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Activating attachment memories affects Default Mode Network in a non-clinical sample with perceived dysfunctional parenting: an EEG functional connectivity study

Mauro Adenzato[1^], Claudio Imperatori[2^], Rita B. Ardito[3*], Enrico Maria Valenti[2,4], Giacomo Della Marca[5], Sofia D'Ari[2], Luigia Palmiero[2], Jacopo S. Penso[1], Benedetto Farina[2]

[1]Department of Psychology, University of Turin, Turin, Italy

[2]Department of Human Sciences, European University of Rome, Rome, Italy

[3]Department of Neuroscience "Rita Levi Montalcini", University of Turin, Turin, Italy

[4]Department of Dynamic and Clinical Psychology, Sapienza University of Rome, Rome, Italy

[5]Sleep Disorders Unit, Institute of Neurology, Catholic University, Rome, Italy

[^] These authors contributed equally to this work

***Corresponding author:**

Rita B. Ardito, Ph.D.

Department of Neuroscience "Rita Levi Montalcini", University of Turin, Turin, Italy

Via Cherasco, 15 – 10126 Turin, Italy

Phone: +39.011.670.3071

E-mail address: rita.ardito@unito.it

Abstract

Dysfunctional parenting constitutes a factor of psychopathological vulnerability affecting development both at neurobiological and psychological level. The default mode network (DMN), a large scale network for brain functional integration, is supposed to play a crucial role in those psychological functions altered by dysfunctional parenting. Here we investigate electroencephalography DMN functional connectivity in relation to perceived dysfunctional parenting (PDP) in a non-clinical sample. We hypothesized that participants with high PDP would exhibit decreased DMN connectivity after the activation of attachment memories. Our results support this hypothesis: participants with high PDP showed a decrease of theta connectivity between left temporoparietal junction and right anterior cingulate cortex after the activation of attachment memories, and, compared to participants with low PDP, showed a decrease of delta connectivity in the same brain areas. We interpret these decreased DMN connectivity in participants with high PDP as the "neurophysiological signature" of the impaired ability to mentalize their own relational experiences with significant others after the activation of early attachment memories. Thus, the activation of attachment memories in individuals exposed to dysfunctional parenting could lead to a transitory failure of functional brain connectivity and consequent disturbance of high integrative mental functions, such as emotional regulation and mentalization.

Keywords: Attachment, Default mode network, Dysfunctional parenting, Electroencephalography, Functional connectivity, Mentalization

Introduction

Dysfunctional parenting, such as very low care, emotional abuse and high overprotection, has been compared to others forms of child maltreatment [1-5]. Consistently, a significant and increasing body of evidence suggests that dysfunctional and/or neglectful parenting, like other forms of early relational trauma, is one of the major risk factor and negative prognostic cause for almost all psychiatric disorders [6-9]. Indeed, regardless of specific diagnosis, it has been reported that dysfunctional parenting constitutes a factor of psychopathological vulnerability affecting development both at neurobiological and psychological level [2, 6-8, 10-13]. Among the most common psychopathological consequences related to dysfunctional parenting there are emotive disorders and dysregulation [14, 15], alterations in inhibitory control and executive functions [10], cognitive and consciousness disturbances [2, 16, 17], self identity and self agency alterations [18, 19], mentalization dysfunctions [5, 20], relational problems and low social competence [21, 22]. Some scholars have hypothesized that a significant amount of these psychopathological disturbances have in common a lack of mental integration produced by dysfunctional parenting [2, 16, 23, 24].

Under a neurobiological point of view it has been supposed that the default mode network (DMN), a crucial large scale network for brain functional integration [25-27], plays an important role in those psychological functions altered by dysfunctional parenting [28-31]. Consistently, the DMN and its subcomponents alterations are frequently reported in people with dysfunctional parenting and other forms of child maltreatment [8, 12].

One of the most intriguing issue in this area is that clinical observations and empirical research data lead to consider that some of these disturbances, such as emotive and behavioural dysregulation, dissociative symptoms, mentalization disruption, relational problems, are not stable symptoms, but may emerge when triggered by socio-emotional stimuli like the activation of early attachment memories [16, 24, 32]. According to attachment theory and its subsequent clinical applications, the automatic and implicit (i.e., unconscious) activation of attachment relational memories in

individuals with histories of neglect or maltreatment in childhood could trigger disintegrative psychopathological process that leads to typical psychopathology related to dysfunctional parenting [2, 23, 24].

For this reason the aim of the present study was to investigate electroencephalography DMN functional connectivity in relation to the quality of the perceived dysfunctional parenting (PDP), i.e. the self-reported experiences of neglect, abuse and/or overprotection within the relationships with one's parents [33-35], both in resting state (RS) and after the activation of attachment memories using the Adult Attachment Interview as a trigger (AAI) [36, 37] in a non-clinical sample. Based on empirical data and clinical grounds, we hypothesized that participants with PDP, compared to participants without PDP, would exhibit decreased DMN connectivity after the activation of attachment memories.

Materials and Methods

Participants

Participants were 50 students (fourteen men, mean age: 22.62 ± 2.41 years) recruited through advertisements posted in the university. The enrollment lasted from October 2017 to May 2018. Study participants contributed voluntarily and anonymously after providing informed consent. They did not receive payment or any other compensation (i.e., academic credit). Inclusion criteria were: age between 18 and 30 years, both genders. Exclusion criteria were: history of psychiatric disease and/or neurologic diseases; head trauma; left handedness; assumption of Central Nervous System active drugs in the two weeks prior to assessment. A checklist with dichotomous items was used to assess inclusion/exclusion criteria and socio-demographic data.

After receiving information about the aim of the study, all participants provided a written consent to participate in the study that was performed according to the Helsinki declaration standards. The research was approved by the European University's ethic review board.

Procedure

After providing the written informed consent, all participants were administered the Measure of Parental Style (MOPS) [38] and the Brief Symptom Inventory (BSI) [39]. Furthermore, in order to identify the presence of past and/or current psychiatric disorders, during the intake visit, participants were asked screening questions according to a checklist prepared for a previous study [40].

On a separate day from the self-report assessment, all participants underwent the Adult Attachment Interview (AAI) [37], a semi-structured interview able to activate the attachment system by the retrieval of childhood emotional and relational memories of past attachment experiences [37, 41]. Rigorous psychometric testing and meta-analyses of the AAI demonstrate stability and discriminant and predictive validity in both clinical and nonclinical populations [42, 43]. In the present study, the AAI was used as "trigger stimulus" of the attachment behavioral system. Indeed, previous studies demonstrated that the AAI is able to alter psychophysiological parameters related to the emotion regulation of people with different attachment styles [36, 44, 45] and to modify the cortical functional connectivity related to the retrieval of early attachment memories in both healthy and clinical subjects [16].

Trained clinical psychologists (LP) administered the AAI in a quiet and comfortable room. EEG recordings were performed before (Pre-AAI condition) and immediately after (i.e., about 8-10 minutes for each participant) the interview (Post-AAI condition). The interviews lasted on average one hour and 30 minutes.

Questionnaire

The MOPS [38] is the redefined version of the Parental Bonding Instrument [46] and it is composed by 30 items which separately investigate mother' (15 items) and father' (15 items) parental styles. Items are scored on a 4-point Likert scale (from "not true at all" to "extremely true") and grouped in three dimensions for each parent, confirmed through principal components analysis

[38]: indifference, over-control, and abuse. Higher scores reflect higher self-reported experiences of neglect, abuse and/or overprotection during the first 16 years of life. The MOPS has been used extensively in clinical research [9, 34, 47] and it was developed to overcome some negative aspects (e.g., low clarity of several items) of the original version [9]. Satisfactory psychometric properties have been reported in the original validation study [38]. Furthermore, good cross-cultural adaptation has been observed [48, 49]. In the present research, the Italian version of the scale has been used [48] and the Cronbach's alpha in the present sample was 0.88 for the 30-item MOPS total scores.

The BSI [39] is the short version of the Symptom Checklist-90R [50] and it is composed by 53 items evaluating a broad range of psychological symptoms during the past seven days. Items are scored on a 4-point Likert scale ranging from 0 (not at all) to 4 (extremely) and grouped in 9 primary symptom dimensions: somatization, obsession-compulsion, interpersonal sensitivity, depression, anxiety, hostility, phobic anxiety, paranoid ideation and psychoticism. A measure of general level of psychopathology, the global severity index (GSI), is also calculated using the sums for the nine symptom dimensions (higher scores reflects more self-reported symptoms). The BSI is widely used in clinical research and it is characterized by good psychometric properties [51]. In the present study, the Italian version of the scale was used [52] and the Cronbach's alpha in the present sample was 0.94 for the GSI.

EEG recordings

Resting State (RS) EEG was recorded using the Micromed System Plus digital EEGraph (Micromed© S.p.A., Mogliano Veneto, TV, Italy) in the European University EEG Lab, with each participants sitting in a comfortable armchair, with his/her eyes closed, in a quiet, semi-darkened silent room for 5 minutes. In order to avoid alcohol and or caffeine effects on EEG data, participants were asked to refrain from drinking alcohol and caffeine for 4 to 6 hours immediately before their EEG recordings.

148 EEG recordings included 31 standard scalp leads, positioned according to the 10-20 system
149 (recording sites: Fp1, AF3, F3, FC1, C3, CP1, P3, PO3, O1, F7, FC5, T3, CP5, T5, Fz, Cz, Pz, Fp2,
150 AF4, F4, FC2, C4, CP2, P4, PO4, O2, F8, FC6, T4, CP6, T6), and the Electrocardiography (ECG).
151 The reference electrodes were placed on the linked mastoids. Impedances were kept below 5KΩ
152 before starting the recording and checked again at the end of the experimental recording. Sampling
153 frequency was 256 Hz; A/D conversion was made at 16 bit; pre-amplifiers amplitude range was
154 ±3200 and low-frequency pre-filters were set at 0.15 Hz. The following band-pass filters were used:
155 HFF= 0.2 Hz; LFF= 128 Hz. In the present study the following frequency bands was considered:
156 delta (0.5–4 Hz); theta (4.5–7.5 Hz); alpha (8–12.5 Hz); beta (13–30 Hz); gamma (30.5–60 Hz).
157 Details about artifact rejection have been described elsewhere [40]. Briefly, visual artifact
158 rejection (e.g., cap adjustment) was firstly performed on the raw EEG trace. These segments were
159 removed, and then independent component analysis (ICA) was applied to EEG recordings to
160 identify and remove non-cerebral artifacts (i.e., eye and muscular movements, cardiac pulses)
161 before data analysis. Although it has been reported that ICA correction may affect EEG
162 connectivity [53], correcting artifacts using this procedure is widely used in EEG phase
163 synchronization studies [54-56]. Furthermore, several reports [54-56] documented no significant
164 modifications of EEG coherence data after ICA correction.
165 The minimum length of the artifact-free EEG recording included in the analysis was 180
166 seconds (even if not consecutive) for each participant for each condition (i.e., pre-AAI and post-
167 AAI).
168
169 *Connectivity analysis*
170 All EEG analysis were performed using the exact Low Resolution Electromagnetic
171 Tomography software (eLORETA), a validated tool for localizing brain electric activity based on
172 multichannel surface EEG recordings [57]. The eLORETA software is characterized by a
173 satisfactory localization agreement with different multi-modal imaging techniques, and it is a

suitable tool for DMN assessment [58]. The connectivity analysis were performed using the lagged phase synchronization formula [59]. This algorithm has been widely used to assess EEG functional connectivity, and it is characterized by several advantages (e.g., it is resistant to non-physiological artifacts) [59]. Although functional magnetic resonance imaging (fMRI) is commonly used to investigate the functional connectivity of DMN, recent studies have shown that EEG is also suitable for investigating the functional properties of this network.

According to previous EEG connectivity studies [29, 30, 40, 60], in order to evaluate functional connectivity in the DMN, 12 Regions of Interest (ROIs) were selected (Figure 1) and the 'single nearest voxel' option (i.e., each ROI consisted of a single voxel, the closest to each seed) was chosen (detailed DMN Montreal Neurological Institute and Talairach coordinates can be found in [40]). Briefly, the "ROI-maker#2 method" available in the eLORETA software has been selected and, starting from 42 Brodmann Areas (BAs) in each hemisphere provided by the software [61], 12 ROIs were defined according Thatcher et al. [58].

The eLORETA calculated the lagged phase synchronization values between all these ROIs (i.e., 144 connections) and the source reconstruction algorithm [57].

Statistical analysis

In order to reveal groups of subjects with high and low PDP (i.e., PDP+ and PDP- groups), a Two Step Cluster Analysis procedure was performed using MOPS sub-scales scores. Cluster solutions was assessed using Schwarz's Bayesian Criterion (BIC) as clustering criterion [62]. Chi-squared tests ($\chi 2$), and Mann–Whitney's U tests were used to investigate differences between clusters, respectively for dichotomous and dimensional variables.

EEG connectivity analysis was performed using the eLORETA software. Between and within comparisons were performed for each frequency band. Specifically the following statistical comparisons were performed: i) Pre-AAI PDP+ vs Pre-AAI PDP-, ii) Post-AAI PDP+ vs Pre-AAI PDP+, iii) Post-AAI PDP- vs Pre-AAI PDP-, iv) Post-AAI PDP+ vs Pre-AAI PDP-. All

comparisons were performed using the statistical non-parametric mapping (SnPM) methodology provided by the eLORETA software (i.e, a Fisher's permutation test) [63]. In order to avoid family-wise type-I errors, the non-parametric randomization procedure (supplied by the eLORETA software), was performed for the correction of multiple comparison [63]. For all comparisons, the eLORETA software provides experimental values of T, corresponding to a significance of $p < 0.01$ and $p < 0.05$.

Finally, Spearman's *rho* correlation coefficients were reported as measures of associations among MOPS subscales scores, GSI, and any significant EEG connectivity data observed in the between comparisons. Cluster Analysis, Chi-squared tests, Mann–Whitney's U tests, correlation analyses were performed using IBM SPSS Statistics for Windows, version 23.0. The use of nonparametric tests was chosen because none of the present variables were normally distributed (Shapiro–Wilk test, $p < 0.05$).

Results

The Two Step Cluster Analysis procedure indicated a 2-group solution (BIC change= −36.01; Ratio of distance measures= 3.38). 34 % of the sample (N= 17) was included in the first cluster, and 66 % (N= 33) was included in cluster 2. Compared to individuals included in cluster 2 (i.e., PDP- group), subjects included in cluster 1 (i.e., PDP+ group) had significantly higher scores in all PDP sub-subscales. Thus, cluster 1 is mostly characterized by individuals reporting higher PDP. Furthermore, compared to the individuals with low PDP, those with high PDP had a significantly higher scores in the GSI and in all BSI subscales, with the exception of interpersonal sensitivity, phobic anxiety and psychoticism subscales. Detailed bivariate analyses are listed in Table 1.

EEG recordings suitable for the analysis were obtained for all participants. Qualitative visual evaluation of the EEG recordings, performed by a trained neurophysiologist, showed no relevant modifications of the background rhythm frequency (e.g., epileptic discharges).

Furthermore, no relevant modifications of EEG signal (e.g., evidence of sleepiness) during the recordings were detected. The average time analyzed for the present sample was 283 ± 14 sec and 276 ± 15 respectively for cluster 1 and cluster 2 subjects in pre-AAI condition and 291 ± 12 sec and 277 ± 19 in post-AAI condition.

Connectivity results

In the between-groups comparison (PDP+ vs PDP-) for the Pre-AAI condition, the thresholds for significance were T= ±2.73 corresponding to $p< 0.05$, and T= ±3.19 corresponding to $p< 0.01$. In this condition, no significant modifications were observed between groups (Figure 2; Panel A).

In the within-group comparison (Post-AAI vs. Pre-AAI) for the PDP+ group, the thresholds for significance were T= ±3.99 corresponding to $p< 0.05$, and T= ±4.63, corresponding to $p< 0.01$. In this comparison, significant modifications were observed in the theta frequency (Figure 3; Panel A). Compared to Pre-AAI condition, PDP+ individuals showed in Post-AAI condition a decrease of theta connectivity between left Temporoparietal Junction (TPJ; ROI 11) and right Anterior Cingulate Cortex (ACC; ROI 8) (T= -4.09, $p= 0.037$). No significant differences were observed in the other frequency bands, although a significant trend was observed between left TPJ and right ACC also in the delta band (T= -3.76, $p= 0.08$).

In the within-group comparison (Post-AAI vs. Pre-AAI) for the PDP- group, the thresholds for significance group were T= ±3.54 corresponding to $p< 0.05$, and T= ±4.03, corresponding to $p< 0.01$. In this comparison, significant modifications were observed in the alpha frequency band (Figure 3; Panel B). Compared to Pre-AAI condition, PDP- individuals showed in Post-AAI condition an increase of alpha connectivity between right TPJ (ROI 12) and both right and left Posterior Cingulate Cortex (PCC; ROI 6 and ROI 5) (respectively T= 3.56; $p= 0.047$ and T= 3.71; $p= 0.030$). No significant differences were observed in the other frequency bands

In the between-groups comparison (PDP+ vs PDP-) for the Post-AAI condition, the thresholds for significance were T= ±2.92 corresponding to $p< 0.05$, and T= ±3.47 corresponding to

$p < 0.01$. Significant modifications were observed in the delta band (Figure 3; Panel B). Compared to PDP- individuals, PDP+ participants showed a decrease of delta connectivity between left TPJ (ROI 11) and right ACC (ROI 8) (T= -3.29; p= 0.018). No significant differences were observed in the other frequency bands.

Association among EEG functional connectivity data, MOPS and GSI scores

MOPS total score was negatively related with the strength of delta connectivity between left TPJ and right ACC (*rho*= -0.28; p= 0.048). Furthermore, the strength of delta connectivity observed after the AAI between left TPJ and right ACC was negatively related with both maternal indifference (*rho*= -0.36; p= 0.010) and maternal over-control (*rho*= -0.33; p= 0.020) sub-scale. Although GSI was positively related with all MOPS sub-scales, no significant correlation was observed between EEG connectivity data and psychopathological score. Detailed correlations are reported in Table 2.

Discussion

The a priori hypothesis of the present study was that participants with high PDP (i.e., PDP+ group), compared to participants with low or without PDP (i.e., PDP- group), would exhibit decreased DMN connectivity after the activation of attachment memories. Our results support this hypothesis. Indeed, after the activation of attachment memories triggered by the AAI, PDP+ participants (within-group comparison) showed a decrease of theta connectivity between left TPJ and right ACC. Furthermore, after the administration of the AAI, compared to PDP- participants, PDP+ individuals showed a decrease of delta connectivity in the same brain areas (i.e., left TPJ and right ACC). Consistently with our hypothesis, these connectivity modifications were observed exclusively after the activation of early attachment memories as no significant DMN connectivity differences were detected in the between-groups comparison before the administration of the AAI.

Our results are in line with previous studies reporting DMN alterations in people with dysfunctional parenting and other forms of early relational adverse experiences [8, 12]. The DMN is thought to be involved in several higher-order integrative mental functions such as self-consciousness, self-processing and episodic memory [58, 64] that are supposed to be impaired by dysfunctional parenting. This network has been conceptualized as a distributed and dynamic brain system composed by a set of interacting hubs and subsystems with specific functions [26, 27]. Specifically, the dorsal medial subsystem, which includes several brain areas such as the TPJ and dorsal medial prefrontal cortex, has been associated with mentalization, social cognition as well as with semantic/conceptual processing. Conversely, the medial temporal subsystem, which involves anatomical regions such as hippocampal formation, the retrosplenial cingulate cortex and ventral medial prefrontal cortex, has been related with autobiographical thought, episodic memory and contextual retrieval. Finally, the midline hubs of the DMN, namely the mPFC, the rostral anterior cingulate and the posterior cingulate cortex, are involved across a wide range of self-related processes integrating the dorsal medial and medial temporal subsystem [26, 27].

Therefore, taking into account DMN related functions and processes as well as the type of mental processes elicited by the AAI, we may speculate that the decreased DMN connectivity observed between left TPJ and right ACC in the PDP+ participants is the "neurophysiological signature" of the impaired ability to mentalize their own relational experiences with significant others after the activation of early attachment memories.

Indeed, the role of both left and right TPJ and of the ACC in mentalization [i.e., the ability to attribute mental states to oneself and to others and to understand that others have mental states independent from one's own; see for example 65, 66, 67] is widely recognized in the literature [68-70].

Our interpretation is also strengthened by the increase of DMN connectivity between right TPJ and both right and left PCC observed in PDP- participants after the administration of the AAI. The PCC is considered the crucial node of the DMN [71] and it is involved in several emotion and cognitive

processing [72], with critical relevance in maintaining a sense of self-consciousness and self-referential thoughts during RS [73]. It is also interesting to note that PDP- participants showed increased connectivity in the alpha frequency band, which is considered to be positively related to DMN activity as well as with spontaneous self-referential processes, such as mentalization [25]. Therefore, this result seems to support our interpretation according to which during the AAI participants are induced to mentalize their own relational experiences, presumably activating a set of self-related processes such as episodic and autobiographical memories.

Taken together, our results support both clinical observations and experimental results according to which the alterations associated with PDP are not stable symptoms but may emerge when triggered by early attachment relational memories [16, 23, 32]. These results seem to be also partially consistent with MRI studies indicating that early-life adversity may be associated with structural alterations in brain white matter, specifically in the cingulate cortex [11]. According to our data, it is possible to hypothesize that, in individuals with early adverse relational experiences, the structural connectivity deficit becomes functionally evident and clinically symptomatic when the system is overloaded by affective and cognitive attachment related stimuli. Moreover, although it was not among the goals of the present research, future work could investigate whether and to what extent the attachment style can influence the DMN functional connectivity. In fact, the same adverse experiences can lead to very different development paths and the possibility that different attachment styles may be associated with different patterns of alteration of functional connectivity cannot be excluded in principle.

Another result rising from this study is the usefulness of MOPS in detecting the psychopathological vulnerability related to the early relational adverse experiences. In our opinion, MOPS based screening could be useful in the clinical practice where the clients report their own perception of parental way to protect and control, and combined with EEG connectivity seems to be a useful and reliable tool to improve our understanding on the psychopathological processes underlying PDP.

In spite of our interesting results, the study has some limitations that should be considered. Firstly, the sample size is limited, which may affect the generalizability of the results. Furthermore, our sample included mostly female participants, and previous studies showed sex differences in EEG brain activity during RS condition [74].

Secondly, we have investigated PDP and psychopathology using self-report measures, which are known to be potentially affected by social desirability. Thirdly, we used scalp EEG recordings, which have an intrinsic limit in space resolution. Finally, we have investigated DMN functional connectivity after AAI, which make our interpretations specific to the activation of attachment memories. It is possible that others not-related attachment triggers (e.g., viewing negative emotional facial expressions) may be associated with different DMN alterations in high PDP individuals. Although these ideas are purely hypothetical, they might be useful in guiding future research.

Despite these limitations, to the best of our knowledge, this is the first study which investigated the association between DMN EEG functional connectivity and PDP both in RS and after the activation of attachment memories. In conclusion, our results seem to support the hypothesis according to which the activation of attachment memories in individuals exposed to dysfunctional parenting and other forms of early relational adverse experiences could lead to a transitory failure of functional brain connectivity and consequent disturbance of high integrative mental functions. These transitory alterations might explain, even partially, the emergence of some typical psychopathological symptoms such as emotional dysregulation, dissociative symptoms, inhibitory control and executive functions disturbances, self identity and self agency and mentalization impairments [10, 16, 23, 24]. Therefore, our result also highlights the possibility of developing new therapeutic approaches focused on the self neuro-modulation, such as alpha/theta neurofeedback, which may increases mentalization and DMN EEG connectivity [60].

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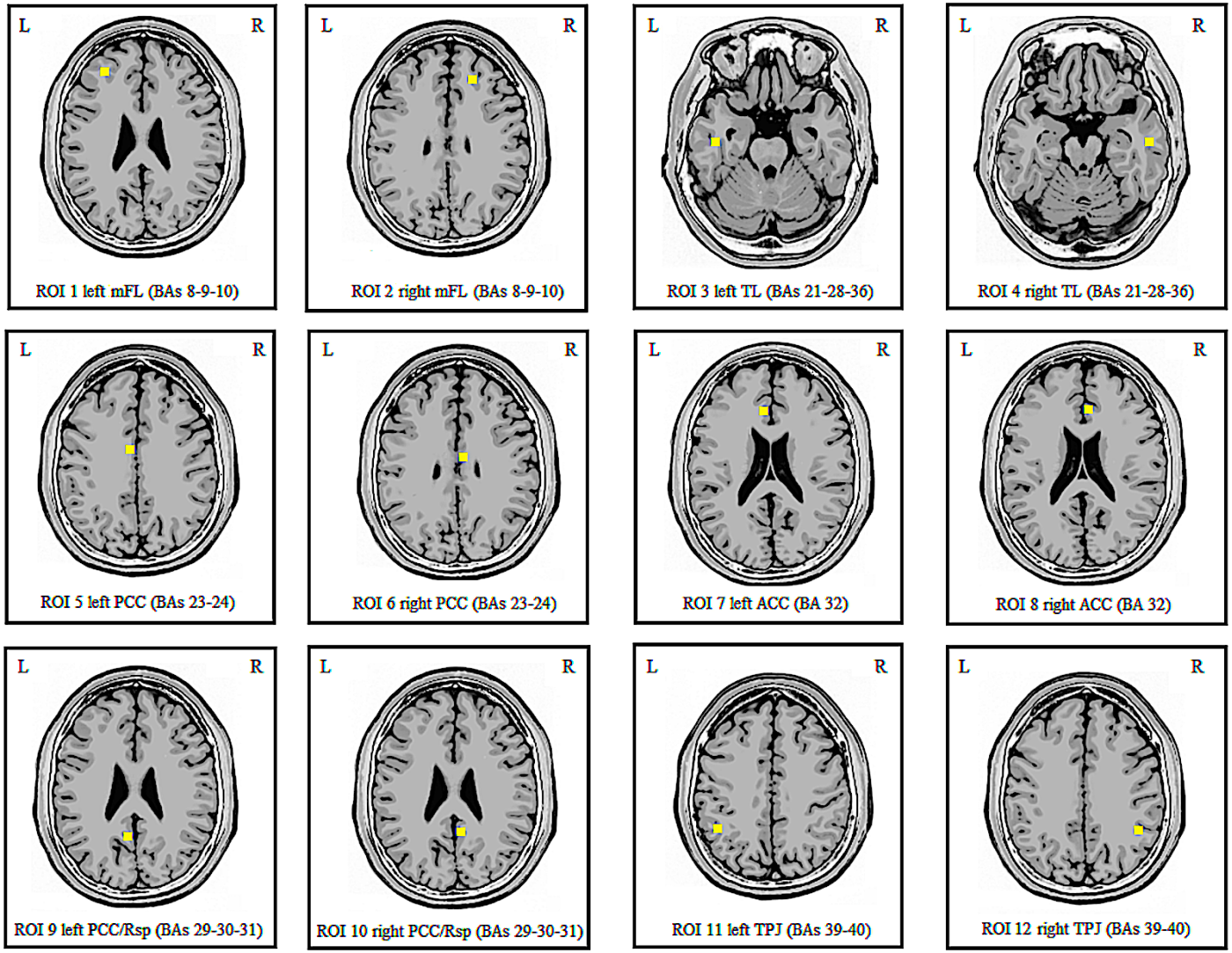
559 **Legend to figure 1.** Axial view of the Default Mode Network regions of interest.



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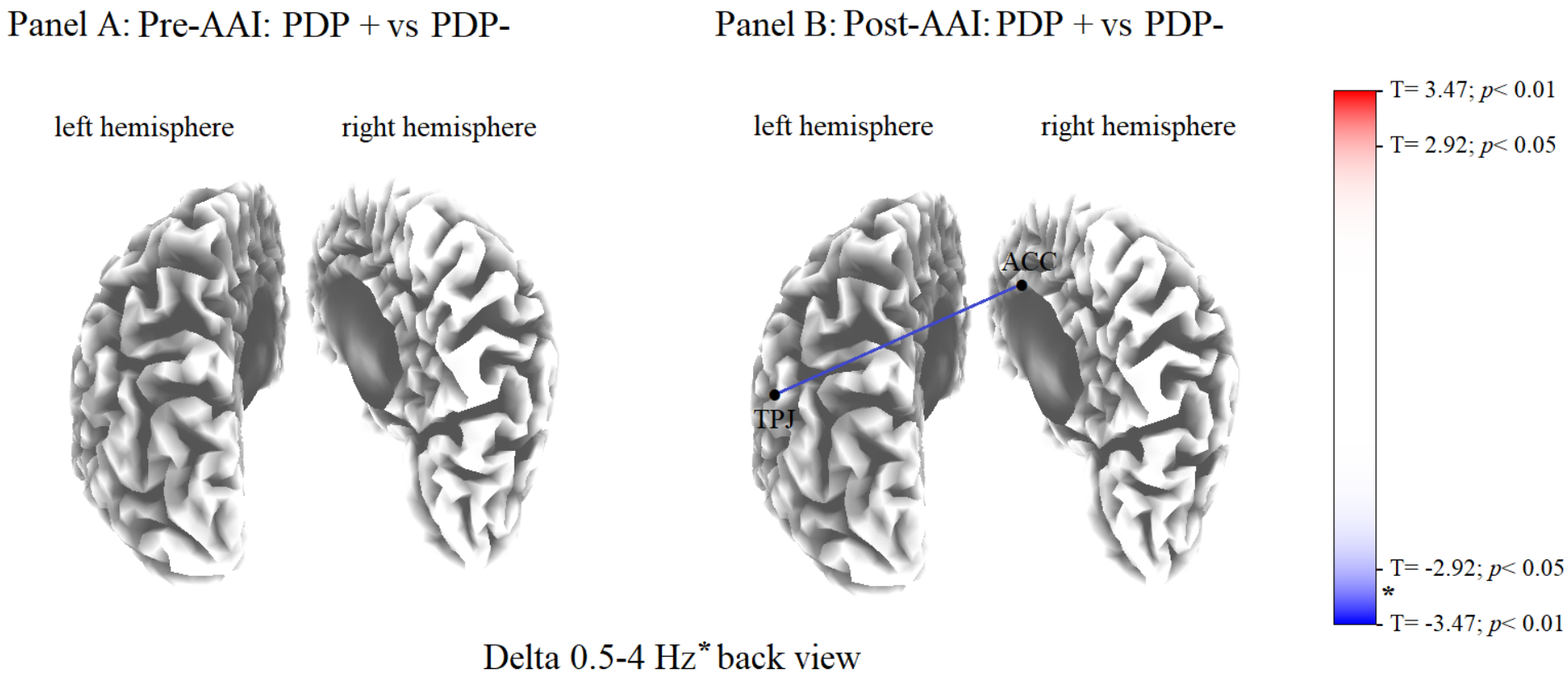
561 Abbreviation:L= left; R= right; ROI= region of interest; mFL= medial Frontal Lobe; BA (Brodmann area); TL= Temporal Lobe; PCC= Posterior Cingulate cortex; ACC=
562 Anterior Cingulate Cortex; PCC/Rsp= Posterior Cingulate/Retrosplenial cortex; TPJ= temporo-parietal junction

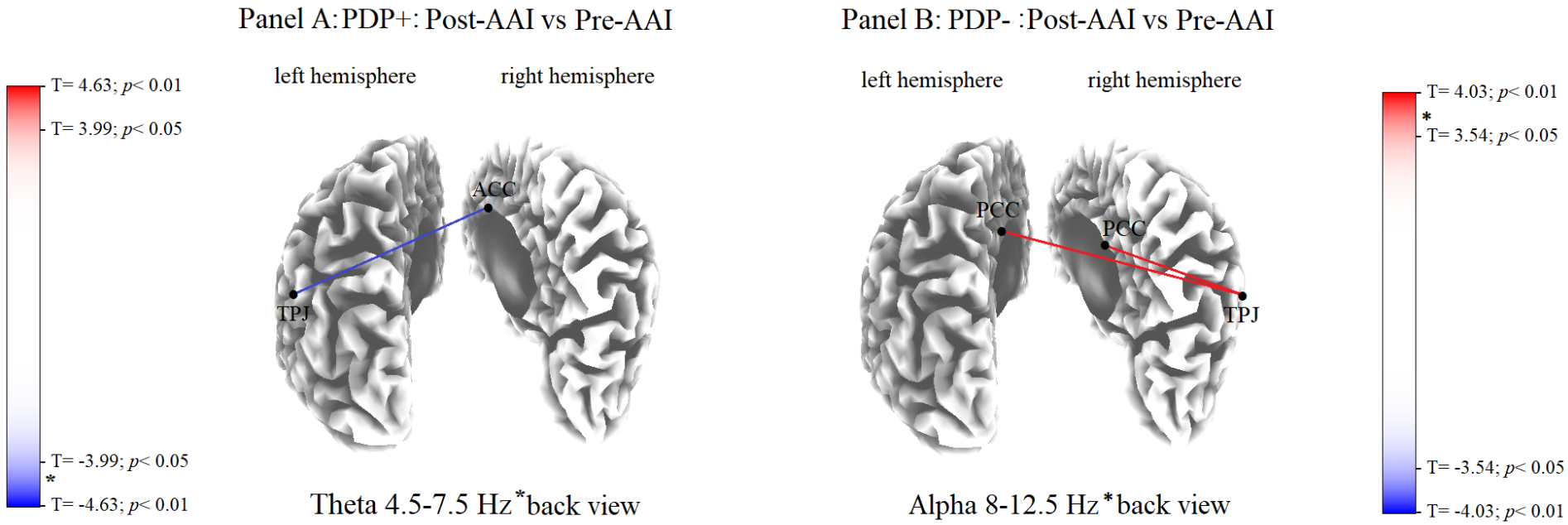
Legend to figure 2. Results of the eLORETA between comparisons of EEG functional connectivity in the delta bands. Panel A: PDP+ vs PDP- in Pre-AAI condition; Panel B: PDP+ vs PDP- in Post-AAI condition. Blue lines indicate connections presenting a significant decrease of EEG functional connectivity. Red lines would indicate increase of EEG functional connectivity (not present). Threshold values (T) for statistical significance (corresponding to $p < 0.05$ and $p < 0.01$) are reported in the right side of the figure. In this comparison, significant modifications (*) were observed in the Post-AAI condition. Compared to PDP- individuals, PDP+ subjects showed a decrease of delta connectivity between left TPJ and right ACC.



Abbreviations: AAI= Adult Attachment Interview; PDP+= high perceived dysfunctional parenting group; PDP-= low perceived dysfunctional parenting group; TPJ= temporoparietal junction; ACC= anterior cingulate cortex

Legend to figure 3.

Results of the eLORETA within comparisons (pre-AAI vs post-AAI) of EEG functional connectivity in PDP+ and PDP- group, respectively for theta and alpha band. Panel A: Post-AAI vs Pre-AAI for PDP+ group; Panel B: Post-AAI vs Pre-AAI for PDP- group. Blue lines indicate connections presenting a significant decrease of EEG functional connectivity. Red lines indicate connections presenting an increase of EEG functional connectivity. Threshold values (T) for statistical significance (corresponding to $p< 0.05$ and $p< 0.01$) are reported in the right and left side of the figure, respectively for PDP+ and PDP- group. In this comparisons, significant modifications were observed for both PDP+ (*) and PDP- group (*). Compared to Pre-AAI condition, PDP+ individuals showed in Post-AAI condition a decrease of theta connectivity between left TPJ and right ACC. Conversely, compared to Pre-AAI condition, PDP- individuals showed in Post-AAI condition an increase of alpha connectivity between right TPJ and both right and left PCC.



Abbreviations:
AAI= Adult Attachment Interview; PDP+= high perceived dysfunctional parenting group; PDP-= low perceived dysfunctional parenting group; TPJ= temporoparietal junction; ACC= Anterior Cingulate Cortex; PCC= Posterior Cingulate Cortex

590 Table 1. **Bivariate analyses**

Variables	**Cluster 1 (PDP+) (N = 17)**	**Cluster 2 (PDP-) (N = 33)**	**Test Statistics**	***p* =**
Age – M ± DS	22.59 ± 2.90	22.64 ± 2.16	*U* = 265	0.748
Educational level (years) – M ± SD	15.71 ± 1.65	15.44 ± 1.84	*U* = 249	0.624
Women - N (%)	13 (76.5)	23 (69.7)	χ^2_1 = 0.26	0.613
MOPS total score – M ± DS	40.65 ± 15.81	9.47 ± 6.32	*U* = 0.05	**< 0.001**
MOPS sub-scales				
Maternal indifference – M ± DS	5.59 ± 5.72	0.94 ± 1.50	*U* = 89	**< 0.001**
Maternal over-control – M ± DS	7.94 ± 2.79	3.39 ± 2.18	*U* = 57.5	**< 0.001**
Maternal abuse – M ± DS	6.24 ± 4.63	0.94 ± 1.48	*U* = 57	**< 0.001**
Paternal indifference – M ± DS	8.47 ± 5.63	1.13 ± 1.34	*U* = 43.5	**< 0.001**
Paternal over-control – M ± DS	5.18 ± 2.38	2.53 ± 2.34	*U* = 113	**< 0.001**
Paternal abuse – M ± DS	7.24 ± 3.82	0.44 ± 0.88	*U* = 22.5	**< 0.001**
BSI-GSI – M ± DS	0.95 ± 0.66	0.56 ± 0.68	*U* = 143.5	**0.005**
Somatization – M ± DS	0.86 ± 0.72	0.44 ± 0.63	*U* = 172	**0.024**
Obsession-Compulsion – M ± DS	1.40 ± 0.86	0.81 ± 0.90	*U* = 153	**0.009**
Interpersonal Sensitivity – M ± DS	0.94 ± 0.88	0.57 ± 0.70	*U* = 203	0.106
Depression – M ± DS	0.98 ± 0.75	0.62 ± 0.79	*U* = 170	**0.023**
Anxiety – M ± DS	1.21 ± 0.82	0.75 ± 0.87	*U* = 168	**0.021**
Hostility – M ± DS	0.98 ± 0.99	0.46 ± 0.67	*U* = 186.5	**0.049**
Phobic Anxiety – M ± DS	0.47 ± 0.72	0.25 ± 0.61	*U* = 238	0.324
Paranoid Ideation – M ± DS	1.01 ± 0.75	0.49 ± 0.79	*U* = 138	**0.003**
Psychoticism – M ± DS	0.76 ± 0.71	0.48 ± 0.71	*U* = 194	0.071

Abbreviations:
PDP+= high perceived dysfunctional parenting group; PDP-= low perceived dysfunctional parenting group; SD = standard deviation; MOPS= Measure of Parental Style; BSI-GSI = Brief Symptom Inventory-Global Severity Index;

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Table 2. Values of Spearman's *rho* correlation coefficient among EEG connectivity data, MOPS and BCSL-GSI scores in all sample (N = 50). Significant correlations are indicated by stars (*).

	MOPS Total	Maternal indifference	Maternal over-control	Maternal abuse	Paternal indifference	Paternal over-control	Paternal abuse	GSI	Delta ROIs 11-8
MOPS total	-								
Maternal indifference	0.78**	-							
Maternal over-control	0.83**	0.57**	-						
Maternal abuse	0.81**	0.73**	0.72**	-					
Paternal indifference	0.79**	0.60**	0.52**	0.59**	-				
Paternal over-control	0.70**	0.38**	0.58**	0.41**	0.41**	-			
Paternal abuse	0.80**	0.51**	0.52**	0.64**	0.66**	0.60**	-		
GSI	0.55**	0.51**	0.46**	0.46**	0.42**	0.34*	0.36*	-	
Delta ROIs 11-8	-0.28*	-0.36*	-0.33*	-0.26	-0.13	-0.08	-0.22	-0.26	-

Abbreviations: ROIs= Regions of interest; MOPS= Measure of Parental Style; = Global Severity Index;
* $p < 0.05$; ** $p < 0.01$