
Region-network hierarchical sparsity priors for high-dimensional inference in brain imaging

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Abstract

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1 Introduction

Many quantitative scientific domains underwent a recent passage from the classical regime (i.e., “long data”) to the high-dimensional regime (i.e., “wide data”) [30]. Also in the brain imaging domain, many contemporary methods for acquiring brain signals yield more variables per observation than total observations per data sample. This high-dimensional scenario challenges various statistical methods from classical statistics. For instance, estimating generalized linear models without additional assumptions yields an underdetermined system of equations. Many such ill-posed estimation problems have benefited from *sparsity* assumptions [11, 24]. They act as a regularizer and can be used for model selection. Sparse supervised and unsupervised learning algorithms have proven to yield statistical relationships that can be readily estimated, reproduced, and interpreted [21]. Generally, *structured sparsity* can impose domain knowledge on the statistical estimation, thus shrinking and selecting variables guided by expected data distributions [3]. Such restrictions to complexity are an attractive plan of attack for the >100,000 variables of brain maps. Yet, what generally accepted neurobiological structure lends itself to harness the *curse of dimensionality* by structured sparsity priors?

Concepts on human brain organization have long been torn between the two extremes *functional specialization* and *functional integration*. Functional specialization emphasizes that microscopically distinguishable brain regions are responsible distinct classes of computational processes [32]. Functional integration, in turn, emphasizes that brain function is enabled by complex connections between these distinct brain regions [45]. These notions were predominantly derived from invasive examination of anatomy (i.e., histological preparation), connectivity (i.e., axonal tracing), and functional properties (i.e., single-cell recordings) in animals. Regarding functional segregation into specialized regions, early histological investigations into the microscopic heterogeneity of the human cerebral cortex have resulted in several detailed anatomical maps [9, 49]. Regarding axonal connections, each such cortical area has been observed to possess a unique set of incoming and outgoing connections [36, 51, 41]. Both local infrastructure and its unique global connectivity profile together are thought to realize brain function. In sum, cortical brain modules versus connections between them reflect functional specialization versus functional integration [20, 35]. Importantly, probably no existing brain analysis method acknowledges that both functional organizations are inextricably involved in the realization of mental operations [47, 40].

Functional specialization has been explored and interpreted based on many different research methods. Single-cell recordings and microscopic examination revealed, for instance, the specialization in the occipital visual cortex into V1, V2, V3, V3A/B, and V4 [25, 53]. Tissue lesion of the mid-fusiform gyrus of the visual system, in turn, was frequently reported to impair recognition of others' identity from faces [26]. The whole-brain localization of sensory, motor, and emotional functions to cortical areas was later enabled by non-invasive brain imaging with functional magnetic resonance imaging (fMRI) and positron emission tomography (PET) [19]. Further, radioactive mapping **Michael: is this the correct term?** of neurotransmitter receptors rendered accessible yet another local characteristic of neuronal populations [54]. In the computational era, automatic clustering methods are increasingly employed to regionally differentiate the cerebral cortex, which can partly be more fine-grained than classical microscopical borders [7, 17]. High-throughput approaches today enable ultrahigh-resolution 3D models of brain anatomy at near-cellular scale [2]. As a crucial common point, all these methodological approaches yield neuroscientific findings that are naturally interpreted according to non-overlapping, discrete region compartments as the basic architecture of brain organization.

It is more recent that the main interpretational focus has shifted from circumscribed regions to network stratifications in systems neuroscience [52, 46]. Invasive axonal tracing studies in monkeys were complemented by diffusion MRI tractography in humans as a now frequently employed method to outline fiber bundles between brain regions [27]. Besides analyses of electrophysiological oscillations [13] and graph-theoretical properties [12], studies of functional connectivity [10] and independent component analysis (ICA) [6] became the workhorses of network discovery in neuroimaging. These revealed the important implication of canonical brain networks across psychological tasks, including the so-called "default-mode network" [38], "salience network" [42], and "dorsal attention network" [14]. Characteristic changes in the configuration of these macroscopical networks were repeatedly observed to be induced by the onset of given psychological tasks [18]. As a common point of all these methods, interpretation of findings naturally embraces cross-regional integration by overlapping network compartments as the basic architecture of brain organization, in stark contrast to methods examining regional specialization.

Building on these two major interpretational streams in systems neuroscience, the present study proposes to incorporate established neurobiological structure underlying functional segregation and integration into supervised estimators by hierarchical structured sparsity. Learning techniques exploiting structured sparsity have recently made much progress in various application domains from processing of auditory signals [16], natural images [23] and videos [31, 34] to astrophysics [48], genetics [39, 33], and conformational dynamics of protein complexes [29]. This is extended by the present work that introduced neuroscience-specific estimators capitalizing on neurobiologically plausible region and network priors. Based on the largest neuroimaging repository, we demonstrated that domain-informed supervised models gracefully tackle the curse of dimensionality, yield more human-interpretable results, and generalize better to new samples than domain-naïve estimators.

2 Methods

Our main contribution is the domain-specific adaptation of sparse structured penalties making it possible to jointly incorporate functional specialization and functional integration priors into statistical estimators. We make use of the hierarchical group lasso as introduced by Jennatton and others in [28] to create a set of convex penalties with which we can probe this interplay between regional specificity and inter-regional functional integration during different cognitive tasks.

Rationale Three-dimensional voxel brain images as obtained by various neuroimaging techniques are very high-dimensional, but also very highly structured. While its explicit dimensionality, the number of voxels, which varies with the resolution of the image is generally of the order of 100000, the number of samples available for analyses never exceeds hundreds or thousands. This $n \ll p$ situation immediately implies an underdeterminedness for any linear model basing its decisions on dot products with the voxel values.

Luckily, there is an abundance structure to be exploited, which can be injected into the estimation as a domain knowledge prior. Here we are interested in incorporating knowledge about both the well-established modules of functional specialization and the interactions of these modules amongst

each other across the brain. Developmentally, such large-scale networks emerge during late fetal growth (Doria et al., 2010), before cognitive capacities mature in childhood. In adults, nodes of a same cohesive network have more similar functional profiles than nodes from different networks (Anderson et al., 2013). On the signal level, functional modules are localized structures implicating neighboring voxels in an image. A variety of known functional modules has been compiled into atlases of the brain. Different types of atlases rely on different ways of segregating modules from another. For instance, anatomical atlases rely on the variation of tissue properties across the brain, which can be at the cellular level or based on higher-order structure. Global functional networks can be discovered by statistical analysis of functional brain imaging data, for example the BOLD contrast in fMRI. A well-established method of estimating these networks is independent component analysis [?]. It robustly identifies the default mode, saliency and attention networks. Global functional networks as estimated by ICA are continuous brain maps attributing weights to each voxel. Since the regions involved in these networks are nevertheless largely disjoint spatially and roughly in correspondence with module boundaries **Michael: verify this!!!**, it is possible to uniquely associate each module from a region atlas to one of the extracted global functional networks.

This correspondence makes it possible to encode our knowledge about the networks and regions into a hierarchy: A brain contains a certain number of networks, which in turn consist of the atlas modules as subregions. We can use this hierarchy to create a structural prior of expected weight maps for classifiers.

The hierarchical group lasso as introduced by [?] lends itself perfectly to represent this structure. It is based on the group lasso [?] but permits groups to contain each other in a tree structure. The first level of the tree are the network groups containing all the voxels of the modules associated to them. Then each network contains as subgroups the modules associated with it, grouping the voxels of each of these regions together. As with the group lasso, it is possible to associate an individual penalty to each group. In trading off the penalties on the network level against the penalties of the region level, we can create different regimes of estimation: Setting a low penalty on the network groups makes it probable that all of them are active in the estimated weight map. If we then select higher penalties on region groups, selection of relevant region groups is forced without the bias of the network maps. Conversely, setting low penalties on the region maps makes it possible for all voxels to be active. Selecting higher penalties on the networks then leads to a selection of networks with all regions associated to it. Taken together and varied smoothly, we can evaluate a tradeoff between the relevance of modular regions and functionally integrated networks in statistical estimation of brain activity.

Problem formulation We formulate our estimation problem in the framework of regularized risk estimation applied to linear models: We would like to estimate a good predictor of cognitive task given a brain image. Let the set $\mathcal{X} \subset \mathbb{R}^p$ represent brain images of $p > 0$ voxels.

Then we would like to minimize the risk $\mathcal{L}(\hat{y}, y)$, where $\hat{y} = X\hat{w} + \hat{b}$, while regularizing to incorporate a useful prior. Taken together, this can be framed as an optimization problem

$$\arg \min_{w,b} \mathcal{L}(Xw + b, y) + \lambda \Omega(w),$$

where $\lambda > 0$ and Ω is the regularizer.

Brain regions are defined as disjoint groups of voxels. Let \mathcal{G} be a partition of $\{1, \dots, p\}$, i.e.

$$\bigcup_i g_i = \{1, \dots, p\} \text{ and } g_i \cap g_j = \emptyset \quad \forall i \neq j$$

Brain networks consist of regions and are thus super-regions or groups of regions. The set of brain networks \mathcal{H} is also a partition of $\{1, \dots, p\}$ and in addition it is consistent with \mathcal{G} in the sense that

$$\text{for all } g \in \mathcal{G}, h \in \mathcal{H}, \quad \text{either } g \subset h \text{ or } g \cap h = \emptyset.$$

This allows a clear association of each region $g \in \mathcal{G}$ to a network $h \in \mathcal{H}$ and thus establishes a tree structure (up to adding a root node containing all voxels).

For a brain image $w \in \mathbb{R}^p$ and a group g , the vector $w_g \in \mathbb{R}^{|g|}$ is defined as the restriction of w to the coordinates in g . The structured penalty incorporating network and region information can then

be written as

$$\Omega(w) = \alpha \sum_{h \in \mathcal{H}} \eta_h \|w_h\|_2 + \beta \sum_{g \in \mathcal{G}} \eta_g \|w_g\|_2.$$

According to [?] we set $\eta_g = 1/\sqrt{|g|}$ to account for varying group size. The hierarchy-level-specific factors $\alpha > 0$ and $\beta > 0$ are used to trade-off region-weighted and network-weighted models against each other.

The prediction problem at hand is a multiclass classification. We choose to attack this using one-vs-rest scheme on a binary logistic regression, whose loss can be written as

$$\sum_{i=1}^n \log(1 + \exp(-y_i \langle x_i, w \rangle)) + \lambda \Omega(w),$$

if $y \in -1, 1$ and with $x_i \in \mathbb{R}^p$ the training sample brain images.

MISSING

- cross-validation scheme
- analysis of penalty tradeoff (CV scheme for that?)
- mentioning of control methods in methods. Trace norm, sparse group sparsity. Anything else? ENET?

Implementation. The analyses were performed in Python. We used *nilearn* to handle the large quantities of neuroimaging data [1] and *Theano* for automatic, numerically stable differentiation of symbolic computation graphs [5, 8]. All Python scripts that generated the results are accessible online for reproducibility and reuse (<http://github.com/banilo/nips2015>).

all algorithm from a same software library - ζ SPAMs

Data. As the currently biggest openly-accessible reference dataset, we chose resources from the Human Connectome Project (HCP) [4]. Neuroimaging task data with labels of ongoing cognitive processes were drawn from 500 healthy HCP participants (cf. Appendix for details on datasets). 18 HCP tasks were selected that are known to elicit reliable neural activity across participants (Table 1). In sum, the HCP task data incorporated 8650 first-level activity maps from 18 diverse paradigms administered to 498 participants (2 removed due to incomplete data). All maps were resampled to a common $60 \times 72 \times 60$ space of 3mm isotropic voxels and gray-matter masked (at least 10% tissue probability). The supervised analyses were thus based on labeled HCP task maps with 79,941 voxels of interest representing z-values in gray matter.

Cognitive Task	Stimuli	Instruction for participants
1 Reward	Card game	Guess the number of a mystery card for gain/loss of money
2 Punish		
3 Shapes	Shape pictures	Decide which of two shapes matches another shape geometrically
4 Faces	Face pictures	Decide which of two faces matches another face emotionally
5 Random		
6 Theory of mind	Videos with objects	Decide whether the objects act randomly or intentionally
7 Mathematics	Spoken numbers	Complete addition and subtraction problems
8 Language	Auditory stories	Choose answer about the topic of the story
9 Tongue movement		Move tongue
10 Food movement	Visual cues	Squeezing of the left or right toe
11 Hand movement		Tapping of the left or right finger
12 Matching		Decide whether two objects match in shape or texture
13 Relations	Shapes with textures	Decide whether object pairs differ both along either shape or texture
14 View Bodies	Pictures	Passive watching
15 View Faces	Pictures	Passive watching
16 View Places	Pictures	Passive watching
17 View Tools	Pictures	Passive watching
18 Two-Back	Various pictures	Indicate whether current stimulus is the same as two items earlier

Table 1: Description of psychological tasks to predict.

These labeled data were complemented by unlabeled activity maps from HCP acquisitions of unconstrained resting-state activity [43]. These reflect brain activity in the absence of controlled thought.

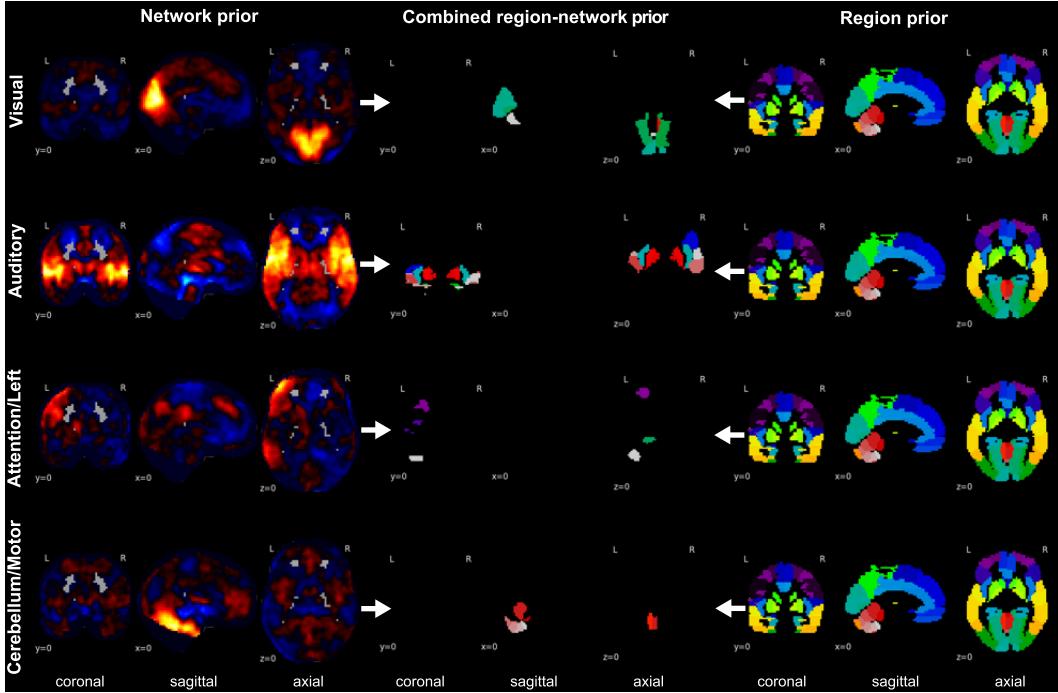


Figure 1: **Building blocks of the region-network tree.** Depicts neurobiological priors introduced into the classification problem by hierarchical structured sparsity. *Left:* Continuous, partially overlapping brain network priors (*hot-colored*, taken from [44]) accommodate the functional integration perspective of brain organization. *Right:* Discrete, non-overlapping brain region priors (*single-colored*, taken from [15]) accommodate the functional segregation perspective. *Middle:* These two types of predefined voxel groups are incorporated into hierarchical priors of parent networks with their descending region nodes. *Top to bottom:* Four exemplary region-network priors are shown, including the early cortex that processes visual and sound information from the environment, a well-known attentional circuit in the left brain hemisphere, and the cerebellum that is involved in motor behavior.

In sum, the HCP rest data concatenated 8000 unlabeled, noise-cleaned rest maps with 40 brain maps from each of 200 randomly selected participants.

We were further interested in the utility of the optimized low-rank projection in one task dataset for dimensionality reduction in another task dataset. To this end, the HCP-derived network decompositions were used as preliminary step in the classification problem of another large sample. The ARCHI dataset [37] provides activity maps from diverse experimental tasks, including auditory and visual perception, motor action, reading, language comprehension and mental calculation. Analogous to HCP data, the second task dataset thus incorporated 1404 labeled, grey-matter masked, and z-scored activity maps from 18 diverse tasks acquired in 78 participants.

sparse statistical models have only few nonzero parameters

3 Experimental Results

Benchmarking hierarchical tree sparsity against common sparsity penalties. Hierarchical region-network priors have been systematically evaluated against other popular choices of sparse classification algorithms in an 18-class scenario (Figure 2). Logistic regression with ℓ_1/ℓ_2 block norm penalization incorporated a hierarchy of previously known region and network neighborhoods for a neurobiological bias of the statistical estimation. Vanilla logistic regression with ℓ_1 -penalization does not assume any previously known special structure. This classification estimator embraces a vision of neural activity structure that expects a minimum of topographically and functionally independent brain voxel to be relevant. Logistic regression with sparse group spar-

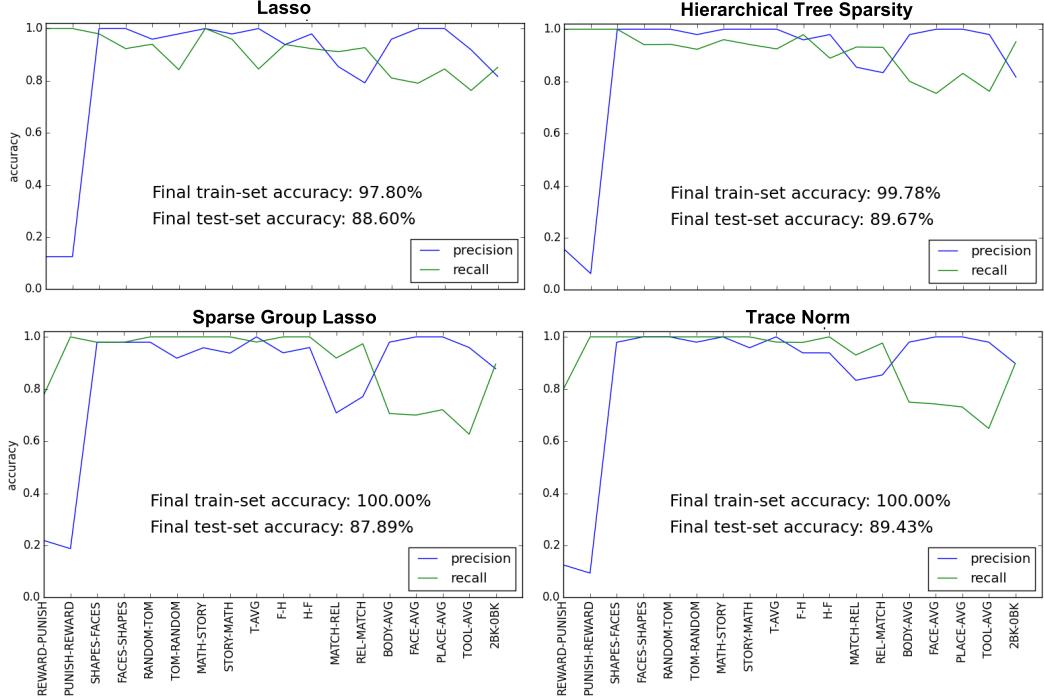


Figure 2: Model performance across sparsity priors. Compares the performance of logistic regression estimators with 4 different structured and unstructured sparsity terms in classifying neural activity from 18 psychological tasks. The class-wise precision and recall metrics were obtained on the same test set. Unstructured ℓ_1 -penalized logistic regression (*upper left*) imposed a minimum of relevant brain voxels without assuming special structure. Structured ℓ_1/ℓ_2 block norm with additional ℓ_1 term (*lower left*) imposed region compartments, but naïve to network structure. Structured trace-norm penalization (*lower right*) imposed low-rank structure with sparsity of network patterns, but naïve to region structure. Structured ℓ_1/ℓ_2 block norm with a hierarchy of both region and network priors (*upper right*) exhibited the best out-of-sample performance. A priori knowledge of both region and network neighborhoods was hence most beneficial for predicting psychological tasks from brain maps.

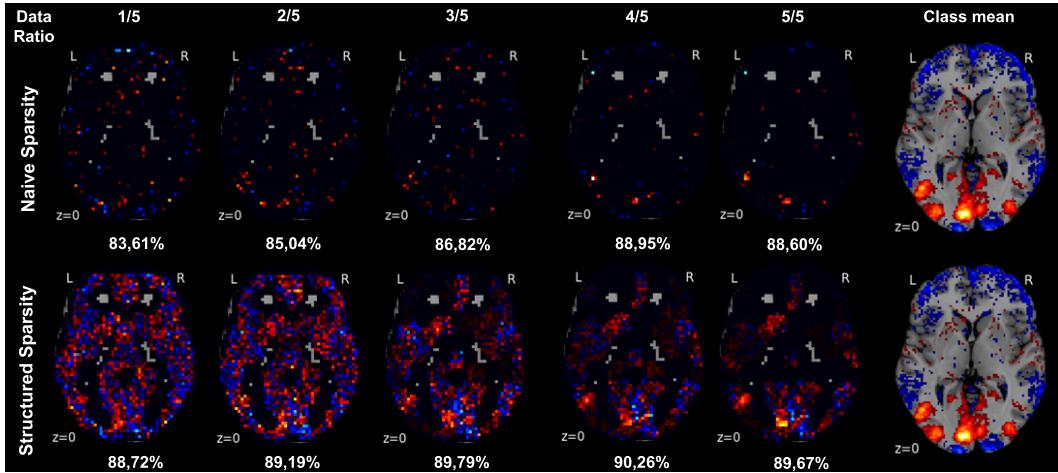


Figure 3: Naïve versus informed sparse model selection across training set sizes. Ordinary ℓ_1 -penalized logistic regression (*upper row*) is compared to hierarchical-tree-penalized logistic regression (*lower row*) with increasing fraction of the available training data (*left to right columns*). For one example (i.e., “View tools”) from 18 psychological tasks, unthresholded axial maps of model weights are shown for comparison against the sample average of that class (*rightmost column*, thresholded at the 75th percentile). The out-of-sample accuracies for predicting all 18 classes is given in percent. In the data-scarce scenario, typical for brain imaging, hierarchical tree sparsity achieves much better support recovery with the biggest difference in model performance. In the data-rich scenario, neurobiologically informed logistic regression profits more from the available information quantities than neurobiologically naive logistic regression.

sity imposes a structured ℓ_1/ℓ_2 block norm with additional ℓ_1 term with a known atlas of region voxel groups onto the statistical estimation process. This supervised estimator shrinks and selects the coefficients of topographically compact voxel groups expected to be relevant together. Logistic regression with trace-norm penalization imposed low-rank structure. This supervised classification algorithm expected a minimum of unknown network patterns to be relevant. The stratified and shuffled training data were submitted to a nested cross-validation scheme for model selection and model assessment. In the inner CV layer, the logistic regression estimators have been trained in a one-versus-rest design that distinguishes each class from the respective 17 other classes (number of maximal iterations=100, tolerance=0.001). In the outer CV layer, grid search selected among candidates for the respective λ parameter by searching between 10^{-2} and 10^1 in 9 steps on a logarithmic scale. Importantly, the thus selected sparse logistic regression classifier was evaluated on an identical test set in all settings. Hierarchical tree sparsity demonstrated the best generalization in distinguishing unseen neural activity maps from 18 tasks (89.67%, mean recall XX.YY, mean precision XX.YY). It was closely followed by logistic regression with trace-norm regularization that is structured but not neurobiologically informed (89.43%, mean recall XX.YY, mean precision XX.YY). Lasso featured an average performance comparing to the other sparse estimators (88.60%, mean recall XX.YY, mean precision XX.YY). Introduced a priori knowledge of brain region compartments by sparse group sparsity performed worst (87.89%, mean recall XX.YY, mean precision XX.YY). In sum, biasing sparse model selection by domain knowledge of region-network hierarchies outperformed other types of frequently used sparse penalization techniques.

Sample complexity of naïve versus informed sparse model selection. Subsequently, the sample complexity of ℓ_1 -penalized and hierarchical-tree-penalized logistic regression were quantitatively compared (Figure 3). Region-network priors should bias model selection towards more neurobiologically plausible classification estimators. This should yield better out-of-sample generalization and support recovery than ℓ_1 -constrained logistic regression naïve to neurobiology in the data-scarce and data-rich scenarios. The HCP task data with examples from 18 classes were first divided into 90% of training set (i.e., 7584 neural activity maps) and 10% of test set (i.e., 842 maps). Both learning algorithms were fitted based on the training set at different subsampling fractions: 20% (1516 maps), 40% (3033 maps), 60% (4550 maps), 80% (6067 maps), and 100% (7584 maps). The

stratified and shuffled training data were submitted to a nested cross-validation scheme for model selection and model assessment. In the inner CV layer, the logistic regression estimators have been trained in a one-versus-rest design that distinguishes each class from the respective 17 other classes (number of maximal iterations=100, tolerance=0.001). In the outer CV layer, grid search selected among candidates for the respective λ parameter by searching between 10^{-2} and 10^1 in 9 steps on a logarithmic scale. Importantly, the thus selected sparse logistic regression classifier was evaluated on an identical test set in all settings. Three observation have been made. In the data-scarce scenario (i.e., 1/5 of actual training data), hierarchical tree sparsity achieved the biggest advantage in out-of-sample performance by 5.11% as well as better support recovery with weight maps already much closer to the training data average. In the case of scarce training data, which is typical for the brain imaging domain, regularization by region-network priors indeed allowed for more effective extraction of classification-relevant structure from the neural activity scans. Across scenarios, the weight maps from ordinary logistic regression exhibit higher variance and many more zero coefficients than hierarchical tree logistic regression. Given the usually high multicollinearity in neuroimaging data, this observation is likely to reflect instable selection of representatives among class-responsive predictor groups due to the ℓ_1 -norm penalization. In the data-rich scenario (i.e., entire training data used for model fitting), neurobiologically informed logistic regression profits more from the increased information quantities than neurobiologically naive logistic regression. That is, the region-network priors actually further enhance the similarity to the weight maps even in abundant input data. This was the case although the maximal classification performance of $\approx 90\%$ has already been reached with small training data fractions by the structured estimator. In contrast, the unstructured estimator reached this generalization performance only with bigger input data quantities.

Support recovery as a function of region and network emphasis. Finally, the relative importance of the region and network priors within the hierarchical tree prior was quantified (Figure 4). The η_g group of region priors was multiplied with a region-network ratio, while the η_g group of network priors was biased by the corresponding network-region ratio. A region-network ratio of 3, for instance, increased the relative importance of known region structure by multiplying $\frac{3}{1}$ to the η_g factor of all region groups and multiplying $\frac{1}{3}$ to all network groups. The data splitting cross-validation scheme was identical to the above modelling experiments. As the most important observation, a range between region-dominant and network-dominant structured penalties yields quantitatively almost identical generalization to new data but qualitatively different decision functions manifested in the weight maps (Figure 4, second and forth column). Classification models with many zero coefficients but high absolute coefficients in either region compartments or network compartments can similarly extrapolation to unseen neural activity maps. Second, these perform similar to equilibrated region-network priors that set less voxel coefficients to zero and spread the probability mass with lower absolute coefficients across the whole brain (Figure 4, third column in the middle). Third, overly strong emphasis on either level of the hierarchical prior can yield the neurobiologically informative maps of the most necessary region or network structure for statistically significant out-of-sample performance (Figure 4, leftmost and rightmost columns). In sum, stratifying the hierarchical tree penalty between region and network emphasis suggests that *class-specific region-network weights* might offer more performant and more interpretable classification models in the future.

4 Discussion

Relevant structure in neuroimaging data has long been investigated according to two separate organizational principles: functional segregation into discrete brain regions and functional integration by interregional brain networks. This paper demonstrates the simultaneous exploitation of both these neurobiological compartments for sparse variable selection and high-dimensional prediction in a reference dataset. Introducing existing domain knowledge into model selection allowed privileging members of the function space that are most neurobiologically plausible. Domain-informed hierarchical structured sparsity is shown to enhance both model interpretability and generalization performance, although these statistical-learning goals are typically in conflict.

The present approach has important advantages over previous neuroimaging studies that capitalized on dimensionality reduction to harness the curse of dimensionality. They often used preliminary region-wise pooling functions or regression against network templates for subsequent supervised

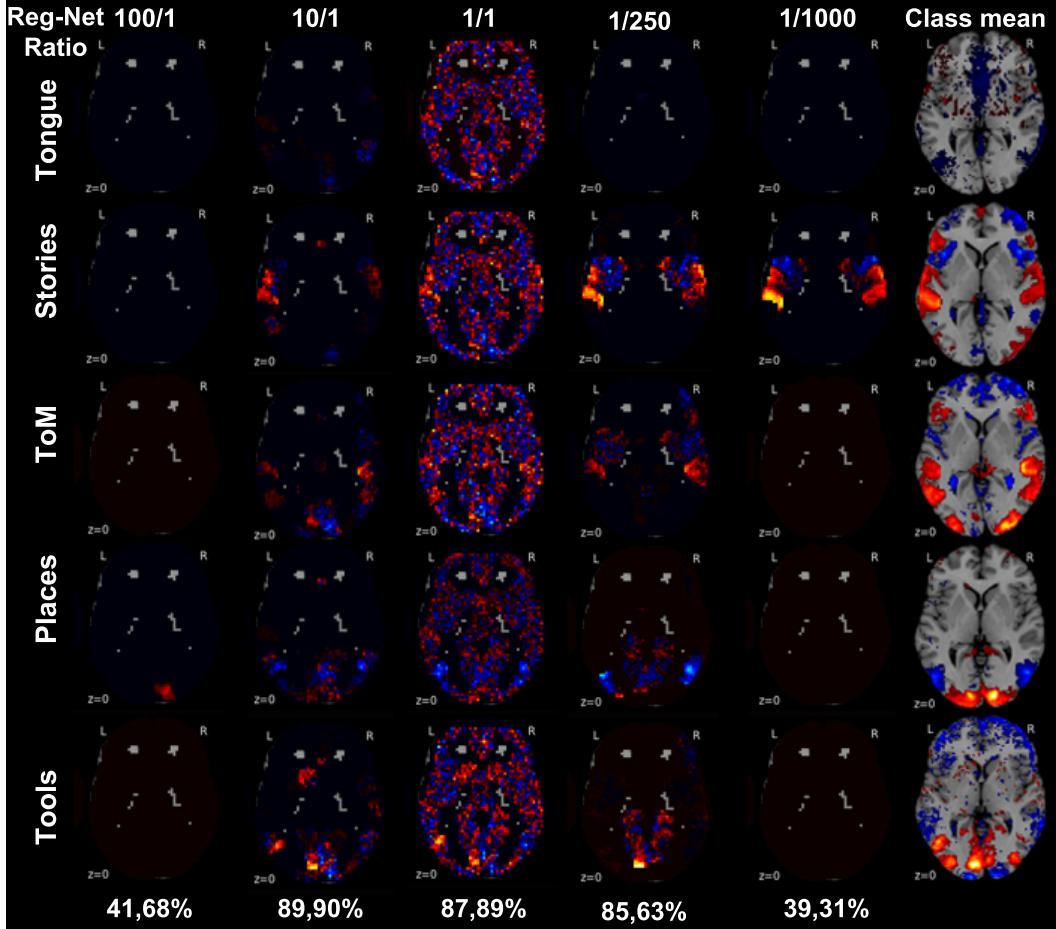


Figure 4: Support recovery as a function of region and network emphasis. The relative impact of the region and network priors on model selection is systematically varied against each other. This region-network ratio (*upper fractions*) weighted voxel groups to privilege sparse models in function space that acknowledge known brain region neighborhoods (*left columns*) or known brain networks neighborhoods (*right columns*). Among the 18 classes, the model weights are shown for the psychological tasks (*from top to bottom*): tongue movement, listening stories, taking somebody else’s perspective (ToM, “theory of mind”), as well as viewing locations and tools. The 18-class out-of-sample accuracy is indicated on the *bottom* and the class-wise mean neural activity (*right-most column*, thresholded at the 75th percentile). Different emphasis on regions versus networks in hierarchical structured sparsity can yield comparable model performance. Favoring region versus network structure during model selection recovers complementary aspects of the neural activity pattern. Equal region and network emphasis yields more dispersed, less interpretable predictive model choices.

learning on the aggregated feature space. Such lossy two-step approaches of feature engineering and inference *i*) can only account for either functional specialization or functional integration of brain organization, *ii*) depend on the ground truth being a region or network effect, and *iii*) cannot issue individual coefficients for every brain voxels. Hierarchical region-network sparsity overcomes these shortcomings by estimating individual voxel contributions while benefitting from their functional segregation and integration to restrict statistical complexity. Viewed from the bias-variance tradeoff, our modification to logistic regression estimators entailed a large decrease in model variance but only a modest increase in model bias. Viewed from the Vapnik-Chervonenkis dimensions, this entailed a healthy decrease in the complexity capacity of the prediction model with a higher chance of generalizing to unobserved data.

In the future, region-network sparsity priors could be incorporated into various pattern-learning methods in systems neuroscience. This includes supervised methods for whole-brain classification and regression with one or several target variables. The principled regularization scheme could even inform unsupervised structure discovery methods, such as principal component analysis [29] and k-means clustering [50]. Additionally, model regularization by hierarchical structured sparsity could be extended from the spatial domain of neural activity to priors of coherent spatiotemporal activity structure [22]. Ultimately, successful high-dimensional inference is an important prerequisite for predicting diagnosis, disease trajectories, and treatment responses in personalized psychiatry and neurology.

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