



# Neuroanatomical correlates of perceptual aberrations in psychosis☆☆☆



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## ABSTRACT

**Background:** Aberrations in body perception are common in psychotic disorders. The insula and temporoparietal junction (TPJ) are involved in body ownership and spatial perception suggesting that abnormal structure of these regions might be related to the expression of perceptual aberrations in psychosis.

**Methods:** 58 individuals with a primary psychotic disorder and 40 healthy subjects completed the Chapman Perceptual Aberration Scale (PAS) and underwent structural magnetic resonance imaging (MRI). Grey matter volume was extracted from a-priori defined TPJ, whole insula, and insula sub-division regions-of-interest (ROIs) and correlated with PAS scores. Additionally, a voxel-based morphometry (VBM) analysis examining the correlation between voxel-wise grey matter volume and PAS scores was conducted.

**Results:** PAS scores in psychosis patients correlated with bilateral whole insula (right:  $r = -0.30, p = 0.026$ ; left:  $r = -0.35, p = 0.011$ ) and right TPJ ( $r = -0.27, p = 0.024$ ) volumes. The correlation between grey matter volume and PAS was strongest for the posterior sub-division of the insula (right:  $r = -0.32, p = 0.017$ ; left:  $r = -0.37, p = 0.006$ ). VBM analyses confirmed the ROI results: negative correlations with PAS were identified in clusters within the posterior and dorsal anterior insula, and the right TPJ. An exploratory, whole-brain analysis also revealed two additional regions located in the left middle orbitofrontal gyrus and left inferior temporal gyrus that inversely correlated with PAS scores.

**Conclusions:** Perceptual aberrations in individuals with psychosis are related to lower grey matter volume in the insula and TPJ. This relationship was strongest in the posterior region of the insula and right TPJ; brain areas that have been implicated in interoception and somesthesia.

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## 1. Introduction

Perceptual aberrations, including the sensation that one's organs are rotting, feeling that the body is unreal or that the shape and size of body parts are changing or merging with external objects, and altered sense of bodily ownership are common in psychosis (Bleuler, 1950; Kraepelin et al., 1919). Patients with a psychotic illness, primarily schizophrenia, and those at-risk for developing psychosis score higher on self-report questionnaires of perceptual aberration such as the Perceptual Aberration Scale (PAS; Brosey and Woodward, 2015; Horan et al., 2008; Kwapil, 1998; Lenzenweger and Loranger, 1989; Schurhoff et al., 2005; Chapman et al., 1978; Katsanis et al., 1990). Similarly, behavioral studies have found that patients exhibit impairments in self-monitoring (Kircher and Leube, 2003), increased tactile illusion

vividness (Thakkar et al., 2011), abnormal sense of self (Hecht, 2010), and deficits in action attribution (Farrer et al., 2004).

Lesion and neuroimaging investigations have repeatedly linked the insula and temporoparietal junction (TPJ) to perceptual aberrations, including body-ownership/agency and sensory perception (Berlucchi and Aglioti, 1997; Ionta et al., 2011; Tsakiris et al., 2010; Baier and Karnath, 2008; Blakemore and Frith, 2003). The insula plays a key role in integrating perceptual experiences, affect, and cognition (Kelly et al., 2012; Makris et al., 2006; Chang et al., 2013). The dorsal anterior insula is associated with chemosensory (Pritchard et al., 1999) and socio-emotional processing (Sanfey et al., 2003; Chang et al., 2011). The posterior insula plays a role in pain and sensorimotor processing (Craig, 2002; Wager and Barrett, 2004). Lesions within the insula are linked to somatoparaphrenia, the belief that part or parts of an individual's body belong to someone else (Baier and Karnath, 2008; Vallar and Ronchi, 2009). Multiple lines of evidence indicate that the right insula in particular is involved in body ownership and agency (Karnath and Baier, 2010; Tsakiris et al., 2010; Moro et al., 2016; Vallar and Ronchi, 2009; Hilti et al., 2013), which is consistent with the right hemisphere's dominance for spatial processing (Corbetta and Shulman, 2002).

The TPJ integrates sensory and spatial signals from the body and environment (Blakemore and Frith, 2003; Jackson and Decety, 2004) and is an

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important neural locus for self-processing that involves cognitive aspects of the self (Blanke et al., 2005) and theory of mind (Samson et al., 2004). The right TPJ in particular has been linked to shifts in spatial attention (Shulman et al., 2010), body ownership (Ionta et al., 2011; Tsakiris et al., 2010), and agency (Karnath and Baier, 2010). Damage to the TPJ can result in asomatognosia (i.e. loss of awareness of a body part or limb), anosognosia (i.e. lack of insight into an illness or disability), somatoparaphrenia, and out of body experiences (Berlucchi and Aglioti, 1997; Ionta et al., 2011; Blanke et al., 2002). In healthy subjects, the duration of task-elicited activity in the right TPJ correlates with PAS scores (Arzy et al., 2007) and interference of TPJ activity by transcranial magnetic stimulation (TMS) impairs mental transformation of one's own body (Blanke et al., 2005).

Reduced insula volume is a consistent finding in psychotic disorders, schizophrenia in particular (Glahn et al., 2008). Insular volume reduction correlates with deficits in social cognition and emotion regulation (Giuliani et al., 2011), facial and prosody affect perception (Li et al., 2009; Mitchell et al., 2004), information processing difficulties (White et al., 2010), and sensory deficits such as pain insensitivity (De la Fuente-Sandoval et al., 2010; Wylie and Tregellas, 2010). Although not as extensively studied, reduced TPJ grey matter volume has also been found in psychosis (Honea et al., 2008; Segall et al., 2009) and linked to aberrant sensory perception (Wible, 2012; Spence et al., 1997). Despite the known role of the insula and TPJ in bodily perception/agency, and considerable evidence that these two regions are abnormal in psychosis, the association between perceptual aberrations and grey matter volume of these regions has not been examined. This investigation was undertaken to test the hypothesis that the severity of perceptual aberrations in psychosis inversely correlates with reduced TPJ and insula volumes, particularly in the right hemisphere. We further hypothesized that this relationship would be especially robust in the posterior insula given this region's involvement in somesthesia and perception.

## 2. Methods

### 2.1. Participants

Forty-two healthy subjects and 56 individuals with a psychotic disorder were included in this study. The psychosis group included 40 individuals with a schizophrenia spectrum disorder (schizophrenia, schizoaffective disorder, and schizophreniform disorder) and 16 individuals with bipolar disorder with psychotic features. Patients were recruited from the Vanderbilt Psychotic Disorders Program at Vanderbilt Psychiatric Hospital in Nashville, TN. Healthy subjects were recruited from Nashville and the surrounding area. The study was approved by the Vanderbilt University Institutional Review Board. All study participants provided written informed consent prior to enrolling in the study. The Structured Clinical Interview for Diagnosing DSM-IV disorders (First and Gibbon, 2004) was used to confirm diagnoses in patients and rule out psychopathology in healthy individuals. Study exclusion criteria included age < 16 or > 65, premorbid intellect estimated using the Wechsler Test of Adult Reading (WTAR: Wechsler, 2001) < 70, head trauma, presence of a systemic medical illness or CNS disorder, active substance abuse within the past month, and psychotropic drug use (healthy individuals only).

### 2.2. Study procedures

Participants completed the PAS, a 35 item true/false self-report questionnaire (Chapman et al., 1978), and underwent an MRI session on a Phillips Intera Achieva 3T scanner, which included collection of a T1-weighted anatomical scan (170 sagittal slices, matrix 256 × 256, 1 mm<sup>3</sup> isotropic resolution, TR = 8.0 ms, TE = 3.7 ms). Individuals in the psychotic disorders group were also evaluated with the Positive

and Negative Syndrome Scales (PANSS: Kay et al., 1987) to quantify severity of psychotic symptoms.

### 2.3. Neuroimaging data analysis

Each T1-weighted structural MRI scan was segmented into grey matter, white matter, and CSF using the VBM8 toolbox (<http://dbm.neuro.uni-jena.de/vbm/>) for Statistical Parametric Mapping 8 (<http://www.fil.ion.ucl.ac.uk/spm/software/spm8/>). Following segmentation, native space grey matter images were normalized to the VBM8 T1 template using the high-dimensional DARTEL approach (Ashburner, 2007). Grey matter images were modulated by the non-linear warping component only in order to preserve the volume of the original images after removing the effects of total brain volume. Given strong evidence that the right insula (Ehrsson et al., 2007; Tsakiris et al., 2007; Baier and Karnath, 2008; Karnath and Baier, 2010; Ionta et al., 2011) and TPJ (Ionta et al., 2011; Tsakiris et al., 2010; Karnath and Baier, 2010) are involved in symptoms of disordered bodily perception and ownership, we focused our analysis on regions-of-interest (ROIs) created for these two structures. The insula ROI was derived from Kelly et al.'s (2012) investigation of the functional architecture of the insula. Briefly, they identified between 2 and 15 insula clusters on the basis of clustering and covariance analysis of multimodal neuroimaging data and meta-analyses of fMRI studies. Their 3 cluster solution, which included posterior, ventral anterior, and dorsal anterior sub-divisions corresponds to other studies of insular sub-regions (Chang et al., 2013; Cauda et al., 2011; Deen et al., 2011). The three sub-divisions were combined to create a right whole insula ROI. For the TPJ, we used [neurosynth.org](http://neurosynth.org) (Yarkoni et al., 2011) to identify the voxel with the highest T-value for the search term "right temporoparietal junction". Briefly, [neurosynth.org](http://neurosynth.org) is a web-based platform for conducting meta-analyses on 1000's of published neuroimaging studies based on key terms within the article. In this instance, a meta-analysis of the search term 'right temporoparietal junction' was performed. We selected the voxel with highest activation value for the TPJ search term. This corresponded to MNI coordinates 58, -48, 16. A 10 mm sphere centered on this coordinate served as the right TPJ ROI. Grey matter volume was extracted from the whole insula and TPJ ROIs, and served as the primary dependent variables in the analyses described below.

### 2.4. Statistical analysis

We first compared right TPJ and insula volumes between psychosis patients and healthy control subjects using ANCOVA analyses with age and sex included as covariates. Our primary hypothesis, that right TPJ and insula volumes would be inversely correlated with PAS scores in patients with psychosis, was tested using partial correlation analysis with age and sex entered as covariates. Given that our a-priori hypotheses were unidirectional, all analyses were one-tailed. The threshold for statistical significance was Bonferroni corrected ( $p = 0.025$ ) to correct for the number of ROIs. Significant findings for the whole insula ROI were followed up with exploratory analyses using the 3 insula sub-divisions.

A voxel-wise analysis was also performed to further localize grey matter volume differences between groups and correlations between grey matter volume and perceptual aberrations in patients. The normalized grey matter images were smoothed with a 4 mm kernel and entered into: 1) a between groups analysis comparing patients to healthy subjects; and 2) a regression analysis with perceptual aberrations entered as a predictor of grey matter volume. The VBM analyses were masked to include only voxels in the LONI insula probabilistic atlas (Shattuck et al., 2008) and TPJ ROI, as defined above. The VBM ROI analysis was followed by an exploratory whole-brain analysis. All VBM analyses were thresholded at the cluster-level Family-wise error (FWE) corrected ( $p_{(FWE-corr)}$ )  $p = 0.05$  (corrected for ROI and whole-brain search volumes) for voxel-wise  $p_{(uncorrected)} = 0.005$ .

**Table 1**  
Sample demographics.

Variable	Healthy subjects		Psychosis		Statistics		
	n = 42		n = 56		t/ $\chi^2$	df	p
Sex (M:F)	30:12		36:20		0.557	1	0.456
Ethnicity (W:AA:O)	23:17:2		36:17:3		5.987	3	0.112
SCZ:BD	–		40:16		–	–	–
	Mean	SD	Mean	SD	Statistics		
					t/ $\chi^2$	df	p
Age	29.4	11.4	27.3	9.0	1.01	96	0.315
Education	15.2	2.0	13.5	2.3	3.89	94	<0.001
Parental education	14.3	2.2	14.7	2.6	0.76	91	0.447
PAS	2.3	1.2	7.3	6.7	5.52	59.6	<0.001
WTAR	109.3	13.6	101.2	14.7	2.51	96	0.014
PANSS positive	–		19.2	7.3	–	–	–
PANSS negative	–		15.1	8.2	–	–	–
PANSS general	–		31.0	8.6	–	–	–
APD dose (CPZ equivalents)	–		336.6	251.4	–	–	–

Abbreviations: AA = African American; APD = Antipsychotic Drug; BD = Psychotic Bipolar Disorder; CPZ = Chlorpromazine; df = Degrees of Freedom; F = Female; O = Other; PANSS: PAS: Perceptual Aberration Scale; Positive and Negative Syndrome Scale; W = White.

### 3. Results

Demographics, PAS scores, and clinical data are presented in Table 1. PAS scores were significantly more variable in individuals with a psychotic disorder (Levene's test of equality of variances:  $F = 33.94$ ,  $p < 0.001$ ). Welch's  $t$ -test, which is robust to violations of the homogeneity of variances assumption, confirmed that PAS scores were significantly higher in patients ( $t(59.6) = 5.52$ ,  $p < 0.001$ ). Among psychosis patients, PAS scores were significantly higher in schizophrenia spectrum disorder compared to bipolar disorder with psychotic features ( $t(48.9) = 2.43$ ,  $p = 0.019$ ).

**Table 2**  
Correlations between perceptual aberrations and brain volumes.

	Healthy		Psychosis (n = 56)			
	Controls (n = 42)		All patients		Schizophrenia	
	$r^a$	$p^b$	$r^a$	$p^b$	$r^a$	$p^b$
Right hemisphere						
Whole insula	0.01	0.482	–0.30	0.013	–0.33	0.021
Anterior	0.06	0.361	–0.26	0.027	–0.32	0.024
Dorsal	0.07	0.327	–0.24	0.041	–0.23	0.079
Posterior	–0.22	0.082	–0.32	0.008	–0.38	0.010
TPJ	0.02	0.460	–0.27	0.024	–0.26	0.056
Left hemisphere						
Whole insula	0.04	0.402	–0.35	0.005	–0.42	0.005
Anterior	0.05	0.375	–0.27	0.024	–0.26	0.061
Dorsal	0.14	0.191	–0.23	0.050	–0.31	0.031
Posterior	–0.08	0.307	–0.37	0.003	–0.45	0.002
TPJ	0.08	0.311	–0.12	0.201	–0.07	0.349

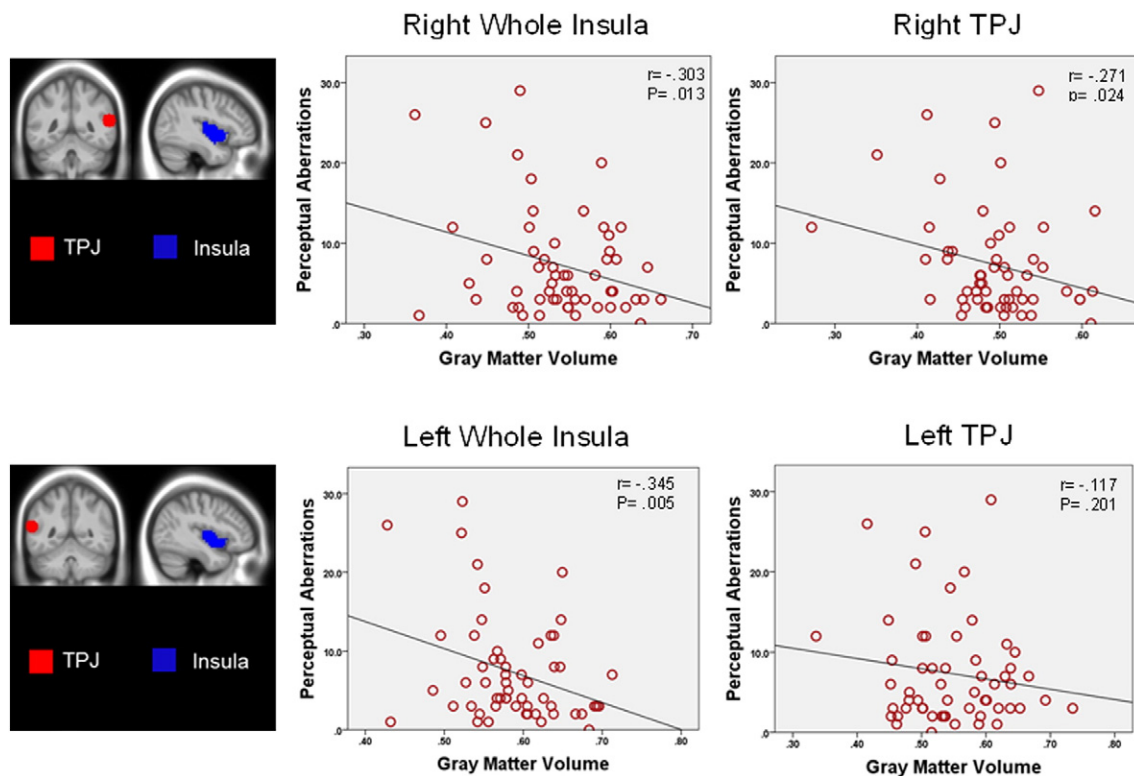
<sup>a</sup> Partial correlation with age and sex entered as covariates.

<sup>b</sup> One-tailed.

#### 3.1. Insula and TPJ ROI analyses

Right TPJ and insula volumes did not differ between patients and healthy subjects (right TPJ:  $F(1,94) = 0.01$ ,  $p = 0.976$ ; right insula:  $F(1,94) = 0.05$ ,  $p = 0.829$ ). Similar results were obtained when the analysis was restricted to patients with a schizophrenia spectrum diagnosis (right TPJ:  $F(1,78) = 0.05$ ,  $p = 0.829$ ; right insula:  $F(1,78) = 0.08$ ,  $p = 0.785$ ).

Scatter plots depicting the correlation between PAS scores and insula and TPJ volumes in psychosis are presented in Fig. 1. Complete results are presented in Table 2. Our primary hypothesis, that right TPJ and insula volumes would inversely correlate with severity of perceptual aberrations in psychosis, was supported (insula:



**Fig. 1.** Relationship between grey matter volume and perceptual aberrations in psychosis. Normalized grey matter volume was extracted from a-priori defined TPJ and insula regions-of-interest (ROIs) shown in red and blue, respectively, and correlated with Perceptual Aberration Scale raw scores.

$r = -0.30, p = 0.013$ ; TPJ:  $r = -0.27, p = 0.024$ ) (see Fig. 1). Results remained unchanged after adding daily dose of antipsychotic medication as an additional covariate (right insula:  $r = -0.27, p = 0.027$ ; right TPJ:  $r = -0.23, p = 0.046$ ). Similar results were obtained when the analysis was restricted to schizophrenia spectrum patients (see Table 2). To determine if the correlations were specific to the right hemisphere, we repeated the partial correlations using left hemisphere ROIs for the TPJ and insula (see Table 2). Briefly, left whole insula volume inversely correlated with PAS scores within the whole sample of psychosis patients ( $r = -0.35, p = 0.005$ ) and the schizophrenia spectrum sub-group ( $r = -0.42, p = 0.005$ ). This

correlation remained significant after controlling for antipsychotic medication ( $r = -0.32, p = 0.010$ ). Left TPJ volume was unrelated to PAS scores in the entire cohort of psychosis patients ( $r = -0.12, p = 0.201$ ) and the schizophrenia spectrum sub-group ( $r = -0.07, p = 0.349$ ).

Results of the insular sub-division analysis are presented in Table 2 and Fig. 2. In brief, PAS scores inversely correlated with all 3 insula sub-divisions in both the left and right hemispheres (all  $r$ 's  $> 0.23, p < 0.050$ ). However, the relationship was strongest for the posterior sub-division of the insula (right:  $r = -0.32, p = 0.008$ ; left:  $r = -0.37, p = 0.003$ ).

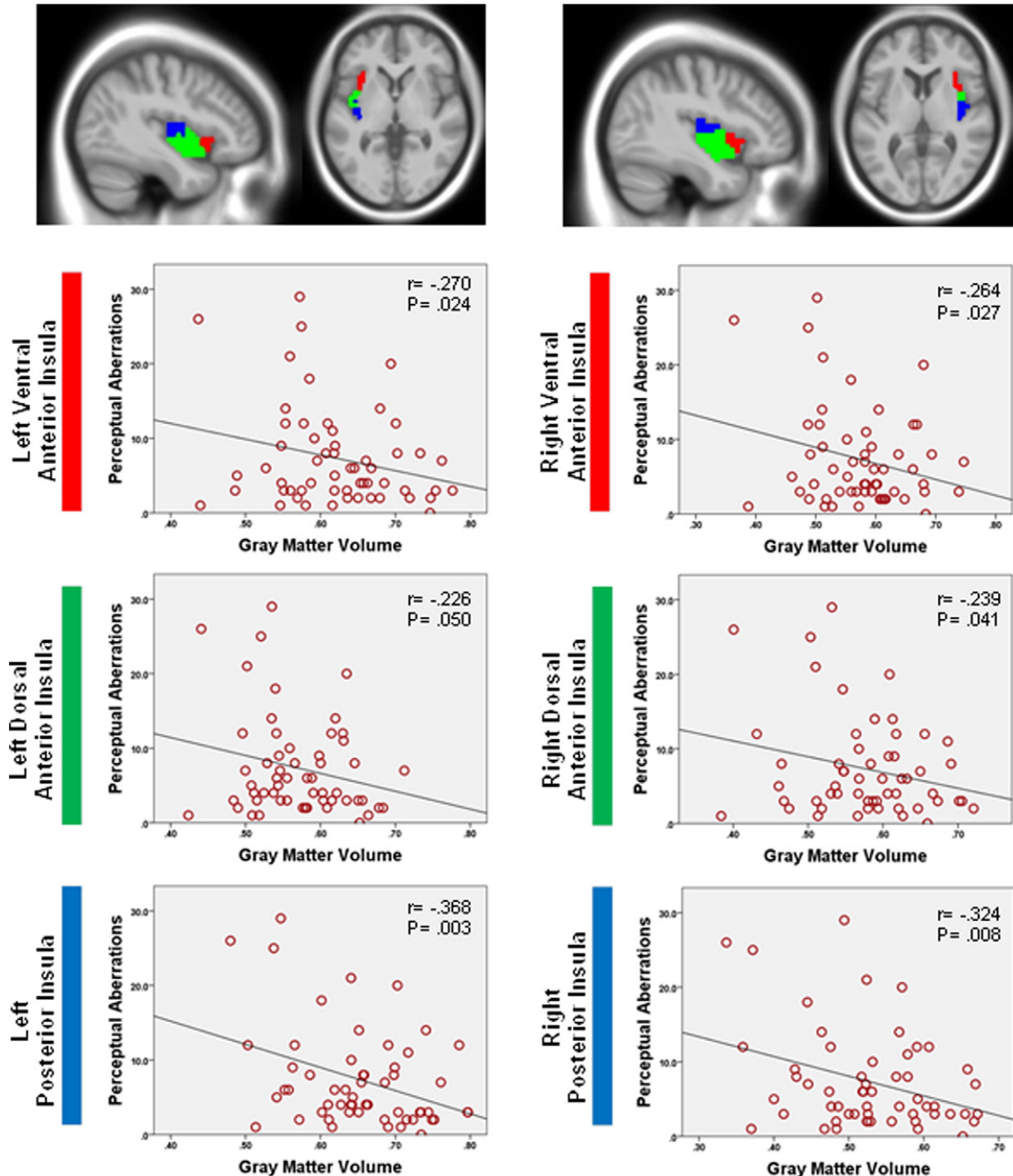


Fig. 2. Scatterplots depicting correlation between perceptual aberrations, as measured with the Perceptual Aberration Scale, and grey matter volume of insula sub-regions in psychosis.

### 3.2. VBM analyses

No significant volumetric differences between patients and healthy subjects were detected within the TPJ and insula ROIs. Within the psychosis patient group, voxel-wise regression analysis restricted to the insula and TPJ ROIs revealed significant inverse correlations between PAS scores and voxel-wise grey matter volume in clusters located in the left insula, including posterior insula, and right TPJ (see Fig. 3). The exploratory, whole-brain voxel-wise regression analysis revealed two clusters located in the left middle orbitofrontal gyrus and posterior region of the left inferior temporal gyrus where reduced grey matter volume correlated with higher PAS scores (see Supplementary Fig. 1). Of note, there were no significant findings for the reverse contrast (i.e. grey matter positively correlated with PAS scores).

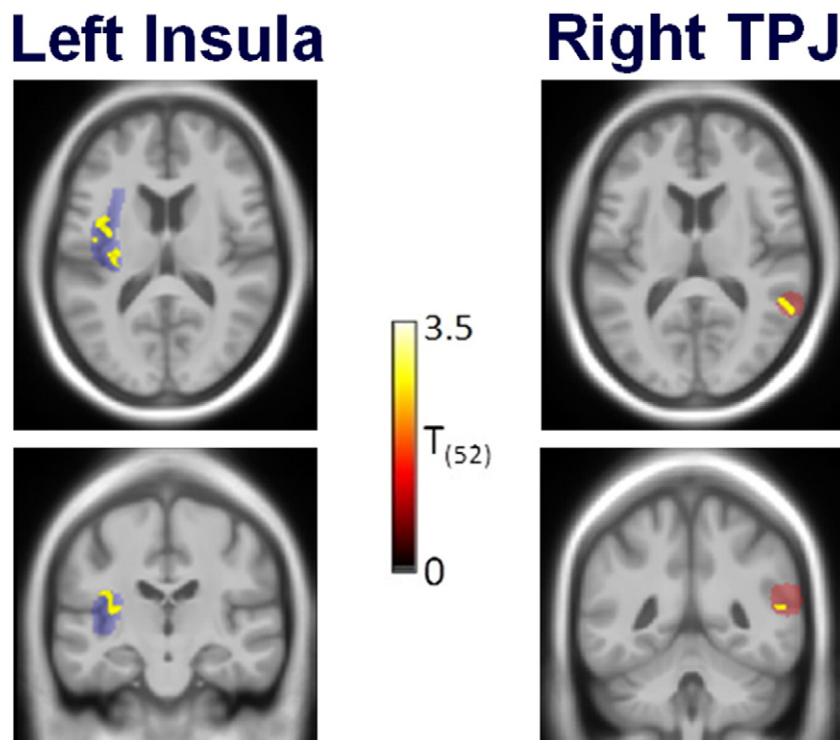
### 4. Discussion

We examined the relationship between self-reported perceptual aberrations and brain structure in psychosis. This investigation was motivated by several factors. First, perceptual aberrations are common in psychosis. Second, structural brain lesions involving the insula and TPJ are associated with disturbances in body ownership and agency. Finally, there is considerable evidence that the structure and function of the insula and, to a lesser extent TPJ, are abnormal in psychosis. Using both ROI and voxel-wise approaches, we identified several correlations between perceptual aberrations and grey matter volume of the insula and TPJ in psychosis. Within the insula, this relationship localized primarily to bilateral posterior insula and left dorsal anterior insula. The right TPJ also showed a significant negative relationship with PAS scores in patients with a psychotic disorder.

The correlation between insula and perceptual aberrations in psychosis is consistent with neuroimaging and lesion studies linking this

brain region to body ownership and awareness (Berlucchi and Aglioti, 1997; Tsakiris et al., 2010; Karnath and Baier, 2010; Ionta et al., 2011; Blanke et al., 2005; Kelly et al., 2012). In particular, the posterior insula is involved in somatosensory, visual, and motor functions (Downar et al., 2002; Francis et al., 2000; Juliano et al., 1983; Karnath et al., 2000; McGlone et al., 2002; Showers and Lauer, 1961; Treede et al., 2000). The dorsal anterior insula integrates sensory and interoceptive feedback from the posterior insula with emotional and cognitive responses to the same stimuli (Craig, 2009). This integrative function of the dorsal anterior insula could indicate its relationship to aberrant somatosensory perception in a region mapping with cognitive functioning (Nelson et al., 2010; Mutschler et al., 2009) and emotional domains (Kelly et al., 2012). VBM analysis revealed significant clusters in the left ventral anterior and posterior insula in patients with a primary psychotic disorder. There is functional overlap between the regions, both are involved in perception, somesthesia, interoception, air hunger, and action (Kurth et al., 2010; Kelly et al., 2012), which may explain why significant effects were found in both areas.

Given the dominant role of the right hemisphere in spatial processing, body ownership, and sensory perception (Tsakiris et al., 2010; Karnath and Baier, 2010), we hypothesized that the relationship between perceptual aberrations and insula volume would be limited to, or at least stronger, in the right hemisphere. Unexpectedly, the correlation between perceptual aberrations and insula volume was marginally stronger for the left hemisphere, both in the ROI and VBM analysis. While lesion studies generally implicate the right hemisphere in body-ownership and agency, this relationship may not be unique to the right hemisphere (Baier and Karnath, 2008; Nathanson et al., 1952; Cutting, 1978; Starkstein et al., 1992; Baier and Karnath, 2005). Studies using intracarotid barbiturate injections in subjects undergoing WADA testing have found that anosognosia, while more frequent following right hemisphere injection, is also sometimes detected after left hemisphere injection (Breier et al., 1995). Moreover, it is important to note



**Fig. 3.** VBM analysis revealed significant clusters in the left insula and right TPJ regions-of-interest where grey matter volume inversely correlated with perceptual aberrations in psychosis. Insula and TPJ ROIs are shown in blue and red, respectively. Left insula clusters located at MNI coordinates  $-30 -19 10$  (cluster size = 166 voxels,  $t(52) = 3.74$ , cluster-level  $p_{(FWE-corrected)} = 0.045$ ) and MNI coordinates  $-35 -4 10$  (cluster size = 216 voxels,  $t(52) = 3.57$ , cluster-level  $p_{(FWE-corrected)} = 0.021$ ). Right TPJ cluster located at MNI coordinates  $59-52 9$  (cluster size = 63 voxels,  $t(52) = 3.65$ , cluster-level  $p_{(FWE-corrected)} = 0.032$ ).

that while functional imaging studies tend to find greater activation of the right insula, hemispheric interactions are rarely explicitly tested.

In contrast, the relationship between perceptual aberrations and TPJ was limited to the right hemisphere. Saxe and Wexler (2005) have theorized that the left and right TPJ may offer different contributions to theory of mind; the right TPJ being more involved in external sensory events and transitive mental states, while the left TPJ is involved in attribution of socially relevant traits to others. Heberlein et al. (2004) found that patients with damage to the left TPJ are selectively impaired in the attribution of personality traits to others. Conversely damage to the right TPJ can result in aberrations of physical perception such as anosognosia, asomatognosia, or somatoparaphrenia (Heberlein et al., 2004). Other studies have found that the right TPJ is active when individuals distinguish themselves from others (Decety and Sommerville, 2003) and is involved in agency, which involves integrating external sensory signals with a self-produced comparison of one's bodily actions (Ruby and Decety, 2001; Farrer and Frith, 2002; Farrer et al., 2003).

Exploratory whole brain VBM analysis revealed significant negative correlations between PAS scores and grey matter volume in the left middle orbitofrontal gyrus and posterior left inferior temporal gyrus. The middle orbitofrontal gyrus receives input from the visual, taste, olfactory, and somatosensory systems (Zald and Kim, 1996b; Zald and Kim, 1996a) and is involved with reward processing and reinforcement related to these sensations (Rolls, 2000). This region has been linked to body ownership (Limanowski and Blankenburg, 2016) and exhibits cortical thinning in schizophrenia (Kuperberg et al., 2003; Convit et al., 2001; Crespo-Facorro et al., 2000). The inferior temporal gyrus also shows significant grey matter reduction in schizophrenia as well (Kuperberg et al., 2003; Shenton et al., 2001) and is associated with vestibular illusions like "dizziness, swinging, and spinning, sinking feeling, levitation, and lightness" (McCabe et al., 2008).

This study has a number of limitations. Contrary to other investigations (Glahn et al., 2008; Honea et al., 2008; Segall et al., 2009) we did not find volumetric differences between healthy controls and patients with a psychotic disorder in the Insula and TPJ. However, a previous investigation of a larger cohort of psychosis patients, from which the subset of patients included in this study was drawn from, did find reduced insula volume in psychosis (Woodward and Heckers, 2015). It's possible our study lacked sufficient power to detect insula volume reduction. Moreover, our psychosis sample also included bipolar patients, and both schizophreniform and chronic patients which may have obscured group differences in insula and TPJ volumes. Another drawback to this study is the lack of a functional task eliciting aberrant perception. Although, beyond the scope of this study, these results provide support for pursuing those methodologies in the future.

Supplementary data to this article can be found online at <http://dx.doi.org/10.1016/j.schres.2016.10.005>.

#### Contributors

Authors EB and NDW conceived and designed the research reported on, analyzed the data, and co-wrote the final draft of the manuscript. Author NDW provided funding for the project.

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#### Conflict of interest

No commercial support was received for this manuscript and the authors have no conflicts of interest to report.

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