

Multiple testing

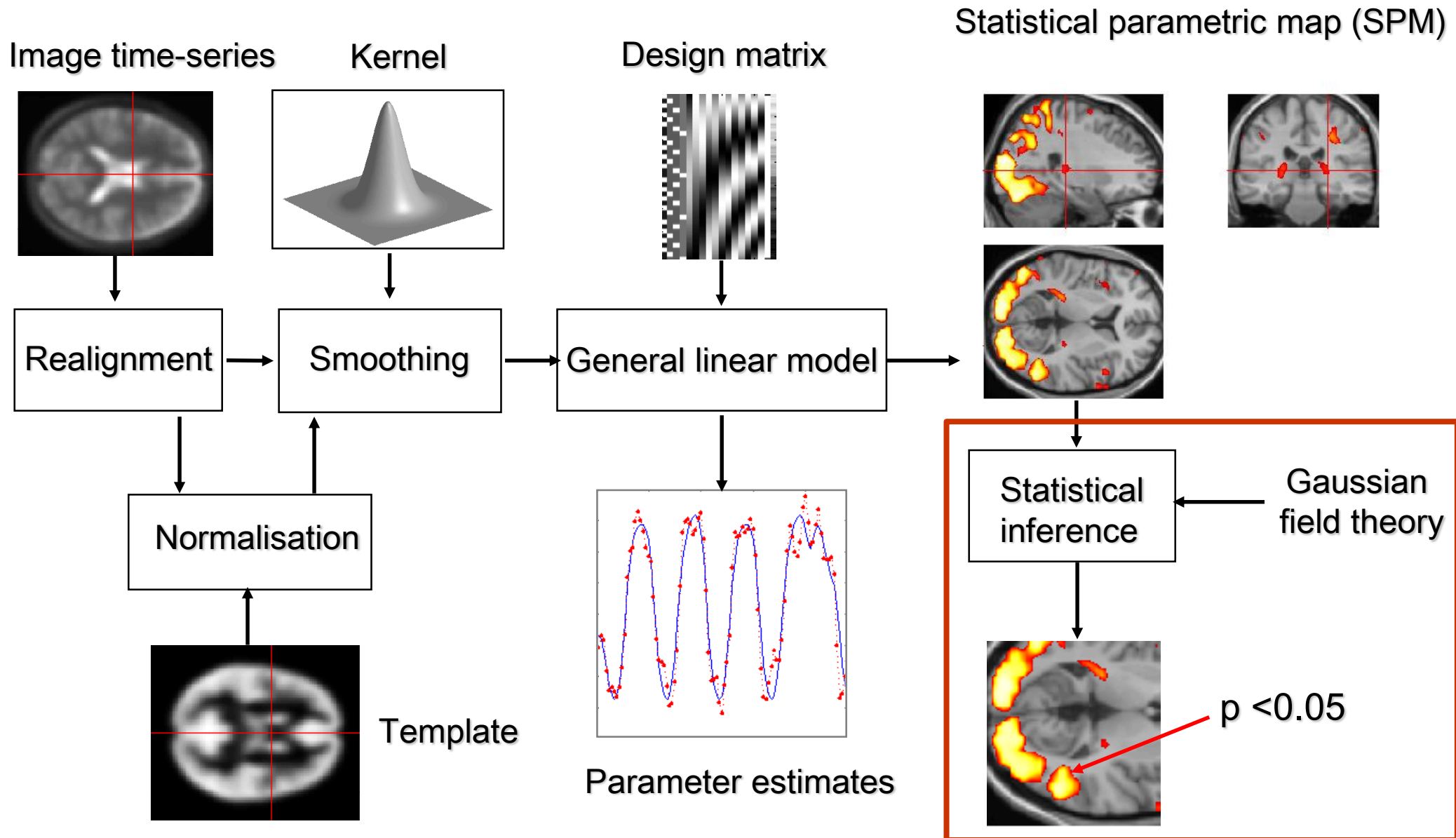
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University of Zurich

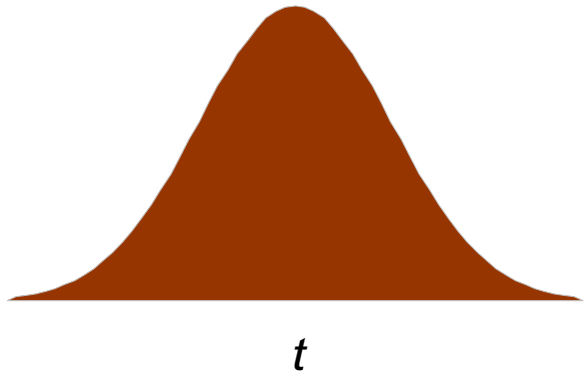
With many thanks for slides & images to:

FIL Methods group

Overview of SPM – Random field theory

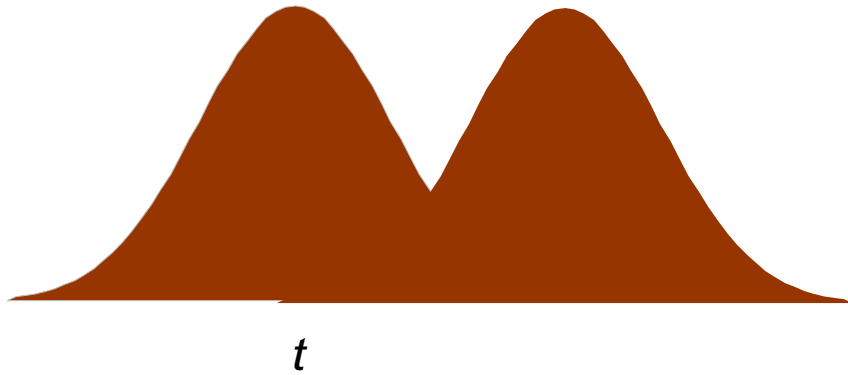


Error at a single voxel



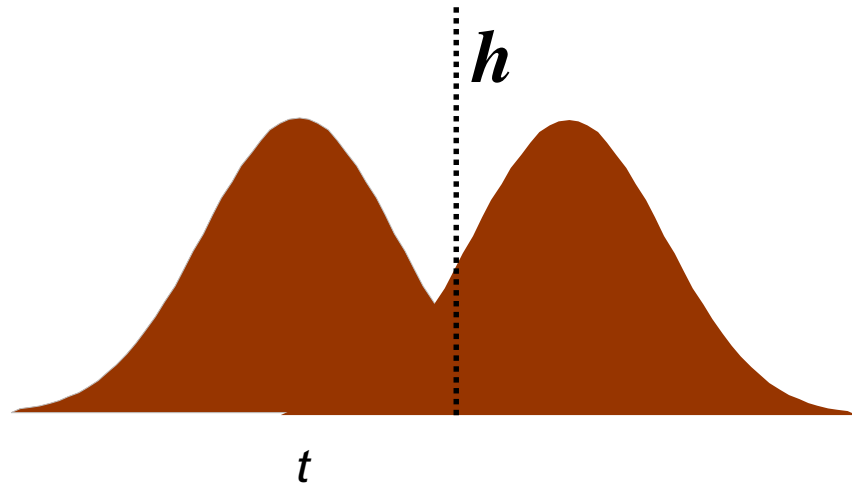
$$t = \frac{\text{contrast of estimated parameters}}{\sqrt{\text{variance estimate}}}$$

Error at a single voxel



$$t = \frac{\text{contrast of estimated parameters}}{\sqrt{\text{variance estimate}}}$$

Error at a single voxel



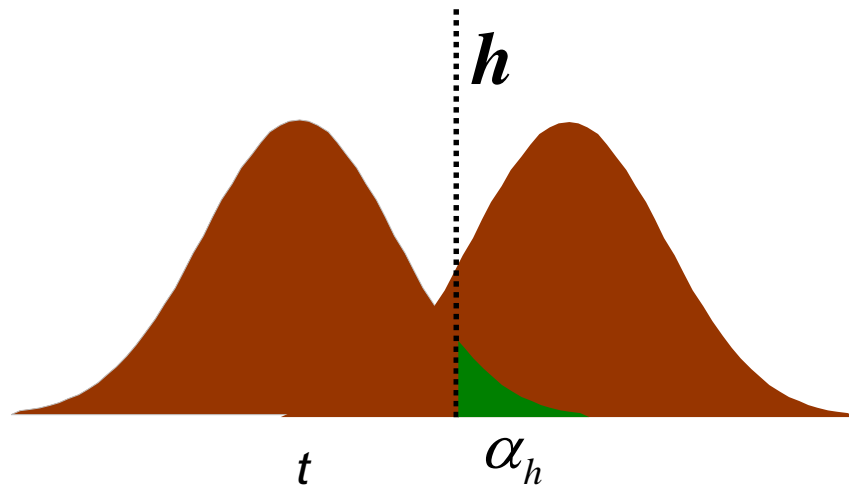
Decision:

H_0 , H_1 : zero/non-zero activation

***contrast of
estimated
parameters***

$$t = \frac{\text{contrast of estimated parameters}}{\sqrt{\text{variance estimate}}}$$

Error at a single voxel



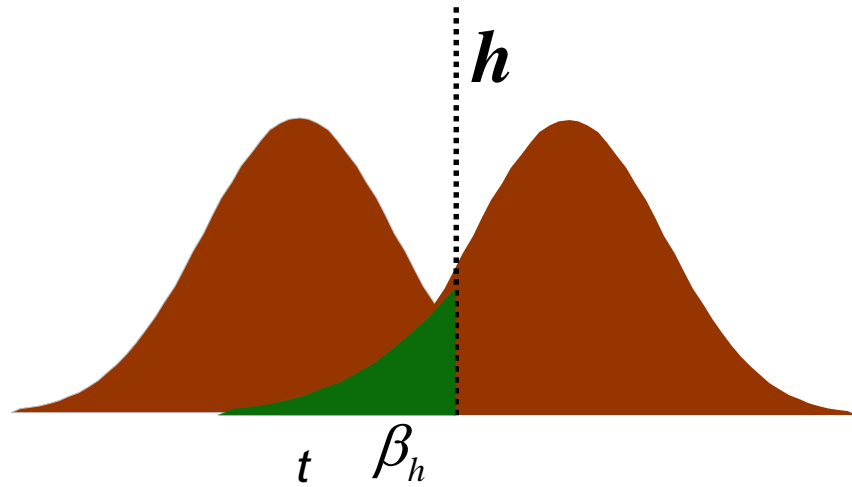
Decision:

H_0 , H_1 : zero/non-zero activation

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$$t = \frac{\text{contrast of estimated parameters}}{\sqrt{\text{variance estimate}}}$$

Error at a single voxel



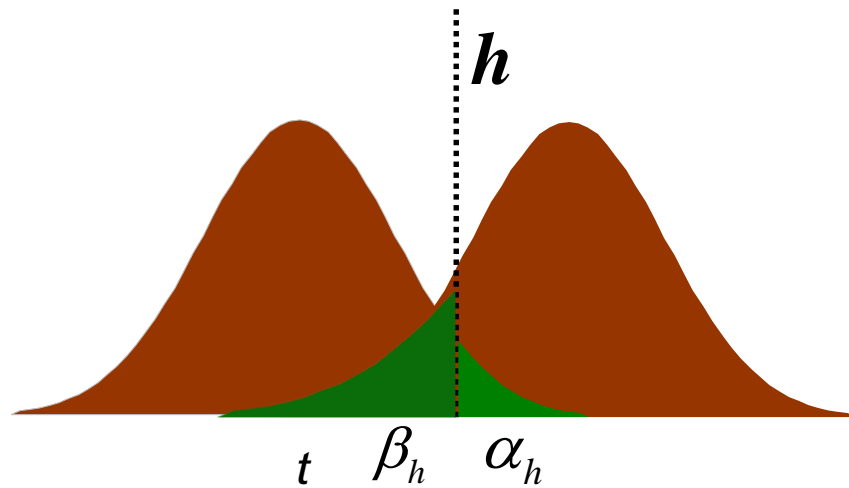
Decision:

H_0 , H_1 : zero/non-zero activation

**contrast of
estimated
parameters**

$$t = \frac{\text{contrast of estimated parameters}}{\sqrt{\text{variance estimate}}}$$

Error at a single voxel



Decision:

H_0 , H_1 : zero/non-zero activation

Decision rule (threshold) h ,
determines related error rates α_h , β_h

Convention: Penalize complexity
Choose h to give acceptable α_h under H_0

**contrast of
estimated
parameters**

$$t = \frac{\text{contrast of estimated parameters}}{\sqrt{\text{variance estimate}}}$$

Types of error

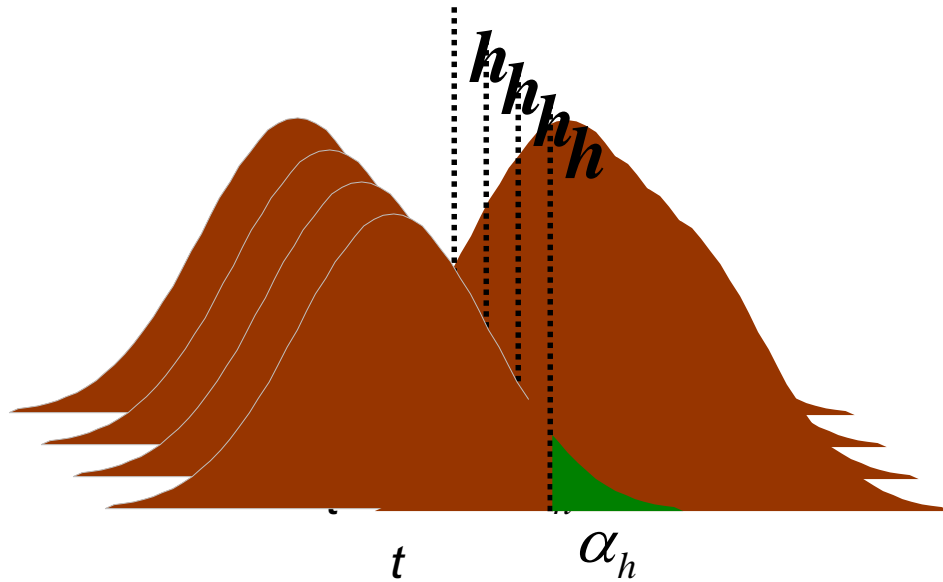
		Reality	
		H ₀	H ₁
Decision	H ₁	False positive (FP) α_h	True positive (TP)
	H ₀	True negative (TN)	False negative (FN) β_h

specificity: $1 - \alpha_h$
= TN / (TN + FP)
= proportion of actual negatives which are correctly identified

sensitivity (power): $1 - \beta_h$
= TP / (TP + FN)
= proportion of actual positives which are correctly identified

Multiple tests

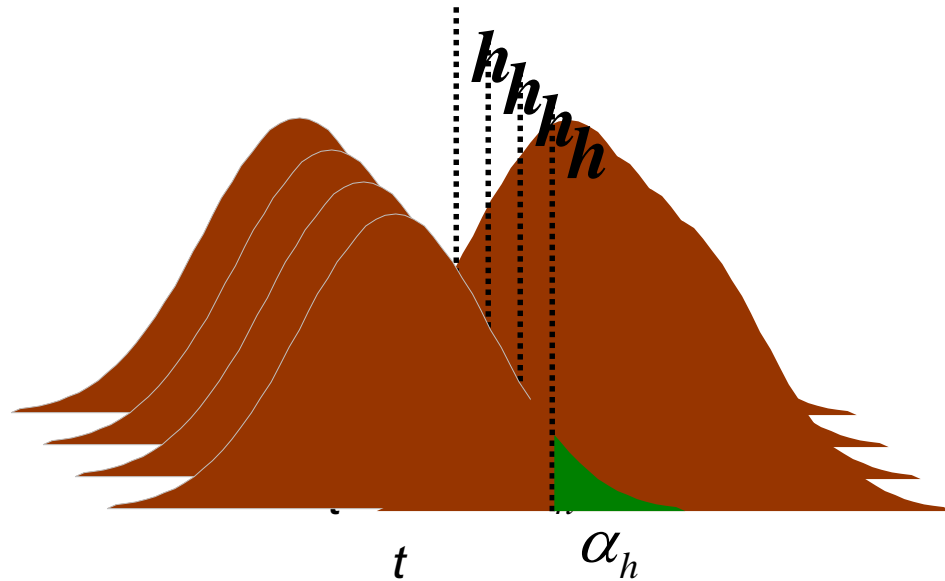
What is the problem?



*contrast of
estimated
parameters*

$$t = \frac{\text{contrast of estimated parameters}}{\sqrt{\text{variance estimate}}}$$

Multiple tests



Penalize each independent opportunity for error.

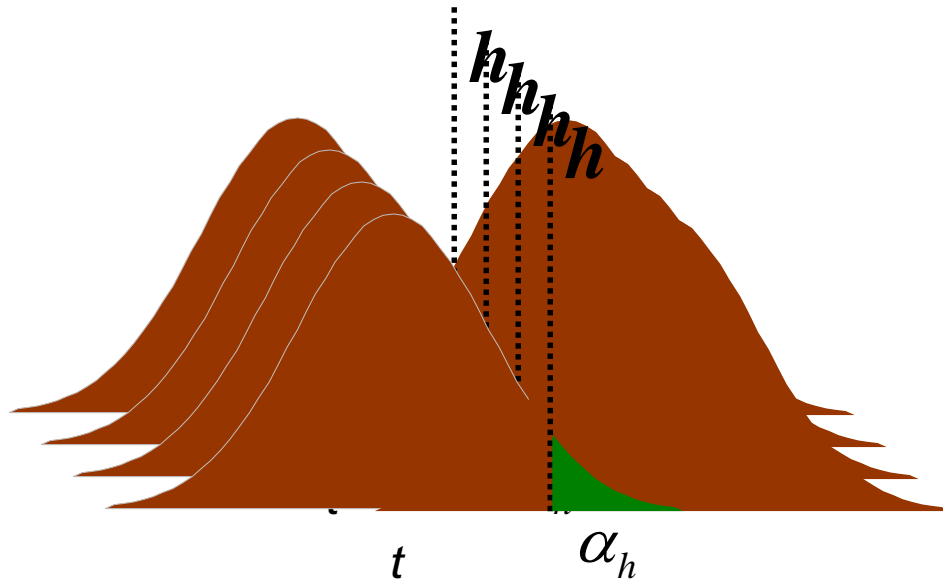
$$p(1 \text{ or more } FP) = FWER_h$$

$$E\left(\frac{FP}{\text{All positives}}\right) = FDR$$

**contrast of
estimated
parameters**

$$t = \frac{\text{contrast of estimated parameters}}{\sqrt{\text{variance estimate}}}$$

Multiple tests



Bonferonni

$$FWER_h \leq N\alpha_h$$
$$FWER_h / N \leq \alpha_h$$

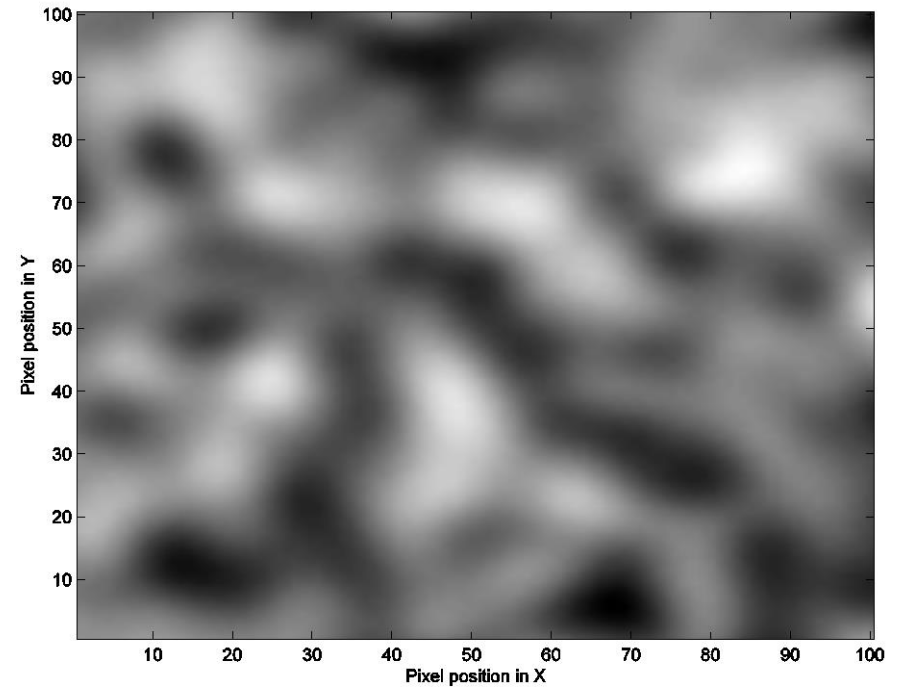
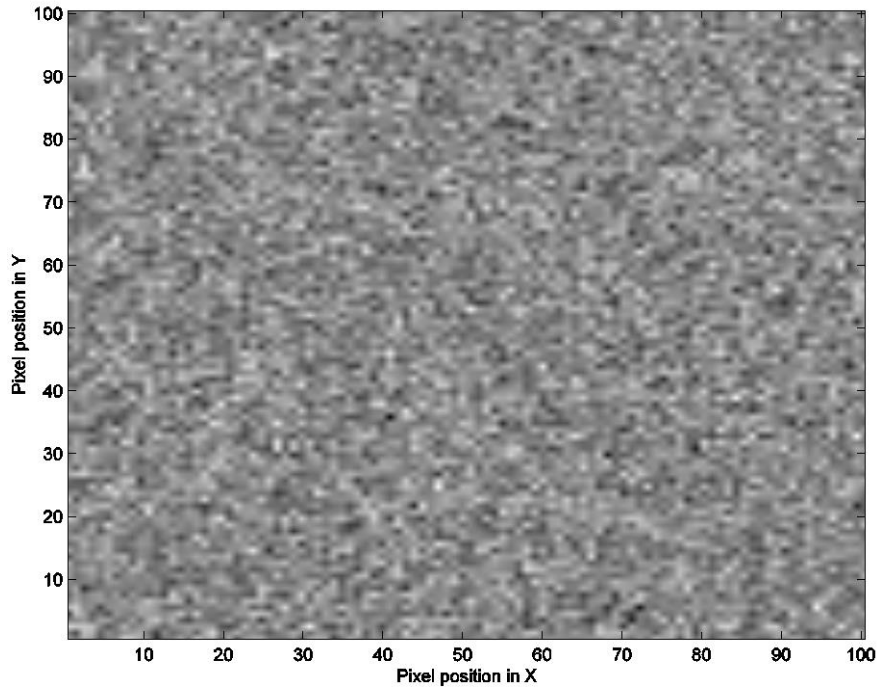
**contrast of
estimated
parameters**

$$t = \frac{\text{contrast of estimated parameters}}{\sqrt{\text{variance estimate}}}$$

Convention: Choose h to limit $FWER_h$
assuming family-wise H_0

Issues

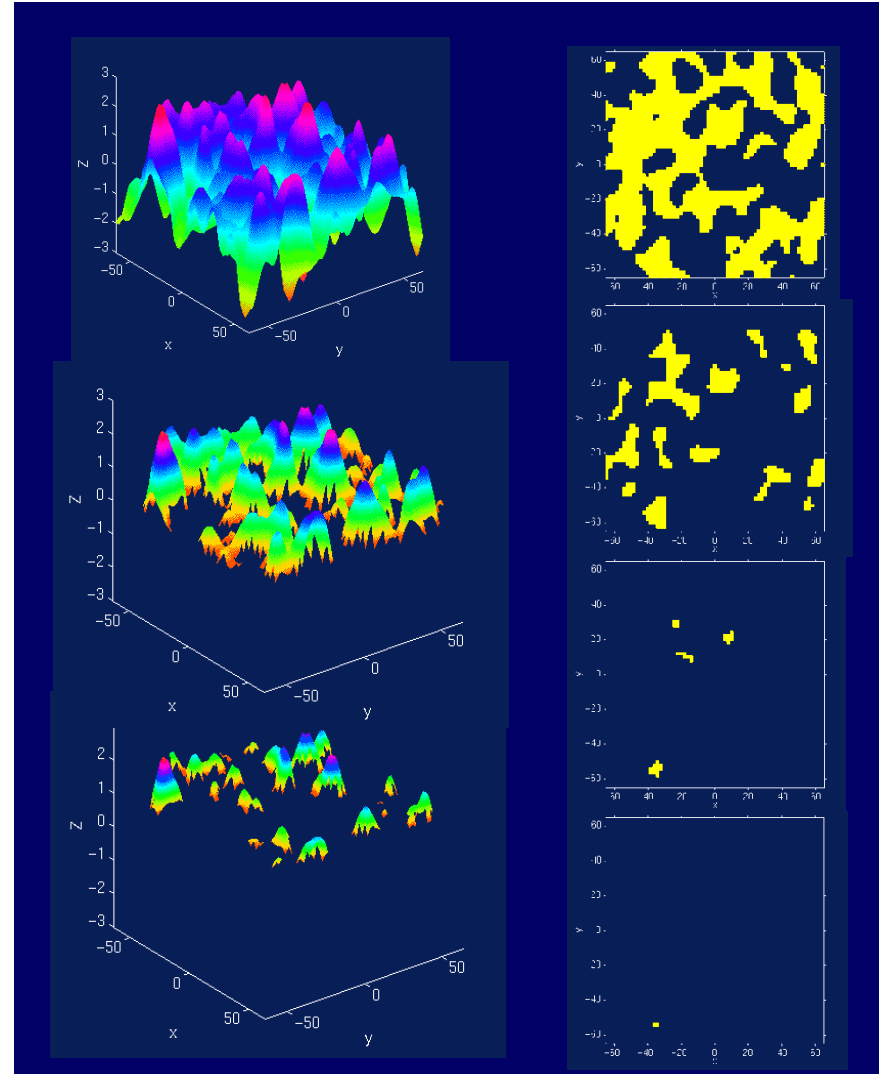
1. Voxels or regions
2. Bonferroni too harsh (insensitive)
 - Unnecessary penalty for sampling resolution (#voxels/volume)
 - Unnecessary penalty for independence



- intrinsic smoothness
 - MRI signals are acquired 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
 - deliberate additional smoothing to increase SNR
 - Robustness to between-subject anatomical differences

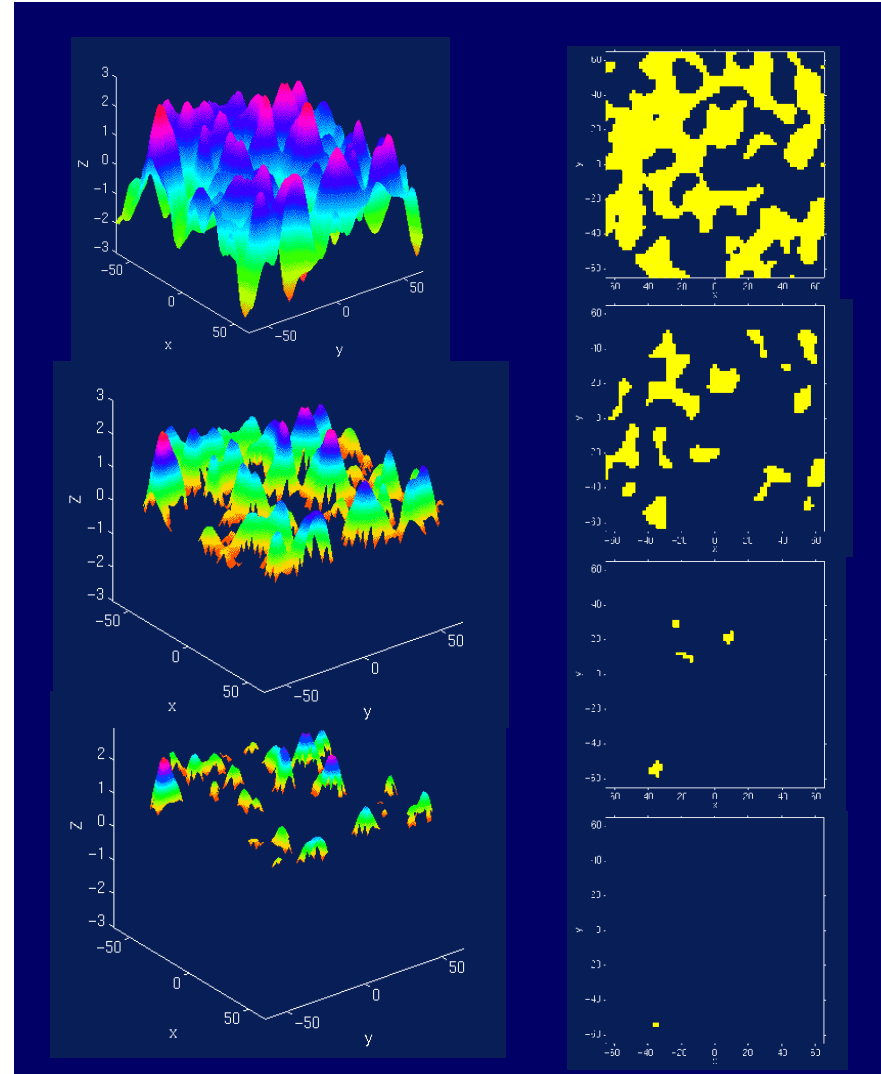
Acknowledge/estimate dependence
Detect effects in smooth landscape, not voxels

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



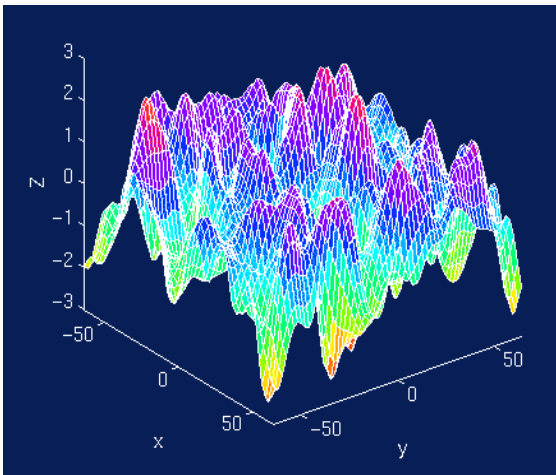
Null distribution?

1. Simulate null experiments
2. Model null experiments

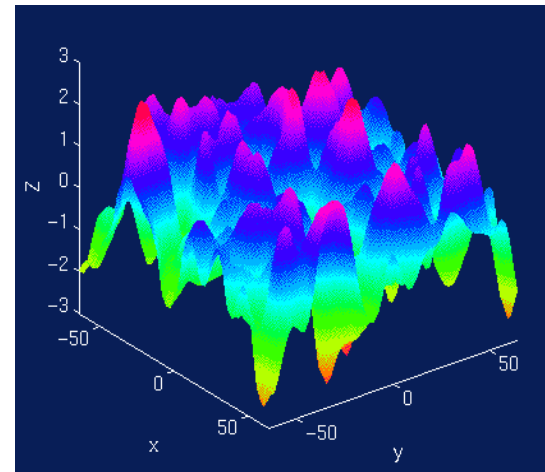


Use continuous random field theory

- image \approx discretised continuous random field



←
Discretisation
("lattice
approximation")



Smoothness quantified: resolution elements ('resels')

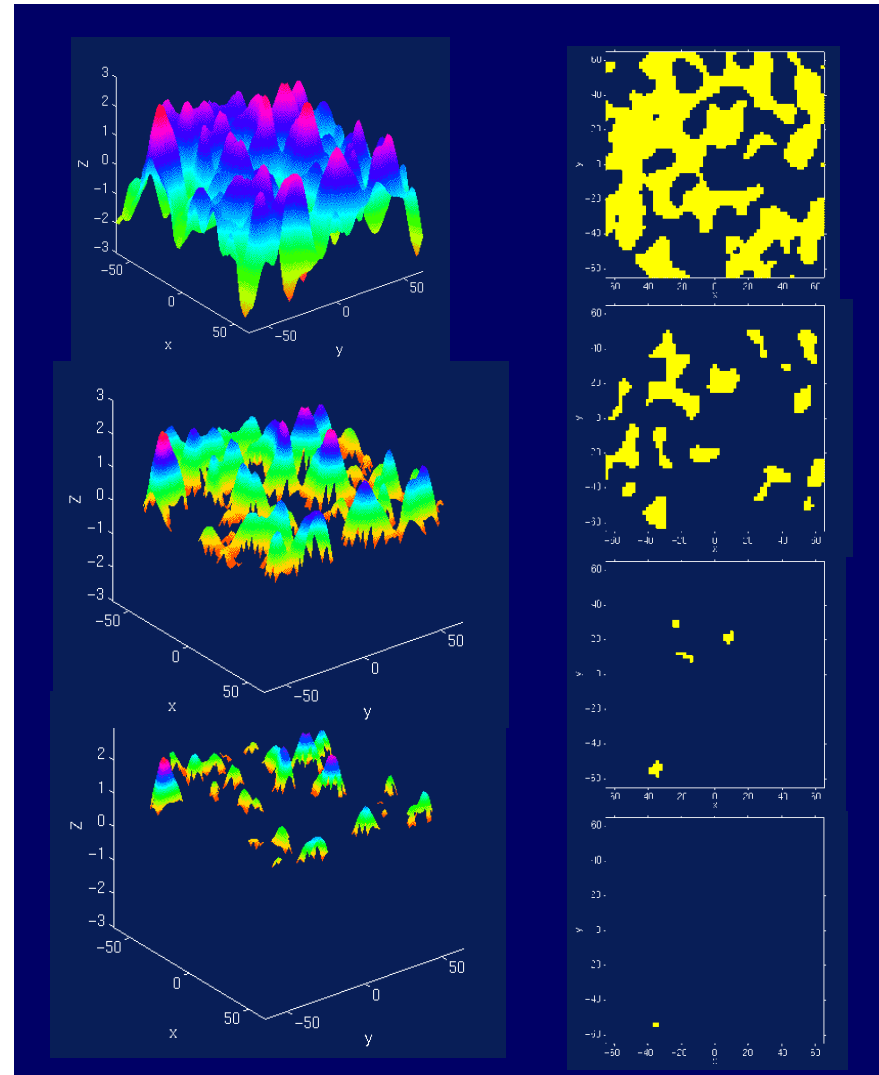
- similar, but not identical to # independent observations
- computed from spatial derivatives of the residuals

Euler characteristic

– threshold an image at high h

blobs = N_h

$$\begin{aligned} \text{FWER} &\approx E [N_h] \\ &= p(\text{blob}) \end{aligned}$$



Unified Formula

- General form for expected Euler characteristic
 - χ^2 , F , & t fields

$$E[N_h(\Omega)] = \sum_d R_d(\Omega) \rho_d(h)$$

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

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

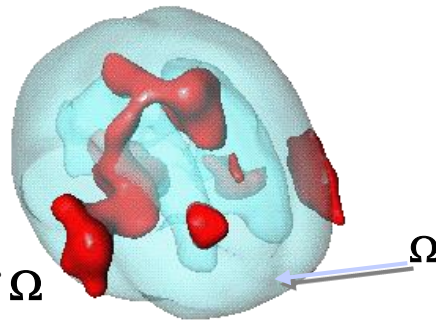
– function of dimension, space Ω and smoothness:

$R_0(\Omega) = N(\Omega)$ 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:

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

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

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

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

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

Euler characteristic (EC) for 2D images

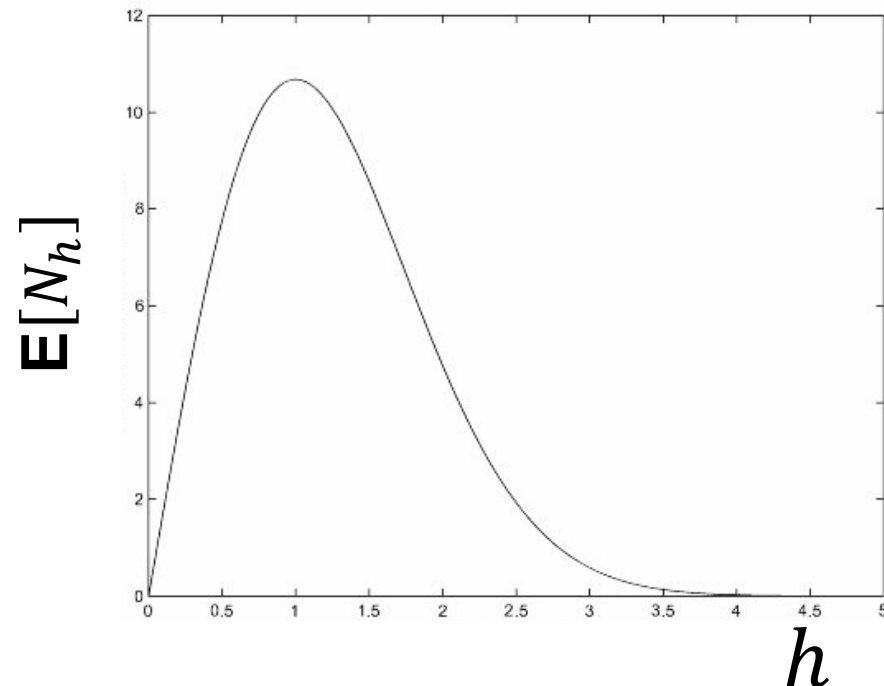
$$E[N_h] = R(4 \log 2)(2\pi)^{-3/2} h \exp(-0.5h^2)$$

R = number of resels

h = threshold

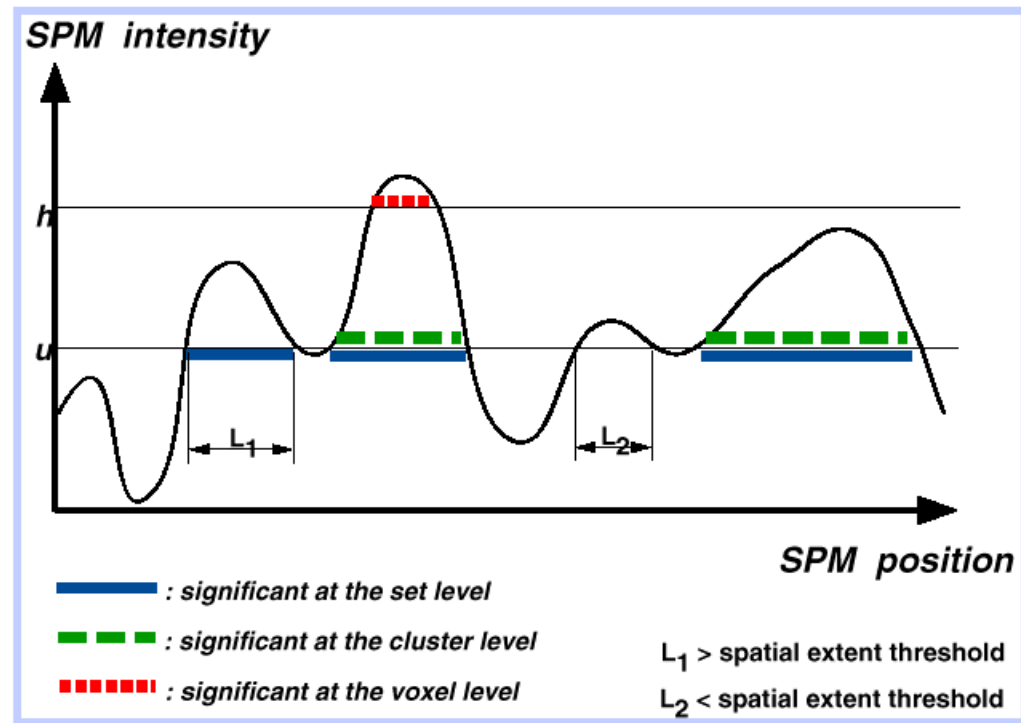
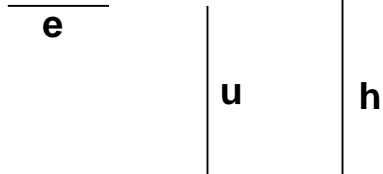
Set h such that $E[N_h] = 0.05$

Example: For 100 resels, $E[N_h] = 0.049$ 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



Statistics: *p*-values adjusted for search volume

set-level		cluster-level				peak-level					mm mm mm		
p	C	$p_{\text{FWE-corr}}$	$q_{\text{FDR-corr}}$	k_E	p_{uncorr}	$p_{\text{FWE-corr}}$	$q_{\text{FDR-corr}}$	T	(Z_{max})	p_{uncorr}			
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	5.90	0.000	-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	0.000	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	6.42	5.38	0.000	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	0.000	4	-78	-24
						0.014	0.472	5.65	4.89	0.000	2	-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	0.008	0.011	0.406	5.76	4.96	0.000	10	-10	8
		0.000	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	4.68	0.000	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

table shows 3 local maxima more than 8.0mm apart

Height threshold: $T = 5.21$, $p = 0.000$ (0.050)
 Extent threshold: $k = 0$ voxels, $p = 1.000$ (0.050)
 Expected voxels per cluster, $\langle k \rangle = 2.519$
 Expected number of clusters, $\langle c \rangle = 0.05$
 FWEp: 5.213, FDRp: 6.702, FWEc: 1, FDRc: 20

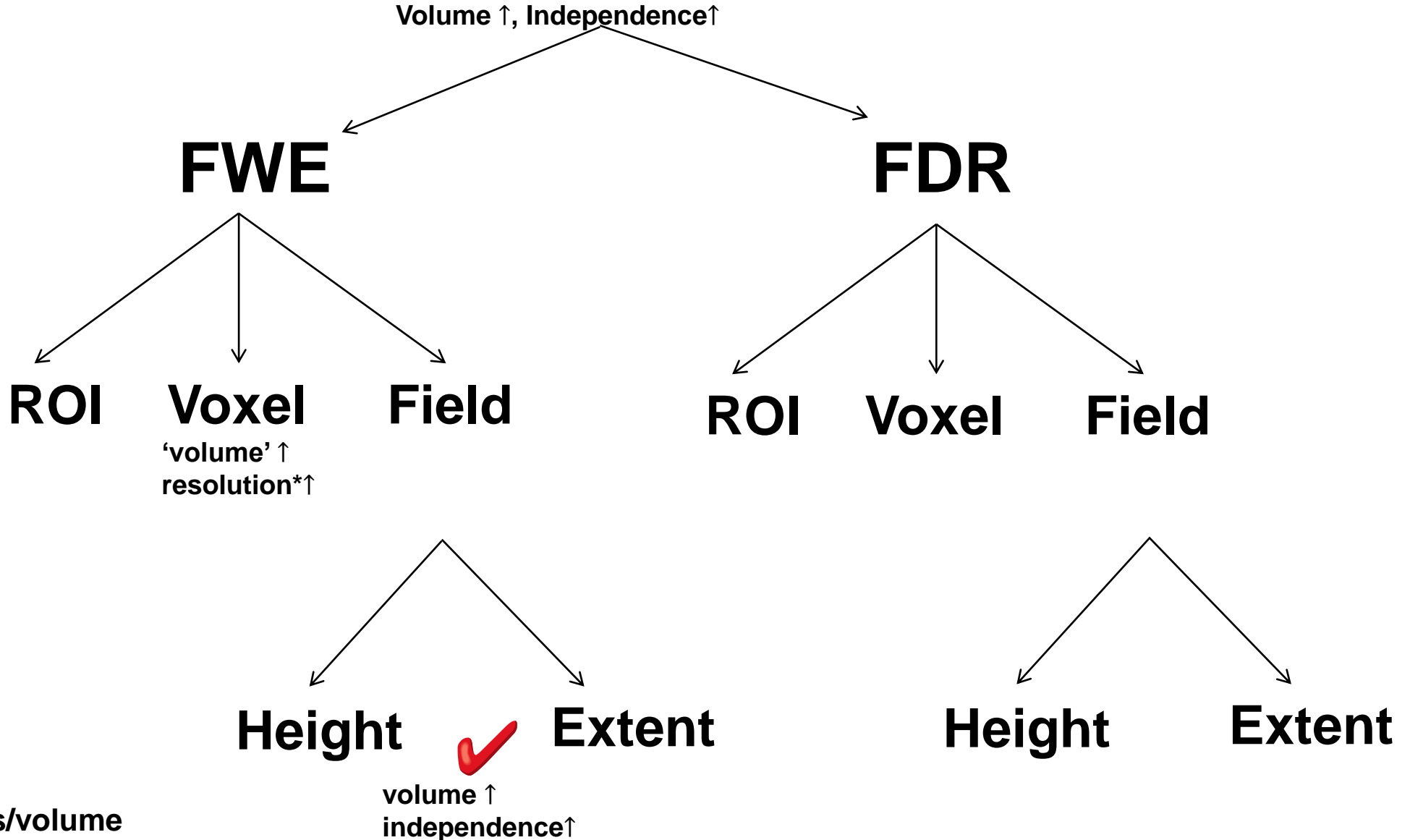
Degrees of freedom = [1.0, 45.0]
 FWHM = 9.8 10.6 15.6 mm mm mm; 4.9 5.3 3.9 (voxels)
 Volume: 880432 = 55027 voxels = 472.2 resels
 Voxel size: 2.0 2.0 4.0 mm mm mm; (resel = 102.26 voxels)
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Detect an effect of *unknown* extent & location

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.