## Investigating Cortical Variability Using a Generic Gyral Model

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## Introduction

Human cortical folding shows a high degree of inter-individual variability which makes inter subject comparisons very difficult.


We investigate inter-subject variability using a set of cortical landmarks. These landmarks are derived from a generic gyral model (GGM) that was introduced at MICCAI 2005. The GGM resulted from gyral maps of 96 subjects. The vertices of the GGM serve as landmarks that can be easily serve as landmarks that can be easily
identified across a large population identified across a large population
of subjects using non-linear registration techniques. Their variances and covariances can then be analyzed and used in a regionalized way.


Generic Gyral Model (GGM) - MICCAI 2005

## Obtaining landmarks

The generic gyral model (GGM) consists of an averaged line rep- We used these vertices as landmarks for inter-subject registration. resentation for some major gyri at eight depth levels. Each depth These landmarks can be identified in each individual via nonlevel contains about 85 vertices so that the entire model has about linear registration. An initial rough match is obtained via a cubic 700 vertices. polynomial, a refined match via thinplate splines

single subject gyral line


GGM-landmarks to be matched

## Correcting for $\mathrm{AC} / \mathrm{PC}$ bias

Landmarks that are closer to AC/PC will tend to be less variable than those that are further away. To correct for this problem, we divided the displacement magnitude of each vertex by its distance from AC/PC.

## Results

1. Variability maps


A displacement map of one individual data set.
The displacement is corrected for the bias introduced by the AC/PC-alignment by dividing the displacement magnitude by the distance from AC/PC. The color-coded values correspond to corrected displacement magnitudes.


Variability map computed as an average of displacement maps of 96 subjects.
2. Variability decreases with depth

|  | perisylvian | precentra | postcentra 1 | parietal | temp oral | middle front al |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: |
| deep | 6.890 | 2.669 | 1.811 | 2.435 | 6.054 | 5.399 |
| shallow | 8.892 | 3.837 | 1.995 | 2.128 | 6.085 | 5.578 |

The decrease in variance with depth.

The table shows the decrease in variance with depth. The estimates correspond to standard deviations averaged across the landmarks within a gyrus and summed over $\mathrm{x}, \mathrm{y}, \mathrm{z}$. Note that in all but one case, the standard deviations are less at the deeper level.

## Discussion

The novelty of our approach lies the usage of the generic gyral model which provides a set of well-defined landmarks that can be identified across may subjects using non-linear registration. These landmarks are particularly useful because they are located on main gyri that can be easily detected in most healthy subjects. Also, these landmarks are separated into different strata of depth so that variability can be investigated as a function of depth.

## Reference:

C. Loolmann,.X. von Cramon, A.C.F. Colcheseser: middle frontal gyrus (c). The color coding is based on the AC/PC-corrected variability measure described above.

