

# Decoding behavioral responses from fMRI without learning behavioral responses from fMRI

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Introduction:

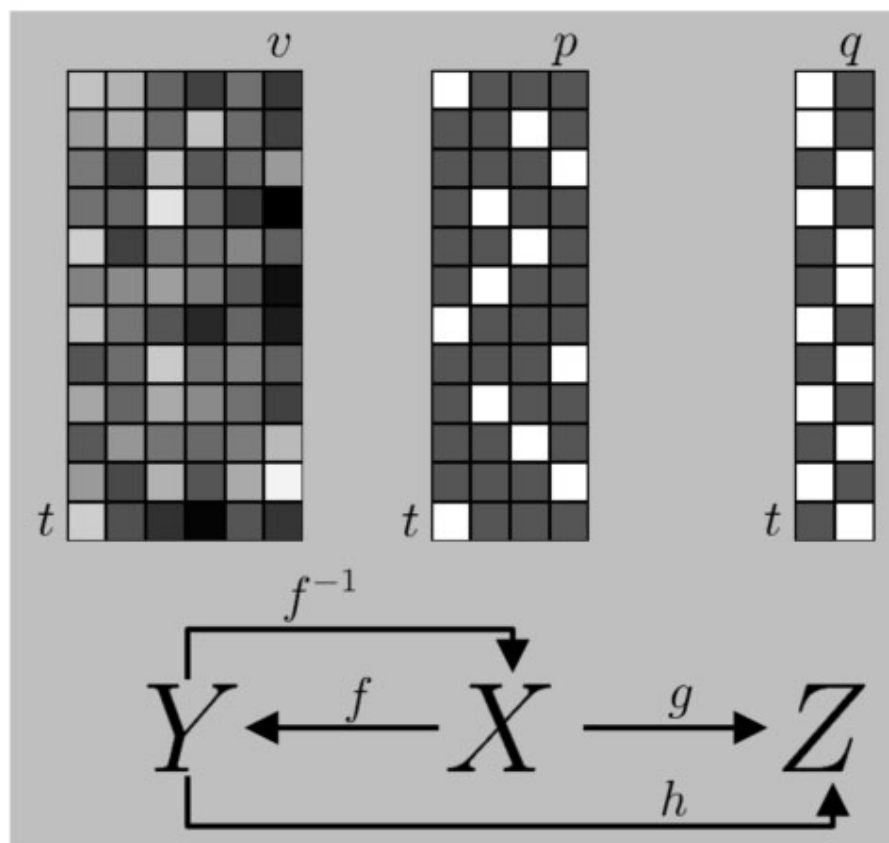
The data acquired during a functional magnetic resonance imaging (fMRI) experiment can usually be categorized into experimental design **X** (e.g. experimental conditions, modulator variables), measured signals **Y** (i.e. BOLD signals in several voxels) and behavioral data **Z** (e.g. button presses, stimulus ratings). In multivariate pattern analysis (MVPA) of fMRI data [1,2], behavioral data are typically decoded by training an algorithm to predict the recorded responses **Z** based on the measured data **Y** (*conventional response decoding*, CRD). Here we show that this can be achieved without constructing an explicit mapping from fMRI signals to behavioral responses. In fact, behavioral data can also be decoded when first reconstructing the experimental design **X** from measured data **Y** and then predicting behavioral responses **Z** from the

reconstructed design  $X$  (*neurobehavioral decoding*, NBD).

## Methods:

Our main focus was on comparing three decoding algorithms: (1) First, behavioral data are directly predicted from the experimental design (*psychobehavioral model*, PBM),  $Z = g(X)$ , e.g. using a logistic regression model predicting discrete button presses from continuous stimulus variables. (2) For CRD, behavioral responses are directly predicted from measured signals,  $Z = h(Y)$ , here using inverse transformed encoding models (ITEM) [3] for searchlight-based trial-wise decoding. (3) For NBD, the experimental design is decoded from measured signals,  $X = f^{-1}(Y)$ , again using trial-wise ITEM, and the estimated behavioral model is used to indirectly predict behavioral responses from fMRI signals:  $Z = g(f^{-1}(Y))$  (see Figure 1).

All these models are estimated in a cross-validated fashion, using leave-one-out cross-validation over fMRI recording sessions. Because behavioral responses and experimental conditions are best decoded from different parts of the brain, the approaches are made comparable by selecting the most informative searchlight based on within-sample decoding accuracy and reporting results as out-of-sample decoding accuracies.



decoding from the design  $\hat{Z} = \hat{g}(X)$

conventional response decoding  $\hat{Z} = \hat{h}(Y)$

neurobehavioral decoding  $\hat{Z} = \hat{g}(\hat{f}^{-1}(Y))$

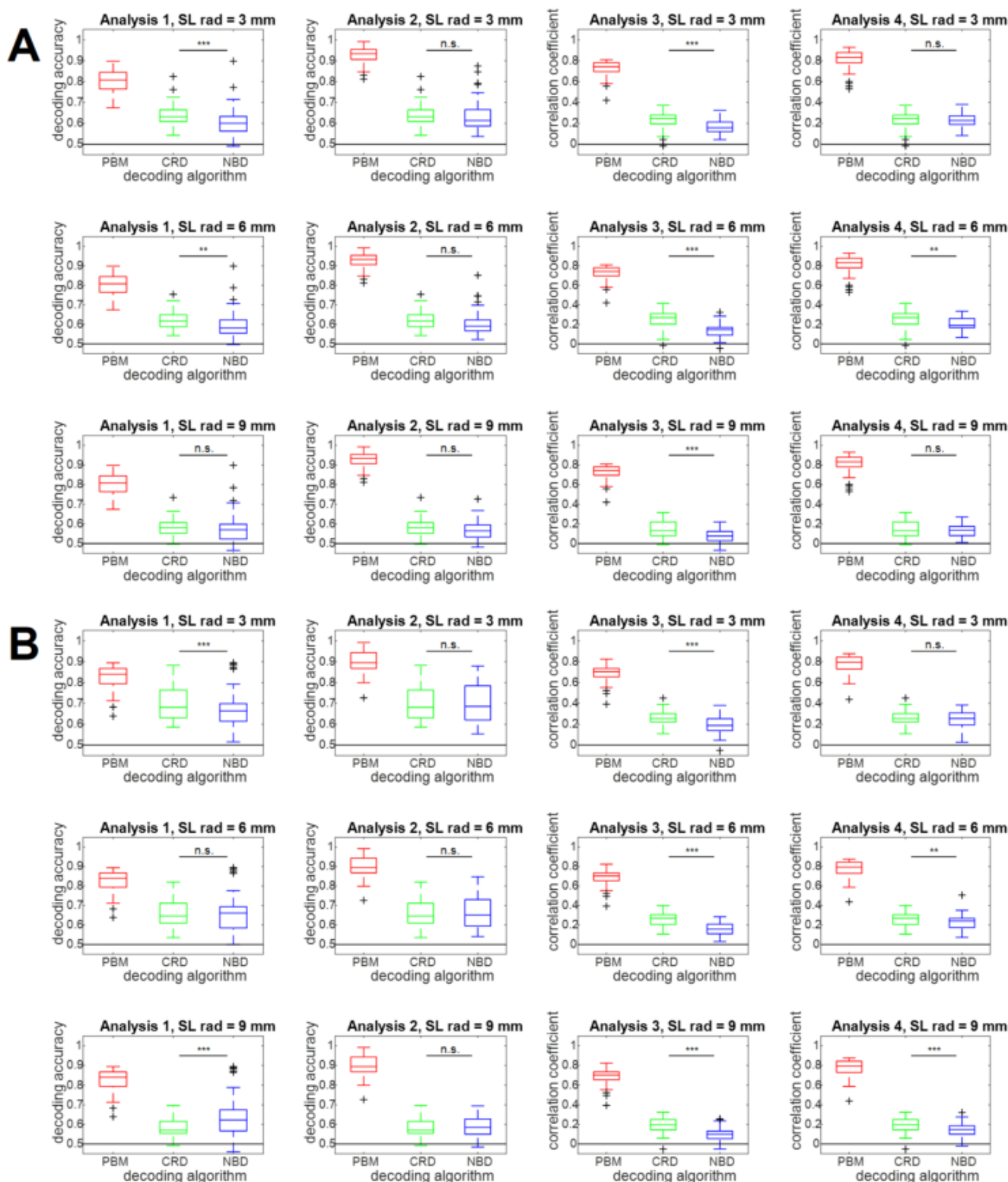
$t$  trials     $p$  conditions     $q$  responses     $v$  voxels

**Figure 1.** *Theory behind neurobehavioral decoding.* The analyses operate on the trial-by-voxel matrix  $Y$  (e.g. fMRI signal amplitudes in a searchlight), the trial-by-condition matrix  $X$  (i.e. experimental design) and the trial-by-response matrix  $Z$  (i.e. behavioral responses). The theory assumes a neurophysiological model  $f$  (from  $X$  to  $Y$ , inverse  $f^{-1}$ ), a psychobehavioral model  $g$  (from  $X$  to  $Z$ ) as well as a mapping for conventional response decoding  $h$  (from  $Y$  to  $Z$ ).

([https://files.aievolution.com/prd/hbm2101/abstracts/abs\\_2192/Figure\\_1.png](https://files.aievolution.com/prd/hbm2101/abstracts/abs_2192/Figure_1.png))

Results:

We analyzed the entire data set ( $N = 108$  subjects) from the *Neuroimaging Analysis and Replication and Prediction Study* (NARPS) [4,5]. In this experiment, subjects were offered a mixed gamble in each trial, with certain amounts of money to win or lose (= experimental design  $X$ ), and then indicated favorability of the bet (= behavioral responses  $Z$ ) using a four-point Likert scale [6]. In our investigations, we tested several analysis strategies (Analysis 1: discrete  $X$  &  $Z$ ; Analysis 2: continuous  $X$ , discrete  $Z$ ; Analysis 3: discrete  $X$ , continuous  $Z$ ; Analysis 4: continuous  $X$  &  $Z$ ) and observed that (i) the PBM clearly outperforms CRD and NBD; (ii) CRD and NBD both allow for above-chance decoding of behavioral responses; and (iii) CRD mildly outperforms NBD, except when decoding discrete  $Z$  from continuous  $X$  (see Figure 2). These results were largely unaffected by searchlight radius ( $r = 3, 6, 9$  mm) and experimental group (a between-subject factor).



**Figure 2.** Decoding accuracies as a function of decoding method. Performances of three decoding algorithms (red: PBM; green: CRD; blue: NBD) are visualized using box plots for four analysis strategies and three searchlight radii, separately for (A) equal range and (B) equal indifference ( $N = 54$  each) condition. CRD and NBD are tested against each other using a two-tailed paired t-test. Abbreviations: PBM, CRD, NBD: see text; n.s. = not significant; \* $p < 0.05$ , \*\* $p < 0.01$ , \*\*\* $p < 0.001$ .

([https://files.aievolution.com/prd/hbm2101/abstracts/abs\\_2192/Figure\\_2.png](https://files.aievolution.com/prd/hbm2101/abstracts/abs_2192/Figure_2.png))

## Conclusions:

In this proof-of-concept study, we have demonstrated that behavioral responses can be decoded without training on neurophysiological data measured during behavioral responses, but rather indirectly by taking a detour via the experimental design. This is particularly interesting, because CRD is commonly seen as a

sanity check, the decoding accuracy of which should not be exceeded by other analyses. It is also worth noting that in our example, just one response dimension (favorability), but two design dimensions (gain and loss) had to be decoded. We hypothesize that decoding the design from the data acts as a feature reduction mechanism which helps NBD predicting behavior using the psychologically most meaningful factors. In the future, we want to investigate the performance of NBD when the mapping from experimental conditions to behavioral responses is more deterministic [7,8] or completely random [9,10].

## Emotion, Motivation and Social Neuroscience:

Reward and Punishment

## Higher Cognitive Functions:

Decision Making

## Modeling and Analysis Methods:

Classification and Predictive Modeling <sup>1</sup>

Multivariate Approaches <sup>2</sup>

## Keywords:

Cognition

Data analysis

FUNCTIONAL MRI

Machine Learning

Multivariate

Statistical Methods

Other - MVPA

<sup>1|2</sup>Indicates the priority used for review

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No

Please indicate below if your study was a "resting state" or "task-activation" study.

Task-activation

Healthy subjects only or patients (note that patient studies may also involve healthy subjects):

Healthy subjects

Was any human subjects research approved by the relevant Institutional Review Board or ethics panel? NOTE: Any human subjects studies without IRB approval will be automatically rejected.

Yes

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Not applicable

Please indicate which methods were used in your research:

Functional MRI

Behavior

Computational modeling

Other, Please specify - multivariate pattern analysis

For human MRI, what field strength scanner do you use?

3.0T

Which processing packages did you use for your study?

SPM

Other, Please list - ITEM toolbox

Provide references using author date format

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