

Variational physiologically informed solution to hemodynamic and perfusion response estimation from ASL fMRI data

Aina Frau-Pascual and Florence Forbes and Philippe Ciuciu

contact: aina.frau-pascual@inria.fr

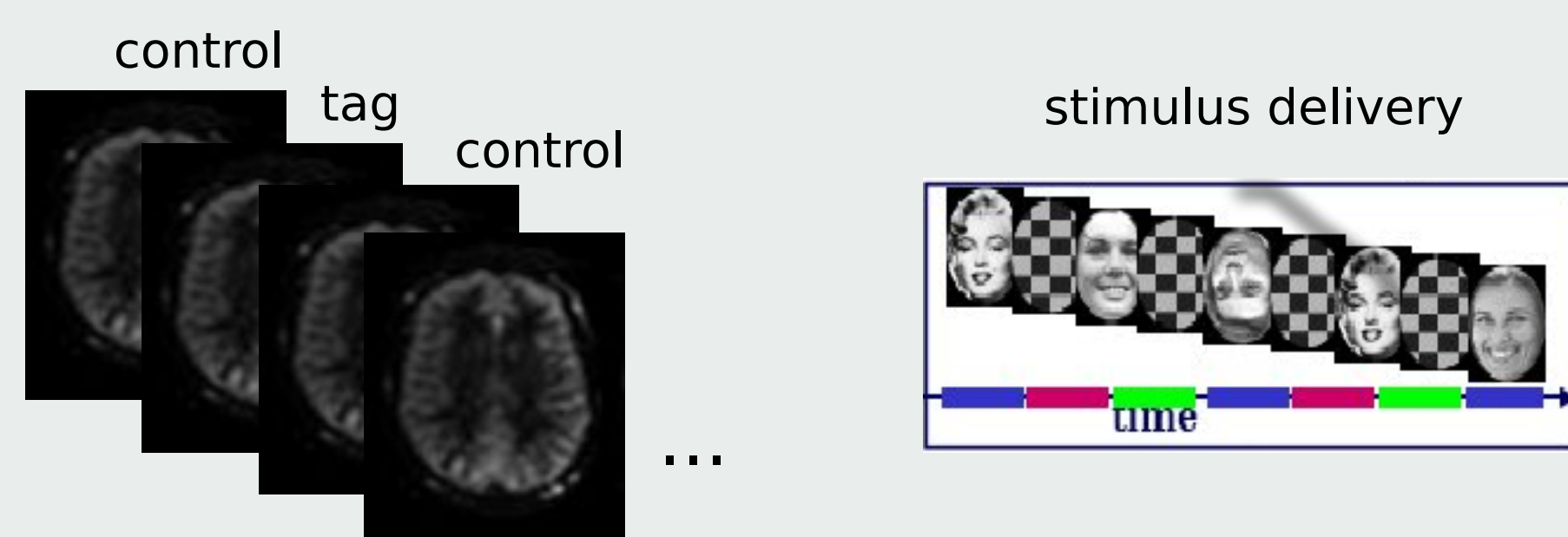
Abstract

Functional Arterial Spin Labeling (fASL) [1] MRI can provide a quantitative measurement of cerebral blood flow and its variations elicited by specific tasks. The statistical analysis of fASL has been done using

- General linear model (GLM) [2] with regressors based on the canonical hemodynamic response function.
 - Joint detection-estimation (JDE) [3] framework which allows the extraction of both task-related perfusion and hemodynamic responses not restricted to canonical shapes. Previous ASL-JDE attempts have been based on Markov Chain Monte Carlo (MCMC) methods, very computationally expensive.
- Contribution: a variational expectation-maximization (VEM) algorithm [4] for hemodynamic and perfusion responses estimation.

Framework

ASL fMRI [1] data provide a quantitative measurement of blood perfusion changes elicited by task performance or stimulus delivery in the brain

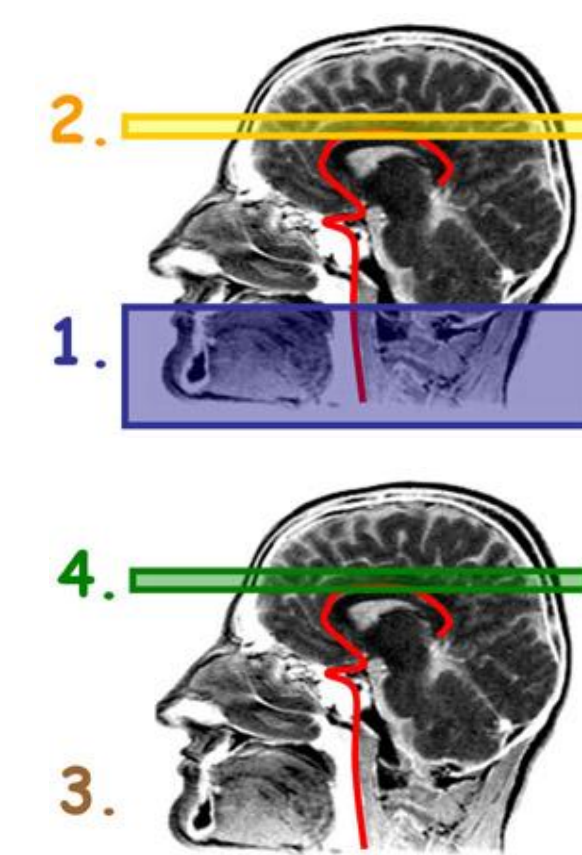


Magnetically tagged image (Tag)

Tag inflowing arterial blood by magnetic inversion

Control image

Repeat acquisition without labeling inflowing blood



Time delay (1) to (2): Labeled water reaches capillary bed and is exchanged with water molecules in the tissue

signal change

Control Image (4) - Tag Image (2)

$$\uparrow - \uparrow = \uparrow \propto \text{CBF}$$

The difference in magnetization is proportional to regional cerebral blood flow

Ref: http://fmri.research.umich.edu/research/main_topics/asl.php

ASL Joint detection estimation (JDE) framework [3]

$$\text{ASL signal} = \text{perfusion baseline} + \sum_{m=1}^M (\text{task-related perfusion } c_j^m \mathbf{W} \mathbf{X}^m \mathbf{g} + \text{task-related BOLD } a_j^m \mathbf{X}^m \mathbf{h}) + \text{drift term } \mathbf{P} \ell_j + \text{noise term } b_j$$

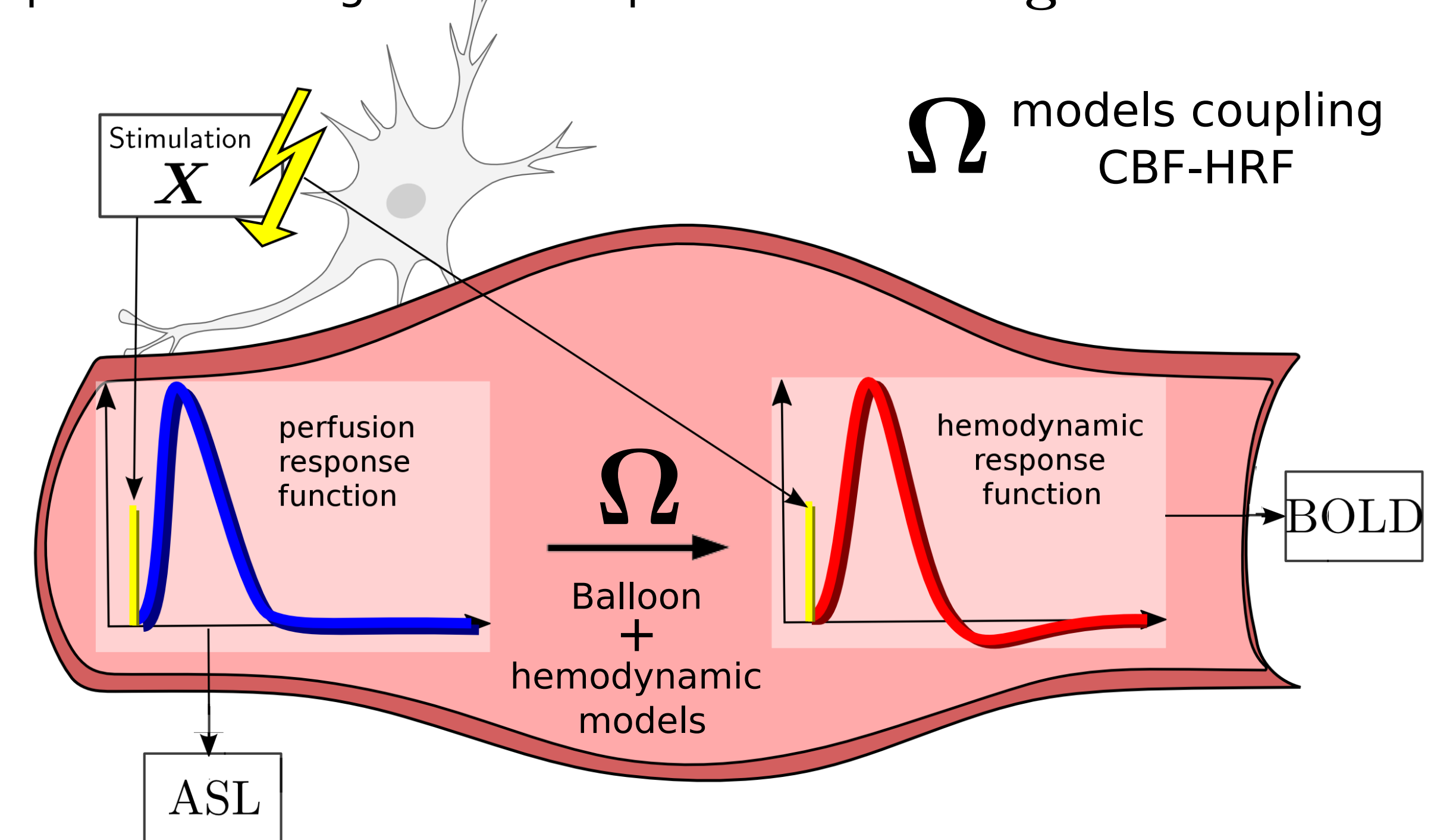
$$\mathbf{y}_j = \alpha_j \mathbf{w} + \sum_{m=1}^M (c_j^m \mathbf{W} \mathbf{X}^m \mathbf{g} + a_j^m \mathbf{X}^m \mathbf{h}) + \mathbf{P} \ell_j + b_j$$

For every voxel in a parcel, ASL signal can be decomposed into different terms. We estimate the parameters of this model.

Parameters: \mathbf{g} (perfusion response function (PRF)), \mathbf{h} (hemodynamic response function (HRF)), \mathbf{c} (perfusion response levels (PRLs)), \mathbf{a} (hemodynamic response levels (HRLs)), and \mathbf{q} (labels (active/non-active)).

Physiologically informed JDE [5]

We consider physiological information in the estimation as a prior knowledge of the response functions $\mathbf{g} = \Omega \mathbf{h}$



Expectation-Maximization

$$\text{E-step: } \tilde{\mathbf{p}}^{(r)} = \arg \max_{\tilde{\mathbf{p}}} F(\tilde{\mathbf{p}}, \boldsymbol{\theta}^{(r)})$$

$$\text{M-step: } \boldsymbol{\theta}^{(r+1)} = \arg \max_{\boldsymbol{\theta}} F(\tilde{\mathbf{p}}^{(r)}, \boldsymbol{\theta})$$

Maximizing function F is equivalent to minimizing the Kullback-Leibler divergence between $\tilde{\mathbf{p}}$ and the true posterior $p(\mathbf{a}, \mathbf{h}, \mathbf{c}, \mathbf{g}, \mathbf{q} | \mathbf{y})$

Variational EM

Restrict solutions to the ones that allow

$$\tilde{p}(\mathbf{a}, \mathbf{h}, \mathbf{c}, \mathbf{g}, \mathbf{q}) = \tilde{p}_a(\mathbf{a}) \tilde{p}_h(\mathbf{h}) \tilde{p}_c(\mathbf{c}) \tilde{p}_g(\mathbf{g}) \tilde{p}_q(\mathbf{q})$$

E and M step can be decomposed in stages corresponding to the different parameters

The E-H step, for example, goes:

$$\tilde{p}_h = \arg \max_{\tilde{p}_h \in \mathcal{D}_H} F(\tilde{p}_a, \tilde{p}_h, \tilde{p}_c, \tilde{p}_g, \tilde{p}_q; \boldsymbol{\theta})$$

Variational Expectation-Maximization

We can constraint the search to pointwise estimates $\tilde{\mathbf{h}}$ and $\tilde{\mathbf{g}}$ by replacing the probabilities on \mathbf{h} and \mathbf{g} by Dirac functions:

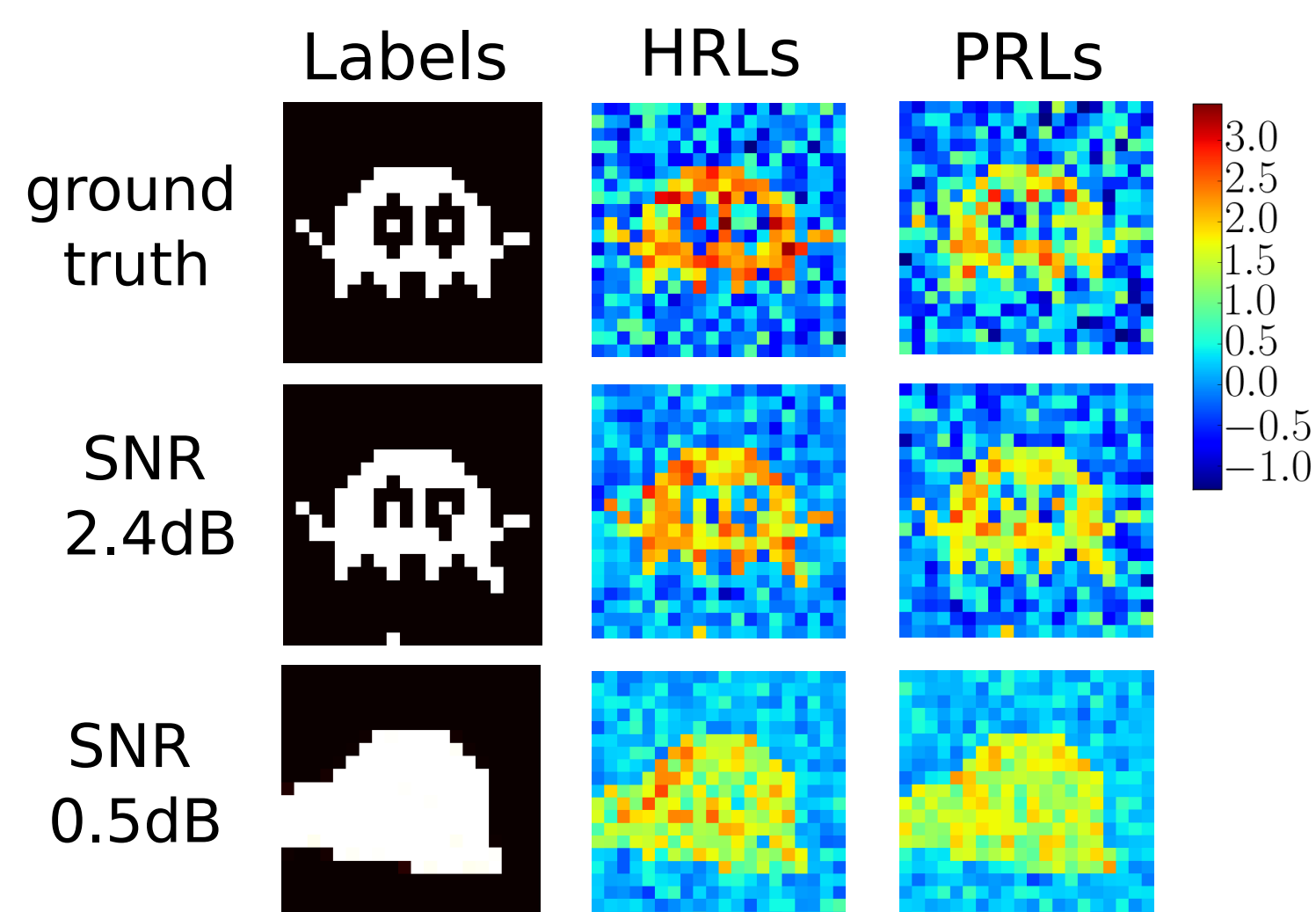
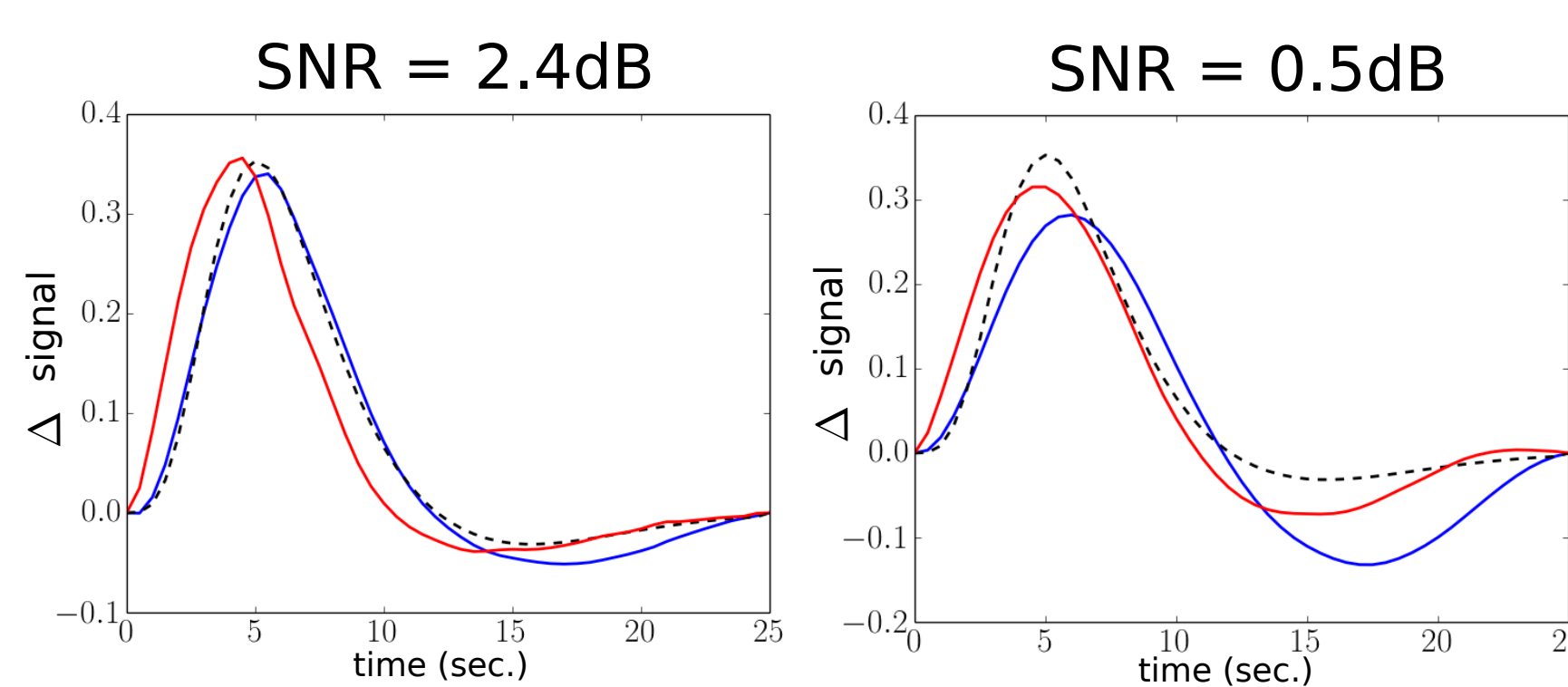
$$\tilde{\mathbf{p}} = \tilde{p}_a \delta_{\tilde{\mathbf{h}}} \tilde{p}_c \delta_{\tilde{\mathbf{g}}} \tilde{p}_q$$

And so: $\tilde{\mathbf{h}} = \arg \max_{\tilde{\mathbf{h}}} F(\tilde{p}_a \delta_{\tilde{\mathbf{h}}} \tilde{p}_c \delta_{\tilde{\mathbf{g}}} \tilde{p}_q; \boldsymbol{\theta})$

We can easily include constraints like $\|\mathbf{h}\|_2^2 = 1$, $\|\mathbf{g}\|_2^2 = 1$

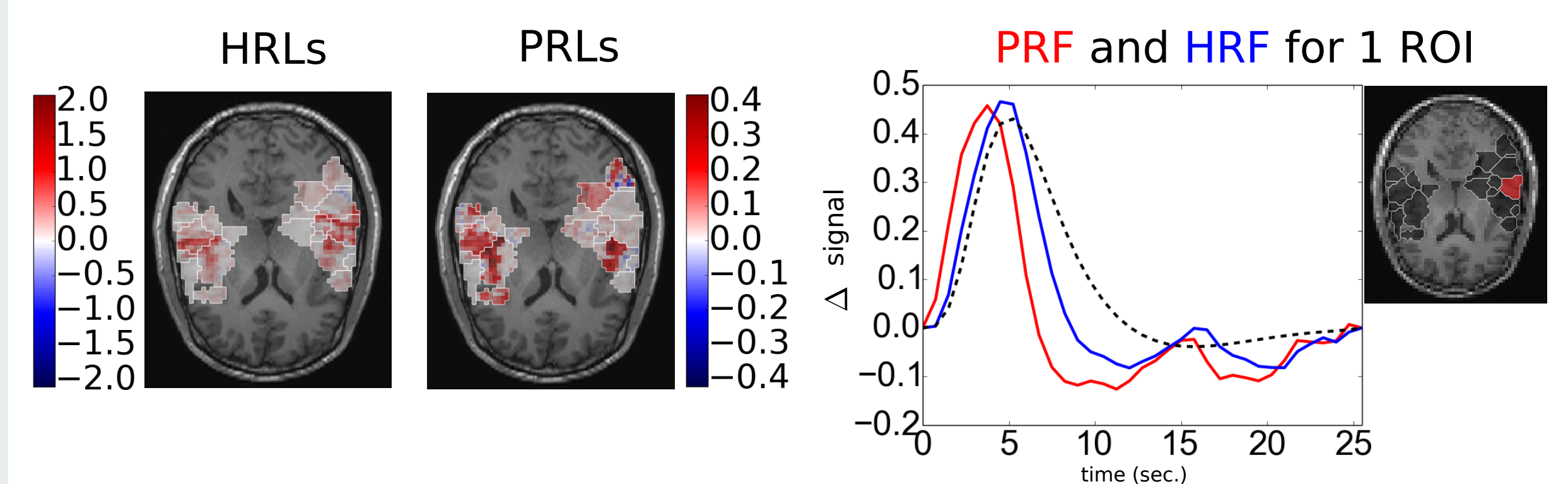
Artificial data

Repetition time: TR = 3s
Number of scans: 288
Fast event-related paradigm: mean ISI = 5s



Real data

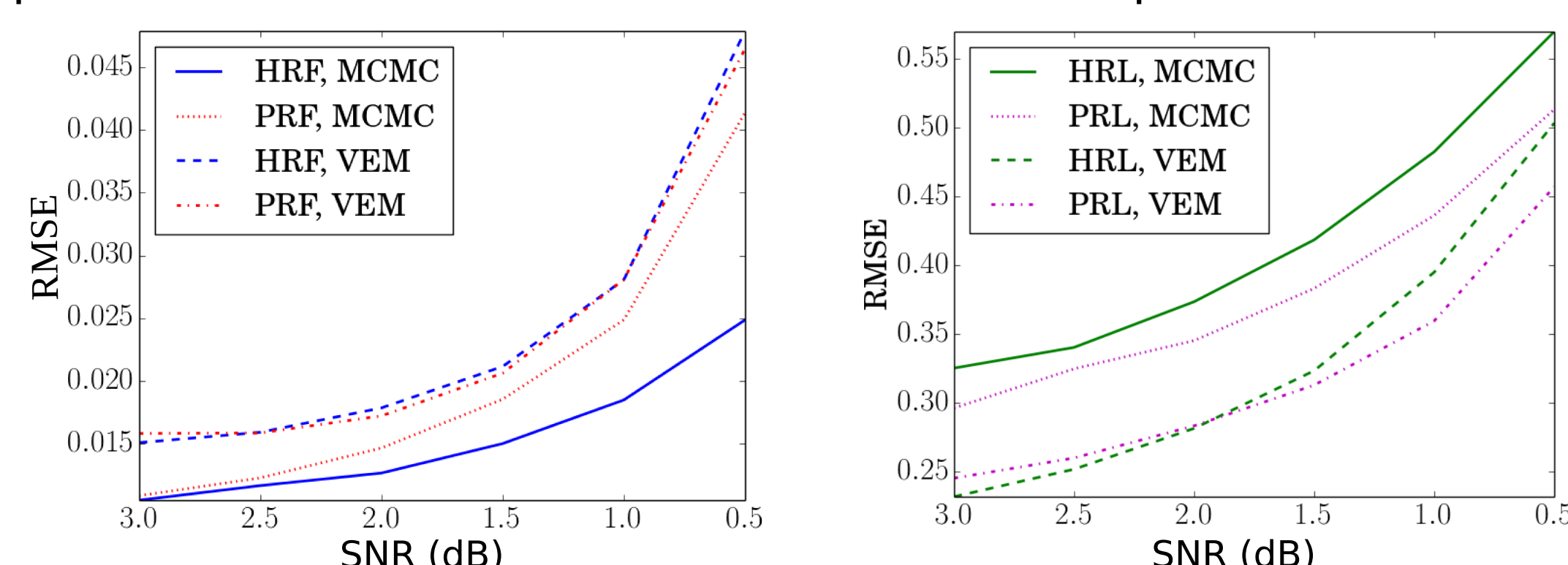
Paradigm: fast event-related design (mean ISI = 5.1s), with 60 auditory and visual stimuli, TR = 3s.



Results

Comparison with stochastic ASL-JDE

Both methods have a similar performance, but VEM recovers better response levels while MCMC recovers better response functions



References

- [1] D. Williams, J. Detre, J. Leigh, and A. Koretsky, "Magnetic resonance imaging of perfusion using spin inversion of arterial water", Proceedings of the National Academy of Sciences, vol. 89, no. 1, pp. 212-216, 1992.
- [2] L. Hernandez-Garcia, H. Jahani, and D. B. Rowe, "Quantitative analysis of arterial spin labeling fMRI data using a general linear model", Magnetic resonance imaging, vol. 28, no. 7, pp. 919-927, 2010.
- [3] T. Vincent, J. Warnking, M. Villien, A. Krainik, P. Ciuciu, and F. Forbes, "Bayesian Joint Detection-Estimation of cerebral vasoreactivity from ASL fMRI data," in 16th Proc. MICCAI, LNCS Springer Verlag, Nagoya, Japan, Sep. 2013, vol. 2, pp. 616-623.
- [4] L. Chaari, T. Vincent, F. Forbes, M. Dojat, and P. Ciuciu, "Fast joint detection-estimation of evoked brain activity in event-related fMRI using a variational approach," IEEE Trans. on Medical Imaging, vol. 32, no. 5, pp. 821-837, May 2013
- [5] A. Frau-Pascual, T. Vincent, J. Sloboda, P. Ciuciu, and F. Forbes, Physiologically informed Bayesian analysis of ASL fMRI data", in Bayesian and Graphical Models for Biomedical Imaging, pp. 37-48. Springer, 2014