Abnormal Angular Gyrus Asymmetry in Schizophrenia

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Abstract

Objective—Few studies have evaluated the parietal lobe in schizophrenia despite the fact that it has an important role in attention, memory, and language—all functions that have been reported to be abnormal in schizophrenia. The inferior parietal lobule, in particular, is of interest because it is not only part of the heteromodal association cortex but also is part of the semantic-lexical network, which also includes the planum temporale. Both the inferior parietal lobule, particularly the angular gyrus of the inferior parietal lobule, and the planum temporale are brain regions that play a critical role as biological substrates of language and thought. The authors compared volume and asymmetry measures of the individual gyri of the parietal lobe by means of magnetic resonance imaging (MRI) scans.

Method—MRI scans with a 1.5-Tesla magnet were obtained from 15 male chronic schizophrenic and 15 comparison subjects matched for age, gender, and parental socioeconomic status.

Results—Inferior parietal lobule volumes showed a leftward asymmetry (left 7.0% larger than right) in comparison subjects and a reversed asymmetry (left 6.3% smaller than right) in schizophrenic subjects. The angular gyrus accounted for this difference in asymmetry, with the left angular gyrus being significantly larger (18.7%) than the right in comparison subjects, a finding that was not observed in schizophrenic patients. A further test of angular gyrus asymmetry showed a reversal of the normal left-greater-than-right asymmetry in the schizophrenic patients.

Conclusions—Patients with schizophrenia showed a reversed asymmetry in the inferior parietal lobule that was localized to the angular gyrus, a structure belonging to the heteromodal association cortex as well as being part of the semantic-lexical network. This finding contributes to a more comprehensive understanding of the neural substrates of language and thought disorder in schizophrenia.
has been proposed as the site of the key abnormality in schizophrenia (7). Further, both the supramarginal gyrus and the angular gyrus have been described as part of a semantic-lexical network that supports “word meanings” represented by a “grid of connectivity” that constitutes a “final pathway for the chunking of words into thought” (1). The role of the inferior parietal lobule, and especially the angular gyrus, in language comprehension has been further confirmed by functional MRI (fMRI) and positron emission tomography studies (8-10). Moreover, the semantic-lexical network proposed by Mesulam (1) includes both the inferior parietal lobule and the planum temporale, the latter being located on the superioposterior surface of superior temporal gyrus, which, in turn, has been correlated with symptoms of thought disorder (5,6,11). More anteriorly, the superior temporal gyrus has been correlated with auditory hallucinations (5,6,12).

Of note, the inferior parietal lobule and neighboring cortical regions often exhibit marked lateral asymmetry (13,14) and belong to structures that support language. The presence of left-greater-than-right asymmetry also appears to be important for normal language development. For example, the absence of normal lateralization in these regions has been reported in subjects with autism (15,16) as well as in other language disorders (17). In schizophrenia, a reversal of normal asymmetry in the superior temporal gyrus, including the planum temporale, has been associated with thought disorder (5,6). Thus, a reversal in normal asymmetry may be importantly related to schizophrenic pathology (18-20). Consistent with these findings, we hypothesized that a reversal of normal asymmetry in the superior temporal gyrus would also be found in the inferior parietal lobule. Furthermore, since cortical asymmetries are present during fetal development (21), finding an absence or a reversal of normal asymmetry might indicate a disruption in neuronal development.

The relevance of the angular gyrus and supramarginal gyrus to schizophrenia stems not only from their functions as part of the heteromodal association cortex but also from their reciprocal neuroanatomical connections to the prefrontal (22) and temporal lobes (23)—brain regions that have been shown to have disease-related abnormalities. We examined these neuroanatomical relationships by capitalizing on the opportunity of having volumetric data on prefrontal and temporal lobe regions in the same subjects (11,24,25). If the hypothesis that schizophrenia is a disorder characterized primarily by heteromodal association cortex abnormalities is correct, then higher correlations between the inferior parietal lobule and other areas interconnected with the heteromodal association cortex would be predicted for schizophrenic patients but not for comparison subjects. Also, in an exploratory analysis, we examined the relationship between the heteromodal inferior parietal lobule and formal thought disorder, as well as neuropsychological measures of attention and memory, all of which are likely to contribute to thinking and judgment—functions commonly associated with the inferior parietal lobule.

METHOD

Subjects

Fifteen right-handed male subjects with chronic schizophrenia were recruited from the Veterans Affairs Medical Center in Brockton, Mass. (11). Diagnoses were determined by using DSM-III-R criteria from the Schedule for Affective Disorders and Schizophrenia (26) and from hospital chart reviews. All patients were receiving neuroleptic medication equivalent to a mean of 881 mg of chlorpromazine per day. The patients had spent a mean of 7.1 years (SD=4.6) in the hospital, with a mean duration of illness of 15.8 years (SD=8.8).

Fifteen normal comparison subjects were matched to the patient group on gender (male), handedness (right), age (20–55 years, matched within 2 years), and parental socioeconomic status. No comparison subjects had a history of mental illness nor did their first-degree relatives. The exclusion criteria for both groups included no major drug or alcohol abuse in the...
previous 5 years, no history of electroconvulsive therapy, no neurological illness, and no
medications known to affect brain MRI scans (e.g., steroids). There were no statistical
differences between the comparison and schizophrenic subjects in height, weight, head
circumference, or scores on the WAIS-R information subscale (27). Written informed consent
was obtained from all 30 subjects after the procedures had been fully explained.

Clinical and Neuropsychological Measures

Eleven patients were categorized as having mostly positive symptoms according to the Scale
for the Assessment of Positive Symptoms (28), none had mostly negative symptoms according
to the Scale for the Assessment of Negative Symptoms (29), and four were rated as having
mixed negative and positive symptoms. The mean score on the Thought Disorder Index (30)
for the patients was 60.4 (SD=61.8), on which normal subjects score no greater than 5.
Schizophrenic patients were evaluated by means of both standardized and experimental
neuropsychological procedures (31).

MRI Methods

Image acquisition—A detailed description of the parcellation rules and anatomical
definitions used in this study is included in appendix 1. The MRI images were acquired on a
Signa 1.5-T system (GE Medical Systems, Milwaukee). A sagittal localizer was used to orient
the images in the proper plane. Total brain volume was obtained by means of a double-echo,
spin-echo acquisition with the following parameters: TR=3000 msec, TE=30 and 80 msec,
field of view=24 cm, acquisition matrix=256 × 256 pixels, and voxel dimensions=0.9 mm ×
0.9 mm × 3 mm, which resulted in 108 contiguous double-echo axial slices (54 levels).

For the inferior parietal lobule brain regions, a three-dimensional Fourier transform spoiled
gradient recall acquisition was used with the following parameters: TR=35 msec, TE=5 msec
with one repetition, flip angle=45°; field of view=24 cm; matrix=256 × 256 × 124 (192 phase-
encoding steps). The dimensions for each voxel were 0.9 mm × 0.9 mm × 1.5 mm. The images
were reformatted into 124 contiguous 1.5-mm coronal slices.

Image processing—Image processing was completed on workstations (Sun Microsystems,
Mountain View, Calif.) with several multi-step computer algorithms. A postprocessing filter
was used to reduce noise and to enhance morphologic details (32), followed by a
semiautomated segmentation algorithm used to separate gray matter, white matter, and CSF
(33). Brain volumes for gray matter, white matter, and CSF were then calculated by summing
the voxels for each of these tissue classes across all brain slices (11,24).

Reliability—Raters were blind to subject diagnosis for all measures. Both inter- and intrarater
reliability were measured for each of the parietal regions by using an intraclass correlation
coefficient. For interrater reliability, three judges (R.D., D.V.I., and J.L.) measured each of the
parietal regions on 10 coronal slices (two sets of five contiguous slices) on three randomly
selected brains, thus producing six measures for each parietal region (i.e., a left and right
measure for each of three brains). For these six measures in each parietal region, intrarater
reliabilities were 0.96 for the inferior parietal lobule, 0.96 for the superior parietal gyrus, and
0.97 for the postcentral gyrus. Intrarater reliabilities, computed by using all of the slices from
one randomly selected brain and measured by one rater (R.D.) at two separate times
(approximately 1 year apart), were 0.97 for the inferior parietal lobule, 0.98 for the superior
parietal gyrus, and 0.94 for the postcentral gyrus.

Statistical Analysis

All measures were corrected for total intracranial volume (unless otherwise mentioned) in order
to control for variations in head size. To test for group volume differences, t tests uncorrected
for multiple comparisons were conducted on the total volume of the parietal lobe, superior parietal gyrus, and the postcentral gyrus. However, to test the main hypothesis of group volume differences within the inferior parietal lobule, a repeated measures analysis of variance (ANOVA) was used with diagnosis as a between-group factor (patient versus normal comparison group) and two within-group factors of laterality (left versus right) and region (angular gyrus and supramarginal gyrus). To follow up on the laterality-by-group interaction, asymmetry scores were also computed for each region of interest by using the following formula: (left-right/\text{left+right}) \times 100.

Pearson correlations for both absolute and relative values were used to examine the relationship between the volumes of parietal lobe gray matter and anatomically connected regions of the prefrontal (24,25) and temporal gray matters (11). Because of the nonnormal distribution of clinical measures, an additional exploratory analysis was performed in which nonparametric Spearman rank order tests were used to test for significant correlations between the parietal lobe gray matter volumes and clinical and neuropsychological data. Again, we note that these correlations were exploratory in nature and not our major focus of interest.

RESULTS

MRI Volume Comparisons

For comparative purposes, table 1 provides volumes for all parietal regions studied. As predicted, no group volume differences were found for the total parietal lobe, superior parietal gyrus, or the postcentral gyrus. However, volume differences were noted between the two groups for the inferior parietal lobule, specifically the angular gyrus.

A repeated measures ANOVA showed no overall group volume differences or an overall laterality effect (table 2). However, there was a significant laterality-by-group interaction that indicated a difference in asymmetry between the two groups. As shown in figure 1, the comparison subjects had a leftward asymmetry (left inferior parietal lobule volume 7.0% larger than the right), and the schizophrenic patients showed a reversed asymmetry (left inferior parietal lobule volume 6.3% smaller than the right).

The volumes of the angular gyrus and the supramarginal gyrus for the two groups were significantly different (table 2). There was also a significant region-by-group interaction. Post hoc tests indicated that schizophrenic patients had a larger right angular gyrus than the normal comparison group (table 1). In the comparison subjects, the left angular gyrus volume was considerably larger (18.4%) than the right (paired t=2.71, df=14, p=0.02), whereas in the patients, the left angular gyrus was not significantly different (4.7% less volume) from the right (t=0.7, df=14, p>0.05).

Left-Right MRI Volume Asymmetries

To evaluate further the aforementioned laterality-by-group interaction of the inferior parietal lobule as well as the laterality of the other parietal regions, we computed asymmetry coefficients by using the formula (left–right)/(\text{left+right}) \times 100 (note that the asymmetry coefficient is dimensionless). A negative value indicates a larger right than left side volume. Student’s t tests were used to compare asymmetry coefficients between the two groups for all of the regions.

There was a significant difference between the two groups in the asymmetry coefficient for the total parietal lobe (table 3). The comparison group exhibited a leftward asymmetry, with the left parietal lobe 6.0% larger than right (paired t=3.18, df=14, p=0.007), while the schizophrenic group exhibited virtually no total parietal asymmetry (t=−1.1, df=14, p>0.30).
Neither group exhibited significant asymmetry of the superior parietal gyrus, and both left and right postcentral gyrus volumes were similar in the schizophrenic and comparison groups. Both groups showed a significant leftward asymmetry, with the left postcentral gyrus being 14.8% larger than the right in the comparison subjects (paired t=2.76, df=14, p=0.02), and 13.4% larger in the schizophrenic patients (paired t=3.00, df=14, p=0.01).

A significant group difference in asymmetry coefficient was evident for the inferior parietal lobule (table 3), which was accounted for by a group difference in the angular gyrus. Note that all of the schizophrenic subjects except two had asymmetry coefficients below the mean (and median) of the subjects in the normal comparison group (figure 2). The supramarginal gyrus did not show a significant asymmetry for either group. Thus, although the post hoc tests based on the ANOVA group-by-region interaction did not show statistically significant group differences between left and right angular gyrus in the schizophrenic patients, the differences between groups in the asymmetry coefficient suggests that there is a reversal of the normal asymmetry in the schizophrenic patients for the angular gyrus.

Correlations Between Parietal Gray Matter Volumes and Anatomically Connected Regions in Prefrontal and Temporal Cortex

For all correlations, significance levels were set at p≤0.05 (two-tailed), which corresponded to r>0.51 for the 15 schizophrenic subjects and r>0.53 for 14 comparison subjects (one comparison subject was dropped because artifact in the prefrontal cortex made this region too difficult to assess the volume accurately). In addition, the values reported here are for absolute volumes, but the correlations were considered significant only if they reached p≤0.05 for both relative and absolute volumes. For all of these correlations, parietal cortex regions included the right and left inferior parietal lobules, superior parietal gyrus, and postcentral gyrus; prefrontal cortex regions included right and left superior frontal, middle frontal, inferior frontal, and orbital frontal gyri; and temporal cortex regions included right and left anterior superior temporal gyrus, posterior superior temporal gyrus, parahippocampal gyrus, and amygdala-hippocampal complex.

High correlations were found between left and right inferior parietal lobule volumes in both schizophrenic (r=0.68, p<0.005) and comparison subjects (r=0.54, p<0.04), as well as between the left and right superior parietal gyrus (schizophrenic subjects: r=0.81, p<0.001; comparison subjects: r=0.75, p<0.001). Additionally, the comparison subjects showed a significant correlation between the left and right postcentral gyrus (r=0.63, p<0.01).

As hypothesized, the schizophrenic group showed several high correlations between gray matter volumes of the inferior parietal lobule and regions of the prefrontal cortex (table 4). Figure 3 provides an illustrative summary of these correlations, with the bottom arrows depicting prefrontal correlations with the left inferior parietal lobule, and the top arrows depicting prefrontal correlations with the right inferior parietal lobule (note: arrows do not imply direction). In addition to these correlations, the left postcentral gyrus correlated significantly with the left superior frontal gyrus (r=0.72, p<0.003) and with the right inferior frontal gyrus (r=0.70, p<0.005). In contrast, the comparison subjects showed no volumetric correlations between inferior parietal lobule volumes and prefrontal volumes at p<0.05.

The differences between groups in the correlations for respective brain areas were tested by using a Fisher’s z transformation. Of note, significant group differences in correlations emerged for the left inferior parietal lobule and prefrontal structures even though the correlations between left and right inferior parietal lobules and the prefrontal lobe structures were comparably high in the schizophrenic group. This result highlights the salience of the left inferior parietal lobule correlations with prefrontal measures in the schizophrenic group.
In the comparison subjects, but not in the schizophrenic subjects, the inferior parietal lobule asymmetry coefficient correlated inversely with all of the prefrontal structures ($r = -0.50$ to $-0.76$, $p = 0.03$ to $\leq 0.001$). In addition, the left postcentral gyrus correlated significantly with several prefrontal structures: the left superior frontal gyrus ($r = 0.83$, $p < 0.001$), left orbital gyrus ($r = 0.70$, $p < 0.004$), right superior frontal gyrus ($r = 0.64$, $p < 0.01$), and the right orbital gyrus ($r = 0.64$, $p < 0.01$).

In the schizophrenic group, both the left and right inferior parietal lobules correlated significantly with the left amygdala ($r = 0.67$, $p < 0.007$) and the left anterior portion of the superior temporal gyrus ($r = 0.66$, $p < 0.008$ (figure 4 and table 5). In the normal comparison group, the one significant correlation was between left inferior parietal lobule and the left amygdala ($r = 0.53$, $p = 0.04$). The group differences in the respective, pairwise correlations, as tested with Fisher’s formula, did not reach statistical significance.

**Correlations With Clinical Status and Clinical Measures and With Neuropsychological Data**

Because of the exploratory nature of the correlations and the issue of multiple tests (i.e., some of the significant results might be due to chance), we report only correlations where $p$ was below 0.01.

No correlations between regions of interest and clinical measures in schizophrenic patients reached the significance level of $p < 0.01$. Exploratory analyses of correlations between neuropsychological measures and brain volume in schizophrenia patients indicated that reduced right inferior parietal lobule volume correlated with lower scores on tests of visual attention and visual memory (Trails B: $r_c = -0.76$, $p < 0.002$; visual reproduction I: $r_c = 0.71$, $p < 0.003$; visual reproduction II: $r_c = 0.84$, $p < 0.001$). Additionally, reduced inferior parietal lobule asymmetry was related to poor performance on Trails A ($r_c = -0.70$, $p < 0.005$) and Trails B ($r_c = -0.69$, $N = 14$, $p < 0.004$).

**DISCUSSION**

The present study examined gray matter volume in individual gyri of the parietal lobe by using high resolution MRI (1.5-mm thick slices) and neuroanatomically based boundary definitions. The major findings from this study were that schizophrenic patients, in contrast to a normal comparison group, showed 1) a reversal of the normal left-greater-than-right asymmetry in the inferior parietal lobule that was localized to the angular gyrus and 2) significant volumetric correlations between parietal lobe regions and regions of the frontal and temporal cortex.

More specifically, gray matter abnormalities observed in the inferior parietal lobule consisted of a reversal of the normal left-greater-than-right asymmetry that was localized to the angular gyrus and further confirmed with a measure of asymmetry. The specificity of this asymmetry finding was underscored by the fact that no differences were observed between schizophrenic and comparison subjects for total parietal lobe, superior parietal gyrus, or postcentral gyrus. Further, there was also an absence of group differences in the lateralization patterns for superior parietal gyrus and postcentral gyrus. This finding supports previous reports of abnormality within the parietal lobe (34-37) and localizes the abnormality to the angular gyrus. Of note, schizophrenic patients did not show leftward lateralization of the total parietal lobe, which was present in the comparison subjects. This result was driven by the reversal of the normal left-greater-than-right asymmetry within the angular gyrus. This finding further underscores the utility of separate volumetric analyses for structures comprising the parietal lobe.

The present study is, to our knowledge, the first to report a reversal of normal asymmetry in the angular gyrus in schizophrenic patients, a brain region belonging to the neural circuitry that supports semantic aspects of language processing. As noted in our introduction, recent fMRI
data suggest the involvement of the inferior parietal lobule, and especially the angular gyrus, in semantic processing. This region, in fact, is regarded as part of a semantic-lexical network that includes the planum temporale and is involved in both assigning meaning to strings of sounds and, at its output stage, in generating associative links responsible for constructing complex meanings and thought processes. Previous studies in schizophrenia have reported abnormal cortical asymmetry in superior temporal gyrus, especially in the region of the planum temporale (38). Thus, the present finding, which extends the finding of abnormal asymmetry to the angular gyrus, enhances our understanding of the neural underpinnings of a core feature of schizophrenic syndrome: disordered thought and language processes.

The relationship between asymmetry of the planum temporale and that of the angular gyrus, the two structures belonging to the language network, has also been previously noted by Eidelberg and Galaburda (13). These investigators reported a correlation between the degree of lateralization of the planum temporale and the angular gyrus. Thus, there appears to be a leftward lateralization for structures specialized for language (7,13,18,19). The presence of such asymmetries might have an evolutionary advantage in developing and supporting language in the human species, in which the specialized function of one hemisphere might confer additional advantage (18-20).

The reversal of normal asymmetry in the angular gyrus, in addition to previous reports of the reversed asymmetry in superior temporal gyrus, provides further support for the relationship between abnormal laterality patterns and the origins of schizophrenic pathology (18-20). Since cortical asymmetries are present during fetal development (13,21), it is thus quite possible that abnormal asymmetries in these two regions in schizophrenia may have a common neurodevelopmental origin.

The affected structures might be abnormal as a result of faulty developmental mechanisms such as gliosis or pruning, the disease process itself, or both. In fact, striking correlations between the inferior parietal lobule and neuroanatomically connected cortical regions of the prefrontal cortex and the temporal lobe, which were observed only in the schizophrenic subjects, provide support for the existence of a pathologic process (such as excitotoxicity) that affects multiple, functionally interconnected brain regions in schizophrenia (39-41). The high correlations between regions that constitute primarily heteromodal association cortex regions, and observed only in the patient group, support the notion that schizophrenia might preferentially affect the association cortex.

Also of interest is the finding that in comparison subjects, the asymmetry coefficient was inversely correlated with frontal lobe volumes, which indicates that a normal brain’s development entails leftward lateralization of language structures (13,18,19), and that this lateralization pattern is correlated in a healthy brain with the frontal lobes, which are intimately involved in mediating processes in parietal language areas.

A limitation of this study is the use of multiple tests, which elevates the risk of type I errors. However, we focused our attention on the inferior parietal lobule, where we had an a priori rationale for evaluating both the angular and supramarginal gyri. Furthermore, we used more conservative criteria for the significance level (i.e., p≤0.01) of the exploratory analyses as a compromise for the multiple correlations performed.

In summary, this study, which employed improved methods of parcellation and measurement of the gray matter of the parietal lobe, provides evidence for the absence of normal left-greater-than-right asymmetry in the angular gyrus, a brain region categorized functionally as part of the heteromodal association cortex and, importantly, linked to aspects of semantic processing. These findings, taken together, provide support for localized volumetric changes in schizophrenia associated with selective cognitive impairments and a possible
neurodevelopmental component to schizophrenic pathogenesis. These findings also afford a more comprehensive understanding of schizophrenic pathology by demonstrating similar pathologic processes that affect functionally related brain areas: the inferior parietal lobule region and the temporal and frontal areas.

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APPENDIX 1. Delineating Regions Within the Parietal Lobe

For parcellating the parietal lobe we first used a reslice editing algorithm to reformat images into the sagittal and axial plane. Specific markings were then made on the sagittal slices in order to define specific neuroanatomical boundaries for delineating the gyri. These markings were then recreated on the original three-dimensional Fourier transform spoiled gradient recall acquisition coronal images, where the regions of interest were completed by using manually guided tracing. The final editing step was a surface-rendering algorithm that was used to create three-dimensional views of the relevant structures (42). The three-dimensional images of the regions of interest could then be viewed individually or within the context of the entire cortex, whereby images could be rotated around x, y, and z axes in order to achieve the best possible visualization of each region of interest. After examining the three-dimensional images, any necessary corrections were made on the original region of interest tracing. Once editing was complete, volumetric measures were derived for each region of interest, as was done for the whole brain volume, by summing the voxels for each region of interest across all relevant slices.

Defining Regions of Interest

The boundaries for the regions of interest in this study were determined with the assistance of an anatomical atlas (43). For all boundaries that involved cutting planes, we corrected for head rotation (tilt) around all three axes. Head rotation about the fronto-occipital axis was measured by a line drawn perpendicular to the interhemispheric fissure on a coronal slice at the level of the parietal lobe. Head rotation about the vertical (z) axis was measured by a line drawn perpendicular to the interhemispheric fissure on an axial slice at the level of the parietal lobe. Head rotation about the bitemporal axis was measured by a line drawn from the most anterior point of the corpus callosum to the most posterior point (illustrated in figure 5) on a midsagittal slice. This reference line was more reliably determined than an anterior to posterior commissure line, which was verified as being virtually parallel with the callosal line (mean difference angle less than 3°) for the 30 cases reported here. After correction for rotation about the fronto-occipital and vertical axes, all brain rotation about the bitemporal axis was corrected to match the brain with the least rotation (brain with the callosal line most nearly horizontal).

Parietal Lobe

Medial surface

The parietal lobe is bounded by the frontal lobe and occipital lobe and by the cingulate gyrus (figure 6). We defined the frontoparietal border by the central sulcus (label 1) and the marginal ramus of the cingulate sulcus (label 2) and, since the sulci do not intersect, by a vertical line that extended from the most posterior portion of the central sulcus to the cingulate sulcus (line A). This line was extended laterally in the coronal plane (perpendicular to the sagittal plane).
The parieto-occipital fissure (label 4) was a clear anatomical boundary that separated the parietal and occipital lobes. The parietal lobe and cingulate gyrus were bounded anteriorly by the subparietal sulcus (label 3). In the absence of a clear anatomical division, we defined the posterior and ventral parietocingulate border by a vertical line (line B) that extended from the subparietal sulcus to the occipitoparietal fissure. This line was drawn on the coronal slice 13.5 mm (nine slices) posterior to the most posterior point of the callosum. This line was extended laterally in the coronal plane (perpendicular to the sagittal plane).

Lateral surface

The lateral boundaries between the parietal lobe and the frontal, temporal, and occipital lobes can be seen in the three-dimensional surface renderings of the cortex in figure 7.

Anteriorly, the central sulcus (label 1) is seen as the parieto-frontal lobe boundary. The Sylvian fissure (label 4) bounded the parietal and temporal lobes anteriorly. More posteriorly, the ventral boundary of the parietal lobe was defined by the three planes (labeled A, B, and C), which are all perpendicular to the sagittal plane. Plane A began at the dorsal level of the Sylvian fissure on the most posterior coronal slice of the postcentral gyrus and continued posteriorly by using the same vertical (z) position for 15 mm (10 coronal slices).

Plane B of the ventral parietal boundary was defined by two parallel lines. The first line was drawn on a midsagittal slice from the most anterior point of the corpus callosum and extending posteriorly at a 9° angle to the callosal reference line from figure 5 (figure 8, left image). The second line that defined this plane was drawn on a more lateral sagittal slice and used the same coordinates as the first boundary line (figure 8, right image). The 9° angle between the reference line and the boundary line was selected so as to include the maximum amount of parietal lobe gray matter without including any (or at least only minimal amounts of) temporal or occipital lobe tissue. The posterior boundary of the parietal lobe was the plane C in figure 7, defined by two parallel lines. The first line was drawn through the parieto-occipital fissure on a midsagittal slice (figure 9, left image). The second line was drawn on a more lateral sagittal slice by using the same coordinates as the first line (figure 9, right image).

Left and right parietal hemispheres were separated by the interhemispheric fissure.

Parcellation of the Parietal Lobe

The regions of interest within the parietal lobe include the postcentral gyrus, superior parietal gyrus, and inferior parietal lobule. The inferior parietal lobule consists of the angular gyrus and the supramarginal gyrus. As can be seen in figure 7, the postcentral gyrus was separated from the superior parietal gyrus and the inferior parietal lobule by the postcentral sulcus (label 2). The inferior parietal lobule was separated from the superior parietal gyrus by the intraparietal sulcus (label 3). The inferior parietal lobule was further subdivided into the angular gyrus and the supramarginal gyrus. In the absence of a clear, consistent anatomical boundary between the angular gyrus and the supramarginal gyrus, the bound between these two inferior parietal lobule structures was defined by the coronal slice midway between the most posterior and most anterior coronal slices of the inferior parietal lobule. For this reason it is possible that some volumetric data points categorized as belonging to the angular gyrus might be a part of the supramarginal gyrus and conversely, some supramarginal gyrus volumetric data could have been categorized as belonging to the angular gyrus. Subtle bias in the final volumetric data is thus conceivable. Data from other laboratories as well as data coming from this laboratory from a different sample should help resolve this issue.
REFERENCES


29. Andreasen, NC. Scale for the Assessment of Negative Symptoms (SANS). University of Iowa; Iowa City: 1981.


FIGURE 1. Differential Left-Versus-Right Asymmetry in Inferior Parietal Lobule Volume in Male Patients With Schizophrenia and Healthy Male Comparison Subjects
FIGURE 2. Asymmetry Coefficients for the Inferior Parietal Lobule in Male Patients With Schizophrenia and Healthy Male Comparison Subjects
FIGURE 3. Gray Matter Volume Correlations Between the Inferior Parietal Lobule and Regions of the Prefrontal Cortex in 14 Male Schizophrenic Subjects\textsuperscript{a}

\textsuperscript{a} Solid lines indicate significance at p<0.01; dotted line indicates significance at p<0.05.

a Solid lines indicate significance at p<0.01; dotted line indicates significance at p<0.05.
FIGURE 5. Midsagittal MRI Slice Depicting Measurement of Head Rotation (Tilt) About the Bitemporal Axis

The reference line (in red), used to measure head tilt, was drawn from the most anterior point of the corpus callosum to the most posterior point. This line was found to be virtually parallel to the bicomissural line (i.e., the white line drawn between the anterior commissure and the posterior commissure).
FIGURE 6. Regions of the Parietal Lobe

a Image depicts a sagittal MRI slice approximately 5 mm lateral to the midsagittal slice, with the superior parietal gyrus traced in yellow and the postcentral gyrus in blue.
FIGURE 7. Three-Dimensional Surface Rendering of the Cortex

Gyri of the parietal lobe are color-coded as follows: postcentral gyrus (blue), superior parietal gyrus (green), supramarginal gyrus (red), and angular gyrus (yellow). See text and subsequent figures for detailed descriptions of the numerically designated boundaries and the alphabetically labeled planes.
FIGURE 8. Boundaries Defining Plane B (from figure 7) of the Cortex

The left side of the figure shows a midsagittal MRI slice with the reference line from figure 5 in red. The white line was drawn from the most anterior to the most posterior point of the corpus callosum, which formed a 9° angle with the reference line. The right side of the figure depicts the second line that defined the boundary of plane B, which was drawn on a lateral (sagittal) MRI slice with the same x, y coordinates as the red reference line.
FIGURE 9. Boundaries Defining Plane C (from figure 7) of the Cortex

The plane was defined by two parallel lines, the first of which is depicted in the midsagittal MRI slice on the left as a straight line drawn through the parieto-occipital fissure. The second line was drawn with identical x, y coordinates as the first line and is shown on the more lateral sagittal MRI slice on the right.
### TABLE 1
Parietal Lobe Region Relative Volumes in Male Patients With Schizophrenia and Healthy Male Comparison Subjects

<table>
<thead>
<tr>
<th>Region</th>
<th>Patients With Schizophrenia (N=15)</th>
<th>Comparison Subjects (N=15)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Relative Volume(^a)</td>
<td>SD</td>
</tr>
<tr>
<td>Total parietal lobe</td>
<td></td>
<td></td>
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<tr>
<td>Left</td>
<td>337.4</td>
<td>37.9</td>
</tr>
<tr>
<td>Right</td>
<td>332.8</td>
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<td>Left</td>
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<td>14.8</td>
</tr>
<tr>
<td>Right</td>
<td>121.9</td>
<td>10.3</td>
</tr>
<tr>
<td>Angular gyrus</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Left</td>
<td>50.6</td>
<td>10.3</td>
</tr>
<tr>
<td>Right(^b)</td>
<td>53.1</td>
<td>13.6</td>
</tr>
<tr>
<td>Supramarginal gyrus</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Left</td>
<td>63.6</td>
<td>9.3</td>
</tr>
<tr>
<td>Right</td>
<td>68.8</td>
<td>12.1</td>
</tr>
<tr>
<td>Superior parietal gyrus</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Left</td>
<td>132.7</td>
<td>21.9</td>
</tr>
<tr>
<td>Right</td>
<td>131.1</td>
<td>18.1</td>
</tr>
<tr>
<td>Postcentral gyrus</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Left</td>
<td>90.5</td>
<td>13.2</td>
</tr>
<tr>
<td>Right</td>
<td>79.8</td>
<td>13.6</td>
</tr>
</tbody>
</table>

\(^a\) Corrected for total intracranial volume to control for variations in head size (regional volume/total intracranial capacity × 100).

\(^b\) Significant difference between groups (t=−2.66, df=28, p<0.01).
## TABLE 2
Inferior Parietal Lobule Volume Differences Between Male Patients With Schizophrenia and Healthy Male Comparison Subjects

<table>
<thead>
<tr>
<th>Effect/Interaction</th>
<th>F $^a$</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group (schizophrenia versus comparison)</td>
<td>0.26</td>
<td>0.61</td>
</tr>
<tr>
<td>Laterality (left versus right)</td>
<td>0.00</td>
<td>0.98</td>
</tr>
<tr>
<td>Region (angular gyrus and supramarginal gyrus)</td>
<td>114.04</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Laterality by group</td>
<td>6.29</td>
<td>0.02</td>
</tr>
<tr>
<td>Region by group</td>
<td>4.25</td>
<td>0.05</td>
</tr>
<tr>
<td>Laterality by region</td>
<td>2.07</td>
<td>0.11</td>
</tr>
<tr>
<td>Laterality by region by group</td>
<td>0.64</td>
<td>0.43</td>
</tr>
</tbody>
</table>

$^a$ Repeated measures ANOVA (df=1, 28).
**TABLE 3**  
Parietal Region Asymmetry in Male Patients With Schizophrenia and Healthy Male Comparison Subjects

<table>
<thead>
<tr>
<th>Region</th>
<th>Asymmetry Coefficient</th>
<th>SD</th>
<th>Asymmetry Coefficient</th>
<th>SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patients With Schizophrenia (N=15)</td>
<td>Comparison Subjects (N=15)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total parietal lobe $^b$</td>
<td>0.60</td>
<td>2.4</td>
<td>2.94</td>
<td>3.5</td>
</tr>
<tr>
<td>Inferior parietal lobule $^c$</td>
<td>−2.98</td>
<td>6.7</td>
<td>3.15</td>
<td>7.5</td>
</tr>
<tr>
<td>Angular gyrus $^d$</td>
<td>−1.81</td>
<td>11.2</td>
<td>7.51</td>
<td>11.1</td>
</tr>
<tr>
<td>Supramarginal gyrus</td>
<td>−3.70</td>
<td>9.6</td>
<td>0.10</td>
<td>7.9</td>
</tr>
<tr>
<td>Superior parietal gyrus</td>
<td>0.45</td>
<td>4.0</td>
<td>0.73</td>
<td>5.2</td>
</tr>
<tr>
<td>Postcentral gyrus</td>
<td>6.34</td>
<td>8.3</td>
<td>8.01</td>
<td>9.0</td>
</tr>
</tbody>
</table>

$^a$ Computed as (left-right)/(left+right) × 100.

$^b$ Significant difference between groups (t=2.11, df=28, p=0.04).

$^c$ Significant difference between groups (t=2.14, df=28, p=0.03).

$^d$ Significant difference between groups (t=2.29, df=28, p<0.03).
TABLE 4
Volume Correlations Between the Left and Right Inferior Parietal Lobules and Frontal Lobe Structures in 15 Male Patients With Schizophrenia

<table>
<thead>
<tr>
<th>Brain Region</th>
<th>r</th>
<th>95% Confidence Interval</th>
</tr>
</thead>
<tbody>
<tr>
<td>Left inferior parietal lobule</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Left frontal lobe</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Middle frontal gyrus</td>
<td>0.56</td>
<td>0.25–0.78</td>
</tr>
<tr>
<td>Inferior frontal gyrus</td>
<td>0.80</td>
<td>0.62–0.90</td>
</tr>
<tr>
<td>Orbital gyrus</td>
<td>0.76</td>
<td>0.55–0.88</td>
</tr>
<tr>
<td>Right frontal lobe</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Orbital gyrus</td>
<td>0.84</td>
<td>0.70–0.92</td>
</tr>
<tr>
<td>Inferior frontal gyrus</td>
<td>0.66</td>
<td>0.39–0.93</td>
</tr>
<tr>
<td>Superior frontal gyrus</td>
<td>0.72</td>
<td>0.49–0.85</td>
</tr>
<tr>
<td>Right inferior parietal lobule</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Left frontal lobe</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Middle frontal gyrus</td>
<td>0.67</td>
<td>0.41–0.83</td>
</tr>
<tr>
<td>Inferior frontal gyrus</td>
<td>0.76</td>
<td>0.55–0.88</td>
</tr>
<tr>
<td>Orbital gyrus</td>
<td>0.84</td>
<td>0.70–0.92</td>
</tr>
<tr>
<td>Right frontal lobe</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Orbital gyrus</td>
<td>0.74</td>
<td>0.52–0.87</td>
</tr>
<tr>
<td>Inferior frontal gyrus</td>
<td>0.68</td>
<td>0.42–0.83</td>
</tr>
<tr>
<td>Superior frontal gyrus</td>
<td>0.80</td>
<td>0.62–0.90</td>
</tr>
</tbody>
</table>
### TABLE 5

Volume Correlations Between the Left and Right Inferior Parietal Lobules and Temporal Lobe Structures in 14 Male Patients With Schizophrenia

<table>
<thead>
<tr>
<th>Brain Region</th>
<th>r</th>
<th>95% Confidence Interval</th>
</tr>
</thead>
<tbody>
<tr>
<td>Left inferior parietal lobule</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Left temporal lobe</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Anterior superior temporal gyrus</td>
<td>0.70</td>
<td>0.45–0.84</td>
</tr>
<tr>
<td>Amygdala</td>
<td>0.66</td>
<td>0.39–0.80</td>
</tr>
<tr>
<td>Right inferior parietal lobule</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Left temporal lobe</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Anterior superior temporal gyrus</td>
<td>0.71</td>
<td>0.47–0.84</td>
</tr>
<tr>
<td>Amygdala</td>
<td>0.67</td>
<td>0.41–0.80</td>
</tr>
<tr>
<td>Right temporal lobe: hippocampus</td>
<td>0.59</td>
<td>0.29–0.78</td>
</tr>
</tbody>
</table>