# **Multiple testing**

#### Justin Chumbley

Laboratory for Social and Neural Systems Research University of Zurich

With many thanks for slides & images to:

FIL Methods group

# Overview of SPM – Random field theory





*contrast* of estimated parameters

*t* =



*contrast* of estimated parameters

*t* =



Decision:  $H_0$ ,  $H_1$ : zero/non-zero activation



*t* =



Decision:  $H_0$ ,  $H_1$ : zero/non-zero activation





Decision:  $H_0$ ,  $H_1$ : zero/non-zero activation

*contrast* of estimated parameters

*t* =



Decision:  $H_0$ ,  $H_1$ : zero/non-zero activation

Decision rule (threshold) *h*, determines related error rates  $\alpha_h$ ,  $\beta_h$ 

*contrast* of estimated parameters



Convention: Penalize complexity Choose *h* to give acceptable  $\alpha_h$  under H<sub>0</sub>



#### Multiple tests



#### What is the problem?



*t* =

## **Multiple tests**



Penalize each independent opportunity for error.

 $p(1 \text{ or more } FP) = FWER_h$  $E(\frac{FP}{All \text{ positives}}) = FDR$ 

*contrast* of estimated parameters

t =

#### **Multiple tests**



#### Bonferonni

 $FWER_{h} \leq N\alpha_{h}$   $FWER_{h} / N \leq \alpha_{h}$ 



Convention: Choose *h* to limit  $FWER_h$ assuming family-wise H<sub>0</sub>

#### Issues

- 1. Voxels or regions
- 2. Bonferroni too harsh (insensitive)
  - Unnecessary penalty for sampling resolution (#voxels/volume)
  - Unnecessary penalty for independence



- intrinsic smoothness
  - MRI signals are aquired in k-space (Fourier space); after projection on anatomical space, signals have continuous support
  - diffusion of vasodilatory molecules has extended spatial support
- extrinsic smoothness
  - resampling during preprocessing
  - matched filter theorem
    - $\rightarrow$  deliberate additional smoothing to increase SNR
  - Robustness to between-subject anatomical differences

Acknowledge/estimate dependence Detect effects in smooth landscape, not voxels

- Apply high threshold: identify improbably high peaks
- 2. Apply lower threshold: identify improbably broad peaks
- 3. Total number of regions?



#### Null distribution?

- 1. Simulate null experiments
- 2. Model null experiments



# Use continuous random field theory

• image ≈ discretised continuous random field



Smoothness quantified: resolution elements ('resels')

- similar, but not identical to # independent observations
- computed from spatial derivatives of the residuals

### **Euler characteristic**

- threshold an image at high h# blobs =  $N_h$ 

FWER  $\approx E[N_h]$ = p (blob)



# **Unified Formula**

- General form for expected Euler characteristic
  - $\chi^2$ , *F*, & *t* fields

$$\mathsf{E}[N_h(\Omega)] = \sum_d \mathsf{R}_d(\Omega) \,\rho_d(h)$$

Small volumes: Anatomical atlas, 'functional localisers', orthogonal contrasts, volume around previously reported coordinates...

# $R_d(\Omega)$ : *d*-dimensional Minkowski functional of Ω

- function of dimension, space  $\Omega$  and smoothness:

 $\mathbf{R}_0(\Omega) = N(\Omega)$  Euler characteristic of Ω

 $R_1(\Omega)$  = resel diameter

 $R_2(\Omega)$  = resel surface area

 $R_3(\Omega) = resel volume$ 



#### $\rho_d(\Omega)$ : *d*-dimensional EC density of $Z(\underline{x})$

 function of dimension and threshold, specific for RF type:

E.g. Gaussian RF:

Ω

$$\rho_0(h) = 1 - \Phi(h)$$

 $\rho_1(h) = (4 \ln 2)^{1/2} \exp(-h^2/2) / (2\pi)$ 

 $\rho_2(h) = (4 \ln 2) \exp(-h^2/2) / (2\pi)^{3/2}$ 

 $\rho_3(h) = (4 \ln 2)^{3/2} (h^2 - 1) \exp(-h^2/2) / (2\pi)^2$ 

 $\rho_4(h) = (4 \ln 2)^2 (h^3 - 3h) \exp(-h^2/2) / (2\pi)^{5/2}$ 

### Euler characteristic (EC) for 2D images

$$E[N_h] = R(4\log 2)(2\pi)^{-3/2}h\exp(-0.5h^2)$$

- R = number of resels
- *h* = threshold

Set *h* such that E[NIh] = 0.05

Example: For 100 resels, E[NIh] = 0.049 for a Z threshold of 3.8. That is, the probability of getting one or more blobs where Z is greater than 3.8, is 0.049.



# Spatial extent: similar

#### Voxel, cluster and set level tests



е

u

h

set-level		cluster-level				peak-level							
ρ	C	$\rho_{\rm FWE-com}$	Q <sub>FDR-001</sub> r	Η <sub>Ε</sub>	$\rho_{\rm uncorr}$	$\rho_{\rm FOE-corr}$	Ø <sub>FDR•com</sub>	T	(Z_)	$P_{\rm uncom}$			
0.000	16	0.000	0.000	138	0.000	0.000	0.000	11.04	7.64	0.000	-34	-70	-28
						0.000	0.009	7.31	5.9D	D.000	-44	-74	-24
		0.000	0.000	452	0.000	0.000	0.000	9.82	7.14	0.000	6	16	40
		0.000	0.000	300	0.000	0.000	0.000	9.14	6.84	0.000	44	16	0
						0.041	0.833	5.29	4.64	D.000	38	12	16
		0.000	0.000	173	0.000	0.000	0.009	7.39	5.95	0.000	44	-58	-28
						0.000	0.009	7.35	5.93	0.000	52	-58	-20
						0.002	0.087	6.4Z	5,3B	D.000	50	-66	-24
		0.000	0.000	112	0.000	0.000	0.025	6.93	5.69	0.000	-2	-66	-24
						0.012	0.418	5.73	4.94	D.000	4	-76	-24
						0.014	0.472	5.65	4.89	D.000	z	-86	-28
		0.013	0.374	3	0.257	0.010	0.406	5.77	4.97	0.000	-52	20	-4
		0.000	0.019	20	800.0	0.011	0.406	5.76	4.96	0.000	10	-10	8
		0.008	0.263	5	0.148	0.016	0.472	5.63	4.87	0.000	-8	-16	12
		0.000	0.012	24	0.004	0.016	0.472	5.61	4.86	0.000	44	4	28
						0.035	0.736	5.34	<b>4.6</b> B	D.000	46	6	20
		0.006	0.231	6	0.116	0.018	0.472	5.59	4.84	0.000	-6	-48	-16
		0.026	0.520	1	0.520	0.021	0.538	5.52	4.80	0.000	-6	-54	-16
		0.026	0.520	1	0.520	0.030	0.713	5.40	4.72	0.000	6	-84	-28

#### Statistics: p-values adjusted for search volume

fable shows 3 local maxima more than 8.0mm apart.

Height threshold: $T = 5.21$ , $p = 0.000$ (0.050)	Degrees of freedom = [1.0, 45.0]
Extent threshold: k = 0 voxels, p = 1.000 (0.050)	FWHM = 9.8 10.6 15.6 mm mm mm; 4.9 5.3 3.9 {voxels}
Expected voxels per cluster, <k> = 2.519</k>	Volume: B80432 = 55027 voxels = 472.2 resels
Expected number of clusters, <c> = 0.05</c>	Voxel size: 2.0 2.0 4.0 mm mm mm; (resel = 102.26 voxels)
FWEp: 5.213, FDRp: 6.702, FWEc: 1, FDRc: 20	Page 1

1

# Detect an effect of *unknown* extent & location

There is a multiple testing problem ('voxel' or 'blob' perspective).

More corrections needed as ...



# Further reading

- Friston KJ, Frith CD, Liddle PF, Frackowiak RS. Comparing functional (PET) images: the assessment of significant change. J Cereb Blood Flow Metab. 1991 Jul;11(4):690-9.
- Genovese CR, Lazar NA, Nichols T. Thresholding of statistical maps in functional neuroimaging using the false discovery rate. Neuroimage. 2002 Apr;15(4):870-8.
- Worsley KJ Marrett S Neelin P Vandal AC Friston KJ Evans AC. A unified statistical approach for determining significant signals in images of cerebral activation. Human Brain Mapping 1996;4:58-73.