

FuncISH: learning a functional representation of neural ISH images

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ABSTRACT

Motivation: High-spatial resolution imaging datasets of mammalian brains have recently become available in unprecedented amounts. Images now reveal highly complex patterns of gene expression varying on multiple scales. The challenge in analyzing these images is both in extracting the patterns that are most relevant functionally and in providing a meaningful representation that allows neuroscientists to interpret the extracted patterns.

Results: Here, we present *FuncISH*—a method to learn functional representations of neural *in situ* hybridization (ISH) images. We represent images using a histogram of local descriptors in several scales, and we use this representation to learn detectors of functional (GO) categories for every image. As a result, each image is represented as a point in a low-dimensional space whose axes correspond to meaningful functional annotations. The resulting representations define similarities between ISH images that can be easily explained by functional categories. We applied our method to the genomic set of mouse neural ISH images available at the Allen Brain Atlas, finding that most neural biological processes can be inferred from spatial expression patterns with high accuracy. Using functional representations, we predict several gene interaction properties, such as protein–protein interactions and cell-type specificity, more accurately than competing methods based on global correlations. We used *FuncISH* to identify similar expression patterns of GABAergic neuronal markers that were not previously identified and to infer new gene function based on image–image similarities.

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Supplementary information: Supplementary data are available at *Bioinformatics* online.

1 INTRODUCTION

In recent years, high-resolution expression data measured in mammalian brains became available in quantities and qualities never witnessed before (Henry and Hohmann, 2012; Lein *et al.*, 2007; Ng *et al.*, 2009), calling for new ways to analyze neural gene expression images. Most existing methods for bio-imaging analysis were developed to handle data with different characteristics, like *Drosophila* embryos (Frise *et al.*, 2010; Peng *et al.*, 2007; Pruteanu-Malinici *et al.*, 2011) or cellular imagery (Coelho *et al.*, 2010; Peng *et al.*, 2010). The mammalian brain, composed of billions of neurons and glia, is organized in highly complex anatomical structures and poses new challenges for analysis. Current approaches for analyzing brain images are based on smooth non-linear transformations to a reference

atlas (Davis and Eddy, 2009; Hawrylycz *et al.*, 2011) and may be insensitive to fine local patterns like those emerging from the layered structure of the cerebellum or the spatial distribution of cortical interneurons.

Another challenge for automatic analysis of biological images lies in providing human interpretable analysis. Most machine-vision approaches are developed for tasks in analysis of natural images, like object recognition. In such tasks, humans can understand the scene effortlessly and infer complex relations between objects easily. In bio-imaging, however, the goal of image analysis is often to reveal features and structures that are hardly seen even by experts. It is, therefore, important that an image analysis approach provides meaningful interpretation to any patterns or structures that it detects.

Here, we develop a method to *learn functional representations of expression images* by using predefined functional ontologies. This approach has two main advantages, accuracy and interpretability, and it builds on a growing body of work in object recognition in natural images, showing how images can be represented using the activations of a large set of detectors (Deng *et al.*, 2011; Li *et al.*, 2010a, b; Malisiewicz, 2012; Malisiewicz *et al.*, 2011; Torresani *et al.*, 2010). For object recognition, the detectors may include common objects, like a detector for the presence of a chair, a mug or a door. Here, we show how to adapt this idea to represent gene expression images, by training a large set of detectors, each corresponding to a known functional category, like *axon guidance* or *glutamatergic receptors*. Once this representation is trained, every gene is represented as a point in a low-dimensional space whose axes correspond to functional *meaningful* categories.

We describe in Section 2.2 how to learn functional representations in a discriminative way and demonstrate the effectiveness of the approach on *in situ* hybridization (ISH) gene expression images of the adult mouse brain collected by the Allen Institute for Brain Science (Lein *et al.*, 2007). ISH image analysis has been used in the past to infer gene biological functions from spatial co-expression in non-neural tissues (Frise *et al.*, 2010). However, inferring functions based on gene expression patterns in the brain is believed to be hard, as several studies found very low variability between transcriptomic patterns of different brain regions, sometimes even lower than between-subject variability for the same area (Khaitovich *et al.*, 2004, 2005). Neural expression patterns are usually studied using methods that average expression values over a brain region, and this averaging removes fine-resolution spatial information that may differentiate between brain regions. Here, we analyze high-resolution ISH images at several scales, taking into account subtle, even cellular resolution, information for functional inference.

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We find that gene function can indeed be inferred from neural ISH images, particularly in biological processes that are related to neural activities. Our approach detects related genes with better accuracy based on the similarity of their functional representations. Furthermore, these similarities can be explained and interpreted using semantic terms.

2 METHODS

2.1 The data

We used whole-brain, expression-masked images of gene expression measured using ISH, publicly available at the Allen Brain Atlas (www.brain-map.org, also see Supplementary Material). Expression was measured for the entire mouse genome. For each gene, a different adult mouse brain was sliced into 100- μ m thick slices, mRNA abundance was measured and the slice was imaged. The database holds image series for >20K transcripts. Most genes have one corresponding image series, containing ~25 imaged brain slices. Some genes were imaged more than once and have several associated image series. In our analysis, we used the most medial slice for each image series, yielding a typical image size of 8 K \times 16 K pixels. In all, 4823 of the available 21 174 images showed no expression in the brain and were ignored in subsequent analysis, leaving

16351 images representing 15612 genes. We also tested our approach on a larger image set constructed by taking three images for each gene: the medial slice, and lateral slices at 30% and 50% of brain size (from one hemisphere). The results with this three-image set were mixed, and all results reported later in the text are for the one-slice dataset (Supplementary Material). Figure 1 shows examples of images, demonstrating the complexity of neural expression patterns across brain regions and multiple scales. The images analyzed in our study were in gray scale but are shown here as color-coded by expression intensity for better visualization.

2.2 A functional representation of images

We present a method to identify similarities between neural ISH images and to explain these similarities in functional terms.

Our method consists of a *visual phase*, where we transform the raw pixel images into a robust visual representation, and a *semantic phase*, where we transform that *visual representation* using a set of 2081 gene-function detectors. The output of these detectors comprises a higher-order *semantic representation* of the images in a gene-functional space (Fig. 2). Similar two-phase systems have recently been proposed and applied successfully for tasks, such as cross-domain image similarity and object detection in natural images (Deng *et al.*, 2011; Li *et al.*, 2010a, b; Malisiewicz, 2012; Malisiewicz *et al.*, 2011; Torresani *et al.*, 2010).

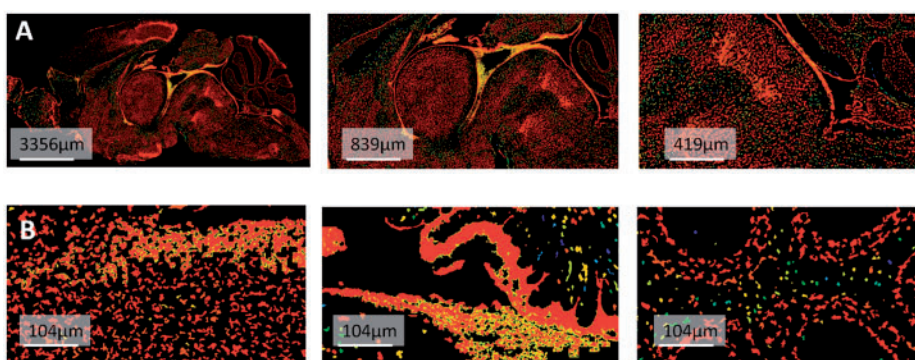


Fig. 1. The raw data. ISH image for the gene *Tuba1* shown (A) at different scales and (B) in three different regions

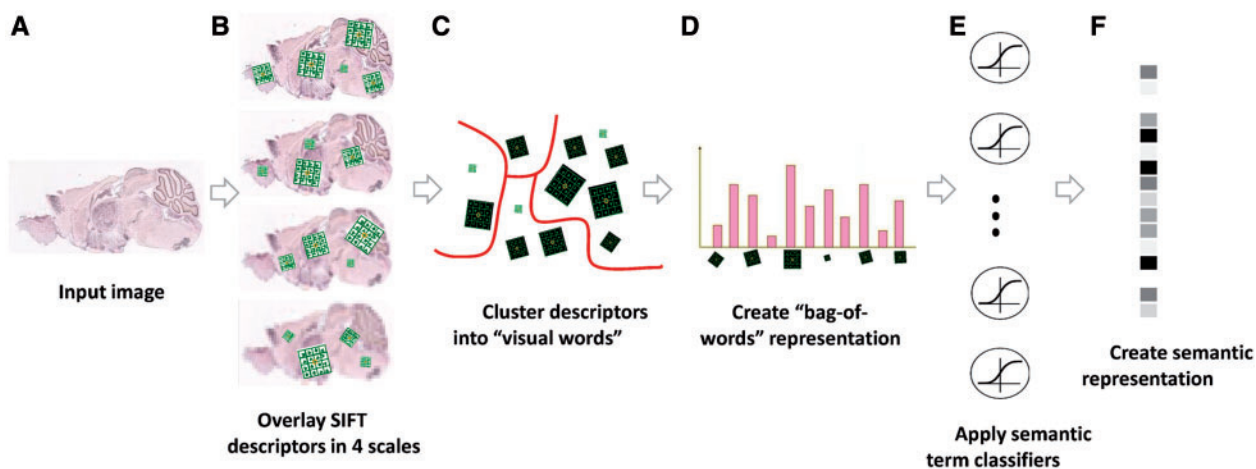


Fig. 2. Illustration of the image processing pipeline. (A) Original image in pixel grayscale indicating level of gene expression. (B) Local SIFT descriptors are extracted from image at 4 resolutions. (C) Descriptors from all 16351 images are clustered into 500 representative ‘visual words’ for each resolution level using k-Means. (D) Each image is represented as a histogram counting the occurrences of visual words. (E) L2-regularized logistic regression classifiers are applied for 2081 GO categories. (F) The final 2081 dimensional image representation

For the first, visual, phase, we first represent each image as a collection of local descriptors using SIFT features (Lowe, 2004). This step aims to address the problem that ISH brain images of the same gene vary significantly in shape and size when measured in different brains (Kirsch *et al.*, 2012). SIFT features are histograms of oriented gradients on a small grid. The resulting image-patch SIFT descriptor is invariant to small rotation and illumination (but not to scale), making imaged-slices from different brains more comparable. We computed SIFT descriptors of dimension 128 extracted on a dense grid spanning the full image (Bosch *et al.*, 2006, 2007; Csurka and Dance, 2004), at four spatial resolutions. In ISH images, different information lies in different descriptor sizes, and we wish that the representation captures spatial patterns both at the level of single cells, micro-circuitry and at the coarser level of distribution of expression across brain layers. To capture information at multiple scales, we used the VLFeat implementation of SIFT (Vedaldi and Fulkerson, 2010), where scale-invariance is not incorporated automatically. Specifically, each image is represented as a collection of ~ 1 M SIFT descriptors, computed by down sampling each image at a factor of 1, 2, 4 and 8. As the descriptors were extracted from high-resolution images, which are mostly dark, many descriptors were completely dark and were discarded.

Next, to achieve a compact non-linear representation of each image, we aggregate the descriptors from all images for a given resolution level and cluster them to form a dictionary of distinct ‘visual words’ per each resolution level. We used the original Lloyd optimization for k-Means with L_2 distance, initializing the centroids by randomly sampling data points. The clustering procedure was repeated multiple times ($n=3$), and the solution with the lowest energy was used. We tested four different dictionary sizes ($k=100, 200, 500$ and 1000), all yielding similar results (Supplementary Material), and we report later in the text results for $k=500$, which obtained slightly higher accuracies. Next, we construct a standard ‘bag-of-words’^{20,21} description of each image. As a result of this process, each image is described by four concatenated 500-dimensional vectors counting how many times each ‘visual word’ appeared in it at a given resolution level. We also added a count of the number of zero descriptors per resolution level, ending up with a 2004-dimensional vector describing each image. Using this approach, similar spatial information from different brain regions is preserved, as opposed to using global correlation-based approaches.

We then turn to the second, ‘semantic’, phase, and represent each image by a set of functional descriptors. Given a set of predefined Gene Ontology (GO) annotations of each gene, we train one separate classifier for each known biological annotation category, using the SIFT bag-of-words representation as an input vector. Specifically, here, we trained a set of 2081 L_2 -regularized logistic regression classifiers [using LIBLINEAR (Fan *et al.*, 2008)] corresponding to biological-processes GO classes that have 15–500 annotated genes (Supplementary Material). We trained the classifiers using two layers of 5-fold cross-validation, performed as follows: the full set of 16 351 gene images was split into five non-overlapping equal sets (without controlling for the number of positives in each split), training the classifiers on four of them and testing performance on the fifth unseen test set of images. This procedure was repeated five times, each time with a different set acting as the test set. All accuracy and other results later in the text are reported for a held-out test set that was not used during training.

To tune the logistic regression regularization hyperparameter, we used a second layer of cross-validation. We repeated the splitting procedure within each of the five training sets, splitting each of them again into five subsets of images, using four for training and the fifth as a validation set. The regularization hyperparameter was selected from the values (0.001, 0.01, 0.1, 1, 10 and 100). At the end of this process, each gene is then represented as a vector of ‘activations’, corresponding to the likelihood that the gene belongs to one functional category, such as ‘forebrain development’ or ‘regulation of fatty acid transport’.

The representation described earlier in the text removes important information about global location in the brain. We, therefore, also tested an approach using spatial pyramids (Lazebnik *et al.*, 2006), where descriptor histograms are computed separately for different parts of the image. Unfortunately, this approach results in feature vectors whose dimensionality was too high for the current dataset and yielded poor classification results (Supplementary Material).

2.3 Similarity between functional profiles

We use two gene–gene similarity measures in this work, taking each gene as a vector of functional category activations. The first, *flat-sim*, is simply the linear correlation of two functional category activation vectors. The second, *GO-sim*, takes into account the known directed acyclic graph (DAG) structure among the functional categories of the GO annotation.

Formally, the *flat-sim* score between a pair of L_2 -normalized feature vectors $a = (a_1 \dots a_m)$ and $b = (b_1 \dots b_m)$ is given by their dot product $flat-sim(a, b) = \sum_{i=1}^m a_i \cdot b_i$. This additive similarity measure allows assessing the contribution of each individual feature to the overall similarity score, by setting the contribution of the feature i (corresponding to GO category i) to $a_i \cdot b_i$. Thus, for each pair of similar images, we can sort the GO categories by order of their contribution to the similarity, providing a semantic interpretation of the correlation.

However, *flat-sim* does not take into account that the activation of some functional categories can be far more informative than others. For example, two genes that share a specific function like ‘negative regulation of systemic arterial blood pressure’ are much more likely to be functionally similar than a pair of genes sharing a more general category like ‘metabolism’. We address this issue by adapting a functional similarity measure between gene products developed by (Schlicker *et al.*, 2006), which we refer to as *GO-sim*. *GO-sim* is designed to give high similarity scores to gene pairs that share many specific and similar functional categories. We treat our model’s functional activations as binary annotations (using a threshold of 0.5) and calculate *GO-sim* as follows.

For each GO category i , we calculate its *information content (IC)* as $IC(i) = -\log_{10} \frac{\#genes\ in\ i}{total\ \#\ of\ genes}$, which measures the specificity of each category. For each pair of categories i and j , we consider the set of their common ancestors $anc(i, j)$ and define $sim_{rel}(i, j) = \max_{k \in anc(i, j)} \frac{2IC(k)}{IC(i)+IC(j)} (1 - 10^{-IC(k)})$. The measure sim_{rel} is symmetric, bounded between 0 and 1, and attains larger values for pairs of categories that are both specific and close to each other in the GO graph.

In our method, each gene is annotated with multiple categories. Naïvely, we could calculate the mean sim_{rel} measure between all pairs of categories, but calculating this mean could give weight to many irrelevant categories and be sensitive to the addition of extra annotations to a gene. Instead, we use a more robust method to measure similarity between two sets of function annotations, developed by (Schlicker *et al.*, 2006). This method relies on the most similar gene pairs, instead of all the pairs. For two **binary** activation vectors $a = (a_1 \dots a_m)$, $b = (b_1 \dots b_m)$ define a matrix $S_{ij} = sim_{rel}(i, j)a_i b_j$. Then we define $sim_{a \rightarrow b} = \frac{1}{m} \sum_{i=1}^m (\max_{j=1 \dots m} S_{ij})$ that measures for each annotation of a its most similar annotation in b and averages across all of a ’s annotations. We similarly define $sim_{b \rightarrow a}$ with the roles of a and b switched, and use it to define $GO-sim = \max(sim_{a \rightarrow b}, sim_{b \rightarrow a})$. To assess the contribution of individual gene functional annotations to the *GO-sim* measure, we look at the category pairs (i, j) corresponding to the highest values of S_{ij} . Each such pair also has its ‘most informative common ancestor’ $MICA(i, j) = \operatorname{argmax}_{k \in anc(i, j)} \frac{2IC(k)}{IC(i)+IC(j)} (1 - 10^{-IC(k)})$. These ancestor functional categories give a succinct interpretation of the similarity between genes a and b .

Computing *GO-sim* for $n=16351$ genes, each with m functional annotations, is computationally burdensome, requiring $O(n^2 m^2)$

operations. In this study, we, therefore, use only 164 brain-related categories of the 2081 functional categories for calculating *GO-sim*.

3 RESULTS

We start with evaluating the quality of the low-dimensional semantic representation that we learned in two aspects: the classification accuracy for individual semantic terms and the precision of our gene–gene similarity measure compared with a spatial correlation-based method. We then take a closer look at discriminative spatial patterns, mapping them back onto raw images. Finally, we use the geometry of the low-dimensional semantic space to infer new gene functions via gene similarities and their interpretations.

3.1 Predicting functional annotations using brain ISH images

We applied FuncISH to 16 K ISH images of 15 K genes, and we mapped each image to a vector corresponding to 2000 GO categories as functional features. We used the area under the ROC curve (AUC) as a measure of classification accuracy. All evaluations were performed on a separate held-out test set. We find that 37% of the GO categories tested yielded a test set AUC value that was significantly above random (permutation test, $P < 0.05$). This was encouraging, as the variability of expression between brain regions was previously shown to be very low (Khaitovich *et al.*, 2004, 2005). This suggests that fine spatial resolution in neural tissues can reveal highly meaningful expression patterns.

Which functional categories can be best predicted by ISH images? Table 1 lists the top 15 GO categories that achieved the best test-set AUC classification scores. Interestingly, these include mostly biosynthesis/metabolism processes and neural processes. To further test whether neural categories achieve higher classification values based on neural expression patterns, Figure 3 compares the AUC scores of 164 categories related to the nervous system with the AUC scores of the remaining categories. As expected, neural GO categories receive significantly higher AUCs (Wilcoxon, $P < 10^{-38}$), with 69% of categories yielding significantly above random AUC values.

These AUC values suggest that when a gene is represented as a feature vector of classifiers activations, many of the features carry a meaningful signal. The axes of the new low-dimensional representation correspond to functional properties of each gene, linking functions of the genes to the geometry of the space in which they are embedded.

3.2 Comparison with Neuroblast, the ABA image-correlation tool

How well does FuncISH compare with other methods suggested for finding similarity between these images? We compared our results with *NeuroBlast*, a method to detect image–image similarities available on the ABA website (Hawrylycz *et al.*, 2011). This method uses a non-linear mapping of the images to a reference anatomical atlas to apply voxel–voxel correlation between the images.

To evaluate the quality of the similarity measure, we used three sets of pairwise relations as evidence of gene relatedness: (i) markers of known cell types (Cahoy *et al.*, 2008), such

Table 1. The GO categories classified with highest test-set AUC values

GO ID	GO category name	No. of genes	AUC
GO:0060311	Negative regulation of elastin catabolic process	17	1
GO:0042759	Long-chain fatty acid biosynthetic process	23	0.98
GO:0009449	γ -Aminobutyric acid biosynthetic process	20	0.96
GO:0009448	γ -Aminobutyric acid metabolic process	23	0.96
GO:0032348	Negative reg. of aldosterone biosynthetic process	21	0.94
GO:2000065	Negative regulation of cortisol biosynthetic process	21	0.94
GO:0043206	Fibril organization	23	0.94
GO:0031947	Negative reg. of glucocorticoid biosynthetic process	22	0.94
GO:0042136	Neurotransmitter biosynthetic process	23	0.94
GO:0022010	Central nervous system myelination	29	0.89
GO:0008038	Neuron recognition	20	0.87
GO:0042220	Response to cocaine	30	0.87
GO:0050919	Negative chemotaxis	16	0.86
GO:0042274	Ribosomal small subunit biogenesis	15	0.86
GO:0016486	Peptide hormone processing	17	0.85

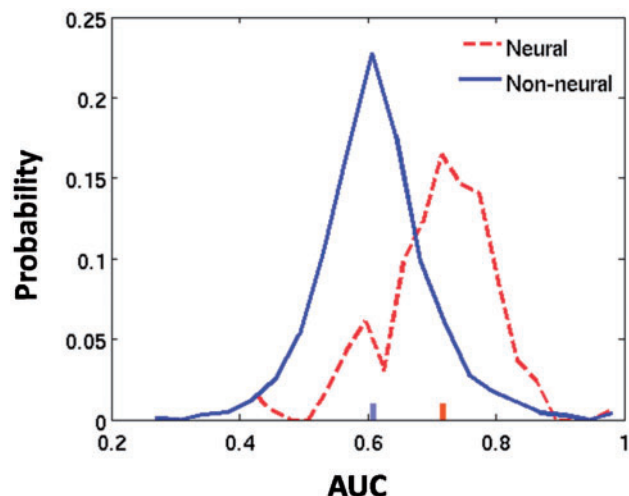


Fig. 3. AUC scores for GO categories related to the nervous system (dashed, red) and the remaining categories (solid, blue). AUC scores are significantly higher for neural categories (Wilcoxon test, $p < 10^{-38}$). The red and blue ticks indicate the median of each set

as astrocytes or oligodendrocytes; (ii) occurrence in the same KEGG pathway (Kanehisa, 2002); and (iii) a set of known protein–protein interactions taken from *IntAct* (Kerrien *et al.*, 2012). For each of the 16 531 genes, we ranked the 100 most similar genes according to four different similarity measures: (i) FuncISH *GO-sim*, (ii) FuncISH *flat-sim*, (iii) cosine similarity between the SIFT bag-of-words representations (Fig. 2D) and (iv) the ABA *NeuroBlast* tool. For each of the pairwise relations (cell-type markers, KEGG pathway and PPIs), we plot the mean fraction of relations retrieved at the top-K most similar genes (precision-at-k), a standard method in information retrieval (Manning and Raghavan, 2009). Figure 4 shows that for all three validation labels, FuncISH *GO-sim* provides superior precision for the top 10 ranked similar genes. The superior precision of *GO-sim* over *flat-sim* is presumably because

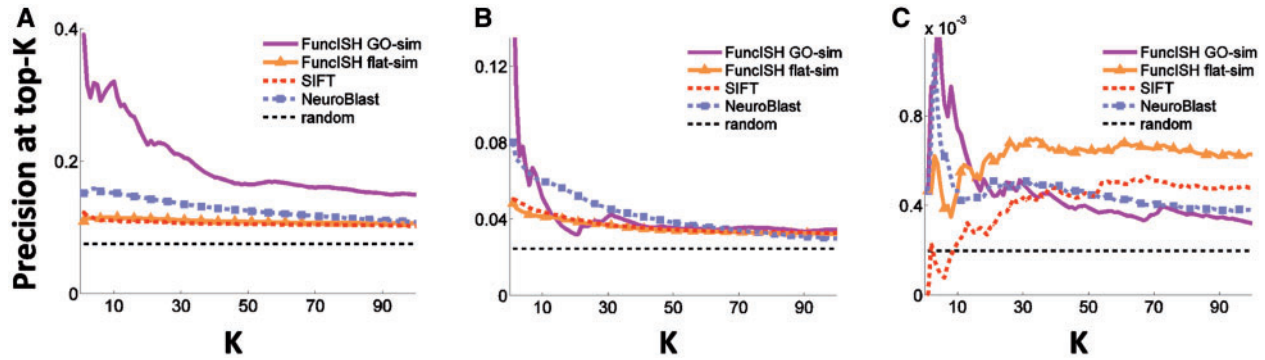


Fig. 4. Precision at top-K for similarity defined by (A) cell type marker (B) KEGG pathways (C) protein-protein interaction. Precision was measured using functional representations (*FuncISH*, purple lines for *GO-sim*, orange for *flat-sim*), SIFT (red) and NeuroBlast (blue)

GO-sim weighs categories more correctly and also possibly because GO-sim was limited to brain-related categories that tend to be more accurately predicted (Fig. 3). On the other hand, we see that NeuroBlast outperforms flat-sim in most cases.

3.3 Identifying and explaining similarities between GABAergic neuron markers

We now turn to a deeper look into the similarity predictions. Interestingly, the highest classification scores were achieved for the neural-related categories *GABA biosynthetic process* and *GABA metabolic process* (shown in Table 1), implying that our algorithm can identify spatial patterns of GABAergic neurons. A prominent member of the GABAergic neuron marker family is *parvalbumin B (Pvalb)*, which encodes for a calcium-binding protein. We examined the genes that are most similar to *Pvalb*, and we found that another GABAergic neuronal marker and a calcium-binding protein, *calbindin D28K (Calb1)*, is at the top 15 most similar gene lists for all associated image series. *Pvalb* and *Calb1* belong to a family of cellular Ca^{2+} buffers in GABAergic interneurons. The third member in this family is calretinin (*Calb2*). Looking at the similarity rank of *Calb1* and *Calb2*, *Calb2* ranks at the top 2 percentile (of 16 351 images in the dataset) at 16 of 17 cases. Similarities between these three genes were not identified by NeuroBlast. This may be because NeuroBlast uses spatial correlation measures that produce results heavily reliant on the spatial location of expression, whereas FuncISH can identify patterns that can appear in different regions of the brain. A major benefit of representing genes in the functional embedding space is that similarities between genes can be ‘explained’ in functional terms. *Calb1*, *Pvalb* and *Calb2* are all involved in regulation of synaptic plasticity (Schwaller, 2012). When looking at the semantic interpretations explaining the similarities between the genes, 6 of the top 10 GO categories are indeed directly related to synaptic plasticity, such as ‘synaptic transmission’, ‘regulation of synaptic plasticity’ and ‘learning’.

3.4 Finding important spatial patterns in different scales using SIFT ‘visual words’

A major advantage of representing ISH images with SIFT descriptors is the ability to point directly to spatial patterns in

these complex images. Although their name suggest differently, SIFT descriptors at several scales capture different types of patterns. Figure 5 shows three visual words for each of the four scales, selected as the visual words that contributed most to classification. Scale invariance is often assumed when analyzing natural images, as objects are photographed at varying distances. ISH images, however, contain distinctive information in the different scales. As Figure 5 demonstrates, the four sizes of visual words correspond to grids capturing different neural entities. The smallest descriptors cover an actual area of $36 \times 36 \mu m^2$ and capture fine-scaled information, such as cell shapes and cell densities; the medium-size discriminative descriptors of $72 \times 72 \mu m^2$ tend to trace thinner cell layers; larger descriptor sizes of $144 \times 144 \mu m^2$ and $288 \times 288 \mu m^2$ can cover large and intricate patterns of a mixture of cells and cell types in a tissue. Interestingly, the four visual words with the highest contribution to classification were the words counting the zero descriptors in each scale. This means that the highest information content lies in ‘least informative’ descriptors, and that overall expression levels (‘sparseness’ of expression) are important factors in functional prediction of genes based on their spatial expression. Our method presents a new representation of ISH imagery as SIFT descriptors, and using multiple scales allows revealing the multi-resolution nature of the images.

Which scale carries the most meaningful signal for functional prediction? Figure 5E shows the mean absolute value of visual words weights in every scale for all GO categories, showing that all scales contribute significantly to the scores, with the medium contributing most.

Figure 5A–D shows descriptors that contributed to classification of **all** the categories. Furthermore, each GO category has its own visual words that are important to its classification, and looking into their details reveals spatial properties that are unique to specific biological processes.

As an interesting example of this effect, we considered the gene *adducin β (Add2)*. *Add2* is annotated to several GO categories, including ‘positive regulation of protein binding’ and ‘actin filament bundle assembly’. Figure 6 overlays the top weighted visual words of the two categories over the *Add2* ISH image. It is easy to see that the descriptors important for classification of ‘actin filament bundle assembly’ are much smaller than those important for classification of the more general

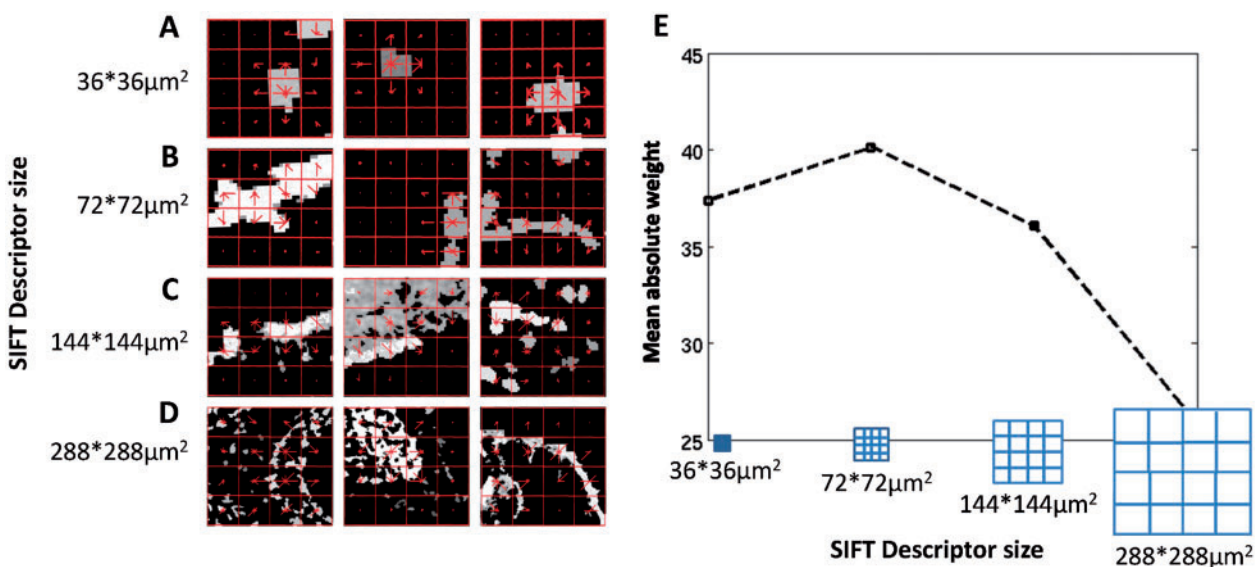


Fig. 5. Representing ISH images with visual words. (A, B, C, D) The three visual words with highest absolute weight (averaged over all categories) at each scale. The SIFT descriptors (red grid) are plotted on top of each panel. The histogram of oriented gradients used in the SIFT descriptor is plotted in the center of each element of the grid, as a set of red lines, where the length of the line correspond to the magnitude of the gradient in its direction. (E) Mean absolute weight for the four scales of visual words calculated over classifiers for all categories

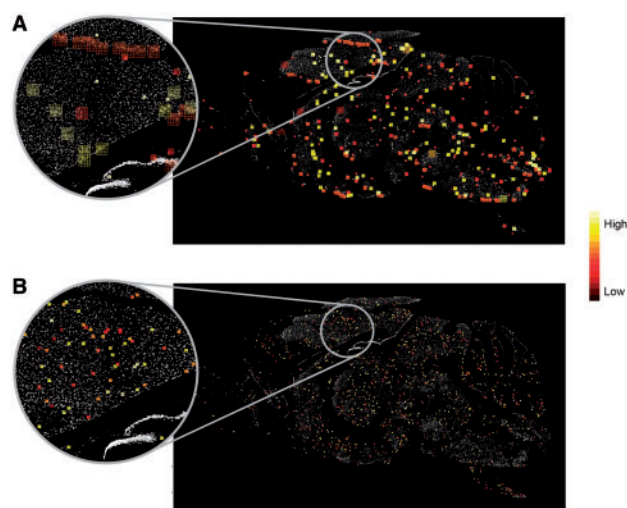


Fig. 6. The visual words important in classifying Add2 GO categories are overlaid on the Add2 ISH image. Larger descriptors are needed for the classification of ‘regulation of protein binding’ (A), while the discriminative visual words for ‘actin filament bundle assembly’ (B) are much smaller, capturing properties such as cell shapes. The descriptors are color-coded by their importance in classification, highest importance is in bright yellow

category ‘positive regulation of protein binding’ (*t*-test, $P < 10^{-17}$). This implies that small-scaled features, such as specific cell shapes, are important to identify genes related to actin filament bundle assembly processes. Actin assemblies are important for the navigation of neural growth cones, by re-orienting growth cones away from inhibitory cues (Challacombe *et al.*, 1996). Representing the images with histograms of oriented gradients could capture tiny differences in cell shapes that

are in the process of synapse formation, a developmental process occurring continuously throughout adulthood (Vidal-Sanz *et al.*, 1987).

3.4 Inferring new gene functions via explainable similarities

We now demonstrate how the semantic representation learned by FuncISH can be used to propose new gene functional annotations. Consider as an example the gene *synaptodin 2* (*Synpo2*) that is known to bind actin, but otherwise has little known associated information. FuncISH can be used to propose functional annotations for *synpo2* by looking at the genes that are similar to *Synpo2* and considering both the GO functions that contribute to this similarity and the spatial pattern of expression.

First, we find that *Synpo2* is similar to two other genes *Npepps* and *Rasa4*, but for different reasons (the list of top five semantic explanations for these similarities is shown in Table 2). *Npepps* is an aminopeptidase that is active specifically in the brain (Hui, 2007), and the similarity between *Synpo2* and *Npepps* is explained by processes related to protein processing, such as ubiquitination and protein proteolysis. At the same time, *Rasa4* is a GTPase-activating protein that suppresses the Ras/mitogen-activated protein kinase pathway in response to Ca^{2+} (Vigil *et al.*, 2010), and the similarity between *Synpo2* and *Rasa4* is explained by high-level neural processes, such as axon guidance or synaptic transmission.

Interestingly, *Synpo2* and *Rasa4* are expressed in different brain regions: looking at their spatial expression patterns reveals that *Synpo2* is expressed exclusively in the thalamus, whereas *Rasa4* is expressed in olfactory areas. Therefore, their similarity is not in their global expression patterns across regions, but rather in local spatial patterns. This could reflect expression in

Table 2. Top 10 GO annotations explaining the similarities between the gene *Synpo2* and *Npepps* (left column) and *Rasa4* (right column)

<i>Synpo2</i> – <i>Npepps</i>		<i>Synpo2</i> – <i>Rasa4</i>	
GO ID	GO name	GO ID	GO name
GO:0070646	Protein modification by small protein removal	GO:0006836	Neurotransmitter transport
GO:0006412	Translation	GO:0051970	Negative regulation of transmission of nerve impulse
GO:0016567	Protein ubiquitination	GO:0050805	Negative regulation of synaptic transmission
GO:0051603	Proteolysis involved in cellular protein catabolic process	GO:0007411	Axon guidance
GO:0032446	Protein modification by small protein conjugation	GO:0031645	Negative regulation of neurological system process

similar cell types or tissues that exhibit similar spatial distribution at different brain regions. *Npepps* is more ubiquitously expressed in the brain, and it is located in the thalamic area where *synpo2* is expressed. The co-location of *Synpo2* and *Npepps* suggests they could be participating in similar biological processes in these areas, possibly in protein-modification processes as suggested by the list of top explanations for the similarity.

4 SUMMARY

We present *FuncISH*—a method to learn functional representations of neural ISH images, yielding an interpretable measure of similarity between complex images that are difficult to analyze and interpret. Using *FuncISH*, we successfully infer ~700 functional annotations from neural ISH images, and we use them to detect gene–gene similarities. This approach reveals similarities that are not captured by previous global correlation-based methods, but it also ignores important global location information. Combining local and global patterns of expression is, therefore, an important topic for further research, as well as the use of more sophisticated non-linear classifiers, such as kernel-SVM, for creating better representations. Importantly, *FuncISH* provides semantic interpretations for similarity, enabling the inference of new gene functions from spatial co-expression.

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