

Investigating the relationship of myeloarchitecture and connectivity in the human cortex using MRI



Julia M Huntenburg¹, Pierre-Louis Bazin¹, Alexandros Goulas², Daniel S Margulies¹

¹ Max Planck Institute for Human Cognitive and Brain Sciences Leipzig, Germany

² University Medical Center Hamburg-Eppendorf, Germany
huntenburg@cbs.mpg.de

MAX PLANCK INSTITUTE FOR HUMAN COGNITIVE AND BRAIN SCIENCES
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Introduction

A link between microarchitecture and connectivity has been demonstrated in macaque cortex [1].

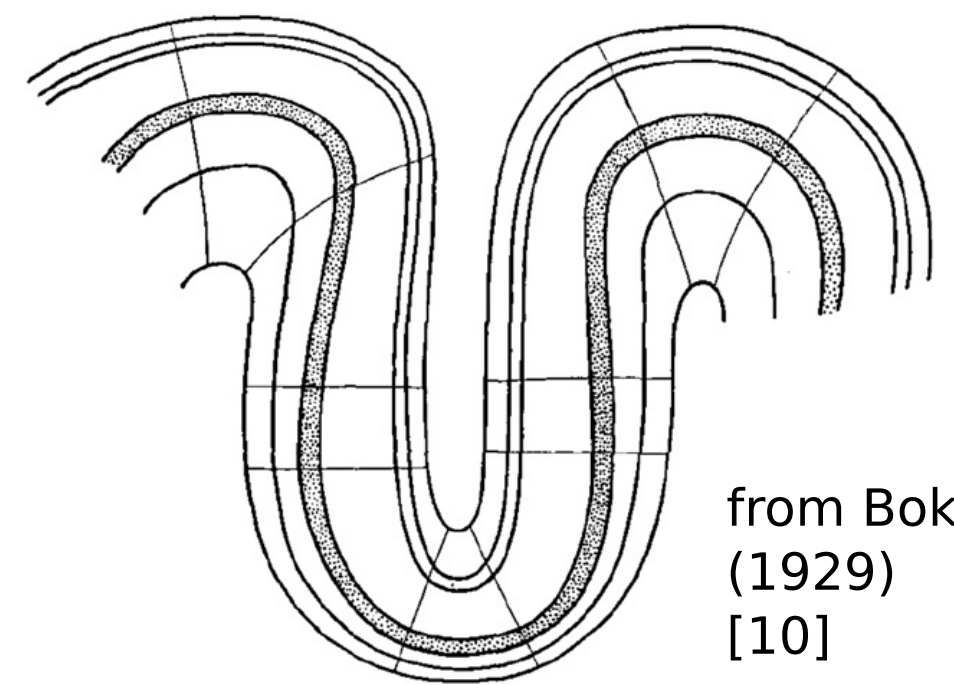
We investigate this link in human cortex using high-resolution MRI:

- Resting state functional connectivity (FC)
- T1 maps as *in vivo* proxy for myeloarchitecture [2].

MRI datasets of 8 subjects acquired at 7T

- 4 resting state scans [3] (300 volumes, 70 slices, voxel=1.5 mm³, TR=3s, TE=17ms, FA=70°, GRAPPA=3)
- Quantitative T1 map [4] (MP2RAGE, voxel=0.5mm³, T1/2=900/2750ms, TR=5s, TE=2.45ms, FA1/2=5/3°)

Data



Preprocessing [5-7], sampling and minimal smoothing (FWHM=1.5/3 mm) of T1 and FC on study-specific surface template [8].

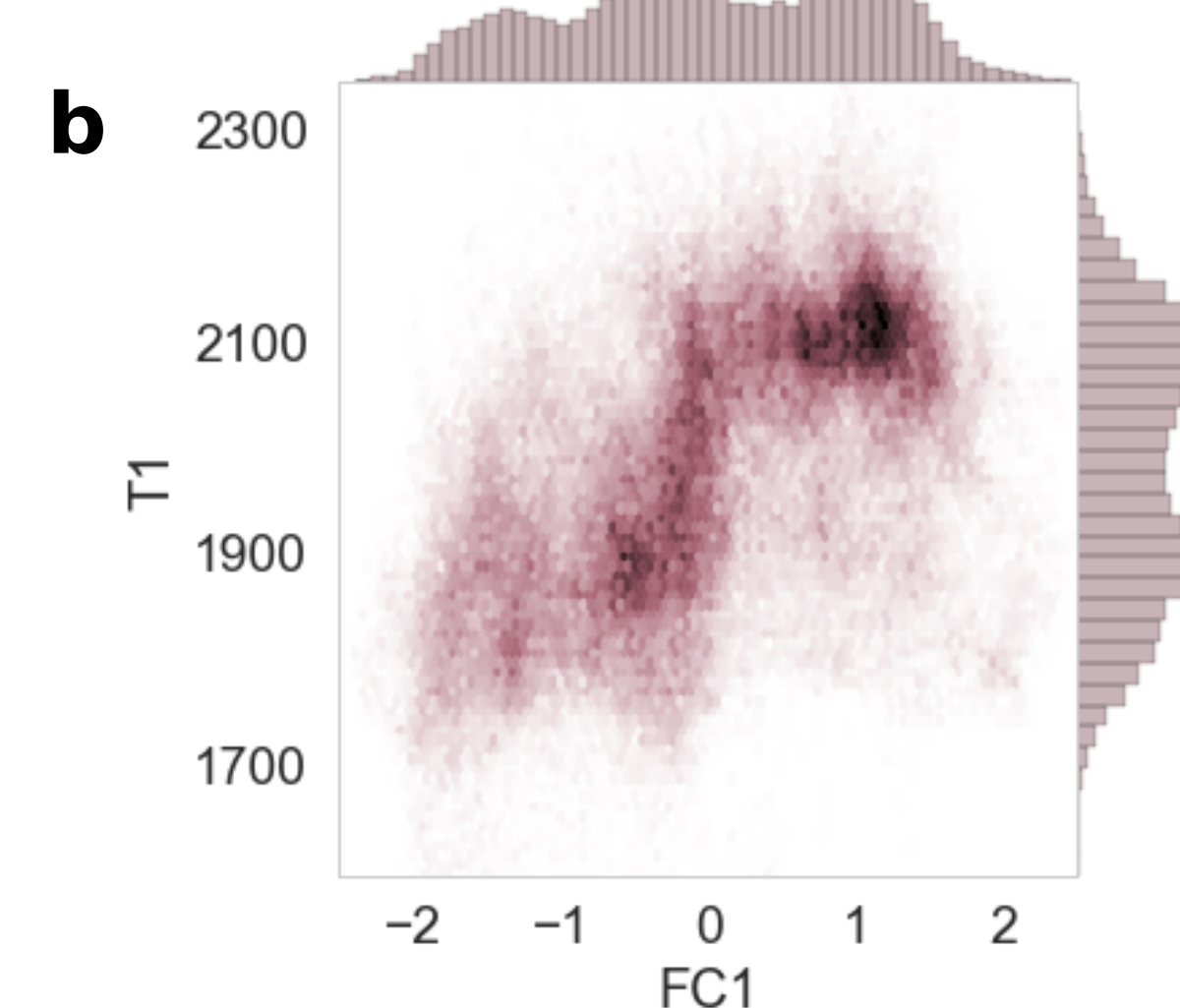
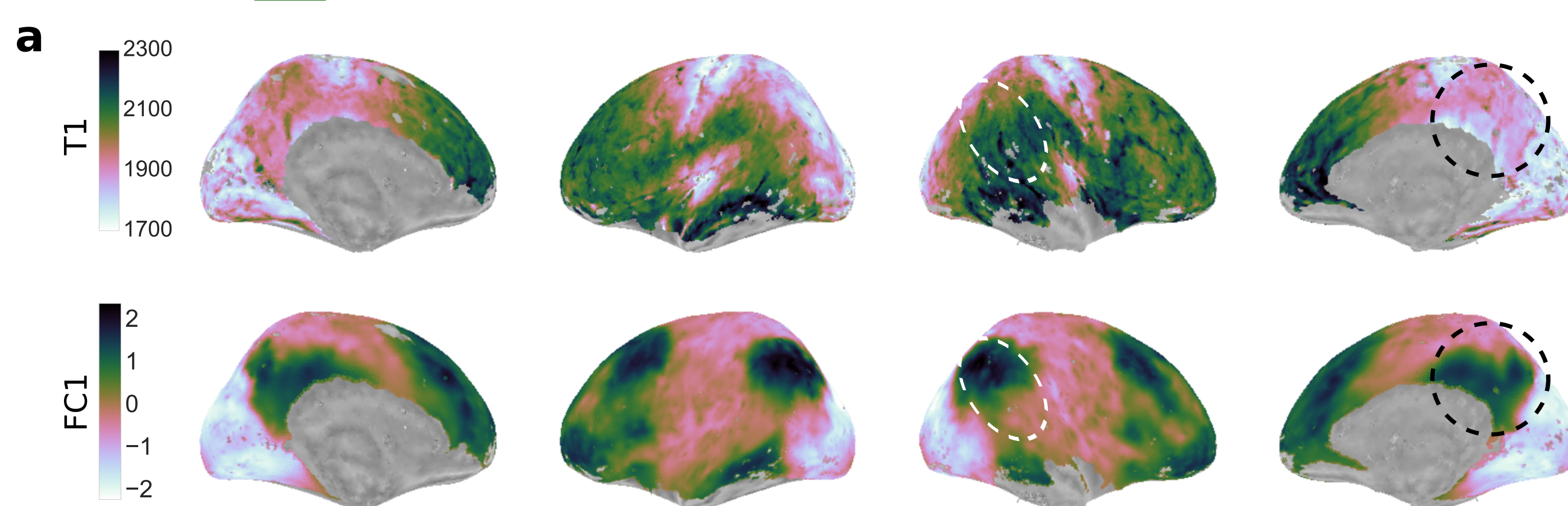
T1 sampled on 11 volume-preserving intracortical surfaces [9,10]. Only 5 central values are used to reduce partial volume effects.

Nonlinear dimensionality reduction of group average FC matrix using diffusion embedding [11].

Methods

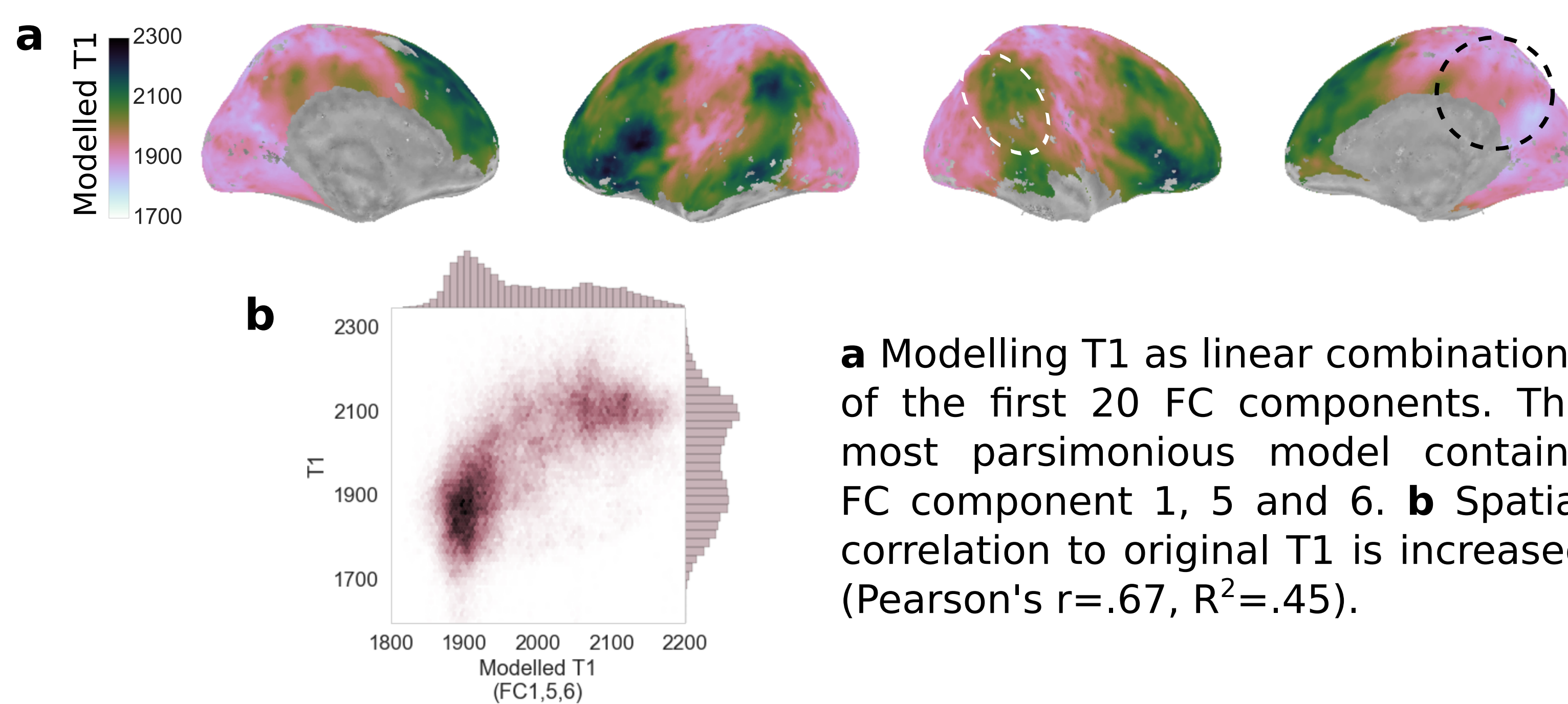
Results

1 Intracortical T1 and FC are systematically related.



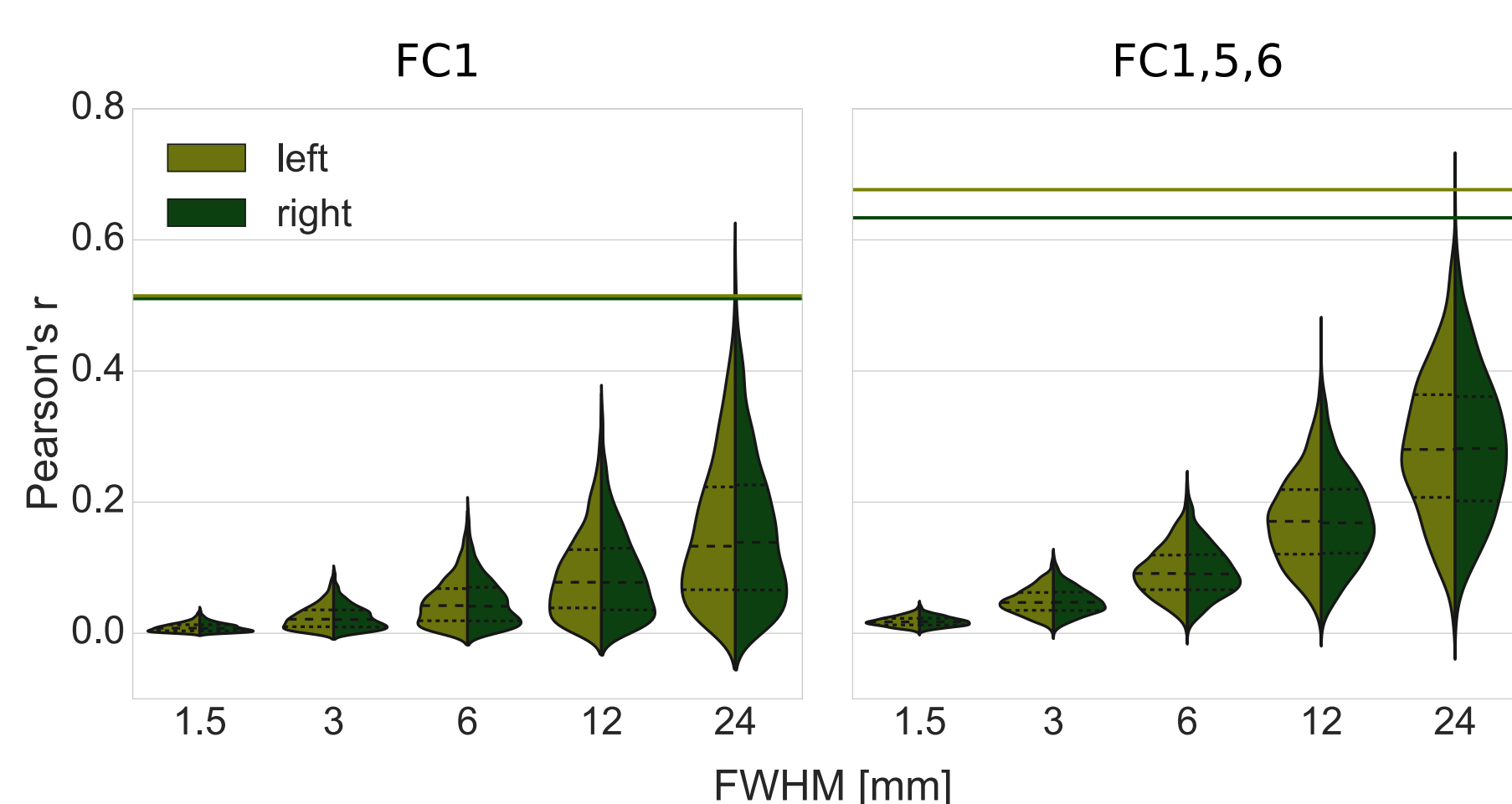
a The T1 map reflects myelin content, with lower T1 indicating higher myelin. The first component of the FC decomposition (FC1) represents the main variance in the FC data (12%). **b** T1 and FC1 are spatially correlated (Pearson's $r=.53$, $R^2=.28$).

2 Modelling T1 with multiple FC components improves fit.



a Modelling T1 as linear combinations of the first 20 FC components. The most parsimonious model contains FC component 1, 5 and 6. **b** Spatial correlation to original T1 is increased (Pearson's $r=.67$, $R^2=.45$).

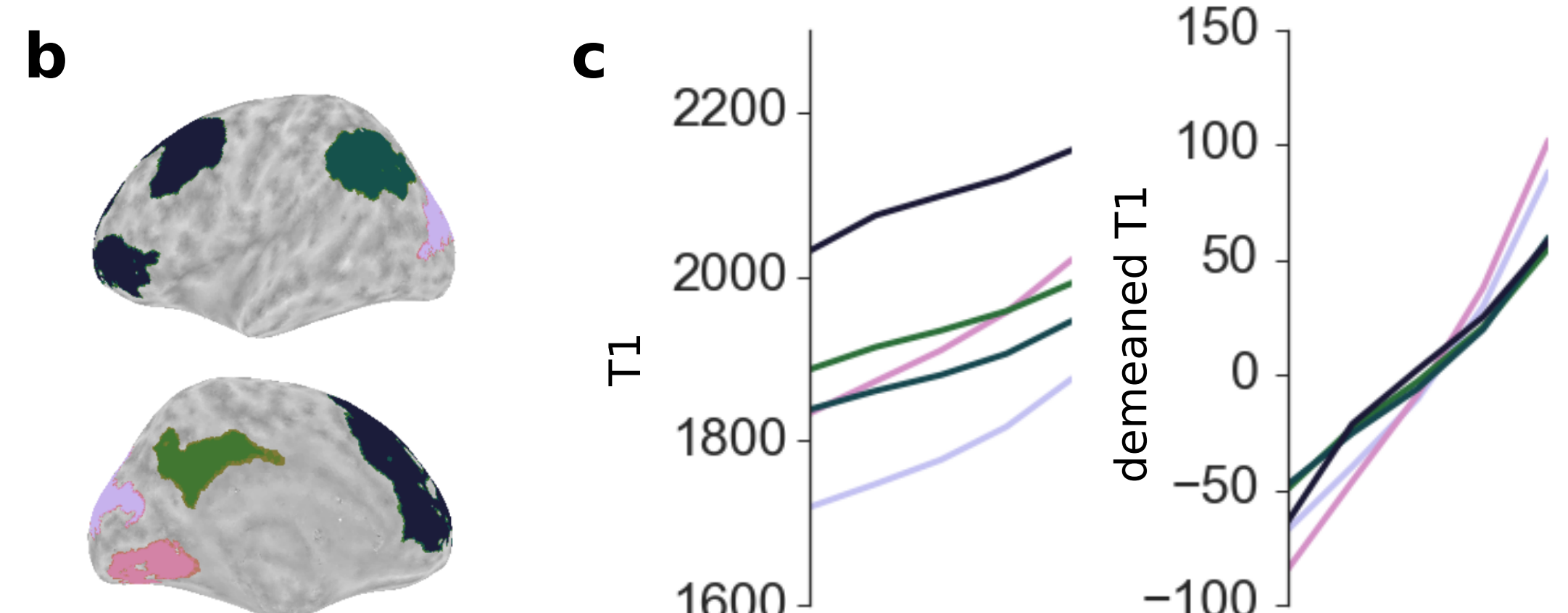
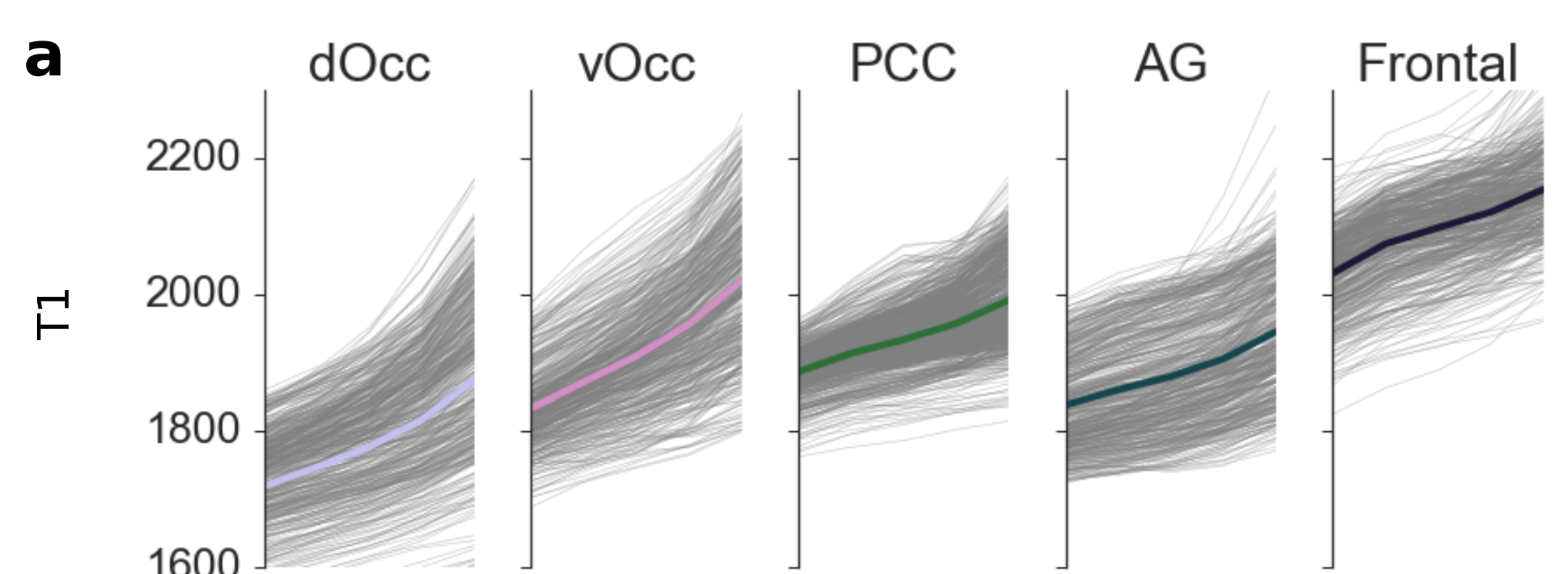
3 Correlation is not explained by data smoothness alone.



1000 random datasets per hemisphere, smoothed with different kernel sizes. When modelling the random maps with FC1 (left) or FC1,5,6 (right) correlation values are much lower than for the real data (horizontal lines).

4 T1 profile shapes link diverging regions.

a T1 values at 5 levels of cortical depth for each vertex in high (PCC/precuneus, AG, frontal) and low (dorsal and ventral occipital) FC1 peak clusters. **b** Cluster locations in the left hemisphere. **c** Average profiles (left) and demeaned average profiles (right) of the five clusters. Shape, but not mean, discriminates clusters from either side of the spectrum.



Discussion

We demonstrate a systematic relationship between *in vivo* measures of intracortical myelin and functional connectivity in human cortex.

Similar myeloarchitecture may reflect simultaneous evolution of distributed areas upon common environmental demands, resulting in interconnected functional entities [1].

Superimposed connectivity gradients and laminar myelin distribution provide additional information but also pose methodological challenges.

Integrating these complex data in an interpretable way will be a crucial next step.

References

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