

Delineating the Superior Temporal Gyrus using Dynamic Programming in Schizophrenia and Bipolar Disorder

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Introduction

The superior temporal gyrus (STG) has been implicated in neurological disorders that disrupt its normal function in, for example, schizophrenia (SCZ) and bipolar disorder (BPD) [1,2]. However, several neuroimaging and post-mortem studies have discovered conflicting differences in the morphometry of the STG in SCZ and BPD [3,4,5,6]. These differences arise from the consideration that the volume of the gray matter of the STG is considered an entity. The highly folded cortical structure of the STG as well as anatomical definitions of its boundaries may confound quantification of morphometric measures. Dynamic Programming (DP) allows for two critical components of cortical analysis: namely, a 3D representation of the grey-white matter surface and a cortical representation that shows the position, variation, and density of the neocortical cells which are currently viewed as gray matter on MRI [7]. DP has been used to analyze such morphometric properties as volume, surface area, and thickness of the planum temporale [8]. In this study, we use DP to delineate the structure of the STG and to test differences in the STG due to gender, laterality, and diagnosis.

Labeled Cortical Distance Maps (LCDM)

The flow chart below describes the pipeline for generating LCDMs [8,9]. Following Miller et al. [9],

Results



LCDMs are generated, which are histograms of labeled tissue compartments of GM, WM, and cerebrospinal fluid computed as a function of distance from the GM/WM isosurface.



Figure 1: Left: distance profile of cortical thickness [red = gray matter (GM), blue = white matter (WM), yellow = gray/white isosurface, green line = distance of laminar layer]. Middle: segmentation of gray and white matter. Right: corresponding LCDM.

Methods

MPRAGE scans of 1mm³ resolution from 60 subjects were included in our study, of which 20 were controls (10 males and 10 females, age: 36.5±11.2, mean±SD), 20 were SCZ patients (10 males and 10 females, age: 36.5 ± 7.82), and 20 were BPD patients (10 males and 10 females, age: 40 ± 9.43) were selected.

DP delineation of the STG was initiated with several landmarks. The posterior landmark of the STG boundary begins at the intersection of the angular gyrus (AG) and the STG at the most posterior extent of the lateral fissure (LF) [Figs. 2B, C; Figs. 3B, D]. The anterior landmark of the STG boundary is located at the superior portion of the temporal pole at the ascending ramus of the LF [Figs. 2B, C; Fig. 3A]. The inferior extent of the STG boundary follows from the posterior landmark along the superior temporal sulcus (STS) all the way to the anterior landmark [Fig. 2B; Fig. 3B, D]. The superior extent of the STG boundary follows from the anterior landmark along the LF to the posterior landmark [Fig. 2B; Fig. 3A, C]. Surface area (SA) was determined from the triangulated manifold while volume (V) and cortical thickness (T) were respectively derived from the total number of voxels and the 95th percentile distance from the LCDM histogram respectively. To account for the large size of the STG, thickness and volume data were retrieved at censored distances of 4 mm, 4.5 mm, 5 mm, 5.5 mm, 6 mm.



Surface Area (mm²)





Figure 2: MRI slices of STG in coronal (A) and sagittal (B, C) views.

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4.5 mm



4.5 mm





Thickness

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Gender	p < 0.05	p < 0.05	p = 0.06	p = 0.0862	p = 0.1486
Laterality	p < 0.005	p < 0.0001	p < 0.0005	p < 0.001	p < 0.0121
Diagnosis	p = 0.559	p = 0.558	p = 0.4958	p = 0.4472	p = 0.5234
Volume					
Gender	p < 0.0005	p < 0.001	p < 0.001	p < 0.005	p < 0.005
Laterality	p < 0.05	p < 0.05	p < 0.05	p < 0.05	p < 0.01
Diagnosis	p < 0.01	p < 0.01	p < 0.05	p < 0.05	p < 0.05

4 mm

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Surface Area				
Gender	p < 0.0001			
Laterality	p < 0.001			
Diagnosis	p < 0.05			

Table: p-values of gender, laterality, and diagnosis.

Discussion

-data was most sensitive at 4 and 4.5 mm (i.e. more voxels is less significant) -between subjects, laterality, gender, and diagnosis play a significant role in SA of STG; diagnosis and gender play a role in V of STG; and gender has a marginally significant difference in T -within subjects, laterality is significantly different for SA, V, and T -SCZ has larger V than normals, and BPD has smaller V than normals -SCZ has larger SA than normals, and BPD has smaller SA than normals -left STG is always larger than right STG; females have thicker STG than males

-LCDM and laminar approximation (LA) are closely similar as validated by surface overlays in Figure 3; future progress will compare both LCDM and LA and validate LCDM

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