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Temporal Relationship Between Change in Subjective Distress and PTSD Symptom Decrease During Prolonged Exposure Therapy for Posttraumatic Stress Disorder

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There is growing evidence that change in distress is an indicator of change during Prolonged Exposure (PE) for post-traumatic stress disorder (PTSD). However, temporal sequencing studies investigating whether change in distress

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precedes PTSD symptom decline are lacking. These studies are essential since the timeline between indicators of change and treatment outcome is a key assumption for mediation. The aim of the present study was to assess the temporal relationship between within- and between-session change in subjective distress and PTSD symptom decrease. We analyzed session data from 86 patients with PTSD. Data were analyzed using dynamic panel models. We distinguished temporal effects (within-persons) from averaged effects (between-persons). Results regarding the temporal effect showed that within-session change in subjective distress preceded PTSD symptom improvement while the reversed effect was absent. Averaged within-session change in subjective distress was also related to PTSD symptom improvement. Results regarding the temporal effect of between-session change in subjective distress

showed that it did not precede PTSD symptom improvement. Averaged between-session change in subjective distress was related to PTSD symptom improvement. This study provides evidence for within- but not between-session change in subjective distress as indicator of change during PE. We also found that the way of modeling potential indicators of change affects results and implications. We recommend future studies to analyze mediators during treatment using temporal rather than averaged effects.

Keywords: PTSD; prolonged exposure; working mechanism; change in distress; temporal sequencing; dynamic panel model

PROLONGED EXPOSURE (PE) is a widely researched and effective psychotherapy for Posttraumatic Stress Disorder (PTSD), but remission rates leave ample room for improvement (Lee et al., 2016; Mavranouzouli et al., 2020; Watts et al., 2013). Investigating indicators of mechanisms of change, i.e. processes responsible for symptom change, will lead to a better understanding of the theoretical underpinnings of PE and may provide directions for further improvements (Kazdin, 2007; Kindt, 2014). Emotional Processing Theory (EPT) has long been the dominant theory on PE's mechanisms of change (Foa & Kozak, 1986). In short, EPT proposes that prolonged exposure to fear-evoking stimuli leads to emotional processing that, in turn, leads to symptom alleviation. Emotional processing is not directly measurable (Foa & McLean, 2016), but within-session change in subjective distress and between-session change in subjective distress are suggested to be indicators of change as they indicate emotional processing taking place (Foa & Kozak, 1986; Foa & McLean, 2016).

A large body of work supports the proposition that between-session change in subjective distress¹ is related to positive treatment outcome in patients with PTSD (e.g., Cooper, Clifton, & Feeny, 2017; see Table 1 for overview), although this work has also been criticized (e.g., Craske et al., 2008). Reasons for this criticism include limited use of complete session data—either by averaging session data or only considering the first and last sessions—and the categorization of outcome in (re-

sponder) categories that do not allow for a direct evaluation of the relationship between the indicators of change and outcome (Craske et al., 2008). Moreover, given that many previous studies had small samples to begin with (see Table 1), results may be unreliable. Most studies found no evidence that within-session change in subjective distress and symptom improvement are related. But note: these studies suffered from the same limitations as studies into between-session change in subjective distress. Importantly, nearly all of the previous studies considered the *averaged* effect of change in subjective distress (across individuals), referring to the relationship between averaged change in subjective distress across all sessions and treatment outcome. The *temporal effect* of change in subjective distress, referring to the relationship between change in subjective distress at timepoint X and outcome at timepoint X+1 within a person, has rarely been investigated (see Table 1). Temporal effects, however, are much more likely to reflect indicators of change than averaged effects, so the omission of temporal effects is problematic (Falkenstrom et al., 2020; Kazdin, 2007).

Establishing a timeline between an indicator of change and symptom change is in fact a crucial *prerequisite* for establishing mediation (Hayes, 2013; Kazdin, 2007; Kumpula et al., 2017) and the direction of the relationship between change in subjective distress and symptom change is as yet unclear. Previous results showing that averaged between-session change in subjective distress and symptom change are related may refer to three different associations: between-session change in subjective distress *precedes* symptom improvement, *co-occurs* with symptom improvement, or *follows* symptom improvement. Only the first is relevant from the perspective of mechanisms of change. Second, temporal relations are clinically relevant as they provide information about change processes on an individual level. In contrast, averaged effects may be influenced by (unchangeable) covariates at the individual level and are therefore less informative for change processes. For example, patients with high intelligence might have more between-session change in subjective distress and more symptom improvement while these are temporally unrelated to each other. Third, using temporal data has statistical advantages as it results in more power than averaged data and takes covariates at the person level into account. When averaged relationships are generalized to temporal relationships, these covariates may result in biased conclusions (Hamaker, 2012). For example, on average, a higher number of PE sessions might be related to worse treatment outcomes

¹ Note that in previous work, the terms *habituation* or *extinction* have been interchangeably used to describe subjective change in distress levels during exposure sessions, while these terms actually refer to theoretically distinct mechanisms. To avoid theoretical confusion, we use the descriptive term *change in subjective distress* throughout this article.

Table 1
Evidence for the Effect of Within- and Between-Session Change in Distress as Mediators of Prolonged Exposure

Study	Year	Sample size	Mechanism of change	Within person data mechanism	Within person data outcome	Within-session	Between-session
Norr et al.	2019	108	Within and Between	Not used	Not used	–	+/-
Reger et al.	2019	96	Between	Used	Not used	NA	+
Rauch et al.	2018	97	Within and Between	Used	Not used	–	+
Hendriks et al.	2018	69	Within and Between	Not used	Not used	–	+
Badour et al.	2017	46	Within and Between	Not used	Not used	–	+
de Kleine et al.	2017	50	Within and Between	Not used	Not used	+	+
Wisco et al.	2016	22	Between	Used	Not used	NA	+
Harned et al.	2015	16	Within and Between	Not used	Not used	–	+
Nacasch et al.	2015	39	Within and Between	Not used	Not used	–	+
Sripada et al.	2015	12	Within and Between	Used	Not used	–	+
de Kleine et al.	2015	67	Within and Between	Used ¹	Used ¹	+	+
Bluett et al.	2014	88	Between	Not used	Not used	NA	+
Gallagher et al.	2012	88	Between	Not used	Not used	NA	+
Van Minnen et al.	2006	92	Within and Between	Not used	Not used	–	+
Rauch et al.	2004	69	Between	Not used	Not used	NA	+
Van Minnen et al.	2002	34	Within and Between	Not used	Not used	–	+
Jaycox et al.	1998	37	Within and Between	Not used	Not used	–	+

Note. ¹Used for within-session but not for between-session change in distress.

NA = Not applicable; + = significant finding; – = non-significant finding; +/- = mixed finding.

(between-persons). However, this might be due to covariates at the person level, e.g., persons who respond well may finish treatment early. If this averaged result is generalized to a temporal effect one might falsely conclude that providing more PE sessions to a patient leads to poorer treatment outcome.

Almost all studies on the effect of change in subjective distress as indicator of change during PE have used averaged-person data, raising doubts about the conclusions. The only exception is a study about the effect of within-session change in subjective distress on symptom change during D-cycloserine- versus placebo-enhanced PE (de Kleine, Smits, Hendriks, Becker, & van Minnen, 2015). This study is one of only two studies (de Kleine et al., 2017; de Kleine et al., 2015) that found a significant relationship between within-session change in subjective distress and PTSD symptom improvement. This raises the question whether earlier null-findings on the effect of within-session change in subjective distress on symptom change might be explained by the data-analytic strategy. Ideally, a study using temporal data would also report on averaged-person relationships as “control analysis,” as this allows a better comparison to previous findings in this field.

The aim of the current study was to investigate whether within- and between-session change in subjective distress is related to PTSD symptom improvement using temporal data. We studied the timeline between change in subjective distress and symptom improvement using dynamic panel models. These models allow for distinguishing temporal effects from averaged effects without violating assumptions (a problem with mixed-model analyses; see Allison et al., 2017; Hamaker & Muthen, 2019; Leszczensky & Wolbring, 2019). Based on the premises of EPT, we expected change in subjective distress, both within- and between-sessions, to predict next session change in PTSD symptoms. To test temporality, we reversed predictors and outcome, and expected that PTSD symptoms would not—or to a lesser extent—predict subsequent changes in subjective distress within- or between-sessions. To allow comparison with previous studies, we also assessed the averaged-person effect of change in subjective distress within- and between-sessions to elucidate whether the use of temporal data leads to different results than the use of averaged data. Based on previous findings (Cooper, Clifton, et al., 2017), we expected averaged change in subjective distress between-sessions, but not within-sessions, to predict PTSD symptom decrease.

Method

PARTICIPANTS

We used the data from the IMPACT study (Opriel et al., 2018), a multicenter randomized controlled trial comparing PE with intensified PE (iPE) and phase-based treatment compromising Skills Training in Affective and Interpersonal Regulation followed by PE (STAIR+PE). The trial is registered at the clinical trials registry, number NCT03194113. All participants (1) met DSM-5 diagnosis of PTSD established with the Clinician Administered PTSD Scale (CAPS-5) with moderate-severe PTSD-symptoms (CAPS-5 score ≥ 26) following repeated interpersonal childhood physical/sexual abuse by a primary caretaker or an authority figure and had at least one specific memory of the traumatic event (Boeschoten et al., 2015), (2) were between 18 and 65 years old, and (3) spoke Dutch. Participants were excluded when they (1) were involved in a compensation case or legal procedures concerning admission or stay in The Netherlands, (2) were pregnant, (3) engaged in severe nonsuicidal self-injury (NSSI) that required hospitalization during the past 3 months, (4) engaged in severe suicidal behavior defined by either a suicide attempt during the past 3 months or acute suicidal ideations with serious intent to die with a specific plan for suicide and preparatory acts, (5) had a severe disorder in the use of alcohol or drugs (≥ 6 symptoms) in the last 3 months according to the Mini-International Neuropsychiatric Interview version 7.0.2 for DSM-5 (MINI; Sheehan et al., 1998), (6) suffered from cognitive impairment (estimated IQ < 70), (7) changed psychotropic medication in the 2 months prior to inclusion, or (8) engaged in any current psychological treatment. Informed consent was obtained prior to randomization from all participants. For this article, we included participants from the exposure only conditions²: PE ($n = 48$) and iPE ($n = 51$). Patients also had to complete at least two PE sessions with measurements of subjective distress levels and PTSD symptoms, such that a timeline could be established ($n_{PE} = 44$, $n_{iPE} = 42$). Most patients were female (79%) and patients had an age between 20 and 60 years old ($M = 36.8$, $SD = 11.5$). Almost half (40%) of the patients had a non-Western cultural background, 20% of the patients were highly educated

² The STAIR+PE condition is excluded because it is based on the notion that skills training in the first phase of treatment will increase the tolerability of PE and therefore influences the proposed working mechanism of PE. This precludes conclusions about the working mechanism of PE.

(i.e. higher vocational education or university), 43% of the patients were employed and 51% of the patients used psychotropic medication. Patients suffered on average from 3.0 comorbid Axis-1 diagnoses ($SD = 1.9$) in addition to the PTSD diagnosis and 47% of the patients suffered from severe suicidality according to the MINI (Sheehan, et al., 1998). Moreover, 62% of the patients met criteria for a personality disorder according to the Structured Clinical Interview for DSM-IV Personality Disorders (SCID-2; Weertman et al., 2003). We refer to the design paper for detailed information about the design, recruitment, participants, procedure or therapy (Oprel et al., 2018) and to the main outcome paper for detailed information about the study sample (Oprel, et al., 2021). The study was approved by the Medical Ethical Committee of Leiden University Medical Center (NL57984.058.16).

PROCEDURE

After enrollment, patients were randomized to PE, iPE and STAIR+PE (1:1:1 ratio) by an independent researcher based on a computerized randomization sequence of permuted blocks of six participants stratified by gender. Prolonged exposure (PE) was delivered in 16 weekly sessions of 90 minutes. Intensive prolonged exposure (iPE) was delivered in 14 sessions of 90 minutes starting with three weekly sessions for four weeks followed by two sessions after one and two months. For practical reasons, iPE was alternately provided by two therapists. The treatment manual of PE and iPE was identical and largely based on the protocol by Foa et al. (2007). The exposure sessions involved psychoeducation in the first session and 60 minutes imaginal exposure and exposure in vivo from the second session onwards. During imaginal exposure, patients were instructed to repeatedly and vividly recount the most disturbing traumatic memories. Between sessions, patients listened to recordings of the imaginal exposure and performed in-vivo homework assignments. The exposure sessions involved psychoeducation in the first session and 60 minutes of imaginal exposure from the second session onwards. Therapists' adherence to the PE protocol was ensured through training, an exam with pilot patients graded by supervisors and weekly group supervision (supervisors: RAdK and AvM). A random selection of the PE sessions (135 sessions; ~10% of the total sessions) was rated by independent observers for treatment adherence based on the Dutch translation of the original adherence rater checklist scale by Foa and colleagues. Protocol adherence was

high (M session elements completed = 90%, $SD = 18\%$).

For this paper, data from session 15 and 16 of the PE condition were omitted, because these sessions did not include sufficient observations for the temporal models (only 18 patients [21%] completed session 15 and 15 patients [17%] completed session 16).

MEASURES

Weekly changes in PTSD symptoms were assessed during every session of PE and during session 1, 4, 7, 10, 12, 13, and 14 of iPE. Subjective distress levels were assessed during in-session exposure, every session from the second session onwards.

PTSD symptoms

The primary outcome of this study was self-reported PTSD symptom severity measured with the weekly version of the PTSD checklist for DSM-5: PCL-5 (Blevins, Weathers, Davis, Witte, & Domino, 2015). The PCL-5 consists of 20 items scored on a five-point Likert scale, ranging from 0 (not at all) to 4 (extremely), with total scores ranging from 0-80. The PCL-5 demonstrated high internal consistency in previous studies, high test-retest reliability and convergent and divergent validity with other measures (Blevins et al., 2015; Van Praag et al., 2020) and showed substantial agreement with a clinical interview for assessing PTSD in a Dutch population (van der Meer et al., 2017). The PCL-5 demonstrated high internal consistency in previous studies (Cronbach's $\alpha = .94$; Blevins, et al., 2015). In the current sample, the PCL-5 had a high internal consistency at the first session (Cronbach's $\alpha = .89$). For the standard PE condition, data was available for 44 patients who completed on average 12.07 sessions (range 3–16, total sum of sessions = 531). The PCL-5 was assessed at the start of every session and completed in 98.5% of the sessions ($n = 523$). For the iPE condition, data was available for 42 patients who completed on average 12.83 sessions (range 4–14, total sum of sessions = 539). The PCL-5 was assessed at the start of session 1, 4, 7, 10, 12, 13 and 14 (total sum of sessions with PCL-5 = 265) and completed in 97.7% of the sessions ($n = 259$).

Change in Subjective Distress Within and Between Sessions

During the 60 minutes of imaginal exposure of PE (every session except the first session), participants' subjective distress was assessed with subjective units of distress (SUDs). Every 10 minutes, the participants rated their subjective distress on a scale from 0 (*no distress*) to 100 (*maximum dis-*

ress). The SUD peak was indicated by the highest subjective distress score within a session and SUD end was indicated by the last observed subjective distress score within a session. In line with EPT (Foa & McLean, 2016) and previous work (e.g., Harned et al., 2015; Hendriks, et al., 2018; Nacasch et al., 2015), change in subjective distress within-session was indicated by the difference between the SUD peak and SUD end of a session. Change in subjective distress between sessions was indicated by the change in SUD peak ratings over two subsequent sessions. In 21 sessions (2.1% of the total exposure sessions) the therapist or patient refrained from performing any exposure in-session, so there was no SUDs data available for those sessions. Of all sessions wherein exposure took place for PE ($n = 474$), SUDs data were available for 96.0% ($n = 455$) of the sessions. Of all sessions wherein exposure took place for iPE ($n = 489$), SUD data were available for 97.8% of the sessions ($n = 478$). For the temporal analyses, we used the data per session for within- and between-session change in subjective distress. For the averaged analyses, data of within- and between-session change in subjective distress was averaged over all sessions per person.

STATISTICAL ANALYSES

The data analysis plan was preregistered at OSF (Center for Open Science; Hoeboer, et al., 2020). We used dynamic panel models based on maximum likelihood estimation (Allison et al., 2017) following recent recommendations for models with lagged dependent variables (Falkenstrom et al., 2020; Xu et al., 2019). Models were fitted using structural equation models (SEM) with R package Lavaan and dpm (Rosseel, 2012). In these models, results are corrected for stable, unobserved heterogeneity between persons and reverse causation (Allison et al., 2017). We corrected for the autoregressive effect of the outcome variable (the effect of the outcome at time point X-1 on the same outcome at time point X) and used cross-lagged effects of predictors (the effect of the predictor at time point X-1 on outcome at time point X). We used fixed effect models, which included a random intercept that was allowed to correlate with predictors, thereby correcting for the effect of clustering without violating the assumption of independent errors. Missing data was handled using full information maximum likelihood (FIML). The temporal relationship between mediators and outcome is by default estimated with the fixed effect model of the dynamic panel model. We included bootstrapped standard errors in all analyses to account for violations to the nor-

mal distribution of the data. This was especially relevant for the analyses with change in subjective distress as dependent variable. The assumptions of all models were met.

Temporal Analyses

In the first analysis, we assessed a dynamic panel model with the PCL-5 scores as dependent variable and with the autoregressive effect of the PCL-5 and cross-lagged *within*-session change in subjective distress as independent variables. For example, PCL-5 scores in session 4 were predicted by PCL-5 scores in session 3 and within-session change in subjective distress ($SUD_{\text{peak}} - SUD_{\text{end}}$) during session 3. In the iPE condition, participants had multiple sessions per week, while the PCL-5 was administered once per week. Therefore, only the SUDs data that was directly linked to PCL-5 assessment was used from this condition (e.g. session 3 included no PCL-5 score so within-session change in subjective distress from session 2 was not used).

In the second analysis, we assessed a dynamic panel model with PCL-5 scores as dependent variable and with the autoregressive effect of the PCL-5 and cross-lagged *between*-session change in subjective distress as independent variables. To illustrate, PCL-5 scores at session 4 were predicted by PCL-5 scores at session 3 and the change in peak distress between session 2 and 3 ($SUD_{\text{peak session2}} - SUD_{\text{peak session3}}$).

As the two exposure conditions differed in their delivery format (weekly vs. intensive) and the delivery format might affect change mechanisms, we ran two additional analyses to investigate the effect of condition on the relationship between change in distress and PCL-5 outcomes. These analyses were carried out using the same model as for the primary analyses, but additionally included condition (PE versus iPE) and the interaction effect between condition and mediators. If condition proved to affect outcomes, analyses were carried out per condition.

To test temporality, we next ran dynamic panel models testing effects in the opposite direction. In the third analysis, we included *within*-session change in subjective distress as dependent variable and the autoregressive effect of *within*-session change in subjective distress and cross-lagged change in PCL-5 scores as independent variables. In the fourth analysis, we included *between*-session change in subjective distress as dependent variable and the autoregressive effect of *between*-session change in subjective distress and cross-lagged change in PCL-5 scores as independent variables.

Averaged Analyses

To test whether using temporal data would lead to different results than using averaged-person data, we performed two analyses with averaged-person effects. The averaged-person effect was estimated using a fixed-effect model including person-averaged mediators. In the first analysis, we assessed a dynamic panel model with PCL-5 scores as dependent variable and with the autoregressive effect of PCL-5 scores and averaged change in subjective distress within-sessions as independent variables. In other words, we assessed the effect of the *average* change in subjective distress on PTSD symptom change over the course of treatment. In the second analysis, we assessed a dynamic panel model with PCL-5 score as dependent variable and with the autoregressive effect of PCL-5 scores and averaged change in subjective distress between sessions as independent variables.

Results

Fifty-five (64%) of the 86 patients who were included in this study completed 14 sessions. The PCL-5 scores decreased during the course of treatment, from on average 54.24 ($SD = 12.72$) in the first session to on average 30.80 ($SD = 23.10$) in session 14. Within-session change in subjective distress showed a large variation between patients and was larger at the start of treatment ($M_{\text{session } 2} = 25.48$; $SD_{\text{session } 2} = 23.82$) compared to the end of treatment ($M_{\text{session } 14} = 15.07$; $SD_{\text{session } 14} = 19.22$). Between-session change in subjective distress also showed a large variation between patients without clear pattern over the course of treatment ($M_{\text{session } 3} = 7.01$; $SD_{\text{session } 3} = 14.90$ to $M_{\text{session } 14} = 5.44$; $SD_{\text{session } 14} = 21.97$; see [Table 2](#) for more details).

TEMPORAL ANALYSES

We found that within-session change in subjective distress was significantly related to lower PTSD symptoms in the next session (i.e. the temporal effect): $b = -.04$, $SE = .02$, $z = -2.17$, $p = .03$, Cohen's $d = .48$, while correcting for the autoregressive effect of PTSD symptoms (see [Table 3](#)). This effect was not different for iPE compared to PE ($b = .01$, $SE = .05$, $z = .27$, $p = .79$). The reversed temporal effect of PTSD symptom change on next session's within-session change in subjective distress was not significant: $b = -.08$, $SE = .09$, $z = -.85$, $p = .40$, while correcting for the autoregressive effect of within-session change in subjective distress.

We found that between-session change in subjective distress was not significantly related to lower PTSD symptoms in the next session (i.e.,

the temporal effect): $b = .003$, $SE = .02$, $z = .17$, $p = .86$, while correcting for the autoregressive effect of PTSD symptoms (see [Table 4](#)). This effect was not different for iPE compared to PE ($b = -.03$, $SE = .04$, $z = -.73$, $p = .47$). The reversed temporal effect of PTSD symptom change on between-session change in subjective distress in the next session was also not significant $b = .05$, $SE = .12$, $z = .39$, $p = .70$, while correcting for the autoregressive effect of between-session change in subjective distress.

AVERAGED ANALYSES

Averaged within-session ($b = -.16$, $SE = .05$, $z = -3.06$, $p = .002$, Cohen's $d = .70$) and between-session ($b = -.53$, $SE = .20$, $z = -2.71$, $p = .007$, Cohen's $d = .61$) change in subjective distress were both related to lower PTSD symptoms over the course of treatment while correcting for the autoregressive effect of PTSD symptoms.

Discussion

The main goal of this study was to test the effect of change in subjective distress during prolonged exposure (PE) therapy on PTSD symptom improvement using temporal analyses. The results indicated that within- and not between-session change in subjective distress preceded symptom improvement. These findings stand in contrast to the commonly expressed finding that between- and not within-session change in subjective distress is related to better treatment response (e.g., [Asnaani et al., 2016](#); [Brown et al., 2019](#); [Cooper, Clifton, et al., 2017](#); [Foa & McLean, 2016](#)). Importantly, in the current work we used a new-analytic framework ([Allison et al., 2017](#)) and distinguished temporal from averaged effects ([Falkenstrom et al., 2020](#); [Hamaker, 2012](#); [Hamaker & Muthen, 2019](#)), which probably explains the divergent findings.

Our first hypothesis, that within-session change in subjective distress would predict change in PTSD symptoms to the next session, was confirmed. Crucially, we did not find the reversed effect. Our findings thus point to within-session change in subjective distress as an indicator of change during PE, as it precedes and predicts symptom improvement ([Kazdin, 2007](#)). This finding is in line with EPT, but stands in contrast with most previous studies that examined the effect of within-session subjective change in distress on PE outcome (see [Table 1](#)). Notably, these studies used data-analytic strategies that only considered averaged effects. The only other study using temporal data for both within-session change in subjective distress and PTSD symptom change during PE

Table 2
Descriptive Information About Mechanisms of Change and Outcome as a Function of Session

Session	PCL-5			Within-session change in distress			Between-session change in distress		
	N	M	SD	N	M	SD	N	M	SD
1	85	54.29	12.72						
2	43	55.93	12.59	86	25.48	23.82			
3	44	54.25	15.74	85	24.94	26.54	85	7.01	14.90
4	83	50.61	15.32	83	22.35	21.00	82	1.77	15.64
5	42	46.95	17.87	79	23.99	21.72	79	5.72	14.74
6	40	46.10	18.42	73	21.63	20.73	74	3.95	21.51
7	73	42.93	18.83	73	20.41	18.76	72	-0.10	16.90
8	35	38.03	21.71	69	18.96	17.63	68	2.81	17.77
9	33	34.94	21.17	64	18.91	19.51	66	2.18	16.28
10	66	36.50	20.42	64	21.20	21.67	63	-0.05	14.96
11	27	32.93	23.60	63	19.52	17.89	62	4.21	19.30
12	62	32.08	20.07	60	19.67	21.30	59	5.24	17.42
13	62	30.35	20.95	56	15.02	16.58	55	6.69	23.53
14	55	30.80	23.10	46	15.07	19.22	48	5.44	21.97

Note. PCL-5 = PTSD checklist for DSM-5.

Table 3
Temporal Effect of Within-Session Change in Subjective Distress on Next Session's PTSD Symptoms and Reversed Effect of PTSD Symptom Change on Next Session's Within-Session Change in Subjective Distress

Temporal effects	Estimate	SE	z-value	p-value
Lagged within-session change in subjective distress	-.04	.02	-2.17	.03
Autoregressive effect PCL-5 score	.70	.06	12.37	<.001
Reversed effects				
Lagged change in PCL-5 score	-.08	.09	-.85	.40
Autoregressive effect within-session change in subjective distress	.11	.07	1.75	.08

Note. PTSD = Posttraumatic Stress Disorder; PCL-5 = PTSD checklist for DSM-5.

Table 4
Temporal Effect of Between-Session Change in Subjective Distress on Next Session's PTSD Symptoms and Reversed Effect of PTSD Symptom Change on Next Session's Between-Session Change in Subjective Distress

Temporal effects	Estimate	SE	z-value	p-value
Lagged between-session change in subjective distress	.003	.02	.17	.86
Autoregressive effect PCL-5 score	.66	.09	7.78	<.001
Reversed effects				
Lagged change in PCL-5 score	.05	.12	.39	.70
Autoregressive effect between-session change in subjective distress	-.41	.06	-7.56	<.001

Note. PTSD = Posttraumatic Stress Disorder; PCL-5 = PTSD checklist for DSM-5.

found similar results (de Kleine et al., 2015). Our findings imply that within-session reduction of subjective distress precedes PTSD symptom change during PE. This is of clinical relevance, as in-session indices of change can guide clinicians in their implementation of PE.

In contrast to our expectations, we found that averaged within-session change in subjective distress was also related to change in PTSD symptoms. This is remarkable as the data-analytic strategy for this analysis was in line with earlier work, yet leading to a different outcome. Our find-

ing implies that those with, on average, more within-session change in subjective distress showed more change in PTSD symptoms. One important factor that might explain our divergent findings is a difference in statistical power. Notably, about half of the previous studies that assessed within-session change in distress included small sample sizes with less than 40 patients (Harned et al., 2015; Jaycox et al., 1998; Nacasch et al., 2015; Sripada & Rauch, 2015; van Minnen & Hageraars, 2002). Moreover, these studies mostly defined outcome as a pre-post difference rather

than utilizing the repeated measurements per patient (resulting in far less power; e.g., [Morgan & Case, 2013](#)). Therefore, these studies lacked adequate power resulting in increased false positive and false negative findings (see for rationale: [Button et al., 2013](#)). In line, a recent meta-analysis on change in subjective distress on symptom improvement during PE concluded that there was insufficient power to establish the effect of within-session change in subjective distress on outcome ([Rupp et al., 2017](#)).

Our second hypothesis, that between-session change in subjective distress predicts change in PTSD symptoms in the next session, was not confirmed, nor did we find the reversed effect. This finding contradicts previous studies that consistently found between-session change in subjective distress to be related to PTSD symptom change (see [Table 1](#)). However, this difference might be explained by our different data-analytic method. Previous studies did not use temporal analyses but assessed averaged effects. Indeed, in line with previous work, we found that averaged between-session change in subjective distress predicted change in PTSD symptoms. As these analyses omit the temporal relationship between indicators of change and outcome, this relationship might be driven by a third factor related to both the indicator of change and outcome (i.e., personal characteristics such as learning ability) or time-congruency of both factors. The latter would imply that between-session change in distress might be a *proxy* of treatment response, rather than an indicator of change ([Cooper, Clifton, et al., 2017](#)). To conclude, our results indicate that between-session reduction in distress does not precede PTSD symptom decline. These results are supported by previous work that showed that patients without between-session change in distress also improved over the course of treatment (e.g., [Bluett, et al., 2014](#)).

This is the first temporal sequencing study about within- and between-session change in subjective distress as indicators of change during PE. Although temporal precedence is a key assumption often overlooked when studying change processes ([Kazdin, 2007](#)), it does not in itself suggest a mechanistic relationship. To establish mechanisms of change, additional evidence is required such as experimental evidence of cause (see [Tryon, 2018](#)). Note that our results also do not imply that within-session reduction of subjective distress is the only indicator of change during PE, as it is likely that multiple change mechanisms explain treatment outcome ([Kredlow et al., 2020](#); [Vervliet et al., 2013](#)). Based on novel insights from

emotional learning research, the inhibitory learning theory (ILT; [Craske et al., 2014](#)) postulates that the learning and retrieval of inhibitory non-threat associations is crucially important for successful treatment outcome. Both EPT and ILT are rooted in extinction theory and partially overlap in theoretical mechanisms ([Cooper, Clifton, et al., 2017](#)), but the theories differ with respect to their view on distress reduction as an index of meaningful change. In short, ILT proposes that distress reduction may be a by-product of inhibitory learning. ILT proposes new indices of meaningful change during exposure therapy such as expectancy violation or enhanced tolerance of distress ([Bluett et al., 2014](#); [Craske et al., 2008](#); [Knowles & Olatunji, 2019](#); [Sripada et al., 2016](#)). Future studies might test whether these indices also precede and fuel PTSD symptom decrease, and how they relate to distress reduction. Moreover, EPT also proposes other indices of emotional processing such as emotional engagement. Strong empirical evidence for the relevance of emotional engagement is lacking ([Cooper, Clifton, et al., 2017](#)), but so are temporal studies assessing its relevance. Thus, future studies might also examine such indices with temporal models.

An already previously established indicator of change during PE is the reduction of maladaptive trauma-related cognitions ([Cooper, Clifton, et al., 2017](#)). In studies focusing on trauma-related cognitions (e.g. “the world is dangerous” or “I have no future”), mixed-effect models including temporal data have already been successfully used to establish the timeline between these cognitions and PTSD symptom improvement ([Cooper, Zoellner, et al., 2017](#); [Kumpula et al., 2017](#); [Zalta et al., 2014](#)). Changes in trauma-related cognitions were found to be related to symptom improvement during PE and to precede symptom improvement. Our current findings add to these findings as within-session change in subjective distress also predicted and preceded symptom improvement during PE. An important next step is to test several indicators of change simultaneously in one model, to better understand how they (interactively) lead to symptom PTSD change. In light of the recent developments in the availability of statistical algorithms to adequately model temporal data and lagged effects (e.g. using dynamic panel models; [Rosseel, 2012](#)), we also urge future studies into mechanisms of change to take temporality into account and distinguish averaged relationships (between-persons) from temporal relationships (within-persons). Note that already collected data might also be re-analyzed using temporal sequencing models to improve understand-

ing about within- and between-session change in subjective distress as indicators of change during PE. Future studies might also consider the use of experience sample and ecological momentary assessments to establish a timeline between indicators of change and symptom change more precisely (see, for example, [Padovano & Miranda, 2018](#)).

The current study has several limitations. First, the intensified PE condition in our study did not have session data available for every exposure session and included only fourteen PE sessions. This resulted in less temporal precision in this condition, less data and consequently less power. Second, the panel data in our study was unbalanced due to missing data which is inherent to clinical trials but reduces statistical power ([Moral-Benito et al., 2019](#)). This was especially problematic for sessions 15 and 16 of the PE condition, which were therefore omitted for the analyses. Related to this, the current sample size did not allow for assessing multiple indicators of change in one dynamic panel model. Future studies may consider including other relevant predictors of symptom improvement in dynamic panel models such as homework adherence ([Cooper, Kline, et al., 2017](#)). Third, exposure sessions in our study included a total of 60 minutes of exposure, which is similar to some previous studies (e.g., [de Kleine et al., 2017](#); [Gallagher & Resick, 2012](#); [Hendriks et al., 2018](#)), but diverges from others that included a total of 30–45 minutes of in-session exposure (e.g., [Bluett et al., 2014](#); [Harned et al., 2015](#); [Norr et al., 2019](#)). As already noted by [Cooper, Clifton, et al. \(2017\)](#) in their review, duration of the in-session exposure may affect patterns of change in distress, and therefore results have to be replicated in studies using the same data-analytic strategy and different exposure lengths. Finally, the assessment method of change in distress in the current study (subjective self-reportage) differs from methods used in controlled laboratory research on underlying mechanisms of fear extinction that commonly include physiological indicators of distress ([Carpenter et al., 2019](#)). Physiological measures of distress might, therefore, be an important additional indicator of change in distress and have already been shown to relate to treatment response in previous research ([Wangelin & Tuerk, 2015](#)).

To conclude, we found that within, but not between-session change in subjective distress predicted next session's change in PTSD symptoms using temporal data. Against contemporary belief, these results indicate that within-session change in subjective distress is an indicator of change during PE. This suggests that within-session change in

subjective distress could be used to monitor treatment progress. Since this is the first study to investigate temporal relationships between change in subjective distress and PTSD symptom change, more research is needed to replicate these findings.

Conflict of Interest Statement

The authors declare that there are no conflicts of interest.

References

- Allison, P. D., Williams, R., & Moral-Benito, E. (2017). Maximum likelihood for cross-lagged panel models with fixed effects 2378023117710578. *Socius*, 3. <https://doi.org/10.1177/2378023117710578>.
- Asnaani, A., McLean, C. P., & Foa, E. B. (2016). Updating Watson & Marks (1971): How has our understanding of the mechanisms of extinction learning evolved and where is our field going next? *Behavior Therapy*, 47, 654–668. <https://doi.org/10.1016/j.beth.2016.02.003>.
- Badour, C. L., Flanagan, J. C., Gros, D. F., Killeen, T., Pericot-Valverde, I., Korte, K. J., Allan, N. P., & Back, S. E. (2017). Habituation of distress and craving during treatment as predictors of change in PTSD symptoms and substance use severity. *Journal of Consulting and Clinical Psychology*, 85, 274–281. <https://doi.org/10.1037/ccp0000180>.
- Blevins, C. A., Weathers, F. W., Davis, M. T., Witte, T. K., & Domino, J. L. (2015). The posttraumatic stress disorder checklist for DSM-5 (PCL-5): Development and initial psychometric evaluation. *Journal of Traumatic Stress*, 28, 489–498. <https://doi.org/10.1002/jts.22059>.
- Bluett, E. J., Zoellner, L. A., & Feeny, N. C. (2014). Does change in distress matter? Mechanisms of change in prolonged exposure for PTSD. *Journal of Behavior Therapy and Experimental Psychiatry*, 45, 97–104. <https://doi.org/10.1016/j.jbtep.2013.09.003>.
- Boeschoten, M. A., Bakker, A., Jongedijk, R. A., Van Minnen, A., Elzinga, B. M., Rademaker, A. R., & Olff, M. (2015). *Clinician administered PTSD scale for DSM-5—Dutch version*. Arq Psychotrauma Expert Groep Diemen. <https://doi.org/10.1080/20008198.2018.1546085>.
- Brown, L. A., Zandberg, L. J., & Foa, E. B. (2019). Mechanisms of change in prolonged exposure therapy for PTSD: Implications for clinical practice. *Journal of Psychotherapy Integration*, 29, 6–14. <https://doi.org/10.1037/int0000109>.
- Button, K. S., Ioannidis, J. P. A., Mokrysz, C., Nosek, B. A., Flint, J., Robinson, E. S. J., & Munafò, M. R. (2013). Power failure: Why small sample size undermines the reliability of neuroscience (vol 14, pg 365–376, 2013) 444–444. *Nature Reviews Neuroscience*, 14. <https://doi.org/10.1038/nrn3475>.
- Carpenter, J. K., Pinaire, M., & Hofmann, S. G. (2019). From extinction learning to anxiety treatment: Mind the gap. *Brain Sciences*, 9. <https://doi.org/10.3390/brainsci9070164>.
- Cooper, A. A., Clifton, E. G., & Feeny, N. C. (2017). An empirical review of potential mediators and mechanisms of prolonged exposure therapy. *Clinical Psychology Review*, 56, 106–121. <https://doi.org/10.1016/j.cpr.2017.07.003>.
- Cooper, A. A., Kline, A. C., Graham, B., Bedard-Gilligan, M., Mello, P. G., Feeny, N. C., & Zoellner, L. A. (2017).

- Homework “dose,” type, and helpfulness as predictors of clinical outcomes in prolonged exposure for PTSD. *Behavior Therapy*, 48, 182–194. <https://doi.org/10.1016/j.beth.2016.02.013>.
- Cooper, A. A., Zoellner, L. A., Roy-Byrne, P., Mavissakalian, M. R., & Feeny, N. C. (2017). Do changes in trauma-related beliefs predict PTSD symptom improvement in prolonged exposure and sertraline? *Journal of Consulting and Clinical Psychology*, 85, 873–882. <https://doi.org/10.1037/ccp0000220>.
- Craske, M. G., Kircanski, K., Zelikowsky, M., Mystkowski, J., Chowdhury, N., & Baker, A. (2008). Optimizing inhibitory learning during exposure therapy. *Behaviour Research and Therapy*, 46, 5–27. <https://doi.org/10.1016/j.brat.2007.10.003>.
- Craske, M. G., Treanor, M., Conway, C. C., Zbozinek, T., & Vervliet, B. (2014). Maximizing exposure therapy: An inhibitory learning approach. *Behaviour Research and Therapy*, 58, 10–23. <https://doi.org/10.1016/j.brat.2014.04.006>.
- de Kleine, R. A., Hendriks, L., Becker, E. S., Broekman, T. G., & van Minnen, A. (2017). Harm expectancy violation during exposure therapy for posttraumatic stress disorder. *Journal of Anxiety Disorders*, 49, 48–52. <https://doi.org/10.1016/j.janxdis.2017.03.008>.
- de Kleine, R. A., Smits, J. A., Hendriks, G. J., Becker, E. S., & van Minnen, A. (2015). Extinction learning as a moderator of d-cycloserine efficacy for enhancing exposure therapy in posttraumatic stress disorder. *Journal of Anxiety Disorders*, 34, 63–67. <https://doi.org/10.1016/j.janxdis.2015.06.005>.
- Falkenstrom, F., Solomonov, N., & Rubel, J. (2020). Using time-lagged panel data analysis to study mechanisms of change in psychotherapy research: Methodological recommendations. *Counselling & Psychotherapy Research*, 20, 435–441. <https://doi.org/10.1002/capr.12293>.
- Foa, E. B., Hembree, E. A., & Rothbaum, B. O. (2007). *Prolonged exposure therapy for PTSD: Emotional processing of traumatic experiences: Therapist guide*. Oxford University Press.
- Foa, E. B., & Kozak, M. J. (1986). Emotional Processing of Fear - Exposure to Corrective Information. *Psychological Bulletin*, 99, 20–35. <https://doi.org/10.1037/0033-2909.99.1.20>.
- Foa, E. B., & McLean, C. P. (2016). The efficacy of exposure therapy for anxiety-related disorders and its underlying mechanisms: The case of OCD and PTSD. *Annual Review of Clinical Psychology*, 12, 1–28. <https://doi.org/10.1146/annurev-clinpsy-021815-093533>.
- Gallagher, M. W., & Resick, P. A. (2012). Mechanisms of change in cognitive processing therapy and prolonged exposure therapy for PTSD: Preliminary evidence for the differential effects of hopelessness and habituation. *Cognitive Therapy and Research*, 36, 750–755. <https://doi.org/10.1007/s10608-011-9423-6>.
- Hamaker, E. L. (2012). Why researchers should think “within-person”: A paradigmatic rationale. In M. R. Mehl & T. S. Conner (Eds.), *Handbook of research methods for studying daily life*. Guilford Press. Guilford Press, pp. xxvii, p. 676.
- Hamaker, E. L., & Muthen, B. (2019). The fixed versus random effects debate and how it relates to centering in multilevel modeling. *Psychological Methods*, 25, 365–379.
- Harned, M. S., Ruork, A. K., Liu, J., & Tkachuck, M. A. (2015). Emotional activation and habituation during imaginal exposure for PTSD among women with borderline personality disorder. *Journal of Traumatic Stress*, 28, 253–257. <https://doi.org/10.1002/jts.22013>.
- Hayes, A. F. (2013). *Introduction to mediation, moderation, and conditional process analysis: A regression-based approach*. The Guilford Press.
- Hendriks, L., de Kleine, R. A., Broekman, T. G., Hendriks, G. J., & van Minnen, A. (2018). Intensive prolonged exposure therapy for chronic PTSD patients following multiple trauma and multiple treatment attempts. *European Journal of Psychotraumatology*, 9, 1425574. <https://doi.org/10.1080/20008198.2018.1425574>.
- Hoeboer, C., van der Does, W., de Kleine, R., Oprel, D. A. C., School, M., van Minnen, A., & Kooistra, M. (2020). Mechanism of change of exposure therapy. *In*.
- Jaycox, L. H., Foa, E. B., & Morral, A. R. (1998). Influence of emotional engagement and habituation on exposure therapy for PTSD. *Journal of Consulting and Clinical Psychology*, 66, 185–192. <https://doi.org/10.1037/0022-006x.66.1.185>.
- Kazdin, A. E. (2007). Mediators and mechanisms of change in psychotherapy research. *Annual Review of Clinical Psychology*, 3, 1–27. <https://doi.org/10.1146/annurev.clinpsy.3.022806.091432>.
- Kindt, M. (2014). A behavioural neuroscience perspective on the aetiology and treatment of anxiety disorders. *Behaviour Research and Therapy*, 62, 24–36. <https://doi.org/10.1016/j.brat.2014.08.012>.
- Knowles, K. A., & Olatunji, B. O. (2019). Enhancing inhibitory learning: The utility of variability in exposure. *Cognitive and Behavioral Practice*, 26, 186–200. <https://doi.org/10.1016/j.cbpra.2017.12.001>.
- Kredlow, M. A., de Voogd, L., & Phelps, E. A. (2020). Laboratory analogues and therapy procedures: A case for translation from the clinic to the laboratory. *In*.
- Kumpula, M. J., Pentel, K. Z., Foa, E. B., LeBlanc, N. J., Bui, E., McSweeney, L. B., Knowles, K., Bosley, H., Simon, N. M., & Rauch, S. A. M. (2017). Temporal sequencing of change in posttraumatic cognitions and PTSD symptom reduction during prolonged exposure therapy. *Behavior Therapy*, 48, 156–165. <https://doi.org/10.1016/j.beth.2016.02.008>.
- Lee, D. J., Schnitzlein, C. W., Wolf, J. P., Vythilingam, M., Rasmusson, A. M., & Hoge, C. W. (2016). Psychotherapy versus pharmacotherapy for posttraumatic stress disorder: Systemic review and meta-analyses to determine first-line treatments. *Depress Anxiety*, 33, 792–806. <https://doi.org/10.1002/da.22511>.
- Leszczensky, L., & Wolbring, T. (2019). How to deal with reverse causality using panel data? Recommendations for researchers based on a simulation study. *Sociological Methods & Research*. <https://doi.org/10.1177/0049124119882473>.
- Mavranzouli, I., Megnin-Viggars, O., Daly, C., Dias, S., Welton, N. J., Stockton, S., Bhutani, G., Grey, N., Leach, J., Greenberg, N., Katona, C., El-Leithy, S., & Pilling, S. (2020). Psychological treatments for post-traumatic stress disorder in adults: A network meta-analysis. *Psychological Medicine*, 1–14. <https://doi.org/10.1017/S0033291720000070>.
- Moral-Benito, E., Allison, P., & Williams, R. (2019). Dynamic panel data modelling using maximum likelihood: An alternative to Arellano-Bond. *Applied Economics*, 51, 2221–2232. <https://doi.org/10.1080/00036846.2018.1540854>.

- Morgan, T. M., & Case, L. D. (2013). Conservative sample size determination for repeated measures analysis of covariance. *Annals of Biometry & Biostatistics*, 1.
- Nacasch, N., Huppert, J. D., Su, Y. J., Kivity, Y., Dinshtein, Y., Yeh, R., & Foa, E. B. (2015). Are 60-minute prolonged exposure sessions with 20-minute imaginal exposure to traumatic memories sufficient to successfully treat PTSD? A randomized noninferiority clinical trial. *Behavior Therapy*, 46, 328–341. <https://doi.org/10.1016/j.beth.2014.12.002>.
- Norr, A. M., Bourassa, K. J., Stevens, E. S., Hawrilenko, M. J., Michael, S. T., & Reger, G. M. (2019). Relationship between change in in-vivo exposure distress and PTSD symptoms during exposure therapy for active duty soldiers. *Journal of Psychiatric Research*, 116, 133–137. <https://doi.org/10.1016/j.jpsychires.2019.06.013>.
- Oprel, D. A. C., Hoeboer, C. M., Schoorl, M., De Kleine, R. A., Wigard, I. G., Cloitre, M., Van Minnen, A., & Van der Does, W. (2018). Improving treatment for patients with childhood abuse related posttraumatic stress disorder (IMPACT study): Protocol for a multicenter randomized trial comparing prolonged exposure with intensified prolonged exposure and phase-based treatment. *BMC Psychiatry*, 18. <https://doi.org/10.1186/s12888-018-1967-5>.
- Oprel, D. A. C., Hoeboer, C. M., Schoorl, M., Kleine, R. A. d., Cloitre, M., Wigard, I. G., van Minnen, A., & van der Does, W. (2021). Effect of prolonged exposure, intensified prolonged exposure and STAIR+Prolonged Exposure in patients with PTSD related to childhood abuse: A randomized controlled trial. *European Journal of Psychotraumatology*, 12, 1851511. <https://doi.org/10.1080/20008198.2020.1851511>.
- Padovano, H. T., & Miranda, R. (2018). Using ecological momentary assessment to identify mechanisms of change: An application from a pharmacotherapy trial with adolescent cannabis users. *Journal of Studies on Alcohol and Drugs*, 79, 190–198. <https://doi.org/10.15288/jsad.2018.79.190>.
- Rauch, S. A. M., Koola, C., Post, L., Yasinski, C., Norrholm, S. D., Black, K., & Rothbaum, B. O. (2018). In session extinction and outcome in virtual reality exposure therapy for PTSD. *Behaviour Research and Therapy*, 109, 1–9. <https://doi.org/10.1016/j.brat.2018.07.003>.
- Reger, G. M., Smolenski, D., Norr, A., Katz, A., Buck, B., & Rothbaum, B. O. (2019). Does virtual reality increase emotional engagement during exposure for PTSD? Subjective distress during prolonged and virtual reality exposure therapy. *Journal of Anxiety Disorders*, 61, 75–81. <https://doi.org/10.1016/j.janxdis.2018.06.001>.
- Rossee, Y. (2012). lavaan: An R package for structural equation modeling. *Journal of Statistical Software*, 48, 1–36. <https://doi.org/10.18637/jss.v048.i02>.
- Rupp, C., Doebler, P., Ehring, T., & Vossbeck-Elsebusch, A. N. (2017). Emotional processing theory put to test: A meta-analysis on the association between process and outcome measures in exposure therapy. *Clinical Psychology & Psychotherapy*, 24, 697–711. <https://doi.org/10.1002/cpp.2039>.
- Sheehan, D. V., Lecrubier, Y., Sheehan, K. H., Amorim, P., Janavs, J., Weiller, E., Hergueta, T., Baker, R., & Dunbar, G. C. (1998). The mini-international neuropsychiatric interview (MINI): The development and validation of a structured diagnostic psychiatric interview for DSM-IV and ICD-10. *Journal of Clinical Psychiatry*, 59, 22–33.
- Sripada, R. K., & Rauch, S. A. M. (2015). Between-session and within-session habituation in prolonged exposure therapy for posttraumatic stress disorder: A hierarchical linear modeling approach. *Journal of Anxiety Disorders*, 30, 81–87. <https://doi.org/10.1016/j.janxdis.2015.01.002>.
- Sripada, R. K., Rauch, S. A. M., & Liberzon, I. (2016). Psychological mechanisms of PTSD and Its treatment. *Current Psychiatry Reports*, 18. <https://doi.org/10.1007/s11920-016-0735-9>.
- Tryon, W. W. (2018). Mediators and mechanisms. *Clinical Psychological Science*, 6, 619–628. <https://doi.org/10.1177/2167702618765791>.
- van der Meer, C. A. I., Bakker, A., Schrieken, B. A. L., Hoofwijk, M. C., & Olff, M. (2017). Screening for trauma-related symptoms via a smartphone app: The validity of smart assessment on your mobile in referred police officers. *International Journal of Methods in Psychiatric Research*, 26. <https://doi.org/10.1002/mpr.1579>.
- van Minnen, A., & Foa, E. B. (2006). The effect of imaginal exposure length on outcome of treatment for PTSD. *Journal of Traumatic Stress*, 19, 427–438. <https://doi.org/10.1023/A:1020177023209>.
- van Minnen, A., & Hagens, M. (2002). Fear activation and habituation patterns as early process predictors of response to prolonged exposure treatment in PTSD. *Journal of Trauma Stress*, 15, 359–367.
- Van Praag, D. L. G., Fardzadeh, H. E., Covic, A., Maas, A. I. R., & von Steinbuchel, N. (2020). Preliminary validation of the Dutch version of the Posttraumatic stress disorder checklist for DSM-5 (PCL-5) after traumatic brain injury in a civilian population. *PLoS One*, 15. <https://doi.org/10.1371/journal.pone.0231857>.
- Vervliet, B., Craske, M. G., & Hermans, D. (2013). Fear extinction and relapse: State of the art. *Annual Review of Clinical Psychology*, 9(9), 215–248. <https://doi.org/10.1146/annurev-clinpsy-050212-185542>.
- Wangelin, B. C., & Tuerk, P. W. (2015). Taking the pulse of prolonged exposure therapy: Physiological reactivity to trauma imagery as an objective measure of treatment response. *Depression and Anxiety*, 32, 927–934. <https://doi.org/10.1002/da.22449>.
- Watts, B. V., Schnurr, P. P., Mayo, L., Young-Xu, Y., Weeks, W. B., & Friedman, M. J. (2013). Meta-analysis of the efficacy of treatments for posttraumatic stress disorder. *Journal of Clinical Psychiatry*, 74, e541–550. <https://doi.org/10.4088/JCP.12r08225>.
- Weertman, A., Arntz, A., Dreessen, L., van Velzen, C., & Vertommen, S. (2003). Short-interval test-retest interrater reliability of the Dutch version of the structured clinical interview for DSM-IV personality disorders (SCID-II). *Journal of Personality Disorders*, 17, 562–567. <https://doi.org/10.1521/pedi.17.6.562.25359>.
- Wisco, B. E., Baker, A. S., & Sloan, D. M. (2016). Mechanisms of change in written exposure treatment of posttraumatic stress disorder. *Behavior Therapy*, 47, 66–74. <https://doi.org/10.1016/j.beth.2015.09.005>.
- Xu, R., DeShon, R. P., & Dishop, C. R. (2019). Challenges and opportunities in the estimation of dynamic models. *Organizational Research Methods*, 1–25. <https://doi.org/10.1177/1094428119842638>.
- Zalta, A. K., Gillihan, S. J., Fisher, A. J., Mintz, J., McLean, C. P., Yehuda, R., & Foa, E. B. (2014). Change in negative cognitions associated with PTSD predicts symptom reduction in prolonged exposure. *Journal of Consulting and Clinical Psychology*, 82, 171–175. <https://doi.org/10.1037/a0034735>.

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