based on scientific data or expert opinion. A  
150 complete bibliography is included in the guideline.

151

#### 152 **Lack of Evidence – Consensus of Experts**

153 Where existing literature was ambiguous or conflicting, or where scientific data was lacking on an issue,  
154 recommendations were based on the clinical experience of the Working Group.

155 This update of the Stroke Rehabilitation Guideline is the product of many months of diligent effort and  
156 consensus building among knowledgeable individuals from the VA, DoD, and academia, as well as  
157 guideline facilitators from the private sector. An experienced moderator facilitated the multidisciplinary  
158 Working Group. The list of participants is included in [Appendix D](#)

159

#### 160 **REFERENCES**

161 Agency for Health Care Policy and Research (AHCPR). Manual for conducting systematic review. August  
162 1996. Prepared by Steven H. Woolf.

163 Harris RP, Helfand M, Woolf SH, Lohr KN, Mulrow CD, Teutsch SM, Atkins D; Methods Work Group,  
164 Third US Preventive Services Task Force, Current methods of the U.S. Preventive Services Task  
165 Force: a review of the process. Am J Prev Med 2001 Apr;20(3 Suppl):21-35. Available at;  
166 <http://www.ahrq.gov/clinic/ajpmsuppl/harris1.htm>

167 Society for Medical Decision-Making Committee (SMDMC). Proposal for clinical algorithm standards,  
168 SMDMC on Standardization of Clinical Algorithms. Med Decis Making 1992 Apr-Jun;  
169 12(2):149-54.

170 United States Preventive Service Task Force (USPSTF). Guide to clinical preventive services. 2nd edition.  
171 Washington, DC: US Department of Health and Human Services, Office of Disease Prevention  
172 and Health Promotion, 1996.

173 Woolf SH. Practice guidelines, a new reality in medicine II; Methods of developing guidelines. Arch  
174 Intern Med 1992 May; 152(5):946-52.

175

**Management of Post-Traumatic Stress****UPDATE 2010 PICO QUESTIONS****A. Acute Intervention / Prevention in adults with recent exposure to trauma or diagnosed with ASD**

1. Is **debriefing** more effective than no intervention or any other intervention for prevention of PTS disorder?
2. Is **pharmacotherapy** more effective than no intervention or any other intervention for prevention of full PTS disorder?
  - a. **alpha-blockers**
  - b. **beta-blockers**
  - c. **Sympatolitic**
  - d. **DCS and CBT**
3. Are any **psychotherapy techniques** more effective than no intervention or any other intervention for prevention of full PTS disorder?
4. Is **psychoeducation** more effective than no intervention or any other intervention for prevention of full PTS disorder?
5. Are any **Complimentary Alternative Medicine (CAM) approaches** more effective than no intervention or any other intervention for prevention of full PTS disorder?
6. Is **early intervention** more effective than **later intervention** for prevention of full PTS disorder?
7. Is **combination** of pharmacotherapy and psychotherapy more effective than no intervention or any other intervention for prevention of full PTS disorder?
8. Is **peer counseling** more effective than **counseling** by an outside team for prevention of full PTS disorder?
9. Is **outreach (screening, repeated screening)** more effective than no intervention or any other intervention for prevention of full PTS disorder?

202

203 **B. Treatment for PTSD**

204

205

206 **Which of the following treatment interventions for adult patients with PTSD lead to achieve Resolution**  
 207 **of symptoms and Functional outcome?** (Consider effectiveness in special population (e.g., Gender, Combat  
 208 veterans, Elderly)

209

**10. Psychotherapy Techniques:**

210

Is **prolonged exposure** more effective interventions in the treatment of PTSD?

211

Is EMDR more effective than other interventions in the treatment of PTSD?

212

Is cognitive processing therapy more effective than other interventions in the treatment of  
 213 PTSD?

214

Is DBT, MBCT, ACT or mindfulness more effective than other interventions in the treatment  
 215 of PTSD?

216

Psychoeducation (Battlemind, stress control) more effective than other interventions in the  
 217 treatment of PTSD?

218

**11. Pharmacotherapy Classes:**

219

-MAOI and TCAs

220

-SSRIs

221

-SNRIs

222

-DNRI

223

-Novel antidepressant (trazodone, nefazodone)

224

-Conventional antipsychotics

225

-Atypical antipsychotics

226

-Anticonvulsants

227

-Anxiolytic (Benzodiazepine)

228

-Sedative hypnotics (for sleep)

229

-Antiadrenergics

230

231

**12. Somatic:**

232

-ECT

233

-rTMS

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**13. Complementary Alternative Medicine (CAM)**

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-Acupuncture

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-Meditation

237

-Herbal, food suppl.,

238

-Yoga, Tai Chi

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242**APPENDIX B**  
**Acronym List**

|        |   |
|--------|---|
| ABCs   | Airway, breathing, circulation                                  |
| AHCPR  | Agency for Healthcare Policy and Research                       |
| APA    | American Psychiatric Association                                |
| ASD    | Acute stress disorder   |
| ASR    | Acute stress reaction   |
| AUDIT  | Alcohol Use Disorders Identification Test                       |
| BEP    | Brief Eclectic Psychotherapy                                    |
| BFT    | Behavioral Family Therapy                                       |
| BL     | baseline;   |
| CAGE   | Alcohol abuse/dependence screening test mnemonic                |
| CAPS   | Clinician-Administered PTSD Scale;                              |
| CAPS   | Clinician Administered PTSD Scale                               |
| CAPS-1 | Clinician-Administered PTSD Scale1-month version;               |
| CAPS-2 | Clinician-Administered PTSD Scale1-week version;                |
| CAPS-2 | Clinician-Administered PTSD Scale Part 2;                       |
| CAPS-D | Clinician-Administered PTSD Scale hyperarousal subscale;        |
| CBC    | Complete blood count  |
| CBT    | Cognitive Behavioral Therapy                                    |
| CCTR   | Cochrane Central Register of Controlled Trials                  |
| CDR    | Commander   |
| CGI    | Clinical Global Impression;                                     |
| CGI-I  | Clinical Global Impression-Improvement;                         |
| CGI-S  | Clinical Global Assessment of Severity;                         |
| CGIC   | Clinical Global Impression of Change;                           |
| CI     | confidence interval;  |
| CISD   | Critical Incident Stress Debriefing                             |
| CNS    | Central nervous system  |
| COSR   | Combat and operational stress reactions                         |
| CPT    | Cognitive Processing Therapy                                    |
| CPT-C  | CPT-Cognitive   |
| CT     | Cognitive Therapy   |
| CTT    | Cognitive Trauma Therapy  |
| CV     | Cardiovascular  |
| DARE   | Database of Abstracts of Reviews of Effectiveness               |
| DAST   | Drug Abuse/Dependence Screener                                  |
| DBT    | Dialectical Behavioral Therapy                                  |
| DCS    | D-cycloserine;  |
| DoD    | Department of Defense   |
| DSM-IV | Diagnostic and Statistical Manual of Mental Disorders (4th ed.) |
| DTS    | Davidson Trauma Scale;  |
| dx     | diagnosis   |
| EBM    | Evidence-based medicine   |
| EBPTU  | Evaluation and Brief PTSD Treatment Unit                        |
| ED     | Emergency Department;   |
| EEG    | Electroencephalography  |

|                    |   |
|--------------------|---|
| EKG                | Electrocardiogram   |
| EMDR               | Eye Movement Desensitization and Reprocessing               |
| EMTs               | Emergency Medical Teams                                     |
| ES                 | effect size   |
| ESRT               | Emotional Self-Regulation Therapy                           |
| ET                 | Exposure Therapy  |
| EtoH               | Ethanol   |
| FDA                | U. S. Food and Drug Administration                          |
| GAF                | Global Assessment of Function                               |
| GI                 | Gastrointestinal  |
| grp(s)             | group(s);   |
| GT                 | group therapy   |
| GU                 | Genitourinary   |
| HIV                | Human immunodeficiency virus                                |
| Interapy           | Internet Therapy  |
| IRT                | Imagery Rehearsal Therapy                                   |
| IRT                | Image Rehearsal Therapy                                     |
| ITT                | Intention to Treat  |
| LOC                | Level of consciousness                                      |
| LOF                | Level of function   |
| MAOIs              | Monoamine oxidase inhibitors                                |
| MAST               | Michigan Alcohol Screening Test                             |
| MDD                | Major Depressive Disorder                                   |
| MHP                | Mental health providers                                     |
| MI                 | Myocardial infarction                                       |
| MISS               | Mississippi Scale for Combat-Related PTST-civilian version; |
| MMSE               | Mini-Mental State Examination                               |
| MRI                | Magnetic resonance imaging                                  |
| MSE                | Mental status examination                                   |
| MVA                | motor vehicle accident;                                     |
| N/R                | not reported;   |
| NET                | Narrative Exposure Therapy (a form of Exposure Therapy)     |
| NIMH               | National Institute of Mental Health                         |
| NS                 | Nervous system  |
| OMO                | Ongoing military operations                                 |
| OTC                | Over-the-counter  |
| PCL                | Posttraumatic Stress Disorder Checklist;                    |
| PCL-C              | PTSD Checklist – Civilian Version                           |
| PCL-M              | Patient Checklist for PTSD-Military Version;                |
| PCL-M              | PTSD Checklist – Military Version                           |
| PCL-S              | PTSD Checklist – Stressor Specific Version                  |
| PCP                | Primary care provider                                       |
| PE                 | Physical examination  |
| PE (Interventions) | Prolonged Exposure  |
| PIES               | Proximity, Immediacy, Expectancy, Simplicity                |
| PSQI               | Pittsburgh Sleep Quality Index                              |
| PsychEd            | Psychological Education                                     |
| pt(s)              | patient(s);   |
| PTSD               | posttraumatic stress disorder;                              |

|          |  |
|----------|--|
| PTSD     | Post-traumatic Stress Disorder                 |
| QE       | Quality of evidence                            |
| RA       | Repeated Assessment                            |
| RCS      | Readjustment Counseling Services               |
| RCT      | Randomized controlled trial                    |
| RLX      | Relaxation Training                            |
| RTD      | Return-to-duty                                 |
| SC       | Supportive Counseling                          |
| SC       | Supportive Counseling                          |
| SIADH    | Syndrome of inappropriate antidiuretic hormone |
| SIPU     | Specialized Inpatient PTSD Unit                |
| SIT      | Stress Inoculation Therapy                     |
| SM       | Service member                                 |
| SR       | Strength of recommendation                     |
| SSRI     | Selective Serotonin Reuptake Inhibitors        |
| SUD      | Substance Use Disorder                         |
| SUNY     | State University of New York                   |
| TAU      | Treatment as Usual                             |
| TCAs     | Tricyclic Antidepressants                      |
| TOP-8    | Treatment Outcome PTSD rating scale;           |
| TSH      | Thyroid Stimulating Hormone                    |
| Tx or RX | treatment                                      |
| USPSTF   | U.S. Preventive Service Task Force             |
| VA       | Veterans Affairs                               |
| VAMC     | Veterans Affairs Medical Center                |
| Vets     | Veterans                                       |
| VHA      | Veterans Health Administration                 |
| WL       | wait list                                      |

**APPENDIX C**  
**PTSD Screening Tools**

**Primary Care PTSD Screen (PC-PTSD)**

The table below shows the Primary Care PTSD Screen (PC-PTSD) that has been designed for use in primary care and other medical settings. The PC-PTSD is brief and problem-focused. The screen does *not* include a list of potentially traumatic events. There are two reasons for this:

- Studies on trauma and health in both male and female patients suggest that the active mechanism linking trauma and physical health is the diagnosis of PTSD. In other words, the relationship between trauma and health appears to be mediated through a current PTSD diagnosis.
- A symptom-driven screen, rather than a trauma-focused screen, is attractive to primary care staff who may not be able to address a patient's entire trauma history during their visit with the patient. Such a trauma inquiry might be especially problematic with a VA population where the average number of traumatic events meeting criterion A for PTSD is over four.

A positive response to the screen does not necessarily indicate that a patient has Post-traumatic Stress Disorder. However, a positive response does indicate that a patient *may* have PTSD or trauma-related problems and further investigation of trauma symptoms by a mental-health professional may be warranted.

| <b>Primary Care PTSD Screen</b>   |                                |
|---|--------------------------------|
| <b>In your life, have you ever had any experience that was so frightening, horrible, or upsetting that, <i>in the past month</i>, you...</b>  |                                |
| 1. Have had nightmares about it or thought about it when you did not want to?   | <b>YES                  NO</b> |
| 2. Tried hard not to think about it or went out of your way to avoid situations that reminded you of it?                                      | <b>YES                  NO</b> |
| 3. Were constantly on guard, watchful, or easily startled?  | <b>YES                  NO</b> |
| 4. Felt numb or detached from others, activities, or your surroundings?   | <b>YES                  NO</b> |
| <i>Current research suggests that the results of the PC-PTSD should be considered "positive" if a patient answers "yes" to any two items.</i> |                                |

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## PTSD CheckList – Civilian Version (PCL-C)

**Patient’s Name:** \_\_\_\_\_

Instruction to patient: Below is a list of problems and complaints that veterans sometimes have in response to stressful life experiences. Please read each one carefully, put an “X” in the box to indicate how much you have been bothered by that problem *in the last month*.

| No. | Response:   | Not at all (1) | A little bit (2) | Moderately (3) | Quite a bit (4) | Extremely (5) |
|-----|---|----------------|------------------|----------------|-----------------|---------------|
| 1.  | Repeated, disturbing <i>memories, thoughts, or images</i> of a stressful experience from the past?  |                |                  |                |                 |               |
| 2.  | Repeated, disturbing <i>dreams</i> of a stressful experience from the past?   |                |                  |                |                 |               |
| 3.  | Suddenly <i>acting or feeling</i> as if a stressful experience <i>were happening again</i> (as if you were reliving it)?  |                |                  |                |                 |               |
| 4.  | Feeling <i>very upset</i> when <i>something reminded</i> you of a stressful experience from the past?   |                |                  |                |                 |               |
| 5.  | Having <i>physical reactions</i> (e.g., heart pounding, trouble breathing, or sweating) when <i>something reminded</i> you of a stressful experience from the past? |                |                  |                |                 |               |
| 6.  | Avoid <i>thinking about</i> or <i>talking about</i> a stressful experience from the past or avoid <i>having feelings</i> related to it?                             |                |                  |                |                 |               |
| 7.  | Avoid <i>activities or situations</i> because <i>they remind you</i> of a stressful experience from the past?   |                |                  |                |                 |               |
| 8.  | Trouble <i>remembering important parts</i> of a stressful experience from the past?   |                |                  |                |                 |               |
| 9.  | Loss of <i>interest in things that you used to enjoy</i> ?  |                |                  |                |                 |               |
| 10. | Feeling <i>distant</i> or <i>cut off</i> from other people?   |                |                  |                |                 |               |
| 11. | Feeling <i>emotionally numb</i> or being unable to have loving feelings for those close to you?   |                |                  |                |                 |               |
| 12. | Feeling as if your <i>future</i> will somehow be <i>cut short</i> ?   |                |                  |                |                 |               |
| 13. | Trouble <i>falling or staying asleep</i> ?  |                |                  |                |                 |               |
| 14. | Feeling <i>irritable</i> or having <i>angry outbursts</i> ?   |                |                  |                |                 |               |
| 15. | Having <i>difficulty concentrating</i> ?  |                |                  |                |                 |               |
| 16. | Being “ <i>super alert</i> ” or watchful on guard?  |                |                  |                |                 |               |
| 17. | Feeling <i>jumpy</i> or easily startled?  |                |                  |                |                 |               |

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Weathers, F.W., Huska, J.A., Keane, T.M. *PCL-C for DSM-IV*. Boston: National Center for PTSD – Behavioral Science Division, 1991.

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## PTSD CheckList – Military Version (PCL-M)

Patient's Name: \_\_\_\_\_

Instruction to patient: Below is a list of problems and complaints that veterans sometimes have in response to stressful military experiences. Please read each one carefully, put an "X" in the box to indicate how much you have been bothered by that problem in *the last month*.

| No. | Response:  | Not at<br>all (1) | A little<br>bit (2) | Moderately<br>(3) | Quite a<br>bit (4) | Extremely<br>(5) |
|-----|--|-------------------|---------------------|-------------------|--------------------|------------------|
| 1.  | Repeated, disturbing <i>memories, thoughts, or images</i> of a stressful military experience?  |                   |                     |                   |                    |                  |
| 2.  | Repeated, disturbing <i>dreams</i> of a stressful military experience?   |                   |                     |                   |                    |                  |
| 3.  | Suddenly <i>acting or feeling</i> as if a stressful military experience <i>were happening again</i> (as if you were reliving it)?                              |                   |                     |                   |                    |                  |
| 4.  | Feeling <i>very upset</i> when <i>something reminded</i> you of a stressful military experience?   |                   |                     |                   |                    |                  |
| 5.  | Having <i>physical reactions</i> (e.g., heart pounding, trouble breathing, or sweating) when <i>something reminded</i> you of a stressful military experience? |                   |                     |                   |                    |                  |
| 6.  | Avoid <i>thinking about</i> or <i>talking about</i> a stressful military experience or avoid <i>having feelings</i> related to it?                             |                   |                     |                   |                    |                  |
| 7.  | Avoid <i>activities</i> or <i>situations</i> because <i>they remind you</i> of a stressful military experience?  |                   |                     |                   |                    |                  |
| 8.  | Trouble <i>remembering important parts</i> of a stressful military experience?   |                   |                     |                   |                    |                  |
| 9.  | Loss of <i>interest in things that you used to enjoy</i> ?   |                   |                     |                   |                    |                  |
| 10. | Feeling <i>distant</i> or <i>cut off</i> from other people?  |                   |                     |                   |                    |                  |
| 11. | Feeling <i>emotionally numb</i> or being unable to have loving feelings for those close to you?  |                   |                     |                   |                    |                  |
| 12. | Feeling as if your <i>future</i> will somehow be <i>cut short</i> ?  |                   |                     |                   |                    |                  |
| 13. | Trouble <i>falling</i> or <i>staying asleep</i> ?  |                   |                     |                   |                    |                  |
| 14. | Feeling <i>irritable</i> or having <i>angry outbursts</i> ?  |                   |                     |                   |                    |                  |
| 15. | Having <i>difficulty concentrating</i> ?   |                   |                     |                   |                    |                  |
| 16. | Being " <i>super alert</i> " or watchful on guard?   |                   |                     |                   |                    |                  |
| 17. | Feeling <i>jumpy</i> or easily startled?   |                   |                     |                   |                    |                  |

Weathers, F.W., Huska, J.A., Keane, T.M. *PCL-M for DSM-IV*. Boston: National Center for PTSD – Behavioral Science Division, 1991.

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**PTSD CheckList – Stressor Specific Version (PCL-S)**

The event you experienced was: \_\_\_\_\_ on: \_\_\_\_\_

Instruction to patient: Below is a list of problems and complaints that veterans sometimes have in response to stressful military experiences. Please read each one carefully, put an “X” in the box to indicate how much you have been bothered by that problem in *the last month*.

| No. | Response:   | Not at all (1) | A little bit (2) | Moderately (3) | Quite a bit (4) | Extremely (5) |
|-----|---|----------------|------------------|----------------|-----------------|---------------|
| 1.  | Repeated, disturbing <i>memories, thoughts, or images</i> of the stressful experience?  |                |                  |                |                 |               |
| 2.  | Repeated, disturbing <i>dreams</i> of the stressful experience?   |                |                  |                |                 |               |
| 3.  | Suddenly <i>acting or feeling</i> as if the stressful experience <i>were happening again</i> (as if you were reliving it)?                              |                |                  |                |                 |               |
| 4.  | Feeling <i>very upset</i> when <i>something reminded</i> you of the stressful experience?   |                |                  |                |                 |               |
| 5.  | Having <i>physical reactions</i> (e.g., heart pounding, trouble breathing, or sweating) when <i>something reminded</i> you of the stressful experience? |                |                  |                |                 |               |
| 6.  | Avoid <i>thinking about or talking about the</i> stressful experience or avoid <i>having feelings</i> related to it?                                    |                |                  |                |                 |               |
| 7.  | Avoid <i>activities or situations</i> because <i>they remind you</i> of the stressful experience?   |                |                  |                |                 |               |
| 8.  | Trouble <i>remembering important parts</i> of the stressful experience?   |                |                  |                |                 |               |
| 9.  | Loss of <i>interest in things that you used to enjoy</i> ?  |                |                  |                |                 |               |
| 10. | Feeling <i>distant or cut off</i> from other people?  |                |                  |                |                 |               |
| 11. | Feeling <i>emotionally numb</i> or being unable to have loving feelings for those close to you?   |                |                  |                |                 |               |
| 12. | Feeling as if your <i>future</i> will somehow be <i>cut short</i> ?   |                |                  |                |                 |               |
| 13. | Trouble <i>falling or staying asleep</i> ?  |                |                  |                |                 |               |
| 14. | Feeling <i>irritable</i> or having <i>angry outbursts</i> ?   |                |                  |                |                 |               |
| 15. | Having <i>difficulty concentrating</i> ?  |                |                  |                |                 |               |
| 16. | Being “ <i>super alert</i> ” or watchful on guard?  |                |                  |                |                 |               |
| 17. | Feeling <i>jumpy</i> or easily startled?  |                |                  |                |                 |               |

Weathers, F.W., Huska, J.A., Keane, T.M. *PCL-S for DSM-IV*. Boston: National Center for PTSD – Behavioral Science Division, 1991.*This is a Government document in the public domain.*

**APPENDIX D**  
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**APPENDIX E**  
**Bibliography**

*[Will be completed in the final version]*