

Simplified power and sample size calculations using prevalence & magnitude of active peaks.

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Introduction

- There is increasing concern about statistical power in neuroscience research:
 - Low power decreases the chance of detecting a true effect
 - Low power reduces the chance that a statistically significant result indicates a true effect (Ioannidis, 2005).
- ⇒ **A power analysis is a critical component of any study**
- Power analyses for fMRI are difficult: need to specify magnitude, spatial extent and location of a hypothesized effect.
- We present a simple way to characterize the spatial signal in a fMRI study, and a direct way to estimate power based on an existing pilot study.
- We must estimate (or have prior knowledge of):
 - the volume of the brain that is activated
 - the average effect size in activated brain regions
- This allows the calculation of power for given sample size, brain volume and smoothness.

Methods

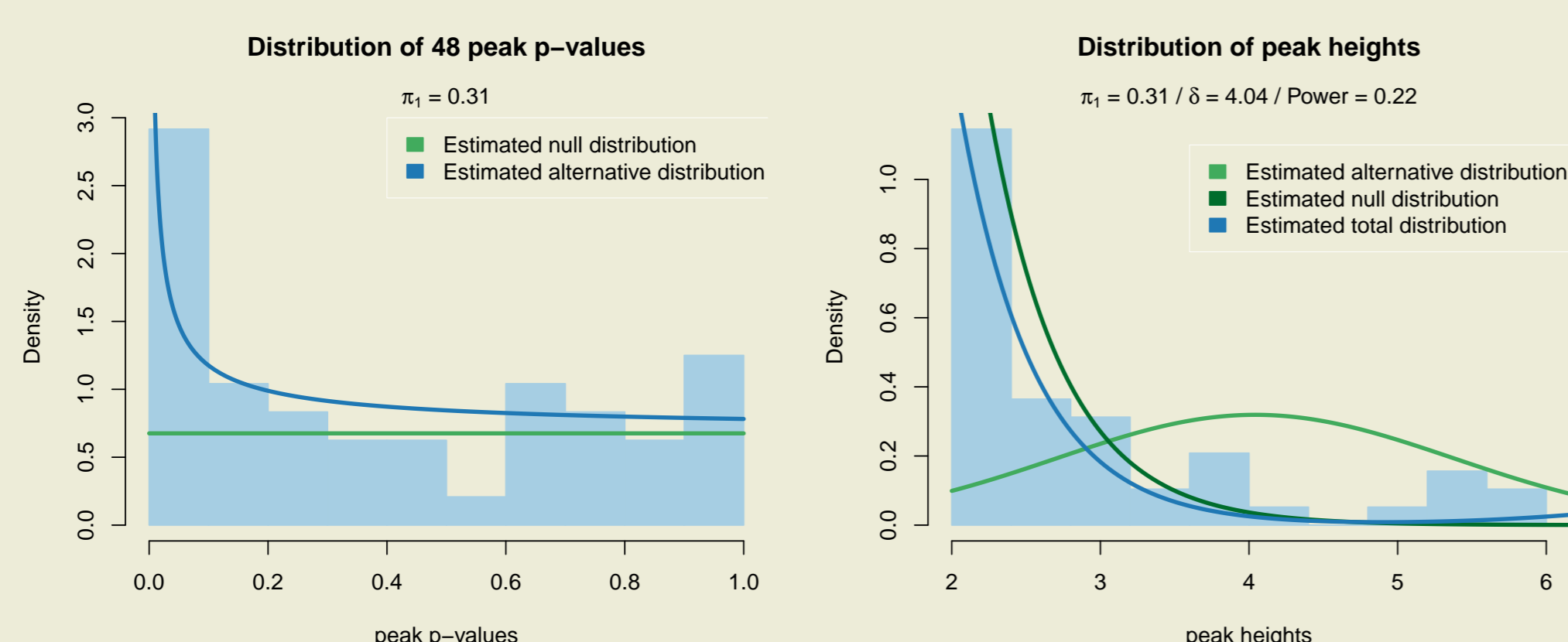
- We start from peak p -values in a group level analysis. We estimate π_1 , proportion of peak p -values that are non-null, as described in Durnez, Moerkerke & Nichols (2014).
- We assume that the null distribution of peak values is an exponential distribution (Worsley, 2007).
- We assume that the alternative distribution of peak values is a truncated normal distribution (truncated at excursion threshold u).
- Therefore, the distribution of peak values can be written as a mixture:

$$f(x|\pi_0, \mu_1, \sigma_1, u) = (1 - \pi_1)u \exp(-u(x - u)) + \pi_1 \frac{\frac{1}{\sigma_1} \varphi\left(\frac{x - \mu_1}{\sigma_1}\right)}{1 - \Phi\left(\frac{u - \mu_1}{\sigma_1}\right)}$$

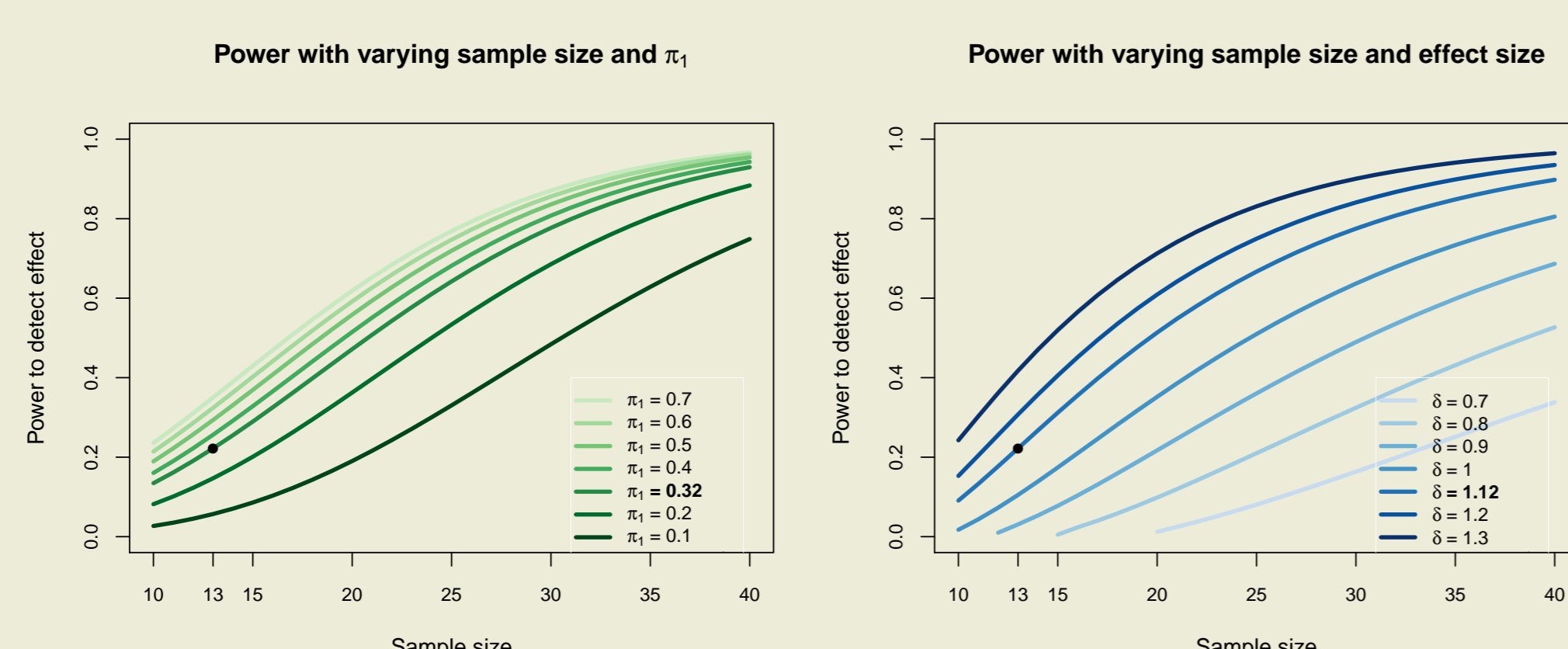
- μ_1 and σ_1 can be estimated using maximum likelihood, where μ_1 is the expected peak height in activated regions.
- Power can be estimated for a given threshold t as $P(T > t|H_a)$ with T the T -statistic of the peak.

Example

- Data from Seurinck et al. (2011). The original study was analyzed voxelwise with FWER-control at the 5% significance level. We reanalyzed the data peakwise with FDR-control at the 5% significance level.
- Sample size: 13
- Step 1: compute prevalence of activation
- Step 2: estimate truncated distributions

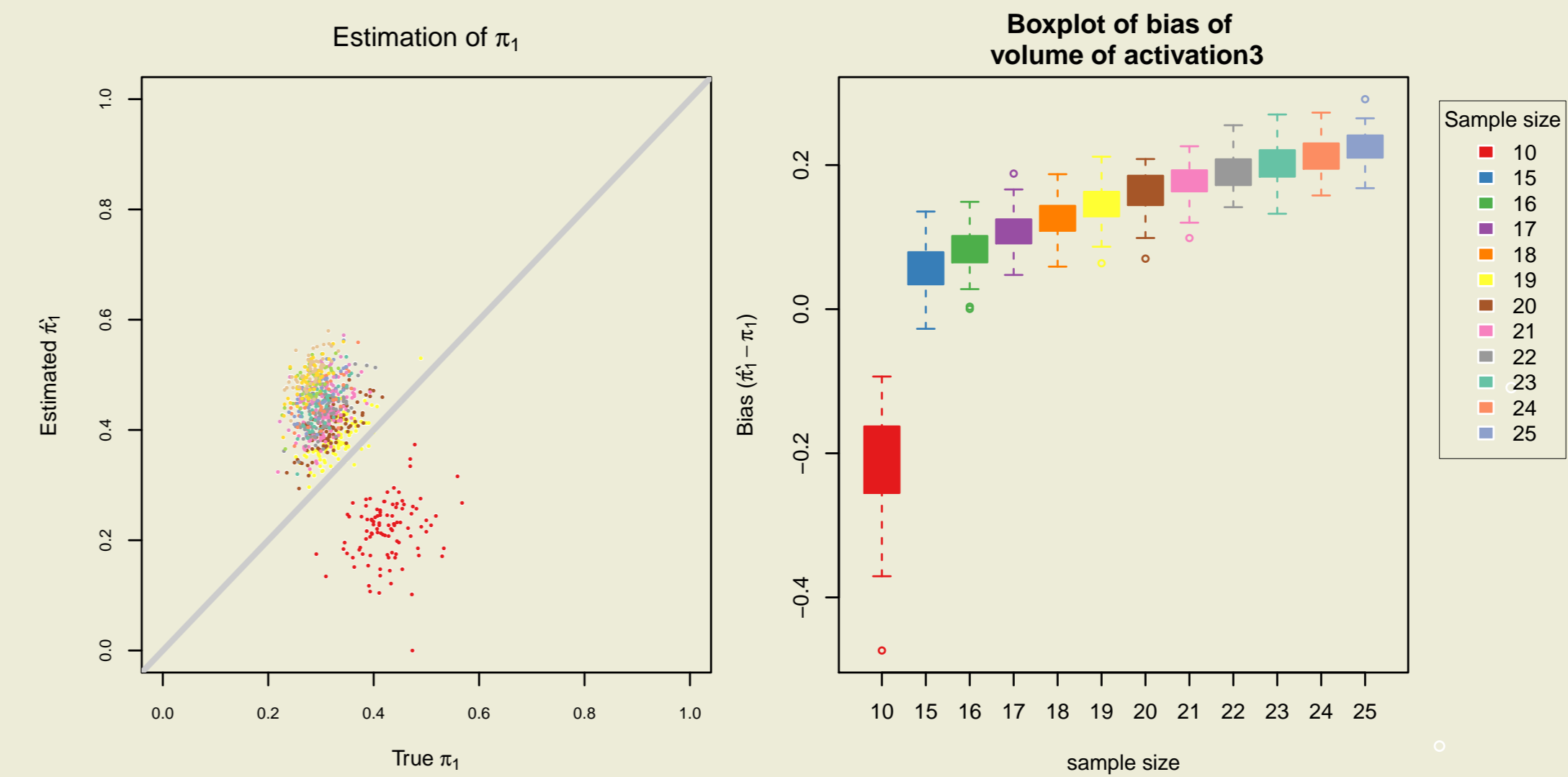


- Expected peak height under activation $\mu_1 = \delta = 4.04$
- Effect size $\mu/\sigma = \delta/\sqrt{n} = 1.13$
- Step 3: estimate power for given sample size, estimated peak height under activation and prevalence of activation (π_1)

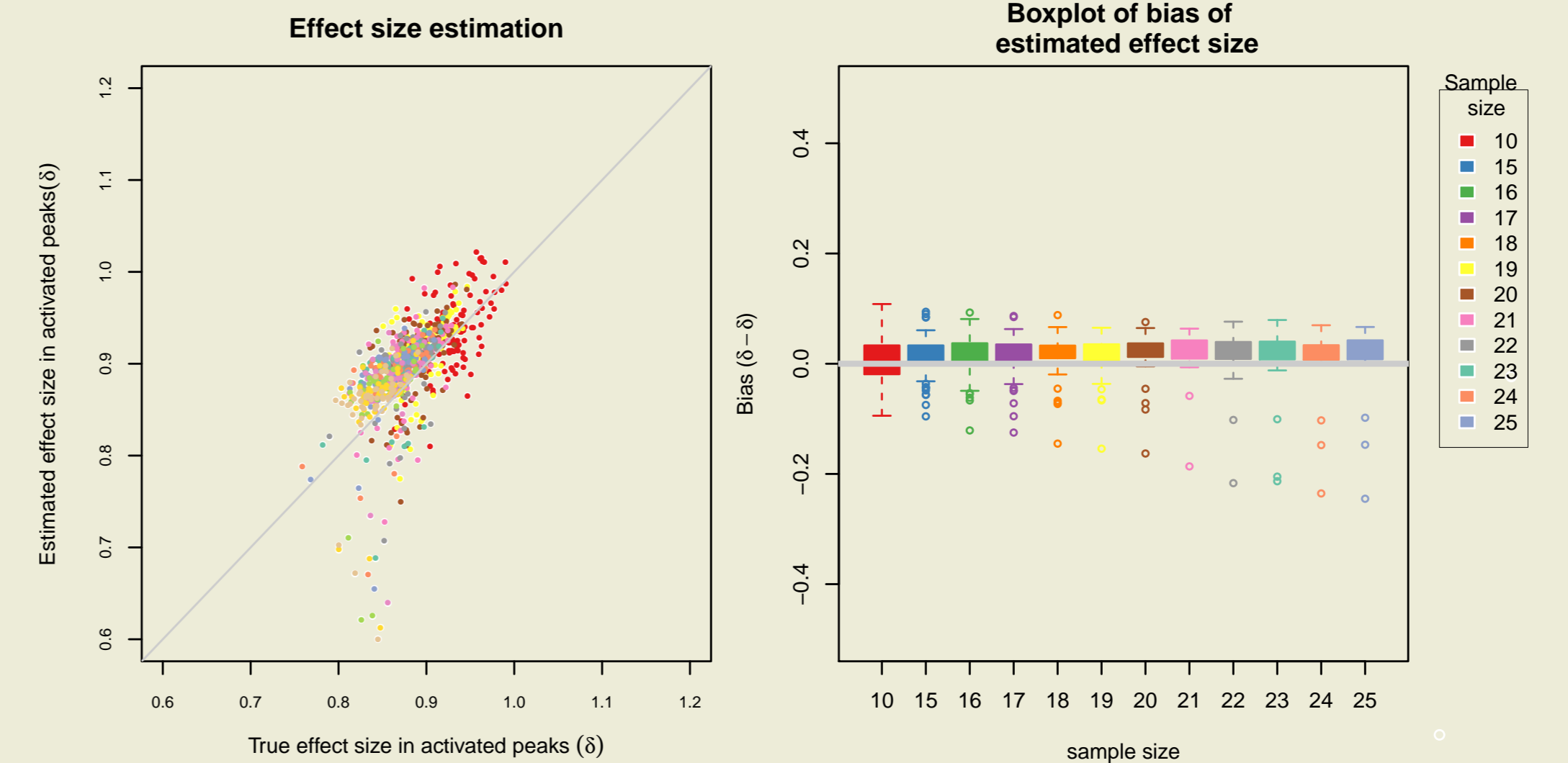


Simulations

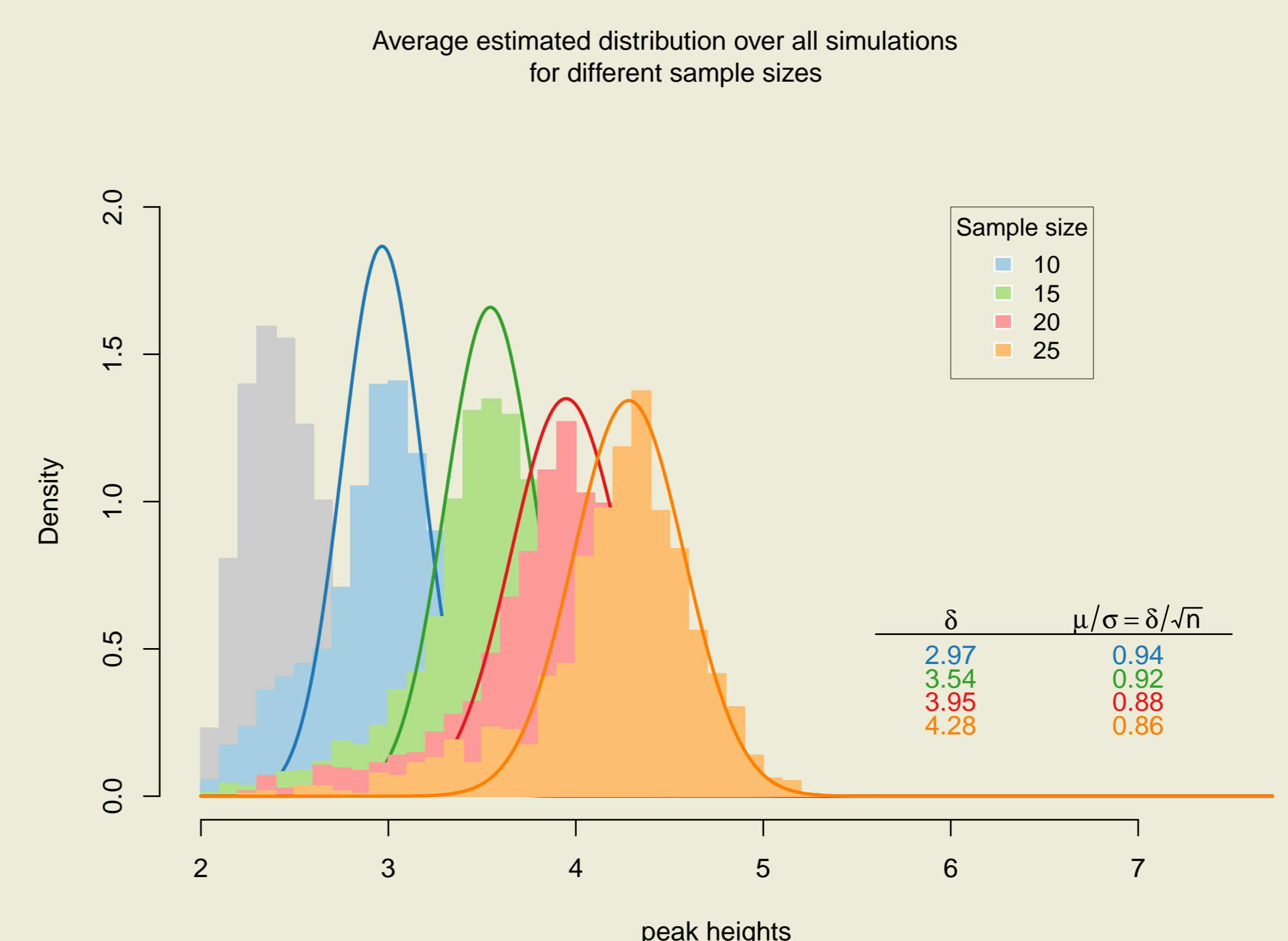
- 100 full-brain datasets with smooth Gaussian noise (3 voxels) superimposed with activation (3 % BOLD change) in 4 foci (3% of total brain volume)
- Prevalence of activation:** For small sample sizes: very conservative estimates (close to 0).



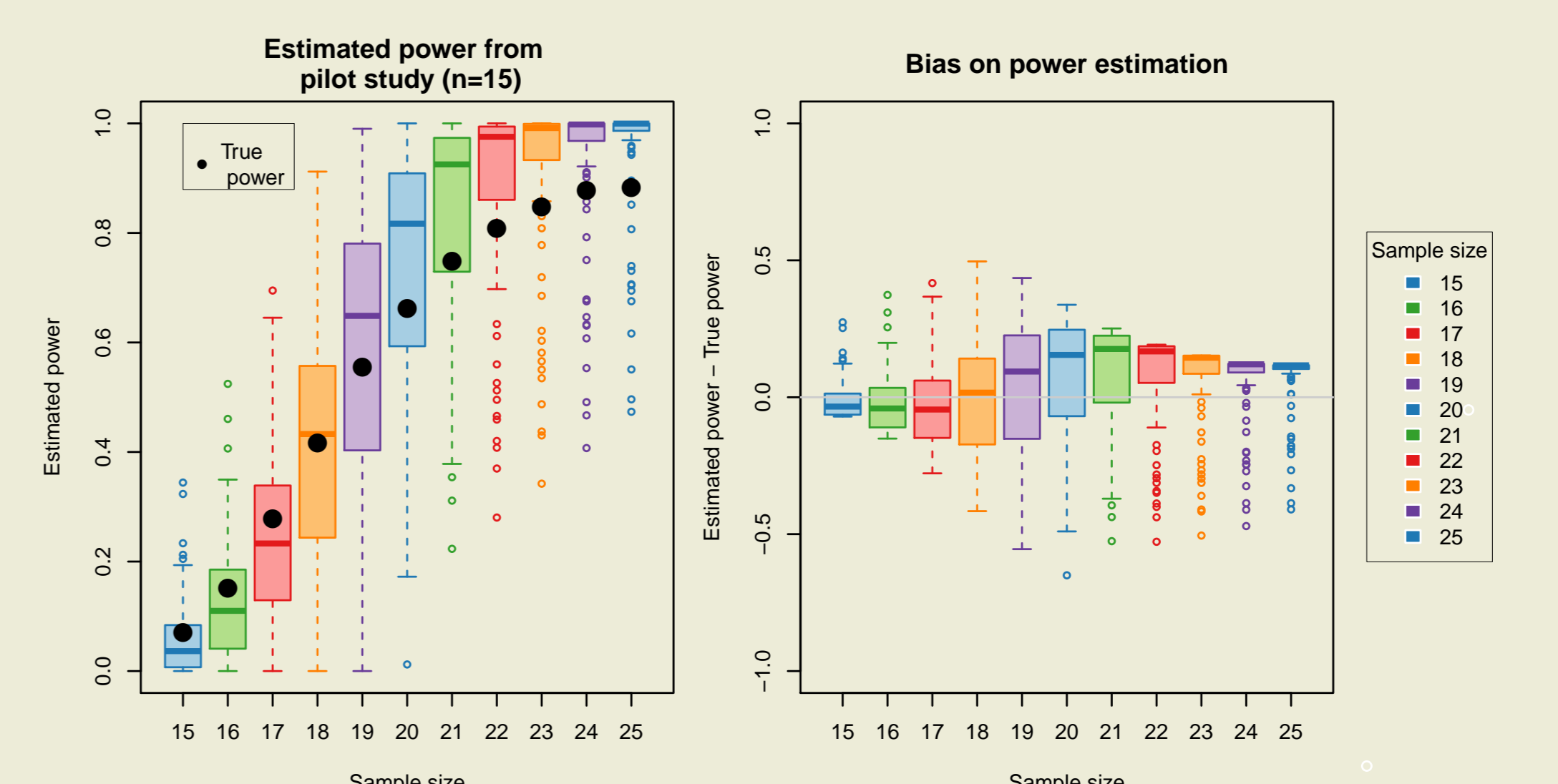
- Effect size estimation:** Rather conservative estimates.



- Alternative distribution:** Constant effect size (μ/σ) over different sample sizes to be used in power analyses.



- Power calculations:** Good estimates for realistic values of power with FWER control at the 5% significance level.



References and acknowledgements

- Durnez, Moerkerke, Nichols (2014). NeuroImage, 84.
 Ioannidis (2005). PLOS Medicine, 2:6.
 Seurinck, de Lange, Achten, Vingerhoets (2011). Journal of Cognitive Neuroscience, 23:6.
 Worsley (2006). Random Field Theory in Friston, Ashburner, Kiebel, Nichols and Penny, Statistical Parametric Mapping. Elsevier, London
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