

# Methods & Models for fMRI Analysis 2019

# GROUP ANALYSIS

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*With many thanks for slides & images to Guillaume Flandin*



University of  
Zurich UZH

**ETH**

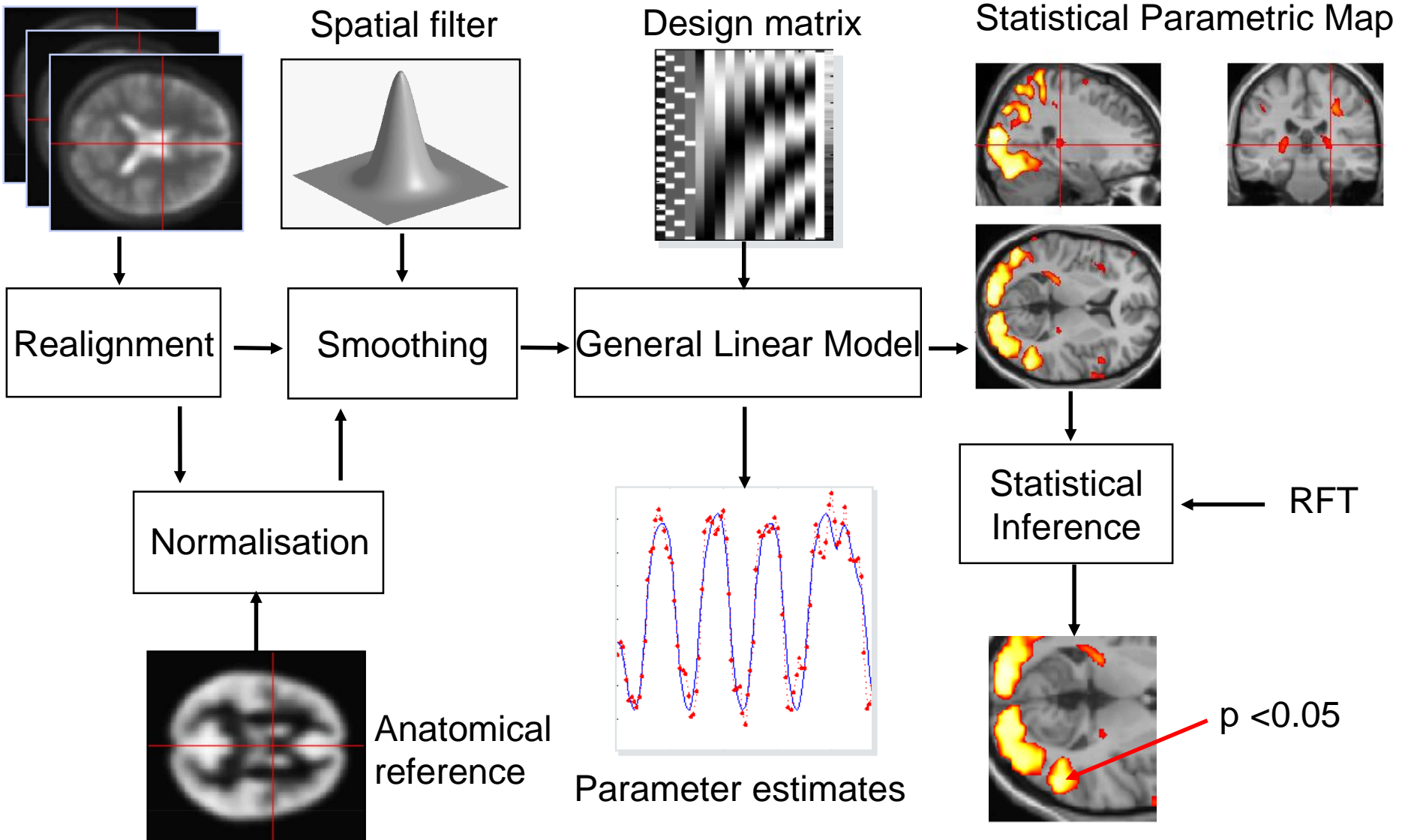
Eidgenössische Technische Hochschule Zürich  
Swiss Federal Institute of Technology Zurich



Translational Neuromodeling Unit

# Overview of SPM Steps

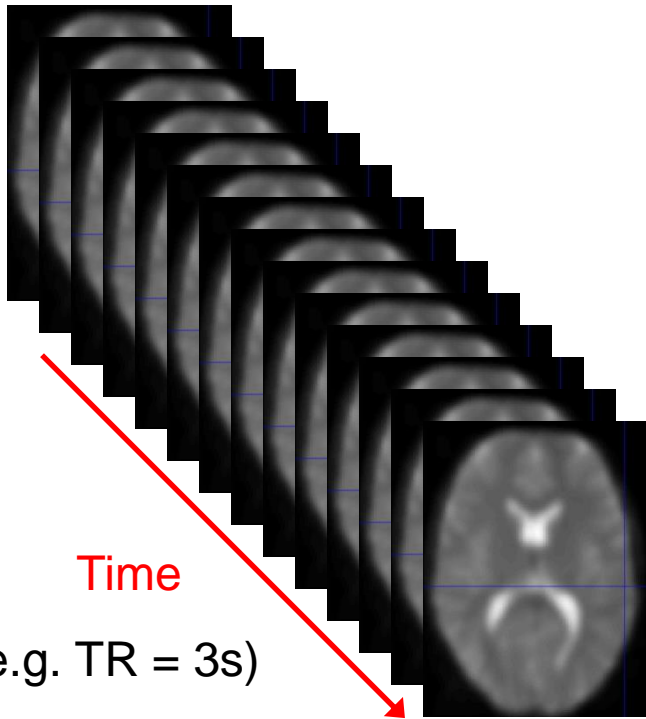
Image time-series



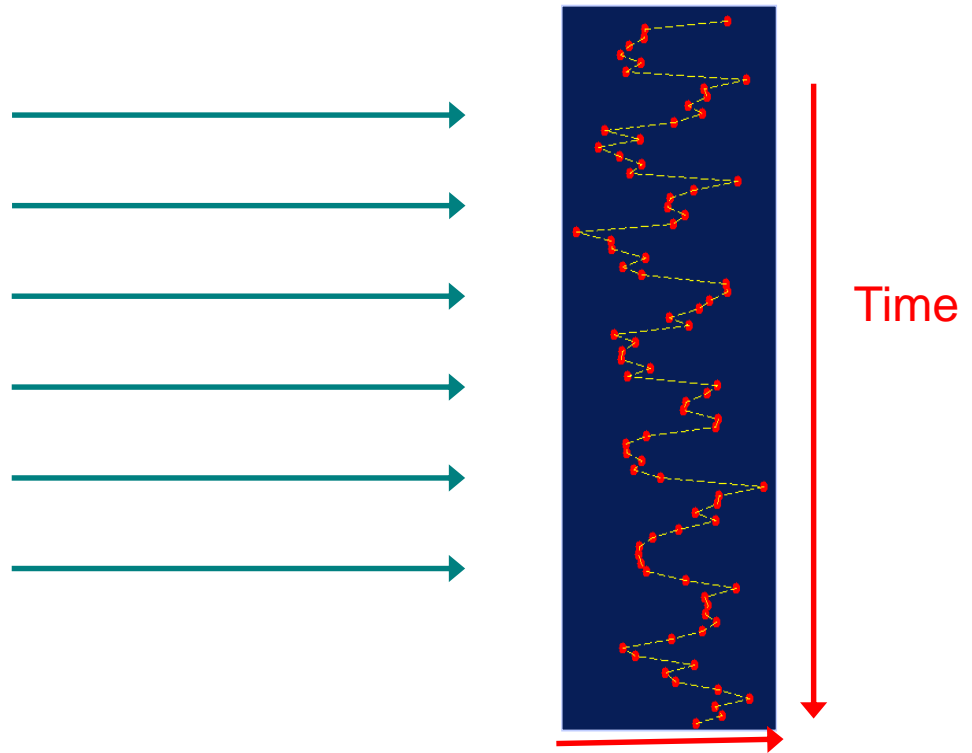
# 1<sup>st</sup> Level Analysis is within subject

$$y = X\beta + e$$

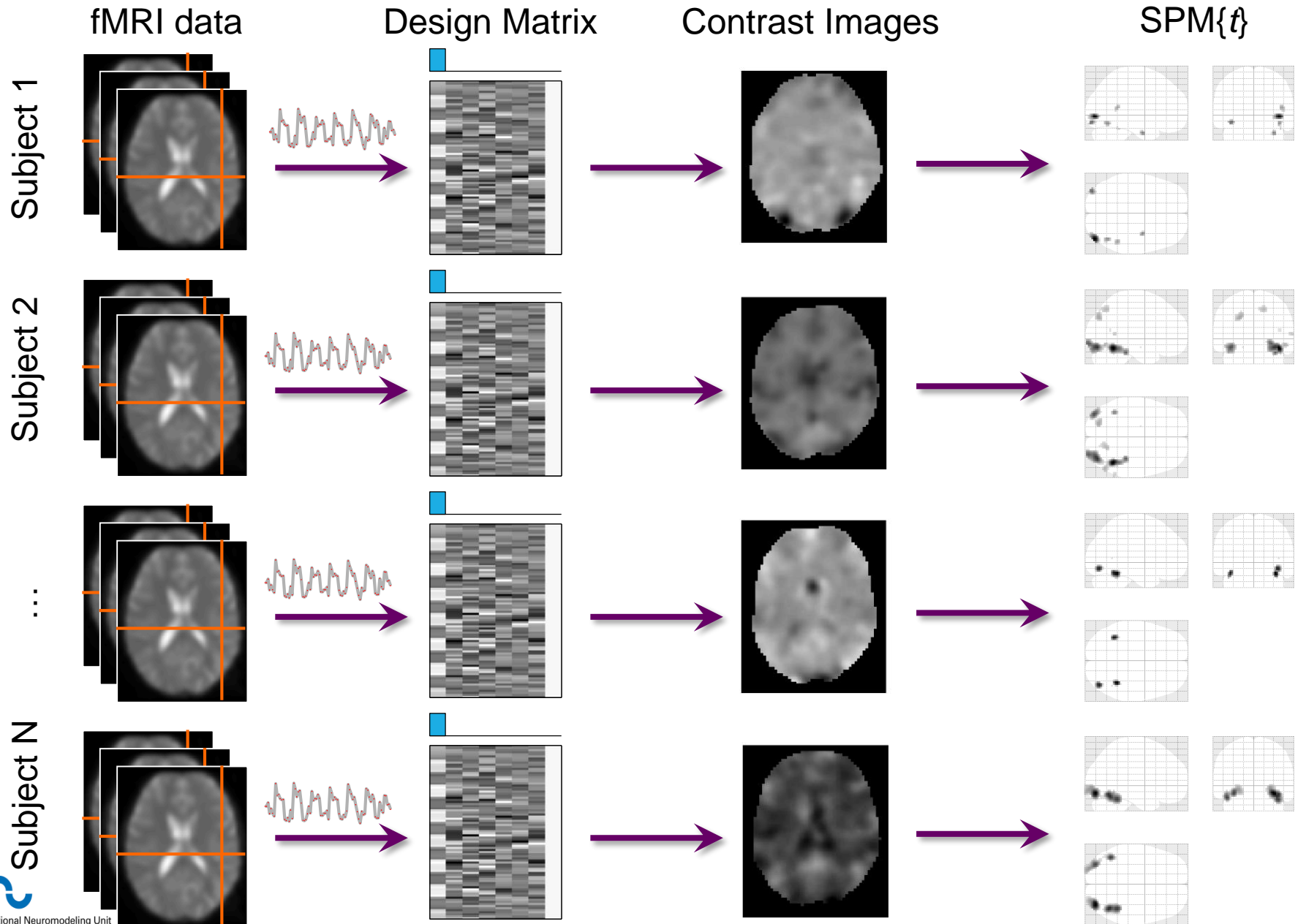
fMRI scans



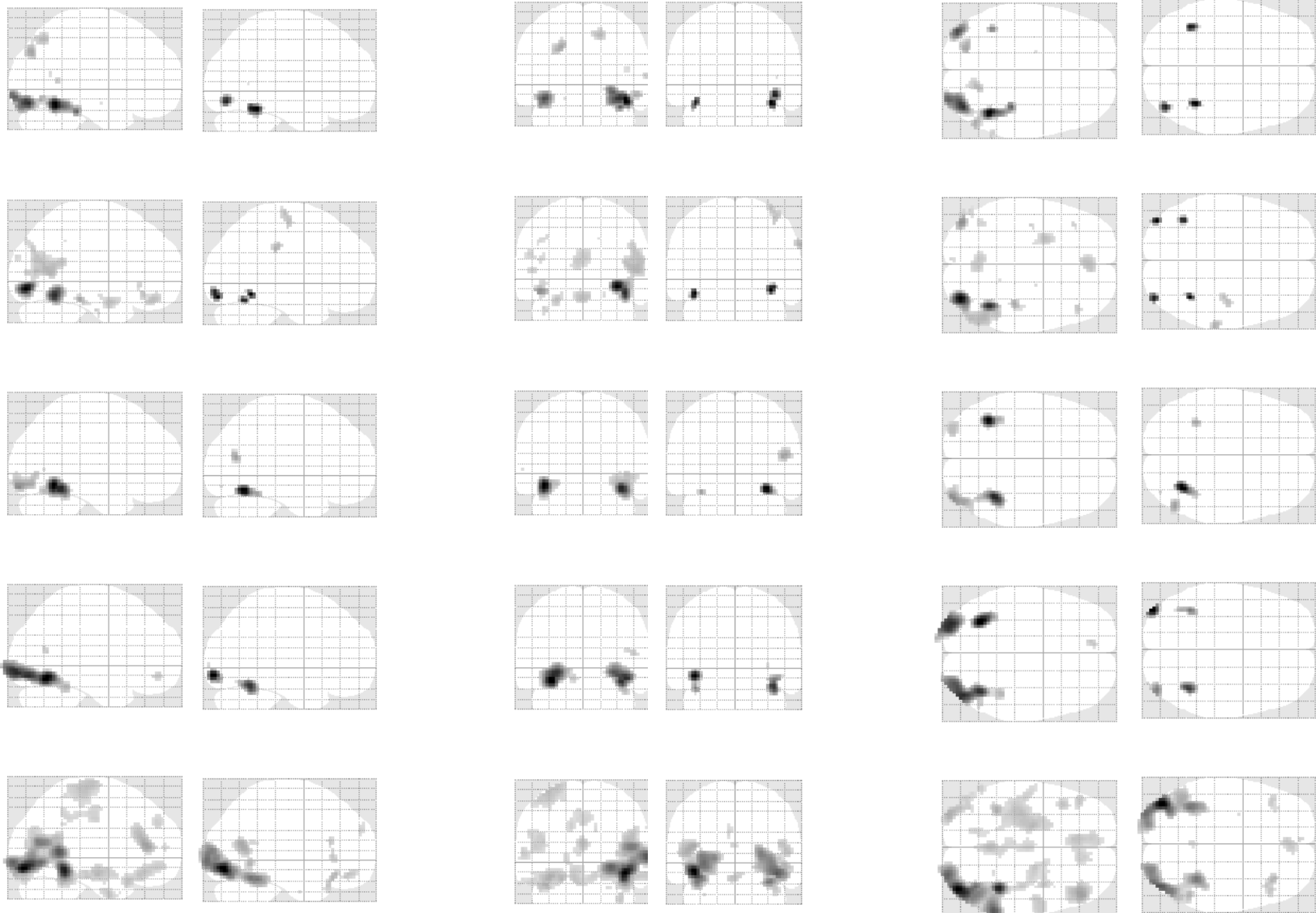
Voxel time course



# GLM: repeat over subjects

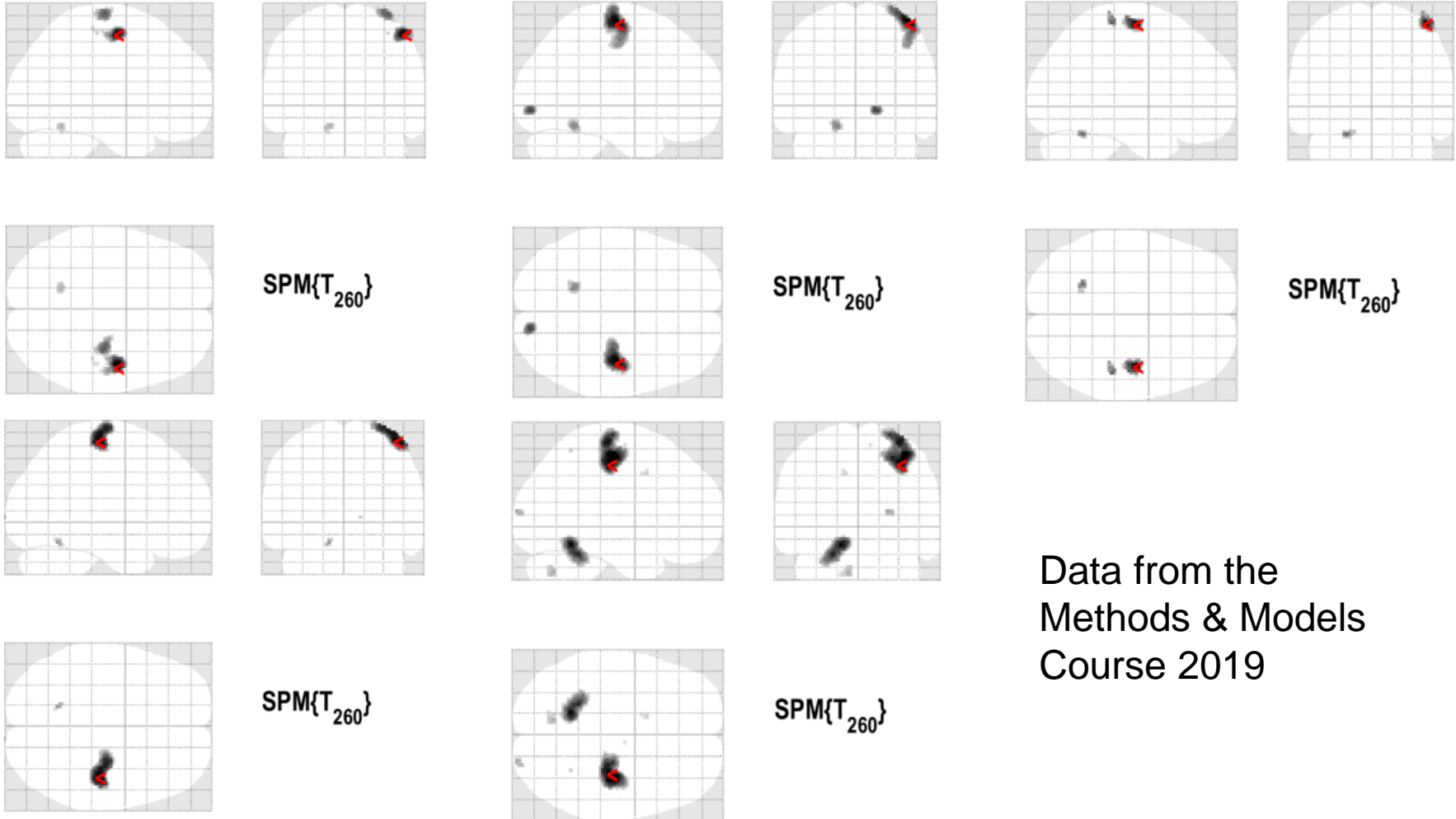


# First level analyses ( $p < 0.05$ FWE):



# First level analyses ( $p < 0.05$ FWE):

## Left Press > Right Press



Data from the  
Methods & Models  
Course 2019

# 2<sup>nd</sup> level analysis – across subjects

- It isn't enough to look just at individuals.
- So, we need to look at which voxels are showing a significant activation difference between levels of X consistently within a group.
  1. Average contrast effect across sample
  2. Variation of this contrast effect
  3. T-tests

# Group Analysis: Fixed vs Random

Does the group activate on average?

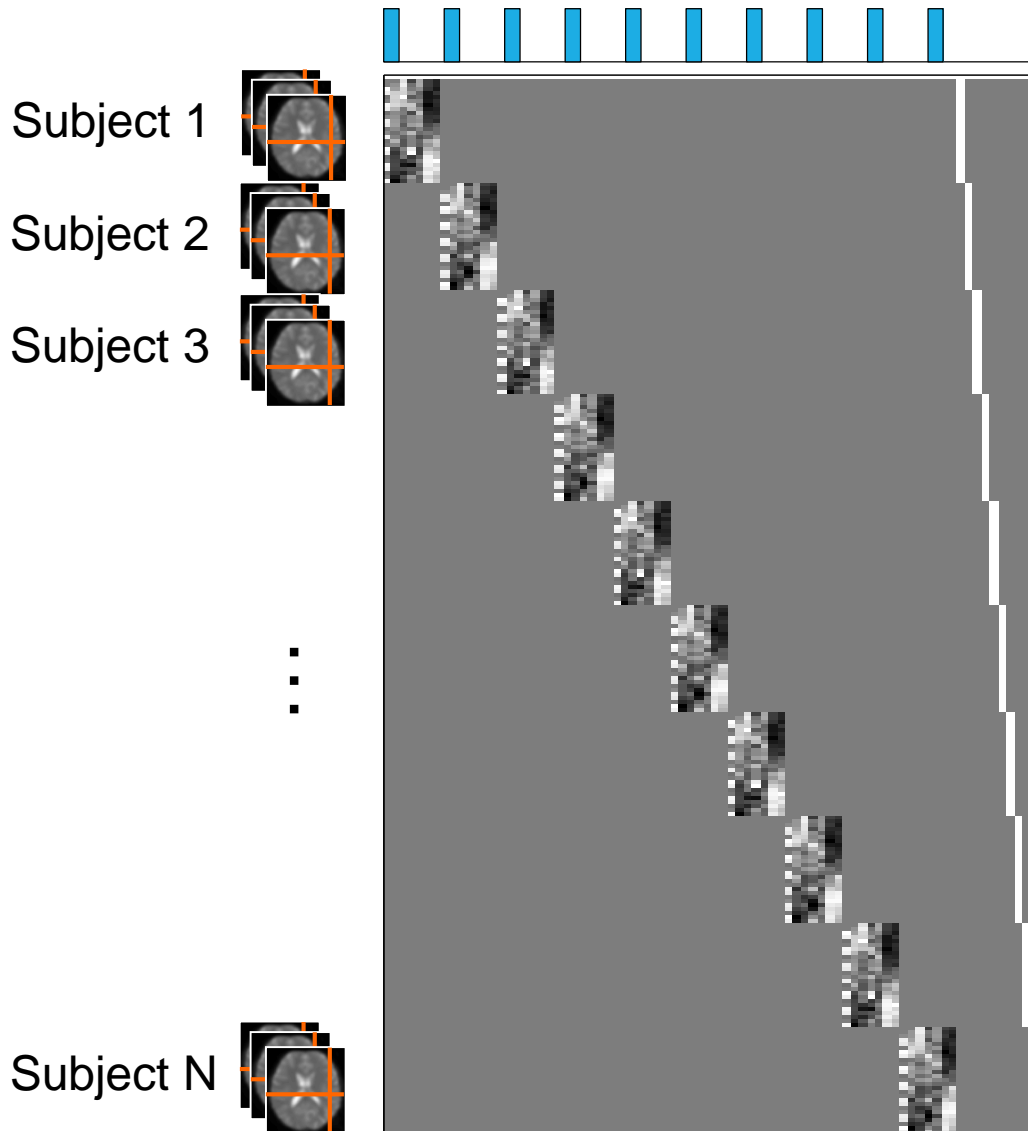


What group mean are we after?

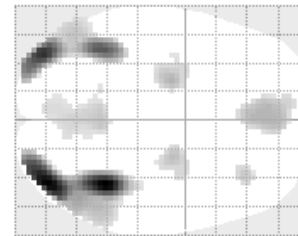
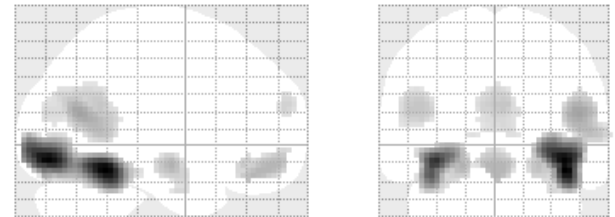
- The group mean for those exact 7 subjects?  
→ **Fixed effects analysis (FFX)**
- The group mean for the population from which these 7 subjects were drawn?  
→ **Random effects analysis (RFX)**



# Fixed effects analysis (FFX)



Modelling all subjects at once



variance over subjects at each voxel

# Fixed effects analysis (FFX)

$$y = X^{(1)}\beta^{(1)} + \varepsilon^{(1)}$$

Modelling all subjects at once

$$y = \begin{matrix} X_1^{(1)} & & \\ & X_2^{(1)} & \\ & & X_3^{(1)} \end{matrix} \beta^{(1)} + \varepsilon^{(1)}$$

- ✓ Simple model
- ✓ Lots of degrees of freedom
- ✗ Large amount of data
- ✗ Assumes common variance over subjects at each voxel

# Fixed effects

$$y = X^{(1)} \beta^{(1)} + \varepsilon^{(1)}$$



- Only one source of random variation (over sessions):

→ measurement error

Within-subject Variance

- True response magnitude is *fixed*.

# Whole Group – FFX calculation

- N subjects = 12 with each 50 scans = 600 scans

$$c = [4, 3, 2, 1, 1, 2, 3, 3, 3, 2, 4, 4]$$

Within subject variability:

$$s_w = [0.9, 1.2, 1.5, 0.5, 0.4, 0.7, 0.8, 2.1, 1.8, 0.8, 0.7, 1.1]$$

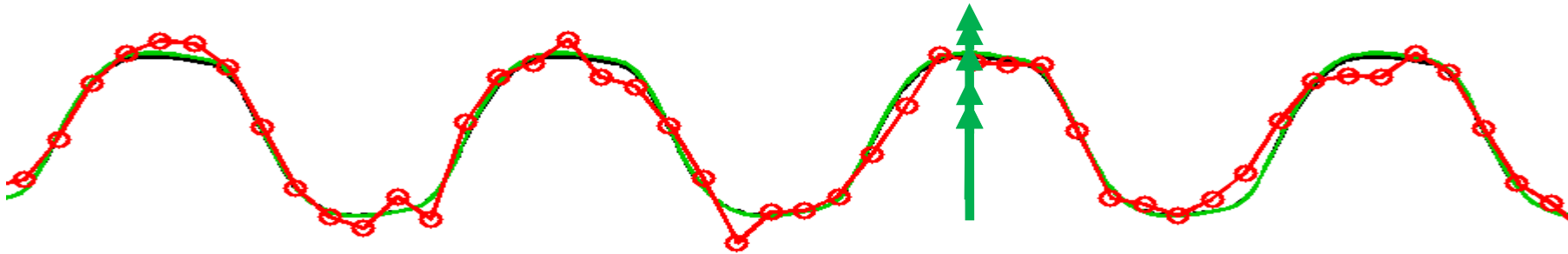
- Mean group effect = 2.67
- Mean  $s_w = 1.04$
- Standard Error Mean (SEM) =  $s_w / (\text{sqrt}(N)) = 0.04$

$$t = M / \text{SEM} = 62.7, p = 10^{-51}$$

# Random effects

$$y = X^{(1)} \beta^{(1)} + \varepsilon^{(1)}$$

$$\beta^{(1)} = X^{(2)} \beta^{(2)} + \varepsilon^{(2)}$$



- Two sources of random variation:

→ measurement errors

→ response magnitude (over subjects)

Within-subject Variance

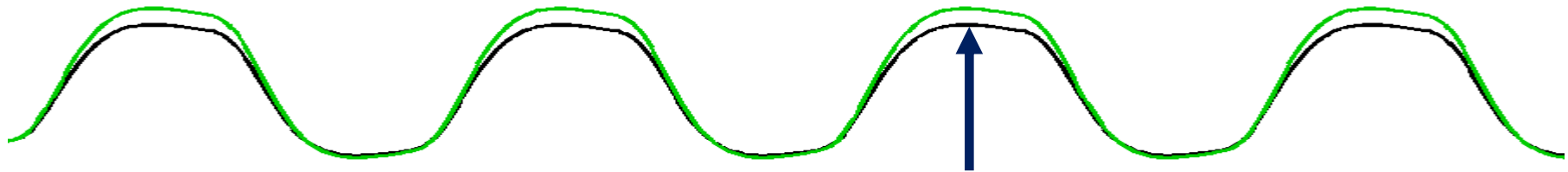
Between-subject Variance

- Response magnitude is *random*

→ each subject/session has random magnitude

# Random effects

$$y = X^{(1)} \beta^{(1)} + \varepsilon^{(1)}$$
$$\beta^{(1)} = X^{(2)} \beta^{(2)} + \varepsilon^{(2)}$$



- Two sources of random variation:

→ measurement errors

→ response magnitude (over subjects)

Within-subject Variance

Between-subject Variance

- Response magnitude is *random*

→ each subject/session has random magnitude

→ but population mean magnitude is *fixed*.

# Whole Group – RFX calculation

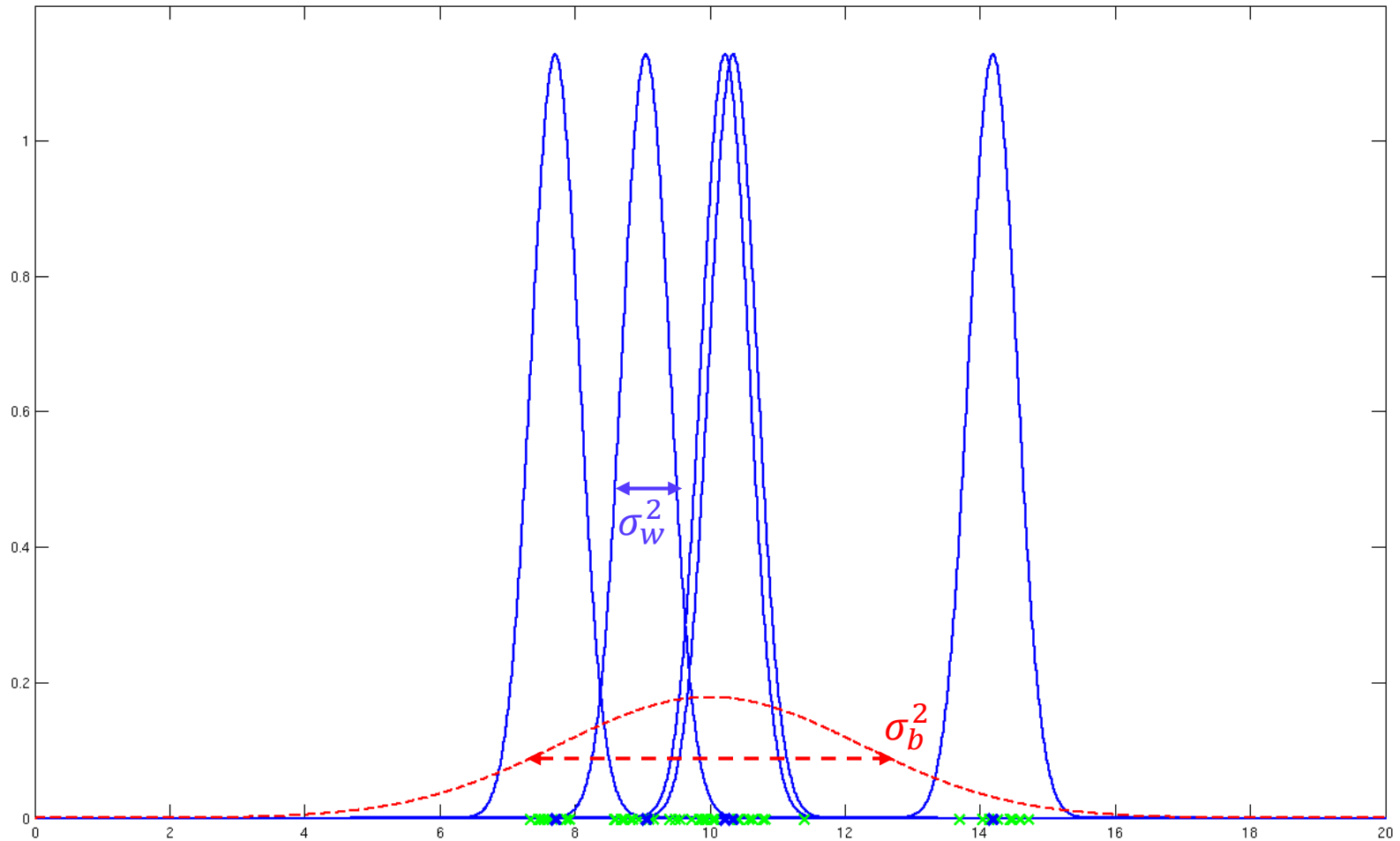
- N subjects = 12

$c = [4, 3, 2, 1, 1, 2, 3, 3, 3, 2, 4, 4]$

- Mean group effect = 2.67
- Mean  $s_b$  (SD) = 1.07
- Standard Error Mean (SEM) =  $s_b / (\text{sqrt}(N)) = 0.31$

$t = M / \text{SEM} = 8.61, p = 10^{-6}$

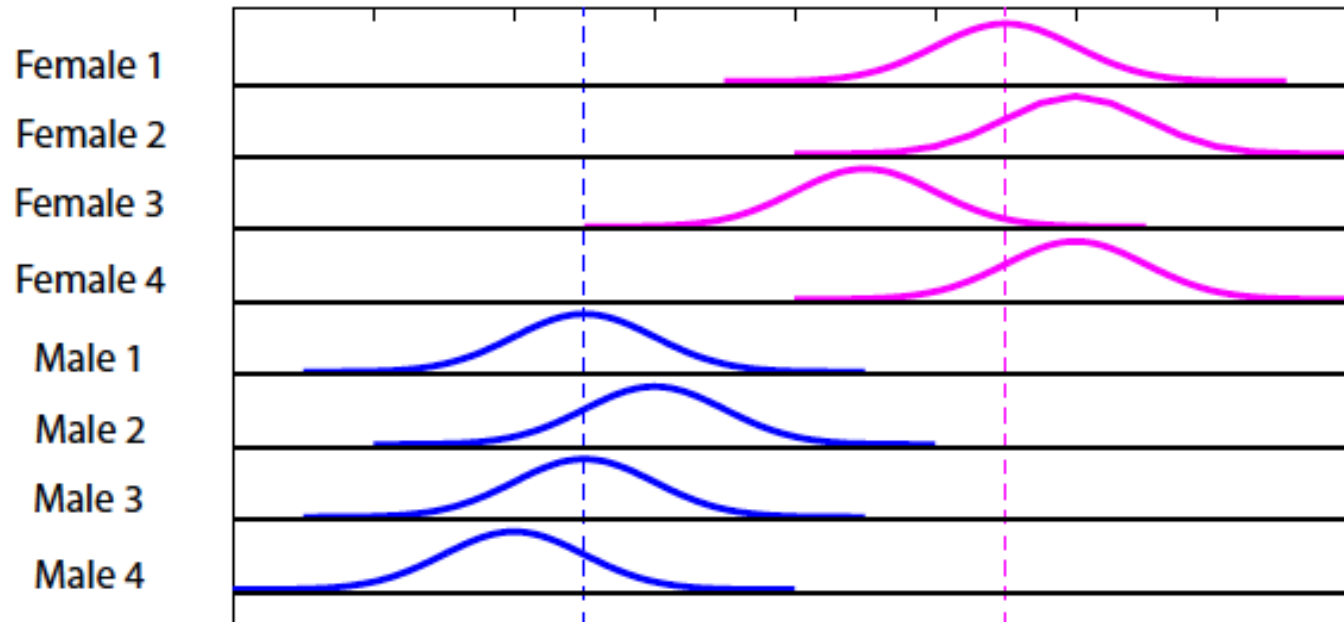
# Random effects



Probability model underlying random effects analysis



# Fixed vs random effects



*Handbook of functional MRI data analysis.* Poldrack, R. A., Mumford, J. A., & Nichols, T. E. Cambridge University Press, 2011

# Fixed vs random effects

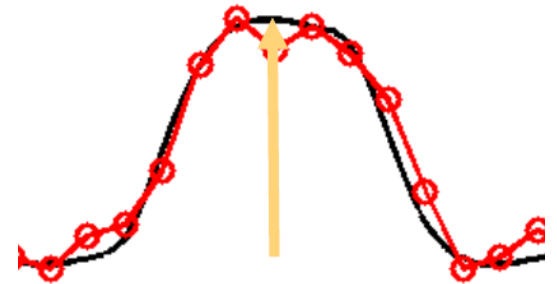
With **Fixed Effects Analysis (FFX)** we compare the group effect to the *within-subject variability*. It is not an inference about the population from which the subjects were drawn.

With **Random Effects Analysis (RFX)** we compare the group effect to the *between-subject variability*. It is an inference about the population from which the subjects were drawn. If you had a new subject from that population, you could be confident they would also show the effect.

# Fixed vs random effects

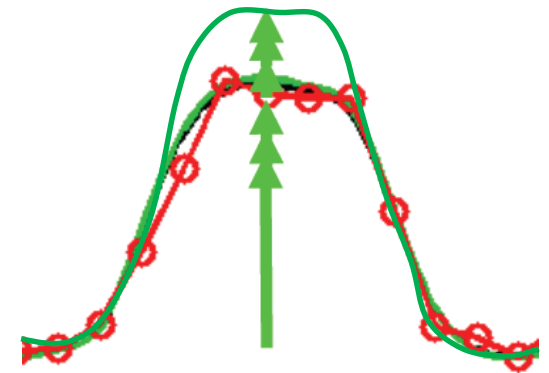
## Fixed-effects

- Is not of interest across a population
- Used for a case study
- Only source of variation is measurement error (Response magnitude is **fixed**)



## Random-effects

- If I have to take another sample from the population, I would get the same result
- Two sources of variation
  - Measurement error
  - Response magnitude is **random** (population mean magnitude is fixed)



## **Hierarchical linear models:**

- Random effects models
- Mixed effects models
- Nested models
- Variance components models

... all the same

... all alluding to multiple sources of variation  
(in contrast to fixed effects)

# Linear hierarchical models

## Hierarchical Model

$$\begin{aligned}y &= X^{(1)}\theta^{(1)} + \varepsilon^{(1)} \\ \theta^{(1)} &= X^{(2)}\theta^{(2)} + \varepsilon^{(2)} \\ &\vdots \\ \theta^{(n-1)} &= X^{(n)}\theta^{(n)} + \varepsilon^{(n)}\end{aligned}$$

## Multiple variance components at each level

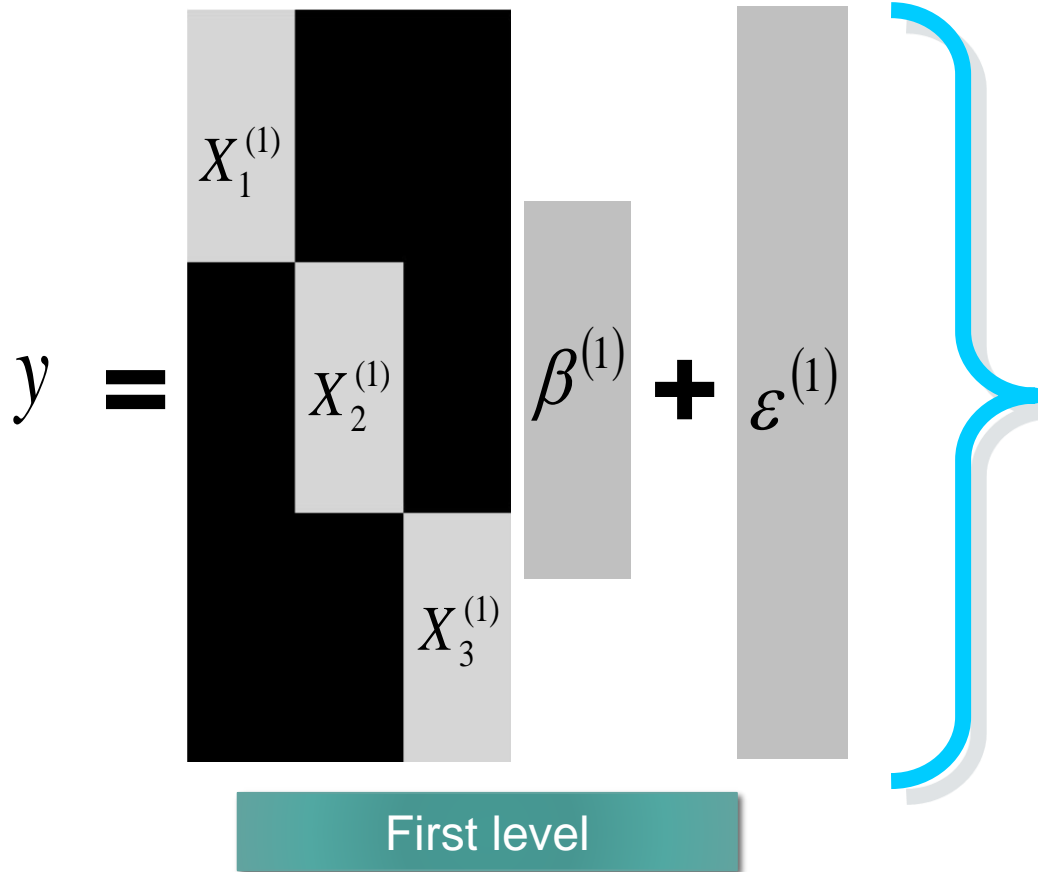
$$C_{\varepsilon}^{(i)} = \sum_k \lambda_k^{(i)} Q_k^{(i)}$$

At each level, distribution of parameters is given by level above.

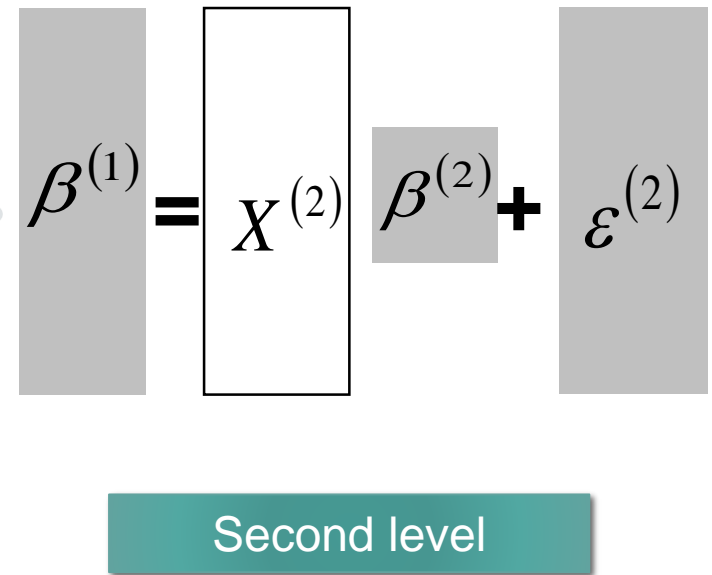
What we don't know: distribution of parameters and variance parameters (hyperparameters).

# Hierarchical models

## Example: Two level model

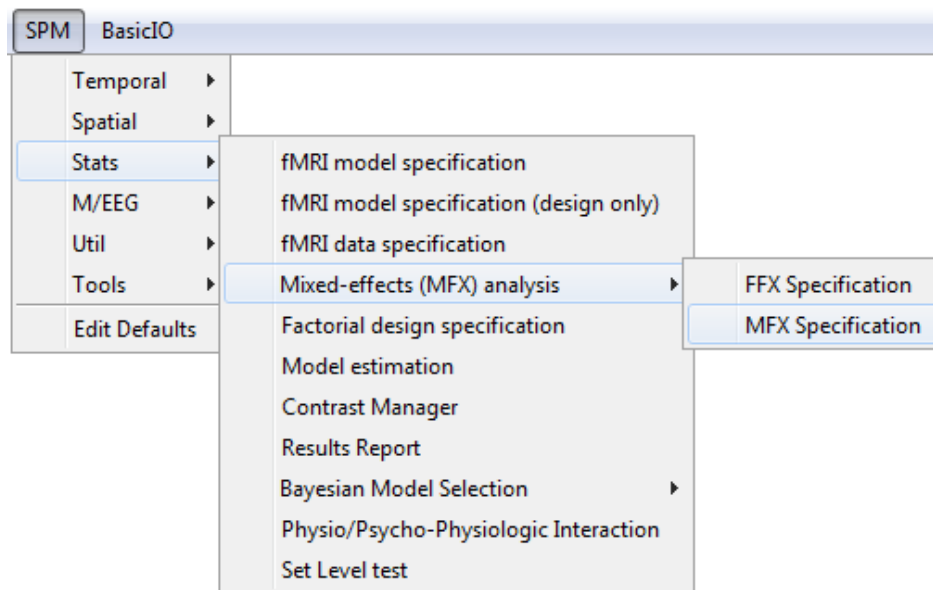


$$y = X^{(1)} \beta^{(1)} + \varepsilon^{(1)}$$
$$\beta^{(1)} = X^{(2)} \beta^{(2)} + \varepsilon^{(2)}$$



# Hierarchical models

- Restricted Maximum Likelihood (ReML)
- Parametric Empirical Bayes
- Expectation-Maximisation Algorithm



`spm_mfx.m`

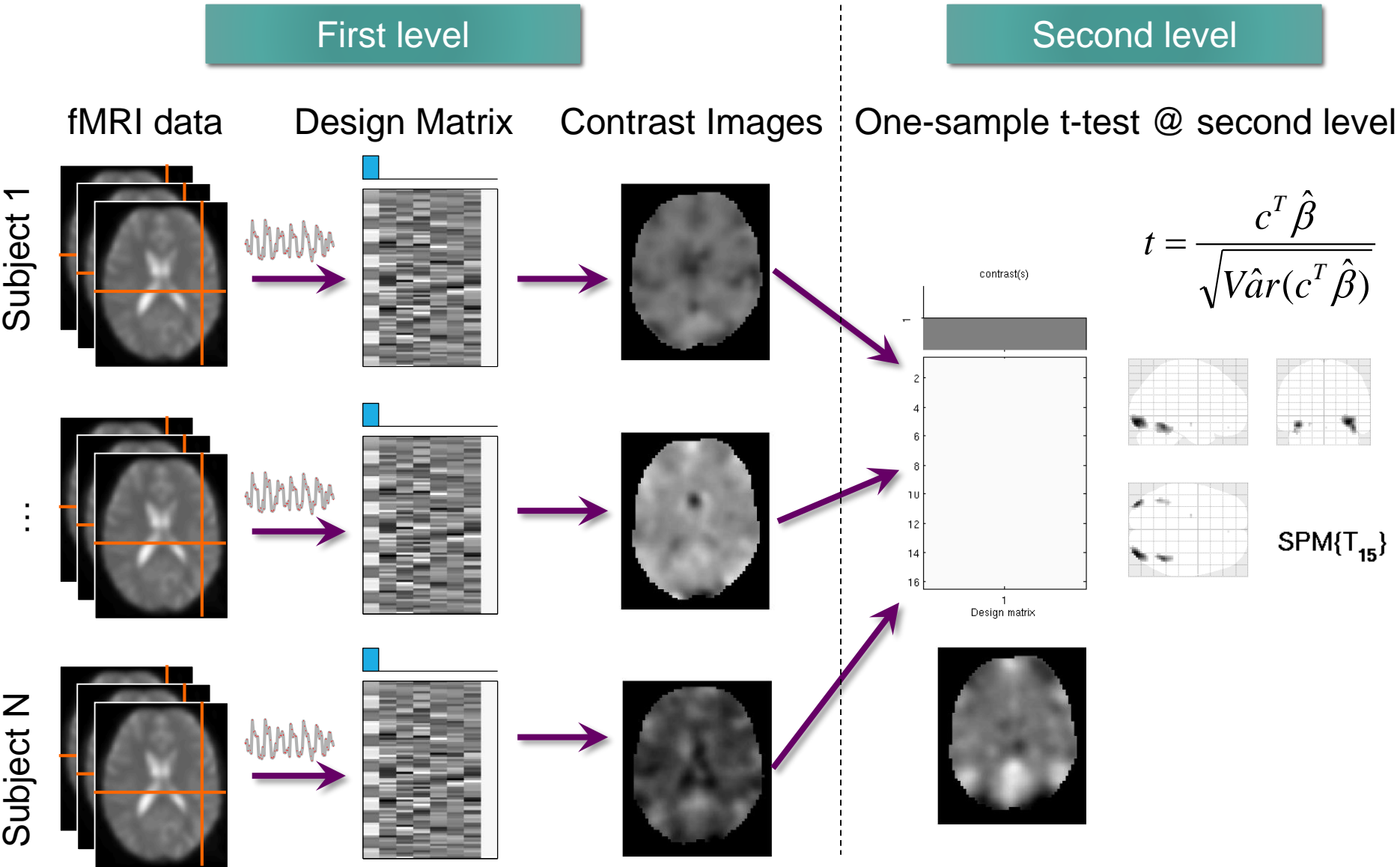
*Mixed-effects and fMRI studies.* Friston et al., NeuroImage, 2005.

# Practical problems

- Full MFX inference using REML or EM for a whole-brain 2-level model has enormous computational costs
  - for many subjects and scans, covariance matrices become extremely large
  - nonlinear optimisation problem for each voxel
- Moreover, sometimes we are only interested in one specific effect and do not want to model all the data.
- Is there a fast approximation?



# Summary Statistics RFX Approach



## Assumptions

- The summary statistics approach is exact if for each session/subject:
  - Within-subjects variances the same
  - First level design the same (e.g. number of trials)
- However, summary statistics approach is robust against typical violations

*Mixed-effects and fMRI studies.* Friston et al., NeuroImage, 2005.

*Statistical Parametric Mapping: The Analysis of Functional Brain Images.* Elsevier, 2007.

*Simple group fMRI modeling and inference.* Mumford & Nichols. NeuroImage, 2009.

# Summary Statistics RFX Approach

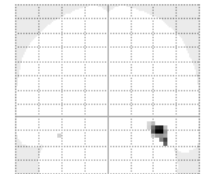
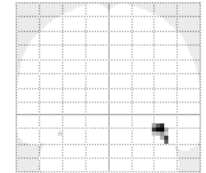
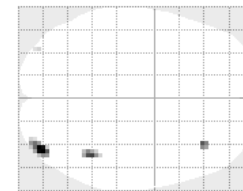
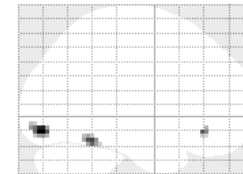
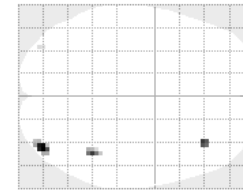
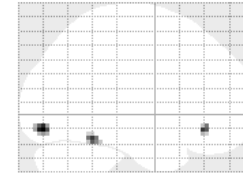
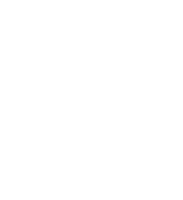
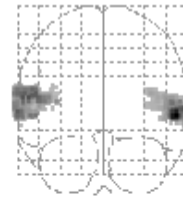
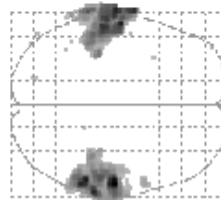
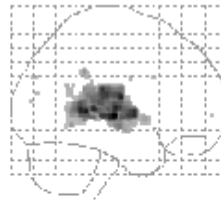
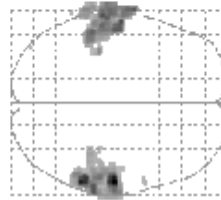
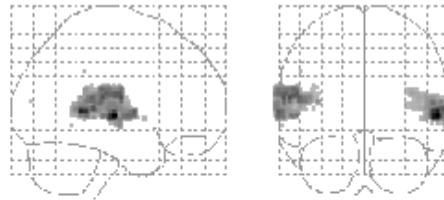
## Robustness

Summary statistics



SPM uses this!

Hierarchical Model



Listening to words

Viewing faces

- **One effect per subject:**
  - Summary statistics approach
  - One-sample t-test at the second level
- **More than one effect per subject or multiple groups:**
  - Non-sphericity modelling
  - Covariance components and ReML

# Reminder: sphericity

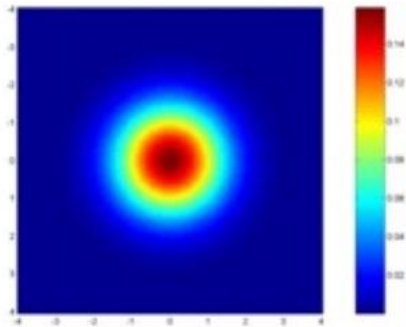
$$y = X\theta + \varepsilon$$

„sphericity“ means:

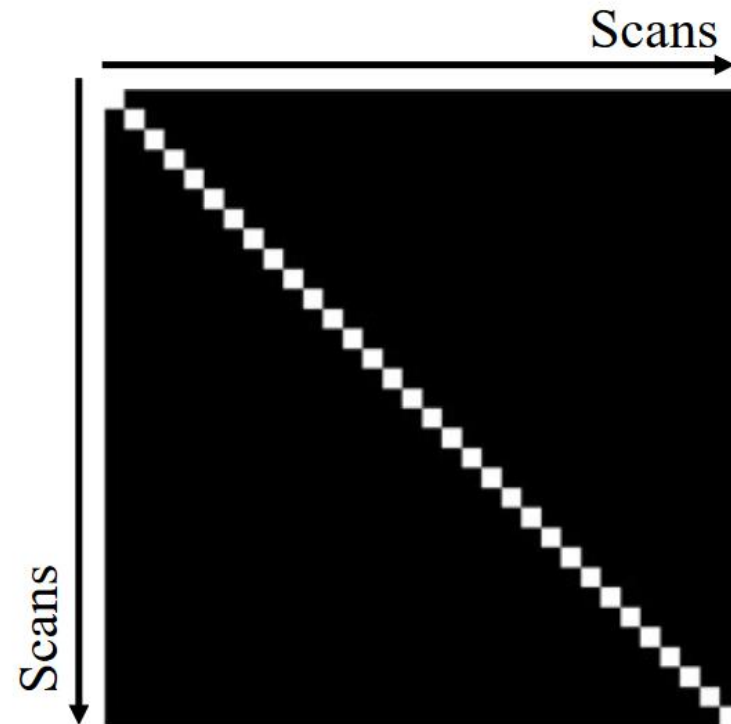
$$\text{Cov}(\varepsilon) = \sigma^2 I$$

i.e.  $\text{Var}(\varepsilon_i) = \sigma^2$

$$C_\varepsilon = \text{Cov}(\varepsilon) = E(\varepsilon\varepsilon^T)$$



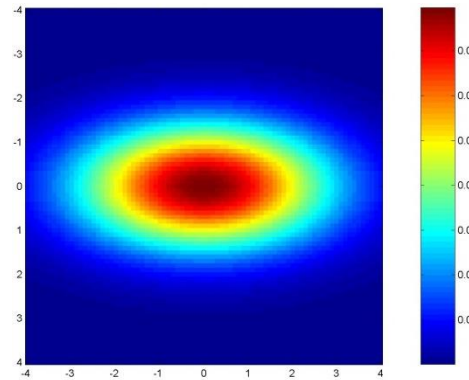
$$\text{Cov}(e) = \begin{bmatrix} 1 & 0 \\ 0 & 1 \end{bmatrix}$$



# GLM assumes Gaussian “spherical” (i.i.d.) errors

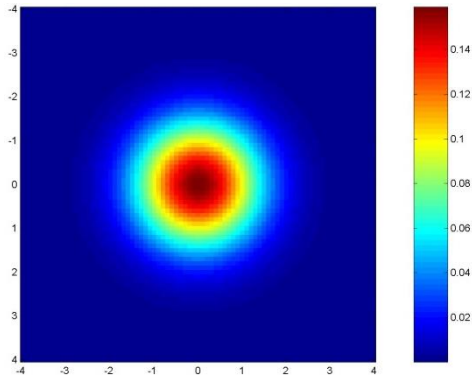
**sphericity = iid:**  
error covariance is  
scalar multiple of  
identity matrix:  
 $\text{Cov}(e) = \sigma^2 \mathbf{I}$

Examples for non-sphericity:

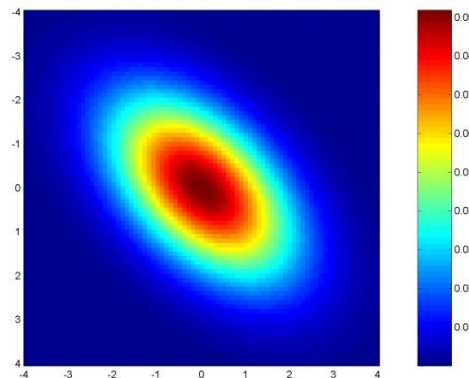


$$\text{Cov}(e) = \begin{bmatrix} 4 & 0 \\ 0 & 1 \end{bmatrix}$$

non-identically  
distributed



$$\text{Cov}(e) = \begin{bmatrix} 1 & 0 \\ 0 & 1 \end{bmatrix}$$



$$\text{Cov}(e) = \begin{bmatrix} 2 & 1 \\ 1 & 2 \end{bmatrix}$$

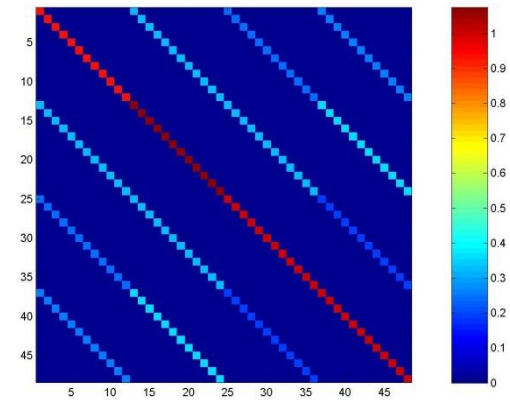
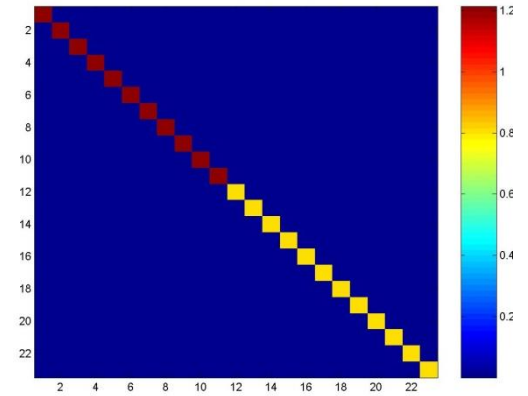
non-independent

# 2nd level: Non-sphericity

Errors are independent  
but not identical  
(e.g. different groups (patients, controls))

Errors are not independent  
and not identical  
(e.g. repeated measures for each subject  
(multiple basis functions, multiple  
conditions, etc.))

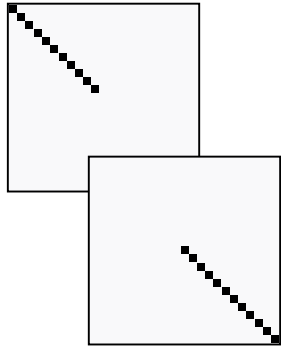
Error covariance matrix



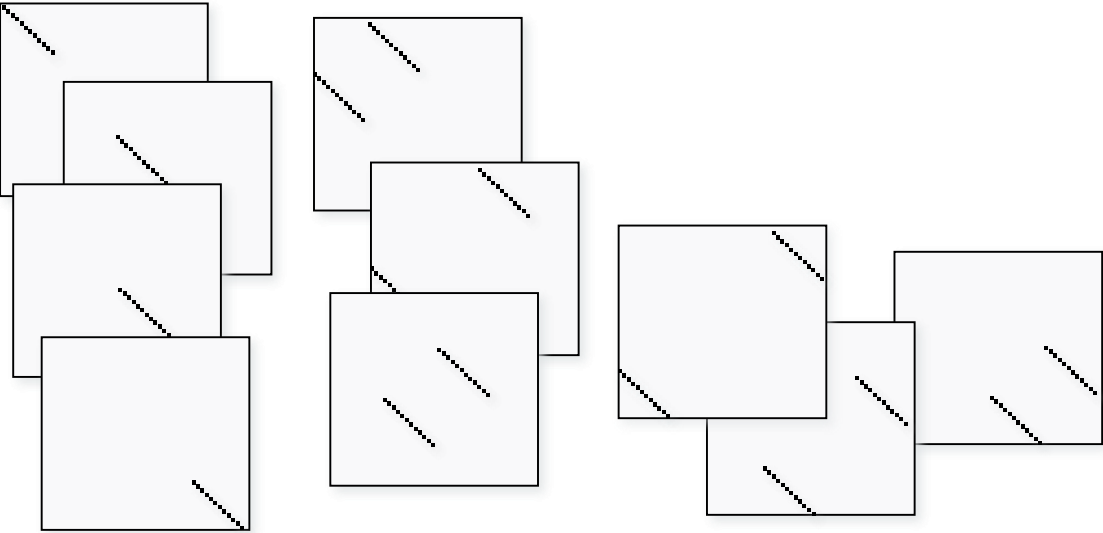
# 2nd level: Variance components

$$\text{Cov}(\varepsilon) = \sum_k \lambda_k Q_k$$

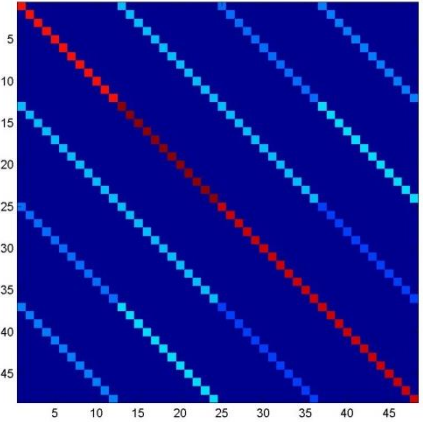
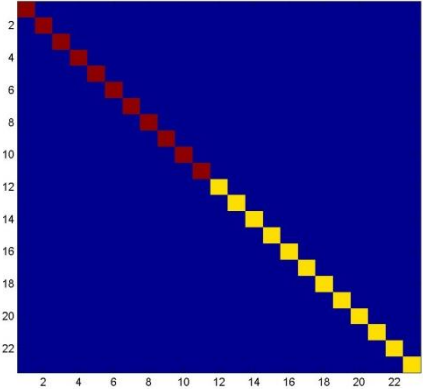
$Q_k$ 's:



$Q_k$ 's:



Error covariance matrix



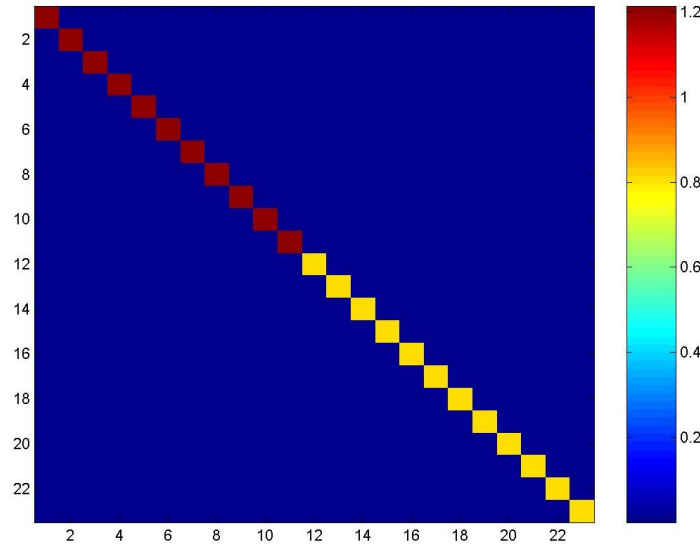
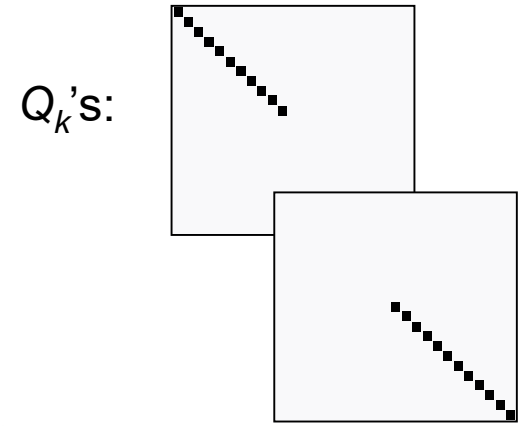


# Example 1: between-subjects ANOVA

- Stimuli:
  - Auditory presentation (SOA = 4 sec)
  - 250 scans per subject, block design
  - 2 conditions
    - Words, e.g. “book”
    - Words spoken backwards, e.g. “koob”
- Subjects:
  - 12 controls
  - 11 blind people

# Example 1: Covariance components

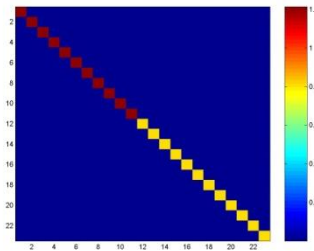
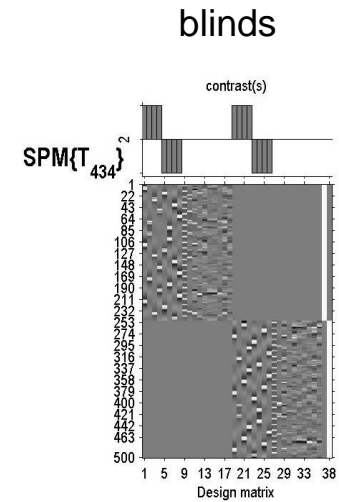
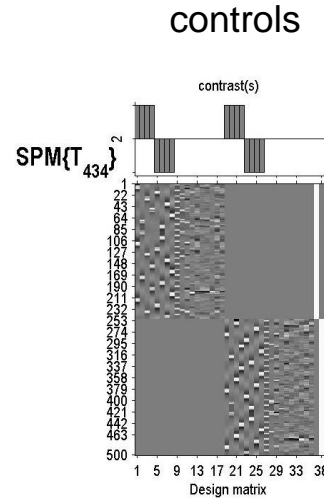
- Two-sample t-test:
  - Errors are independent but not identical.
  - 2 covariance components



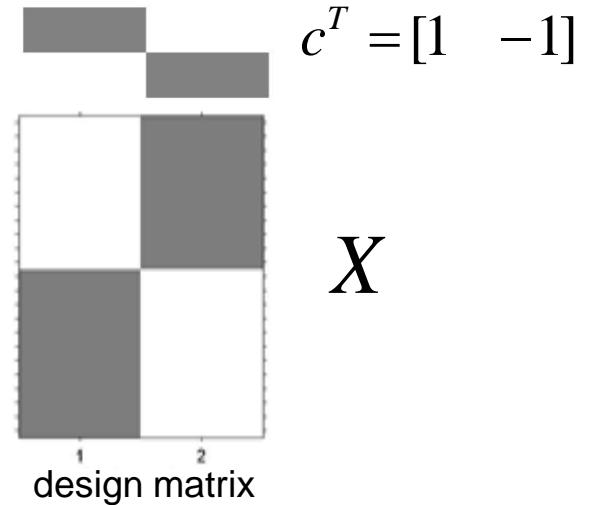
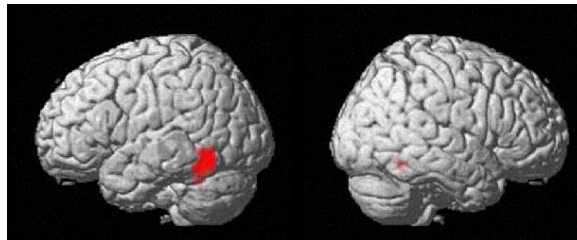
Error covariance matrix

# Example 1: Group differences

First Level



$$Cov(\epsilon)$$



Second Level

# Example 2: within-subjects ANOVA

## – Stimuli:

- Auditory presentation (SOA = 4 sec)
- 250 scans per subject, block design

## • Words:

Motion	Sound	Visual	Action
“jump”	“click”	“pink”	“turn”

## – Subjects:

- 12 controls

## – Question:

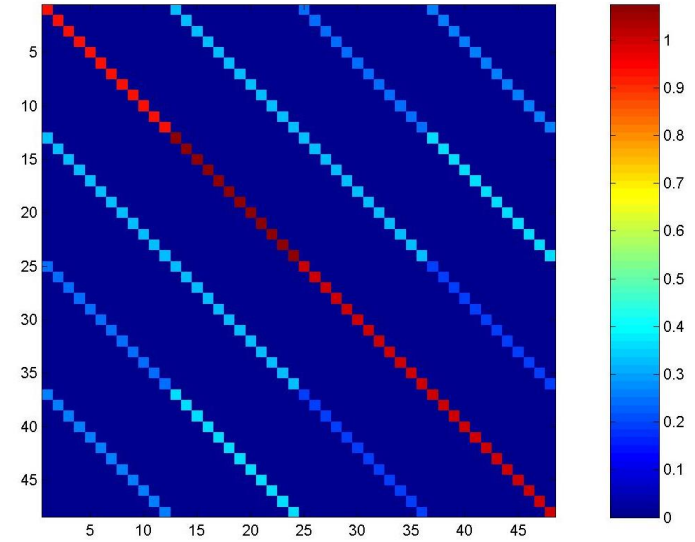
- What regions are generally affected by the semantic content of the words?

Noppeney et al., Brain, 2003.

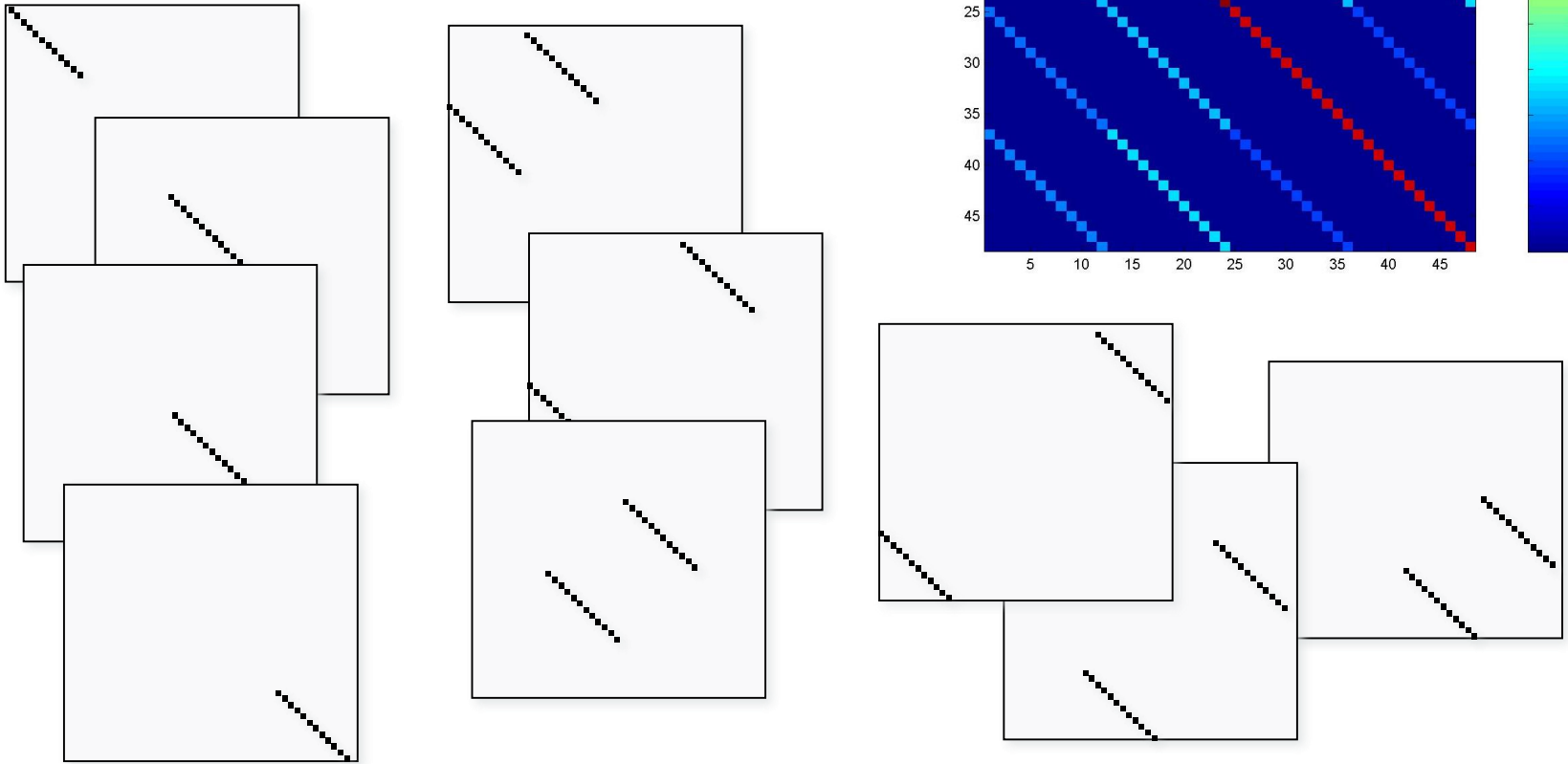
# Example 2: Covariance components

→ Errors are not independent and not identical

Error covariance matrix



$Q_k$ 's:



# Example 2: Repeated measures ANOVA

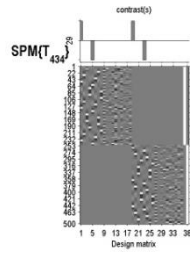
First Level

Motion

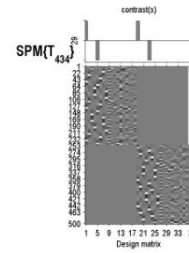
Sound

Visual

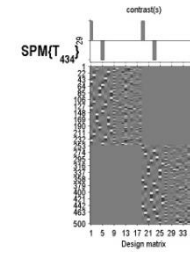
Action



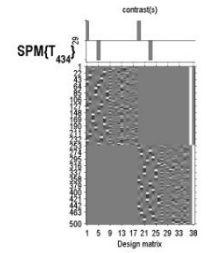
?



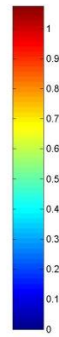
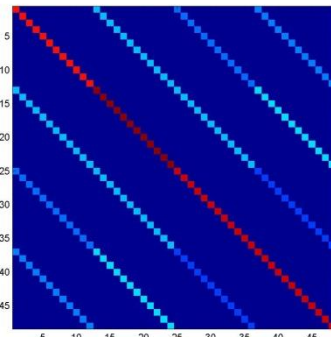
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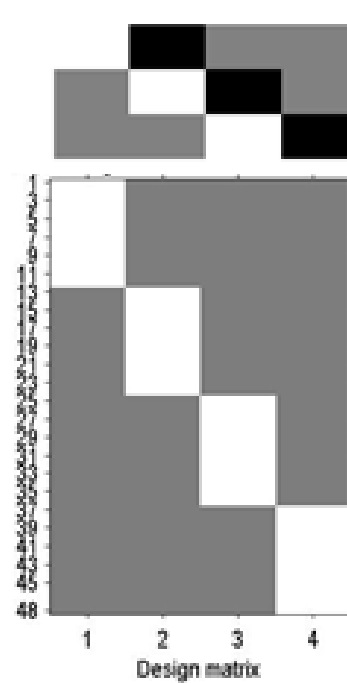
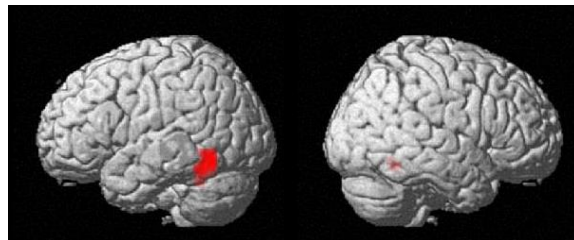
?



Second Level



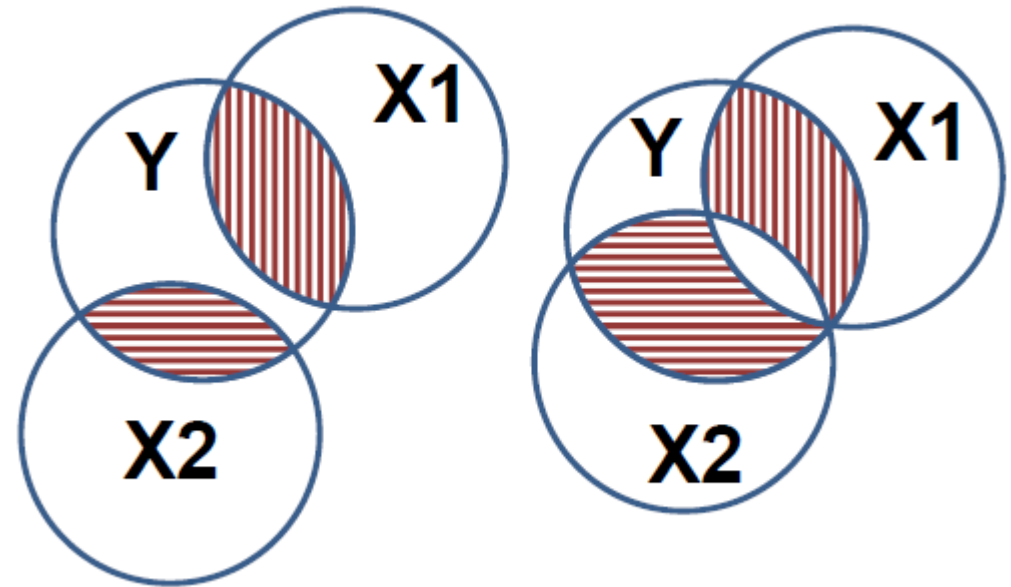
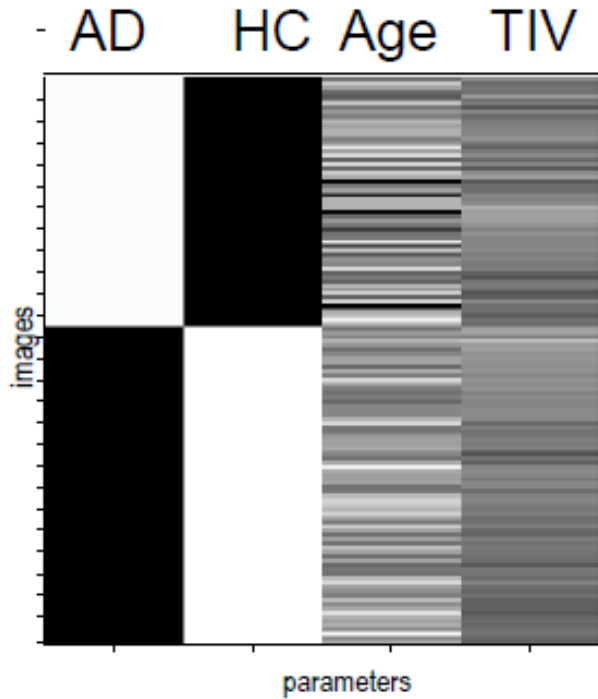
$Cov(\varepsilon)$



$$c^T = \begin{pmatrix} 1 & -1 & 0 & 0 \\ 0 & 1 & -1 & 0 \\ 0 & 0 & 1 & -1 \end{pmatrix}$$

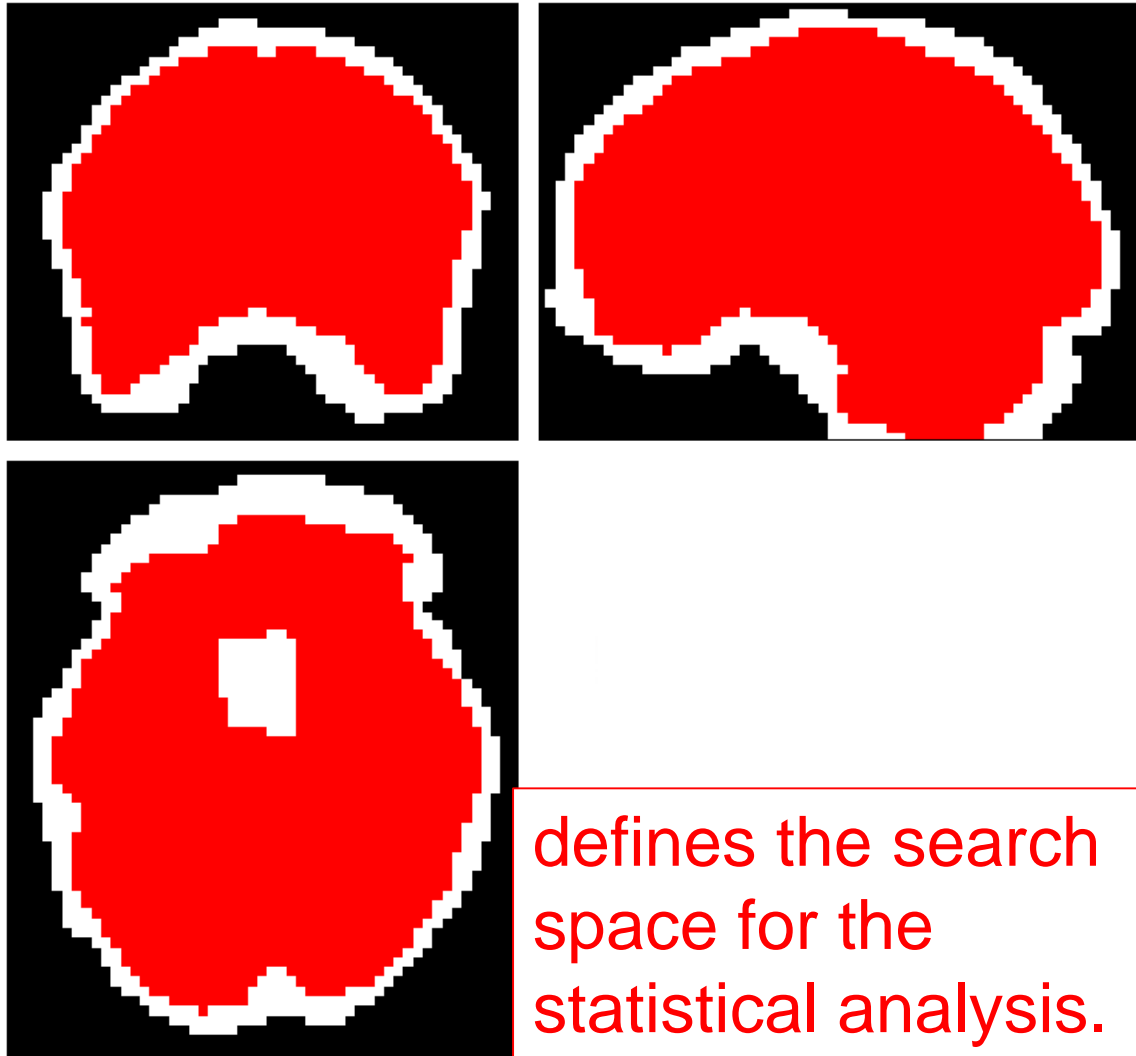
$X$

# ANCOVA model



Mean centering continuous covariates for a group fMRI analysis, by J. Mumford:  
[http://mumford.fmripower.org/mean\\_centering/](http://mumford.fmripower.org/mean_centering/)

# Analysis mask: logical AND

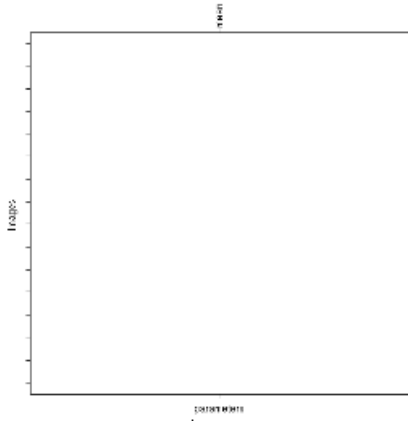




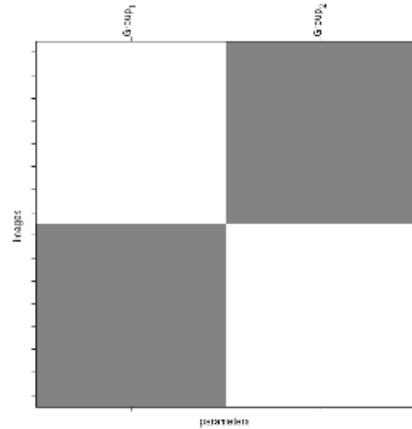
# SPM interface: factorial design specification

## Options:

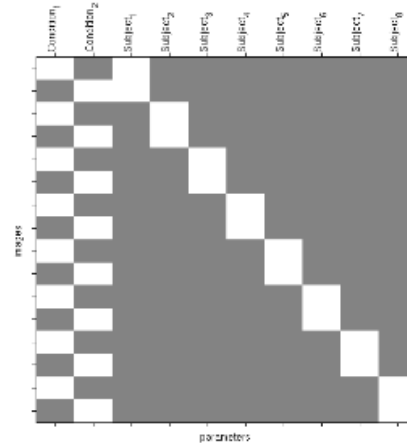
One-sample t-test



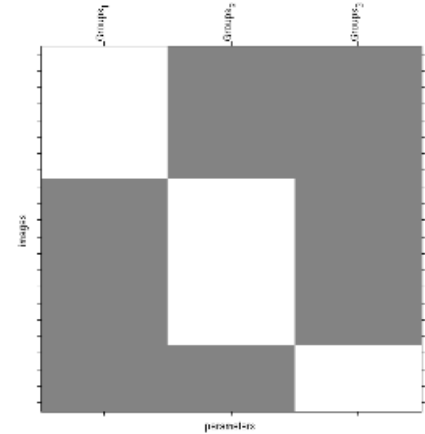
Two-sample t-test



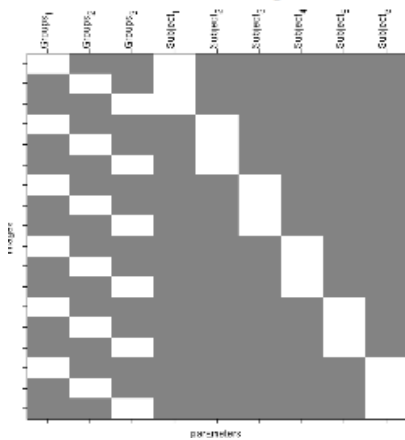
Paired t-test



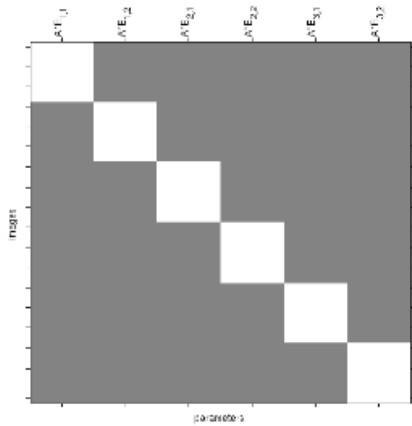
One-way ANOVA



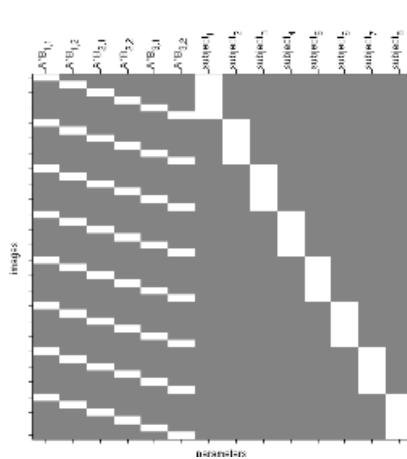
One-way ANOVA within-subject



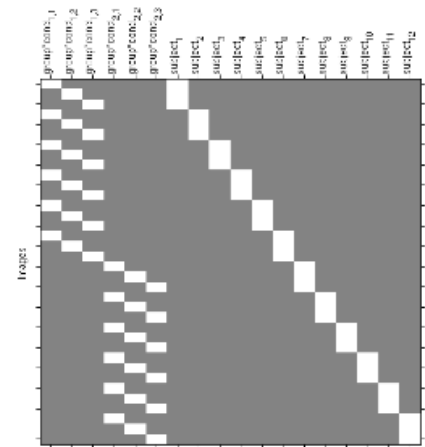
Full Factorial



Flexible Factorial



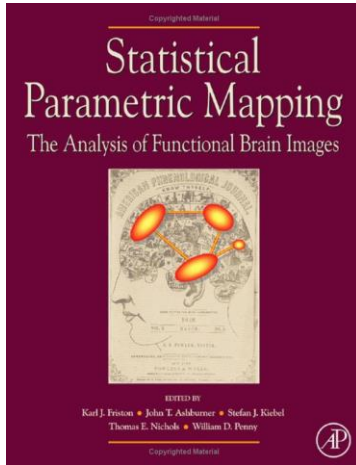
Flexible Factorial



# Summary

- Group inference usually proceeds with **RFX analysis**, not FFX. Group effects are compared to between rather than within subject variability.
- **Hierarchical models** provide a gold-standard for RFX analysis but are computationally intensive.
- **Summary statistics** approach is a robust method for RFX group analysis.
- Can also use '**ANOVA**' or '**ANOVA within subject**' at second level for inference about multiple experimental conditions or multiple groups.

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