Relationship of screen-based symptoms for mild traumatic brain injury and mental health problems in Iraq and Afghanistan veterans: Distinct or overlapping symptoms?

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Abstract—This study used factor analytic techniques to differentiate distinct from overlapping screen-based symptoms of traumatic brain injury (TBI), posttraumatic stress disorder (PTSD), and depression in Iraq and Afghanistan veterans. These symptoms were derived from screen results of 1,549 veterans undergoing Department of Veterans Affairs postdeployment screening between April 2007 and January 2010. Veterans with positive TBI screens were approximately twice as likely to also screen positive for depression and PTSD (adjusted relative risks = 1.9 and 2.1, respectively). Irritability was a shared symptom between TBI and PTSD, and emotional numbing was a shared symptom between PTSD and depression. Symptoms unique to TBI included dizziness, headaches, memory problems, and light sensitivity. Four separate constructs emerged: TBI, PTSD, depression, and a fourth construct consisting of hypervigilance and sleep problems. These findings illuminate areas of overlap between TBI and common postdeployment mental health problems. Discriminating symptoms of TBI from mental health problems may facilitate diagnosis, triage to specialty care, and targeted symptom management. The emergence of a fourth factor consisting of sleep problems and hypervigilance highlights the need to attend to specific symptoms in the postdeployment screening process.

Key words: depression, factor analysis, hypervigilance, insomnia, mental health, OIF/OEF, posttraumatic stress disorder, screen, traumatic brain injury, veteran.

INTRODUCTION

Traumatic brain injury (TBI) has received greater public attention because of the conflicts in Iraq and Afghanistan (Operation Iraqi Freedom [OIF]/Operation Enduring Freedom [OEF]) [1]. An estimated 22 percent of servicemembers returning from OIF have reported experiencing TBIs and concussions [2], and of injured OIF/OEF servicemembers, 31 percent have been diagnosed with a TBI [3]. Although TBIs range in severity, mild TBI comprises roughly 77 percent of all head injuries.
among OIF/OEF veterans and is most challenging to diagnose because of symptom overlap with mental health disorders: posttraumatic stress disorder (PTSD) and depression, in particular [4–5].

In an attempt to increase early detection of mild TBI in returning combat veterans entering the Department of Veterans Affairs (VA) system, the VA implemented the first-level TBI screen in April 2007 [6]. The VA considers OIF/OEF veterans to be at risk for having had a possible mild TBI if, on the first-level VA TBI screen, they report a head injury with loss of consciousness or altered mental status and postconcussive symptoms (PCSs, immediately following the exposure and within 1 wk prior to screening) [7]. This definition is sensitive and identifies at-risk OIF/OEF veterans but is not necessarily specific for a TBI diagnosis because mental health conditions, such as PTSD and depression, have also led to positive responses on the VA TBI screen [8]. OIF/OEF veterans who screen positive on the VA first-level TBI screen are therefore referred for second-level TBI screening to conduct a more in-depth assessment for a history of a TBI exposure.

Given that TBI results from exposure to a traumatic event, such as a blast, symptoms of TBI commonly co-occur with both PTSD and depression [9]. TBI can increase the risk of developing PTSD and other mental health conditions because the event associated with the head injury was potentially life-threatening and was associated with other traumatic war zone exposures [10–11]. In recent studies among OIF/OEF veterans with mild TBI, an estimated 33 to 44 percent also met criteria for PTSD and 33 to 62 percent also met criteria for depression [12–14].

Particularly challenging is the fact that many of the cardinal features of mild TBI, such as memory problems, diminished attention and concentration, irritability, and insomnia, are also hallmark symptoms of PTSD and depression [10,12]. Schneiderman et al. found that PTSD was one of the factors most strongly associated with TBI-related PCSs [15]. Indeed, in one published study of OIF servicemembers with mild TBI, after controlling for the effects of PTSD, headaches were the only distinguishing feature associated with loss of consciousness in the battlefield [12]. Nevertheless, a gap exists in identifying other symptoms that differentiate TBI, PTSD, and depression.

The purpose of this study was to address this gap by identifying distinguishing features of each disorder in order to more accurately discriminate between screen-based symptoms of TBI, PTSD, and depression in OIF/OEF veterans. Because the symptoms of TBI, PTSD, and depression can overlap clinically, problems with triage and appropriate referral for specialty care can ensue, creating delays in diagnosis and treatment [15–17]. Specifically, using factor analytic techniques, we sought to identify symptoms from TBI, PTSD, and depression screens that were overlapping and, in contrast, symptoms that were uniquely associated with a positive TBI screen (and not better accounted for by PTSD or depression). We focused on VA screen results because they represent the level of information that clinicians use to make initial decisions about diagnosis and treatment of their patients. To our knowledge, no study to date has used the VA postdeployment mental health and TBI screens to evaluate symptom discrimination in OIF/OEF veterans. A better understanding of the symptoms that differentiate among TBI, PTSD, and depression in the initial screening stage may facilitate triage, more accurate diagnosis, and early symptom management of OIF/OEF veterans.

**METHODS**

**Data Source**

We used VA administrative data for OIF/OEF veterans who were screened for TBI at a VA medical center and five affiliated VA community-based outpatient clinics. We extracted responses to the VA TBI screen and the OIF/OEF postdeployment screen for PTSD and depression from the Veterans Health Information Systems and Technology Architecture. We derived demographic and military service information from the national VA OEF/OIF Roster [18].

**Sample Selection**

The study population consisted of 1,713 OIF/OEF veterans who received a first-level TBI screen from April 1, 2007, through January 8, 2010. Of these, we excluded 164 because their screening was completed at another facility and the results were unavailable (n = 86), they had previously received a TBI diagnosis (n = 67), they had refused screening (n = 10), or they had an incomplete screen (n = 1). Our final sample included 1,549 veterans. We found no significant demographic and military service characteristic differences between veterans who were included compared with those who were excluded.
from the study sample. For veterans who had completed more than one TBI screen on different dates \((n = 127)\), we used only the most recent and complete screen for this study. When examining the prevalence of mental health conditions, we further restricted the sample to veterans who completed a PTSD screen and depression screen \((n = 1,400 \text{ and } 1,407, \text{ respectively})\) within 1 year of their most recent TBI screen.

Next, in order to perform the more in-depth statistical analyses described later, we selected a subsample of veterans who had completed all four questions of the TBI screen (see “Measures” section for TBI screen structure) and who had completed both a PTSD and depression screen \((n = 358)\).

Measures

**Traumatic Brain Injury Screen**

The VA TBI screen is a five-section assessment tool modified from the Brief TBI Screen [19] to evaluate possible TBI in OIF/OEF veterans [20]. The first section of the TBI screen identifies veterans who were previously diagnosed with TBI. Veterans with prior TBI diagnoses were not administered the remainder of the TBI screen. Veterans without a prior TBI diagnosis were subsequently asked four sets of questions [7]. A positive screen for TBI (TBI+) consisted of endorsing one or more questions in each of the four sections: (1) a qualifying TBI event (blast or explosion, vehicular accident and/or crash, fragment wound or bullet wound above the shoulders, or fall), (2) immediate symptoms following the event (losing consciousness/knocked out; being dazed, confused, or seeing stars; not remembering the event; concussion; or head injury), (3) new or worsening TBI-related symptoms following the event (memory problems or lapses, balance problems or dizziness, light sensitivity, irritability, headaches, or sleep problems), and (4) current symptoms (same as in section 3). A negative screen for TBI (TBI–) occurred when a veteran failed to endorse at least one item in any of the four sections. Once a veteran failed to endorse any items in a particular section of the VA TBI screen, the screen was terminated.

**Posttraumatic Stress Disorder Screen**

The Primary Care Posttraumatic Stress Disorder Screen is a brief four-item screen designed to detect possible PTSD symptoms in primary care and other medical settings [21–22]. The screen yields binary responses (“yes” or “no”) for each of four PTSD symptom clusters: re-experiencing, avoidance, emotional numbing, and arousal. Endorsing three or more symptoms constituted a positive screen for PTSD [23].

**Depression Screen**

The Patient Health Questionnaire-2 [24] is a two-item depression screen adapted from the nine-item depression module of the Patient Health Questionnaire [25]. This shorter screen assessed symptoms of depressed mood (down and/or hopeless) and anhedonia (little interest) on a four-point Likert scale ranging from 0 (“not at all”) to 3 (“nearly every day”). Total scores range from 0–6. Veterans who obtained a total score of ≥3 screened positive for depression [24].

**Statistical Analysis**

We first compared characteristics of TBI+ and TBI– veterans using \(t\)-tests for continuous variables and Pearson chi-square tests for categorical variables. To compare PTSD and depression screens between TBI+ and TBI– veterans, we used log-binomial regression to estimate risk ratios and associated confidence intervals. We adjusted models for potential confounders, including sociodemographics and military factors. We conducted all analyses, other than factor analyses, using SAS version 9.2 (SAS Institute Inc; Cary, North Carolina).

**Exploratory Factor Analysis**

We conducted an exploratory factor analysis (EFA) in Mplus, version 5 (Muthén & Muthén; Los Angeles, California) in order to identify a viable factor structure of the 12 different symptoms included in the three screens: TBI screen (memory problems, balance problems or dizziness, light sensitivity, irritability, headaches, and sleep problems), PTSD screen (nightmares, avoidance, emotional numbing, and hypervigilance), and depression screen (little interest and down and/or hopeless). We used a robust weighted least-squares with mean and variance adjustment (WLSMV) method estimator and treated all 12 items as binary or ordinal variables. The WLSMV method estimator provided weighted least-square parameter estimates using a diagonal weight matrix and robust standard errors and a mean- and variance-adjusted chi-square test statistic. We considered the maximum number of factors (7) that were technically possible. We rotated the raw factor loadings using oblique (promax) rotation. We based model selection on consideration of eigenvalues,
which were the estimated residual variances, as well as the a priori hypothesis that the factor structure would reflect the clinical screens from which the items originated. We retained items with primary factor loadings of ≥0.4.

Confirmatory Factor Analysis

We then conducted a confirmatory factor analysis (CFA) to confirm the exploratory factor model and to assess whether the factor structure required modification. We used Mplus to estimate the confirmatory factor model, using the WLSMV method estimator. We used the derivatives for parameter estimates as modification indices in order to determine which parameters were most likely to improve model fit. We compared nested models using WLSMV estimates of chi-square and the derivatives difference test for change in model fit, which is a robust goodness-of-fit test where the chi-square is adjusted to obtain an accurate p-value [26]. We used the following goodness-of-fit indices to assess the degree of fit between the model and the data: Tucker Lewis Index (TLI; acceptable: >0.90; excellent: >0.95), Comparative Fit Index (CFI; acceptable: >0.90; excellent: >0.95) [27], and root mean square error of approximation (RMSEA; acceptable: <0.08; excellent: <0.05) [28].

RESULTS

Characteristics of Study Population

Among the 1,549 OIF/OEF veterans who received the brief TBI screen, 89.6 percent were male (n = 1,388) and the mean age was 30.20 ± 8.39 yr (mean ± standard deviation). The majority of veterans were white (42.3%). Most veterans were Active Duty (65.1%), of Army service (50.5%), and enlisted (92.3%) (Table 1).

Demographic and Military Service Comparisons

Of the 1,549 veterans who were screened, 25 percent (n = 387) were TBI+. A comparison of TBI+ and TBI– veterans showed that TBI+ veterans were more likely to be male (χ² = 18.3, degree of freedom [df] = 1, p < 0.001), in the Army (χ² = 51, df = 4, p < 0.001), and enlisted (χ² = 13.6, df = 1, p < 0.001) compared with TBI– veterans (Table 2).

Prevalence of Traumatic Brain Injury and Mental Health Screens

A TBI+ result was associated with a positive PTSD screen (χ² = 188.4, df = 1, p < 0.001) and depression screen (χ² = 94.6, df = 1, p < 0.001) result. Among those who received both mental health screens within 12 months of their TBI screen (n = 1,359), 58 percent of the TBI+ veterans screened positive for at least PTSD or depression (χ² = 133.9, df = 1, p < 0.001), with 21.3 percent screening positive for both PTSD and depression (χ² = 37.9, df = 1, p < 0.001).

When we examined the adjusted relative risks of positive PTSD and depression screens among TBI+ and TBI– veterans, TBI+ veterans were 2.1 times more likely to screen positive for PTSD, 2.2 times more likely to screen positive for depression, 1.7 times more likely to screen positive for both PTSD and depression, and 2.2 times more likely to screen positive for either PTSD and/or depression compared with TBI– veterans (Table 2).

Relationships Among Traumatic Brain Injury, Posttraumatic Stress Disorder, and Depression Screen-Based Symptoms

A four-factor solution emerged from an EFA considering all 12 symptoms included in the TBI, PTSD, and depression screens and allowing the maximum number of factors permitted by the data (7). The fit indices, lack of negative residual variances, and theory all provided support for a four-factor solution. Table 3 presents a summary of the EFA results, including factor loadings (i.e., probit regression coefficients) for the individual items, eigenvalues, and factor correlations. Based on these results, the first factor consisted of five TBI symptoms (excluding sleep problems), the second factor was comprised of the two depression symptoms, the third factor was comprised of the four PTSD symptoms, and a fourth factor consisted of hypervigilance and sleep problems.

Based on the results of the EFA, we specified a model in which five of the six symptoms on the TBI screen loaded onto the latent variable TBI, the two symptoms on the depression screen loaded onto the latent variable depression and the four symptoms on the PTSD screen loaded onto the latent variable PTSD; for the fourth factor, a correlated error was specified between hypervigilance and sleep problems (i.e., the three other factors did not account for the correlation between these two items). The fit statistics from this measurement model indicated a fairly poor fit (χ² = 165.32, df = 26, p <
Table 1.
Demographic characteristics of veterans screened for traumatic brain injury (TBI).

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>TBI+ (n = 387)</th>
<th>TBI– (n = 1,162)</th>
<th>Total (n = 1,549)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (mean ± SD)</td>
<td>29.60 ± 7.89</td>
<td>30.40 ± 8.55</td>
<td>30.20 ± 8.39</td>
</tr>
<tr>
<td>Sex, * n (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Female</td>
<td>18 (4.7)</td>
<td>143 (12.3)</td>
<td>161 (10.4)</td>
</tr>
<tr>
<td>Male</td>
<td>369 (95.3)</td>
<td>1,019 (87.7)</td>
<td>1,388 (89.6)</td>
</tr>
<tr>
<td>Race, n (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Black</td>
<td>14 (3.6)</td>
<td>40 (3.4)</td>
<td>54 (3.5)</td>
</tr>
<tr>
<td>Hispanic</td>
<td>60 (15.5)</td>
<td>147 (12.7)</td>
<td>207 (13.4)</td>
</tr>
<tr>
<td>White</td>
<td>160 (41.3)</td>
<td>495 (42.6)</td>
<td>655 (42.3)</td>
</tr>
<tr>
<td>Other</td>
<td>46 (11.9)</td>
<td>152 (13.1)</td>
<td>198 (12.8)</td>
</tr>
<tr>
<td>Unknown</td>
<td>107 (27.6)</td>
<td>328 (28.2)</td>
<td>435 (28.1)</td>
</tr>
<tr>
<td>Marital Status, n (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Divorced</td>
<td>11 (2.8)</td>
<td>36 (3.1)</td>
<td>47 (3.0)</td>
</tr>
<tr>
<td>Married</td>
<td>114 (29.5)</td>
<td>318 (27.4)</td>
<td>432 (27.9)</td>
</tr>
<tr>
<td>Never Married</td>
<td>262 (67.7)</td>
<td>808 (69.5)</td>
<td>1,070 (69.1)</td>
</tr>
<tr>
<td>Military Status, n (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Active Duty</td>
<td>267 (69.0)</td>
<td>741 (63.8)</td>
<td>1,008 (65.1)</td>
</tr>
<tr>
<td>Reserves</td>
<td>120 (31.0)</td>
<td>421 (36.2)</td>
<td>541 (34.9)</td>
</tr>
<tr>
<td>Military Branch, * n (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Air Force</td>
<td>14 (3.6)</td>
<td>115 (9.9)</td>
<td>129 (8.3)</td>
</tr>
<tr>
<td>Army</td>
<td>242 (62.5)</td>
<td>540 (46.5)</td>
<td>782 (50.5)</td>
</tr>
<tr>
<td>Coast Guard</td>
<td>0 (0.0)</td>
<td>4 (0.3)</td>
<td>4 (0.3)</td>
</tr>
<tr>
<td>Marine Corps</td>
<td>104 (26.9)</td>
<td>302 (26.0)</td>
<td>406 (26.2)</td>
</tr>
<tr>
<td>Navy</td>
<td>27 (7.0)</td>
<td>201 (17.3)</td>
<td>228 (14.7)</td>
</tr>
<tr>
<td>Rank, * n (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Enlisted</td>
<td>374 (96.6)</td>
<td>1,056 (90.9)</td>
<td>1,430 (92.3)</td>
</tr>
<tr>
<td>Officer</td>
<td>13 (3.4)</td>
<td>106 (9.1)</td>
<td>119 (7.7)</td>
</tr>
</tbody>
</table>

Note: Comparisons are between TBI+ and TBI–.
*p < 0.001.
SD = standard deviation, TBI+ = positive screen for TBI, TBI– = negative screen for TBI.

Table 2.
Cumulative prevalence rates of positive mental health (MH) screen outcomes among veterans screened for traumatic brain injury (TBI).

<table>
<thead>
<tr>
<th>MH Screen</th>
<th>Full Sample</th>
<th>TBI+ n (%)</th>
<th>TBI– n (%)</th>
<th>Total Group</th>
<th>Unadjusted Model</th>
<th>Adjusted Model†</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Risk Ratio</td>
<td>95% CI</td>
</tr>
<tr>
<td>PTSD</td>
<td>1,400†</td>
<td>254 (72.6)</td>
<td>350</td>
<td>324 (30.9)</td>
<td>1,050</td>
<td>2.40†</td>
</tr>
<tr>
<td>Depression</td>
<td>1,407†</td>
<td>154 (44.5)</td>
<td>346</td>
<td>196 (18.5)</td>
<td>1,061</td>
<td>2.40†</td>
</tr>
<tr>
<td>Depression and</td>
<td>1,359§</td>
<td>112 (33.1)</td>
<td>338</td>
<td>177 (17.3)</td>
<td>1,021</td>
<td>1.91‡</td>
</tr>
<tr>
<td>PTSD</td>
<td>1,359§</td>
<td>196 (58.0)</td>
<td>338</td>
<td>245 (24.0)</td>
<td>1,021</td>
<td>2.40‡</td>
</tr>
</tbody>
</table>

†Adjusted for age, sex, marital status, military status, military branch, and rank.
‡Veterans who received particular MH screen within 1 yr of initial TBI screen.
§Veterans who received both MH screens within 1 yr of initial TBI screen.
CI = confidence interval, PTSD = posttraumatic stress disorder, TBI+ = positive screen for TBI, TBI– = negative screen for TBI.
Table 3.
Four-factor solution from exploratory factor analysis * for screen-based symptoms of traumatic brain injury, posttraumatic stress disorder, and depression (n = 358).

<table>
<thead>
<tr>
<th>Symptom</th>
<th>n (%)</th>
<th>Factor Loadings†</th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Factor 1‡</td>
<td>Factor 2§</td>
<td>Factor 3¶</td>
<td>Factor 4**</td>
<td></td>
</tr>
<tr>
<td>Current Dizziness</td>
<td>105 (29)</td>
<td>0.69</td>
<td>−0.13</td>
<td>0.06</td>
<td>0.15</td>
<td></td>
</tr>
<tr>
<td>Current Headaches</td>
<td>211 (59)</td>
<td>0.58</td>
<td>0.02</td>
<td>−0.15</td>
<td>0.28</td>
<td></td>
</tr>
<tr>
<td>Current Irritability</td>
<td>243 (68)</td>
<td>0.48</td>
<td>0.09</td>
<td>0.09</td>
<td>0.22</td>
<td></td>
</tr>
<tr>
<td>Current Memory Problems</td>
<td>183 (51)</td>
<td>0.70</td>
<td>0.04</td>
<td>0.07</td>
<td>−0.08</td>
<td>0.53</td>
</tr>
<tr>
<td>Current Sleep Problems</td>
<td>276 (77)</td>
<td>0.28</td>
<td>0.03</td>
<td>−0.05</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Current Visual Problem</td>
<td>117 (33)</td>
<td>0.78</td>
<td>0.02</td>
<td>−0.01</td>
<td>−0.14</td>
<td></td>
</tr>
<tr>
<td>Nightmares</td>
<td>267 (75)</td>
<td>0.04</td>
<td>−0.02</td>
<td></td>
<td>0.87</td>
<td>0.06</td>
</tr>
<tr>
<td>Avoidance</td>
<td>259 (72)</td>
<td>0.08</td>
<td>0.13</td>
<td></td>
<td></td>
<td>0.92</td>
</tr>
<tr>
<td>Hypervigilance</td>
<td>284 (79)</td>
<td>−0.11</td>
<td>−0.15</td>
<td>0.66</td>
<td>0.61</td>
<td></td>
</tr>
<tr>
<td>Emotional Numbing</td>
<td>264 (74)</td>
<td>−0.09</td>
<td></td>
<td>0.41</td>
<td>0.40</td>
<td>0.31</td>
</tr>
<tr>
<td>Little Interest††</td>
<td>235 (66)</td>
<td>0.04</td>
<td>0.85</td>
<td>0.01</td>
<td>0.10</td>
<td></td>
</tr>
<tr>
<td>Down/Hopelessness</td>
<td>242 (68)</td>
<td>0.02</td>
<td></td>
<td>1.01</td>
<td>0.07</td>
<td>−0.11</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Factor Correlations</th>
<th>Factor 1</th>
<th>Factor 2</th>
<th>Factor 3</th>
<th>Factor 4</th>
</tr>
</thead>
<tbody>
<tr>
<td>Factor 1</td>
<td>1.00</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Factor 2</td>
<td>0.37</td>
<td>1.00</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Factor 3</td>
<td>0.10</td>
<td>0.53</td>
<td>1.00</td>
<td></td>
</tr>
<tr>
<td>Factor 4</td>
<td>0.35</td>
<td>0.43</td>
<td>0.32</td>
<td>1.00</td>
</tr>
</tbody>
</table>

**Fit Statistics**

- Chi-Square Test of Model Fit: $p = 0.48$
- RMSEA: 0.00
- Weighted Root Mean Square Residual: 0.027

*Weighted least-squares with mean and variance adjustment estimation; oblique (promax) rotation.
†Factor loadings are probit regression coefficients. Interpretation of positive probit regression coefficient is that increase in value of latent variable leads to increase in predicted probability in having positive response in case of binary variable or of having increased expected value of ordinal variable. Bold numbers represent symptoms that had factor loadings >0.40.
‡Eigenvalue = 4.89.
§Eigenvalue = 2.35.
¶Eigenvalue = 1.20.
**Eigenvalue = 0.86.
††Number and percentage shown are for those who scored greater than zero on that item, but in confirmatory factor analysis, these variables were modeled as ordinal with possible values of 0–4.
RMSEA = root mean square error of approximation.

Based on a review of the parameter derivatives (modification indices), we made several modifications: (1) sleep was added to the depression factor, (2) sleep was added to the TBI factor, (3) emotional numbing was added to the depression factor, and (4) irritability was cross-loaded onto the PTSD factor. This final model resulted in a favorable chi-square goodness of fit test ($\chi^2 = 28.95$, $df = 27$, $p = 0.36$); the other fit indices also indicated a very good fit ($\chi^2$ difference test = 94, $df = 4$, $p < 0.001$; CFI = 0.999; TLI = 0.999; RMSEA = 0.014). The Figure shows the final CFA with standardized estimates (probit regression coefficients) and their standard errors. All freely estimated parameters estimates were statistically significant ($p < 0.05$).

The correlation between the TBI factor and the PTSD factor was weak (0.2), indicating that the factor structure distinguished the TBI latent variable from the PTSD latent variable fairly well (Figure). This suggested that there were particular symptoms that were unique to the latent TBI variable (dizziness/balance problems, headache, memory problems, and light sensitivity), while other symptoms (e.g., irritability) overlapped with the PTSD factor. In addition, sleep problems overlapped with the depression factor. Certain symptoms (nightmares, avoidance) were unique to the PTSD factor, but one
Figure.
Confirmatory factor analysis for symptoms of traumatic brain injury (TBI), posttraumatic stress disorder (PTSD), and depression ($n = 358$). Note: Estimates shown are standardized estimates and corresponding robust standard errors. $\chi^2 = 30.076$, degree of freedom $= 27$, $p = 0.31$, Tucker Lewis Index $= 0.999$, Comparative Fit Index $= 0.999$, root mean square error of approximation $= 0.018$. * $p < 0.001$, † $p = 0.01$, ‡ $p < 0.05$.

Symptom, emotional numbing, overlapped with the depression factor. We specified the correlated error term between sleep problems and hypervigilance because results from EFA indicated that these symptoms comprised a fourth factor.

$R^2$ estimates (squared multiple correlations) provided information on how much variance the factors accounted for in the observed variables. Nightmares and avoidance were well explained by the model ($R^2 = 0.74$ and $0.99$, respectively), as were the symptoms of little interest and...
feeling depressed ($R^2 = 0.87$ and $0.94$, respectively). However, the model did not account well for the six TBI-related symptoms (all $R^2 < 0.50$), particularly sleep problems ($R^2 = 0.31$). This suggested that the variability in the distribution of the TBI-related symptoms was not readily explained by co-occurring symptoms of PTSD or depression. In particular, these results supported the idea that sleep problems, the most prevalent TBI-related symptom, were not strongly associated with any of the factors but were still commonly experienced.

**DISCUSSION**

Military personnel are returning home with multiple postdeployment conditions and identifying distinguishing features of each condition is critical in order to appropriately triage, diagnose, and provide targeted early interventions. We found that while postdeployment screens accurately identified symptoms associated with TBI, PTSD, and depression, hypervigilance and sleep problems likely constituted a fourth factor that may have represented common postdeployment symptoms due to the high frequency with which these symptoms occurred (79% and 77%, respectively). Furthermore, we found that while some screen-based symptoms were shared (e.g., irritability), others were unique (e.g., nightmares).

While the TBI screen assessed six symptoms after a TBI injury mechanism had been established, we found that only dizziness/balance problems, light sensitivity, headaches, and memory problems uniquely identified a veteran with mild TBI-related symptoms. In contrast, endorsement of the two other symptoms (sleep problems and irritability) may have led to an inflated false positive rate on the TBI screen because they overlapped with mental health symptoms of PTSD and depression. Similarly, while nightmares and avoidance best captured unique PTSD screen symptoms, hypervigilance and sleep problems constituted common co-occurring postdeployment symptoms. Emotional numbing is an overlapping screen-based symptom shared by PTSD and depression.

These findings suggest that if clinicians want to initially assess for TBI and minimize false positives, focusing on distinguishing rather than shared TBI screen symptoms will be most informative. As always, referral for second-level TBI evaluation by a neurologist or rehabilitation medicine specialist will help determine a veteran’s ultimate diagnosis.

Although these distinguishing features suggest possible ways for these screens to be more specific when administered together (i.e., by focusing on distinguishing features of each screen), it will also be important to ensure that doing so will not significantly reduce sensitivity. This is an important area for future investigation.

These findings also have implications for treatment planning. Often, mental health, neurology, and rehabilitation teams must reach consensus concerning the best course of treatment for a particular veteran, and little empirical evidence exists for how to best treat veterans with multiple co-occurring postdeployment symptoms. One possibility is to shift from a diagnosis-based to more of a symptom-based approach, with a particular emphasis on those symptoms associated with functional impairment and problems with reintegration. Our findings suggest that among the most prevalent postdeployment conditions, there are common screen symptoms that may represent shared pathways that can be treated collaboratively across disciplines. For example, there are excellent treatments for irritability and anger and a growing understanding of how to best tailor treatments to target particular types of anger [29]. Given that irritability and anger are shared symptoms of TBI and PTSD, focusing on a symptom-based approach to treatment may help ameliorate both conditions. In addition, there should be more research on the efficacy of treatments that focus on shared symptomatology. For example, treatments that focus on shared symptoms of insomnia, emotional regulation, and cognitive dysfunction may be the best first-line treatments. This shift is consistent with other findings that have stressed the need for integrated care and have highlighted the clinical challenges of addressing PTSD and TBI as separate conditions [17].

Some may argue that symptoms within a common cluster should be treated together as a whole. Indeed, in cases where there is a strong evidence-base for doing so, this is another option: to consider similar symptoms together, such as the distinguishing symptoms of PTSD. These can be targeted through evidence-based treatments (EBTs) such as prolonged exposure and cognitive processing therapy. The same is true for the distinguishing features of depression; EBTs for depression have received excellent support and should be used when these are the presenting symptoms. Once these treatments are administered, symptoms should be re-evaluated and any residual symptoms treated accordingly. Whether a symptom-based or diagnosis-based mental health specialty
approach is taken also largely depends on contextual factors such as patient preferences, provider training, and systemic resources.

Several limitations should be noted. We used population-based screen data and, as a result, these data should not be generalized to veterans with confirmed diagnoses of TBI, PTSD, and depression. Future investigations should be conducted to better understand whether these findings based on screen-derived symptoms can be replicated in those who receive TBI, PTSD, and depression diagnoses. It is also important to note that these screens are based on self-report rather than direct clinician observation and/or interview or standardized measurement of posttraumatic amnesia and loss of consciousness. Self-report may also result in the over- or under-estimation of specific symptoms. For example, it may be that because these results are based on self-reported screens, we may have false negatives caused by underreporting or minimizing of symptoms that are consequently not included in these data. Additionally, endorsed symptoms may vary from person to person (e.g., many different types of headaches, with some more severe and disabling) and do not yield any information about severity or intensity (e.g., insomnia once a week vs daily). Screens also may lack sensitivity to detect the level of depression that may include other associated symptoms, such as sleep and memory problems. Because of the way in which the VA TBI screen is structured, those who are TBI– do not complete the entire TBI screen. Consequently, when conducting factor analysis, we could only include veterans who had completed the TBI screen in its entirety (those who screened out did not have the opportunity to respond to the question about current TBI symptoms). Although all screens were done within 1 year of each other, not all were necessarily done at the same time, and in some cases, these screens were conducted several years after the veteran returned home. Despite these limitations, VA screen results represent the level of information that clinicians use to make initial decisions about assessment, triage, and early symptom management of their patients.

CONCLUSIONS

Overall, we found that TBI+ veterans were approximately twice as likely to also screen positive for depression and PTSD. Four distinct constructs emerged: TBI, PTSD, depression, and a fourth construct consisting of hypervigilance and sleep problems. Screen symptoms unique to TBI include dizziness, headaches, memory problems, and light sensitivity. We found an overlapping screen-based symptom between TBI and PTSD (irritability) and between PTSD and depression (emotional numbing). A symptom-focused, integrated care approach will help to promote more expedient and efficient care for our returning veterans.

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REFERENCES


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