Involvement of the cerebellum in EMDR efficiency: a metabolic connectivity PET study in PTSD

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The aim of the present study was to investigate the metabolic changes of precuneus connectivity through PET study in PTSD.

ABSTRACT

Background: We recently reported an improvement of precuneus PET metabolism after EMDR therapy in military participants suffering from PTSD.

Objective: The aim of the present study was to investigate the metabolic changes of precuneus connectivity in these participants after such treatment.

Method: Fifteen participants with PTSD performed a brain 18F-FDG-PET sensitized by virtual reality exposure to war scenes, before and after EMDR treatment. Inter-regional correlation analysis was performed to study metabolic changes of precuneus connectivity through SPM-T maps at whole-brain level (p < 0.005 for the voxel, p < 0.05 for the cluster).

Results: A decrease of connectivity was observed after EMDR between the precuneus and two significant bilateral clusters of the cerebellum (bilateral Crus I and VI cerebellar lobules, Tmax voxel of 5.8 and 5.3, and cluster size of 343 and 314 voxels respectively). Moreover, higher cerebellar metabolism before treatment was associated with reduced clinical PTSD scores after EMDR (p = 0.03).

Conclusions: The posterior cerebellum and its metabolic connectivity with the precuneus are involved in the clinical efficiency of EMDR in PTSD.

KEYWORDS

PTSD; EMDR; PET; connectivity; cerebellum

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HIGHLIGHTS

- The connectivity between the precuneus and the posterior cerebellum is decreased in PTSD after EMDR.
- Higher cerebellar metabolism before EMDR is associated with reduced PTSD scores after EMDR.
- The posterior cerebellum and its metabolic connectivity with the precuneus are involved in the clinical efficiency of EMDR in PTSD.
1. Introduction

Post-traumatic stress disorder (PTSD) is the re-experiencing of specific aspects of the traumatic memory in cases of exposure to traumatic events (Hoppen & Morina, 2019). This psychiatric condition is related to a failure in extinguishing a fear-conditioned process (“DSM-5,” 2013). This leads to several symptoms involving both emotional under- and over-modulation (Lanius, Frewen, Vermetten, & Yehuda, 2010). Emotional under-modulation refers to decreased regulation of prefrontal regions resulting from a hyperactivity in limbic regions associated to re-experiencing of the traumatic moment with intrusive thoughts, flashbacks, and nightmares, avoidance of situations related to the trauma and hyper-arousal. By contrast, over-modulation of these same prefrontal areas leads to diminished limbic activity and is consequently responsible for negative alterations in cognition and mood, with also attempts to restrict unwanted emotional experiences including in dissociation, numbing, and analgesia (Kolk, Burbridge, & Suzuki, 1997; Lanius et al., 2010; Van Der Kolk, 1998). In fact, PTSD could be the consequence of a different way to encode events of a traumatic memory as compared to ordinary events (Van Der Kolk, 1998).

Eye movement desensitization and reprocessing (EMDR) is now a well-established treatment for PTSD (Chen et al., 2014). EMDR, which consists in alternate bilateral stimulations (ABS) while the patient is asked to maintain their attention on traumatic memories, aims at re-elaborating memory from traumatic events in order to lead them to an adaptive resolution (Landin-Romero, Moreno-Alcazar, Pagani, & Amann, 2018; Shapiro Francine, 2001). The exact physiological mechanism for reaching therapeutic efficiency in EMDR is nonetheless still unknown (Sara Carletto, Borsato, & Pagani, 2017).

18F-FDG PET is a functional imaging technique that could help to better understanding this efficiency. Indeed, it allows for studying neuronal glucose metabolism. In the framework of military PTSD, it can be sensitized by virtual reality exposure to investigate metabolic changes during stressful tasks (Verger et al., 2018) involving war-related visual, auditory, and proprioceptive stimuli. We recently showed an increased precuneus metabolism upon virtual reality exposure of war scenes after EMDR treatment in military participants suffering from PTSD (Rousseau et al., 2019). The metabolic connectivity can be explored through the interregional correlation analysis (IRCA) developed by Lee et al. (Lee et al., 2008), to underline the networks resulting from EMDR treatment in these participants. Briefly, it pivots on the fact that different brain areas are metabolically connected when they share similar spatial variance in radiotracer uptake. The precuneus is, in fact, a key region for the well-known ‘default mode network’ in resting-state brain, and it has important functions for self-related mental representations and integration of past and present information. The DMN involves core hub areas including the posterior cingulate, the precuneus, the medial prefrontal cortex, and the bilateral inferior parietal lobes. These hub areas are connected to other regions of the DMN including among others the temporal cortex and the medial temporal lobe (Fransson & Marrelec, 2008). A relative decreased of precuneus activation in PTSD patients has been shown in resting-state MRI studies, as well as a negative correlation between preunecal activity and PTSD severity, similarly to what was reported in our PET study (Rousseau et al., 2019), suggesting disruptions in self-referential thought (Geuze, Vermetten, de Kloet, & Westenberg, 2008; Ke et al., 2016). By studying functional connectivity of the precuneus, a decreased of connectivity has been found in key regions of the DMN, particularly prefrontal areas. (Akiki et al., 2018). By contrast, in a real-time fMRI study after amygdala neurofeedback treatment, an increased connectivity between the precuneus and the left dorsolateral prefrontal cortex was associated with a decrease in hyper-arousal symptoms of PTSD (Misaki et al., 2018).

Therefore, the aim of the present study was to investigate the precuneus metabolic PET connectivity changes in military participants suffering from PTSD before and after EMDR.

2. Methods

2.1. Sample

Fifteen military male participants (36.8 ± 8.9 years old) suffering from PTSD since their returns from the Afghanistan or Mali wars were included. The mean disease duration was 5.8 ± 0.8 years. These participants were recruited in the military hospital Sainte-Anne at Toulon, France. Diagnosis of PTSD was established according to the DSM-IV TR (American Psychiatric Association, 2000). Patients with present and/or past neurological or psychiatric conditions, with the exception of anxiety and depressive disorders if their occurrence was connected with PTSD, and with an addictive disorder were excluded. Detailed characteristics of these patients are available in a previous study involving the same sample of patients (Table 1 in Rousseau et al., 2019).
2019). For patient’s selection, the Posttraumatic Stress Checklist Scale (PCLS) (Ventureyra, Yao, Cottriaux, Note, & De Mey-Guillard, 2002), which is a brief and self-report questionnaire for evaluating the severity of the three main syndromes of PTSD, and the Clinician-Administered PTSD Scale (CAPS) (Blake et al., 1995), which is a structured interview providing a categorical diagnosis, were used as defined by the DSM. The mean scores for the previous tests at inclusion were 62.1 ± 8.2 for the PCLS and 78.1 ± 12.3 for the CAPS, respectively. All these participants performed a brain $^{18}$F-FDG PET sensitized by virtual reality interactive exposure of war scenes before and after EMDR therapy. The second $^{18}$F-FDG PET was performed during the month following the end of the EMDR therapy. Subjects gave written informed consent for their participation in accordance with the Declaration of Helsinki. The study was approved by the Institutional Review Board CPP Sud Méditerranée (Ref: 2014-002126-12).

### 2.2. EMDR

None of the participant had received formal exposure or cognitive-behavioural therapy before the EMDR procedure. EMDR therapy was performed according to the standard protocol (Landin-Romero et al., 2018). All participants were symptom-free and no longer diagnosed with PTSD after EMDR therapy, as assessed by a psychiatric diagnosis according to DSM-IV criteria and clinical PTSD scales.

### 2.3. $^{18}$F-FDG-PET acquisition and analysis

$^{18}$F-FDG PET sensitized by virtual reality exposure was performed using an integrated PET/CT camera (Discovery 710, GE Healthcare, Waukesha, WI) with parameters previously described (Verger et al., 2018). Briefly, each subject was confronted with intense trauma cues, consisting of the attack on their group by insurgents during a patrol in an artificially created village in Afghanistan, with several virtual soldiers wounded. Patients were exposed to the environment approximately 10 min before the injection. $^{18}$F-FDG (150 MBq) was injected intravenously 1 min before their unit was assaulted, and the virtual exposure immersion was maintained after injection for approximately 7 min. Thereafter, patients were placed in a quiet environment with their eyes closed but continued to feel the stress of the VRE. PET images, acquired in a lying position as recommended in standard practice, started 30 min after the injection and ended 15 min later.

Whole-brain statistical analysis was performed at the voxel level using SPM8 software (Wellcome Department of Cognitive Neurology, University College, London, UK). PET images were spatially normalized onto the PET template of the Montreal National Institute (MNI) space, smoothed with a Gaussian filter (8 mm full-width at half-maximum), resulting in $2 \times 2 \times 2$ mm voxel images. Proportional scaling was applied. The bilateral precuneus was selected since it was shown to be the sole area to be involved in PTSD participants responsive to EMDR after SPM T-maps comparison before and after EMDR therapy using paired t-tests ($p < 0.005$, uncorrected, $k > 180$) (Rousseau et al., 2019). Its mask was derived from PickAtlas (https://www.nitrc.org/projects/wfu_pickatlas/) using the Automated Anatomical Labelling (AAL) atlas (Tzourio-Mazoyer et al., 2002), and volumes of interest were individually extracted with Marsbar software (http://marsbar.sourceforge.net/). To evaluate metabolic connectivity from the precuneus mask in the intergroup comparison, IRCA was performed as previously described (Lee et al., 2008). Briefly, mean values of precuneus cerebral metabolic rate of glucose were used as interacted covariates to find regions showing significant voxel-wise negative/positive correlations for the direct comparison between PTSD patients before and after EMDR therapy. Results were expressed as an increase of positive or negative connectivity. A threshold of $p < 0.005$ uncorrected for the voxel, and $p < 0.05$ uncorrected for the cluster was applied for SPM (T) maps. These thresholds of significance lead to a cluster volume of more than 300 voxels, well beyond the ‘expected voxels per cluster’ volume provided by SPM ($k > 51$), used to calculate the probability density function for cluster size based on random field theory to avoid type I and II errors (Lieberman & Cunningham, 2009). The precise identification of each structure located by its MNI coordinates, its respective volume, and T-max intensity were extracted by using the report provided by the SPM xjView toolbox (http://www.alivelearn.net/xjview). Mean values of metabolism were extracted at the individual level for precuneus and significant cluster(s) to calculate correlations with Clinician-Administered PTSD Scale CAPS, and Posttraumatic Stress Checklist Scale, PCLS. A canonical analysis correlating cerebellar metabolic values before EMDR to the combination of CAPS and PCLS after therapy was performed. A $p$-value <0.05 was considered as significant.

### 3. Results

All the participants were free of PTSD symptoms at the end of the EMDR therapy (Table 1 in Rousseau et al., 2019). The mean PCLS and CAPS scores after EMDR were respectively at 24.4 ± 8.1 and 13.9 ± 12.7 ($p < 0.001$ for the comparison between scores before and after EMDR).

As shown in Figure 1 and as compared to $^{18}$F-FDG PET scans before EMDR, a decrease of connectivity was noticed after EMDR between the precuneus and the cerebellum in two bilateral significant clusters (343 voxels, MNI coordinates of x: 40; y: -66; z: -20, and $T_{\text{max}}$ voxel of 5.8 for the right cerebellar cluster involving the Crus I and VI lobules, 314 voxels, MNI...
coordinates of x: -30; y: -86; z: -20, and T\textsubscript{max} voxel of 5.3 for the left cerebellar cluster involving Crus I and VI lobules).

Moreover, higher cerebellar metabolic values before EMDR were correlated with better responses to clinical PTSD scales after EMDR (canonical analysis correlating cerebellar metabolic values before EMDR to the combination of CAPS and PCLS after therapy; rho = 0.67; \( p = 0.03 \))

4. Discussion
The current study shows that PTSD remission after EMDR therapy is reflected by a decrease of connectivity between the precuneus and the posterior cerebellum (Crus I and lobule VI). In addition, higher cerebellar metabolism before treatment was correlated to the significantly reduced clinical PTSD scores after EMDR.

Previous studies exploring the physiological mechanisms involved in PTSD were based on the top-down theory (Nicholson et al., 2017). This theory states that in patients with PTSD, the mesial prefrontal cortex, a structure implicated in emotion regulation, is not able to correctly play its role of normalizing the excess of activity of limbic structures (Nicholson et al., 2017), and especially an hyperactivation of amygdalae (Harper, Rasolkhani-Kalhorn, & Drozd, 2009). This insufficiency leads to an incorrect encoding of traumatic memory events and thus to clinical symptoms of the PTSD (Van Der Kolk, 1998).

In this line, EMDR efficiency has been shown of resulting from a shift of sub-cortical to neo-cortical transfer information in an EEG study (Pagani et al., 2012), with the reactivation of the mesial prefrontal cortex to correctly encode the traumatic events.

However, a bottom-up theory, involving the cerebellum, has been recently proposed in PTSD (S. Carletto & Borsato, 2017). The cerebellum is a structure implicated in fear emotional regulation (Schmahmann, 2010) and is strongly connected to the whole brain (Zhu, Yung, Kwok-Chong Chow, Chan, & Wang, 2006). The cerebellum presents altered functions in patients with PTSD in a perfusion PET study (Pissiota et al., 2002). Our results support the role of the cerebellum in PTSD.

Our participants with PTSD exhibited a decrease of connectivity between the precuneus and the posterior cerebellum after clinical response to EMDR. This result shows that the posterior cerebellum and its metabolic connectivity with the precuneus is likely involved in PTSD symptoms and traumatic memory evolution after EMDR therapy. The precuneus, which has already been shown to be the sole structure with increased metabolism after EMDR (Rousseau et al., 2019), is implicated in fear-conditioning and extinction regulation, a mechanism that would be disrupted in PTSD (Fullana et al., 2018). The precuneus and the cerebellum have also been described together as two brain areas involved in PTSD in an fMRI study, as their activation at baseline was correlated with improvement of PTSD symptoms (Ke et al., 2016). The posterior cerebellum (Crus I and II, lobules VI and VII) is devoted to cognitive functions, including working memory, language, visuospatial, and executive functions (Rabellino, Densmore, Théberge, McKinnon, & Lanius, 2018). This area is associated to the Cerebellar Cognitive Affective Syndrome, characterized by affect in emotion regulation associated with either passivity, blunted affect, and withdrawal, or disinhibition, irritability, emotional lability (Schmahmann, 2004; Stoodley & Schmahmann, 2010). Interestingly, these symptoms are those also

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*Figure 1. Significant right (left panel) and left (right panel) cerebellar clusters showing decreased $^{18}$F-FDG PET metabolic connectivity with precuneus after EMDR, projected onto MRI slices, spatially normalized into the standard SPM template. The right and left cerebellar cluster involve both the Crus I and VI cerebellar lobules at respective T-max voxel of 5.8 and 5.3.*
exhibited by individuals with PTSD (“DSM-5,” 2013). The posterior cerebellum has already been found to be involved in PTSD as an increased activity in this area has been shown in PTSD as compared to healthy controls in MRI at rest (Wang et al., 2016), and an increased regional cerebral blood flow (rCBF) has been reported in lobule VI and Crus I in PTSD as compared to healthy controls (Bonne et al., 2003). In addition, the posterior cerebellum was shown to participate in the DMN in resting-state fMRI study (Kucyi, Hove, Biederman, Van Dijk, & Valera, 2015). Indeed, lateral cerebellar areas including Crus I/II were shown to be functionally connected with the DMN, and neurostimulation of these cerebellar areas was shown to modulate functional connectivity specifically with and between core DMN hubs (Kucyi et al., 2015). Interestingly, the DMN is known to count for the long-term clinical outcome of PTSD since it is associated with PTSD symptom severity (Reuveni et al., 2016). Regards to the posterior cerebellum, a decreased functional connectivity between this cerebellar part and the central autonomic network was found to correlate positively with PTSD symptomatology in a resting-state fMRI (Thome et al., 2017), this decrease of cerebellar connectivity associated to better PTSD symptomatology being in line with our PET results. These findings suggest that in PTSD, as state anxiety symptoms increase, the posterior cerebellum demonstrates increased connectivity with other cortical areas (Remington & Remington, 2012). By contrast, in a resting-state fMRI study comparing PTSD to healthy controls, Rabellino et al. found a decreased functional connectivity between the posterior cerebellum and prefrontal regions involved in PTSD (Rabellino et al., 2018). This observation strengthens nevertheless that the posterior cerebellum is involved in emotional dysregulation. It should be kept in mind that PTSD is a complex physio-pathological entity exceeding the top-down and bottom-up theories by implementing notions of emotion under- or over-modulation (Lanius et al., 2010). In line with these previous observations, we hypothesize that the decreased of connectivity between the precuneus, an area involved in emotion regulation through self-referential thought and participating in the DMN, and the posterior cerebellum, the cognitive entity part of the cerebellum could explain the better outcome of PTSD patients after EMDR. These connectivity changes are part of the DMN connectivity which is known to be involved in PTSD (Reuveni et al., 2016).

Interestingly, in MRI studies, the volume of the cerebellum is inversely correlated to the clinical PTSD symptoms (Baldaçara et al., 2011), but is also positively correlated to the ability to experience positive affect in such patients (Frewen, Dozois, & Lanius, 2012). In addition, it has been postulated that its activation was a potential explanation of EMDR therapy (Calancie, Khalid-Khan, Booij, & Munoz, 2018). This is also in line with our results showing that higher cerebellar metabolism before treatment was correlated with better clinical response to EMDR therapy.

Further studies should include waitlist and more larger PTSD participant groups with other treatments to verify that our results are reproducible and specifically related to EMDR therapy (Rousseau et al., 2019). Moreover, this metabolic connectivity analysis has not been performed for the whole brain but by using an a priori hypothesis based on previous reported results (Rousseau et al., 2019). Further analyses exploring the whole-brain connectivity should be set up to better apprehending physiological mechanisms implicated in EMDR efficiency.

Altogether, the present study highlights the involvement of the posterior cerebellum in participants with PTSD evolution. Its metabolic connectivity with the precuneus underlines networks related to EMDR therapy efficiency.

**Disclosure statement**

No potential conflict of interest was reported by the authors.

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