DCM Advanced, Part I: Model Selection

Will Penny

SPM short course for fMRI, London 2014

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Model Compariso
Model Evidence
Complexity
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Families

FFX Mode Inference

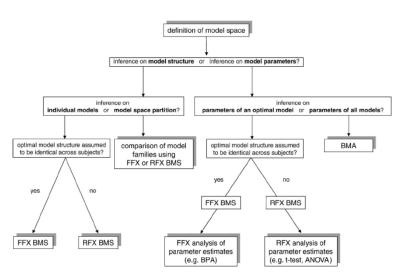
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Ten Simple Rules



Stephan et al. Neuroimage, 2010

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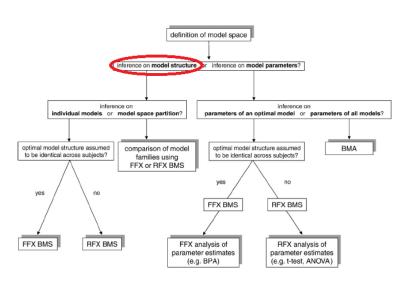
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The model evidence is given by integrating out the dependence on model parameters

$$p(y|m) = \int p(y,\theta|m)d\theta$$

= $\int p(y|\theta,m)p(\theta|m)d\theta$

Because we have marginalised over θ the evidence is also known as the marginal likelihood.

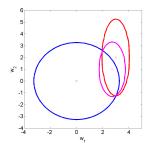
For linear Gaussian models there is an analytic expression for the model evidence.

$$y = Xw + e$$

where X is a design matrix and w are now regression coefficients. For prior mean μ_w , prior covariance C_w , observation noise covariance C_y the posterior distribution is given by

$$S_w^{-1} = X^T C_y^{-1} X + C_w^{-1}$$

$$m_w = S_w \left(X^T C_y^{-1} y + C_w^{-1} \mu_w \right)$$



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The log model evidence comprises sum squared precision weighted prediction errors and Occam factors

$$\log p(y|m) = -\frac{1}{2}e_y^T C_y^{-1} e_y - \frac{1}{2}\log|C_y| - \frac{N_y}{2}\log 2\pi$$
$$- \frac{1}{2}e_w^T C_w^{-1} e_w - \frac{1}{2}\log\frac{|C_w|}{|S_w|}$$

where prediction errors are the difference between what is expected and what is observed

$$e_y = y - Xm_w$$

 $e_w = m_w - \mu_w$

Bishop, Pattern Recognition and Machine Learning, 2006

$$\log p(y|m) = Accuracy(m) - Complexity(m)$$

where

$$Accuracy(m) = -\frac{1}{2}e_y^T C_y^{-1} e_y - \frac{1}{2}\log|C_y| - \frac{N_y}{2}\log 2\pi$$

and

Complexity(m) =
$$\frac{1}{2}e_w^T C_w^{-1} e_w + \frac{1}{2}\log\frac{|C_w|}{|S_w|}$$

 $\approx KL(prior||posterior)$

The Kullback-Leibler divergence measures the distance between probability distributions.

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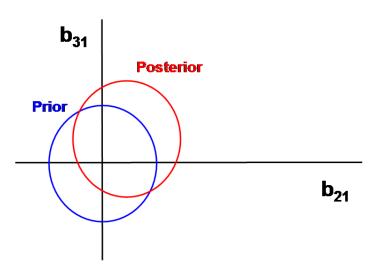
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Small KL



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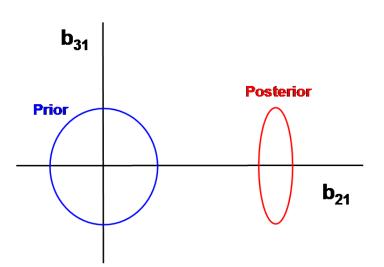
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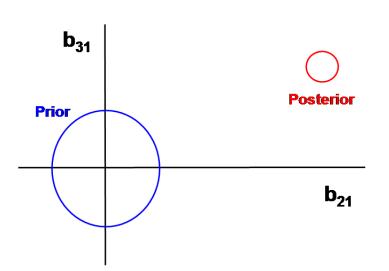
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For nonlinear models, we replace the true posterior with the approximate posterior (m_w, S_w) , and the previous expression becomes an approximation to the log model evidence called the (negative) Free Energy

$$F = -\frac{1}{2}e_{y}^{T}C_{y}^{-1}e_{y} - \frac{1}{2}\log|C_{y}| - \frac{N_{y}}{2}\log 2\pi$$
$$- \frac{1}{2}e_{w}^{T}C_{w}^{-1}e_{w} - \frac{1}{2}\log\frac{|C_{w}|}{|S_{w}|}$$

where

$$e_y = y - g(m_w)$$

 $e_w = m_w - \mu_w$

and $g(m_w)$ is the DCM prediction. This is used to approximate the model evidence for DCMs. *Penny, Neuroimage, 2011*.

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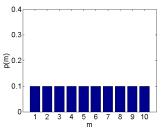
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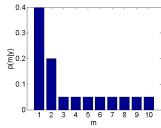
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Bayes rule for models

A prior distribution over model space p(m) (or 'hypothesis space') can be updated to a posterior distribution after observing data y.





This is implemented using Bayes rule

$$p(m|y) = \frac{p(y|m)p(m)}{p(y)}$$

where p(y|m) is referred to as the evidence for model m and the denominator is given by

$$p(y) = \sum_{m'} p(y|m')p(m')$$

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The Bayes factor for model *j* versus *i* is the ratio of model evidences

$$B_{ji} = \frac{p(y|m=j)}{p(y|m=i)}$$

We have

$$B_{ij} = \frac{1}{B_{ii}}$$

Hence

$$logB_{ji} = log p(y|m=j) - log p(y|m=i)$$
$$= F_j - F_i$$

Posterior Model Probability

Given equal priors, p(m = i) = p(m = j) the posterior model probability is

$$p(m = i|y) = \frac{p(y|m = i)}{p(y|m = i) + p(y|m = j)}$$

$$= \frac{1}{1 + \frac{p(y|m = j)}{p(y|m = i)}}$$

$$= \frac{1}{1 + B_{ji}}$$

$$= \frac{1}{1 + \exp(\log B_{ji})}$$

$$= \frac{1}{1 + \exp(-\log B_{ij})}$$

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Posterior Model Probability

Hence

$$p(m=i|y)=\sigma(\log B_{ij})$$

where is the Bayes factor for model i versus model j and

$$\sigma(x) = \frac{1}{1 + \exp(-x)}$$

is the sigmoid function.

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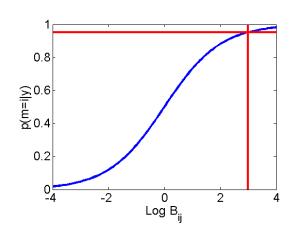
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The posterior model probability is a sigmoidal function of

the log Bayes factor

$$p(m=i|y) = \sigma(\log B_{ij})$$



Bayes factors

The posterior model probability is a sigmoidal function of the log Bayes factor

$$p(m=i|y)=\sigma(\log B_{ij})$$

Table 1 Interpretation of Bayes factors

B_{ij}	$p(m=i y) \ (\%)$	Evidence in favor of model
1-3	50-75	Weak
3-20	75-95	Positive
20-150	95-99	Strong
≥150	≥99	Very strong

Bayes factors can be interpreted as follows. Given candidate hypotheses i and j, a Bayes factor of 20 corresponds to a belief of 95% in the statement 'hypothesis i is true'. This corresponds to strong evidence in favor of i.

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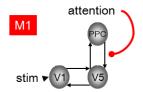
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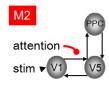
Kass and Raftery, JASA, 1995.

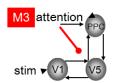
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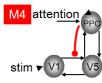
Example

Modelling visual attention responses with DCM for fMRI. Modulatory connections shown in red.









How does attention affect visual processing network?

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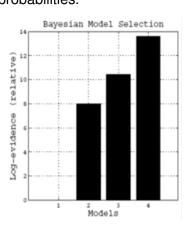
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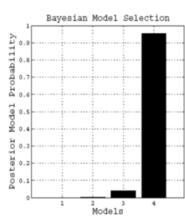
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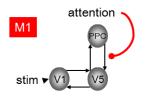
Relative Model evidences and equivalent posterior probabilities.

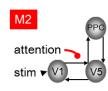


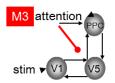


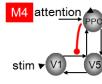
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How does attention affect visual processing network?









Model 2 is better than model 1 by (Bayes) factor 2966 Model 3 is better than model 2 by (Bayes) factor 12 Model 4 is better than model 3 by (Bayes) factor 23 DCM Advanced, Part I: Model Selection

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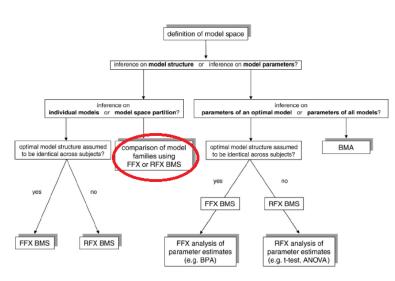
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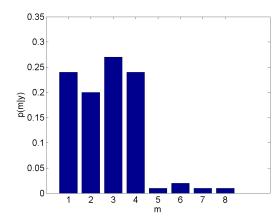
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Posterior Model Probabilities

Say we've fitted 8 DCMs and get the following distribution over models



Similar models share probability mass (dilution). The probability for any single model can become very small esp. for large model spaces.

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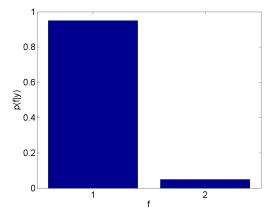
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Model Families

Assign model m to family f eg. first four to family one, second four to family two. The posterior family probability is then

$$p(f|y) = \sum_{m \in S_f} p(m|y)$$



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$$p(f)=\frac{1}{K}$$

The prior family probability is related to the prior model probability

$$p(f) = \sum_{m \in S_f} p(m)$$

where the sum is over all N_f models in family f. So we set

$$p(m) = \frac{1}{KN_f}$$

for all models in family f before computing p(m|y). This allows us to have families with unequal numbers of models. *Penny et al. PLOS-CB*, 2010.

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Different Sized Families

So say we have two families. We want a prior for each family of p(f) = 0.5.

If family one has $N_1=2$ models and family two has $N_2=8$ models, then we set

$$p(m) = \frac{1}{2} \times \frac{1}{2} = 0.25$$

for all models in family one and

$$p(m) = \frac{1}{2} \times \frac{1}{8} = 0.0625$$

for all models in family two.

These are then used in Bayes rule for models

$$p(m|y) = \frac{p(y|m)p(m)}{p(y)}$$

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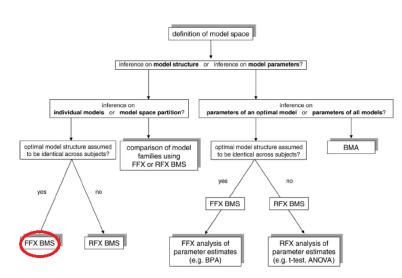
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Fixed Effects BMS



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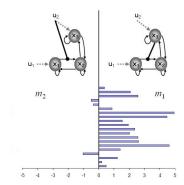
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Fixed Effects BMS

Two models, twenty subjects.

$$\log p(Y|m) = \sum_{n=1}^{N} \log p(y_n|m)$$



The Group Bayes Factor (GBF) is

$$B_{ij}=\prod_{n=1}^N B_{ij}(n)$$

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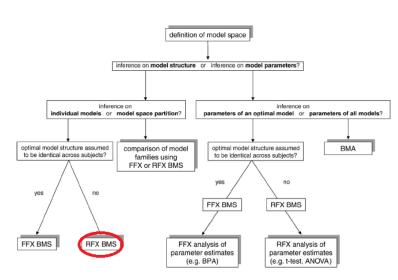
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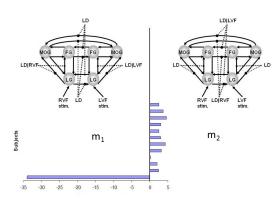
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Stephan et al. J. Neurosci, 2007



11/12=92% subjects favour model 2.

GBF = 15 in favour of model 1. FFX inference does not agree with the majority of subjects.

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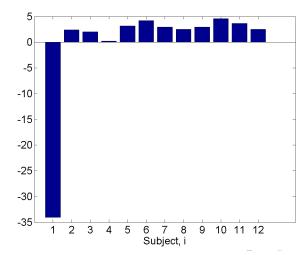
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Log Bayes Factor in favour of model 2

$$\log \frac{p(y_i|m_i=2)}{p(y_i|m_i=1)}$$



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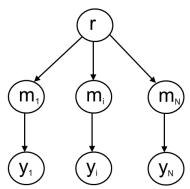
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Model frequencies r_k , model assignments m_i , subject data y_i .



Approximate posterior

$$q(r,m|Y) = q(r|Y)q(m|Y)$$

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Stephan et al, Neuroimage, 2009.

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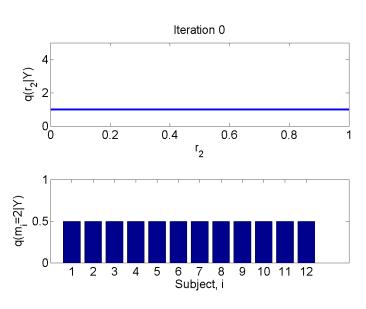
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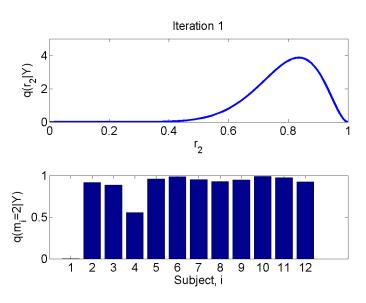
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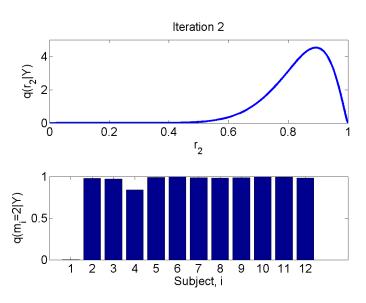
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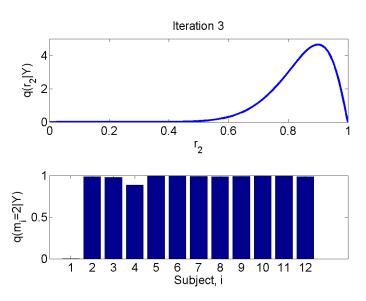
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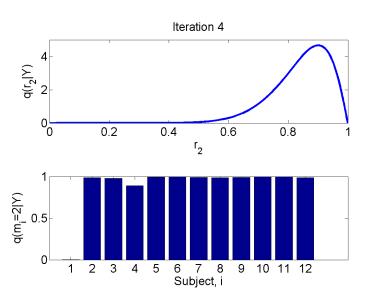
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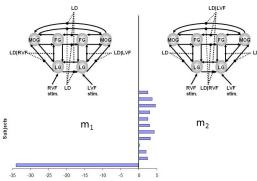
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Random Effects

11/12=92% subjects favoured model 2.



$$E[r_2|Y] = 0.84$$

 $p(r_2 > r_1|Y) = 0.99$

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where the latter is called the exceedance probability.



The use of Exceedance Probabilities (xp's) assumes the frequencies are different for each model.

But what if the model frequencies are all the same ? (H_0 : omnibus hypothesis)

Let $p_0 = p(H_0|Y)$. Then the (posterior) probability that frequencies are different is $1 - p_0$.

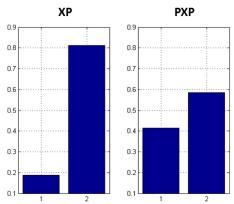
Rigoux et al. (*Neuroimage*, 2014) show how to compute p_o and then define Protected Exceedance Probabilities as

$$pxp = xp(1-p_o) + \frac{1}{K}p_o$$

where K is the number of models.

 p_o also referred to as 'Bayes Omnibus Risk (BOR)'.

Synthetic data (K = 2 models, N = 12 subjects, mean log evidence difference=0) .



We have $p_0 = 0.72$.

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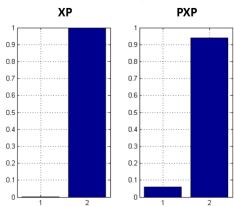
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Protected Exceedance Probabilities

Synthetic data (K = 2 models, N = 12 subjects, mean log evidence difference=1).



We have $p_0 = 0.11$.

PXPs also very useful for large *K*.

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PXPs

The ranking of models from RFX inference can depend on the comparison set.

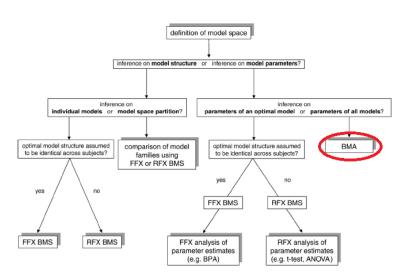
Say we have two models with 7 subjects prefering model 1 and 10 ten subjects preferring model 2. The model frequencies are $r_1 = 7/17 = 0.41$ and $r_2 = 10/17 = 0.59$.

Now say we add a third model which is similar to the second, and that 4 of the subjects that used to prefer model 2 now prefer model 3. The model frequencies are now $r_1 = 7/17 = 0.41$, $r_2 = 6/17 = 0.35$ and $r_3 = 4/17 = 0.24$.

This is like voting in elections.

Penny et al. PLOS-CB, 2010.

Model Averaging



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FFX Mode Inference

> FX Model ference

Model Averaging

RFX Paramet Inference

FFX Paramet nference

This can then be combined with the posterior model probabilities p(m|y) to compute a posterior over parameters

$$p(\theta|y) = \sum_{m} p(\theta, m|y)$$
$$= \sum_{m} p(\theta|m, y) p(m|y)$$

which is independent of model assumptions (within the chosen set). Here, we marginalise over *m*.

The sum over *m* could be restricted to eg. models within the winning family.

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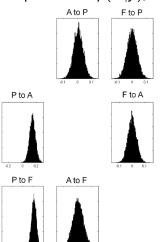
Model Averaging

RFX Paramete Inference

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The distribution $p(\theta|y)$ can be gotten by sampling; sample m from p(m|y), then sample θ from $p(\theta|m,y)$.



If a connection doesn't exist for model m the relevant samples are set to zero.

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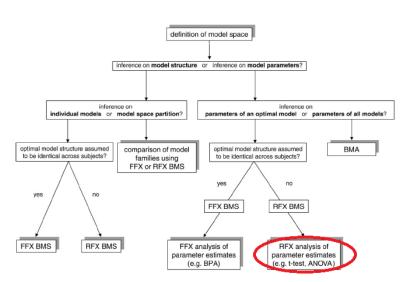
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Model Averaging

RFX Parameter Inference

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RFX Parameter Inference

If ith subject has posterior mean value m_i we can use these in Summary Statistic approach for group parameter inference (eg two-sample t-tests for control versus patient inferences).

eg P to A connection in controls: 0.20, 0.12, 0.32, 0.11, 0.01, ...

eg P to A connection in patients: 0.50, 0.42, 0.22, 0.71, 0.31, ...

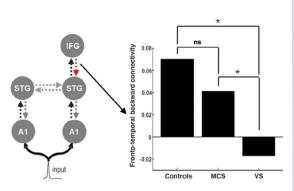
Two sample t-test shows the P to A connection is stronger in patients than controls (p < 0.05). Or one sample t-tests if we have a single group.

RFX is more conservative than BPA.

RFX Parameter Inference

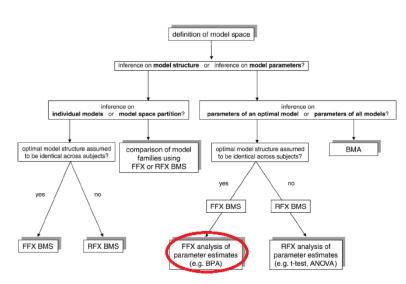
T-tests on backward connection from IFG to STG

Fig. 4. Quantitative effective connectivity analvsis revealed that the only significant difference between VS patients and controls was an impairment of backward connectivity from frontal to temporal cortex. MCS subjects showed significantly preserved connectivity compared with VS subjects and were not significantly different from controls.



Boly et al. Science, 2011

FFX Parameter Inference



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FFX Parameter Inference

FFX Parameter Inference

RFX parameter inference (eg. t-tests, F-tests) - allow for variability over eg. subjects.

FFX parameter inference - assumes no variability over eg. subjects/sessions.

FFX parameter inference - implemented using 'Bayesian Parameter Averaging' (BPA)

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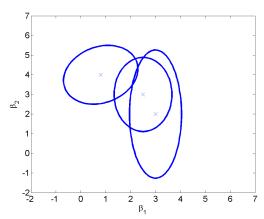
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Bayesian Parameter Averaging

If for the *i*th subject the posterior mean and precision are μ_i and Λ_i



Three subjects shown.

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If for the *i*th subject the posterior mean and precision are μ_i and Λ_i then the posterior mean and precision for the group are

$$\Lambda = \sum_{i=1}^{N} \Lambda_{i}$$

$$\mu = \Lambda^{-1} \sum_{i=1}^{N} \Lambda_{i} \mu_{i}$$

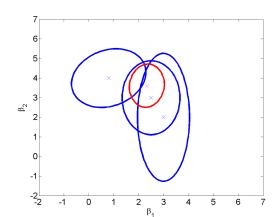
Kasses et al, Neuroimage, 2010.

This is a FFX analysis where each subject adds to the posterior precision.

Bayesian Parameter Averaging

$$\Lambda = \sum_{i=1}^{N} \Lambda_{i}$$

$$\mu = \Lambda^{-1} \sum_{i=1}^{N} \Lambda_{i} \mu_{i}$$



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If for the *i*th subject the posterior mean and precision are μ_i and Λ_i then the posterior mean and precision for the group are

$$\Lambda = \sum_{i=1}^{N} \Lambda_i - (N-1)\Lambda_0$$

$$\mu = \Lambda^{-1} \left(\sum_{i=1}^{N} \Lambda_i \mu_i - (N-1)\Lambda_0 \mu_0 \right)$$

Formulae augmented to accomodate non-zero priors Λ_0 and μ_0 .

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