DCM workshop

Part 1

Timerseries extraction First-level DCM model

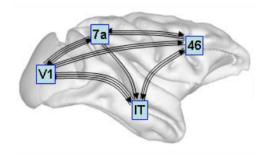
David Willinger

Workshop

Today (Nov 10)

- Some backgound on effective connectivity analysis
- How to identify functional networks
- How to extract time series information
- How to build a first-level DCM
- Review model fit

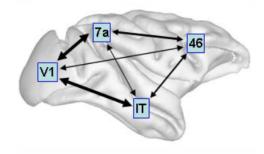
structural connectivity



- presence of physical connections
- Diffusion weighted imaging (DWI), tractography, tracer studies

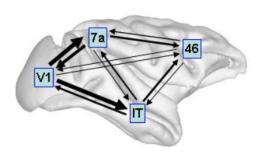
Sporns, 2007, Scholarpedia

functional connectivity



- statistical dependencies between regional time series
- correlations, Independent Component Analysis (ICA)

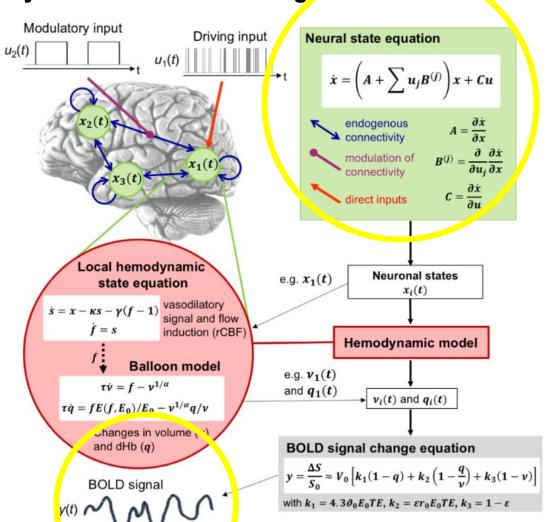
effective connectivity



- causal (directed) influences between neuronal populations
- Dynamic causal modeling (DCM)

DCM analysis looks at this

Dynamic causal modelling



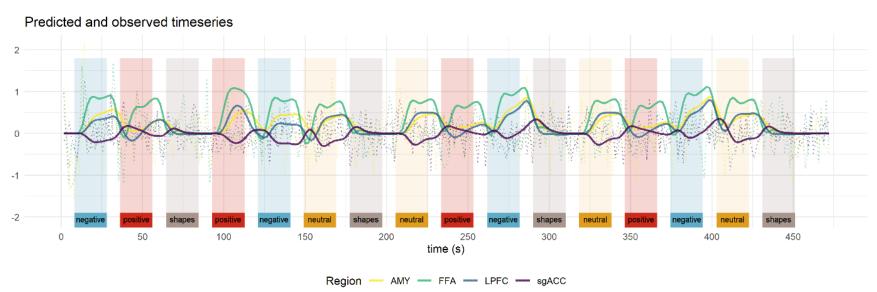
"How does change of activity in region A affect region B?"

Parameter matrices A, B and C

GLM analysis usually looks at this

Background Dynamic causal modelling

Example (face matching):



Dynamic causal modelling

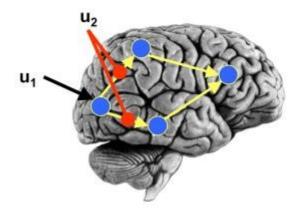
$$\dot{x} = \left(A + \sum u_j B^{(j)}\right) x + Cu$$

- A ... "intrinsic / average connectivtiy"
- → Represents the average connectivity during the task ("intercept")
- B ... "modulatory parameters"
- → B matrix can model additive connectivity changes for a **condition**
- C ... driving input
- → Generally, we will use an "all events" regressor as input
- D ... non-linear connectivity changes of one region between others
- → extension of B-matrix to explicitly model "source region"

This is typically supplied via an SPM.mat (GLM model)

Dynamic causal modelling

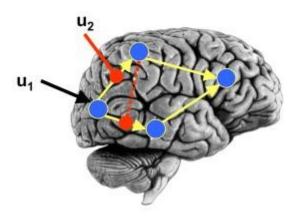
bilinear DCM



Bilinear state equation

$$\frac{dx}{dt} = \left(A + \sum_{i=1}^{m} u_i B^{(i)}\right) x + Cu$$

nonlinear DCM

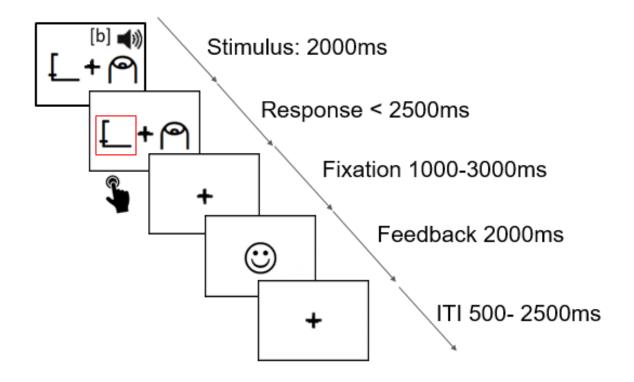


Nonlinear state equation

$$\frac{dx}{dt} = \left(A + \sum_{i=1}^{m} u_i B^{(i)}\right) x + Cu \qquad \frac{dx}{dt} = \left(A + \sum_{i=1}^{m} u_i B^{(i)} + \sum_{j=1}^{n} x_j D^{(j)}\right) x + Cu$$

Stephan et al., 2008

DCM Example: Task



AllRead

- Associative learning task
- Analysis contains two runs
- Reinforcement learning model captures learned association strength (values 0-1)

Sample

from the 99 kids:

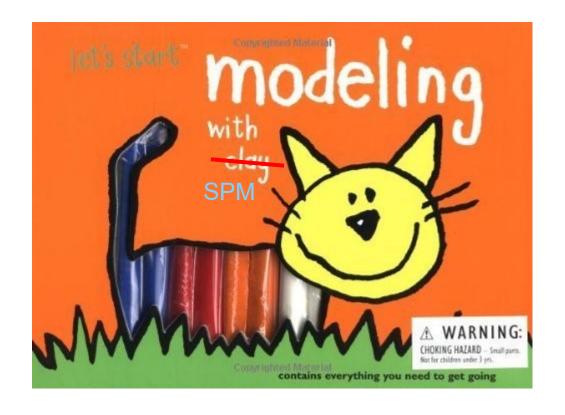
Movement 4 no good block, 10 only 1 good block => -14, To good performers (0 error): -5
 →80

Groups

- 27 poor (<16 SLRT)
- 42 typical (>25 SLRT)
- 11 gap (in between)

Demographics

- aged 8.9 ± 0.74 years (group difference)
- 44 male, 36 female
- SLRT whole group: mean 40.35 ± 42.12
- 27 poor SLRT mean: 5.01 ± 4.25, 42 typical SLRT mean: 58.57 ± 21.53
- 8 left handed
- nonverbal IQ: 105.1 ± 7.01 (no significant groups difference)



Prerequisites 1: Identify your hypothesis

- DCM is not good in exploratory analysis
- We need to formulate hypotheses and networks we are interested in a priori
- Typically:
 - How is connectivity modulated by condition A vs B
 - How do groups differ in connectivity
 - How does connectivity vary with a covariate

Prerequisites 2: Identify your network

- Goal: reveal the brain regions that are related to the task, such that we can extract the timecourse
- If a GLM analysis does not show activity in a region for any contrast, there is **no motivation** to include it in a DCM analysis → nothing to explain by our model
- Our example:
 - Is connectivity between regions supporting audiovisual integration affected by learning?
 - Do poor readers and typical readers differ?

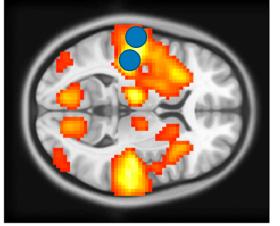
DCM Example: Identify your network

Main effect of audiovisual stimulus

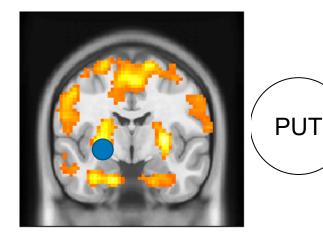


AUC

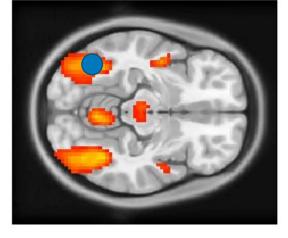


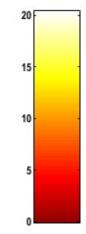


Effect of associative strength (AS)









den Ouden et al. (2010); Li et al. (2017, 2019)

Step 1: Timeseries extraction

- Goal: Extract the timeseries from all regions for all subjects
- This multivariate timeseries will be data to be modeled!
- Important:
 - Contains only signal from voxels with at least some degree of experimental effect (puncorr. < 0.05)
 - Can be constrained functionally and/or anatomically
 - Size of regions depends on structures investigated and smoothing kernel! Rule of thumb (radius ~ smoothing kernel ~ 6-8mm)
- After extraction check for subjects that do not show activity in single regions → will have to be excluded

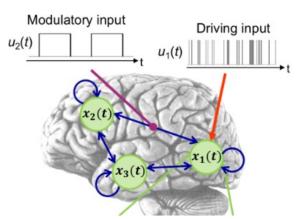
Step 2: Create GLM for DCM

- Goal: Create a model for DCM analysis (separate SPM.mat)
- Includes all regressors that enter the DCM
- Usually simpler than GLMs of activity analysis
- If we have multiple sessions, we need to concatenate them

Step 3: Create DCMs

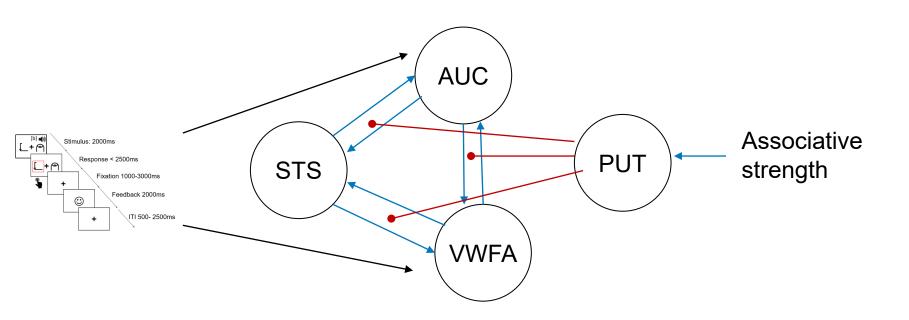
- Goal: Create a model for DCM analysis based on previously created SPM.mat (results in a DCM.mat)
- Stores information about model inputs, priors of connectivity and haemondynamic parameters, timing, onsets, ...
- Connectivity priors → do I have prior knowledge of connections?
- Task modulations
- Model input = where does TASK enter? ("ping the network")

$$\dot{x} = \left(A + \sum u_j B^{(j)}\right) x + Cu$$



DCM example: start with a fully connected graph

 Unless you have a very good reason to assume no connections between two regions



DCM example: Assessment of first-level results

– spm_dcm_fmri_check

DCM example: Assessment of first-level results

spm_dcm_review