



## Neuroskeptic



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## Ban The Blob?

Common pitfalls in interpreting neuroimaging data





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- You should be able to interpret (in terms of localized function) blobs.



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- You should be able to interpret (in terms of localized function) blobs.
- Blobs are the *first* thing you should look for, and the *final* goal of your analysis

## What is an (fMRI) "Blob"?

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- An area activated by a task?
- An area where task-related activity fits a model?
- An area where task-related activity fits a model well enough to pass an arbitrary threshold.

## **Belgian Beer Lakes**



#### Imagine A Study...

- We sample 100 people in each of various cities around Belgium: 50 men and 50 women.
- Each person completes an alcoholism questionnaire: "BLoB" (Belgian Liking of Beer) scale.
- We want to know:
  - Do Belgian men like drinking beer more or less than women?
  - If so, where in Belgium this difference is seen?

## Would this be the **first way** you'd inspect those results?





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#### Why not?

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- It conceals the raw data you only see (effectively) p-values.
- It imposes the arbitrary p < 0.05 cutoff and censors all nonsignificant points (even if they are p = 0.051).
- We know that blobs are significantly different to some null hypothesis, but we don't know whether each blob is *significantly more significant* than any non-blob point.

### **Erroneous Analysis of Interaction**





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 Nieuwenhuis, Forstmann & Wagenmakers (2011) Nat Neurosci 14 (9) Erroneous analysis of interactions in neuroscience

#### What About The Rest of the Brain?



- Thyreau et al 2012 *Neuroimage*
- N=1326 fMRI study of a face processing task (emotional faces vs. grey circles)
- Multicenter IMAGEN consortium.
- o Results:

## With Enough Subjects, The Whole Brain is A Blob



Fig. 2. Relationship between the effect statistics and the anatomical structure, for different group sizes (100, 200, 500, 1326). Top: Tissue probability as a function of the *t*-statistic. Red is gray-matter, blue is white-matter. Plain color lines are the averages over the ROIs, dashed lines are their 25–75% quantiles. Bottom: Average effect *t*-value as a function of the white/gray probability ratio.

#### So What?

- This is not a *surprising* result it is elementary that t-scores / p-values are dependent on sample size.
- But this means that in using t/p-score based thresholding, we are applying a threshold based on our sample size.
- Should the practical limitation of sample size determine *which areas we think are activated*?

o But also...

#### Blobs are not Representative



 The "voodoo correlations" problem – aka circularity, double dipping, non-independence

#### Blobs are not Representative

 Blobs did not create this problem, but they exacerbate it.

- Vul et al showed the error of treating significantly activated blobs (or, worse, peaks within blobs) as representative of anything (they're not).
- Note that blobs might get more representative as the sample size increases.

#### Avoid Voodoo Blobs





Vul et al 2009 Perspect Psychol Sci

### On the other hand...

 Blobs (thresholding) serve a very important purpose.

 Whole-brain corrected blobs (FDR or FWE corrected) are evidence that 'something is going on'.

 To adopt ar uncorrected (Bennet et



The fish that launched a thousand 'skeptics'

## What we need is Diversity

 Blobs should not be the Alpha and the Omega of neuroimaging analysis. They should be one part of a comprehensive approach.

 Look at the unthresholded statistical parametric maps *alongside* the thresholded ones.

• E.g. In FSL you can find these in the *stats*/ directory of FEAT output for fMRI.

# Post-Blob Visualization? Or not quite?



 Allen, Erhardt, Calhoun (2012) Neuron Data visualization in the neurosciences: overcoming the curse of dimensionality