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The Effect of PTSD Symptom Change on Suicidal Ideation in a Combined Military and Civilian Sample Engaged in Cognitive Processing Therapy

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In light of the well-established relationship between posttraumatic stress disorder (PTSD) and suicidal ideation (SI), there has been a push for treatments that simultaneously improve symptoms of PTSD and decrease SI. Using randomized controlled from a implementation-effectiveness trial, the current study investigated the effectiveness of Cognitive Processing Therapy (CPT; Resick, Monson, & Chard, 2016) on PTSD and SI. The patient sample (N = 188) was diverse in military and veteran status, gender, and comorbidity, and 73% of the sample endorsed SI at one or more points during CPT. Participants demonstrated significant improvement in SI over the course of CPT. Multilevel growth curve modeling revealed a significant association between PTSD symptom change and change in SI. Results from crosslagged multilevel regressions indicated that PTSD symptoms predicted SI in the next session, yet SI in a given session did not predict PTSD symptoms in the next session. Potentially relevant clinical factors (i.e., military status, gender, depression diagnosis, baseline SI, study consultation condition) were not associated with the relationship between PTSD symptoms and SI. These results add to the burgeoning literature suggesting that evidence-based treatments for PTSD, like CPT, reduce suicidality in a range of individuals with PTSD, and that this reduction is predicted by improvements in PTSD symptoms.

Keywords: PTSD; suicidality; cognitive processing therapy; treatment process and outcome measures

WITHIN THE UNITED STATES, approximately 123 individuals die by suicide each day (Drapeau & McIntosh, 2017). As such, mental health care systems prioritize suicide prevention. Although the United States has made strides in increasing awareness for suicide prevention, and a body of literadeveloped to identify has treatments to reduce suicide risk and suicidal ideation (SI), emphasis on posttraumatic stress disorder (PTSD), a key risk factor, is often neglected. Thus, the current study aims to expand on previous literature by analyzing the overall and temporal relationship between PTSD symptomatology and SI throughout an evidence-based psychotherapy (EBP) for PTSD.

Extensive research indicates that PTSD is a risk factor for SI and suicide attempts among veteran (Arenson et al., 2018; Guerra & Calhoun, 2011; Pompili et al., 2013), active duty (Bryan & Corso, 2011; McLean et al., 2017) and civilian samples (Boffa et al., 2017; Gradus et al., 2010; Kratovic et al., 2020; Wilcox et al., 2009). A

possible explanation for the elevated suicide risk is the high prevalence of comorbid depression individuals diagnosed with PTSD among (Arenson et al., 2018; Kimbrel et al., 2016; O'Donnell et al., 2004). Comorbid depression and PTSD can result in an increased risk for SI versus PTSD alone (Brown, Zang, et al., 2019; Bryan & Corso, 2011; Calabrese et al., 2011; Ramsawh et al., 2014). However, Guerra and Calhoun (2011) and Wilcox et al. (2009) found that PTSD symptoms and SI remain significantly correlated even when controlling for symptoms of major depressive disorder. Although the specific mechanisms that explain the relationship between PTSD and SI remain unclear, the well-established link between PTSD and SI presents an opportunity to identify strategies to address highly comorbid symptoms within a single treatment (Krysinska & Lester, 2010).

SI has been shown to decrease after engagement in EBPs for PTSD, including individual and group Cognitive Processing Therapy (CPT; Resick et al., 2016; Gradus et al., 2013; Stayton et al., 2019; Horwitz et al., 2019; Resick et al., 2017) and Prolonged Exposure (PE; Foa et al., 2007; Brown, Belli, et al., 2019; Brown, McLean, et al., 2019; Cox et al., 2016; Norr et al., 2018). Moreover, previous studies demonstrate that declines in PTSD symptoms are correlated with reductions in SI during CPT (Gradus et al., 2013) and PE (Brown, Belli, et al., 2019; Brown, McLean, et al., 2019; Cox et al., 2016). Gradus et al. (2013) studied female rape survivors' endorsement of PTSD symptoms and SI before, after, and at various time points during CPT and PE in a randomized controlled trial (RCT) that compared the two treatments. Although reductions in PTSD symptoms were correlated with reductions of SI in both PE and CPT, patients enrolled in CPT saw greater reductions in SI. However, PTSD and SI were only measured every other session in the study, limiting the ability to understand potential session-tosession changes and the temporal precedence of changes in each. In another study, of predominantly male veterans who received PE in three U.S. Veterans' Health Administration centers, Cox et al. (2016) found that PTSD symptoms in one session predicted SI in the next session of PE, as measured by a single item on the Depression Inventory (Beck et al., 1988), in routine care. They did not, however, find the reverse relationship to be true. That is to say, SI in a particular session did not predict PTSD symptoms in the next session. Norr et al. (2018) similarly looked at the

relationships between PTSD symptoms and SI in an RCT among active-duty military members enrolled in PE, Virtual Reality Exposure, and waitlist control using cross-lagged panel models. In contrast to prior findings, SI measured at midtreatment predicted the PTSD symptom change in the subsequent session. These findings suggest that EBPs for PTSD result in a decrease of SI and that there may be a reciprocal relationship among SI and PTSD symptoms.

Although the extant literature has established an association and potential temporal relationships between reductions in PTSD symptoms and SI in the course of EBPs for PTSD, this paper aims to address some of the existing gaps in the literature. First, much of the previous research has been in efficacy trials. To examine the feasibility and scalability of using EBPs in routine care, it is important to understand whether the same effects can be expected among therapists working in community settings and with patients who are more heterogeneous. Furthermore, clinical trials often examine PTSD treatment in specific populations in terms of gender, trauma type, or military status, decreasing the generalizability to the other trauma types and populations despite empirical evidence suggesting that these factors can be impactful. For example, Gradus et al. (2013) focused on female rape survivors, Brown, Belli, et al. (2019) investigated adolescent females with a history of sexual assault, Cox et al. (2016) studied mostly middle-aged male veterans, and Norr et al. (2018) analyzed data from largely male activeduty military members. Zimmerman et al. (2015) found gender to moderate the relationship between PTSD and SI such that women were more likely to experience SI as PTSD symptoms increased, emphasizing the need to study mixedgender samples. Because research has indicated that certain types of traumatic events have stronger associations with suicidal thoughts and behaviors than others (Stein et al., 2010), it is also critical to explore whether these findings regarding changes in PTSD symptoms and SI during treatment remain constant within a broad range of individuals, including active military personnel, veterans, and civilians.

To our knowledge, no previous research has investigated if PTSD symptoms predict SI (along with the converse relationship) within a community-based sample of individuals engaged in CPT for PTSD. Gradus et al. (2013) examined the relative efficacy of CPT versus PE in improving suicidality, but did not examine the directionality of changes in PTSD and suicidality. Cox et al. (2016) and Norr et al. (2018) explored the directions.

tionality of associations between PTSD and SI among patients in PE. As CPT is another first-line treatment for individuals with PTSD and may reduce SI more than PE (Gradus et al., 2013), it is critical to determine whether results from PE trials (e.g., Cox et al., 2016; Norr et al., 2018) translate to CPT. Another limitation to existing literature is infrequent assessment. For example, Norr et al. (2018) measured PTSD symptoms and SI at baseline, Session 5, posttreatment, and 3- and 6-month follow-up. To better examine the relationship between PTSD and SI at a session-to-session level, it is important to consistently collect and analyze symptom measurements at each session.

The current study used data from an implementation trial of CPT (Monson et al., 2018) to explore the relationship between PTSD symptom change and SI change over the course of CPT within a mixed active-duty, veteran, and civilian sample with varied trauma types that were served in diverse routine care settings across Canada. We first sought to replicate previous findings by analyzing change in PTSD symptoms and SI over the course of CPT. Consistent with previous literature (Gradus et al., 2013; Horwitz et al., 2019; Resick et al., 2017; Stayton et al., 2019), we hypothesized that patients would experience reductions in both PTSD and SI over the course of CPT (Hypothesis 1). We also assessed the relationship between changes in PTSD symptoms and SI during treatment. We hypothesized that changes in SI would be positively correlated with changes in PTSD during treatment (Hypothesis 2) based on existing research (Brown, Belli, et al., 2019; Brown, McLean, et al., 2019; Cox et al., 2016; Gradus et al., 2013). To better understand the temporal relationship, we examined whether PTSD symptoms predicted SI and vice versa at a session-tosession level. We hypothesized that changes in PTSD symptoms would predict changes in SI in future sessions, but, based on previous findings (Cox et al., 2016), that the reverse relationship would not be significant (Hypothesis 3). As an exploratory aim, we examined whether clinical depression diagnosis, baseline SI, military/veteran status, gender, and type of consultation received in the implementation trial moderated the relationships described in our third hypothesis (Exploratory Aim 1).

Method

For detailed procedural and demographic information, please refer to the parent study (Monson et al., 2018). To briefly summarize the procedure, the study enrolled therapists (N = 134) with

varying levels of education and experience with EBPs from Veterans Affairs Canada Operational Stress Injury Clinics, Canadian Forces mental health services, and the broader Canadian community. They were randomly assigned to one of three consultation conditions following a 2-day CPT training workshop: (a) standard CPT consultation by expert CPT consultants without session audio review (Standard Consultation); (b) CPT consultation by expert CPT consultants that included review of session audio segments (Consultation Including Audio Review); and (c) no consultation, with delayed feedback on fidelity (i.e., adherence to, and competence in, CPT). All procedures were approved by university Research Ethics Boards (REB) and Institutional Review Boards along with REBs from each site involved in the study.

Therapists identified patients with a Diagnostic and Statistical Manual of Mental Disorders (4th edition) diagnosis of PTSD (DSM-IV; American Psychiatric Association, 2000) with whom they intended to conduct CPT. The parent study began the DSM-5 (American Psychiatric Association, 2013) was published, so the current study used measures of the DSM-IV. As the therapists were recruited in routine care, the study team did not require a certain diagnostic approach, but rather asked therapists to use their organization's routine intake procedures to diagnose disorders including PTSD and depression. Intake may have varied across settings involved in the study, but some health systems use less structured intake procedures than is expected in a randomized clinical trial. Two participants without a clinical diagnosis of PTSD were included in the study because they experienced a trauma, scored higher than the cutoff (50) on the PTSD Checklist (Weathers et al., 1993), and their therapists believed CPT to be potentially helpful. Patients (N = 188) were eligible for participation if they consented to audio recording of therapy sessions and were eligible for CPT based on the standards set by CPT developers (i.e., did not have current untreated symptoms of bipolar disorder or psychosis, substance dependence requiring detoxification, risk of imminent suicide or homicide or significant cognitive impairments; Resick et al., 2016). Therapists were instructed to use their organization's risk assessment protocols and their clinical judgment to determine whether there was a substantial risk of imminent suicide or homicide in determining patient eligibility.

Fifty-two percent of the sample identified as female, with a mean age of 39.4 years (*SD* = 11.3). The sample included 57.6% nonmilitary/nonveteran patients, 23.3% veterans, and 19.2%

patients on active duty. Patients were predominantly White, (88.0%), with the others identifying as Black (2%), Hispanic (1%), Asian/Pacific Islander (3%), Native Canadian (4%), and "other" (4%). Most participants were in committed relationships (59.0%), while the rest were single, widowed, or divorced. Therapists reported that 50.5% of the patients experienced comorbid depressive disorders. For further demographic information, see the parent study (Monson et al., 2018).

MEASURES

The Posttraumatic Stress Disorder Checklist (PCL-IV) is a 17-item, self-report survey that measures DSM-IV PTSD symptom severity on a 5-point Likert scale from "not at all" [1] to "extremely likely" [5] (Weathers et al., 1993). The cut-off score of 50 indicates probable PTSD. The PCL-IV has strong internal consistency in the current study (α = .94) and was completed at baseline, before each therapy session, and posttreatment. Therapists were trained by study staff to start each therapy session with symptom measures so that the therapist and patient could use the scores clinically during the session, particularly for the PCL-IV.

The Outcomes Questionnaire-45 (OQ-45) monitors functioning in routine care (Lambert et al., 1996). Only 1 of the 45 items, which assesses whether the respondent has thoughts of ending one's life, was used to measure SI in the current study on a 5-point Likert scale from "never" [0] to "almost always" [4]. Patients completed the OQ-45 at baseline, before each session, and posttreatment.

Therapists reported each patient's clinical diagnosis of depression to the study team prior to the start of treatment.

ANALYTIC PLAN

Hypothesis 1: Change in PTSD and SI Over Time Using Mplus (Muthén & Muthen, 2017), we conducted multilevel growth curve models to examine overall change in PTSD symptoms and SI during treatment. Prior to examining change, we evaluated two unconditional models (i.e., models with no predictors) to decompose the variance between three possible levels (Level 1: within patient participants/SI and PTSD symptoms across 12 therapy sessions; Level 2: between-patient participants; Level 3: between-therapist participants). Next, we evaluated separate unconditional change models (i.e., time included as the only predictor) to identify the best way to model time for PTSD symptoms and SI (linear versus log linear function of time). Table 2 includes the model comparisons based on change in deviance score

(-2 log-likelihood value). The best fitting model for both PTSD symptoms and SI included the linear model of time, with time as the number of days since baseline assessment.

Hypothesis 2: Association Between PTSD Change and SI Change

We conducted a 2-level multivariate growth model producing initial status and slope estimates for both PTSD and SI in the same model.

Hypothesis 3: Temporal Dynamics of the Association Between PTSD and SI

We conducted cross-lagged multilevel regression analyses using the lmer and lmerTest statistical packages in R Statistical Software (R Core Team, 2018) to evaluate the directionality of the relationship of PTSD and SI. We retained the nesting structure from the analysis used to test Hypotheses 1 and 2. Our first model included SI, assessed at a single session, as the dependent variable and controlled for auto-correlated SI from the previous session. We compared model fit indices to examine which model of time to include in the current model (linear or linear plus quadratic). Our cross-lagged predictor of interest was PTSD symptoms from the previous session. We tested the reverse relationship in our second model. We included cross-lagged SI score as a predictor of PTSD symptoms from a particular session controlled by auto-correlated PTSD symptoms. Consistent with the parent study, we included the linear and quadratic model in the PTSD symptom predictor model.

Exploratory Aim 1: Potential Moderators

We then evaluated the cross-lagged models with each construct (baseline SI, clinical depression diagnosis, military/veteran status, gender, and type of consultation) added and included a variable-bytime interaction term. Missing demographic data were imputed using missForest r package (Stekhoven, 2013).²

Results

HYPOTHESIS I

Table 1 includes descriptive statistics of PTSD symptoms, SI, and days passed since baseline assessment for each session of CPT. As the parent study excluded patients who were at imminent risk

for suicide, the levels of SI were relatively low (M = 0.96, SD = 1.16). However, only 27% of patients consistently reported never having thoughts of ending their lives. Thirty-six percent of the patients consistently reported at least some level of SI, with the rest reporting SI at some weeks and no SI at others.

Unconditional Models

Thirty-seven percent (p < .001) of the total variance in PTSD symptoms occurred at Level 1 (within participants), 62.27% (p < .001) was attributed to Level 2 (between-patient participants), and 0.69% (p = .888) of the variance was attributed to Level 3 (between therapists). Twenty-seven percent (p < .001) of the total SI variance was attributed to Level 1, 63.41% (p < .001) was attributed to Level 2, and 9.77% (p = .127) of the variance was attributed to Level 3. Since Level 3 variance was not statistically significant for either variable we evaluated two-level models for subsequent analyses.

Multilevel Growth Curve Models

SI decreased over time during CPT, b = -0.01, 95% CI [-0.02, -0.001], p = .026, pr = -.12. Consistent with findings from the parent study, PTSD symptoms also exhibited significant decreases during CPT, b = -0.90, 95% CI [-1.06, -0.74], p < .001, pr = -.46. Of those who completed CPT, 52.4% patients experienced reliable clinically significant improvement of PTSD symptoms (Holmes et al., 2019).

HYPOTHESIS 2

PTSD Symptoms and SI

Consistent with Hypothesis 2, the slopes of PTSD symptoms and SI change were highly correlated, r (188) = .64, p < .001, 95% CI [.42, .81] (see Figure 1), indicating that individuals showing larger decreases in PTSD symptoms tended to exhibit larger decreases in SI relative to participants showing smaller decreases in PTSD.

HYPOTHESIS 3

PTSD Symptoms Predicting SI

We included only a linear time coefficient for the lagged analyses with SI as the outcome, as adding a quadratic time coefficient did not significantly improve model fit ($\Delta DEV = 1.10$, $\Delta Parms = 1$, p = .338, pr = .16). When controlling for the prior session SI and the SI symptom trajectory, the PTSD symptoms predicted SI at the subsequent session, b = 0.01, t(730) = 2.65, p = .008, pr = .10 (see Table 3). The effect size of this cross-lagged PTSD symptom was small to medium. Cross-lagged models with each possible potential moderator

² Given that the parent study found dierences in PTSD symptom change across consultation conditions, we included type of consultation as a moderator in analyses for Hypotheses 2 and 3. There was no main eect of consultation type on the relationship between PTSD and SI, so we did not separate the analysis by consultation condition.

Table 1
Descriptive Statistics for Number of Days Since Baseline Assessment, PTSD Symptoms, and SI

Session	Number of Days Since Baseline		PTSD Symptoms			SI		
	M	SD	n	М	SD	n	М	SD
0	0.00	0.00	149	61.12	11.26	143	1.14	1.21
1	13.68	15.94	167	59.37	12.02	158	1.08	1.18
2	22.61	17.75	168	57.56	13.15	161	0.98	1.14
3	35.20	37.30	155	57.31	12.49	146	1.00	1.20
4	45.78	42.72	141	56.87	13.83	134	1.05	1.18
5	54.70	42.87	127	55.47	14.66	123	0.86	1.08
6	64.56	45.48	126	52.92	15.69	118	0.85	1.13
7	77.13	49.71	121	50.90	14.57	115	0.94	1.14
8	86.36	51.05	120	49.93	15.99	120	0.82	1.17
9	95.97	53.16	115	48.62	16.32	113	0.86	1.13
10	109.5	59.58	113	47.32	15.33	112	0.97	1.19
11	118.3	61.19	107	46.77	15.69	101	0.85	1.15
12	122.3	48.32	108	43.64	16.18	105	0.80	1.12

Note. The data comes from the sample that has completed the PTSD symptom and SI measures from a given session. n = sample; M = mean; SD = standard deviation; SI = suicidal ideation; PTSD = posttraumatic stress disorder; Session 0 = pre-session baseline.

Table 2
Growth Curve Estimates for Each Model of Time and Baseline to Session 12 Changes for PTSD Symptoms and SI

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Parameter	Condition	b	95% CI	р
Intercept/Baseline				
	PTSD	61.12	[51.27, 70.97]	<.001
	SI	1.14	[0.96, 1.32]	<.001
Linear Time (Days/30)				
	PTSD	-0.90	[-1.06, -0.74]	<.001
	SI	-0.01	[021, -0.001]	.026
Log function of time [Lr	n(time+1)]			
	PTSD	-1.81	[-3.31, -2.33]	<.001
	SI	-0.05	[-0.08, -0.02]	.003
Baseline - S12 Change	!	<u>Est</u>	<u>d</u>	
	PTSD	-17.35	-1.56	
	SI	-0.33	-0.27	

Note. Initial estimates from PTSD and SI symptom growth curve models did not fully display in the Mplus output due to program limitations. We divided the temporal variable in all models by a constant (30) to change timing from daily to monthly. Dividing both variables by a constant does not change the results other than altering the coefficient metric. B = unstandardized regression coefficient; 95% CI = 95 percent confidence interval; p = p-value; Est = estimate; d = Cohen's d effect size (0.20 small, 0.50 medium, 0.80; (Cohen, 1977); S12 = session 12; Days = days since baseline therapy session; SI = suicidal ideation; PTSD = posttraumatic stress disorder.

(baseline SI, clinical depression diagnosis, military/ veteran status, gender, and type of consultation) by time interaction term, included separately, revealed no significant interactions.

SI Predicting PTSD Symptoms

We included linear and quadratic time coefficients in the cross-lagged models with PTSD symptoms as the outcome ($\Delta DEV = 44.15$, $\Delta Parms = 1$, p < .001). When controlling for the autocorrelated PTSD symptom score and the symptom trajectory of PTSD symptoms, the cross-lagged path from SI to the PTSD score at the next session was not statistically significant, b = 0.13, t(1389) = 0.66, p = .510, pr = .02 (see Table 3). The effect size of this cross-lagged path was small. None of

the interaction terms (time by baseline SI, clinical depression diagnosis, military/veteran status, gender, and type of consultation) were significant when included individually in the cross-lagged SI model.

Discussion

The current study expands on the existing literature that examines the relationship between PTSD symptom change and SI in EBPs for PTSD in two important ways. First, research that has explored the relationship between PTSD and SI, to date, has typically been in the context of efficacy trials and has utilized homogeneous samples (e.g., female rape survivors, male veterans). Conse-

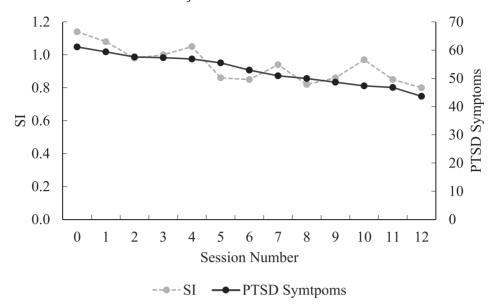


FIGURE 1 Comparison of the 2-level growth curves (PTSD symptom trajectory vs. SI trajectory). We depicted this graph using an index chart to show relative change since baseline of PTSD symptoms and SI. The slopes are highly correlated r(188) = .64, p < .001. SI = suicidal ideation; PTSD = posttraumatic stress disorder

Table 3 Fixed-Effect Estimates for the Multilevel Cross-Lagged Analyses of PTSD Symptoms and SI

Parameter	b	SE	p	pr
SI				
Intercept	0.24	0.12	.057	.09
Auto-Correlated SI	0.45	0.03	<.001	.55
Cross-Lagged PTSD Symptoms	0.01	0.00	.008	.10
Time (Linear)	0.00	0.00	.208	03
PTSD Symptoms				
Intercept	5.74	0.98	<.001	.15
Auto-Correlated PTSD Symptoms	0.88	0.02	<.001	.84
Cross-Lagged SI	0.13	0.19	.510	.01
Time (Linear)	-0.01	0.01	.220	03
Time (Quadratic)	0.00	0.00	.443	.02

Note. b = unstandardized coefficient; SE = standard error; p = p-value; pr = partial regression coefficient (small = .10, medium = .24, large = .37); PTSD = posttraumatic stress disorder; SI = suicidal ideation.

quently, researchers have called for additional inquiry to assess the generalizability of this effect (Gradus et al., 2013). The current study addressed this question by exploring the relationship between change in PTSD symptoms and SI over the course of CPT in routine care settings and by testing the effects of various potential moderators of the relationship in a heterogeneous sample. The results regarding the association between PTSD symptom change and changes in SI are consistent with our hypothesis, and with the results of previous studies (e.g., Gradus et al., 2013). These results, taken with the lack of significant moderators, support the robustness and generalizability of the relationship between PTSD and SI within the context of CPT.

The current study adds to the existing literature by examining the temporal dynamics of PTSD and SI over the course of CPT delivered in routine care settings. Previous studies that have employed cross-lagged models to examine the temporal relationship between PTSD symptom and SI changes involved PE efficacy trial data. The current study corroborates the unidirectional relationship found in PE (Cox et al., 2016) in CPT. Thus, PTSD symptom improvements appear to drive the change in SI, rather than the reverse.

Although specific mechanisms were not tested in the current study, there are several potential explanations for the association between PTSD symptom and SI reductions. Given that previous research has found greater reductions in both guilt and SI (Gradus et al., 2013; Nishith et al., 2005) in CPT than PE and that shame affects the relationship between PTSD and SI (e.g., Bryan et al., 2013; Cunningham et al., 2019), it would be important to investigate the temporal relationship between shame, guilt, and SI. Less frequent and intense reexperiencing and hyperarousal symptoms over the course of treatment may also result in less distress. Additionally, as avoidance resolves, and participants begin to re-engage in activities and reconnect with people, they may feel an increased sense of meaning and life satisfaction—all of which may be relevant to SI. Future research should assess these potential mechanisms.

The current study indicates that CPT can address factors associated with SI, and that decreases in PTSD symptoms result in SI reduction. This is an important message to highlight, given persistent apprehension regarding the administration of trauma-focused therapies with individuals who experience SI. Some have expressed concerns that processing traumatic events will result in PTSD symptom exacerbation (Becker et al., 2004; Larsen et al., 2016). It should be noted that imminent risk of suicide is a contraindication of CPT, though researchers are currently testing CPT with patients with imminent suicidality and with patients who endorse nonsuicidal self-injury (Monson et al., 2018). Withholding CPT due to moderate levels of SI until SI has been reduced may mean delaying an intervention that could actually address this very important clinical concern. Consistent with previous research (e.g., Gradus et al., 2013), the results of the current study suggest that PTSD and SI can be effectively treated simultaneously with CPT and other evidence-based, trauma-focused therapies for PTSD. Although SI significantly decreased over time, patients experienced a minor increase in SI at Session 10. The increase in SI may indicate that as patients near the end of therapy, they begin to consider the idea of navigating life's challenges after leaving therapy and thus might experience a slight increase in SI. Further research should explore predictors of a spike in SI to help therapists prepare for SI increases later in therapy.

The findings of the current study are particularly salient given the prioritization of suicide prevention by health care systems. The Department of Veterans Affairs published the "National Strategy for Preventing Veteran Suicide 2018–2028," which includes guidelines for therapists to follow when working with veterans with SI. While the creation and proliferation of policy and resources is necessary, the discussion of the role of PTSD treatments in addressing SI is missing, which has

serious clinical implications. PTSD is a wellestablished risk factor for suicide attempts (e.g., Calabrese et al., 2011) and treatment for PTSD can result in decreases in SI regardless of gender, military/veteran status, depression diagnosis, and extent of suicidality. Consequently, discussion of PTSD should be explicitly integrated into psychoeducational resources targeting suicide risk, PTSD screening should be considered in conjunction with suicide risk assessments, and CPT and PE might be highlighted as potentially effective suicide prevention strategies for individuals with comorbid PTSD who are not at imminent risk of suicide. Further research should examine whether PTSD treatment is as effective as other strategies for suicide prevention in this population.

The current study exhibits a number of strengths, including its heterogeneous sample, longitudinal design, and collection of data at each session. Additionally, CPT occurred in routine care settings by newly trained CPT therapists, which allowed for greater generalization than previous randomized controlled trials with more restricted sample characteristics. There are, however, some limitations. First, SI was assessed using a single item. Although this approach has been used by similar studies (e.g., Gradus et al., 2013; Horwitz et al., 2018; Norr et al., 2018), it does not allow for a comprehensive assessment of suicidality (e.g., plan, intent, means). It is also possible that the variation in the single item measure of SI overtime may have influenced the overall effect of CPT on SI that was found. Nonetheless, psychotherapy assessments with one-item and treatment measures with fewer items reduce survey burden (Hornsey et al., 2012). Additionally, the current sample endorsed relatively low levels of SI. Replication in samples with higher levels of SI will be important in terms of generalizability and power. SI was also collected at the beginning of each session, which may indicate that the SI may have been tied to anxiety towards starting a new session. If SI were collected upon completion of a therapy session, this relationship between SI and PTSD may be different. Nonetheless, as this study was within a routine care sample, the procedures aligned with the way in which therapists commonly collect symptom measures in order to assist clinically in the session. As the current study did not systematically collect index traumas from each patient, it would be important to look at index trauma as a moderator of the relationship between PTSD and SI in future research. Finally, while the sample was diverse in several ways (i.e., gender, military/ veteran status), it lacked racial, ethnic, and sexual diversity. Consequently, future research should

assess EBPs for PTSD as a means of addressing SI with more comprehensive measures of suicidality among a more diverse sample. Additionally, future research should explore mechanisms that may explain the association between PTSD symptom and SI reductions.

Conclusions

The current study demonstrates that (a) during a course of CPT, PTSD symptoms and SI decrease, (b) reductions in PTSD symptoms were associated with reductions in SI, (c) PTSD symptom improvement in a particular session predicts SI in the next session, and (d) gender, military/veteran status, consultation condition, and depression diagnosis do not affect the relationship between PTSD and SI. These results provide invaluable information for the ongoing large-scale efforts in health care systems to prevent suicide. Specifically, suicide awareness and prevention resources should make explicit the link between PTSD and SI, PTSD screens should be incorporated into suicide risk assessments, and CPT should be considered as a potential strategy to address suicide risk for patients with PTSD.

Conflict of Interest Statement

Dr. Monson receives book royalties from Guilford Press related to authoring the treatment manual of interest to this publication. There are no other interests to disclose.

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