

Bayes Factors for time-resolved neuroimaging data

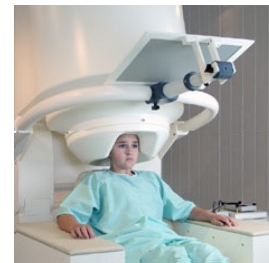
Lina Teichmann

Laboratory of Brain and Cognition, NIMH

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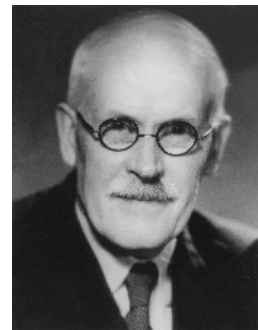
Time-resolved neuroimaging

- M/EEG allows us to study regional and large-scale dynamics of brain activity
- Many different analysis techniques
 - Event-related analyses
 - Time-domain
 - Frequency-domain
 - Source analyses
 - Oscillations
 - Connectivity
 - ...
- Statistics important to draw meaningful conclusions



Using Bayes Factors for M/EEG results

- Based on Harold Jeffreys perspectives and philosophy of scientific learning (*cf. Ly et al., 2016*)
 - Relationship between probability & inference: How probable is one hypothesis in comparison to another when considering the data? (*Jeffreys, 1935, 1939*)
- Practical advantages for M/EEG analyses
 1. Allow us to directly compare two hypotheses
 2. Measure of strength that is interpretable
 3. Enables us to collect data iteratively



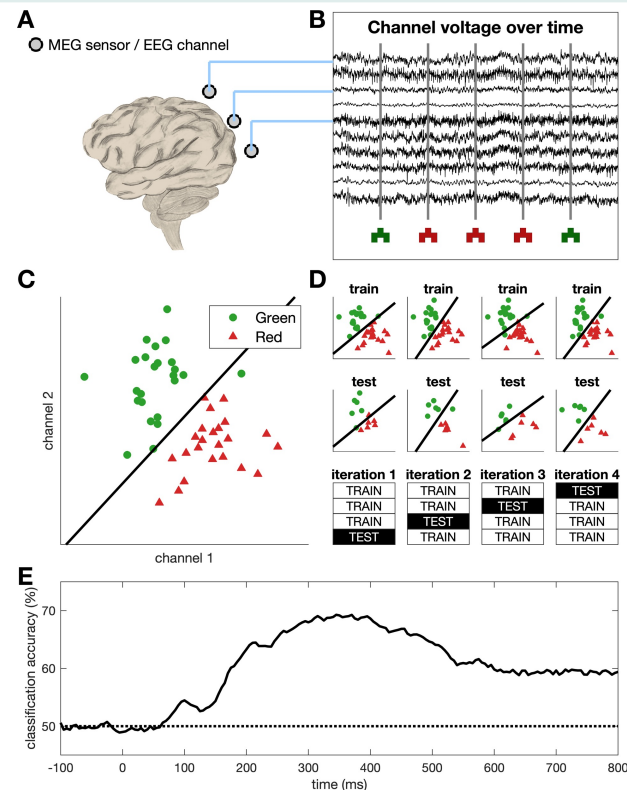
Sir Harold Jeffreys

Time-resolved classification analysis

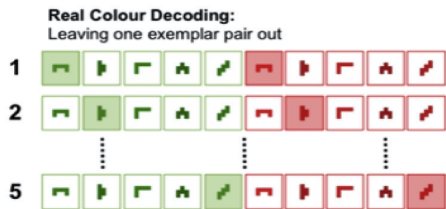
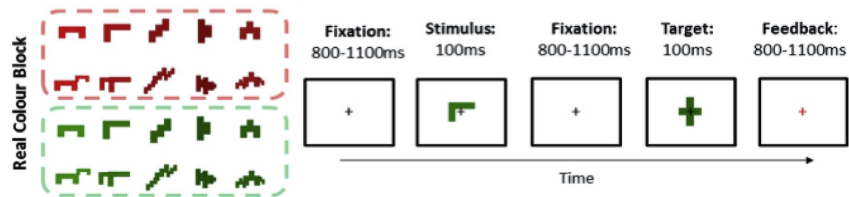
- Example: Colour processing



- Activation- vs information-based framework (*cf. Hebart & Baker, 2018*)
 - Activation: Does red evoke a stronger/weaker signal than green? (univariate)
 - Information: Is there information about green/red in the signal? (multivariate)
- Classification analysis hypotheses
 - H_0 : mean decoding equals chance decoding
 - H_a : mean decoding is larger than chance decoding
- **Bayes Factors**: the probability of one hypothesis versus another given the data

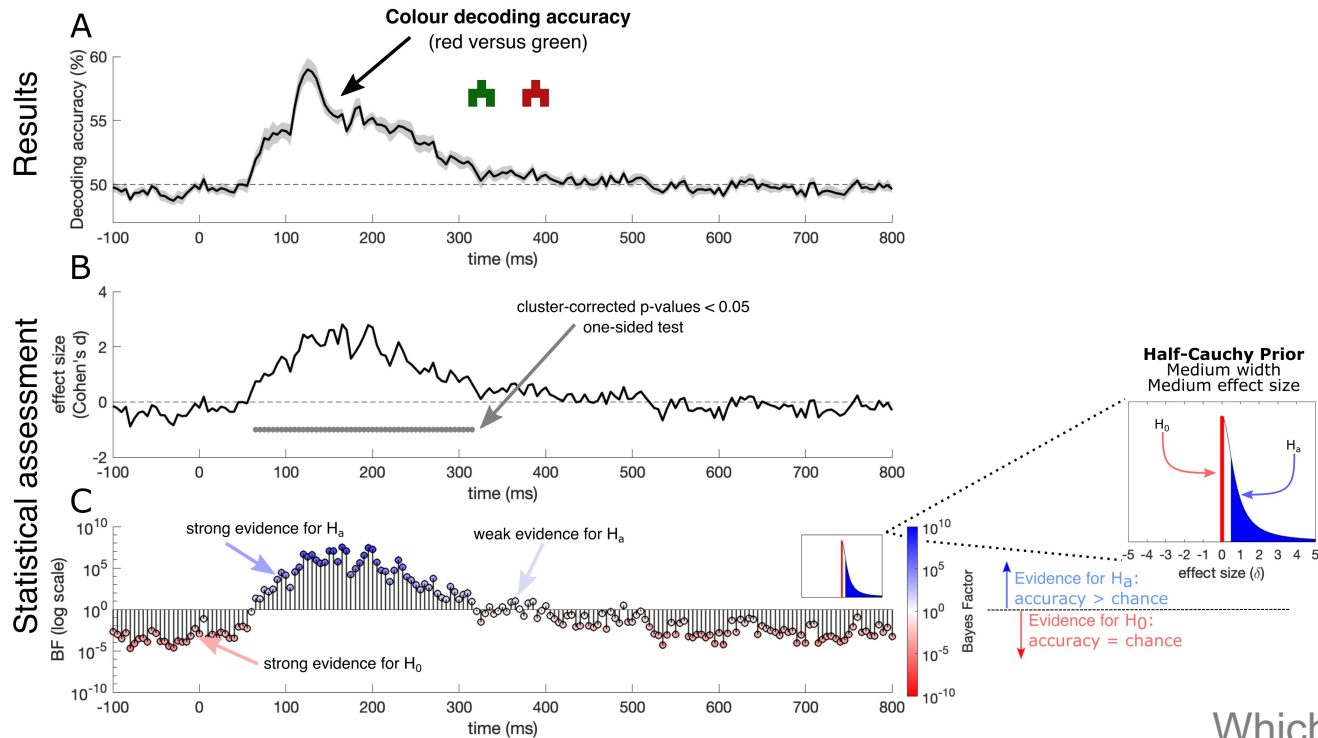


Example dataset



- 18 participants
- 1600 trials
- Epoched -100 to 800ms
- 200 Hz resolution
- Target-detection task (targets excluded from analysis)
- 5-fold cross-validation
- Originally, permutation tests and cluster-corrected p-values

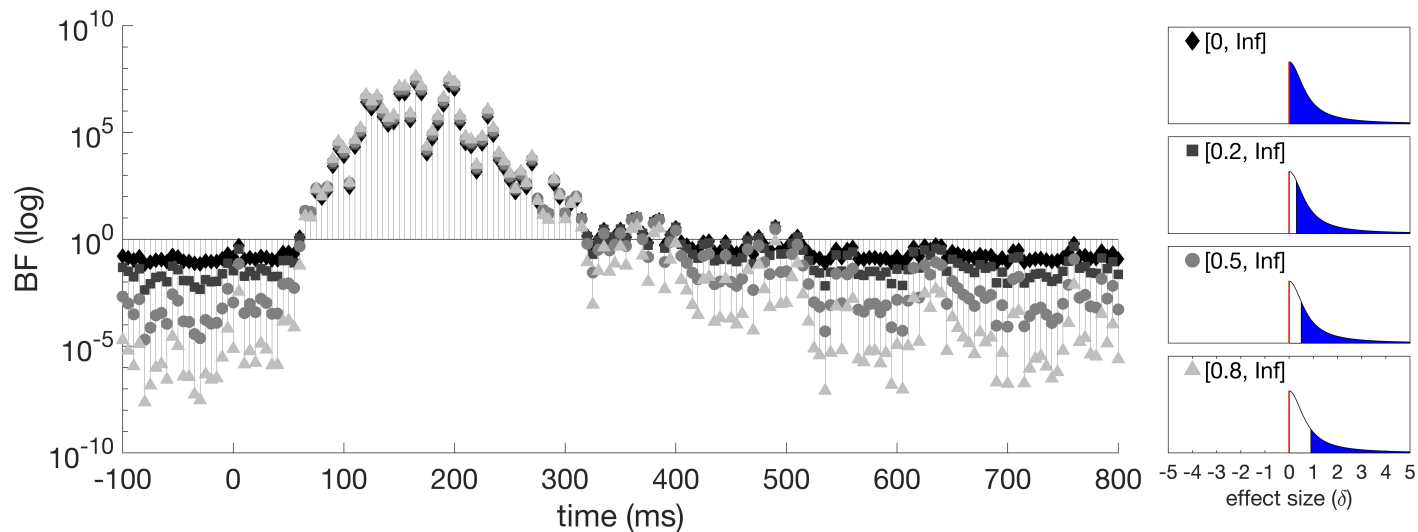
Example dataset results



Which parameters make the most sense?

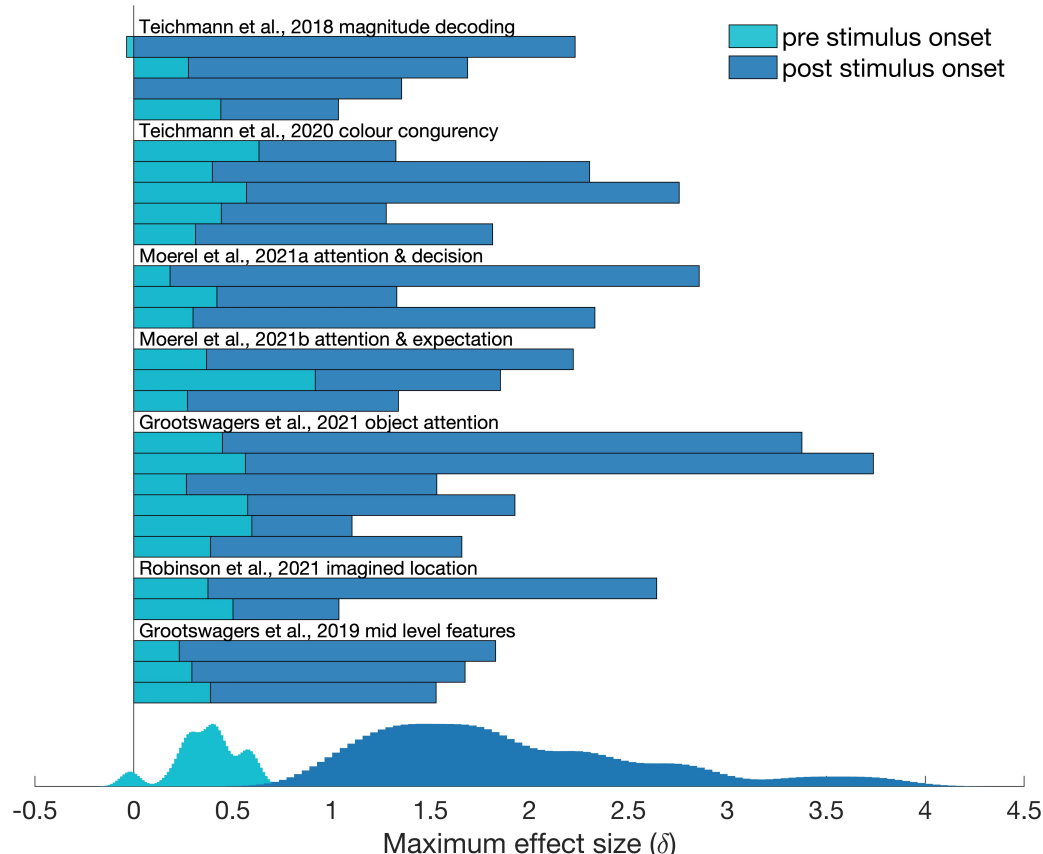
Adjustment of prior range (null interval)

- Increasing the lower bound means allowing small effects under H_0
- Observed chance is often different than theoretical chance



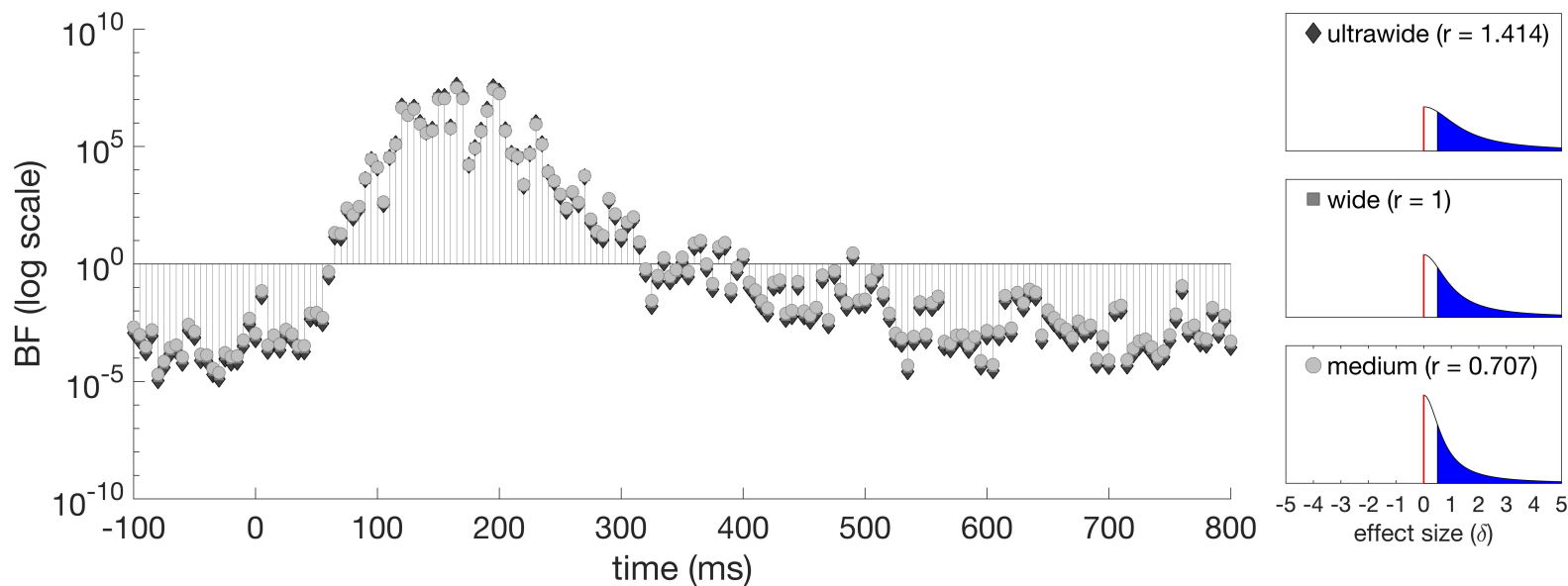
What lower bound makes sense?

- Baseline period: What values can be expected “by chance”



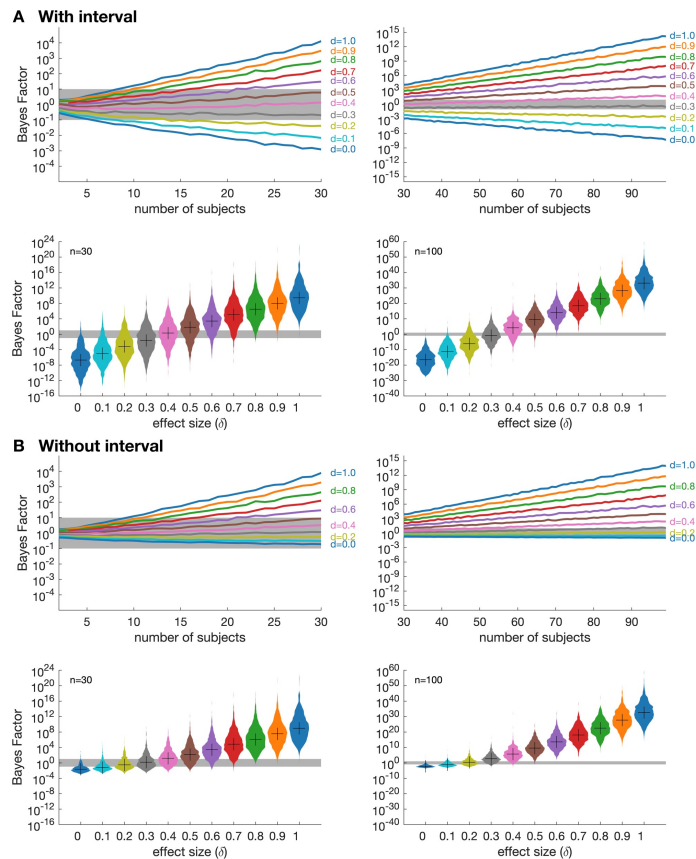
Variation of prior width

- Widths capture expected effect sizes
- Changing prior width has no influence on our data (effect sizes large)



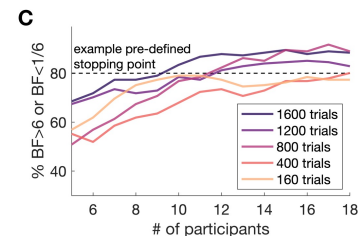
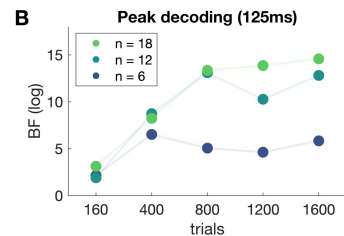
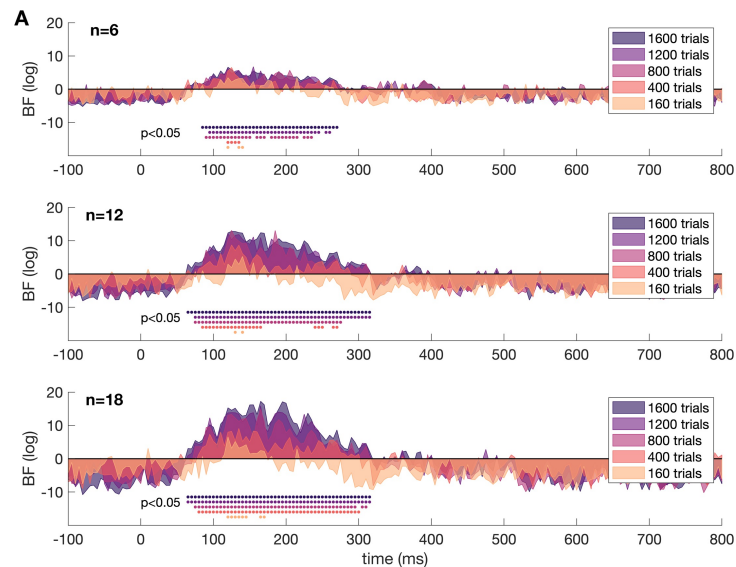
Sensitivity to small effects given parameters

- Simulated classification results for different sample sizes and varied the effect sizes
 - Even if effect is small (and lies within the pre-defined null interval) we find evidence for H_a after a while if the effect is consistently there
 - Large number of observations does not automatically lead to conclusive evidence (if the effect is truly in between the hypotheses)
 - Interval is critical to find evidence for the null



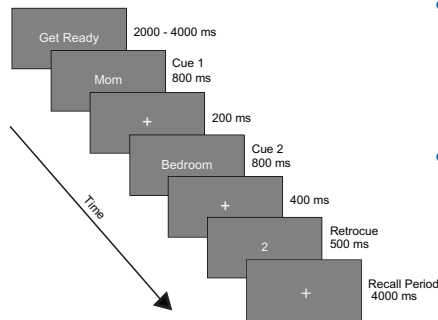
Flexible sampling plans

- Bayes Factors allow us to collect data iteratively
- More trials or more participants?
- Safer to overpower number of trials



*Example studies: making use of Bayes Factor
advantages*

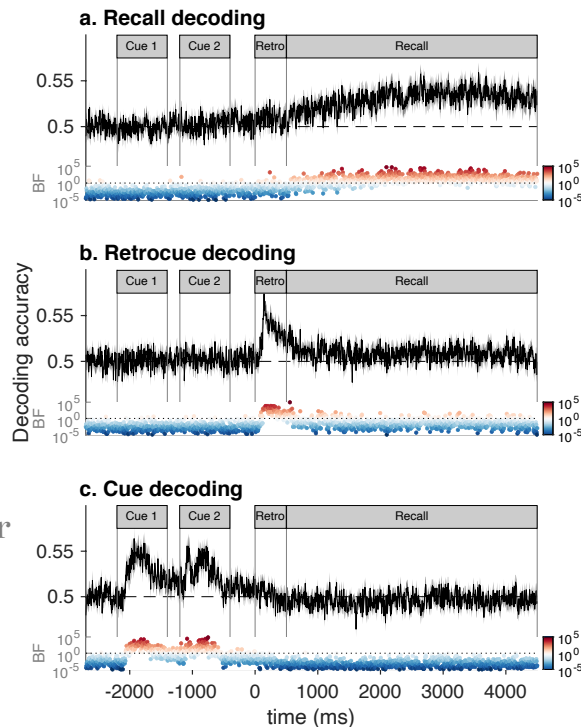
Mental imagery: when do we have the strongest/weakest evidence?



- Stimuli: personally-relevant people and places
- Is there information about stimulus class during mental imagery?

- Long epochs / recall period
- Is there a time-window where we have strong evidence for accurate recall?

[Interpretability of Bayes Factors]



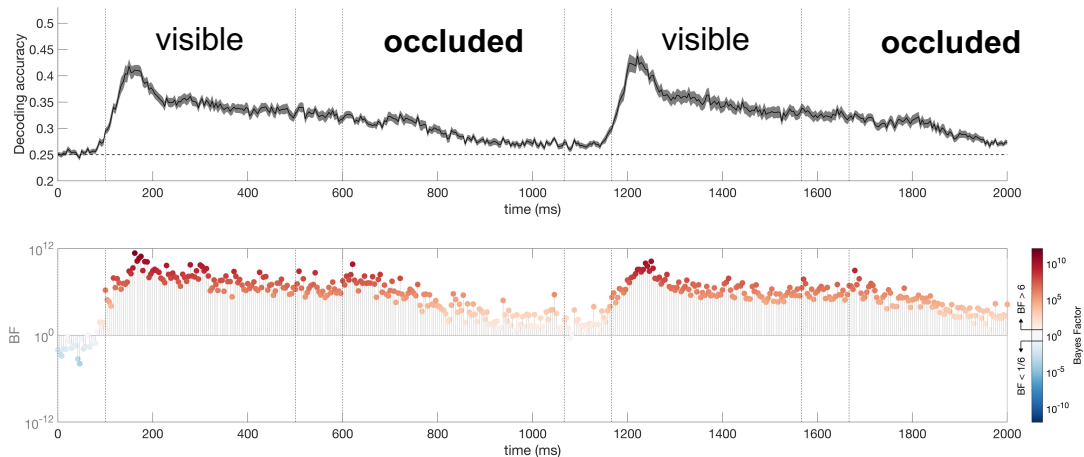
Occlusion: Are there timepoints with **no** information?

- How are object properties represented during occlusion?
- Possible that there is **no** information about the object during occlusion



[Contrasting H_0 and H_a]

In which quadrant is the object?



Thank you 😊



Denise Moerel



Chris Baker



Tijl Grootswagers

Paper on bioRxiv: *An empirically-driven guide on using Bayes Factors for M/EEG decoding*



WESTERN SYDNEY
UNIVERSITY



The MARCS Institute for Brain,
Behaviour and Development