

# Parcellation-Independent Framework for Analysing Developing Brain Networks Using Reparametrisation

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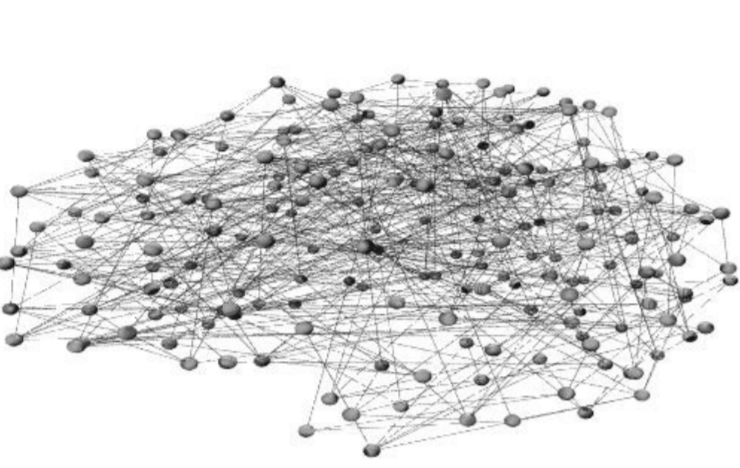
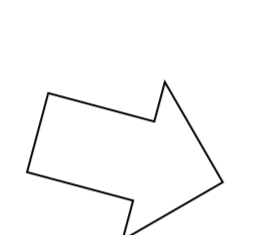
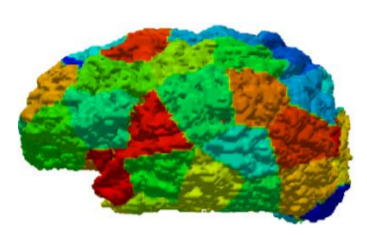
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## Abstract

Brain connectivity may be studied with diffusion MR (dMR), tractography and network theory. However, the lack of a standard for parcellating the neonatal brain leads to the use of atlas- and random-based methods, and thus to the unresolved challenge of comparing graphs with varying numbers and an unknown correspondence of nodes. We propose a parcellation-independent multi-scale analysis of network measures and show its potential in describing developmental changes in neonatal serial dMRI data.

## Network theory in brains

Divide brain into set of regions



Network theory is becoming more prevalent in neuroscience, as it:

- Allows to analyse complex systems
- Finds connections in data
- Defines properties of data points

Find connectivity, e.g. diffusion MRI

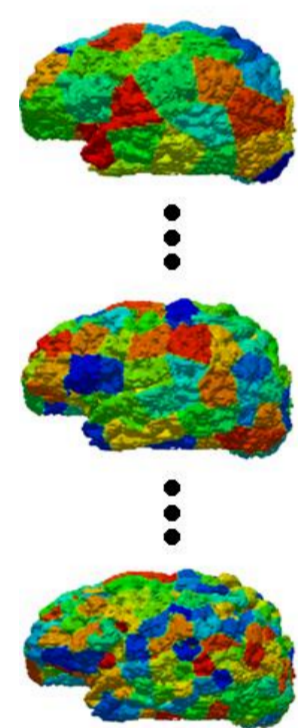
## Preterm and neonatal subjects



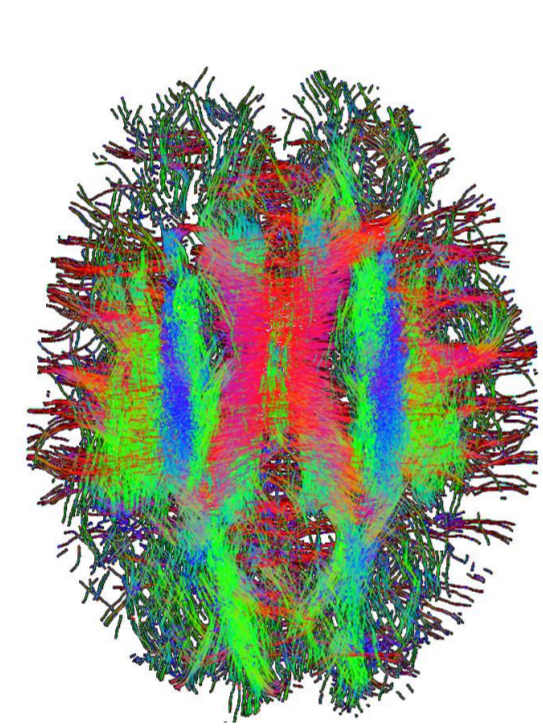
- High prematurity rate (worldwide ~ 10%)
  - Prematurity linked to adverse developmental outcome (~50%)
  - Early intervention and targeted support is desirable
- However, no standard set of regions exist for neonates.
- Stochastic parcellation approaches
  - Rely on fewer assumptions
  - Need to be repeated multiple times for each subject

## The challenge of comparing networks...

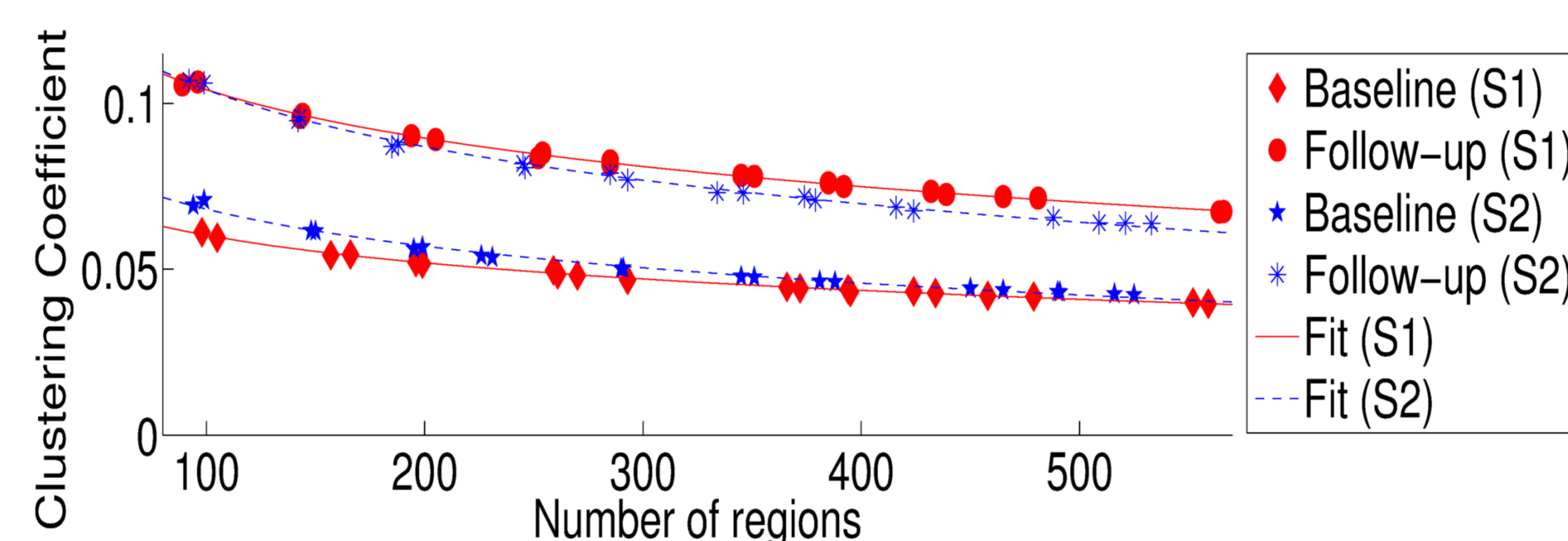
Different sets of regions of the same brain



Same structural connectivity



Varying results depending on number of nodes



- Network measures are highly dependent on set of regions/parcellation method.

Parcellation scheme	Number of nodes
AAL	70-90
Voxel-based	10 <sup>3</sup> -10 <sup>5</sup>
Stochastic	Specified number ± variation

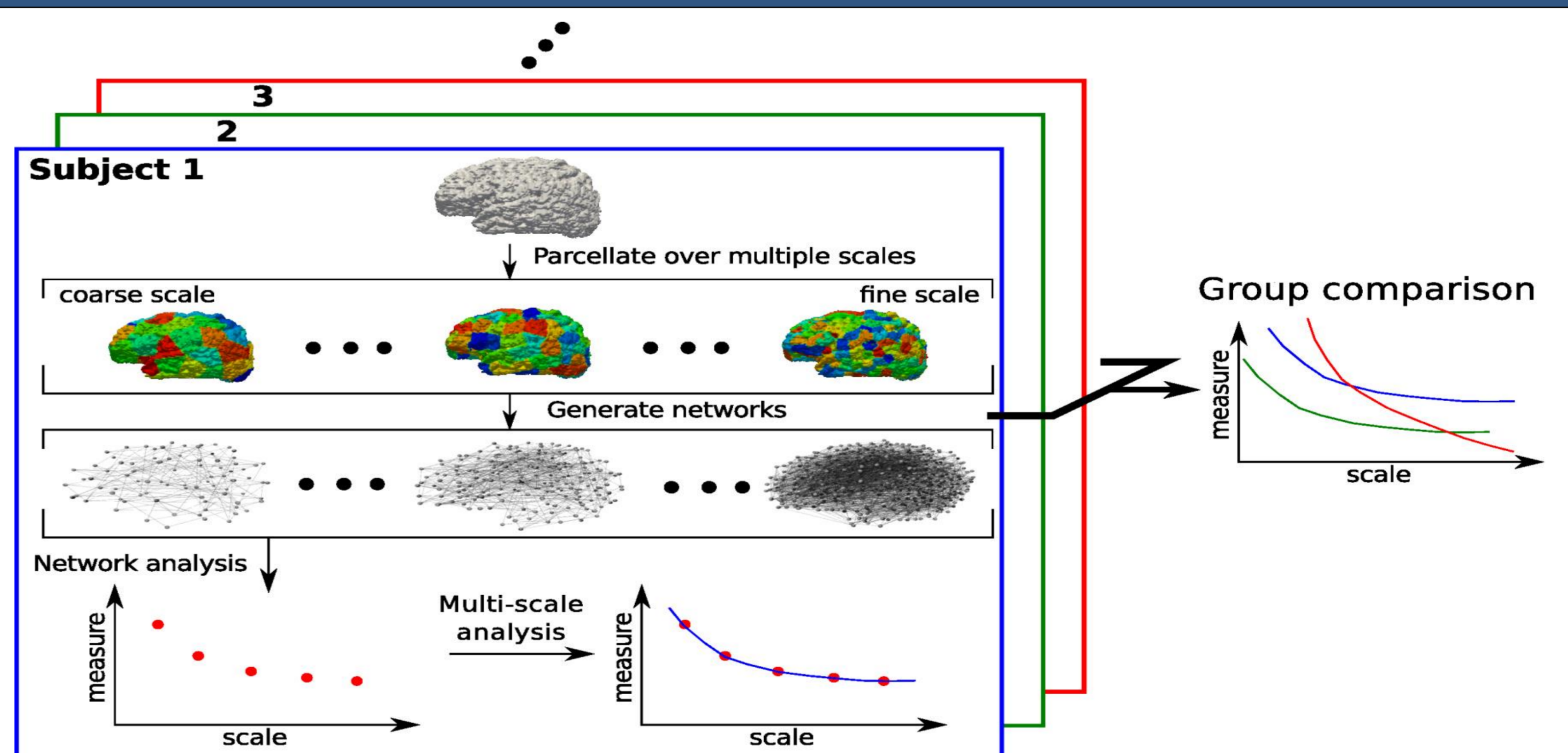
- As a result, comparing across subjects and across studies is difficult.

## Parcellation-independent Framework

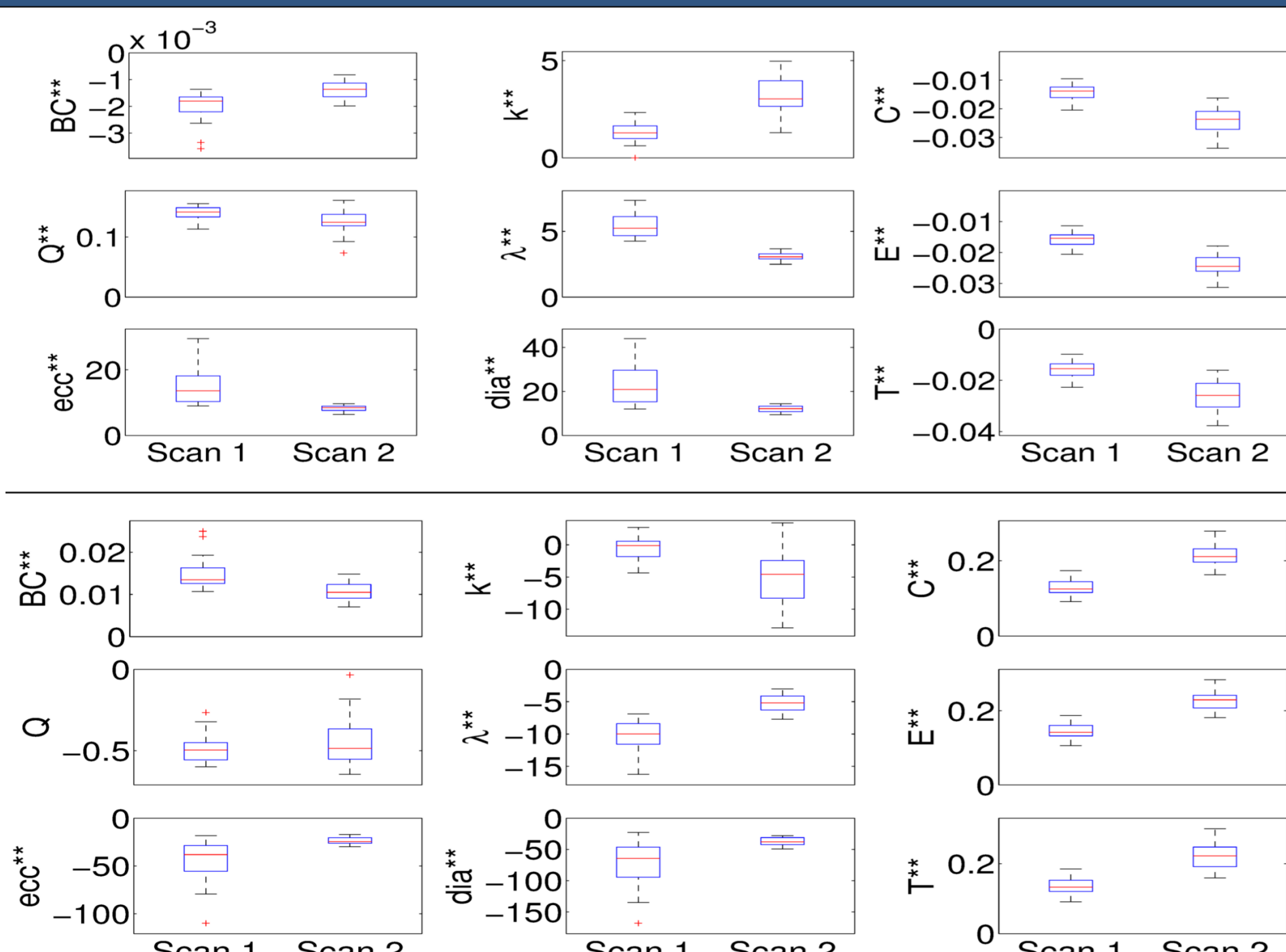
**Figure:** A framework for parcellation-independent multi-scale analysis. Each brain image is parcellated at multiple scales from a coarse (larger regions) to a fine scale. Subsequent estimates of structural networks are based on the subjects' dMRI data. For each network, the fitted models for global network measures over multiple scales, given by

$$m(G) = a * \log(N) + b,$$

where  $m(G)$  is a network measure taken on graph  $G$ ,  $N$  is the number of nodes and  $a$  and  $b$  are the model parameters which are used for group comparison.



## Results



**Figure:** Box-plots for the model parameters  $a$  (left; top) and  $b$  (left; bottom) for each measure at both time points. Values represent the group at each scan, where the subjects of scan one and two were  $30.8 \pm 1.0$  and  $41.2 \pm 1.2$  weeks old, correspondingly.

One and two stars next to the measure name represents P -values of  $P < 0.01$  and  $P < 0.001$ , correspondingly.

## Conclusion

- + Results independent of specific parcellation
- + Multi-scale analysis, circumventing number of nodes bias
- Fitting function is parameter of framework
- Direct interpretation of results is difficult