

[Download Article](#)

9,333 total views

[View Article Impact >](https://www.altmetric.com/details.php?domain=www.frontiersin.org&citation_id=52086454)
domain=www.frontiersin.org&citation_id=52086454**REVIEW article**

Front. Behav. Neurosci., 20 November 2018

Sec. Learning and Memory

<https://doi.org/10.3389/fnbeh.2018.00281><https://doi.org/10.3389/fnbeh.2018.00281>This article is part of the Research Topic
Neurobiological Models of PsychotherapyView all 10 Articles
(/research-topics/7227#articles)

Neuroscience Informed Prolonged Exposure Practice: Increasing Efficiency and Efficacy Through Mechanisms

Monika M. Stojek (<https://www.frontiersin.org/people/u/641933>)^{1,2},Lauren B. McSweeney (<https://www.frontiersin.org/people/u/609723>)^{1,2} andSheila A. M. Rauch (<https://www.frontiersin.org/people/u/570202>)^{1,2,3*}¹ Department of Psychiatry, Emory University School of Medicine, Atlanta, GA, United States² Emory Healthcare Veterans Program, Atlanta, GA, United States³ Atlanta VA Medical Center, Atlanta, GA, United States

Prolonged exposure (PE) is an empirically supported efficacious treatment for posttraumatic stress disorder (PTSD). In this focused review, we briefly review the neurobiological networks in PTSD relevant to PE, discuss the theoretical basis of PE, review the neurobiological mechanisms underlying the effectiveness of PE and identify the enhancements that can be applied to increase treatment response and retention. Based on the reviewed studies, it is clear that PTSD results in disrupted network of interconnected regions, and PE has been shown to increase the connectivity within and between these regions. Successful extinction recall in PE is related to increased functional coherence between the ventromedial prefrontal cortex (vmPFC), amygdala and the hippocampus. Increased connectivity within the dorsolateral PFC (dlPFC) following PE is associated with more effective downregulation of emotional responses in stressful situations. Pre-existing neural connectivity also in some cases predicts response to exposure treatment. We consider various enhancements that have been used with PE, including serotonin reuptake inhibitors (SSRIs), D-cycloserine (DCS), allopregnanolone (ALLO) and propranolol, repetitive transcranial magnetic stimulation (rTMS), oxytocin and MDMA. Given that neural connectivity appears to be crucial in mechanisms of action of PE, rTMS is a logical target for further research as an enhancement of PE. Additionally, exploring the effectiveness and mechanisms of action of oxytocin and MDMA in conjunction with PE may lead to improvement in treatment engagement and retention.

Posttraumatic stress disorder (PTSD) has been increasingly recognized as a public health concern, with increased visibility as military service members return from combat deployments. Due to the psychological burden associated with its symptoms, PTSD is associated with significant physical, psychosocial and economic hardships (Ramchand et al., 2015). The hallmark symptoms of PTSD include: (1) re-experiencing the memory of the traumatic event through thoughts, images or nightmares; (2)

avoidance of the reminders of trauma; (3) negative self-image and/or deterioration in mood; and (4) physiological hyperarousal (American Psychiatric Association, 2013). Trauma focused therapies, such as exposure-based and cognitive therapies, particularly prolonged exposure (PE) and cognitive processing therapy (CPT), are efficacious for the treatment of PTSD (Watts et al., 2013) and are recommended in professional treatment guidelines (American Psychological Association, 2017; Department of Veterans Affairs & Department of Defense (VA & DoD), 2017) and the Institute of Medicine Report (Institute of Medicine, 2014). PE has been studied extensively and has consistently demonstrated causal reduction of symptoms of PTSD, anxiety and depression (Watts et al., 2013). In the era of precision medicine (Shukla et al., 2015) and objective (e.g., neurobiological and psychophysiological) measures of treatment outcome, it is crucial to move beyond establishing efficacy of treatment approaches and to investigation of the mechanisms of action of such efficacious treatments. A focus on mechanisms of action will contribute to increasing efficacy (remission and response), efficiency (time to response) and access (moving the active ingredients to new models of application).

In this review article, we will bridge two bodies of knowledge: the clinical mechanisms of action of an empirically-supported treatment for PTSD and the neurobiological underpinnings of such intervention. As such, we are bridging the bench and the bedside by writing for clinicians who provide these interventions and for the researchers designing future studies. We will begin by briefly reviewing neural circuits implicated in PTSD that are relevant in PE, followed by a description of PE and its theoretical underpinnings. We will examine the state of current knowledge on how PE impacts the neural circuits related to PTSD. We will provide a synthesis of findings to date and discuss neurobiological enhancements that have been or may be used in conjunction with PE to enhance its effectiveness. Finally, we will offer some neurobiologically-informed directions for future research and practice.

Neurobiology of PTSD

In recent years, several reviews have summarized the state of knowledge regarding neural circuits implicated in PTSD (Duval et al., 2015; Liberzon and Abelson, 2016; Sheynin and Liberzon, 2017). In the current article, we briefly review the circuits most relevant to understanding therapeutic mechanisms of action in PE.

The neural circuit underlying fear-related learning is most commonly implicated in models of PTSD (Rauch et al., 2006; Shin and Handwerker, 2009; Jovanovic and Ressler, 2010; Shvil et al., 2013). The fear neurocircuitry consists of several brain structures including the amygdala, anterior cingulate cortex (ACC) and the ventromedial prefrontal cortex (vmPFC; Shin and Liberzon, 2010). The fear network is implicated in evaluating whether a stimulus should be approached or avoided, and activity in this network is correlated with anxiety (Shin and Liberzon, 2010). The evidence from fMRI studies indicates that the amygdala, which receives sensory input and orchestrates the response to threatening signals, is overactive in PTSD, likely contributing to the exaggerated fear response and re-experiencing symptoms (Rauch et al., 2006; Shin et al., 2006; Milad et al., 2009). The vmPFC downregulates the amygdala and appears to play a critical role in extinction recall (Quirk et al., 2000). In PTSD, vmPFC is hypoactive, thus projecting less inhibitory input and contributing to the hyperactivation of the amygdala (Rauch et al., 2006; Shin et al., 2006; Liberzon and Abelson, 2016). ACC which, along with the amygdala, processes aversive stimuli and projects to the peripheral nervous system to trigger a response, has been shown to be hyperactive during extinction recall in individuals with PTSD (Hayes et al., 2012; Koch et al., 2016). This dysregulation in the fear neurocircuitry is purported to underlie the failure to extinguish the fear response over time (Rauch et al., 2006; Jovanovic and Ressler, 2010; Liberzon and Abelson, 2016) and possibly the overgeneralization of fear to non-threatening cues (Stevens et al., 2013; Lopresto et al., 2016).

The neurocircuitry implicated in context processing has also received attention in relation to PTSD etiology and maintenance (Liberzon and Abelson, 2016). Within the fear neurocircuitry, hippocampus is involved in the process of contextualization, or accurately discriminating threat in the environment (Maren and Fanselow, 1995). Hippocampal inputs provide contextual information to the amygdala and to the vmPFC, thus downregulating the amygdala and facilitating extinction learning as the network normally functions (LeDoux, 2000). In the overgeneralized conditioned fear response present in PTSD, hippocampus and vmPFC are hypoactive in environments that are safe thus projecting dampened inputs to the amygdala and failing to downregulate amygdala's functioning in those contexts (Garfinkel et al., 2014). Hypoactivity of the vmPFC and the hippocampus may contribute to the re-experiencing symptoms via difficulties in extinction learning, a process further reinforced by avoidance (Pitman et al., 2012). Individuals with PTSD have demonstrated difficulty in maintaining learned fear extinction, or extinction recall (Milad et al., 2008, 2009). In fMRI studies, patients with PTSD have reduced hippocampal and vmPFC activation, and increased ACC activation during extinction recall (Milad et al., 2009; Hayes et al., 2012; Shvil et al., 2014; Koch et al., 2016) and the contextual processing period of extinction recall (Rougemont-Bücking et al., 2011). Previous studies have found that smaller hippocampal volume is associated with PTSD (Gilbertson et al., 2002). Hippocampus volume does not appear to change longitudinally over the course of PTSD in adults (Lindgren et al., 2016) but there is some evidence for impaired hippocampus development during childhood maltreatment (Dannlowski et al., 2012; Keding and Herringa, 2015). This suggests that hippocampal volume is vulnerability factor for PTSD that is likely epigenetically shaped.

Finally, emotion regulation deficits, or difficulties in awareness and modulation of intense negative emotional states, may be a transdiagnostic factor for the development and maintenance of many psychological disorders, including PTSD (Bradley et al., 2011; Sheynin and Liberzon, 2017). Based on neuroimaging findings, two broad types of emotion regulation include explicit and implicit emotion regulation (Gyurak et al., 2011). Explicit emotion regulation is effortful and requires some level of insight and awareness (Etkin et al., 2015). The most known example is reappraisal, or an alteration of the meaning of an emotion-inducing stimulus. The brain regions implicated in reappraisal include dorsolateral PFC (dlPFC), ventrolateral PFC (vlPFC) and parietal cortex (Buhle et al., 2014; Kohn et al., 2014). Implicit emotion regulation is automatic in response to a stimulus and can occur without insight or awareness (Gyurak et al., 2011). The vmPFC is the brain region primarily implicated in implicit emotion regulation (Gyurak et al., 2011). Therefore, a neurobiological emotion regulation model posits that the PFC regions are responsible for cognitive control via stimulus interpretation (either explicit or implicit), thus downregulating the amygdala activation in response to emotionally salient stimuli (Ochsner et al., 2012; Buhle et al., 2014). Only a handful of studies examined neuropsychological and neurobiological correlates of emotion regulation in PTSD (New et al., 2009; Aupperle et al., 2012; Rabinak et al., 2014; Shepherd and Wild, 2014). The results point to the pattern of suppressing emotions, using less cognitive reappraisal (Shepherd and Wild, 2014), deficits in inhibitory control (Aupperle et al., 2012), and diminished downregulation of emotional response including hypoactivation of dlPFC (New et al., 2009; Rabinak et al., 2014). While these three broad neural circuits are not an exhaustive representation of neurobiological underpinnings of PTSD, they comprehensively represent the regions of interest when considering the effects of PE on neural activity.

Prolonged Exposure and its Theoretical Underpinnings

PE (Foa et al., 1991) is an exposure-based psychological intervention designed to treat PTSD following trauma. The main goal of PE is to promote emotional processing through deliberate systematic confrontation with trauma-related stimuli (Foa, 2011). The key components of PE are: (1) repeated imaginal exposure (IE), which requires the individual to revisit their trauma memory in a therapeutic context; (2) *in vivo* exposure (IVE) to places and situations that are avoided because they evoke stress and

anxiety; and (3) emotional processing that focuses on reviewing the experience of exposure and its impact on thoughts related to the trauma. A significant body of evidence has demonstrated the efficacy and effectiveness of PE in the treatment of PTSD and related depression, general anxiety, guilt, anger, and physical health concerns (e.g., [Foa and Rauch, 2004](#); [Rauch et al., 2009](#)). In relation to PTSD, PE has reliably established clinically significant reduction in symptoms, with large effect sizes among various treatment samples with a variety of trauma histories ([Powers et al., 2010](#)). Based on intent to treat analyses, on average, 53% of those who initiate PE no longer meet diagnostic criteria for the disorder, and the rate of diagnostic change increases to 68% among individuals who complete treatment ([Bradley et al., 2005](#)).

Development of PE was based on Emotional Processing Theory ([Foa and Kozak, 1986](#)). The fundamental tenet of EPT is that there are fear structures (expanded later to include other emotions as well as fear; [Rauch and Foa, 2006](#); expanded later to include other emotions as well as fear; [Rauch and Foa, 2006](#)) that include stimulus, response, and meaning elements, and that these structures are there to assist in response to situations of danger or threat. Activation of the adaptive fear structure is viewed as a normal and rational response to a stimulus (e.g., a car racing towards me), meaning (dangerous) that elicits a fear response (increased heart rate), followed by an action (moving out of the way) to remain safe ([Foa and Kozak, 1986](#)). Following trauma, additional unhelpful fear (or other emotion) structures develop that represent the stimulus, response, and meaning elements from the time of the trauma but that may not represent actual threat or danger outside of the specific trauma context. [Rauch and Foa \(2006\)](#) state that an optimal level of activation including all elements of the trauma structure is necessary for successful treatment. When the full memory, including all the emotional and cognitive responses, is activated, updated information that is incompatible with the trauma memory can be incorporated (reconsolidated) into the memory structure. Extinction¹ occurs in the context of repeated exposure to the feared stimulus and is marked by a reduction in physiological and emotional intensity of response to that stimulus ([Sripada and Rauch, 2015](#)). They argue that the trauma structure of individuals with PTSD contains maladaptive cognitions that underlie the maintenance of PTSD ([Foa et al., 1991](#)).

Theoretically, the learning principles of classical conditioning explain the acquisition of the fear response ([Rothbaum and Davis, 2003](#); [Blechert et al., 2007](#)). Specifically, the traumatic event represents the unconditional stimulus (UCS), which produces the unconditioned response (UCR; e.g., fear, helplessness or horror). Neutral stimuli that were present during traumatic event become conditioned stimuli (CS) that can elicit a conditioned response (CR) similar to the reaction during the initial traumatic event. In an effort to reduce the experience of fear (or other negative emotions), individuals will avoid stimuli which evoke the emotional response. Consistent with the operant conditioning model ([Mowrer, 1960](#)), this avoidance serves as a negative reinforcement strategy, reducing the experience of negative emotions associated with the CS. In patients with PTSD, the overgeneralization of conditioned fear maintains and exacerbates PTSD symptoms ([Lissek and Grillon, 2012](#)).

While EPT and the learning theory are two separate theories of PTSD, there is considerable overlap in the concepts from both theories, and the mechanisms of PE have been conceptualized using both theories ([Rauch and Liberzon, 2016](#)). The EPT concept of extinction in PE can be better described in terms of extinction and relearning including contextualization of learning and memory ([Rauch and Liberzon, 2016](#)). Extinction occurs when new inhibitory associations are formed on top of the fear associations, and is marked by a reduction in subjective fear response to the feared memory and its reminders. This relearning is facilitated through the process of contextualization, or learning to discriminate between safety and threat cues, depending in the context in which they occur ([Maren et al., 2013](#)). The cognitive and emotional processing changes that occur in tandem with extinction and relearning are marked by increased sense of competence, reduced sense that the world is dangerous, and reduction in social and emotional withdrawal ([Rothbaum et al., 2005](#); [Rauch and Liberzon, 2016](#)). Although change in self-efficacy and trauma-related beliefs is not central to learning models, this learning of increased

competence to cope with negative affect and reduced sense of a dangerous world can be viewed as a form of inhibitory learning (Rauch and Liberzon, 2016), which has been speculated to be a one of the key mechanisms of action in exposure-based treatment for PTSD (Craske et al., 2008).

Neurobiology of PE

Development of PE was not rooted in a neurocircuitry-based framework but its purported mechanisms of action and effectiveness may be examined using that framework. The fear and contextualization neurocircuitry is implicated in extinction learning and recall, one of the putative active components of PE (Jovanovic and Ressler, 2010; Liberzon and Abelson, 2016; Lopresto et al., 2016). A few early studies which used exposure-based therapy (but not PE specifically) have found increased activation in the prefrontal regions in individuals with PTSD following a course of psychotherapy (Felmingham et al., 2007; Peres et al., 2007). Following therapy, increased activation in the left PFC was correlated with decreased activation of the amygdala and increased activation in the hippocampus during retrieval of the traumatic memory by individuals with PTSD (Peres et al., 2007). During processing of threatening stimuli, individuals with PTSD demonstrated increased vmPFC (particularly rostral ACC) activation and decreased amygdala activation from pre- to post-treatment (Felmingham et al., 2007). Recent studies examined the effect of PE on extinction in laboratory settings. During fear extinction recall paradigm, individuals who underwent PE demonstrated a decrease in rostral ACC activation from pre- to post-treatment (Helpman et al., 2016a). Structurally, those who remitted from PTSD following a course of PE demonstrated volume reduction and thinning in the left rostral ACC, compared to those who did not remit and to controls (Helpman et al., 2016b). No between-group differences in ACC volume and thickness were observed prior to treatment (Helpman et al., 2016b).

One putative explanation of these functional and structural brain changes following PE is the extinction of maladaptive cognitive-emotional connections resulting from extinction learning (Helpman et al., 2016b). In PTSD, fear neurocircuitry may reinforce existing connections and contribute to the formation of new ones (Johansen et al., 2011). These results suggest that effective extinction that occurs during PE contributes not only to a more balanced feedback loop between the vmPFC and the amygdala (Felmingham et al., 2007; Peres et al., 2007) but also to the thinning of the ACC via decreased activation and pruning of the connectivity (Helpman et al., 2016b). This reciprocal relationship between neural connectivity and treatment response may also work the other way. There is some evidence that individual's capacity to benefit from PE may be modulated by the degree of spontaneous PFC downregulation of the amygdala when processing threat cues prior to treatment (Fonzo et al., 2017). Specifically, patients who before receiving a course of PE had greater activation of the dlPFC as well as less amygdala activation during an emotional reactivity task (detection and processing of threatening cues), showed the biggest gains from PE (Fonzo et al., 2017). This finding potentially suggests that extinction learning and recall may be more difficult for some individuals to achieve based on the extent to which their vmPFC is able to downregulate the amygdala during exposure. Hippocampus is another structure integral to the fear neurocircuitry. Structural differences in the contextualization neurocircuitry, particularly the hippocampus, have been shown to be associated with vulnerability to PTSD (Gilbertson et al., 2002; Lindgren et al., 2016). PE responders and controls had greater baseline hippocampal volume compared to treatment non-responders (Rubin et al., 2016), indicating that hippocampal volume may not only confer risk for PTSD development (Gilbertson et al., 2002; Lindgren et al., 2016) but also be related to better outcome in PE. PE does not affect hippocampal volume (Rubin et al., 2016). Recent research linked deficits in accurately discriminating context between threatening and safe situations to a smaller hippocampus (Negash et al., 2015), thus extinction recall in PE may be affected by it. While a few studies using other exposure-based treatments for PTSD have demonstrated increased hippocampal activation in patients with PTSD following psychotherapy (Felmingham et al., 2007; Peres et al., 2007), no study to date has demonstrated increased activation with PE (Lindauer et al., 2005; van Rooij et al., 2015;

Rubin et al., 2016). Overall, the fear and contextualization neurocircuitries (amygdala, vmPFC, ACC and the hippocampus) appear to be heavily involved in the processes of extinction learning and recall in PE. As expected, PE restores the balance in the vmPFC-amygdala loop and decreases the activation in the ACC during extinction recall. Interestingly, the functioning and volume of some of these structures may also be a prognostic indicator for PE's efficiency and a target for enhancement interventions.

Emotion regulation neurocircuitry is not a unified neurocircuitry but a set of circuits that share regions with the fear neurocircuitry (Ochsner et al., 2002; Sheynin and Liberzon, 2017) but there is some evidence to suggest that activation in and connectivity between these circuits may underlie treatment outcomes in PE. In one neuroimaging study, patients underwent an implicit (unintentional) emotion regulation task and an explicit (intentional and deliberate) emotion regulation task before receiving a course of PE (Fonzo et al., 2017). Individuals with greater vmPFC activation during the implicit emotion regulation task showed larger PTSD symptom reduction at the end of treatment (Fonzo et al., 2017). This points to the possibility that certain individuals' brains may have diminished capacity to reduce interference from an emotionally-salient cue in the environment, possibly making it more difficult to fully engage in PE. Interestingly, activation during the explicit emotion regulation task at baseline did not predict symptom change (Fonzo et al., 2017). This is consistent with EPT's emphasis on emotional engagement during exposures to facilitate extinction and inhibitory learning insofar that efforts at attenuating emotional responses during exposure are counterindicated and interfere with learning (Craske et al., 2008; Foa, 2011). Increased competence to cope with negative affect is a type of inhibitory learning speculated to be an active ingredient in PE's effectiveness (Rauch et al., 2001; Foa and Rauch, 2004; Zalta et al., 2014). These emotion regulation skills are acquired both through successful exposures and through processing that occurs following exposures. In a study examining the effect of PE on emotion regulation skills in a sample of individuals with PTSD (Jerud et al., 2016), PE was associated with clinically meaningful improvements in emotion regulation skills, but a course of sertraline had a similar effect (Jerud et al., 2016). Therefore, it is difficult to determine whether the PE effects on emotion regulation are specific to PE or more generalized to any effective PTSD intervention. One neuroimaging study examined connectivity in the "default mode network" (mPFC, parietal cortex) which has been implicated in attentional control (Fox et al., 2015) before and following a mindfulness based exposure therapy (King et al., 2016). Following exposure therapy (but not present-centered therapy), increased connectivity of the default mode network to the dlPFC was observed, and this increased connectivity was associated with decreased avoidance and hyperarousal symptoms of PTSD (King et al., 2016). This pattern of neural activation suggests increased attentional control, one of the components of emotion regulation (Thompson, 2008; Wilcox et al., 2016), following mindfulness based exposure therapy. Correlation with decreased avoidance and hyperarousal also points to greater deployment of emotion regulation skills at the behavioral level. However, no study has examined the effects of PE on the emotion regulation neurocircuitry, therefore, it is impossible to know how much of the effect is due to mindfulness training and how much is to the exposure component. As suggested previously, emotion regulation deficits may be a transdiagnostic factor contributing to the development and maintenance of various types of psychopathology (Bradley et al., 2011; Sheynin and Liberzon, 2017), therefore, increased connectivity in that region may reflect an alleviation in symptoms due to an effective treatment in general, rather than due to a mechanism specific to PE. PTSD results in a disrupted network of interconnected brain regions. The neuroimaging studies to date suggest that changes in some neurocircuitries are more unique to the putative mechanisms of PE (e.g., the fear and contextualization neurocircuitries are affected by the extinction learning and recall component of PE) while changes in others may be more generally related to mechanisms not unique to PE (e.g., inhibitory learning related to increased emotion regulation skills). Successful extinction recall in PE appears to be related to increased functional coherence between vmPFC, amygdala, and the hippocampus (Helpman et al., 2016a). Similarly, increased connectivity between areas implicated in attentional control (default mode network) and areas implicated in explicit emotion regulation (dlPFC) appears to be indicative of more effective coping with negative affect and downregulation of emotional responses in stressful situations (King et al., 2016; Fonzo et al., 2017). Therefore, while the functioning of individual brain structures is important and

clearly impacted by the active components of PE, it appears that increased and more efficient communication between various structures that regulate each other, is of greater importance in PTSD remission.

Neurobiological Enhancements of PE Treatment

The purpose of this focused review is to create a bridge between neuroscience and practice of PE therapy by examining the effects of exposure therapy on the neural circuits implicated in PTSD. Further, we aim to use the advances in neuroscientific treatment outcome research in order to propose potential enhancement to the practice of PE based on the neurocircuits that have been shown to be affected by it. The neurobiological findings to date may be applied in two ways: (1) to identify potential PE treatment enhancements in order to facilitate emotional engagement, extinction and emotion regulation/inhibitory learning; and (2) to identify individuals who may be more likely to respond to certain enhancement, in order to provide personalized treatment.

Selective serotonin reuptake inhibitors (SSRIs) are recommended in treatment guidelines as treatment for PTSD, following the evidence-based psychotherapies such as PE (Institute of Medicine, 2014; American Psychological Association, 2017). Preliminary evidence suggests that facilitation of serotonergic transmission produced by SSRIs results in increased activation in the vmPFC regions (Brady et al., 2000; Davidson et al., 2001). A recent study compared the effects of PE alone or sertraline alone on attentional inhibition (as measured by a laboratory task) in individuals with PTSD to examine the effects of each of these therapies on one of the purported main mechanism of change in treatment of PTSD, inhibitory learning (Echiverri-Cohen et al., 2016). The authors found that those who showed more symptom improvement with PE treatment showed greater improvements in inhibitory processes from pre- to post-treatment. In contrast, those who showed greater symptom reductions on sertraline made less improvement in their inhibitory processes (Echiverri-Cohen et al., 2016). This discrepancy may point to different mechanism of action of each of these treatment and support the hypothesis that SSRIs bring about more bottom-up neurochemical changes in the fear circuitry, vs. the top-down changes produced through extinction and inhibitory learning in PE. In addition, another study found emotion dysregulation was improved equally as a result of PE or sertraline in individuals with PTSD from pre- to post-treatment but the mechanisms of action of each treatment were not tested (Jerud et al., 2016). Given that SSRIs may have a suspected different path of action than PE on restoring the balance in the limbic-prefrontal system, many have speculated that combining these interventions may augment each alone or alternatively that different people may respond to each treatment. In two studies using reverse designs (one augmenting PE on SSRI non-responders and the other augmenting SSRI for PE non-responders), results supported that for partial responders to SSRI augmentation with PE was effective (Rothbaum et al., 2006). However, when SSRI was added for those who only partially or did not respond to PE, there was no added benefit (Simon et al., 2008). Identification of genetic variants associated with more robust response to SSRIs or PE in PTSD could allow for more personalized and effective treatments. Ongoing biomarkers studies of treatment response (e.g., project PROGRESS; Rauch et al., 2018) promise to inform our field in these areas over the next several years.

Glutamatergic and GABAergic neurotransmission has been implicated in fear conditioning and extinction (Riaza Bermudo-Soriano et al., 2012), and D-cycloserine (DCS), a partial agonist at the *N*-methyl-D-aspartate (NMDA), has been implicated in fear extinction through the modulation of NMDA receptors in the amygdala (Norberg et al., 2008). While preclinical studies showed promising results (Walker et al., 2002; Ledgerwood et al., 2005; Yang and Lu, 2005) when DCS was used to facilitate extinction learning in rodents, human studies using exposure therapy yielded mixed results. Of studies that examined DCS as an augmentation agent in exposure therapy, only one found improved treatment outcomes with DCS (Difede et al., 2014). Two studies found no noticeable added benefit from

supplementing exposure therapy with DCS (de Kleine et al., 2012; Rothbaum et al., 2014), while one study found poorer treatment outcomes in the group who received DCS-augmented exposure therapy compared to placebo (Litz et al., 2012). These mixed findings suggest that DCS may be an effective exposure enhancer for a certain subgroup of individuals with PTSD. For example, de Kleine et al. (2012) found that patients with high initial scores and who completed all treatment sessions actually benefited from DCS augmentation. DCS has also been shown to facilitate reconsolidation (or updating) of fear memory in animal studies (Lee et al., 2006). Therefore, it is possible that in the case of an exposure in which extinction is insufficient, fear memory is reconsolidated in a more intense form (Litz et al., 2012), basically making an unsuccessful exposure worse. To date, the majority of studies using DCS and exposure therapy for PTSD have failed to find a clear enhancing value of DCS. Studies with individuals with fear of heights as well as social anxiety found that administering DCS following a successful exposure, did indeed augment those exposures (Smits et al., 2013a,b; Tart et al., 2013). It appears that DCS may be a very specific intervention and has to be carefully tailored to individual's symptom severity and their response to IE, and more studies of moderators and mediators of DCS impact on extinction and learning are needed before recommending it as an augment to PE.

Exogenous administration of the neurosteroids DHEA(S) and pregnenolone that modulate GABA action in the brain has also been studied in PTSD interventions. Dehydroepiandrosterone (DHEA) and its sulfated metabolite (DHEAS) are endogenous neurosteroids with negative modulatory effects (GABA antagonist) on the GABAergic system (Maninger et al., 2009). DHEA(S) levels have been shown to be inversely related to depression (Barrett-Connor et al., 1999; Wong et al., 2011) and positively related to executive function (Alhaj et al., 2006; Davis et al., 2008). Allopregnanolone (ALLO), an endogenous neurosteroid, is one of the most potent GABA agonists (Lambert et al., 2003) and has been shown to have anxiolytic effects (Paul and Purdy, 1992). In a sample of healthy men, single-dose DHEA administration was associated with decreased activation in the amygdala, and increased connectivity between the amygdala and the hippocampus during emotion regulation laboratory task (Sripada et al., 2013b). Administration of pregnenolone was associated with decreased amygdala activation and with increased connectivity between prefrontal cortical regions and the amygdala during that same task (Sripada et al., 2013c). Therefore, it appears that DHEA(S) and ALLO may be involved in the emotion regulation neurocircuitry and affect communication between the amygdala and the prefrontal regions related to executive functioning. Of note, ALLO has been shown to have positive effects on pain tolerance (Scioli-Salter et al., 2016), symptoms of traumatic brain injury (Marx et al., 2016), and depression and bipolar disorder (Osuji et al., 2010; Brown et al., 2014). The same way emotion dysregulation may be a transdiagnostic indicator of emotional disorders, DHEA(S) and ALLO may have a transdiagnostic therapeutic effect, independent of the mechanisms of action of PE and therefore not specific to PTSD (Rasmusson et al., 2017).

As increased neural connectivity within and between different neurocircuits is emerging as an important mechanism of action in psycho- and pharmacotherapies, repetitive transcranial magnetic stimulation (rTMS) has been of interest as a stand-alone and add-on treatment for PTSD (Karsen et al., 2014; Yan et al., 2017). rTMS uses an electromagnetic field to non-invasively stimulate cortical neurons through repeated changes in the coil's magnetic field (George et al., 2002; George and Post, 2011) and has been approved by the Food and Drug Administration for the treatment of drug-resistant depression. The most common target for these studies has been broadly the dlPFC (Karsen et al., 2014), with its projections to the fear and contextualization circuits (i.e., the amygdala, the hippocampus, and the vmPFC). To date, several reviews and/or meta-analyses have demonstrated the effectiveness of rTMS for treatment of PTSD by targeting the right dlPFC regions (Karsen et al., 2014; Clark et al., 2015; Yan et al., 2017). Fonzo et al. (2017) found that when the right dlPFC was stimulated via rTMS, it downregulated the inhibition of the left amygdala; the magnitude of that effect was a predictor of PE response. A study that used TMS as a stand-alone treatment (no exposure therapy) found that the pre-existing connectivity between the ACC and the default mode network responsible for attentional control as well as connectivity between the amygdala and the vmPFC predicted patient's response in rTMS treatment (Philip et al., 2018). Therefore, assessing patient's brain activation patterns pre-treatment may be used as a predictor of treatment

response. More importantly, the rTMS studies to date identify neurostimulation-accessible brain regions that may serve as targets for enhancing exposure therapy either prior to or during the course of PE (Fonzo et al., 2017).

One novel candidate enhancement to PE that is purported to target the actual engagement in treatment and the quality of therapeutic alliance is a neuropeptide oxytocin (Olff et al., 2010). Oxytocin's properties of enhancing prosocial behavior, trust and warmth may be useful in facilitating extinction and inhibitory learning in exposure therapy through successful therapeutic alliance (McLaughlin et al., 2014) and decreasing dropout rates (Tuerk, 2014). One unpublished small study found that a single administration of oxytocin to individuals with PTSD decreased anxiety scores, irritability, intensity of intrusive symptoms, and increased the desire for social contact (Yatzkar and Klein, 2010). Thus far, only one pilot randomized controlled trial examined the efficacy of administering intranasal oxytocin 45 min prior to each IE in patients with PTSD who were undergoing PE (Flanagan et al., 2018). The group who received oxytocin demonstrated lower PTSD and depression symptoms during PE and had higher therapeutic alliance scores but these differences were not statistically significant, potentially due to low power (Flanagan et al., 2018). Therefore, larger RCTs are warranted to further explore the efficacy of oxytocin as a supplement to PE. Another promising novel augmentation to PE is MDMA (Thal and Lommen, 2018), a substituted amphetamine (i.e., a class of compounds based on the amphetamine structure derived by replacing one or more hydrogen atoms in the amphetamine core structure) with properties similar to mescaline, psilocybin, and other psychedelic compounds. The cognitive effects of MDMA in clinical studies have included enhanced mood, happiness, physical and mental relaxation, increased emotional responsiveness, increased openness and extraversion, and increased prosocial behaviors such as trust and feelings of closeness to other people (Harris et al., 2002; Vollenweider et al., 2002). In addition, MDMA has been demonstrated to facilitate extinction retention in mice (Young et al., 2017). The exact pharmacological mechanisms of MDMA's action are not well-understood. MDMA is known to acutely facilitate the release of serotonin and oxytocin (Dumont et al., 2009; van Wel et al., 2012), potentially contributing to decreased avoidance and greater engagement in exposure-based therapy.

To date, there have been three clinical trials examining the effectiveness of MDMA as an adjunct to psychotherapy for treating PTSD (Mithoefer et al., 2011, 2018; Oehen et al., 2013). In all three of these studies, MDMA was administered shortly before the psychotherapy session and participants demonstrated significant decreases in PTSD symptoms at the end of treatment and at follow up (Mithoefer et al., 2011, 2018; Oehen et al., 2013). Of note, psychotherapy administered in these studies was not one of the empirically supported exposure-based treatments for PTSD and was instead non-directive and focused more on experiencing than on verbal exchanges (Mithoefer, 2011). MDMA remains to be tested as an enhancement to PE. Given the proposed mechanism of action in MDMA (i.e., increases in serotonin and oxytocin, increased trust and openness to new ideas), it is possible that it would be particularly helpful to apply it during the processing portion of the session, when patients' beliefs about themselves, others, and the world are reflected and become less rigid. Alternatively, MDMA may be useful only for those patients who exhibit slow or no extinction during IE. One recent study found that, on a personality trait level, patients who received MDMA during psychotherapy exhibited an increase in the Openness trait post-treatment and at follow-up, while their Neuroticism trait remained unchanged (Wagner et al., 2017). This suggests that indeed the mechanism of effective action in the case of MDMA may be greater openness to new ideas and decreased cognitive rigidity rather than decreased negative emotionality. While promising, the efficacy and effectiveness of supplementing PE with MDMA remains an empirical question.

Future Directions and Conclusions

The mechanisms of action of PE have been of great interest in the past decade and neuroimaging studies followed suit. Currently, it is clear that various neural circuits are impacted in the course of PE but it also appears that certain patterns of neural activation and connectivity predict patient's response in PE. Generally, exposure therapy appears to have an impact on the fear and contextualization neurocircuitry by facilitating improved communication between the vmPFC, hippocampus, and the amygdala, leading to downregulation of the fear response in the amygdala (Helpman et al., 2016a). PE has been shown to decrease the activity in and the volume of the ACC. This increased coherence between these particular regions may be the mechanisms unique to PE (or exposure therapy in general) given that successful exposures lead to extinction learning and recall, thus extinguishing the overgeneralized fear response. There appears to be a more general process of emotional regulation that is of importance as well. Similarly, increased connectivity between areas implicated in attentional control (default mode network) and areas implicated in explicit emotion regulation (dlPFC) appears to be indicative of more effective coping with negative affect and downregulation of emotional responses in stressful situations (King et al., 2016; Fonzo et al., 2017). This process has been conceptualized as more transdiagnostic and less specific to PTSD or PE. Finally, it appears that neural connectivity *prior* to treatment may have profound impact on treatment response, offering some directions for future use of prognostic indicators as well as enhancements.

One future direction concerns identifying the types of patients who would most benefit from PE (and those who might not). Larger hippocampal volume has been shown to be associated with better outcomes in PE (Rubin et al., 2016). Increased connectivity between the fear and the emotion regulation neurocircuitries during emotionally salient tasks is also a predictor of treatment success in PE (Fonzo et al., 2017). Therefore, neuroimaging or electroencephalography should be used in future studies to not only corroborate these findings but also potentially establish certain cut-off benchmark for the magnitude of connectivity or the relative size of the hippocampus in optimal PE response. Given consistently positive findings regarding the effectiveness of rTMS in PTSD symptom improvement, it is a priority to continue to research this potential PE enhancement. Specifically, future studies should focus on neurobiological and psychological moderators and mediators of rTMS effects on PE response, as well as on identifying those individuals who would most benefit from rTMS. Currently, it appears that rTMS may be of particular importance for those people whose brains do not "let" them engage fully in exposure therapy, i.e., those with pre-existing decreased connectivity between the key neurocircuitries, however, that is an empirical question.

Further examination of the enhancements that promote the sense of connectedness and self-compassion (i.e., oxytocin and MDMA) is also warranted. Currently, it is unclear whether these agents affect neural circuits that are more unique to exposure therapy (such as the fear or the contextualization circuits) or the transdiagnostic emotion regulation circuitry, or another circuitry altogether. They may be promoting decreased behavioral avoidance (Dumont et al., 2009; van Wel et al., 2012) but neuroimaging studies examining these agents' effects on neurocircuitries that are of most importance in PE would clarify that conjecture. Concurrently, studies examining the effects of these agents on PE response would be helpful in clarifying whether these agents offer any added benefit above and beyond regular PE. If so, then identifying patients who would be appropriate candidates for such enhancements would be the logical next steps, especially given the heterogeneity of PTSD symptoms.

In order to tap into the change in the ability to handle and regulate negative emotions and identify individuals most likely to benefit from treatment, future studies should employ various emotion regulation tasks during neuroimaging scans pre- and post-treatment. For instance in the PROGrESS trial (Rauch et al., 2018), participants engage in an emotional faces matching task aimed at isolating the amygdala reactivity to threat (i.e., angry and fearful faces) and non-threat (i.e., happy and neutral faces; Hariri et al., 2002). Additionally, participants engage in the Emotion Regulation Task designed to measure both, the explicit emotion regulation (Gyurak et al., 2011) activation in the prefrontal cortical regions including dlPFC and vmPFC, as well as the implicit emotion regulation manifested by the activation in

vmPFC and the amygdala (Costafreda et al., 2008; Buhle et al., 2014). Finally, the implicit emotional regulation processes (Gyurak et al., 2011) are also measured using attentional control with emotional faces task (SEAT; Sripada et al., 2013a). Administration of such tasks while undergoing neuroimaging pre- and post-PE is essential in identifying not only connectivity changes resulting from treatment in the fear, contextualization, and emotion regulation neurocircuitries but also pre-existing connectivity patterns that may be indicative of individuals who might require enhancements in order to fully benefit from PE. It would be useful to develop and utilize neuropsychological measures of hippocampus activity in order to evaluate PE's effect on its function. Additionally, salivary and plasma concentrations of DHEA(S) and ALLO have been shown to be associated with increased communication between prefrontal cortical regions and amygdala (Sripada et al., 2013c). Therefore, establishing benchmarks for the extent of connectivity related to certain concentrations of DHEA(S) and ALLO, and measuring these concentrations pre and during PE may be a potentially less burdensome method of identifying increased communication between the relevant brain structures.

While most of our focus has been on improving response to PE, retention, as previously mentioned, is a key area for improvement across PTSD interventions (Tuerk, 2014). Neurobiological advances can also provide insights that drive better retention through more personalized, more efficient, and more effective care. Dropout rates across populations and treatment setting are approximately 30% or more (Bradley et al., 2005; Eftekhari et al., 2013), which is not surprising given that PTSD is characterized by behavioral avoidance. Therefore, further exploring the effectiveness and the mechanisms of action of oxytocin and MDMA as enhancements to PE is warranted in an effort to improve retention in PE. For instance, for those who have higher potential for dropout, administration of intranasal oxytocin may be particularly effective in retaining them in treatment through increased connectedness with their PE provider. Identifying neurobiological biomarkers of those who are at risk of dropping out would aid in clinical decision-making process regarding patients who are good candidates for such an enhancement.

Conclusions

In this focused review, we reviewed the neurobiological mechanisms underlying the effectiveness of PE, an empirically-supported efficacious treatment for PTSD, and the enhancements that can be applied to increase treatment response and retention. One of the proposed mechanisms of action in PE is exposure which facilitates extinction of a fear response and new adaptive learning. Neurobiologically, PE and successful extinction recall has been associated with increased neural connectivity between vmPFC, amygdala and the hippocampus. Increased connectivity in regions implicated in emotion regulation has also been shown to result from PE although it appears that this change in activation is more transdiagnostic and not unique to PE. Since neural connectivity and communication seem to be at the heart of symptom alleviation in PTSD, treatment enhancements that promote such connectivity, particularly rTMS, offer the most appropriate targets for future research into effectiveness of PE. Further research into the effects of oxytocin and MDMA on treatment response and retention is also warranted. Finally, establishing neurobiological benchmarks for identifying individuals who are less likely to benefit from treatment would be an exciting development to help guide clinical decision-making as to who should receive enhancements to PE.

Author Contributions

MS involved in planning, lead author, primary writer and editor. LM involved in planning, led the literature review, wrote sections and edited. SR senior author involved in planning, review, writing and editing.

Conflict of Interest Statement

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

The reviewer JA and handling Editor declared their shared affiliation at the time of the review.

Footnotes

1. ^ We have chosen to use extinction rather than the term habituation used by Foa and Kozak (1986) based on more recent theorists and neuroscience research that aligns more with extinction processes (see Rauch and Liberzon, 2016 for review).

References

Alhaj, H. A., Massey, A. E., and McAllister-Williams, R. H. (2006). Effects of DHEA administration on episodic memory, cortisol and mood in healthy young men: a double-blind, placebo-controlled study. *Psychopharmacology* 188, 541–551. doi: 10.1007/s00213-005-0136-y

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=16231168) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=16231168) | [CrossRef Full Text](https://doi.org/10.1007/s00213-005-0136-y) (https://doi.org/10.1007/s00213-005-0136-y) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Effects+of+DHEA+administration+on+episodic+memory,+cortisol+and+mood+in+healthy+young+men%3A+a+double-blind,+placebo-controlled+study&author=Alhaj+H.+A.&author=Massey+A.+E.&author=McAllister-Williams+R.+H.&publication_year=2006&journal=Psychopharmacology&volume=188&pages=541-551) (http://scholar.google.com/scholar_lookup?title=Effects+of+DHEA+administration+on+episodic+memory,+cortisol+and+mood+in+healthy+young+men%3A+a+double-blind,+placebo-controlled+study&author=Alhaj+H.+A.&author=Massey+A.+E.&author=McAllister-Williams+R.+H.&publication_year=2006&journal=Psychopharmacology&volume=188&pages=541-551)

American Psychiatric Association. (2013). *The Diagnostic and Statistical Manual of Mental Disorders: DSM 5*. Washington, D.C.: American Psychiatric Publishing.

[Google Scholar](http://scholar.google.com/scholar_lookup?title=The+Diagnostic+and+Statistical+Manual+of+Mental+Disorders%3A+DSM+5&American+Psychiatric+Association&publication_year=2013) (http://scholar.google.com/scholar_lookup?title=The+Diagnostic+and+Statistical+Manual+of+Mental+Disorders%3A+DSM+5&American+Psychiatric+Association&publication_year=2013)

American Psychological Association. (2017). *Clinical Practice Guideline for the Treatment of Post-traumatic Stress Disorder (PTSD) in Adults*. Available online at: www.apa.org/ptsd-guideline/ptsd.pdf (https://www.apa.org/ptsd-guideline/ptsd.pdf)

Aupperle, R. L., Melrose, A. J., and Paulus, M. P. (2012). Executive function and PTSD: disengaging from trauma. *Neuropharmacology* 62, 686–694. doi: 10.1016/j.neuropharm.2011.02.008

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=21349277) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=21349277) | [CrossRef Full Text](https://doi.org/10.1016/j.neuropharm.2011.02.008) (https://doi.org/10.1016/j.neuropharm.2011.02.008) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Executive+function+and+PTSD%3A+disengaging+from+trauma&author=Aupperle+R.+L.&author=Melrose+A.+J.&author=Paulus+M.+P.&publication_year=2012&journal=Neuropharmacology&volume=62&pages=686-694) (http://scholar.google.com/scholar_lookup?title=Executive+function+and+PTSD%3A+disengaging+from+trauma&author=Aupperle+R.+L.&author=Melrose+A.+J.&author=Paulus+M.+P.&publication_year=2012&journal=Neuropharmacology&volume=62&pages=686-694)

Barrett-Connor, E., von Mühlen, D., Laughlin, G. A., and Kripke, A. (1999). Endogenous levels of dehydroepiandrosterone sulfate, but not other sex hormones, are associated with depressed mood in older women: the rancho bernardo study. *J. Am. Geriatr. Soc.* 47, 685–691. doi: 10.1111/j.1532-5415.1999.tb01590.x

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=10366167) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=10366167) | [CrossRef Full Text](https://doi.org/10.1111/j.1532-5415.1999.tb01590.x) (https://doi.org/10.1111/j.1532-5415.1999.tb01590.x) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Endogenous+levels+of+dehydroepiandrosterone+sul-) (http://scholar.google.com/scholar_lookup?title=Endogenous+levels+of+dehydroepiandrosterone+sul-

fate,+but+not+other+sex+hormones,+are+associated+with+depressed+mood+in+older+women%3A+the+rancho+bernardo+study&author=Barrett-Connor+E.&author=von+Mühlen+D.&author=Laughlin+G.+A.&author=Kripke+A.&publication_year=1999&journal=J.+Am.+Geriatr.+Soc.&volume=47&pages=685-691)

Blechert, J., Michael, T., Vriends, N., Margraf, J., and Wilhelm, F. H. (2007). Fear conditioning in posttraumatic stress disorder: evidence for delayed extinction of autonomic, experiential and behavioural responses. *Behav. Res. Ther.* 45, 2019–2033. doi: 10.1016/j.brat.2007.02.012

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=17442266) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=17442266) | [CrossRef Full Text](https://doi.org/10.1016/j.brat.2007.02.012) (https://doi.org/10.1016/j.brat.2007.02.012) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Fear+conditioning+in+posttraumatic+stress+disorder%3A+evidence+for+delayed+extinction+of+autonomic,+experiential+and+behavioural+responses&author=Blechert+J.&author=Michael+T.&author=Vriends+N.&author=Margraf+J.&author=Wilhelm+F.+H.&publication_year=2007&journal=Behav.+Res.+Ther.&volume=45&pages=2019-2033) (http://scholar.google.com/scholar_lookup?title=Fear+conditioning+in+posttraumatic+stress+disorder%3A+evidence+for+delayed+extinction+of+autonomic,+experiential+and+behavioural+responses&author=Blechert+J.&author=Michael+T.&author=Vriends+N.&author=Margraf+J.&author=Wilhelm+F.+H.&publication_year=2007&journal=Behav.+Res.+Ther.&volume=45&pages=2019-2033)

Bradley, B., DeFife, J. A., Guarnaccia, C., Phifer, J., Fani, N., Ressler, K. J., et al. (2011). Emotion dysregulation and negative affect: association with psychiatric symptoms. *J. Clin. Psychiatry* 72, 685–691. doi: 10.4088/jcp.10m06409blu

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=21658350) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=21658350) | [CrossRef Full Text](https://doi.org/10.4088/jcp.10m06409blu) (https://doi.org/10.4088/jcp.10m06409blu) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Emotion+dysregulation+and+negative+affect%3A+association+with+psychiatric+symptoms&author=Bradley+B.&author=DeFife+J.+A.&author=Guarnaccia+C.&author=Phifer+J.&author=Fani+N.&author=Ressler+K.+J.&publication_year=2011&journal=J.+Clin.+Psychiatry&volume=72&pages=685-691) (http://scholar.google.com/scholar_lookup?title=Emotion+dysregulation+and+negative+affect%3A+association+with+psychiatric+symptoms&author=Bradley+B.&author=DeFife+J.+A.&author=Guarnaccia+C.&author=Phifer+J.&author=Fani+N.&author=Ressler+K.+J.&publication_year=2011&journal=J.+Clin.+Psychiatry&volume=72&pages=685-691)

Bradley, R., Greene, J., Russ, E., Dutra, L., and Westen, D. (2005). A multidimensional meta-analysis of psychotherapy for PTSD. *Am. J. Psychiatry* 162, 214–227. doi: 10.1176/appi.ajp.162.2.214

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=15677582) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=15677582) | [CrossRef Full Text](https://doi.org/10.1176/appi.ajp.162.2.214) (https://doi.org/10.1176/appi.ajp.162.2.214) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=A+multidimensional+meta-analysis+of+psychotherapy+for+PTSD&author=Bradley+R.&author=Greene+J.&author=Russ+E.&author=Dutra+L.&author=Westen+D.&publication_year=2005&journal=Am.+J.+Psychiatry&volume=162&pages=214-227) (http://scholar.google.com/scholar_lookup?title=A+multidimensional+meta-analysis+of+psychotherapy+for+PTSD&author=Bradley+R.&author=Greene+J.&author=Russ+E.&author=Dutra+L.&author=Westen+D.&publication_year=2005&journal=Am.+J.+Psychiatry&volume=162&pages=214-227)

Brady, K., Pearlstein, T., Asnis, G. M., Baker, D., Rothbaum, B., Sikes, C. R., et al. (2000). Efficacy and safety of sertraline treatment of posttraumatic stress disorder. *JAMA* 283, 1837–1844. doi: 10.1001/jama.283.14.1837

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=10770145) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=10770145) | [CrossRef Full Text](https://doi.org/10.1001/jama.283.14.1837) (https://doi.org/10.1001/jama.283.14.1837) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Efficacy+and+safety+of+sertraline+treatment+of+posttraumatic+stress+disorder&author=Brady+K.&author=Pearlstein+T.&author=Asnis+G.+M.&author=Baker+D.&author=Rothbaum+B.&author=Sikes+C.+R.&publication_year=2000&journal=JAMA&volume=283&pages=1837-1844) (http://scholar.google.com/scholar_lookup?title=Efficacy+and+safety+of+sertraline+treatment+of+posttraumatic+stress+disorder&author=Brady+K.&author=Pearlstein+T.&author=Asnis+G.+M.&author=Baker+D.&author=Rothbaum+B.&author=Sikes+C.+R.&publication_year=2000&journal=JAMA&volume=283&pages=1837-1844)

Brown, E. S., Park, J., Marx, C. E., Hynan, L. S., Gardner, C., Davila, D., et al. (2014). A randomized, double-blind, placebo-controlled trial of pregnenolone for bipolar depression. *Neuropsychopharmacology* 39, 2867–2873. doi: 10.1038/npp.2014.138



About us ▾

All journals

All articles

Submit your research

Q Search

👤 Login

Frontiers in Behavioral Neur...

Sections ▾

Articles

Research Topics

Editorial Board

About journal ▾

yy&volume=33&pages=2007-2010)

Buhle, J. T., Silvers, J. A., Wager, T. D., Lopez, R., Onyemekwu, C., Kober, H., et al. (2014). Cognitive reappraisal of emotion: a meta-analysis of human neuroimaging studies. *Cereb. Cortex* 24, 2981–2990. doi: 10.1093/cercor/bht154

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23765157) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23765157) | [CrossRef Full Text](https://doi.org/10.1093/cercor/bht154) (https://doi.org/10.1093/cercor/bht154) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Cognitive+reappraisal+of+emotion%3A+a+meta-analysis+of+human+neuroimaging+studies&author=Buhle+J.+T.&author=Silvers+J.+A.&author=Wager+T.+D.&author=Lopez+R.&author=Onyemekwu+C.&author=Kober+H.&publication_year=2014&journal=Cereb.+Cortex&volume=24&pages=2981-2990) (http://scholar.google.com/scholar_lookup?title=Cognitive+reappraisal+of+emotion%3A+a+meta-analysis+of+human+neuroimaging+studies&author=Buhle+J.+T.&author=Silvers+J.+A.&author=Wager+T.+D.&author=Lopez+R.&author=Onyemekwu+C.&author=Kober+H.&publication_year=2014&journal=Cereb.+Cortex&volume=24&pages=2981-2990)

Clark, C., Cole, J., Winter, C., Williams, K., and Grammer, G. (2015). A review of transcranial magnetic stimulation as a treatment for post-traumatic stress disorder. *Curr. Psychiatry Rep.* 17:83. doi: 10.1007/s11920-015-0621-x

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=26324821) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=26324821) | [CrossRef Full Text](https://doi.org/10.1007/s11920-015-0621-x) (https://doi.org/10.1007/s11920-015-0621-x) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=A+review+of+transcranial+magnetic+stimulation+as+a+treatment+for+post-traumatic+stress+disorder&author=Clark+C.&author=Cole+J.&author=Winter+C.&author=Williams+K.&author=Grammer+G.&publication_year=2015&journal=Curr.+Psychiatry+Rep.&volume=17&pages=83) (http://scholar.google.com/scholar_lookup?title=A+review+of+transcranial+magnetic+stimulation+as+a+treatment+for+post-traumatic+stress+disorder&author=Clark+C.&author=Cole+J.&author=Winter+C.&author=Williams+K.&author=Grammer+G.&publication_year=2015&journal=Curr.+Psychiatry+Rep.&volume=17&pages=83)

Costafreda, S. G., Brammer, M. J., David, A. S., and Fu, C. H. Y. (2008). Predictors of amygdala activation during the processing of emotional stimuli: a meta-analysis of 385 PET and fMRI studies. *Brain Res. Rev.* 58, 57–70. doi: 10.1016/j.brainresrev.2007.10.012

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=18076995) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=18076995) | [CrossRef Full Text](https://doi.org/10.1016/j.brainresrev.2007.10.012) (https://doi.org/10.1016/j.brainresrev.2007.10.012) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Predictors+of+amygdala+activation+during+the+processing+of+emotional+stimuli%3A+a+meta-analysis+of+385+PET+and+fMRI+studies&author=Costafreda+S.+G.&author=Brammer+M.+J.&author=David+A.+S.&author=Fu+C.+H.+Y.&publication_year=2008&journal=Brain+Res.+Rev.&volume=58&pages=57-70) (http://scholar.google.com/scholar_lookup?title=Predictors+of+amygdala+activation+during+the+processing+of+emotional+stimuli%3A+a+meta-analysis+of+385+PET+and+fMRI+studies&author=Costafreda+S.+G.&author=Brammer+M.+J.&author=David+A.+S.&author=Fu+C.+H.+Y.&publication_year=2008&journal=Brain+Res.+Rev.&volume=58&pages=57-70)

Craske, M. G., Kircanski, K., Zelikowsky, M., Mystkowski, J., Chowdhury, N., and Baker, A. (2008). Optimizing inhibitory learning during exposure therapy. *Behav. Res. Ther.* 46, 5–27. doi: 10.1016/j.brat.2007.10.003

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=18005936) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=18005936) | [CrossRef Full Text](https://doi.org/10.1016/j.brat.2007.10.003) (https://doi.org/10.1016/j.brat.2007.10.003) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Optimizing+inhibitory+learning+during+exposure+therapy&author=Craske+M.+G.&author=Kircanski+K.&author=Zelikowsky+M.&author=Mystkowski+J.&author=Chowdhury+N.&author=Baker+A.&publication_year=2008&journal=Behav.+Res.+Ther.&volume=46&pages=5-27) (http://scholar.google.com/scholar_lookup?title=Optimizing+inhibitory+learning+during+exposure+therapy&author=Craske+M.+G.&author=Kircanski+K.&author=Zelikowsky+M.&author=Mystkowski+J.&author=Chowdhury+N.&author=Baker+A.&publication_year=2008&journal=Behav.+Res.+Ther.&volume=46&pages=5-27)

Dannowski, U., Stuhrmann, A., Beutelmann, V., Zwanzger, P., Lenzen, T., Grotegerd, D., et al. (2012). Limbic scars: long-term consequences of childhood maltreatment revealed by functional and structural magnetic resonance imaging. *Biol. Psychiatry* 71, 286–293. doi: 10.1016/j.biopsych.2011.10.021

We use cookies

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23112927) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23112927) | [CrossRef Full Text](https://doi.org/10.1016/j.biopsych.2011.10.021) (https://doi.org/10.1016/j.biopsych.2011.10.021) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Limbic+scars%3A+long-term+consequences+of+childhood+maltreatment+revealed+by+functional+and+structural+magnetic+resonance+imaging&author=Dannowski+U.&author=Stuhrmann+A.&author=Beutelmann+V.&author=Zwanzger+P.&author=Lenzen+T.&author=Grotegerd+D.&publication_year=2012&journal=Biol.+Psychiatry&volume=71&pages=286-293) (http://scholar.google.com/scholar_lookup?title=Limbic+scars%3A+long-term+consequences+of+childhood+maltreatment+revealed+by+functional+and+structural+magnetic+resonance+imaging&author=Dannowski+U.&author=Stuhrmann+A.&author=Beutelmann+V.&author=Zwanzger+P.&author=Lenzen+T.&author=Grotegerd+D.&publication_year=2012&journal=Biol.+Psychiatry&volume=71&pages=286-293)

Our website uses cookies that are necessary for the operation and to improve your experience. You can review and control your cookies by clicking on "Accept Cookies" or on "Cookies Settings".

[Download \(articles/10.3389/fnbeh.2018.00281/pdf\)](#)

Davidson, J. R. T., Rothbaum, B. O., van der Kolk, B. A., Sikes, C. R., and Farfel, G. M. (2001). Multicenter, double-blind comparison of sertraline and placebo in the treatment of posttraumatic stress disorder. *Arch. Gen. Psychiatry* 58, 485–492. doi: 10.1001/archpsyc.58.5.485

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=11343529) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=11343529) | [CrossRef Full Text](https://doi.org/10.1001/archpsyc.58.5.485) (https://doi.org/10.1001/archpsyc.58.5.485) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Multicenter,+double-blind+comparison+of+sertraline+and+placebo+in+the+treatment+of+posttraumatic+stress+disorder&author=Davidson+J.+R.+T.&author=Rothbaum+B.+O.&author=van+der+Kolk+B.+A.&author=Sikes+C.+R.&author=Farfel+G.+M.&publication_year=2001&journal=Arch.+Gen.+Psychiatry&volume=58&pages=485-492) (http://scholar.google.com/scholar_lookup?title=Multicenter,+double-blind+comparison+of+sertraline+and+placebo+in+the+treatment+of+posttraumatic+stress+disorder&author=Davidson+J.+R.+T.&author=Rothbaum+B.+O.&author=van+der+Kolk+B.+A.&author=Sikes+C.+R.&author=Farfel+G.+M.&publication_year=2001&journal=Arch.+Gen.+Psychiatry&volume=58&pages=485-492)

Davis, S. R., Shah, S. M., McKenzie, D. P., Kulkarni, J., Davison, S. L., and Bell, R. J. (2008). Dehydroepiandrosterone sulfate levels are associated with more favorable cognitive function in women. *J. Clin. Endocrinol. Metab.* 93, 801–808. doi: 10.1210/jc.2007-2128

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=18073302) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=18073302) | [CrossRef Full Text](https://doi.org/10.1210/jc.2007-2128) (https://doi.org/10.1210/jc.2007-2128) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Dehydroepiandrosterone+sulfate+levels+are+associated+with+more+favorable+cognitive+function+in+women&author=Davis+S.+R.&author=Shah+S.+M.&author=McKenzie+D.+P.&author=Kulkarni+J.&author=Davison+S.+L.&author=Bell+R.+J.&publication_year=2008&journal=J.+Clin.+Endocrinol.+Metab.&volume=93&pages=801-808) (http://scholar.google.com/scholar_lookup?title=Dehydroepiandrosterone+sulfate+levels+are+associated+with+more+favorable+cognitive+function+in+women&author=Davis+S.+R.&author=Shah+S.+M.&author=McKenzie+D.+P.&author=Kulkarni+J.&author=Davison+S.+L.&author=Bell+R.+J.&publication_year=2008&journal=J.+Clin.+Endocrinol.+Metab.&volume=93&pages=801-808)

de Kleine, R. A., Hendriks, G.-J., Kusters, W. J. C., Broekman, T. G., and van Minnen, A. (2012). A randomized placebo-controlled trial of D-cycloserine to enhance exposure therapy for posttraumatic stress disorder. *Biological Psychiatry* 71, 962–968. doi: 10.1016/j.biopsych.2012.02.033

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=22480663) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=22480663) | [CrossRef Full Text](https://doi.org/10.1016/j.biopsych.2012.02.033) (https://doi.org/10.1016/j.biopsych.2012.02.033) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=A+randomized+placebo-controlled+trial+of+D-cycloserine+to+enhance+exposure+therapy+for+posttraumatic+stress+disorder&author=de+Kleine+R.+A.&author=Hendriks+G.-J.&author=Kusters+W.+J.+C.&author=Broekman+T.+G.&author=van+Minnen+A.&publication_year=2012&journal=Biological+Psychiatry&volume=71&pages=962-968) (http://scholar.google.com/scholar_lookup?title=A+randomized+placebo-controlled+trial+of+D-cycloserine+to+enhance+exposure+therapy+for+posttraumatic+stress+disorder&author=de+Kleine+R.+A.&author=Hendriks+G.-J.&author=Kusters+W.+J.+C.&author=Broekman+T.+G.&author=van+Minnen+A.&publication_year=2012&journal=Biological+Psychiatry&volume=71&pages=962-968)

Department of Veterans Affairs & Department of Defense (VA & DoD). (2017). *VA/DoD Clinical Practice Guideline for Management of Post-Traumatic Stress*. Washington, D.C.: CreateSpace Independent Publishing Platform.

Difede, J., Cukor, J., Wyka, K., Olden, M., Hoffman, H., Lee, F. S., et al. (2014). D-cycloserine augmentation of exposure therapy for post-traumatic stress disorder: a pilot randomized clinical trial. *Neuropsychopharmacology* 39, 1052–1058. doi: 10.1038/npp.2013.317

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24217129) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24217129) | [CrossRef Full Text](https://doi.org/10.1038/npp.2013.317) (https://doi.org/10.1038/npp.2013.317) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=D-cycloserine+augmentation+of+exposure+therapy+for+post-traumatic+stress+disorder%3A+a+pilot+randomized+clinical+trial&author=Difede+J.&author=Cukor+J.&author=Wyka+K.&author=Olden+M.&author=Hoffman+H.&author=Lee+F.+S.&publication_year=2014&journal=Neuropsychopharmacology&volume=39&pages=1052-1058) (http://scholar.google.com/scholar_lookup?title=D-cycloserine+augmentation+of+exposure+therapy+for+post-traumatic+stress+disorder%3A+a+pilot+randomized+clinical+trial&author=Difede+J.&author=Cukor+J.&author=Wyka+K.&author=Olden+M.&author=Hoffman+H.&author=Lee+F.+S.&publication_year=2014&journal=Neuropsychopharmacology&volume=39&pages=1052-1058)

Dumont, G. J. H., Sweep, F. C. G. J., van der Steen, R., Hermsen, R., Donders, A. R. T., Touw, D. J., et al. (2009). Increased oxytocin concentrations and prosocial feelings in humans after ecstasy (3,4-methylenedioxymethamphetamine) administration. *Soc. Neurosci.* 4, 359–366. doi: 10.1080/17470910802649470

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=19562632) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=19562632) | [CrossRef Full Text](https://doi.org/10.1080/17470910802649470) (https://doi.org/10.1080/17470910802649470) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Increased+oxytocin+concentrations+and+prosocial+feelings+in+humans+after+ecstasy+(3,4-methylenedioxymethamphetamine)+administration&author=Dumont+G.+J.+H.&author=Sweep+F.+C.+G.+J.&author=van+der+Steen+R.&author=Hermsen+R.&author=Donders+A.+R.+T.&author=Touw+D.+J.&publication_year=2009&journal=Soc.+Neurosci.&volume=4&pages=359-366) (http://scholar.google.com/scholar_lookup?title=Increased+oxytocin+concentrations+and+prosocial+feelings+in+humans+after+ecstasy+(3,4-methylenedioxymethamphetamine)+administration&author=Dumont+G.+J.+H.&author=Sweep+F.+C.+G.+J.&author=van+der+Steen+R.&author=Hermsen+R.&author=Donders+A.+R.+T.&author=Touw+D.+J.&publication_year=2009&journal=Soc.+Neurosci.&volume=4&pages=359-366)

Duval, E. R., Javanbakht, A., and Liberzon, I. (2015). Neural circuits in anxiety and stress disorders: a focused review. *Ther. Clin. Risk Manag.* 11, 115–126. doi: 10.2147/tcrm.s48528

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=25670901) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=25670901) | [CrossRef Full Text](https://doi.org/10.2147/tcrm.s48528) (https://doi.org/10.2147/tcrm.s48528) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Neural+circuits+in+anxiety+and+stress+disorders%3A+a+focused+review&author=Duval+E.+R.&author=Javanbakht+A.&author=Liberzon+I.&publication_year=2015&journal=Ther.+Clin.+Risk+Manag.&volume=11&pages=115-126) (http://scholar.google.com/scholar_lookup?title=Neural+circuits+in+anxiety+and+stress+disorders%3A+a+focused+review&author=Duval+E.+R.&author=Javanbakht+A.&author=Liberzon+I.&publication_year=2015&journal=Ther.+Clin.+Risk+Manag.&volume=11&pages=115-126)

Echiverri-Cohen, A., Zoellner, L. A., Gallop, R., Feeny, N., Jaeger, J., and Bedard-Gilligan, M. (2016). Changes in temporal attention inhibition following prolonged exposure and sertraline in the treatment of PTSD. *J. Consult. Clin. Psychol.* 84, 415–426. doi: 10.1037/ccp0000080

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=26900894) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=26900894) | [CrossRef Full Text](https://doi.org/10.1037/ccp0000080) (https://doi.org/10.1037/ccp0000080) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Changes+in+temporal+attention+inhibition+following+prolonged+exposure+and+sertraline+in+the+treatment+of+PTSD&author=Echiverri-Cohen+A.&author=Zoellner+L.+A.&author=Gallop+R.&author=Feeny+N.&author=Jaeger+J.&author=Bedard-Gilligan+M.&publication_year=2016&journal=J.+Consult.+Clin.+Psychol.&volume=84&pages=415-426) (http://scholar.google.com/scholar_lookup?title=Changes+in+temporal+attention+inhibition+following+prolonged+exposure+and+sertraline+in+the+treatment+of+PTSD&author=Echiverri-Cohen+A.&author=Zoellner+L.+A.&author=Gallop+R.&author=Feeny+N.&author=Jaeger+J.&author=Bedard-Gilligan+M.&publication_year=2016&journal=J.+Consult.+Clin.+Psychol.&volume=84&pages=415-426)

Eftekhari, A., Ruzek, J. I., Crowley, J. J., Rosen, C. S., Greenbaum, M. A., and Karlin, B. E. (2013). Effectiveness of national implementation of prolonged exposure therapy in veterans affairs care. *JAMA Psychiatry* 70, 949–955. doi: 10.1001/jamapsychiatry.2013.36

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23863892) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23863892) | [CrossRef Full Text](https://doi.org/10.1001/jamapsychiatry.2013.36) (https://doi.org/10.1001/jamapsychiatry.2013.36) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Effectiveness+of+national+implementation+of+prolonged+exposure+therapy+in+veterans+affairs+care&author=Eftekhari+A.&author=Ruzek+J.+I.&author=Crowley+J.+J.&author=Rosen+C.+S.&author=Greenbaum+M.+A.&author=Karlin+B.+E.&publication_year=2013&journal=JAMA+Psychiatry&volume=70&pages=949-955) (http://scholar.google.com/scholar_lookup?title=Effectiveness+of+national+implementation+of+prolonged+exposure+therapy+in+veterans+affairs+care&author=Eftekhari+A.&author=Ruzek+J.+I.&author=Crowley+J.+J.&author=Rosen+C.+S.&author=Greenbaum+M.+A.&author=Karlin+B.+E.&publication_year=2013&journal=JAMA+Psychiatry&volume=70&pages=949-955)

Etkin, A., Büchel, C., and Gross, J. J. (2015). The neural bases of emotion regulation. *Nat. Rev. Neurosci.* 16, 693–700. doi: 10.1038/nrn4044

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=26481098) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=26481098) | [CrossRef Full Text](https://doi.org/10.1038/nrn4044) (https://doi.org/10.1038/nrn4044) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=The+neural+bases+of+emotion+regulation&author=Etkin+A.&author=Büchel+C.&author=Gross+J.+J.&publication_year=2015&journal=Nat.+Rev.+Neurosci.&volume=16&pages=693-700) (http://scholar.google.com/scholar_lookup?title=The+neural+bases+of+emotion+regulation&author=Etkin+A.&author=Büchel+C.&author=Gross+J.+J.&publication_year=2015&journal=Nat.+Rev.+Neurosci.&volume=16&pages=693-700)

Felmington, K., Kemp, A., Williams, L., Das, P., Hughes, G., Peduto, A., et al. (2007). Changes in anterior cingulate and amygdala after cognitive behavior therapy of posttraumatic stress disorder. *Psychol. Sci.* 18, 127–129. doi: 10.1111/j.1467-9280.2007.01860.x

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=17425531) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=17425531) | [CrossRef Full Text](https://doi.org/10.1111/j.1467-9280.2007.01860.x) (https://doi.org/10.1111/j.1467-9280.2007.01860.x) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Changes+in+anterior+cingulate+and+amygdala+after+cognitive+behavior+therapy+of+posttraumatic+stress+disorder&author=Felmington+K.&author=Kemp+A.&author=Williams+L.&author=Das+P.&author=Hughes+G.&author=Peduto+A.&publication_year=2007&journal=Psychol.+Sci.&volume=18&pages=127-129) (http://scholar.google.com/scholar_lookup?title=Changes+in+anterior+cingulate+and+amygdala+after+cognitive+behavior+therapy+of+posttraumatic+stress+disorder&author=Felmington+K.&author=Kemp+A.&author=Williams+L.&author=Das+P.&author=Hughes+G.&author=Peduto+A.&publication_year=2007&journal=Psychol.+Sci.&volume=18&pages=127-129)

Flanagan, J. C., Sippel, L. M., Wahlquist, A., Moran-Santa Maria, M. M., and Back, S. E. (2018). Augmenting prolonged exposure therapy for PTSD with intranasal oxytocin: a randomized, placebo-controlled pilot trial. *J. Psychiatr. Res.* 98, 64–69. doi: 10.1016/j.jpsychires.2017.12.014

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=29294429) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=29294429) | [CrossRef Full Text](https://doi.org/10.1016/j.jpsychires.2017.12.014) (https://doi.org/10.1016/j.jpsychires.2017.12.014) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Augmenting+prolonged+exposure+therapy+for+PTSD+with+intranasal+oxytocin%3A+a+randomized,+placebo-controlled+pilot+trial&author=Flanagan+J.+C.&author=Sippel+L.+M.&author=Wahlquist+A.&author=Moran-Santa+Maria+M.+M.&author=Back+S.+E.&publication_year=2018&journal=J.+Psychiatr.+Res.&volume=98&pages=64-69) (http://scholar.google.com/scholar_lookup?title=Augmenting+prolonged+exposure+therapy+for+PTSD+with+intranasal+oxytocin%3A+a+randomized,+placebo-controlled+pilot+trial&author=Flanagan+J.+C.&author=Sippel+L.+M.&author=Wahlquist+A.&author=Moran-Santa+Maria+M.+M.&author=Back+S.+E.&publication_year=2018&journal=J.+Psychiatr.+Res.&volume=98&pages=64-69)

Foa, E. B. (2011). Prolonged exposure therapy: past, present and future. *Depress. Anxiety* 28, 1043–1047. doi: 10.1002/da.20907

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=22134957) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=22134957) | [CrossRef Full Text](https://doi.org/10.1002/da.20907) (https://doi.org/10.1002/da.20907) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Prolonged+exposure+therapy%3A+past,+present+and+future&author=Foa+E.+B.&publication_year=2011&journal=Depress.+Anxiety&volume=28&pages=1043-1047) (http://scholar.google.com/scholar_lookup?title=Prolonged+exposure+therapy%3A+past,+present+and+future&author=Foa+E.+B.&publication_year=2011&journal=Depress.+Anxiety&volume=28&pages=1043-1047)

Foa, E. B., and Kozak, M. J. (1986). Emotional processing of fear: exposure to corrective information. *Psychol. Bull.* 99, 20–35. doi: 10.1037/0033-2909.99.1.20

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=2871574) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=2871574>) | [CrossRef Full Text](https://doi.org/10.1037/0033-2909.99.1.20) (<https://doi.org/10.1037/0033-2909.99.1.20>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Emotional+processing+of+fear%3A+exposure+to+corrective+information&author=Foa+E.+B.&author=Kozak+M.+J.&publication_year=1986&journal=Psychol.+Bull.&volume=99&pages=20-35) (http://scholar.google.com/scholar_lookup?title=Emotional+processing+of+fear%3A+exposure+to+corrective+information&author=Foa+E.+B.&author=Kozak+M.+J.&publication_year=1986&journal=Psychol.+Bull.&volume=99&pages=20-35)

Foa, E. B., and Rauch, S. A. M. (2004). Cognitive changes during prolonged exposure versus prolonged exposure plus cognitive restructuring in female assault survivors with posttraumatic stress disorder. *J. Consult. Clin. Psychol.* 72, 879–884. doi: 10.1037/0022-006x.72.5.879

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=15482045) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=15482045>) | [CrossRef Full Text](https://doi.org/10.1037/0022-006x.72.5.879) (<https://doi.org/10.1037/0022-006x.72.5.879>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Cognitive+changes+during+prolonged+exposure+versus+prolonged+exposure+plus+cognitive+restructuring+in+female+assault+survivors+with+posttraumatic+stress+disorder&author=Foa+E.+B.&author=Rauch+S.+A.+M.&publication_year=2004&journal=J.+Consult.+Clin.+Psychol.&volume=72&pages=879-884) (http://scholar.google.com/scholar_lookup?title=Cognitive+changes+during+prolonged+exposure+versus+prolonged+exposure+plus+cognitive+restructuring+in+female+assault+survivors+with+posttraumatic+stress+disorder&author=Foa+E.+B.&author=Rauch+S.+A.+M.&publication_year=2004&journal=J.+Consult.+Clin.+Psychol.&volume=72&pages=879-884)

Foa, E. B., Rothbaum, B. O., Riggs, D. S., and Murdock, T. B. (1991). Treatment of posttraumatic stress disorder in rape victims: a comparison between cognitive-behavioral procedures and counseling. *J. Consult. Clin. Psychol.* 59, 715–723. doi: 10.1037//0022-006x.59.5.715

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=1955605) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=1955605>) | [CrossRef Full Text](https://doi.org/10.1037//0022-006x.59.5.715) (<https://doi.org/10.1037//0022-006x.59.5.715>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Treatment+of+posttraumatic+stress+disorder+in+rape+victims%3A+a+comparison+between+cognitive-behavioral+procedures+and+counseling&author=Foa+E.+B.&author=Rothbaum+B.+O.&author=Riggs+D.+S.&author=Murdock+T.+B.&publication_year=1991&journal=J.+Consult.+Clin.+Psychol.&volume=59&pages=715-723) (http://scholar.google.com/scholar_lookup?title=Treatment+of+posttraumatic+stress+disorder+in+rape+victims%3A+a+comparison+between+cognitive-behavioral+procedures+and+counseling&author=Foa+E.+B.&author=Rothbaum+B.+O.&author=Riggs+D.+S.&author=Murdock+T.+B.&publication_year=1991&journal=J.+Consult.+Clin.+Psychol.&volume=59&pages=715-723)

Fonzo, G. A., Goodkind, M. S., Oathes, D. J., Zaiko, Y. V., Harvey, M., Peng, K. K., et al. (2017). PTSD psychotherapy outcome predicted by brain activation during emotional reactivity and regulation. *Am. J. Psychiatry* 174, 1163–1174. doi: 10.1176/appi.ajp.2017.16091072

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=28715908) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=28715908>) | [CrossRef Full Text](https://doi.org/10.1176/appi.ajp.2017.16091072) (<https://doi.org/10.1176/appi.ajp.2017.16091072>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=PTSD+psychotherapy+outcome+predicted+by+brain+activation+during+emotional+reactivity+and+regulation&author=Fonzo+G.+A.&author=Goodkind+M.+S.&author=Oathes+D.+J.&author=Zaiko+Y.+V.&author=Harvey+M.&author=Peng+K.+K.&publication_year=2017&journal=Am.+J.+Psychiatry&volume=174&pages=1163-1174) (http://scholar.google.com/scholar_lookup?title=PTSD+psychotherapy+outcome+predicted+by+brain+activation+during+emotional+reactivity+and+regulation&author=Fonzo+G.+A.&author=Goodkind+M.+S.&author=Oathes+D.+J.&author=Zaiko+Y.+V.&author=Harvey+M.&author=Peng+K.+K.&publication_year=2017&journal=Am.+J.+Psychiatry&volume=174&pages=1163-1174)

Fox, K. C. R., Spreng, R. N., Ellamil, M., Andrews-Hanna, J. R., and Christoff, K. (2015). The wandering brain: meta-analysis of functional neuroimaging studies of mind-wandering and related spontaneous thought processes. *NeuroImage* 111, 611–621. doi: 10.1016/j.neuroimage.2015.02.039

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=25725466) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=25725466>) | [CrossRef Full Text](https://doi.org/10.1016/j.neuroimage.2015.02.039) (<https://doi.org/10.1016/j.neuroimage.2015.02.039>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=The+wandering+brain%3A+meta-analysis+of+functional+neuroimaging+studies+of+mind-wandering+and+related+spontaneous+thought+processes&author=Fox+K.+C.+R.&author=Spreng+R.+N.&author=Ellamil+M.&author=Andrews-Hanna+J.+R.&author=Christoff+K.&publication_year=2015&journal=NeuroImage&volume=111&pages=611-621) (http://scholar.google.com/scholar_lookup?title=The+wandering+brain%3A+meta-analysis+of+functional+neuroimaging+studies+of+mind-wandering+and+related+spontaneous+thought+processes&author=Fox+K.+C.+R.&author=Spreng+R.+N.&author=Ellamil+M.&author=Andrews-Hanna+J.+R.&author=Christoff+K.&publication_year=2015&journal=NeuroImage&volume=111&pages=611-621)

Garfinkel, S. N., Abelson, J. L., King, A. P., Sripada, R. K., Wang, X., Gaines, L. M., et al. (2014). Impaired contextual modulation of memories in PTSD: an fMRI and psychophysiological study of extinction retention and fear renewal. *J. Neurosci.* 34, 13435–13443. doi: 10.1523/jneurosci.4287-13.2014

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=25274821) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=25274821>) | [CrossRef Full Text](https://doi.org/10.1523/jneurosci.4287-13.2014) (<https://doi.org/10.1523/jneurosci.4287-13.2014>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Impaired+contextual+modulation+of+memo-) (http://scholar.google.com/scholar_lookup?title=Impaired+contextual+modulation+of+memo-

ries+in+PTSD%3A+an+fMRI+and+psychophysiological+study+of+extinction+retention+and+fear+renewal&author=Garfinkel+S.+N.&author=Abelson+J.+L.&author=King+A.+P.&author=Sripada+R.+K.&author=Wang+X.&author=Gaines+L.+M.&publication_year=2014&journal=J.+Neurosci.&volume=34&pages=13435-13443)

George, M. S., Nahas, Z., Kozel, F. A., Li, X., Denslow, S., Yamanaka, K., et al. (2002). Mechanisms and state of the art of transcranial magnetic stimulation. *J. ECT* 18, 170–181. doi: 10.1097/00124509-200212000-00002

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=12468991) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=12468991) | [CrossRef Full Text](https://doi.org/10.1097/00124509-200212000-00002) (https://doi.org/10.1097/00124509-200212000-00002) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Mechanisms+and+state+of+the+art+of+transcranial+magnetic+stimulation&author=George+M.+S.&author=Nahas+Z.&author=Kozel+F.+A.&author=Li+X.&author=Denslow+S.&author=Yamanaka+K.&publication_year=2002&journal=J.+ECT&volume=18&pages=170-181) (http://scholar.google.com/scholar_lookup?title=Mechanisms+and+state+of+the+art+of+transcranial+magnetic+stimulation&author=George+M.+S.&author=Nahas+Z.&author=Kozel+F.+A.&author=Li+X.&author=Denslow+S.&author=Yamanaka+K.&publication_year=2002&journal=J.+ECT&volume=18&pages=170-181)

George, M. S., and Post, R. M. (2011). Daily left prefrontal repetitive transcranial magnetic stimulation for acute treatment of medication-resistant depression. *Am. J. Psychiatry* 168, 356–364. doi: 10.1176/appi.ajp.2010.10060864

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=21474597) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=21474597) | [CrossRef Full Text](https://doi.org/10.1176/appi.ajp.2010.10060864) (https://doi.org/10.1176/appi.ajp.2010.10060864) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Daily+left+prefrontal+repetitive+transcranial+magnetic+stimulation+for+acute+treatment+of+medication-resistant+depression&author=George+M.+S.&author=Post+R.+M.&publication_year=2011&journal=Am.+J.+Psychiatry&volume=168&pages=356-364) (http://scholar.google.com/scholar_lookup?title=Daily+left+prefrontal+repetitive+transcranial+magnetic+stimulation+for+acute+treatment+of+medication-resistant+depression&author=George+M.+S.&author=Post+R.+M.&publication_year=2011&journal=Am.+J.+Psychiatry&volume=168&pages=356-364)

Gilbertson, M. W., Shenton, M. E., Ciszewski, A., Kasai, K., Lasko, N. B., Orr, S. P., et al. (2002). Smaller hippocampal volume predicts pathologic vulnerability to psychological trauma. *Nat. Neurosci.* 5, 1242–1247. doi: 10.1038/nn958

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=12379862) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=12379862) | [CrossRef Full Text](https://doi.org/10.1038/nn958) (https://doi.org/10.1038/nn958) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Smaller+hippocampal+volume+predicts+pathologic+vulnerability+to+psychological+trauma&author=Gilbertson+M.+W.&author=Shenton+M.+E.&author=Ciszewski+A.&author=Kasai+K.&author=Lasko+N.+B.&author=Orr+S.+P.&publication_year=2002&journal=Nat.+Neurosci.&volume=5&pages=1242-1247) (http://scholar.google.com/scholar_lookup?title=Smaller+hippocampal+volume+predicts+pathologic+vulnerability+to+psychological+trauma&author=Gilbertson+M.+W.&author=Shenton+M.+E.&author=Ciszewski+A.&author=Kasai+K.&author=Lasko+N.+B.&author=Orr+S.+P.&publication_year=2002&journal=Nat.+Neurosci.&volume=5&pages=1242-1247)

Gyurak, A., Gross, J. J., and Etkin, A. (2011). Explicit and implicit emotion regulation: a dual-process framework. *Cogn. Emot.* 25, 400–412. doi: 10.1080/02699931.2010.544160

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=21432682) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=21432682) | [CrossRef Full Text](https://doi.org/10.1080/02699931.2010.544160) (https://doi.org/10.1080/02699931.2010.544160) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Explicit+and+implicit+emotion+regulation%3A+a+dual-process+framework&author=Gyurak+A.&author=Gross+J.+J.&author=Etkin+A.&publication_year=2011&journal=Cogn.+Emot.&volume=25&pages=400-412) (http://scholar.google.com/scholar_lookup?title=Explicit+and+implicit+emotion+regulation%3A+a+dual-process+framework&author=Gyurak+A.&author=Gross+J.+J.&author=Etkin+A.&publication_year=2011&journal=Cogn.+Emot.&volume=25&pages=400-412)

Hariri, A. R., Tessitore, A., Mattay, V. S., Fera, F., and Weinberger, D. R. (2002). The amygdala response to emotional stimuli: a comparison of faces and scenes. *NeuroImage* 17, 317–323. doi: 10.1006/nimg.2002.1179

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=12482086) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=12482086) | [CrossRef Full Text](https://doi.org/10.1006/nimg.2002.1179) (https://doi.org/10.1006/nimg.2002.1179) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=The+amygdala+response+to+emotional+stimuli%3A+a+comparison+of+faces+and+scenes&author=Hariri+A.+R.&author=Tessitore+A.&author=Mattay+V.+S.&author=Fera+F.&author=Weinberger+D.+R.&publication_year=2002&journal=NeuroImage&volume=17&pages=317-323) (http://scholar.google.com/scholar_lookup?title=The+amygdala+response+to+emotional+stimuli%3A+a+comparison+of+faces+and+scenes&author=Hariri+A.+R.&author=Tessitore+A.&author=Mattay+V.+S.&author=Fera+F.&author=Weinberger+D.+R.&publication_year=2002&journal=NeuroImage&volume=17&pages=317-323)

Harris, D. S., Baggott, M., Mendelson, J. H., Mendelson, J. E., and Jones, R. T. (2002). Subjective and hormonal effects of 3,4-methylenedioxymethamphetamine (MDMA) in humans. *Psychopharmacology* 162, 396–405. doi: 10.1007/s00213-002-1131-1

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=12172693) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=12172693) | [CrossRef Full Text](https://doi.org/10.1007/s00213-002-1131-1) (https://doi.org/10.1007/s00213-002-1131-1) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Subjective+and+hormonal+effects+of+3,4-methylenedioxymethamphetamine) (http://scholar.google.com/scholar_lookup?title=Subjective+and+hormonal+effects+of+3,4-methylenedioxymethamphetamine)

(MDMA)+in+humans&author=Harris+D.+S.&author=Baggott+M.&author=Mendelson+J.+H.&author=Mendelson+J.+E.&author=Jones+R.+T.&publication_year=2002&journal=Psychopharmacology&volume=162&pages=396-405)

Hayes, J. P., Hayes, S. M., and Mikedis, A. M. (2012). Quantitative meta-analysis of neural activity in posttraumatic stress disorder. *Biol. Mood Anxiety Disord.* 2:9. doi: 10.1186/2045-5380-2-9

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=22738125) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=22738125) | [CrossRef Full Text](https://doi.org/10.1186/2045-5380-2-9) (https://doi.org/10.1186/2045-5380-2-9) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Quantitative+meta-analysis+of+neural+activity+in+posttraumatic+stress+disorder&author=Hayes+J.+P.&author=Hayes+S.+M.&author=Mikedis+A.+M.&publication_year=2012&journal=Biol.+Mood+Anxiety+Disord.&volume=2&pages=9) (http://scholar.google.com/scholar_lookup?title=Quantitative+meta-analysis+of+neural+activity+in+posttraumatic+stress+disorder&author=Hayes+J.+P.&author=Hayes+S.+M.&author=Mikedis+A.+M.&publication_year=2012&journal=Biol.+Mood+Anxiety+Disord.&volume=2&pages=9)

Helpman, L., Marin, M.-F., Papini, S., Zhu, X., Sullivan, G. M., Schneier, F., et al. (2016a). Neural changes in extinction recall following prolonged exposure treatment for PTSD: a longitudinal fMRI study. *Neuroimage Clin.* 12, 715–723. doi: 10.1016/j.nicl.2016.10.007

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=27761402) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=27761402) | [CrossRef Full Text](https://doi.org/10.1016/j.nicl.2016.10.007) (https://doi.org/10.1016/j.nicl.2016.10.007) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Neural+changes+in+extinction+recall+following+prolonged+exposure+treatment+for+PTSD%3A+a+longitudinal+fMRI+study&author=Helpman+L.&author=Marin+M.-F.&author=Papini+S.&author=Zhu+X.&author=Sullivan+G.+M.&author=Schneier+F.&publication_year=2016&journal=Neuroimage+Clin.&volume=12&pages=715-723) (http://scholar.google.com/scholar_lookup?title=Neural+changes+in+extinction+recall+following+prolonged+exposure+treatment+for+PTSD%3A+a+longitudinal+fMRI+study&author=Helpman+L.&author=Marin+M.-F.&author=Papini+S.&author=Zhu+X.&author=Sullivan+G.+M.&author=Schneier+F.&publication_year=2016&journal=Neuroimage+Clin.&volume=12&pages=715-723)

Helpman, L., Papini, S., Chhetry, B. T., Shvil, E., Rubin, M., Sullivan, G. M., et al. (2016b). PTSD remission after prolonged exposure treatment is associated with anterior cingulate cortex thinning and volume reduction. *Depress. Anxiety* 33, 384–391. doi: 10.1002/da.22471

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=26864570) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=26864570) | [CrossRef Full Text](https://doi.org/10.1002/da.22471) (https://doi.org/10.1002/da.22471) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=PTSD+remission+after+prolonged+exposure+treatment+is+associated+with+anterior+cingulate+cortex+thinning+and+volume+reduction&author=Helpman+L.&author=Papini+S.&author=Chhetry+B.+T.&author=Shvil+E.&author=Rubin+M.&author=Sullivan+G.+M.&publication_year=2016&journal=Depress.+Anxiety&volume=33&pages=384-391) (http://scholar.google.com/scholar_lookup?title=PTSD+remission+after+prolonged+exposure+treatment+is+associated+with+anterior+cingulate+cortex+thinning+and+volume+reduction&author=Helpman+L.&author=Papini+S.&author=Chhetry+B.+T.&author=Shvil+E.&author=Rubin+M.&author=Sullivan+G.+M.&publication_year=2016&journal=Depress.+Anxiety&volume=33&pages=384-391)

Institute of Medicine. (2014). *Treatment for Posttraumatic Stress Disorder in Military and Veteran Populations: Final Assessment*. Washington, DC: The National Academies Press. doi: 10.17226/18724

[CrossRef Full Text](https://doi.org/10.17226/18724) (https://doi.org/10.17226/18724) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Treatment+for+Posttraumatic+Stress+Disorder+in+Military+and+Veteran+Populations%3A+Final+Assessment&Institute+of+Medicine&publication_year=2014) (http://scholar.google.com/scholar_lookup?title=Treatment+for+Posttraumatic+Stress+Disorder+in+Military+and+Veteran+Populations%3A+Final+Assessment&Institute+of+Medicine&publication_year=2014)

Jerud, A. B., Pruitt, L. D., Zoellner, L. A., and Feeny, N. C. (2016). The effects of prolonged exposure and sertraline on emotion regulation in individuals with posttraumatic stress disorder. *Behav. Res. Ther.* 77, 62–67. doi: 10.1016/j.brat.2015.12.002

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=26723004) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=26723004) | [CrossRef Full Text](https://doi.org/10.1016/j.brat.2015.12.002) (https://doi.org/10.1016/j.brat.2015.12.002) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=The+effects+of+prolonged+exposure+and+sertraline+on+emotion+regulation+in+individuals+with+posttraumatic+stress+disorder&author=Jerud+A.+B.&author=Pruitt+L.+D.&author=Zoellner+L.+A.&author=Feeny+N.+C.&publication_year=2016&journal=Behav.+Res.+Ther.&volume=77&pages=62-67) (http://scholar.google.com/scholar_lookup?title=The+effects+of+prolonged+exposure+and+sertraline+on+emotion+regulation+in+individuals+with+posttraumatic+stress+disorder&author=Jerud+A.+B.&author=Pruitt+L.+D.&author=Zoellner+L.+A.&author=Feeny+N.+C.&publication_year=2016&journal=Behav.+Res.+Ther.&volume=77&pages=62-67)

Johansen, J. P., Cain, C. K., Ostroff, L. E., and LeDoux, J. E. (2011). Molecular mechanisms of fear learning and memory. *Cell* 147, 509–524. doi: 10.1016/j.cell.2011.10.009

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=22036561) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=22036561) | [CrossRef Full Text](https://doi.org/10.1016/j.cell.2011.10.009) (https://doi.org/10.1016/j.cell.2011.10.009) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Molecular+mechanisms+of+fear+learning+and+memory&author=Jo) (http://scholar.google.com/scholar_lookup?title=Molecular+mechanisms+of+fear+learning+and+memory&author=Jo

hansen+J.+P.&author=Cain+C.+K.&author=Ostroff+L.+E.&author=LeDoux+J.+E.&publication_year=2011&journal=Cell&volume=147&pages=509-524)

Jovanovic, T., and Ressler, K. J. (2010). How the neurocircuitry and genetics of fear inhibition may inform our understanding of PTSD. *Am. J. Psychiatry* 167, 648–662. doi: 10.1176/appi.ajp.2009.09071074

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=20231322) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=20231322>) | [CrossRef Full Text](https://doi.org/10.1176/appi.ajp.2009.09071074) (<https://doi.org/10.1176/appi.ajp.2009.09071074>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=How+the+neurocircuitry+and+genetics+of+fear+inhibition+may+inform+our+understanding+of+PTSD&author=Jovanovic+T.&author=Ressler+K.+J.&publication_year=2010&journal=Am.+J.+Psychiatry&volume=167&pages=648-662) (http://scholar.google.com/scholar_lookup?title=How+the+neurocircuitry+and+genetics+of+fear+inhibition+may+inform+our+understanding+of+PTSD&author=Jovanovic+T.&author=Ressler+K.+J.&publication_year=2010&journal=Am.+J.+Psychiatry&volume=167&pages=648-662)

Karsen, E. F., Watts, B. V., and Holtzheimer, P. E. (2014). Review of the effectiveness of transcranial magnetic stimulation for post-traumatic stress disorder. *Brain Stimul.* 7, 151–157. doi: 10.1016/j.brs.2013.10.006

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24486424) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24486424>) | [CrossRef Full Text](https://doi.org/10.1016/j.brs.2013.10.006) (<https://doi.org/10.1016/j.brs.2013.10.006>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Review+of+the+effectiveness+of+transcranial+magnetic+stimulation+for+post-traumatic+stress+disorder&author=Karsen+E.+F.&author=Watts+B.+V.&author=Holtzheimer+P.+E.&publication_year=2014&journal=Brain+Stimul.&volume=7&pages=151-157) (http://scholar.google.com/scholar_lookup?title=Review+of+the+effectiveness+of+transcranial+magnetic+stimulation+for+post-traumatic+stress+disorder&author=Karsen+E.+F.&author=Watts+B.+V.&author=Holtzheimer+P.+E.&publication_year=2014&journal=Brain+Stimul.&volume=7&pages=151-157)

Keding, T. J., and Herringa, R. J. (2015). Abnormal structure of fear circuitry in pediatric post-traumatic stress disorder. *Neuropsychopharmacology* 40, 537–545. doi: 10.1038/npp.2014.239

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=25212487) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=25212487>) | [CrossRef Full Text](https://doi.org/10.1038/npp.2014.239) (<https://doi.org/10.1038/npp.2014.239>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Abnormal+structure+of+fear+circuitry+in+pediatric+post-traumatic+stress+disorder&author=Keding+T.+J.&author=Herringa+R.+J.&publication_year=2015&journal=Neuropsychopharmacology&volume=40&pages=537-545) (http://scholar.google.com/scholar_lookup?title=Abnormal+structure+of+fear+circuitry+in+pediatric+post-traumatic+stress+disorder&author=Keding+T.+J.&author=Herringa+R.+J.&publication_year=2015&journal=Neuropsychopharmacology&volume=40&pages=537-545)

King, A. P., Block, S. R., Sripada, R. K., Rauch, S., Giardino, N., Favorite, T., et al. (2016). Altered default mode network (DMN) resting state functional connectivity following a mindfulness-based exposure therapy for posttraumatic stress disorder (PTSD) in combat veterans of afghanistan and IRAQ. *Depress. Anxiety* 33, 289–299. doi: 10.1002/da.22481

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=27038410) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=27038410>) | [CrossRef Full Text](https://doi.org/10.1002/da.22481) (<https://doi.org/10.1002/da.22481>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Altered+default+mode+network+(DMN)+resting+state+functional+connectivity+following+a+mindfulness-based+exposure+therapy+for+posttraumatic+stress+disorder+(PTSD)+in+combat+veterans+of+afghanistan+and+IRAQ&author=King+A.+P.&author=Block+S.+R.&author=Sripada+R.+K.&author=Rauch+S.&author=Giardino+N.&author=Favorite+T.&&publication_year=2016&journal=Depress.+Anxiety&volume=33&pages=289-299) ([http://scholar.google.com/scholar_lookup?title=Altered+default+mode+network+\(DMN\)+resting+state+functional+connectivity+following+a+mindfulness-based+exposure+therapy+for+posttraumatic+stress+disorder+\(PTSD\)+in+combat+veterans+of+afghanistan+and+IRAQ&author=King+A.+P.&author=Block+S.+R.&author=Sripada+R.+K.&author=Rauch+S.&author=Giardino+N.&author=Favorite+T.&&publication_year=2016&journal=Depress.+Anxiety&volume=33&pages=289-299](http://scholar.google.com/scholar_lookup?title=Altered+default+mode+network+(DMN)+resting+state+functional+connectivity+following+a+mindfulness-based+exposure+therapy+for+posttraumatic+stress+disorder+(PTSD)+in+combat+veterans+of+afghanistan+and+IRAQ&author=King+A.+P.&author=Block+S.+R.&author=Sripada+R.+K.&author=Rauch+S.&author=Giardino+N.&author=Favorite+T.&&publication_year=2016&journal=Depress.+Anxiety&volume=33&pages=289-299))

Koch, S. B. J., van Zuiden, M., Nawijn, L., Frijling, J. L., Veltman, D. J., and Olf, M. (2016). Aberrant resting-state brain activity in posttraumatic stress disorder: a meta-analysis and systematic review. *Depress. Anxiety* 33, 592–605. doi: 10.1002/da.22478

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=26918313) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=26918313>) | [CrossRef Full Text](https://doi.org/10.1002/da.22478) (<https://doi.org/10.1002/da.22478>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Aberrant+resting-state+brain+activity+in+posttraumatic+stress+disorder%3A+a+meta-analysis+and+systematic+review&author=Koch+S.+B.+J.&author=van+Zuiden+M.&author=Nawijn+L.&author=Frijling+J.+L.&author=Veltman+D.+J.&author=Olf+M.&publication_year=2016&journal=Depress.+Anxiety&volume=33&pages=592-605) (http://scholar.google.com/scholar_lookup?title=Aberrant+resting-state+brain+activity+in+posttraumatic+stress+disorder%3A+a+meta-analysis+and+systematic+review&author=Koch+S.+B.+J.&author=van+Zuiden+M.&author=Nawijn+L.&author=Frijling+J.+L.&author=Veltman+D.+J.&author=Olf+M.&publication_year=2016&journal=Depress.+Anxiety&volume=33&pages=592-605)

Kohn, N., Eickhoff, S. B., Scheller, M., Laird, A. R., Fox, P. T., and Habel, U. (2014). Neural network of cognitive emotion regulation — An ALE meta-analysis and MACM analysis. *NeuroImage* 87, 345–355. doi: 10.1016/j.neuroimage.2013.11.001

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24220041) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24220041>) | [CrossRef Full Text](https://doi.org/10.1016/j.neuroimage.2013.11.001) (<https://doi.org/10.1016/j.neuroimage.2013.11.001>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Neural+network+of+cognitive+emotion+regulation+-+An+ALE+meta-analysis+and+MACM+analysis&author=Kohn+N.&author=Eickhoff+S.+B.&author=Scheller+M.&author=Laird+A.+R.&author=Fox+P.+T.&author=Habel+U.&publication_year=2014&journal=NeuroImage&volume=87&pages=345-355) (http://scholar.google.com/scholar_lookup?title=Neural+network+of+cognitive+emotion+regulation+-+An+ALE+meta-analysis+and+MACM+analysis&author=Kohn+N.&author=Eickhoff+S.+B.&author=Scheller+M.&author=Laird+A.+R.&author=Fox+P.+T.&author=Habel+U.&publication_year=2014&journal=NeuroImage&volume=87&pages=345-355)

Lambert, J. J., Belelli, D., Peden, D. R., Vardy, A. W., and Peters, J. A. (2003). Neurosteroid modulation of GABAA receptors. *Prog. Neurobiol.* 71, 67–80. doi: 10.1016/j.pneurobio.2003.09.001

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=14611869) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=14611869>) | [CrossRef Full Text](https://doi.org/10.1016/j.pneurobio.2003.09.001) (<https://doi.org/10.1016/j.pneurobio.2003.09.001>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Neurosteroid+modulation+of+GABAA+receptors&author=Lambert+J.+J.&author=Belelli+D.&author=Peden+D.+R.&author=Vardy+A.+W.&author=Peters+J.+A.&publication_year=2003&journal=Prog.+Neurobiol.&volume=71&pages=67-80) (http://scholar.google.com/scholar_lookup?title=Neurosteroid+modulation+of+GABAA+receptors&author=Lambert+J.+J.&author=Belelli+D.&author=Peden+D.+R.&author=Vardy+A.+W.&author=Peters+J.+A.&publication_year=2003&journal=Prog.+Neurobiol.&volume=71&pages=67-80)

Ledgerwood, L., Richardson, R., and Cranney, J. (2005). D-cycloserine facilitates extinction of learned fear: effects on reacquisition and generalized extinction. *Biol. Psychiatry* 57, 841–847. doi: 10.1016/j.biopsych.2005.01.023

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=15820704) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=15820704>) | [CrossRef Full Text](https://doi.org/10.1016/j.biopsych.2005.01.023) (<https://doi.org/10.1016/j.biopsych.2005.01.023>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=D-cycloserine+facilitates+extinction+of+learned+fear%3A+effects+on+reacquisition+and+generalized+extinction&author=Ledgerwood+L.&author=Richardson+R.&author=Cranney+J.&publication_year=2005&journal=Biol.+Psychiatry&volume=57&pages=841-847) (http://scholar.google.com/scholar_lookup?title=D-cycloserine+facilitates+extinction+of+learned+fear%3A+effects+on+reacquisition+and+generalized+extinction&author=Ledgerwood+L.&author=Richardson+R.&author=Cranney+J.&publication_year=2005&journal=Biol.+Psychiatry&volume=57&pages=841-847)

LeDoux, J. E. (2000). Emotion circuits in the brain. *Annu. Rev. Neurosci.* 23, 155–184. doi: 10.1146/annurev.neuro.23.1.155

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=10845062) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=10845062>) | [CrossRef Full Text](https://doi.org/10.1146/annurev.neuro.23.1.155) (<https://doi.org/10.1146/annurev.neuro.23.1.155>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Emotion+circuits+in+the+brain&author=LeDoux+J.+E.&publication_year=2000&journal=Annu.+Rev.+Neurosci.&volume=23&pages=155-184) (http://scholar.google.com/scholar_lookup?title=Emotion+circuits+in+the+brain&author=LeDoux+J.+E.&publication_year=2000&journal=Annu.+Rev.+Neurosci.&volume=23&pages=155-184)

Lee, J. L. C., Milton, A. L., and Everitt, B. J. (2006). Reconsolidation and extinction of conditioned fear: inhibition and potentiation. *J. Neurosci.* 26, 10051–10056. doi: 10.1523/jneurosci.2466-06.2006

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=17005868) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=17005868>) | [CrossRef Full Text](https://doi.org/10.1523/jneurosci.2466-06.2006) (<https://doi.org/10.1523/jneurosci.2466-06.2006>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Reconsolidation+and+extinction+of+conditioned+fear%3A+inhibition+and+potentiation&author=Lee+J.+L.+C.&author=Milton+A.+L.&author=Everitt+B.+J.&publication_year=2006&journal=J.+Neurosci.&volume=26&pages=10051-10056) (http://scholar.google.com/scholar_lookup?title=Reconsolidation+and+extinction+of+conditioned+fear%3A+inhibition+and+potentiation&author=Lee+J.+L.+C.&author=Milton+A.+L.&author=Everitt+B.+J.&publication_year=2006&journal=J.+Neurosci.&volume=26&pages=10051-10056)

Liberzon, I., and Abelson, J. L. (2016). Context processing and the neurobiology of post-traumatic stress disorder. *Neuron* 92, 14–30. doi: 10.1016/j.neuron.2016.09.039

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=27710783) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=27710783>) | [CrossRef Full Text](https://doi.org/10.1016/j.neuron.2016.09.039) (<https://doi.org/10.1016/j.neuron.2016.09.039>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Context+processing+and+the+neurobiology+of+post-traumatic+stress+disorder&author=Liberzon+I.&author=Abelson+J.+L.&publication_year=2016&journal=Neuron&volume=92&pages=14-30) (http://scholar.google.com/scholar_lookup?title=Context+processing+and+the+neurobiology+of+post-traumatic+stress+disorder&author=Liberzon+I.&author=Abelson+J.+L.&publication_year=2016&journal=Neuron&volume=92&pages=14-30)

Lindauer, R. J. L., Vlioger, E. J., Jalink, M., Olf, M., Carlier, I. V. E., Majoie, C. B. L. M., et al. (2005). Effects of psychotherapy on hippocampal volume in out-patients with post-traumatic stress disorder: a MRI investigation. *Psychol. Med.* 35, 1421–1431. doi: 10.1017/S0033291705005246

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=16164766) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=16164766>) | [CrossRef Full Text](https://doi.org/10.1017/S0033291705005246) (<https://doi.org/10.1017/S0033291705005246>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Effects+of+psychotherapy+on+hippocampal+volume+in+out-pa) (http://scholar.google.com/scholar_lookup?title=Effects+of+psychotherapy+on+hippocampal+volume+in+out-pa)

tients+with+post-traumatic+stress+disorder%3A+a+MRI+investigation&author=Lindauer+R.+J.+L.&author=Vlieger+E.+J.&author=Jalink+M.&author=Olff+M.&author=Carlier+I.+V.+E.&author=Ma-
joie+C.+B.+L.+M.&publication_year=2005&journal=Psychol.+Med.&volume=35&pages=1421-1431)

Lindgren, L., Bergdahl, J., and Nyberg, L. (2016). Longitudinal evidence for smaller hippocampus volume as a vulnerability factor for perceived stress. *Cereb. Cortex* 26, 3527–3533. doi: 10.1093/cercor/bhw154

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=27230217) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=27230217) | [CrossRef Full Text](https://doi.org/10.1093/cercor/bhw154) (https://doi.org/10.1093/cercor/bhw154) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Longitudinal+evidence+for+smaller+hippocampus+volume+as+a+vulnerability+factor+for+perceived+stress&author=Lindgren+L.&author=Bergdahl+J.&author=Nyberg+L.&publication_year=2016&journal=Cereb.+Cortex&volume=26&pages=3527-3533) (http://scholar.google.com/scholar_lookup?title=Longitudinal+evidence+for+smaller+hippocampus+volume+as+a+vulnerability+factor+for+perceived+stress&author=Lindgren+L.&author=Bergdahl+J.&author=Nyberg+L.&publication_year=2016&journal=Cereb.+Cortex&volume=26&pages=3527-3533)

Lissek, S., and Grillon, C. (2012). "Learning models of PTSD," in *The Oxford Handbook of Traumatic Disorder*, eds J. G. Beck and D. M. Sloan (Oxford: Oxford University Press), 175–190.

[Google Scholar](http://scholar.google.com/scholar_lookup?title=Learning+models+of+PTSD&author=Lissek+S.&author=Grillon+C.&publication_year=2012&pages=175-190) (http://scholar.google.com/scholar_lookup?title=Learning+models+of+PTSD&author=Lissek+S.&author=Grillon+C.&publication_year=2012&pages=175-190)

Litz, B. T., Salters-Pedneault, K., Steenkamp, M. M., Hermos, J. A., Bryant, R. A., Otto, M. W., et al. (2012). A randomized placebo-controlled trial of D-cycloserine and exposure therapy for posttraumatic stress disorder. *J. Psychiatr. Res.* 46, 1184–1190. doi: 10.1016/j.jpsychires.2012.05.006

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=22694905) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=22694905) | [CrossRef Full Text](https://doi.org/10.1016/j.jpsychires.2012.05.006) (https://doi.org/10.1016/j.jpsychires.2012.05.006) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=A+randomized+placebo-controlled+trial+of+D-cycloserine+and+exposure+therapy+for+posttraumatic+stress+disorder&author=Litz+B.+T.&author=Salters-Pedneault+K.&author=Steenkamp+M.+M.&author=Hermos+J.+A.&author=Bryant+R.+A.&author=Otto+M.+W.&publication_year=2012&journal=J.+Psychiatr.+Res.&volume=46&pages=1184-1190) (http://scholar.google.com/scholar_lookup?title=A+randomized+placebo-controlled+trial+of+D-cycloserine+and+exposure+therapy+for+posttraumatic+stress+disorder&author=Litz+B.+T.&author=Salters-Pedneault+K.&author=Steenkamp+M.+M.&author=Hermos+J.+A.&author=Bryant+R.+A.&author=Otto+M.+W.&publication_year=2012&journal=J.+Psychiatr.+Res.&volume=46&pages=1184-1190)

Lopresto, D., Schipper, P., and Homberg, J. R. (2016). Neural circuits and mechanisms involved in fear generalization: implications for the pathophysiology and treatment of posttraumatic stress disorder. *Neurosci. Biobehav. Rev.* 60, 31–42. doi: 10.1016/j.neubiorev.2015.10.009

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=26519776) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=26519776) | [CrossRef Full Text](https://doi.org/10.1016/j.neubiorev.2015.10.009) (https://doi.org/10.1016/j.neubiorev.2015.10.009) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Neural+circuits+and+mechanisms+involved+in+fear+generalization%3A+implications+for+the+pathophysiology+and+treatment+of+posttraumatic+stress+disorder&author=Lopresto+D.&author=Schipper+P.&author=Homberg+J.+R.&publication_year=2016&journal=Neurosci.+Biobehav.+Rev.&volume=60&pages=31-42) (http://scholar.google.com/scholar_lookup?title=Neural+circuits+and+mechanisms+involved+in+fear+generalization%3A+implications+for+the+pathophysiology+and+treatment+of+posttraumatic+stress+disorder&author=Lopresto+D.&author=Schipper+P.&author=Homberg+J.+R.&publication_year=2016&journal=Neurosci.+Biobehav.+Rev.&volume=60&pages=31-42)

Maninger, N., Wolkowitz, O. M., Reus, V. I., Epel, E. S., and Mellon, S. H. (2009). Neurobiological and neuropsychiatric effects of dehydroepiandrosterone (DHEA) and DHEA sulfate (DHEAS). *Front. Neuroendocrinol.* 30, 65–91. doi: 10.1016/j.yfrne.2008.11.002

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=19063914) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=19063914) | [CrossRef Full Text](https://doi.org/10.1016/j.yfrne.2008.11.002) (https://doi.org/10.1016/j.yfrne.2008.11.002) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Neurobiological+and+neuropsychiatric+effects+of+dehydroepiandrosterone+(DHEA)+and+DHEA+sulfate+(DHEAS)&author=Maninger+N.&author=Wolkowitz+O.+M.&author=Reus+V.+I.&author=Epel+E.+S.&author=Mellon+S.+H.&publication_year=2009&journal=Front.+Neuroendocrinol.&volume=30&pages=65-91) (http://scholar.google.com/scholar_lookup?title=Neurobiological+and+neuropsychiatric+effects+of+dehydroepiandrosterone+(DHEA)+and+DHEA+sulfate+(DHEAS)&author=Maninger+N.&author=Wolkowitz+O.+M.&author=Reus+V.+I.&author=Epel+E.+S.&author=Mellon+S.+H.&publication_year=2009&journal=Front.+Neuroendocrinol.&volume=30&pages=65-91)

Maren, S., and Fanselow, M. S. (1995). Synaptic plasticity in the basolateral amygdala induced by hippocampal formation stimulation *in vivo*. *J. Neurosci.* 15, 7548–7564. doi: 10.1523/jneurosci.15-11-07548.1995

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=7472506) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=7472506) | [CrossRef Full Text](https://doi.org/10.1523/jneurosci.15-11-07548.1995) (https://doi.org/10.1523/jneurosci.15-11-07548.1995) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Synaptic+plasticity+in+the+basolateral+amygdala+in-) (http://scholar.google.com/scholar_lookup?title=Synaptic+plasticity+in+the+basolateral+amygdala+in-

duced+by+hippocampal+formation+stimulation+in+vivo&author=Maren+S.&author=Fanselow+M.+S.&publication_year=1995&journal=J.+Neurosci.&volume=15&pages=7548-7564)

Maren, S., Phan, K. L., and Liberzon, I. (2013). The contextual brain: implications for fear conditioning, extinction and psychopathology. *Nat. Rev. Neurosci.* 14, 417–428. doi: 10.1038/nrn3492

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23635870) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23635870) | [CrossRef Full Text](https://doi.org/10.1038/nrn3492) (https://doi.org/10.1038/nrn3492) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=The+contextual+brain%3A+implications+for+fear+conditioning,+extinction+and+psychopathology&author=Maren+S.&author=Phan+K.+L.&author=Liberzon+I.&publication_year=2013&journal=Nat.+Rev.+Neurosci.&volume=14&pages=417-428) (http://scholar.google.com/scholar_lookup?title=The+contextual+brain%3A+implications+for+fear+conditioning,+extinction+and+psychopathology&author=Maren+S.&author=Phan+K.+L.&author=Liberzon+I.&publication_year=2013&journal=Nat.+Rev.+Neurosci.&volume=14&pages=417-428)

Marx, C. E., Naylor, J. C., Kilts, J. D., Dunn, C. E., Tupler, L. A., Szabo, S. T., et al. (2016). "Neurosteroids and traumatic brain injury translating biomarkers to therapeutics; overview and pilot investigations in Iraq and Afghanistan era veterans," in *Translational Research in Traumatic Brain Injury*, eds D. Laskowitz and G. Grant (Boca Raton, FL: CRC Press/Taylor and Francis Group), 145–161.

McLaughlin, A. A., Keller, S. M., Feeny, N. C., Youngstrom, E. A., and Zoellner, L. A. (2014). Patterns of therapeutic alliance: rupture-repair episodes in prolonged exposure for posttraumatic stress disorder. *J. Consult. Clin. Psychol.* 82, 112–121. doi: 10.1037/a0034696

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24188510) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24188510) | [CrossRef Full Text](https://doi.org/10.1037/a0034696) (https://doi.org/10.1037/a0034696) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Patterns+of+therapeutic+alliance%3A+rupture-repair+episodes+in+prolonged+exposure+for+posttraumatic+stress+disorder&author=McLaughlin+A.+A.&author=Keller+S.+M.&author=Feeny+N.+C.&author=Youngstrom+E.+A.&author=Zoellner+L.+A.&publication_year=2014&journal=J.+Consult.+Clin.+Psychol.&volume=82&pages=112-121) (http://scholar.google.com/scholar_lookup?title=Patterns+of+therapeutic+alliance%3A+rupture-repair+episodes+in+prolonged+exposure+for+posttraumatic+stress+disorder&author=McLaughlin+A.+A.&author=Keller+S.+M.&author=Feeny+N.+C.&author=Youngstrom+E.+A.&author=Zoellner+L.+A.&publication_year=2014&journal=J.+Consult.+Clin.+Psychol.&volume=82&pages=112-121)

Milad, M. R., Igoe, S. A., Lebron-Milad, K., and Novales, J. E. (2009). Estrous cycle phase and gonadal hormones influence conditioned fear extinction. *Neuroscience* 164, 887–895. doi: 10.1016/j.neuroscience.2009.09.011

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=19761818) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=19761818) | [CrossRef Full Text](https://doi.org/10.1016/j.neuroscience.2009.09.011) (https://doi.org/10.1016/j.neuroscience.2009.09.011) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Estrous+cycle+phase+and+gonadal+hormones+influence+conditioned+fear+extinction&author=Milad+M.+R.&author=Igoe+S.+A.&author=Lebron-Milad+K.&author=Novales+J.+E.&publication_year=2009&journal=Neuroscience&volume=164&pages=887-895) (http://scholar.google.com/scholar_lookup?title=Estrous+cycle+phase+and+gonadal+hormones+influence+conditioned+fear+extinction&author=Milad+M.+R.&author=Igoe+S.+A.&author=Lebron-Milad+K.&author=Novales+J.+E.&publication_year=2009&journal=Neuroscience&volume=164&pages=887-895)

Milad, M. R., Orr, S. P., Lasko, N. B., Chang, Y., Rauch, S. L., and Pitman, R. K. (2008). Presence and acquired origin of reduced recall for fear extinction in PTSD: results of a twin study. *J. Psychiatr. Res.* 42, 515–520. doi: 10.1016/j.jpsychires.2008.01.017

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=18313695) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=18313695) | [CrossRef Full Text](https://doi.org/10.1016/j.jpsychires.2008.01.017) (https://doi.org/10.1016/j.jpsychires.2008.01.017) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Presence+and+acquired+origin+of+reduced+recall+for+fear+extinction+in+PTSD%3A+results+of+a+twin+study&author=Milad+M.+R.&author=Orr+S.+P.&author=Lasko+N.+B.&author=Chang+Y.&author=Rauch+S.+L.&author=Pitman+R.+K.&publication_year=2008&journal=J.+Psychiatr.+Res.&volume=42&pages=515-520) (http://scholar.google.com/scholar_lookup?title=Presence+and+acquired+origin+of+reduced+recall+for+fear+extinction+in+PTSD%3A+results+of+a+twin+study&author=Milad+M.+R.&author=Orr+S.+P.&author=Lasko+N.+B.&author=Chang+Y.&author=Rauch+S.+L.&author=Pitman+R.+K.&publication_year=2008&journal=J.+Psychiatr.+Res.&volume=42&pages=515-520)

Milad, M. R., Pitman, R. K., Ellis, C. B., Gold, A. L., Shin, L. M., Lasko, N. B., et al. (2009). Neurobiological basis of failure to recall extinction memory in posttraumatic stress disorder. *Biol. Psychiatry* 66, 1075–1082. doi: 10.1016/j.biopsych.2009.06.026.

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=19748076) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=19748076) | [CrossRef Full Text](https://doi.org/10.1016/j.biopsych.2009.06.026) (https://doi.org/10.1016/j.biopsych.2009.06.026.) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Neurobiological+basis+of+failure+to+recall+extinction+memory+in+posttraumatic+stress+disorder&author=Milad+M.+R.&author=Pitman+R.+K.&author=Ellis+C.+B.&author=Gold+A.+L.&author=Shin+L.+M.&author=Lasko+N.+B.&publication_year=2009&journal=Biol.+Psychiatry&volume=66&pages=1075-1082) (http://scholar.google.com/scholar_lookup?title=Neurobiological+basis+of+failure+to+recall+extinction+memory+in+posttraumatic+stress+disorder&author=Milad+M.+R.&author=Pitman+R.+K.&author=Ellis+C.+B.&author=Gold+A.+L.&author=Shin+L.+M.&author=Lasko+N.+B.&publication_year=2009&journal=Biol.+Psychiatry&volume=66&pages=1075-1082)

Mithoefer, M. C. (2011). MDMA-assisted psychotherapy for the treatment of posttraumatic stress disorder: a revised teaching manual. Available online at: http://www.maps.org/research/mdma/Manual_MDMAPTSD_30Nov11.pdf (http://www.maps.org/research/mdma/Manual_MDMAPTSD_30Nov11.pdf)

Mithoefer, M. C., Mithoefer, A. T., Feduccia, A. A., Jerome, L., Wagner, M., Wymer, J., et al. (2018). 3,4-methylenedioxymethamphetamine (MDMA)-assisted psychotherapy for post-traumatic stress disorder in military veterans, firefighters and police officers: a randomised, double-blind, dose-response, phase 2 clinical trial. *Lancet Psychiatry* 5, 486–497. doi: 10.1016/s2215-0366(18)30135-4

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=29728331) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=29728331>) | [CrossRef Full Text](https://doi.org/10.1016/s2215-0366(18)30135-4) ([https://doi.org/10.1016/s2215-0366\(18\)30135-4](https://doi.org/10.1016/s2215-0366(18)30135-4)) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=3,4-methylenedioxyamphetamine+(MDMA)-assisted+psychotherapy+for+post-traumatic+stress+disorder+in+military+veterans,+firefighters+and+police+officers%3A+a+randomised,+double-blind,+dose-response,+phase+2+clinical+trial&author=Mithoefer+M.+C.&author=Mithoefer+A.+T.&author=Feduccia+A.+A.&author=Jerome+L.&author=Wagner+M.&author=Wymer+J.&publication_year=2018&journal=Lancet+Psychiatry&volume=5&pages=486-497) ([http://scholar.google.com/scholar_lookup?title=3,4-methylenedioxyamphetamine+\(MDMA\)-assisted+psychotherapy+for+post-traumatic+stress+disorder+in+military+veterans,+firefighters+and+police+officers%3A+a+randomised,+double-blind,+dose-response,+phase+2+clinical+trial&author=Mithoefer+M.+C.&author=Mithoefer+A.+T.&author=Feduccia+A.+A.&author=Jerome+L.&author=Wagner+M.&author=Wymer+J.&publication_year=2018&journal=Lancet+Psychiatry&volume=5&pages=486-497](http://scholar.google.com/scholar_lookup?title=3,4-methylenedioxyamphetamine+(MDMA)-assisted+psychotherapy+for+post-traumatic+stress+disorder+in+military+veterans,+firefighters+and+police+officers%3A+a+randomised,+double-blind,+dose-response,+phase+2+clinical+trial&author=Mithoefer+M.+C.&author=Mithoefer+A.+T.&author=Feduccia+A.+A.&author=Jerome+L.&author=Wagner+M.&author=Wymer+J.&publication_year=2018&journal=Lancet+Psychiatry&volume=5&pages=486-497))

Mithoefer, M. C., Wagner, M. T., Mithoefer, A. T., Jerome, L., and Doblin, R. (2011). The safety and efficacy of \pm 3,4-methylenedioxymethamphetamine-assisted psychotherapy in subjects with chronic, treatment-resistant posttraumatic stress disorder: the first randomized controlled pilot study. *J. Psychopharmacol.* 25, 439–452. doi: 10.1177/0269881110378371

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=20643699) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=20643699>) | [CrossRef Full Text](https://doi.org/10.1177/0269881110378371) (<https://doi.org/10.1177/0269881110378371>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=The+safety+and+efficacy+of+3,4-methylenedioxyamphetamine-assisted+psychotherapy+in+subjects+with+chronic,+treatment-resistant+posttraumatic+stress+disorder%3A+the+first+randomized+controlled+pilot+study&author=Mithoefer+M.+C.&author=Wagner+M.+T.&author=Mithoefer+A.+T.&author=Jerome+L.&author=Doblin+R.&publication_year=2011&journal=J.+Psychopharmacol.&volume=25&pages=439-452) (http://scholar.google.com/scholar_lookup?title=The+safety+and+efficacy+of+3,4-methylenedioxyamphetamine-assisted+psychotherapy+in+subjects+with+chronic,+treatment-resistant+posttraumatic+stress+disorder%3A+the+first+randomized+controlled+pilot+study&author=Mithoefer+M.+C.&author=Wagner+M.+T.&author=Mithoefer+A.+T.&author=Jerome+L.&author=Doblin+R.&publication_year=2011&journal=J.+Psychopharmacol.&volume=25&pages=439-452)

Mowrer, O. (1960). *Learning and its Symbolic Processes*. Hoboken, NJ: John Wiley & Sons.

Negash, S., Kliot, D., Howard, D. V., Howard, J. H., Das, S. R., Yushkevich, P. A., et al. (2015). Relationship of contextual cueing and hippocampal volume in amnesic mild cognitive impairment patients and cognitively normal older adults. *J. Int. Neuropsychol. Soc.* 21, 285–296. doi: 10.1017/s1355617715000223

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=25991413) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=25991413>) | [CrossRef Full Text](https://doi.org/10.1017/s1355617715000223) (<https://doi.org/10.1017/s1355617715000223>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Relationship+of+contextual+cueing+and+hippocampal+volume+in+amnesic+mild+cognitive+impairment+patients+and+cognitively+normal+older+adults&author=Negash+S.&author=Kliot+D.&author=Howard+D.+V.&author=Howard+J.+H.&author=Das+S.+R.&author=Yushkevich+P.+A.&publication_year=2015&journal=J.+Int.+Neuropsychol.+Soc.&volume=21&pages=285-296) (http://scholar.google.com/scholar_lookup?title=Relationship+of+contextual+cueing+and+hippocampal+volume+in+amnesic+mild+cognitive+impairment+patients+and+cognitively+normal+older+adults&author=Negash+S.&author=Kliot+D.&author=Howard+D.+V.&author=Howard+J.+H.&author=Das+S.+R.&author=Yushkevich+P.+A.&publication_year=2015&journal=J.+Int.+Neuropsychol.+Soc.&volume=21&pages=285-296)

New, A. S., Fan, J., Murrough, J. W., Liu, X., Liebman, R. E., Guise, K. G., et al. (2009). A functional magnetic resonance imaging study of deliberate emotion regulation in resilience and posttraumatic stress disorder. *Biol. Psychiatry* 66, 656–664. doi: 10.1016/j.biopsych.2009.05.020

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=19589502) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=19589502>) | [CrossRef Full Text](https://doi.org/10.1016/j.biopsych.2009.05.020) (<https://doi.org/10.1016/j.biopsych.2009.05.020>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=A+functional+magnetic+resonance+imaging+study+of+deliberate+emotion+regulation+in+resilience+and+posttraumatic+stress+disorder&author=New+A.+S.&author=Fan+J.&author=Murrough+J.+W.&author=Liu+X.&author=Liebman+R.+E.&author=Guise+K.+G.&publication_year=2009&journal=Biol.+Psychiatry&volume=66&pages=656-664) (http://scholar.google.com/scholar_lookup?title=A+functional+magnetic+resonance+imaging+study+of+deliberate+emotion+regulation+in+resilience+and+posttraumatic+stress+disorder&author=New+A.+S.&author=Fan+J.&author=Murrough+J.+W.&author=Liu+X.&author=Liebman+R.+E.&author=Guise+K.+G.&publication_year=2009&journal=Biol.+Psychiatry&volume=66&pages=656-664)

Norberg, M. M., Krystal, J. H., and Tolin, D. F. (2008). A meta-analysis of D-cycloserine and the facilitation of fear extinction and exposure therapy. *Biol. Psychiatry* 63, 1118–1126. doi: 10.1016/j.biopsych.2008.01.012

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=18313643) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=18313643>) | [CrossRef Full Text](https://doi.org/10.1016/j.biopsych.2008.01.012) (<https://doi.org/10.1016/j.biopsych.2008.01.012>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=A+meta-analysis+of+D-cycloserine+and+the+facilita-) (http://scholar.google.com/scholar_lookup?title=A+meta-analysis+of+D-cycloserine+and+the+facilita-

tion+of+fear+extinction+and+exposure+therapy&author=Norberg+M.+M.&author=Krystal+J.+H.&author=Tolin+D.+F.&publication_year=2008&journal=Biol.+Psychiatry&volume=63&pages=1118-1126)

Ochsner, K. N., Bunge, S. A., Gross, J. J., and Gabrieli, J. D. E. (2002). Rethinking feelings: an fMRI study of the cognitive regulation of emotion. *J. Cogn. Neurosci.* 14, 1215–1229. doi: 10.1162/089892902760807212

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=12495527) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=12495527>) | [CrossRef Full Text](https://doi.org/10.1162/089892902760807212) (<https://doi.org/10.1162/089892902760807212>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Rethinking+feelings%3A+an+fMRI+study+of+the+cognitive+regulation+of+emotion&author=Ochsner+K.+N.&author=Bunge+S.+A.&author=Gross+J.+J.&author=Gabrieli+J.+D.+E.&publication_year=2002&journal=J.+Cogn.+Neurosci.&volume=14&pages=1215-1229) (http://scholar.google.com/scholar_lookup?title=Rethinking+feelings%3A+an+fMRI+study+of+the+cognitive+regulation+of+emotion&author=Ochsner+K.+N.&author=Bunge+S.+A.&author=Gross+J.+J.&author=Gabrieli+J.+D.+E.&publication_year=2002&journal=J.+Cogn.+Neurosci.&volume=14&pages=1215-1229)

Ochsner, K. N., Silvers, J. A., and Buhle, J. T. (2012). Functional imaging studies of emotion regulation: a synthetic review and evolving model of the cognitive control of emotion. *Ann. N Y Acad. Sci.* 1251, E1–E24. doi: 10.1111/j.1749-6632.2012.06751.x

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23025352) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23025352>) | [CrossRef Full Text](https://doi.org/10.1111/j.1749-6632.2012.06751.x) (<https://doi.org/10.1111/j.1749-6632.2012.06751.x>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Functional+imaging+studies+of+emotion+regulation%3A+a+synthetic+review+and+evolving+model+of+the+cognitive+control+of+emotion&author=Ochsner+K.+N.&author=Silvers+J.+A.&author=Buhle+J.+T.&publication_year=2012&journal=Ann.+N+Y+Acad.+Sci.&volume=1251&pages=E1-E24) (http://scholar.google.com/scholar_lookup?title=Functional+imaging+studies+of+emotion+regulation%3A+a+synthetic+review+and+evolving+model+of+the+cognitive+control+of+emotion&author=Ochsner+K.+N.&author=Silvers+J.+A.&author=Buhle+J.+T.&publication_year=2012&journal=Ann.+N+Y+Acad.+Sci.&volume=1251&pages=E1-E24)

Oehen, P., Traber, R., Widmer, V., and Schnyder, U. (2013). A randomized, controlled pilot study of MDMA (\pm 3,4-Methylenedioxymethamphetamine)-assisted psychotherapy for treatment of resistant, chronic post-traumatic stress disorder (PTSD). *J. Psychopharmacol.* 27, 40–52. doi: 10.1177/0269881112464827

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23118021) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23118021>) | [CrossRef Full Text](https://doi.org/10.1177/0269881112464827) (<https://doi.org/10.1177/0269881112464827>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=A+randomized,+controlled+pilot+study+of+MDMA+(+3,4-Methylene-dioxymethamphetamine)-assisted+psychotherapy+for+treatment+of+resistant,+chronic+post-traumatic+stress+disorder+(PTSD)&author=Oehen+P.&author=Traber+R.&author=Widmer+V.&author=Schnyder+U.&publication_year=2013&journal=J.+Psychopharmacol.&volume=27&pages=40-52) ([http://scholar.google.com/scholar_lookup?title=A+randomized,+controlled+pilot+study+of+MDMA+\(+3,4-Methylene-dioxymethamphetamine\)-assisted+psychotherapy+for+treatment+of+resistant,+chronic+post-traumatic+stress+disorder+\(PTSD\)&author=Oehen+P.&author=Traber+R.&author=Widmer+V.&author=Schnyder+U.&publication_year=2013&journal=J.+Psychopharmacol.&volume=27&pages=40-52](http://scholar.google.com/scholar_lookup?title=A+randomized,+controlled+pilot+study+of+MDMA+(+3,4-Methylene-dioxymethamphetamine)-assisted+psychotherapy+for+treatment+of+resistant,+chronic+post-traumatic+stress+disorder+(PTSD)&author=Oehen+P.&author=Traber+R.&author=Widmer+V.&author=Schnyder+U.&publication_year=2013&journal=J.+Psychopharmacol.&volume=27&pages=40-52))

Olf, M., Langeland, W., Witteveen, A., and Denys, D. (2010). A psychobiological rationale for oxytocin in the treatment of posttraumatic stress disorder. *CNS Spectr.* 15, 522–530. doi: 10.1017/s109285290000047x

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=20703199) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=20703199>) | [CrossRef Full Text](https://doi.org/10.1017/s109285290000047x) (<https://doi.org/10.1017/s109285290000047x>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=A+psychobiological+rationale+for+oxytocin+in+the+treatment+of+posttraumatic+stress+disorder&author=Olf+M.&author=Langeland+W.&author=Witteveen+A.&author=Denys+D.&publication_year=2010&journal=CNS+Spectr.&volume=15&pages=522-530) (http://scholar.google.com/scholar_lookup?title=A+psychobiological+rationale+for+oxytocin+in+the+treatment+of+posttraumatic+stress+disorder&author=Olf+M.&author=Langeland+W.&author=Witteveen+A.&author=Denys+D.&publication_year=2010&journal=CNS+Spectr.&volume=15&pages=522-530)

Osuji, I. J., Vera-Bolaños, E., Carmody, T. J., and Brown, E. S. (2010). Pregnenolone for cognition and mood in dual diagnosis patients. *Psychiatry Res.* 178, 309–312. doi: 10.1016/j.psychres.2009.09.006

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=20493557) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=20493557>) | [CrossRef Full Text](https://doi.org/10.1016/j.psychres.2009.09.006) (<https://doi.org/10.1016/j.psychres.2009.09.006>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Pregnenolone+for+cognition+and+mood+in+dual+diagnosis+patients&author=Osuji+I.+J.&author=Vera-Bolaños+E.&author=Carmody+T.+J.&author=Brown+E.+S.&publication_year=2010&journal=Psychiatry+Res.&volume=178&pages=309-312) (http://scholar.google.com/scholar_lookup?title=Pregnenolone+for+cognition+and+mood+in+dual+diagnosis+patients&author=Osuji+I.+J.&author=Vera-Bolaños+E.&author=Carmody+T.+J.&author=Brown+E.+S.&publication_year=2010&journal=Psychiatry+Res.&volume=178&pages=309-312)

Paul, S. M., and Purdy, R. H. (1992). Neuroactive steroids. *FASEB J.* 6, 2311–2322. doi: 10.1096/fasebj.6.6.1347506

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=1347506) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=1347506>) | [CrossRef Full Text](https://doi.org/10.1096/fasebj.6.6.1347506) (<https://doi.org/10.1096/fasebj.6.6.1347506>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Neuroactive+steroids&author=Paul+S.+M.&author=Purdy+R.+H.&publication_year=1992&journal=FASEB+J.&volume=6&pages=2311-2322) (http://scholar.google.com/scholar_lookup?title=Neuroactive+steroids&author=Paul+S.+M.&author=Purdy+R.+H.&publication_year=1992&journal=FASEB+J.&volume=6&pages=2311-2322)

Peres, J. F. P., Newberg, A. B., Mercante, J. P., Simao, M., Albuquerque, V. E., Peres, M. J. P., et al. (2007). Cerebral blood flow changes during retrieval of traumatic memories before and after psychotherapy: a SPECT study. *Psychol. Med.* 37, 1481–1491. doi: 10.1017/s003329170700997x

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=17288648) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=17288648) | [CrossRef Full Text](https://doi.org/10.1017/s003329170700997x) (https://doi.org/10.1017/s003329170700997x) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Cerebral+blood+flow+changes+during+retrieval+of+traumatic+memories+before+and+after+psychotherapy%3A+a+SPECT+study&author=Peres+J.+F.+P.&author=Newberg+A.+B.&author=Mercante+J.+P.&author=Simao+M.&author=Albuquerque+V.+E.&author=Peres+M.+J.+P.&publication_year=2007&journal=Psychol.+Med.&volume=37&pages=1481-1491) (http://scholar.google.com/scholar_lookup?title=Cerebral+blood+flow+changes+during+retrieval+of+traumatic+memories+before+and+after+psychotherapy%3A+a+SPECT+study&author=Peres+J.+F.+P.&author=Newberg+A.+B.&author=Mercante+J.+P.&author=Simao+M.&author=Albuquerque+V.+E.&author=Peres+M.+J.+P.&publication_year=2007&journal=Psychol.+Med.&volume=37&pages=1481-1491)

Philip, N. S., Barredo, J., van't Wout-Frank, M., Tyrka, A. R., Price, L. H., and Carpenter, L. L. (2018). Network mechanisms of clinical response to transcranial magnetic stimulation in posttraumatic stress disorder and major depressive disorder. *Biol. Psychiatry* 83, 263–272. doi: 10.1016/j.biopsych.2017.07.021

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=28886760) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=28886760) | [CrossRef Full Text](https://doi.org/10.1016/j.biopsych.2017.07.021) (https://doi.org/10.1016/j.biopsych.2017.07.021) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Network+mechanisms+of+clinical+response+to+transcranial+magnetic+stimulation+in+posttraumatic+stress+disorder+and+major+depressive+disorder&author=Philip+N.+S.&author=Barredo+J.&author=van't+Wout-Frank+M.&author=Tyrka+A.+R.&author=Price+L.+H.&author=Carpenter+L.+L.&publication_year=2018&journal=Biol.+Psychiatry&volume=83&pages=263-272) (http://scholar.google.com/scholar_lookup?title=Network+mechanisms+of+clinical+response+to+transcranial+magnetic+stimulation+in+posttraumatic+stress+disorder+and+major+depressive+disorder&author=Philip+N.+S.&author=Barredo+J.&author=van't+Wout-Frank+M.&author=Tyrka+A.+R.&author=Price+L.+H.&author=Carpenter+L.+L.&publication_year=2018&journal=Biol.+Psychiatry&volume=83&pages=263-272)

Pitman, R. K., Rasmusson, A. M., Koenen, K. C., Shin, L. M., Orr, S. P., Gilbertson, M. W., et al. (2012). Biological studies of post-traumatic stress disorder. *Nat. Rev. Neurosci.* 13, 769–787. doi: 10.1038/nrn3339

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23047775) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23047775) | [CrossRef Full Text](https://doi.org/10.1038/nrn3339) (https://doi.org/10.1038/nrn3339) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Biological+studies+of+post-traumatic+stress+disorder&author=Pitman+R.+K.&author=Rasmusson+A.+M.&author=Koenen+K.+C.&author=Shin+L.+M.&author=Orr+S.+P.&author=Gilbertson+M.+W.&publication_year=2012&journal=Nat.+Rev.+Neurosci.&volume=13&pages=769-787) (http://scholar.google.com/scholar_lookup?title=Biological+studies+of+post-traumatic+stress+disorder&author=Pitman+R.+K.&author=Rasmusson+A.+M.&author=Koenen+K.+C.&author=Shin+L.+M.&author=Orr+S.+P.&author=Gilbertson+M.+W.&publication_year=2012&journal=Nat.+Rev.+Neurosci.&volume=13&pages=769-787)

Powers, M. B., Halpern, J. M., Ferenschak, M. P., Gillihan, S. J., and Foa, E. B. (2010). A meta-analytic review of prolonged exposure for posttraumatic stress disorder. *Clin. Psychol. Rev.* 30, 635–641. doi: 10.1016/j.cpr.2010.04.007

[CrossRef Full Text](https://doi.org/10.1016/j.cpr.2010.04.007) (https://doi.org/10.1016/j.cpr.2010.04.007) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=A+meta-analytic+review+of+prolonged+exposure+for+posttraumatic+stress+disorder&author=Powers+M.+B.&author=Halpern+J.+M.&author=Ferenschak+M.+P.&author=Gillihan+S.+J.&author=Foa+E.+B.&publication_year=2010&journal=Clin.+Psychol.+Rev.&volume=30&pages=635-641) (http://scholar.google.com/scholar_lookup?title=A+meta-analytic+review+of+prolonged+exposure+for+posttraumatic+stress+disorder&author=Powers+M.+B.&author=Halpern+J.+M.&author=Ferenschak+M.+P.&author=Gillihan+S.+J.&author=Foa+E.+B.&publication_year=2010&journal=Clin.+Psychol.+Rev.&volume=30&pages=635-641)

Quirk, G. J., Russo, G. K., Barron, J. L., and Lebron, K. (2000). The role of ventromedial prefrontal cortex in the recovery of extinguished fear. *J. Neurosci.* 20, 6225–6231. doi: 10.1523/jneurosci.20-16-06225.2000

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=10934272) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=10934272) | [CrossRef Full Text](https://doi.org/10.1523/jneurosci.20-16-06225.2000) (https://doi.org/10.1523/jneurosci.20-16-06225.2000) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=The+role+of+ventromedial+prefrontal+cortex+in+the+recovery+of+extinguished+fear&author=Quirk+G.+J.&author=Russo+G.+K.&author=Barron+J.+L.&author=Lebron+K.&publication_year=2000&journal=J.+Neurosci.&volume=20&pages=6225-6231) (http://scholar.google.com/scholar_lookup?title=The+role+of+ventromedial+prefrontal+cortex+in+the+recovery+of+extinguished+fear&author=Quirk+G.+J.&author=Russo+G.+K.&author=Barron+J.+L.&author=Lebron+K.&publication_year=2000&journal=J.+Neurosci.&volume=20&pages=6225-6231)

Rabinak, C. A., MacNamara, A., Kennedy, A. E., Angstadt, M., Stein, M. B., Liberzon, I., et al. (2014). Focal and aberrant prefrontal engagement during emotion regulation in veterans with posttraumatic stress disorder. *Depress. Anxiety* 31, 851–861. doi: 10.1002/da.22243

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24677490) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24677490) | [CrossRef Full Text](https://doi.org/10.1002/da.22243) (https://doi.org/10.1002/da.22243) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Focal+and+aberrant+prefrontal+engagement+during+emotion+regulation+in+veterans+with+posttraumatic+stress+disorder&author=Rabinak+C.+A.&author=MacNamara+A.&author=Kennedy+A.+E.&author=Angstadt+M.&author=Stein+M.+B.&author=Liberzon+I.&publication_year=2014&journal=Depress.+Anxiety&volume=31&pages=851-861) (http://scholar.google.com/scholar_lookup?title=Focal+and+aberrant+prefrontal+engagement+during+emotion+regulation+in+veterans+with+posttraumatic+stress+disorder&author=Rabinak+C.+A.&author=MacNamara+A.&author=Kennedy+A.+E.&author=Angstadt+M.&author=Stein+M.+B.&author=Liberzon+I.&publication_year=2014&journal=Depress.+Anxiety&volume=31&pages=851-861)

Ramchand, R., Rudavsky, R., Grant, S., Tanielian, T., and Jaycox, L. (2015). Prevalence of, risk factors for and consequences of posttraumatic stress disorder and other mental health problems in military populations deployed to Iraq and Afghanistan. *Curr. Psychiatry Rep.* 17:37. doi: 10.1007/s11920-015-0575-z

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=25876141) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=25876141) | [CrossRef Full Text](https://doi.org/10.1007/s11920-015-0575-z) (https://doi.org/10.1007/s11920-015-0575-z) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Prevalence+of,+risk+factors+for+and+consequences+of+posttraumatic+stress+disorder+and+other+mental+health+problems+in+military+populations+deployed+to+Iraq+and+Afghanistan&author=Ramchand+R.&author=Rudavsky+R.&author=Grant+S.&author=Tanielian+T.&author=Jaycox+L.&publication_year=2015&journal=Curr.+Psychiatry+Rep.&volume=17&pages=37) (http://scholar.google.com/scholar_lookup?title=Prevalence+of,+risk+factors+for+and+consequences+of+posttraumatic+stress+disorder+and+other+mental+health+problems+in+military+populations+deployed+to+Iraq+and+Afghanistan&author=Ramchand+R.&author=Rudavsky+R.&author=Grant+S.&author=Tanielian+T.&author=Jaycox+L.&publication_year=2015&journal=Curr.+Psychiatry+Rep.&volume=17&pages=37)

Rasmusson, A. M., Marx, C. E., Pineles, S. L., Locci, A., Scioli-Salter, E. R., Nillni, Y. I., et al. (2017). Neuroactive steroids and PTSD treatment. *Neurosci. Lett.* 649, 156–163. doi: 10.1016/j.neulet.2017.01.054

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=28215878) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=28215878) | [CrossRef Full Text](https://doi.org/10.1016/j.neulet.2017.01.054) (https://doi.org/10.1016/j.neulet.2017.01.054) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Neuroactive+steroids+and+PTSD+treatment&author=Rasmusson+A.+M.&author=Marx+C.+E.&author=Pineles+S.+L.&author=Locci+A.&author=Scioli-Salter+E.+R.&author=Nillni+Y.+I.&+&publication_year=2017&journal=Neurosci.+Lett.&volume=649&pages=156-163) (http://scholar.google.com/scholar_lookup?title=Neuroactive+steroids+and+PTSD+treatment&author=Rasmusson+A.+M.&author=Marx+C.+E.&author=Pineles+S.+L.&author=Locci+A.&author=Scioli-Salter+E.+R.&author=Nillni+Y.+I.&+&publication_year=2017&journal=Neurosci.+Lett.&volume=649&pages=156-163)

Rauch, S. A. M., Defever, E., Favorite, T., Duroe, A., Garrity, C., Martis, B., et al. (2009). Prolonged exposure for PTSD in a veterans health administration PTSD clinic. *J. Trauma. Stress* 22, 60–64. doi: 10.1002/jts.20380

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=19145643) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=19145643) | [CrossRef Full Text](https://doi.org/10.1002/jts.20380) (https://doi.org/10.1002/jts.20380) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Prolonged+exposure+for+PTSD+in+a+veterans+health+administration+PTSD+clinic&author=Rauch+S.+A.+M.&author=Defever+E.&author=Favorite+T.&author=Duroe+A.&author=Garrity+C.&author=Martis+B.&+&publication_year=2009&journal=J.+Trauma.+Stress&volume=22&pages=60-64) (http://scholar.google.com/scholar_lookup?title=Prolonged+exposure+for+PTSD+in+a+veterans+health+administration+PTSD+clinic&author=Rauch+S.+A.+M.&author=Defever+E.&author=Favorite+T.&author=Duroe+A.&author=Garrity+C.&author=Martis+B.&+&publication_year=2009&journal=J.+Trauma.+Stress&volume=22&pages=60-64)

Rauch, S. A. M., Simon, N. M., Kim, H. M., Acierno, R., King, A. P., Norman, S. B., et al. (2018). Integrating biological treatment mechanisms into randomized clinical trials: design of PROGRESS (PROlonged ExpoSure and Sertraline Trial). *Contemp. Clin. Trials* 64, 128–138. doi: 10.1016/j.cct.2017.10.013

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=29081351) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=29081351) | [CrossRef Full Text](https://doi.org/10.1016/j.cct.2017.10.013) (https://doi.org/10.1016/j.cct.2017.10.013) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Integrating+biological+treatment+mechanisms+into+randomized+clinical+trials%3A+design+of+PROGRESS+(PROlonged+ExpoSure+and+Sertraline+Trial)&author=Rauch+S.+A.+M.&author=Simon+N.+M.&author=Kim+H.+M.&author=Acierno+R.&author=King+A.+P.&author=Norman+S.+B.&+&publication_year=2018&journal=Contemp.+Clin.+Trials&volume=64&pages=128-138) (http://scholar.google.com/scholar_lookup?title=Integrating+biological+treatment+mechanisms+into+randomized+clinical+trials%3A+design+of+PROGRESS+(PROlonged+ExpoSure+and+Sertraline+Trial)&author=Rauch+S.+A.+M.&author=Simon+N.+M.&author=Kim+H.+M.&author=Acierno+R.&author=King+A.+P.&author=Norman+S.+B.&+&publication_year=2018&journal=Contemp.+Clin.+Trials&volume=64&pages=128-138)

Rauch, S. A., Hembree, E. A., and Foa, E. B. (2001). Acute psychosocial preventive interventions for posttraumatic stress disorder. *Adv. Mind Body Med.* 17, 187–191. doi: 10.1054/ambm.2000.0312

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=11572846) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=11572846) | [CrossRef Full Text](https://doi.org/10.1054/ambm.2000.0312) (https://doi.org/10.1054/ambm.2000.0312) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Acute+psychosocial+preventive+interventions+for+posttraumatic+stress+disorder&author=Rauch+S.+A.&author=Hembree+E.+A.&author=Foa+E.+B.&publication_year=2001&journal=Adv.+Mind+Body+Med.&volume=17&pages=187-191) (http://scholar.google.com/scholar_lookup?title=Acute+psychosocial+preventive+interventions+for+posttraumatic+stress+disorder&author=Rauch+S.+A.&author=Hembree+E.+A.&author=Foa+E.+B.&publication_year=2001&journal=Adv.+Mind+Body+Med.&volume=17&pages=187-191)

Rauch, S. L., Shin, L. M., and Phelps, E. A. (2006). Neurocircuitry models of posttraumatic stress disorder and extinction: human neuroimaging research—past, present and future. *Biol. Psychiatry* 60, 376–382. doi: 10.1016/j.biopsych.2006.06.004

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=16919525) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=16919525) | [CrossRef Full Text](https://doi.org/10.1016/j.biopsych.2006.06.004) (https://doi.org/10.1016/j.biopsych.2006.06.004) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Neurocircuitry+models+of+posttraumatic+stress+disor-) (http://scholar.google.com/scholar_lookup?title=Neurocircuitry+models+of+posttraumatic+stress+disor-

der+and+extinction%3A+human+neuroimaging+research—past,+present+and+future&author=Rauch+S.+L.&author=Shin+L.+M.&author=Phelps+E.+A.&publication_year=2006&journal=Biol.+Psychiatry&volume=60&pages=376-382)

Rauch, S., and Foa, E. (2006). Emotional processing theory (EPT) and exposure therapy for PTSD. *J. Contemp. Psychother.* 36, 61–65. doi: 10.1007/s10879-006-9008-y

[CrossRef Full Text](https://doi.org/10.1007/s10879-006-9008-y) (<https://doi.org/10.1007/s10879-006-9008-y>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Emotional+processing+theory+(EPT)+and+exposure+therapy+for+PTSD&author=Rauch+S.&author=Foa+E.&publication_year=2006&journal=J.+Contemp.+Psychother.&volume=36&pages=61-65) ([http://scholar.google.com/scholar_lookup?title=Emotional+processing+theory+\(EPT\)+and+exposure+therapy+for+PTSD&author=Rauch+S.&author=Foa+E.&publication_year=2006&journal=J.+Contemp.+Psychother.&volume=36&pages=61-65](http://scholar.google.com/scholar_lookup?title=Emotional+processing+theory+(EPT)+and+exposure+therapy+for+PTSD&author=Rauch+S.&author=Foa+E.&publication_year=2006&journal=J.+Contemp.+Psychother.&volume=36&pages=61-65))

Rauch, S., and Liberzon, I. (2016). "Mechanisms of action in psychotherapy," in *The Biological Basis of Posttraumatic Stress Disorder: From Brain to Mind*, eds I. Liberzon, and K. Ressler (New York, NY: Oxford University Press), 353–372.

Riaza Bermudo-Soriano, C., Perez-Rodriguez, M. M., Vaquero-Lorenzo, C., and Baca-Garcia, E. (2012). New perspectives in glutamate and anxiety. *Pharmacol. Biochem. Behav.* 100, 752–774. doi: 10.1016/j.pbb.2011.04.010

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=21569789) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=21569789>) | [CrossRef Full Text](https://doi.org/10.1016/j.pbb.2011.04.010) (<https://doi.org/10.1016/j.pbb.2011.04.010>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=New+perspectives+in+glutamate+and+anxiety&author=Riaza+Bermudo-Soriano+C.&author=Perez-Rodriguez+M.+M.&author=Vaquero-Lorenzo+C.&author=Baca-Garcia+E.&publication_year=2012&journal=Pharmacol.+Biochem.+Behav.&volume=100&pages=752-774) (http://scholar.google.com/scholar_lookup?title=New+perspectives+in+glutamate+and+anxiety&author=Riaza+Bermudo-Soriano+C.&author=Perez-Rodriguez+M.+M.&author=Vaquero-Lorenzo+C.&author=Baca-Garcia+E.&publication_year=2012&journal=Pharmacol.+Biochem.+Behav.&volume=100&pages=752-774)

Rothbaum, B. O., Astin, M. C., and Marsteller, F. (2005). Prolonged exposure versus eye movement desensitization and reprocessing (EMDR) for PTSD rape victims. *J. Trauma. Stress* 18, 607–616. doi: 10.1002/jts.20069

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=16382428) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=16382428>) | [CrossRef Full Text](https://doi.org/10.1002/jts.20069) (<https://doi.org/10.1002/jts.20069>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Prolonged+exposure+versus+eye+movement+desensitization+and+reprocessing+(EMDR)+for+PTSD+rape+victims&author=Rothbaum+B.+O.&author=Astin+M.+C.&author=Marsteller+F.&publication_year=2005&journal=J.+Trauma.+Stress&volume=18&pages=607-616) ([http://scholar.google.com/scholar_lookup?title=Prolonged+exposure+versus+eye+movement+desensitization+and+reprocessing+\(EMDR\)+for+PTSD+rape+victims&author=Rothbaum+B.+O.&author=Astin+M.+C.&author=Marsteller+F.&publication_year=2005&journal=J.+Trauma.+Stress&volume=18&pages=607-616](http://scholar.google.com/scholar_lookup?title=Prolonged+exposure+versus+eye+movement+desensitization+and+reprocessing+(EMDR)+for+PTSD+rape+victims&author=Rothbaum+B.+O.&author=Astin+M.+C.&author=Marsteller+F.&publication_year=2005&journal=J.+Trauma.+Stress&volume=18&pages=607-616))

Rothbaum, B. O., Cahill, S. P., Foa, E. B., Davidson, J. R., Compton, J., Connor, K. M., et al. (2006). Augmentation of sertraline with prolonged exposure in the treatment of posttraumatic stress disorder. *J. Traumatic Stress* 19, 625–638. doi: 10.1002/jts.20170

[CrossRef Full Text](https://doi.org/10.1002/jts.20170) (<https://doi.org/10.1002/jts.20170>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Augmentation+of+sertraline+with+prolonged+exposure+in+the+treatment+of+posttraumatic+stress+disorder&author=Rothbaum+B.+O.&author=Cahill+S.+P.&author=Foa+E.+B.&author=Davidson+J.+R.&author=Compton+J.&author=Connor+K.+M.&publication_year=2006&journal=J.+Traumatic+Stress&volume=19&pages=625-638) (http://scholar.google.com/scholar_lookup?title=Augmentation+of+sertraline+with+prolonged+exposure+in+the+treatment+of+posttraumatic+stress+disorder&author=Rothbaum+B.+O.&author=Cahill+S.+P.&author=Foa+E.+B.&author=Davidson+J.+R.&author=Compton+J.&author=Connor+K.+M.&publication_year=2006&journal=J.+Traumatic+Stress&volume=19&pages=625-638)

Rothbaum, B. O., and Davis, M. (2003). Applying learning principles to the treatment of post-trauma reactions. *Ann. N Y Acad. Sci.* 1008, 112–121. doi: 10.1196/annals.1301.012

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=14998877) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=14998877>) | [CrossRef Full Text](https://doi.org/10.1196/annals.1301.012) (<https://doi.org/10.1196/annals.1301.012>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Applying+learning+principles+to+the+treatment+of+post-trauma+reactions&author=Rothbaum+B.+O.&author=Davis+M.&publication_year=2003&journal=Ann.+N+Y+Acad.+Sci.&volume=1008&pages=112-121) (http://scholar.google.com/scholar_lookup?title=Applying+learning+principles+to+the+treatment+of+post-trauma+reactions&author=Rothbaum+B.+O.&author=Davis+M.&publication_year=2003&journal=Ann.+N+Y+Acad.+Sci.&volume=1008&pages=112-121)

Rothbaum, B. O., Price, M., Jovanovic, T., Norrholm, S. D., Gerardi, M., Dunlop, B., et al. (2014). A randomized, double-blind evaluation of D-cycloserine or alprazolam combined with virtual reality exposure therapy for posttraumatic stress disorder in Iraq and Afghanistan war veterans. *Am. J. Psychiatry* 171, 640–648. doi: 10.1176/appi.ajp.2014.13121625

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24743802) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24743802>) | [CrossRef Full Text](https://doi.org/10.1176/appi.ajp.2014.13121625) (<https://doi.org/10.1176/appi.ajp.2014.13121625>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=A+randomized,+double-blind+evaluation+of+D-cycloserine+or+alprazolam+combined+with+virtual+reality+exposure+therapy+for+posttraumatic+stress+disor) (http://scholar.google.com/scholar_lookup?title=A+randomized,+double-blind+evaluation+of+D-cycloserine+or+alprazolam+combined+with+virtual+reality+exposure+therapy+for+posttraumatic+stress+disor

der+in+Iraq+and+Afghanistan+war+veterans&author=Rothbaum+B.+O.&author=Price+M.&author=Jovanovic+T.&author=Norrholm+S.+D.&author=Gerardi+M.&author=Dunlop+B.&+&publication_year=2014&journal=Am.+J.+Psychiatry&volume=171&pages=640-648)

Rougemont-Bücking, A., Linnman, C., Zeffiro, T. A., Zeidan, M. A., Lebron-Milad, K., Rodriguez-Romaguera, J., et al. (2011). Altered processing of contextual information during fear extinction in PTSD: an fMRI study. *CNS Neurosci. Ther.* 17, 227–236. doi: 10.1111/j.1755-5949.2010.00152.x

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=20406268) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=20406268>) | [CrossRef Full Text](https://doi.org/10.1111/j.1755-5949.2010.00152.x) (<https://doi.org/10.1111/j.1755-5949.2010.00152.x>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Altered+processing+of+contextual+information+during+fear+extinction+in+PTSD%3A+an+fMRI+study&author=Rougemont-B%C3%BCcking+A.&author=Linnman+C.&author=Zeffiro+T.+A.&author=Zeidan+M.+A.&author=Lebron-Milad+K.&author=Rodriguez-Romaguera+J.&+&publication_year=2011&journal=CNS+Neurosci.+Ther.&volume=17&pages=227-236) ([http://scholar.google.com/scholar_lookup?title=Altered+processing+of+contextual+information+during+fear+extinction+in+PTSD%3A+an+fMRI+study&author=Rougemont-Bücking+A.&author=Linnman+C.&author=Zeffiro+T.+A.&author=Zeidan+M.+A.&author=Lebron-Milad+K.&author=Rodriguez-Romaguera+J.&+&publication_year=2011&journal=CNS+Neurosci.+Ther.&volume=17&pages=227-236](http://scholar.google.com/scholar_lookup?title=Altered+processing+of+contextual+information+during+fear+extinction+in+PTSD%3A+an+fMRI+study&author=Rougemont-B%C3%BCcking+A.&author=Linnman+C.&author=Zeffiro+T.+A.&author=Zeidan+M.+A.&author=Lebron-Milad+K.&author=Rodriguez-Romaguera+J.&+&publication_year=2011&journal=CNS+Neurosci.+Ther.&volume=17&pages=227-236))

Rubin, M., Shvil, E., Papini, S., Chhetry, B. T., Helpman, L., Markowitz, J. C., et al. (2016). Greater hippocampal volume is associated with PTSD treatment response. *Psychiatry Res. Neuroimaging* 252, 36–39. doi: 10.1016/j.pscychresns.2016.05.001

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=27179314) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=27179314>) | [CrossRef Full Text](https://doi.org/10.1016/j.pscychresns.2016.05.001) (<https://doi.org/10.1016/j.pscychresns.2016.05.001>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Greater+hippocampal+volume+is+associated+with+PTSD+treatment+response&author=Rubin+M.&author=Shvil+E.&author=Papini+S.&author=Chhetry+B.+T.&author=Helpman+L.&author=Markowitz+J.+C.&+&publication_year=2016&journal=Psychiatry+Res.+Neuroimaging&volume=252&pages=36-39) (http://scholar.google.com/scholar_lookup?title=Greater+hippocampal+volume+is+associated+with+PTSD+treatment+response&author=Rubin+M.&author=Shvil+E.&author=Papini+S.&author=Chhetry+B.+T.&author=Helpman+L.&author=Markowitz+J.+C.&+&publication_year=2016&journal=Psychiatry+Res.+Neuroimaging&volume=252&pages=36-39)

Scioli-Salter, E., Forman, D. E., Otis, J. D., Tun, C., Allsup, K., Marx, C. E., et al. (2016). Potential neurobiological benefits of exercise in chronic pain and posttraumatic stress disorder: Pilot study. *J. Rehabil. Res. Dev.* 53, 95–106. doi: 10.1682/jrrd.2014.10.0267

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=27006290) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=27006290>) | [CrossRef Full Text](https://doi.org/10.1682/jrrd.2014.10.0267) (<https://doi.org/10.1682/jrrd.2014.10.0267>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Potential+neurobiological+benefits+of+exercise+in+chronic+pain+and+posttraumatic+stress+disorder%3A+Pilot+study&author=Scioli-Salter+E.&author=Forman+D.+E.&author=Otis+J.+D.&author=Tun+C.&author=Allsup+K.&author=Marx+C.+E.&+&publication_year=2016&journal=J.+Rehabil.+Res.+Dev.&volume=53&pages=95-106) (http://scholar.google.com/scholar_lookup?title=Potential+neurobiological+benefits+of+exercise+in+chronic+pain+and+posttraumatic+stress+disorder%3A+Pilot+study&author=Scioli-Salter+E.&author=Forman+D.+E.&author=Otis+J.+D.&author=Tun+C.&author=Allsup+K.&author=Marx+C.+E.&+&publication_year=2016&journal=J.+Rehabil.+Res.+Dev.&volume=53&pages=95-106)

Shepherd, L., and Wild, J. (2014). Emotion regulation, physiological arousal and PTSD symptoms in trauma-exposed individuals. *J. Behav. Ther. Exp. Psychiatry* 45, 360–367. doi: 10.1016/j.jbtep.2014.03.002

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24727342) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24727342>) | [CrossRef Full Text](https://doi.org/10.1016/j.jbtep.2014.03.002) (<https://doi.org/10.1016/j.jbtep.2014.03.002>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Emotion+regulation,+physiological+arousal+and+PTSD+symptoms+in+trauma-exposed+individuals&author=Shepherd+L.&author=Wild+J.&+&publication_year=2014&journal=J.+Behav.+Ther.+Exp.+Psychiatry&volume=45&pages=360-367) (http://scholar.google.com/scholar_lookup?title=Emotion+regulation,+physiological+arousal+and+PTSD+symptoms+in+trauma-exposed+individuals&author=Shepherd+L.&author=Wild+J.&+&publication_year=2014&journal=J.+Behav.+Ther.+Exp.+Psychiatry&volume=45&pages=360-367)

Sheynin, J., and Liberzon, I. (2017). Circuit dysregulation and circuit-based treatments in posttraumatic stress disorder. *Neurosci. Lett.* 649, 133–138. doi: 10.1016/j.neulet.2016.11.014

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=27845239) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=27845239>) | [CrossRef Full Text](https://doi.org/10.1016/j.neulet.2016.11.014) (<https://doi.org/10.1016/j.neulet.2016.11.014>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Circuit+dysregulation+and+circuit-based+treatments+in+posttraumatic+stress+disorder&author=Sheynin+J.&author=Liberzon+I.&+&publication_year=2017&journal=Neurosci.+Lett.&volume=649&pages=133-138) (http://scholar.google.com/scholar_lookup?title=Circuit+dysregulation+and+circuit-based+treatments+in+posttraumatic+stress+disorder&author=Sheynin+J.&author=Liberzon+I.&+&publication_year=2017&journal=Neurosci.+Lett.&volume=649&pages=133-138)

Shin, L. M., and Handwerker, K. (2009). Is posttraumatic stress disorder a stress-induced fear circuitry disorder? *J. Trauma. Stress* 22, 409–415. doi: 10.1002/jts.20442

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=19743481) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=19743481) | [CrossRef Full Text](https://doi.org/10.1002/jts.20442) (https://doi.org/10.1002/jts.20442) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Is+posttraumatic+stress+disorder+a+stress-induced+fear+circuitry+disorder%3F&author=Shin+L.+M.&author=Handwerker+K.&publication_year=2009&journal=J.+Trauma.+Stress&volume=22&pages=409-415) (http://scholar.google.com/scholar_lookup?title=Is+posttraumatic+stress+disorder+a+stress-induced+fear+circuitry+disorder%3F&author=Shin+L.+M.&author=Handwerker+K.&publication_year=2009&journal=J.+Trauma.+Stress&volume=22&pages=409-415)

Shin, L. M., and Liberzon, I. (2010). The neurocircuitry of fear, stress, and anxiety disorders. *Neuropsychopharmacology* 35, 169–191. doi: 10.1038/npp.2009.83

[CrossRef Full Text](https://doi.org/10.1038/npp.2009.83) (https://doi.org/10.1038/npp.2009.83) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=The+neurocircuitry+of+fear,+stress,+and+anxiety+disorders&author=Shin+L.+M.&author=Liberzon+I.&publication_year=2010&journal=Neuropsychopharmacology&volume=35) (http://scholar.google.com/scholar_lookup?title=The+neurocircuitry+of+fear,+stress,+and+anxiety+disorders&author=Shin+L.+M.&author=Liberzon+I.&publication_year=2010&journal=Neuropsychopharmacology&volume=35)

Shin, L. M., Rauch, S. L., and Pitman, R. K. (2006). Amygdala, medial prefrontal cortex and hippocampal function in PTSD. In *Ann. N Y Acad. Sci.* 1071, 67–79. doi: 10.1196/annals.1364.007

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=16891563) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=16891563) | [CrossRef Full Text](https://doi.org/10.1196/annals.1364.007) (https://doi.org/10.1196/annals.1364.007) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Amygdala,+medial+prefrontal+cortex+and+hippocampal+function+in+PTSD.+In&author=Shin+L.+M.&author=Rauch+S.+L.&author=Pitman+R.+K.&publication_year=2006&journal=Ann.+N+Y+Acad.+Sci.&volume=1071&pages=67-79) (http://scholar.google.com/scholar_lookup?title=Amygdala,+medial+prefrontal+cortex+and+hippocampal+function+in+PTSD.+In&author=Shin+L.+M.&author=Rauch+S.+L.&author=Pitman+R.+K.&publication_year=2006&journal=Ann.+N+Y+Acad.+Sci.&volume=1071&pages=67-79)

Shukla, S. K., Murali, N. S., and Brilliant, M. H. (2015). Personalized medicine going precise: from genomics to microbiomics. *Trends Mol. Med.* 21,461–462. doi: 10.1016/j.molmed.2015.06.002

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=26129865) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=26129865) | [CrossRef Full Text](https://doi.org/10.1016/j.molmed.2015.06.002) (https://doi.org/10.1016/j.molmed.2015.06.002) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Personalized+medicine+going+precise%3A+from+genomics+to+microbiomics&author=Shukla+S.+K.&author=Murali+N.+S.&author=Brilliant+M.+H.&publication_year=2015&journal=Trends+Mol.+Med.&volume=21&pages=461-462) (http://scholar.google.com/scholar_lookup?title=Personalized+medicine+going+precise%3A+from+genomics+to+microbiomics&author=Shukla+S.+K.&author=Murali+N.+S.&author=Brilliant+M.+H.&publication_year=2015&journal=Trends+Mol.+Med.&volume=21&pages=461-462)

Shvil, E., Rusch, H. L., Sullivan, G. M., and Neria, Y. (2013). Neural, psychophysiological and behavioral markers of fear processing in PTSD: a review of the literature. *Curr. Psychiatry Rep.* 15:358. doi: 10.1007/s11920-013-0358-3

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23619614) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23619614) | [CrossRef Full Text](https://doi.org/10.1007/s11920-013-0358-3) (https://doi.org/10.1007/s11920-013-0358-3) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Neural,+psychophysiological+and+behavioral+markers+of+fear+processing+in+PTSD%3A+a+review+of+the+literature&author=Shvil+E.&author=Rusch+H.+L.&author=Sullivan+G.+M.&author=Neria+Y.&publication_year=2013&journal=Curr.+Psychiatry+Rep.&volume=15&pages=358) (http://scholar.google.com/scholar_lookup?title=Neural,+psychophysiological+and+behavioral+markers+of+fear+processing+in+PTSD%3A+a+review+of+the+literature&author=Shvil+E.&author=Rusch+H.+L.&author=Sullivan+G.+M.&author=Neria+Y.&publication_year=2013&journal=Curr.+Psychiatry+Rep.&volume=15&pages=358)

Shvil, E., Sullivan, G. M., Schafer, S., Markowitz, J. C., Campeas, M., Wager, T. D., et al. (2014). Sex differences in extinction recall in posttraumatic stress disorder: a pilot fMRI study. *Neurobiol. Learn. Mem.* 113, 101–108. doi: 10.1016/j.nlm.2014.02.003

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24560771) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24560771) | [CrossRef Full Text](https://doi.org/10.1016/j.nlm.2014.02.003) (https://doi.org/10.1016/j.nlm.2014.02.003) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Sex+differences+in+extinction+recall+in+posttraumatic+stress+disorder%3A+a+pilot+fMRI+study&author=Shvil+E.&author=Sullivan+G.+M.&author=Schafer+S.&author=Markowitz+J.+C.&author=Campeas+M.&author=Wager+T.+D.&publication_year=2014&journal=Neurobiol.+Learn.+Mem.&volume=113&pages=101-108) (http://scholar.google.com/scholar_lookup?title=Sex+differences+in+extinction+recall+in+posttraumatic+stress+disorder%3A+a+pilot+fMRI+study&author=Shvil+E.&author=Sullivan+G.+M.&author=Schafer+S.&author=Markowitz+J.+C.&author=Campeas+M.&author=Wager+T.+D.&publication_year=2014&journal=Neurobiol.+Learn.+Mem.&volume=113&pages=101-108)

Simon, N. M., Connor, K. M., Lang, A. J., Rauch, S., Krulewicz, S., LeBeau, R. T., et al. (2008). Paroxetine CR augmentation for posttraumatic stress disorder refractory to prolonged exposure therapy. *J. Clin. psychiatry* 69, 400–405.

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=18348595) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=18348595) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Paroxetine+CR+augmentation+for+posttraumatic+stress+disorder+refractory+to+prolonged+exposure+therapy&author=Simon+N.+M.&au) (http://scholar.google.com/scholar_lookup?title=Paroxetine+CR+augmentation+for+posttraumatic+stress+disorder+refractory+to+prolonged+exposure+therapy&author=Simon+N.+M.&au

thor=Connor+K.+M.&author=Lang+A.+J.&author=Rauch+S.&author=Krulwicz+S.&author=LeBeau+R.+T.&&publication_year=2008&journal=J.+Clin.+psychiatry&volume=69&pages=400-405)

Smits, J. A. J., Rosenfield, D., Otto, M. W., Marques, L., Davis, M. L., Meuret, A. E., et al. (2013a). D-cycloserine enhancement of exposure therapy for social anxiety disorder depends on the success of exposure sessions. *J. Psychiatr. Res.* 47, 1455–1461. doi: 10.1016/j.jpsychires.2013.06.020

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23870811) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23870811) | [CrossRef Full Text](https://doi.org/10.1016/j.jpsychires.2013.06.020) (https://doi.org/10.1016/j.jpsychires.2013.06.020) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=D-cycloserine+enhancement+of+exposure+therapy+for+social+anxiety+disorder+depends+on+the+success+of+exposure+sessions&author=Smits+J.+A.+J.&author=Rosenfield+D.&author=Otto+M.+W.&author=Marques+L.&author=Davis+M.+L.&author=Meuret+A.+E.&&publication_year=2013a&journal=J.+Psychiatr.+Res.&volume=47&pages=1455-1461) (http://scholar.google.com/scholar_lookup?title=D-cycloserine+enhancement+of+exposure+therapy+for+social+anxiety+disorder+depends+on+the+success+of+exposure+sessions&author=Smits+J.+A.+J.&author=Rosenfield+D.&author=Otto+M.+W.&author=Marques+L.&author=Davis+M.+L.&author=Meuret+A.+E.&&publication_year=2013a&journal=J.+Psychiatr.+Res.&volume=47&pages=1455-1461)

Smits, J. A. J., Rosenfield, D., Otto, M. W., Powers, M. B., Hofmann, S. G., Telch, M. J., et al. (2013b). D-cycloserine enhancement of fear extinction is specific to successful exposure sessions: evidence from the treatment of height phobia. *Biol. Psychiatry* 73, 1054–1058. doi: 10.1016/j.biopsych.2012.12.009

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23332511) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23332511) | [CrossRef Full Text](https://doi.org/10.1016/j.biopsych.2012.12.009) (https://doi.org/10.1016/j.biopsych.2012.12.009) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=D-cycloserine+enhancement+of+fear+extinction+is+specific+to+successful+exposure+sessions%3A+evidence+from+the+treatment+of+height+phobia&author=Smits+J.+A.+J.&author=Rosenfield+D.&author=Otto+M.+W.&author=Powers+M.+B.&author=Hofmann+S.+G.&author=Telch+M.+J.&&publication_year=2013b&journal=Biol.+Psychiatry&volume=73&pages=1054-1058) (http://scholar.google.com/scholar_lookup?title=D-cycloserine+enhancement+of+fear+extinction+is+specific+to+successful+exposure+sessions%3A+evidence+from+the+treatment+of+height+phobia&author=Smits+J.+A.+J.&author=Rosenfield+D.&author=Otto+M.+W.&author=Powers+M.+B.&author=Hofmann+S.+G.&author=Telch+M.+J.&&publication_year=2013b&journal=Biol.+Psychiatry&volume=73&pages=1054-1058)

Sripada, R. K., Garfinkel, S. N., and Liberzon, I. (2013a). Avoidant symptoms in PTSD predict fear circuit activation during multimodal fear extinction. *Front. Hum. Neurosci.* 7:672. doi: 10.3389/fnhum.2013.00672

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24146643) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24146643) | [CrossRef Full Text](https://doi.org/10.3389/fnhum.2013.00672) (https://doi.org/10.3389/fnhum.2013.00672) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Avoidant+symptoms+in+PTSD+predict+fear+circuit+activation+during+multimodal+fear+extinction&author=Sripada+R.+K.&author=Garfinkel+S.+N.&author=Liberzon+I.&publication_year=2013a&journal=Front.+Hum.+Neurosci.&volume=7&pages=672) (http://scholar.google.com/scholar_lookup?title=Avoidant+symptoms+in+PTSD+predict+fear+circuit+activation+during+multimodal+fear+extinction&author=Sripada+R.+K.&author=Garfinkel+S.+N.&author=Liberzon+I.&publication_year=2013a&journal=Front.+Hum.+Neurosci.&volume=7&pages=672)

Sripada, R. K., Marx, C. E., King, A. P., Rajaram, N., Garfinkel, S. N., Abelson, J. L., et al. (2013b). DHEA enhances emotion regulation neurocircuits and modulates memory for emotional stimuli. *Neuropsychopharmacology* 38, 1798–1807. doi: 10.1038/npp.2013.79

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23552182) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23552182) | [CrossRef Full Text](https://doi.org/10.1038/npp.2013.79) (https://doi.org/10.1038/npp.2013.79) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=DHEA+enhances+emotion+regulation+neurocircuits+and+modulates+memory+for+emotional+stimuli&author=Sripada+R.+K.&author=Marx+C.+E.&author=King+A.+P.&author=Rajaram+N.&author=Garfinkel+S.+N.&author=Abelson+J.+L.&&publication_year=2013b&journal=Neuropsychopharmacology&volume=38&pages=1798-1807) (http://scholar.google.com/scholar_lookup?title=DHEA+enhances+emotion+regulation+neurocircuits+and+modulates+memory+for+emotional+stimuli&author=Sripada+R.+K.&author=Marx+C.+E.&author=King+A.+P.&author=Rajaram+N.&author=Garfinkel+S.+N.&author=Abelson+J.+L.&&publication_year=2013b&journal=Neuropsychopharmacology&volume=38&pages=1798-1807)

Sripada, R. K., Marx, C. E., King, A. P., Rampton, J. C., Ho, S. S., and Liberzon, I. (2013c). Allopregnanolone elevations following pregnenolone administration are associated with enhanced activation of emotion regulation neurocircuits. *Biol. Psychiatry* 73, 1045–1053. doi: 10.1016/j.biopsych.2012.12.008

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23348009) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23348009) | [CrossRef Full Text](https://doi.org/10.1016/j.biopsych.2012.12.008) (https://doi.org/10.1016/j.biopsych.2012.12.008) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Allopregnanolone+elevations+following+pregnenolone+administration+are+associated+with+enhanced+activation+of+emotion+regulation+neurocircuits&author=Sripada+R.+K.&author=Marx+C.+E.&author=King+A.+P.&author=Rampton+J.+C.&author=Ho+S.+S.&author=Liberzon+I.&publication_year=2013c&journal=Biol.+Psychiatry&volume=73&pages=1045-1053) (http://scholar.google.com/scholar_lookup?title=Allopregnanolone+elevations+following+pregnenolone+administration+are+associated+with+enhanced+activation+of+emotion+regulation+neurocircuits&author=Sripada+R.+K.&author=Marx+C.+E.&author=King+A.+P.&author=Rampton+J.+C.&author=Ho+S.+S.&author=Liberzon+I.&publication_year=2013c&journal=Biol.+Psychiatry&volume=73&pages=1045-1053)

Sripada, R. K., and Rauch, S. A. M. (2015). Between-session and within-session habituation in prolonged exposure therapy for posttraumatic stress disorder: a hierarchical linear modeling approach. *J. Anxiety Disord.* 30, 81–87. doi: 10.1016/j.janxdis.2015.01.002

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=25613235) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=25613235) | [CrossRef Full Text](https://doi.org/10.1016/j.janxdis.2015.01.002) (https://doi.org/10.1016/j.janxdis.2015.01.002) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Between-session+and+within-session+habituation+in+prolonged+exposure+therapy+for+posttraumatic+stress+disorder%3A+a+hierarchical+linear+modeling+approach&author=Sripada+R.+K.&author=Rauch+S.+A.+M.&publication_year=2015&journal=J.+Anxiety+Disord.&volume=30&pages=81-87) (http://scholar.google.com/scholar_lookup?title=Between-session+and+within-session+habituation+in+prolonged+exposure+therapy+for+posttraumatic+stress+disorder%3A+a+hierarchical+linear+modeling+approach&author=Sripada+R.+K.&author=Rauch+S.+A.+M.&publication_year=2015&journal=J.+Anxiety+Disord.&volume=30&pages=81-87)

Stevens, J. S., Jovanovic, T., Fani, N., Ely, T. D., Glover, E. M., Bradley, B., et al. (2013). Disrupted amygdala-prefrontal functional connectivity in civilian women with posttraumatic stress disorder. *J. Psychiatr. Res.* 47, 1469–1478. doi: 10.1016/j.jpsychires.2013.05.031

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23827769) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23827769) | [CrossRef Full Text](https://doi.org/10.1016/j.jpsychires.2013.05.031) (https://doi.org/10.1016/j.jpsychires.2013.05.031) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Disrupted+amygdala-prefrontal+functional+connectivity+in+civilian+women+with+posttraumatic+stress+disorder&author=Stevens+J.+S.&author=Jovanovic+T.&author=Fani+N.&author=Ely+T.+D.&author=Glover+E.+M.&author=Bradley+B.&&publication_year=2013&journal=J.+Psychiatr.+Res.&volume=47&pages=1469-1478) (http://scholar.google.com/scholar_lookup?title=Disrupted+amygdala-prefrontal+functional+connectivity+in+civilian+women+with+posttraumatic+stress+disorder&author=Stevens+J.+S.&author=Jovanovic+T.&author=Fani+N.&author=Ely+T.+D.&author=Glover+E.+M.&author=Bradley+B.&&publication_year=2013&journal=J.+Psychiatr.+Res.&volume=47&pages=1469-1478)

Tart, C. D., Handelsman, P. R., Deboer, L. B., Rosenfield, D., Pollack, M. H., Hofmann, S. G., et al. (2013). Augmentation of exposure therapy with post-session administration of d-cycloserine. *J. Psychiatr. Res.* 47, 168–174. doi: 10.1016/j.jpsychires.2012.09.024

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23098672) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23098672) | [CrossRef Full Text](https://doi.org/10.1016/j.jpsychires.2012.09.024) (https://doi.org/10.1016/j.jpsychires.2012.09.024) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Augmentation+of+exposure+therapy+with+post-session+administration+of+d-cycloserine&author=Tart+C.+D.&author=Handelsman+P.+R.&author=Deboer+L.+B.&author=Rosenfield+D.&author=Pollack+M.+H.&author=Hofmann+S.+G.&&publication_year=2013&journal=J.+Psychiatr.+Res.&volume=47&pages=168-174) (http://scholar.google.com/scholar_lookup?title=Augmentation+of+exposure+therapy+with+post-session+administration+of+d-cycloserine&author=Tart+C.+D.&author=Handelsman+P.+R.&author=Deboer+L.+B.&author=Rosenfield+D.&author=Pollack+M.+H.&author=Hofmann+S.+G.&&publication_year=2013&journal=J.+Psychiatr.+Res.&volume=47&pages=168-174)

Thal, S. B., and Lommen, M. J. J. (2018). Current perspective on MDMA-assisted psychotherapy for posttraumatic stress disorder. *J. Contemp. Psychother.* 48, 99–108. doi: 10.1007/s10879-018-9382-2

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=29720767) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=29720767) | [CrossRef Full Text](https://doi.org/10.1007/s10879-018-9382-2) (https://doi.org/10.1007/s10879-018-9382-2) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Current+perspective+on+MDMA-assisted+psychotherapy+for+posttraumatic+stress+disorder&author=Thal+S.+B.&author=Lommen+M.+J.+J.&publication_year=2018&journal=J.+Contemp.+Psychother.&volume=48&pages=99-108) (http://scholar.google.com/scholar_lookup?title=Current+perspective+on+MDMA-assisted+psychotherapy+for+posttraumatic+stress+disorder&author=Thal+S.+B.&author=Lommen+M.+J.+J.&publication_year=2018&journal=J.+Contemp.+Psychother.&volume=48&pages=99-108)

Thompson, R. A. (2008). Emotion regulation: a theme in search of definition. *Monogr. Soc. Res. Child Dev.* 59, 25–52. doi: 10.1111/j.1540-5834.1994.tb01276.x

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=7984164) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=7984164) | [CrossRef Full Text](https://doi.org/10.1111/j.1540-5834.1994.tb01276.x) (https://doi.org/10.1111/j.1540-5834.1994.tb01276.x) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Emotion+regulation%3A+a+theme+in+search+of+definition&author=Thompson+R.+A.&publication_year=2008&journal=Monogr.+Soc.+Res.+Child+Dev.&volume=59&pages=25-52) (http://scholar.google.com/scholar_lookup?title=Emotion+regulation%3A+a+theme+in+search+of+definition&author=Thompson+R.+A.&publication_year=2008&journal=Monogr.+Soc.+Res.+Child+Dev.&volume=59&pages=25-52)

Tuerk, P. W. (2014). Starting from something: augmenting exposure therapy and methods of inquiry. *Am. J. Psychiatry* 171, 1034–1037. doi: 10.1176/appi.ajp.2014.14070880

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=25272341) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=25272341) | [CrossRef Full Text](https://doi.org/10.1176/appi.ajp.2014.14070880) (https://doi.org/10.1176/appi.ajp.2014.14070880) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Starting+from+something%3A+augmenting+exposure+therapy+and+methods+of+inquiry&author=Tuerk+P.+W.&publication_year=2014&journal=Am.+J.+Psychiatry&volume=171&pages=1034-1037) (http://scholar.google.com/scholar_lookup?title=Starting+from+something%3A+augmenting+exposure+therapy+and+methods+of+inquiry&author=Tuerk+P.+W.&publication_year=2014&journal=Am.+J.+Psychiatry&volume=171&pages=1034-1037)

van Rooij, S. J. H., Kennis, M., Sjouwerman, R., van den Heuvel, M. P., Kahn, R. S., and Geuze, E. (2015). Smaller hippocampal volume as a vulnerability factor for the persistence of post-traumatic stress disorder. *Psychol. Med.* 45, 2737–2746. doi: 10.1017/s0033291715000707

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=25936409) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=25936409) | [CrossRef Full Text](https://doi.org/10.1017/s0033291715000707) (https://doi.org/10.1017/s0033291715000707) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Smaller+hippocampal+volume+as+a+vulnerability+factor+for+the+persistence+of+post-traumatic+stress+disorder&author=van+Rooij+S.+J.+H.&author=Kennis+M.&author=Sjouwerman+R.&author=van+den+Heuvel+M.+P.&author=Kahn+R.+S.&author=Geuze+E.&publication_year=2015&journal=Psychol.+Med.&volume=45&pages=2737-2746) (http://scholar.google.com/scholar_lookup?title=Smaller+hippocampal+volume+as+a+vulnerability+factor+for+the+persistence+of+post-traumatic+stress+disorder&author=van+Rooij+S.+J.+H.&author=Kennis+M.&author=Sjouwerman+R.&author=van+den+Heuvel+M.+P.&author=Kahn+R.+S.&author=Geuze+E.&publication_year=2015&journal=Psychol.+Med.&volume=45&pages=2737-2746)

van Wel, J. H. P., Kuypers, K. P. C., Theunissen, E. L., Bosker, W. M., Bakker, K., and Ramaekers, J. G. (2012). Effects of acute MDMA intoxication on mood and impulsivity: role of the 5-HT₂ and 5-HT₁ receptors. *PLoS One* 7:e40187. doi: 10.1371/journal.pone.0040187

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=22808116) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=22808116) | [CrossRef Full Text](https://doi.org/10.1371/journal.pone.0040187) (https://doi.org/10.1371/journal.pone.0040187) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Effects+of+acute+MDMA+Intoxication+on+mood+and+impulsivity%3A+role+of+the+5-HT2+and+5-HT1+receptors&author=van+Wel+J.+H.+P.&author=Kuypers+K.+P.+C.&author=Theunissen+E.+L.&author=Bosker+W.+M.&author=Bakker+K.&author=Ramaekers+J.+G.&publication_year=2012&journal=PLoS+One&volume=7&pages=e40187) (http://scholar.google.com/scholar_lookup?title=Effects+of+acute+MDMA+Intoxication+on+mood+and+impulsivity%3A+role+of+the+5-HT2+and+5-HT1+receptors&author=van+Wel+J.+H.+P.&author=Kuypers+K.+P.+C.&author=Theunissen+E.+L.&author=Bosker+W.+M.&author=Bakker+K.&author=Ramaekers+J.+G.&publication_year=2012&journal=PLoS+One&volume=7&pages=e40187)

Vollenweider, F. X., Liechti, M. E., Gamma, A., Greer, G., and Geyer, M. (2002). Acute psychological and neurophysiological effects of MDMA in humans. *J. Psychoactive Drugs* 34, 171–184. doi: 10.1080/02791072.2002.10399951

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=12691207) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=12691207) | [CrossRef Full Text](https://doi.org/10.1080/02791072.2002.10399951) (https://doi.org/10.1080/02791072.2002.10399951) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Acute+psychological+and+neurophysiological+effects+of+MDMA+in+humans&author=Vollenweider+F.+X.&author=Liechti+M.+E.&author=Gamma+A.&author=Greer+G.&author=Geyer+M.&publication_year=2002&journal=J.+Psychoactive+Drugs&volume=34&pages=171-184) (http://scholar.google.com/scholar_lookup?title=Acute+psychological+and+neurophysiological+effects+of+MDMA+in+humans&author=Vollenweider+F.+X.&author=Liechti+M.+E.&author=Gamma+A.&author=Greer+G.&author=Geyer+M.&publication_year=2002&journal=J.+Psychoactive+Drugs&volume=34&pages=171-184)

Wagner, M. T., Mithoefer, M. C., Mithoefer, A. T., MacAulay, R. K., Jerome, L., Yazar-Klosinski, B., et al. (2017). Therapeutic effect of increased openness: investigating mechanism of action in MDMA-assisted psychotherapy. *J. Psychopharmacol.* 31, 967–974. doi: 10.1177/0269881117711712

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=28635375) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=28635375) | [CrossRef Full Text](https://doi.org/10.1177/0269881117711712) (https://doi.org/10.1177/0269881117711712) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Therapeutic+effect+of+increased+openness%3A+investigating+mechanism+of+action+in+MDMA-assisted+psychotherapy&author=Wagner+M.+T.&author=Mithoefer+M.+C.&author=Mithoefer+A.+T.&author=MacAulay+R.+K.&author=Jerome+L.&author=Yazar-Klosinski+B.&publication_year=2017&journal=J.+Psychopharmacol.&volume=31&pages=967-974) (http://scholar.google.com/scholar_lookup?title=Therapeutic+effect+of+increased+openness%3A+investigating+mechanism+of+action+in+MDMA-assisted+psychotherapy&author=Wagner+M.+T.&author=Mithoefer+M.+C.&author=Mithoefer+A.+T.&author=MacAulay+R.+K.&author=Jerome+L.&author=Yazar-Klosinski+B.&publication_year=2017&journal=J.+Psychopharmacol.&volume=31&pages=967-974)

Walker, D. L., Ressler, K. J., Lu, K.-T., and Davis, M. (2002). Facilitation of conditioned fear extinction by systemic administration or intra-amygdala infusions of D-cycloserine as assessed with fear-potentiated startle in rats. *J. Neurosci.* 22, 2343–2351. doi: 10.1523/jneurosci.22-06-02343.2002

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=11896173) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=11896173) | [CrossRef Full Text](https://doi.org/10.1523/jneurosci.22-06-02343.2002) (https://doi.org/10.1523/jneurosci.22-06-02343.2002) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Facilitation+of+conditioned+fear+extinction+by+systemic+administration+or+intra-amygdala+infusions+of+D-cycloserine+as+assessed+with+fear-potentiated+startle+in+rats&author=Walker+D.+L.&author=Ressler+K.+J.&author=Lu+K.-T.&author=Davis+M.&publication_year=2002&journal=J.+Neurosci.&volume=22&pages=2343-2351) (http://scholar.google.com/scholar_lookup?title=Facilitation+of+conditioned+fear+extinction+by+systemic+administration+or+intra-amygdala+infusions+of+D-cycloserine+as+assessed+with+fear-potentiated+startle+in+rats&author=Walker+D.+L.&author=Ressler+K.+J.&author=Lu+K.-T.&author=Davis+M.&publication_year=2002&journal=J.+Neurosci.&volume=22&pages=2343-2351)

Watts, B. V., Schnurr, P. P., Mayo, L., Young-Xu, Y., Weeks, W. B., and Friedman, M. J. (2013). Meta-analysis of the efficacy of treatments for posttraumatic stress disorder. *J. Clin. Psychiatry* 74, e541–e550. doi: 10.4088/JCP.12r08225

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23842024) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=23842024) | [CrossRef Full Text](https://doi.org/10.4088/JCP.12r08225) (https://doi.org/10.4088/JCP.12r08225) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Meta-analysis+of+the+efficacy+of+treatments+for+posttraumatic+stress+disorder&author=Watts+B.+V.&author=Schnurr+P.+P.&author=Mayo+L.&author=Young-Xu+Y.&author=Weeks+W.+B.&author=Friedman+M.+J.&publication_year=2013&journal=J.+Clin.+Psychiatry&volume=74&pages=e541-e550) (http://scholar.google.com/scholar_lookup?title=Meta-analysis+of+the+efficacy+of+treatments+for+posttraumatic+stress+disorder&author=Watts+B.+V.&author=Schnurr+P.+P.&author=Mayo+L.&author=Young-Xu+Y.&author=Weeks+W.+B.&author=Friedman+M.+J.&publication_year=2013&journal=J.+Clin.+Psychiatry&volume=74&pages=e541-e550)

Wilcox, C. E., Pommy, J. M., and Adinoff, B. (2016). Neural circuitry of impaired emotion regulation in substance use disorders. *Am. J. Psychiatry* 173, 344–361. doi: 10.1176/appi.ajp.2015.15060710

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=26771738) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=26771738) | [CrossRef Full Text](https://doi.org/10.1176/appi.ajp.2015.15060710) (https://doi.org/10.1176/appi.ajp.2015.15060710) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Neural+circuitry+of+impaired+emotion+regulation+in+substance+use+disorders&author=Wilcox+C.+E.&author=Pommy+J.+M.&author=Adinoff+B.&publication_year=2016&journal=Am.+J.+Psychiatry&volume=173&pages=344-361) (http://scholar.google.com/scholar_lookup?title=Neural+circuitry+of+impaired+emotion+regulation+in+substance+use+disorders&author=Wilcox+C.+E.&author=Pommy+J.+M.&author=Adinoff+B.&publication_year=2016&journal=Am.+J.+Psychiatry&volume=173&pages=344-361)

Wong, S. Y. S., Leung, J. C., Kwok, T., Ohlsson, C., Vandenput, L., Leung, P. C., et al. (2011). Low DHEAS levels are associated with depressive symptoms in elderly Chinese men: results from a large study. *Asian J. Androl.* 13, 898–902. doi: 10.1038/aja.2011.116

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=21874029) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=21874029) | [CrossRef Full Text](https://doi.org/10.1038/aja.2011.116) (https://doi.org/10.1038/aja.2011.116) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Low+DHEAS+levels+are+associated+with+depressive+symptoms+in+elderly+Chinese+men%3A+results+from+a+large+study&author=Wong+S.+Y.+S.&author=Leung+J.+C.&author=Kwok+T.&author=Ohlsson+C.&author=Vandenput+L.&author=Leung+P.+C.+&publication_year=2011&journal=Asian+J.+Androl.&volume=13&pages=898-902) (http://scholar.google.com/scholar_lookup?title=Low+DHEAS+levels+are+associated+with+depressive+symptoms+in+elderly+Chinese+men%3A+results+from+a+large+study&author=Wong+S.+Y.+S.&author=Leung+J.+C.&author=Kwok+T.&author=Ohlsson+C.&author=Vandenput+L.&author=Leung+P.+C.+&publication_year=2011&journal=Asian+J.+Androl.&volume=13&pages=898-902)

Yan, T., Xie, Q., Zheng, Z., Zou, K., and Wang, L. (2017). Different frequency repetitive transcranial magnetic stimulation (rTMS) for posttraumatic stress disorder (PTSD): a systematic review and meta-analysis. *J. Psychiatr. Res.* 89, 125–135. doi: 10.1016/j.jpsychires.2017.02.021

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=28278422) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=28278422) | [CrossRef Full Text](https://doi.org/10.1016/j.jpsychires.2017.02.021) (https://doi.org/10.1016/j.jpsychires.2017.02.021) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Different+frequency+repetitive+transcranial+magnetic+stimulation+(rTMS)+for+posttraumatic+stress+disorder+(PTSD)%3A+a+systematic+review+and+meta-analysis&author=Yan+T.&author=Xie+Q.&author=Zheng+Z.&author=Zou+K.&author=Wang+L.&publication_year=2017&journal=J.+Psychiatr.+Res.&volume=89&pages=125-135) (http://scholar.google.com/scholar_lookup?title=Different+frequency+repetitive+transcranial+magnetic+stimulation+(rTMS)+for+posttraumatic+stress+disorder+(PTSD)%3A+a+systematic+review+and+meta-analysis&author=Yan+T.&author=Xie+Q.&author=Zheng+Z.&author=Zou+K.&author=Wang+L.&publication_year=2017&journal=J.+Psychiatr.+Res.&volume=89&pages=125-135)

Yang, Y. L., and Lu, K. T. (2005). Facilitation of conditioned fear extinction by D-cycloserine is mediated by mitogen-activated protein kinase and phosphatidylinositol 3-kinase cascades and requires *de novo* protein synthesis in basolateral nucleus of amygdala. *Neuroscience* 134, 247–260. doi: 10.1016/j.neuroscience.2005.04.003

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=15951121) (http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=15951121) | [CrossRef Full Text](https://doi.org/10.1016/j.neuroscience.2005.04.003) (https://doi.org/10.1016/j.neuroscience.2005.04.003) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Facilitation+of+conditioned+fear+extinction+by+D-cycloserine+is+mediated+by+mitogen-activated+protein+kinase+and+phosphatidylinositol+3-kinase+cascades+and+requires+de+novo+protein+synthesis+in+basolateral+nucleus+of+amygdala&author=Yang+Y.+L.&author=Lu+K.+T.&publication_year=2005&journal=Neuroscience&volume=134&pages=247-260) (http://scholar.google.com/scholar_lookup?title=Facilitation+of+conditioned+fear+extinction+by+D-cycloserine+is+mediated+by+mitogen-activated+protein+kinase+and+phosphatidylinositol+3-kinase+cascades+and+requires+de+novo+protein+synthesis+in+basolateral+nucleus+of+amygdala&author=Yang+Y.+L.&author=Lu+K.+T.&publication_year=2005&journal=Neuroscience&volume=134&pages=247-260)

Yatzkar, U., and Klein, E. (2010). P.3.026 Intranasal oxytocin in patients with post traumatic stress disorder: a single dose, pilot double blind crossover study. *Eur. Neuropsychopharmacol.* 20:S84. doi: 10.1016/s0924-977x(10)70098-5

[CrossRef Full Text](https://doi.org/10.1016/s0924-977x(10)70098-5) (https://doi.org/10.1016/s0924-977x(10)70098-5) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=P.3.026+Intranasal+oxytocin+in+patients+with+post+traumatic+stress+disorder%3A+a+single+dose,+pilot+double+blind+crossover+study&author=Yatzkar+U.&author=Klein+E.&publication_year=2010&journal=Eur.+Neuropsychopharmacol.&volume=20&pages=S84) (http://scholar.google.com/scholar_lookup?title=P.3.026+Intranasal+oxytocin+in+patients+with+post+traumatic+stress+disorder%3A+a+single+dose,+pilot+double+blind+crossover+study&author=Yatzkar+U.&author=Klein+E.&publication_year=2010&journal=Eur.+Neuropsychopharmacol.&volume=20&pages=S84)

Young, M. B., Norrholm, S. D., Khoury, L. M., Jovanovic, T., Rauch, S. A. M., Reiff, C. M., et al. (2017). Inhibition of serotonin transporters disrupts the enhancement of fear memory extinction by 3,4-methylenedioxymethamphetamine (MDMA). *Psychopharmacology* 234, 2883–2895. doi: 10.1007/s00213-017-4684-8

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=28741031) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=28741031>) | [CrossRef Full Text](https://doi.org/10.1007/s00213-017-4684-8) (<https://doi.org/10.1007/s00213-017-4684-8>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Inhibition+of+serotonin+transporters+disrupts+the+enhancement+of+fear+memory+extinction+by+3,4-methylenedioxymethamphetamine+(MDMA)&author=Young+M.+B.&author=Norrholm+S.+D.&author=Khoury+L.+M.&author=Jovanovic+T.&author=Rauch+S.+A.+M.&author=Reiff+C.+M.&publication_year=2017&journal=Psychopharmacology&volume=234&pages=2883-2895) ([http://scholar.google.com/scholar_lookup?title=Inhibition+of+serotonin+transporters+disrupts+the+enhancement+of+fear+memory+extinction+by+3,4-methylenedioxymethamphetamine+\(MDMA\)&author=Young+M.+B.&author=Norrholm+S.+D.&author=Khoury+L.+M.&author=Jovanovic+T.&author=Rauch+S.+A.+M.&author=Reiff+C.+M.&publication_year=2017&journal=Psychopharmacology&volume=234&pages=2883-2895](http://scholar.google.com/scholar_lookup?title=Inhibition+of+serotonin+transporters+disrupts+the+enhancement+of+fear+memory+extinction+by+3,4-methylenedioxymethamphetamine+(MDMA)&author=Young+M.+B.&author=Norrholm+S.+D.&author=Khoury+L.+M.&author=Jovanovic+T.&author=Rauch+S.+A.+M.&author=Reiff+C.+M.&publication_year=2017&journal=Psychopharmacology&volume=234&pages=2883-2895))

Zalta, A. K., Gillihan, S. J., Fisher, A. J., Mintz, J., McLean, C. P., Yehuda, R., et al. (2014). Change in negative cognitions associated with PTSD predicts symptom reduction in prolonged exposure. *J. Consult. Clin. Psychol.* 82,171–175. doi: 10.1037/a0034735

[PubMed Abstract](http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24188512) (<http://www.ncbi.nlm.nih.gov/sites/entrez?Db=pubmed&Cmd=ShowDetailView&TermToSearch=24188512>) | [CrossRef Full Text](https://doi.org/10.1037/a0034735) (<https://doi.org/10.1037/a0034735>) | [Google Scholar](http://scholar.google.com/scholar_lookup?title=Change+in+negative+cognitions+associated+with+PTSD+predicts+symptom+reduction+in+prolonged+exposure&author=Zalta+A.+K.&author=Gillihan+S.+J.&author=Fisher+A.+J.&author=Mintz+J.&author=McLean+C.+P.&author=Yehuda+R.&publication_year=2014&journal=J.+Consult.+Clin.+Psychol.&volume=82&pages=171-175) (http://scholar.google.com/scholar_lookup?title=Change+in+negative+cognitions+associated+with+PTSD+predicts+symptom+reduction+in+prolonged+exposure&author=Zalta+A.+K.&author=Gillihan+S.+J.&author=Fisher+A.+J.&author=Mintz+J.&author=McLean+C.+P.&author=Yehuda+R.&publication_year=2014&journal=J.+Consult.+Clin.+Psychol.&volume=82&pages=171-175)

Keywords: PTSD, prolonged exposure, psychotherapy, exposure therapy, neuroscience, extinction learning

Citation: Stojek MM, McSweeney LB and Rauch SAM (2018) Neuroscience Informed Prolonged Exposure Practice: Increasing Efficiency and Efficacy Through Mechanisms. *Front. Behav. Neurosci.* 12:281. doi: 10.3389/fnbeh.2018.00281

Received: 31 May 2018; **Accepted:** 02 November 2018;

Published: 20 November 2018.

Edited by:

Arash Javanbakht (<https://loop.frontiersin.org/people/200798/overview>), Wayne State University, United States

Reviewed by:

James L. Abelson (<https://loop.frontiersin.org/people/214717/overview>), Wayne State University and University of Michigan, United States
Stuart F. White (<https://loop.frontiersin.org/people/110651/overview>), Boys Town National Research Hospital, United States

Copyright © 2018 Stojek, McSweeney and Rauch. This is an open-access article distributed under the terms of the [Creative Commons Attribution License \(CC BY\)](http://creativecommons.org/licenses/by/4.0/) (<http://creativecommons.org/licenses/by/4.0/>). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.

***Correspondence:** Sheila A. M. Rauch, sheila.a.m.rauch@emory.edu (<mailto:sheila.a.m.rauch@emory.edu>)

Disclaimer: All claims expressed in this article are solely those of the authors and do not necessarily represent those of their affiliated organizations, or those of the publisher, the editors and the reviewers. Any product that may be evaluated in this article or claim that may be made by its manufacturer is not guaranteed or endorsed by the publisher.

People also looked at

Treating PTSD: A Review of Evidence-Based Psychotherapy Interventions ([/articles/10.3389/fnbeh.2018.00258/full](https://www.frontiersin.org/articles/10.3389/fnbeh.2018.00258/full))

Laura E. Watkins (<https://loop.frontiersin.org/people/566363/overview>), Kelsey R. Sprang (<https://loop.frontiersin.org/people/632779/overview>) and Barbara O. Rothbaum (<https://loop.frontiersin.org/people/632797/overview>)

Neural Mechanism of Altered Limb Perceptions Caused by Temporal Sensorimotor Incongruence (/articles/10.3389/fnbeh.2018.00282/full)

Osamu Katayama (<https://loop.frontiersin.org/people/484884/overview>), Tatsuya Tsukamoto, Michihiro Osumi (<https://loop.frontiersin.org/people/116015/overview>), Takayuki Kodama (<https://loop.frontiersin.org/people/484900/overview>) and Shu Morioka (<https://loop.frontiersin.org/people/130240/overview>)

Age Effects on Spatiotemporal Dynamics of Response Inhibition: An MEG Study (/articles/10.3389/fnagi.2018.00386/full)

Mei-Yin Lin (<https://loop.frontiersin.org/people/503091/overview>), Yi-Jhan Tseng (<https://loop.frontiersin.org/people/619081/overview>) and Chia-Hsiung Cheng (<https://loop.frontiersin.org/people/174131/overview>)

A Phase II Study to Evaluate the Safety and Efficacy of Prasinezumab in Early Parkinson's Disease (PASADENA): Rationale, Design, and Baseline Data (/articles/10.3389/fneur.2021.705407/full)

Gennaro Pagano (<https://loop.frontiersin.org/people/83103/overview>), Frank G. Boess (<https://loop.frontiersin.org/people/617781/overview>), Kirsten I. Taylor, Benedicte Ricci (<https://loop.frontiersin.org/people/1341199/overview>), Brit Mollenhauer, Werner Poewe (<https://loop.frontiersin.org/people/851270/overview>), Anne Boulay, Judith Anzures-Cabrera, Annamarie Vogt, Maddalena Marchesi, Anke Post, Tania Nikolcheva, Gene G. Kinney, Wagner M. Zago, Daniel K. Ness, Hanno Svoboda, Markus Britschgi (<https://loop.frontiersin.org/people/1111661/overview>), Susanne Ostrowitzki, Tanya Simuni, Kenneth Marek, Martin Koller, Jeff Sevigny, Rachel Doody, Paulo Fontoura (<https://loop.frontiersin.org/people/602335/overview>), Daniel Umbricht (<https://loop.frontiersin.org/people/416802/overview>), Azad Bonni, PASADENA Investigators, Prasinezumab Study Group, Claudia Altendorf, Chareyna Anandan, Giulia Andrews, Solène Ansquer, Raphaelae Arrouasse, Sana Aslam, Jean-Philippe Azulay, Jeanette Baker, Ernest Balaguer Martinez, Shadi Barbu, Kara Bardram, Danny Bega, Helena Bejr-Kasem Marco, Isabelle Benatru, Eve Benchetrit, Felix Bernhard, Amir Besharat, Sagari Bette, Amelie Bichon, Andrew Billnitzer, Sophie Blondeau, Thomas Boraud, Freiderike Borngärber, James Boyd, Kathrin Brockmann, Matthew Brodsky, Ethan Brown, Christof Bruecke, Fabienne Calvas, Monica Canelo, Federico Carbone, Claire Carroll, Laura Casado Fernandez, Catherine Cassé-Perrot, Anna Castrioto, Helene Catala, Justine Chan, Samia Cheriet, Anthony Ciabarra, Joseph Classen, Juliana Coleman, Robert Coleman, Yaroslau Compta, Anne-Gaëlle Corbillé, Jean-Christophe Corvol, Mariana Cosgaya, Nabila Dahodwala, Philippe Damier, Elodie David, Thomas Davis, Marissa Dean, Berengere Debilly, Janell DeGiorgio, Andres Deik, Laure Delaby, Marie-Helene Delfini, Pascal Derkinderen, Philipp Derost, Maria de Toledo, Lisa Deuel, Ann Marie Diaz-Hernandez, Cameron Dietiker, Karina Dimenshteyn, Julio Dotor, Franck Durif, Jens Ebentheuer, Karla Maria Eggert, Sara Eichau Madueño, Claudia Eickhoff, Aaron Ellenbogen, Philipp Ellmerer, Ines Esparragosa Vazquez, Alexandre Eusebio, Siobhan Ewert, John Fang, Danielle Feigenbaum, Frederique Fluchere, Alexandra Foubert-Samier, Marie Fournier, Anne Fradet, Valerie Fraix, Samuel Frank, Franca Fries, Monique Galitzky, Marisol Gallardó Pérez, Jose Manuel García Moreno, Carmen Gasca, Thomas Gasser, Joyce Gibbons, Caroline Giordana, Alicia Gonzalez Martinez, Ira Goodman, Arantza Gorospe, Marie Goubeaud, David Grabli, Mangone Graziella, Stephan Grimaldi, Jeffrey Gross, Raquel Guimaraes-Costa, Andreas Hartmann, Christian Hartmann, Travis Hassell, Robert Hauser, Antonio Hernandez, Jorge Hernandez-Vara, Günter Höglinger, Christian Homedes, Andrea Horta-Barba, Jean-Luc Houeto, Julius Huebl, Jennifer Hui, Stuart Isaacson, Joseph Jankovic, Annette Janzen, Junior Jauregui, Jocelyne Jiao, Maria Jose Marti Domenech, Xavier Joseph, Srinath Kadimi, Pat Kaminski, Silja Kannenberg, Jan Kassubek, Maya Katz, Kevin Klos, Shannon Klos, Christopher Kobet, Jennifer Koeber, Patricia Krause, Andrea Kuehn, Jaime Kulisevsky Bojarsky, Rajeev Kumar, Martin Kunz, Lilla Kurvits, Kimberly Kwei, Simon Laganieri, Brice Laurens, Johannes Levin, Oren Levy, Peter LeWitt, Gurutz Linazasoro Cristóbal, Irene Litvan, Karlo Lizarraga, Katherine Longardner, Rocio Lopez, Lydia Lopez Manzanares, Sara Lucas del Pozo, Maria Rosario Luquin Puido, Nijee Luthra, Kelly Lyons, Sylvia Maass, Gerrit Machetanz, Yolanda Macias, David Maltete, Jorge Uriel Manes Miro, Louise-Laure Mariani, Juan Marin, Kathrin Marini, Ana Marques, Gloria Marti, Maria Jose Marti Domenech, Saul Martinez, Wassilios Meissner, Sara Meoni, Brit Mollenhauer, Dunia Mon Martinez, Johnson Moon, Elena Moro, Peter Morrison, Christoph Muehlberg, Manpreet Multani, Christine Murphy, Anthony

Nicholas, Rajesh Pahwa, Antonio Palasi, Heidi Pape, Neepa Patel, Prity Patel, Marina Peball, Elizabeth Peckham, Terry Peery, Rafael Perez, Jesus Perez, Alisa Petit, Elmar Pinkhardt, Werner Poewe, Elsa Pomies, Cecile Preterre, Joseph Quinn, Olivier Rascol, Philippe Remy, Irene Richard, Benjamin Roeben, Emily Ruether, Jost-Julian Rumpf, David Russell, Hayet Salhi, Daniela Samaniego-Toro, Alexandra Samier-Foubert, Antonio Sanchez, Emmanuelle Schmitt, Alfons Schnitzler, Oliver Schorr, Julie Schwartzbard, Kerstin Schweyer, Klaus Seppi, Victoria Sergo, Holly Shill, Andrew Siderowf, Tanya Simuni, Umberto Spampinato, Ashok Sriram, Natividad Stover, Caroline Tanner, Arjun Tarakad, Carolyn Taylor, Claire Thalamus, Thomas Toothaker, Nadege Van Blercom, Nora Vanegas-Arrogave, Lydia Vela, Sylvian Vergnet, Tiphaine Vidal, Jonathan Vöglein, Ryan Walsh, Cheryl Waters, Mirko Wegscheider, Endy Weidinger, Caroline Weill, Gregor Wenzel, Tatiana Witjas, Isabel Wurster, Brenton Wright, Milan Zimmermann, Rafael Zuzuarregui, Markus Abt, Atieh Bamdadian, Teresa Barata, Nicholas Barbet, Sara Belli, Frank Boess, Azad Bonni, Edilio Borroni, Anne Boulay, Markus Britschgi, Jerome Chague, Valerie Cosson, Christian Czech, Dennis Deptula, Cheikh Diack, Rachelle Doody, Juergen Dukart, Giulia D'Urso, Sebastian Dziadek, Hannah Eddleston, Chris Edgar, Laurent Essieux, Morgan Farrell, Rebecca Finch, Paulo Fontoura, Waltraud Gruenbauer, Andrea Hahn, Stefan Holiga, Michael Honer, Shirin Jadidi, Kelly Johnson-Wood, Markus Keller, Timothy Kilchenmann, Martin Koller, Thomas Kremer, Thomas Kustermann, Claire Landsdall, Michael Lindemann, Florian Lipsmeier, Cecile Luzy, Marianne Manchester, Maddalena Marchesi, Ferenc Martenyi, Meret Martin-Facklam, Katerina Mironova, Annabelle Monnet, Emma Moore, Daniel K Ness, Markus Niggli, Tania Nikolcheva, Susanne Ostrowitzki, Gennaro Pagano, Benedicte Passmard, Agnes Poirier, Anke Post, Megana Prasad, Nathalie Pross, Tiffany Quock, Benedicte Ricci, Ellen Rose, Christoph Sarry, Christine Schubert, Dennis Selkoe, Jeff Sevigny, Kaycee Sink, Hannah Staunton, Tim Steven, Alexander Strasak, Hanno Svoboda, Kirsten Taylor, Radhika Tripuraneni, Dylan Trundell, Daniel Umbricht, Lynne Verselis, Annamarie Vogt, Ekaterina Volkova-Volkmar, Cornelia Weber, Silke Weber and Wagner Zago

Moving from Virtual Reality Exposure-Based Therapy to Augmented Reality Exposure-Based Therapy: A Review (/articles/10.3389/fnhum.2014.00112/full)

Oliver Baus (<https://loop.frontiersin.org/people/118981/overview>) and Stéphane Bouchard (<https://loop.frontiersin.org/people/139499/overview>)

Guidelines



Explore



Outreach



Connect



Follow us



© 2022 Frontiers Media S.A. All rights reserved. [Privacy policy](#) | [Terms and conditions](#)