

Anatomical connectivity patterns predict face selectivity in the fusiform gyrus

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A fundamental assumption in neuroscience is that brain structure determines function. Accordingly, functionally distinct regions of cortex should be structurally distinct in their connections to other areas. We tested this hypothesis in relation to face selectivity in the fusiform gyrus. By using only structural connectivity, as measured through diffusion-weighted imaging, we were able to predict functional activation to faces in the fusiform gyrus. These predictions outperformed two control models and a standard group-average benchmark. The structure–function relationship discovered from the initial participants was highly robust in predicting activation in a second group of participants, despite differences in acquisition parameters and stimuli. This approach can thus reliably estimate activation in participants who cannot perform functional imaging tasks and is an alternative to group-activation maps. Additionally, we identified cortical regions whose connectivity was highly influential in predicting face selectivity within the fusiform, suggesting a possible mechanistic architecture underlying face processing in humans.

A fundamental assumption in neuroscience is that function is deeply rooted in anatomical structure, such as extrinsic connectivity. A region's connectivity pattern determines both the information available as inputs from other regions and its output and influence on other areas. Indeed, changes in connectivity have been shown to occur at the boundaries of functionally defined regions that can be identified through cytoarchitectonics (supplementary motor area versus pre-supplementary motor area)¹. If anatomical connectivity is important for functional operations, then variation in connectivity should correspond with and predict variation in function, even in regions that are not at present anatomically definable or not spatially consistent across the population. This intuitive claim has not yet been formally explored, though various frameworks for such an analysis have been suggested².

In the absence of any extra information, can structural connectivity accurately predict the location and degree of the functional response in the brain? The extrinsic connectivity pattern of a structure may contain sufficient information to predict the extent to which each voxel will respond to a given functional contrast. This hypothesis could be tested using a functional contrast that consistently elicits robust responses, and constrained to an anatomical structure that reliably encapsulates such responses across participants, even if they vary spatially within the region.

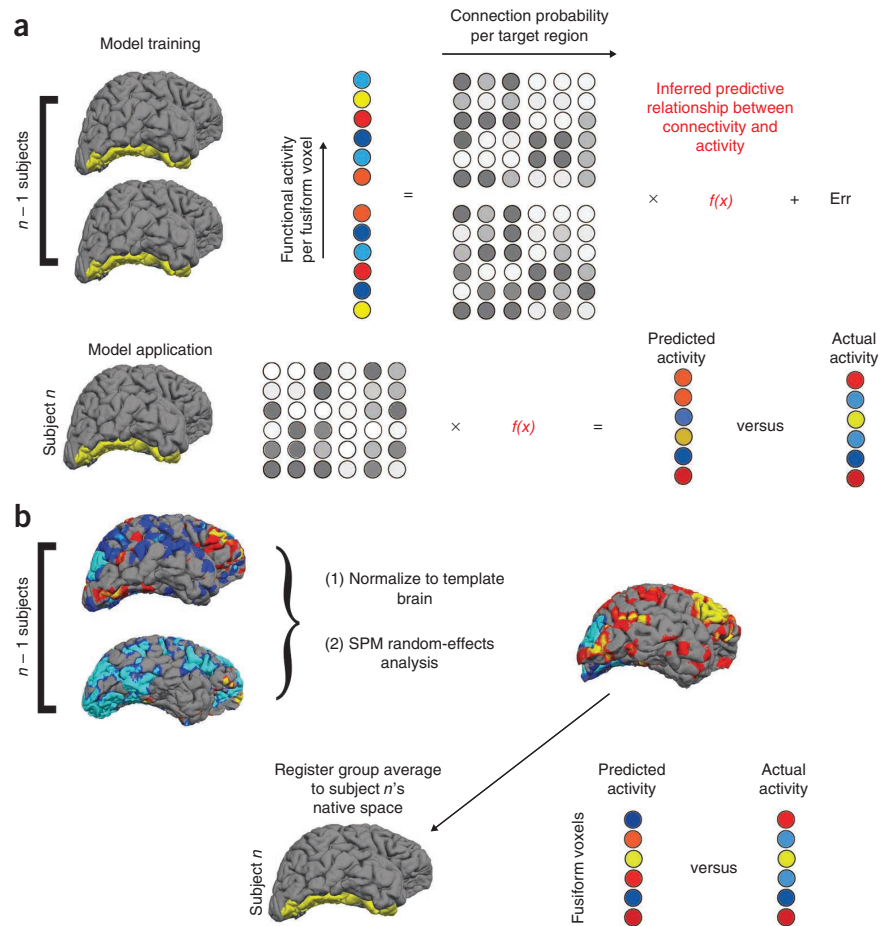
Regions involved in face processing may be well suited for directly testing this conjecture, given their posited specificity of function and replicability across brain imaging techniques, participants and species. A dedicated network of brain regions has been consistently reported to selectively respond to faces, as revealed by functional

magnetic resonance imaging (fMRI)^{3,4}, single-unit recordings^{5,6} and microstimulation⁷. The most robust and selective component of this network is in the fusiform gyrus⁸, in a functionally defined region that is selectively activated in response to faces relative to objects⁹ or scenes¹⁰. This region is typically larger and more reliably observed in the right fusiform and is known as the fusiform face area (FFA). This is consistent with a wide range of evidence that most aspects of face perception are right-hemisphere dominant in the human brain^{8,11,12}. Further, damage to the right fusiform disproportionately impairs face recognition, sometimes even without disturbing other stimulus categories^{13,14}. Given that it is the right fusiform that best responds to faces across participants (for example, refs. 8,15), we chose this region as a testing ground for modeling brain activity as a function of structural connectivity.

A purely structural substrate of face-selective cortices has not yet been established, possibly owing to complications in relating classic approaches of connectivity (such as histological tract-tracing) with functional localization in the same individual. However, diffusion-weighted imaging (DWI), an MRI technique that measures the propensity of water to travel along myelinated axons, can be used to estimate brain connectivity *in vivo*^{16,17}, which can be analyzed alongside fMRI data for the same individual. Using a probabilistic tractography algorithm, we defined the connection probability of each right fusiform voxel (seeds) to all other anatomically parcellated regions (targets) (see **Supplementary Figs. 1 and 2** for exemplar pathways¹⁸⁻²⁰). For the same participants, we calculated the functional activation of faces relative to scenes for each voxel in the fusiform. We then analyzed the relationship between functional activation in the fusiform

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Figure 1 Schematic model design. (a) Linear regression models were trained on all but one participant's data in group 1. The 22 participants' fMRI data for each voxel in the fusiform gyrus (show in yellow on the brain images) are depicted by circles that are color-coded from red to blue, illustrating their responses to the contrast of faces > scenes. Each voxel's corresponding connection probabilities (for the connectivity model) or Euclidian distances (for the distance model) to each target brain region are depicted by the grayscale circles. The fMRI data and connectivity or distance data from each fusiform voxel for the 22 participants were used to train the model, and the resulting model, $f(x)$, was applied to the remaining participant's connectivity or distance data, resulting in predicted fMRI values for each fusiform voxel. The predicted values were then compared to that participant's observed fMRI values and the MAE calculated for each participant. This LOOCV was done iteratively through all the participants, such that each participant had a predicted fMRI image based on a regression from all the other participants. Err, model residual error. (b) Similarly, a LOOCV procedure was performed for the group-average model, but rather than training a linear regression, each participant's whole-brain fMRI data were spatially normalized into MNI space and the data from participants were superimposed to create composite maps; a t -statistic image was then generated from the random-effects analysis. This image was registered to the remaining participant's native space, and the fusiform gyrus was extracted. This activation predicted on the basis of a group analysis was then compared to that participant's observed activation, and an MAE was computed per voxel. SPM, statistical parametric mapping.



and its connection probabilities with the rest of the brain, through a multivariate, voxel-by-voxel approach. This approach allowed us to directly test the conjecture that although the locations of face-selective voxels are variable across the population, their extrinsic connections vary systematically with function in each individual, such that the connection patterns alone can predict functional activation.

Specifically, we used a least-squares linear regression to model the relationship between each fusiform voxel's connection probabilities and its functional activation by means of a leave-one-subject-out cross-validation approach, or LOOCV (Fig. 1). The resulting model was then applied to the remaining participants' connectivity data, and prediction accuracies were tested against two control models and a benchmark model built from a functional group average. The group average is commonly used as a way to build face-selective regions of interest (ROIs) in fMRI studies²¹ and thus provides a standard that a connectivity-based method should meet. The control models, designed from random permutations and Euclidian distance (see Online Methods), were implemented to evaluate against potential confounds.

To assure that a model is not overly fit to the population it was built from, it is good practice to design a model built from all the participants in the LOOCV and apply it to a separate pool of observations naive to the model-building procedure²². We applied such a model to an independent group of participants from a separate study. Analysis of this second group provided further examination of the generalizability of the models, as their data were acquired

with different DWI scan parameters and a different functional task from the first group of participants.

RESULTS

Comparisons between connectivity and control models

After an initial analysis determined that the data possessed sufficient structure for its use in prediction (Supplementary Table 1), we proceeded to build the connectivity models and their controls. A linear regression was trained on the connectivity and fMRI data (contrast for faces > scenes) for all participants but one, and the model was applied to the remaining participant's connectivity data to make predictions of this participant's fMRI data in the right fusiform gyrus; this was done iteratively across all participants. We calculated the absolute error per voxel as the difference between the predicted and actual fMRI images, and mean absolute error (MAE; Table 1) as a measure of accuracy.

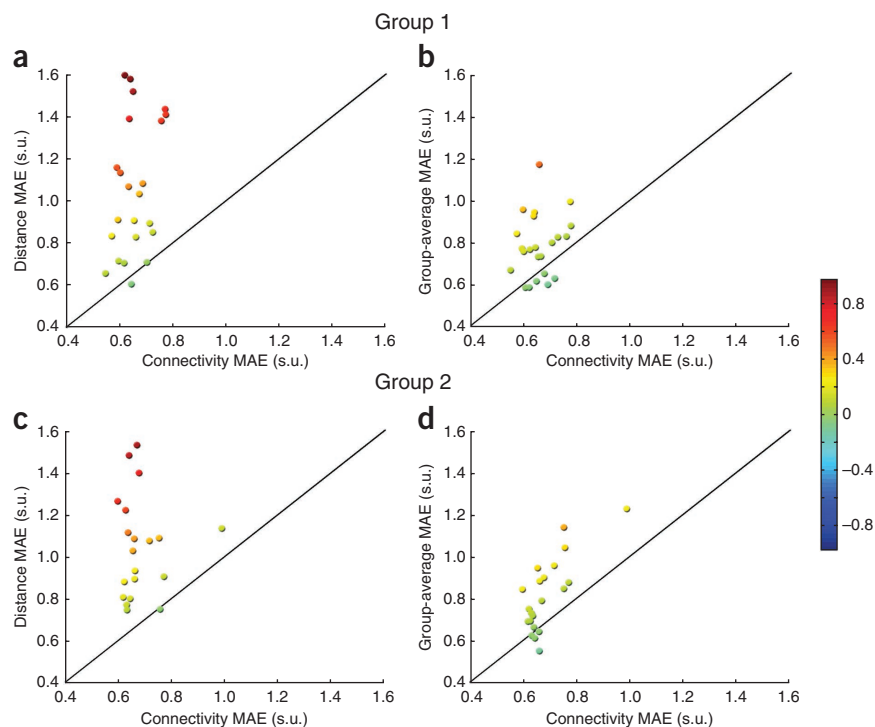
Next, we performed random permutation tests²³ to statistically assess the performance of the connectivity model. We built models

Table 1 MAE for voxels in the fusiform gyrus across subjects

	Group 1	Group 2
Connectivity	0.65 ± 0.013	0.68 ± 0.019
Permutation	0.77 ± 0.008	NA
Distance	1.06 ± 0.066	1.05 ± 0.051
Group average	0.78 ± 0.031	0.82 ± 0.039

Prediction errors for models based on connectivity, mean random permutations, distance and group average are given ± s.e.m. in standard units. NA, not applicable.

Figure 2 Benchmark comparisons per participant. MAEs in standardized units (s.u.) from the connectivity-based predictions are plotted against distance or group-average MAEs for each participant. Participants above the unity line thus have higher (worse) MAEs for the benchmark than for the connectivity-based model. Colors reflect the difference between the connectivity-based model and the benchmark; hotter colors indicate better performance of the connectivity-based model. (a) For 21 of 23 participants in group 1, the distance-based predictions had higher (worse) MAEs than connectivity-based predictions, and no participants' functional activation was better predicted by distance than by connectivity. (b) The connectivity-based model predicted actual fMRI activation with fewer errors than the group average for 17 of 23 participants, whereas 2 participants' functional activation was better predicted by the group average than by connectivity. (c) For 18 of 21 participants in group 2, connectivity better predicted actual activations than distance, whereas no participant's functional activation was better predicted by distance than by connectivity. (d) For group 2, 16 of 21 participants had lower MAEs with the connectivity model, whereas 1 participant had lower MAEs with the group-average model.



designed from the same data but with shuffled pairings between connectivity and functional responses, and, by repeating this process 5,000 times, we generated a distribution of accuracies from random models for each individual. Relative to this distribution, the connectivity models successfully predicted functional selectivity across voxels in 22 out of 23 participants' fusiform gyri at a threshold of $P < 0.001$.

The distance from a seed voxel to a target region may potentially bias the connection estimates because local connections are believed to be more probable than distant ones^{24,25}. In addition, the lateral wall of the fusiform gyrus tends to be face selective, whereas the medial wall tends to be more scene selective. The connectivity model could therefore rely on the relative distance of each voxel to each target, which is basically a high-dimensional spatial coordinate frame. To ensure that the results of the connectivity model were not driven by such unintended relationships, we generated distance control models using the same LOOCV method. These models were designed identically to the connectivity model, with the exception that they used Euclidian distance of the fusiform voxels to other brain regions' center

of mass, rather than their connectivity. The distance models thus use the same number of predictors as the connectivity models and serve as controls for possible overfitting.

We directly compared the performance of the connectivity and distance models, both across participants (on the basis of MAE) and within participants (on the basis of absolute error). Across participants, the connectivity model was significantly more accurate than the distance model (two-tailed t -test of connectivity MAE versus distance MAE, $t_{22} = -6.44$, $P = 1.75 \times 10^{-6}$). A direct comparison of the error per voxel at the individual subject level revealed that the connectivity-based predictions were significantly better than distance in 21 of 23 participants at a threshold of $P < 0.001$ (Fig. 2a).

Comparisons to group-average models

We also performed a group analysis on the whole-brain fMRI data in an iterative LOOCV fashion: a random effects test was performed on the contrast images for faces > scenes for all but one participant (Fig. 1b). The resulting group average was registered to the native anatomical coordinates of the participant left out of the group analysis, and prediction errors were calculated for the right fusiform. Because group analyses are standard in neuroimaging, we chose them as benchmark models that connectivity-based predictions should meet or exceed to be considered useful.

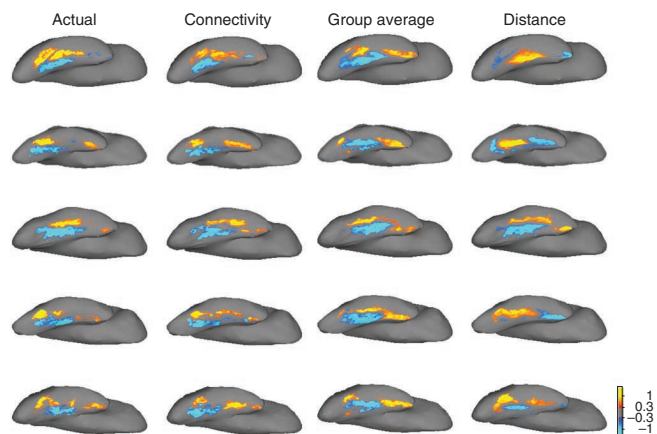


Figure 3 Actual and predicted fMRI activation to faces > scenes in the fusiform gyrus of five example participants. For each participant, actual and predicted activation images (t -statistic values for faces > scenes) were up-sampled from the DWI structural image (where all the analyses were performed) to the same participant's structural scan and were projected onto the participant's inflated brain surface. Each row is a single participant; the leftmost column displays the actual fMRI activation pattern in the right fusiform gyrus. The remaining columns illustrate, from left to right, predicted fMRI images from connectivity, group average and distance. Color scale in standardized units (s.u.).

Table 2 List of target regions that made a statistically significant contribution to the final connectivity model

Target	Coefficient	C.I.	Target	Coefficient	C.I.
R inferior temporal	0.1490	0.137 0.161	R lingual	-0.3868	-0.399 -0.375
R lateral occipital	0.0978	0.085 0.111	R parahippocampal	-0.1373	-0.149 -0.125
R cerebellum	0.0883	0.076 0.100	L fusiform	-0.0638	-0.086 -0.041
R superior temporal	0.0809	0.062 0.100	R inferior parietal	-0.0636	-0.081 -0.046
L cerebellum	0.0714	0.050 0.093	L lingual	-0.0634	-0.077 -0.050
L inferior temporal	0.0635	0.049 0.078	L parahippocampal	-0.0425	-0.058 -0.027
R entorhinal	0.0466	0.036 0.058	L isthmus cingulate	-0.0417	-0.055 -0.028
R middle temporal	0.0293	0.015 0.043	R post central	-0.0396	-0.059 -0.020
R pars opercularis	0.0271	0.007 0.048	R isthmus cingulate	-0.0378	-0.051 -0.025
R thalamus	0.0249	0.007 0.043	R lateral orbitofrontal	-0.0340	-0.052 -0.016
L pericalcarine	0.0215	0.002 0.041	R pars triangularis	-0.0289	-0.055 -0.002
L middle temporal	0.0199	0.003 0.036	R hippocampus	-0.0266	-0.041 -0.012
L temporal pole	0.0145	0.002 0.027	L hippocampus	-0.0240	-0.039 -0.009
L lateral orbitofrontal	0.0130	0.001 0.025	R caudal anterior cingulate	-0.0234	-0.043 -0.004
			L superior temporal	-0.0232	-0.042 -0.004
			R amygdala	-0.0219	-0.033 -0.010
			L paracentral	-0.0206	-0.033 -0.008
			Brain stem	-0.0200	-0.035 -0.005
			L amygdala	-0.0152	-0.027 -0.004

Regions are listed along with their model coefficients and 95% confidence intervals (C.I.). Positive predictors are listed on the left, negative predictors on the right. L, left; R, right.

We compared model performance and found that the connectivity-based predictions were statistically better than the group average, across participants (two-tailed t -test of connectivity MAE versus group average MAE, $t_{22} = -4.01$, $P = 5.94 \times 10^{-4}$). The connectivity model was significantly more accurate than the group average for 17 of 23 participants at $P < 0.001$, whereas the converse was true for only 2 participants (Figs. 2 and 3). For the remaining 4 participants, the models were not significantly different.

Final connectivity models

We then applied the connectivity and distance models generated by group 1 to a separate group of 21 participants, whose connectivity and functional data were naive to the models. These analyses were performed in a similar manner, except that the regressions were trained on all the participants in group 1 (23 of 23) and applied to the connectivity data from each participant in group 2 to produce images of predicted activation. We compared these predictions to each participant's observed fMRI image (Table 1 and Fig. 3). The connectivity model was significantly more accurate across participants than the distance model ($t_{20} = -6.72$, two-tailed t -test, $P = 1.53 \times 10^{-6}$). The connectivity-based predictions were significantly better than distance-based predictions in 18 of 21 participants at $P < 0.001$ (Fig. 2c). The models were not significantly different for the remaining 3 participants.

A group average was generated from all participants' contrast images to faces > scenes in group 1 and registered to each participant's own anatomy in group 2. Across participants, the group-average predictions were significantly less accurate than the connectivity-based predictions ($t_{20} = -4.80$, two-tailed t -test, $P = 1.08 \times 10^{-4}$). Comparing the absolute error within each participant, we found that functional

activation was better predicted by connectivity than by the group average-based model in 16 of 21 participants at $P < 0.001$. Only 1 participant's fusiform profile was more accurately predicted by the group average than by the connectivity model, and the models were not significantly different for the remaining 4 participants (Fig. 2d). The analyses above were repeated for face and scene selectivity in the left fusiform with the same results (Supplementary Results and Discussion).

To investigate which targets made a significant ($P < 0.05$) contribution to the final model (Table 2), we applied a model built from only those significant predictors (with all other model coefficients set to 0) to the structural connectivity data of group 2. The MAE across participants was significantly better than the original connectivity model's MAE (new model's MAE, 0.683 ± 0.02 ; $P = 0.038$), demonstrating the predictive impact of these regions. Some of the highest

positive predictors were right inferotemporal, lateral occipital and superior temporal regions, whereas right lingual and parahippocampal cortices were among the highest negative predictors (Fig. 4).

Spatial relationship of function and connectivity

We calculated the center of mass to the best face (inferotemporal) and scene (lingual) predictors in each participant to visualize the spatial relationship between connectivity and function (Fig. 5a). More subject variability was observed in the medial-lateral dimension for the positive functional activation and in the anterior-posterior dimension for the negative functional activation; we therefore calculated correlations between functional values and connectivity strengths along those dimensions respectively. Across participants, centroid locations for face responses significantly correlated with the centroid locations of connectivity to inferotemporal cortex along the medial-lateral dimension (Fig. 5b, $r = 0.46$, $P = 0.002$). That is, individual participants who had a more medial center of functional activation to faces than other individuals also had a more medial center of connectivity to the inferotemporal target region. Similarly, lingual centroids significantly correlated with scene centroids along the anterior-posterior dimension (Fig. 5c, $r = 0.41$, $P = 0.005$).

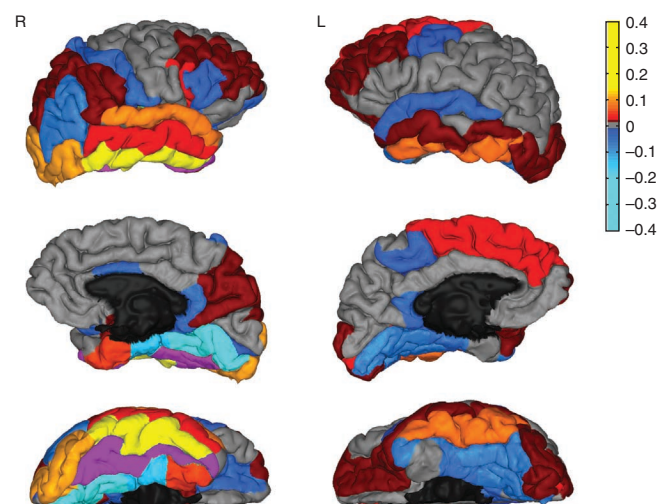


Figure 4 Model coefficients for each target region from the final connectivity model. Target regions are color-coded from hot to cold to reflect positive or negative model coefficients and are projected onto the right and left pial surfaces of an example participant, with the lateral view on the top row, medial view on the second row and ventral view on the bottom. The significant predictors of face-selective voxels are regions labeled from red to yellow, and the scene-selective predictors are labeled from blue to light blue. The seed region is highlighted in purple. Medial walls are shown in black and were not included in the analyses. See Results for the anatomical nomenclature of the target regions.

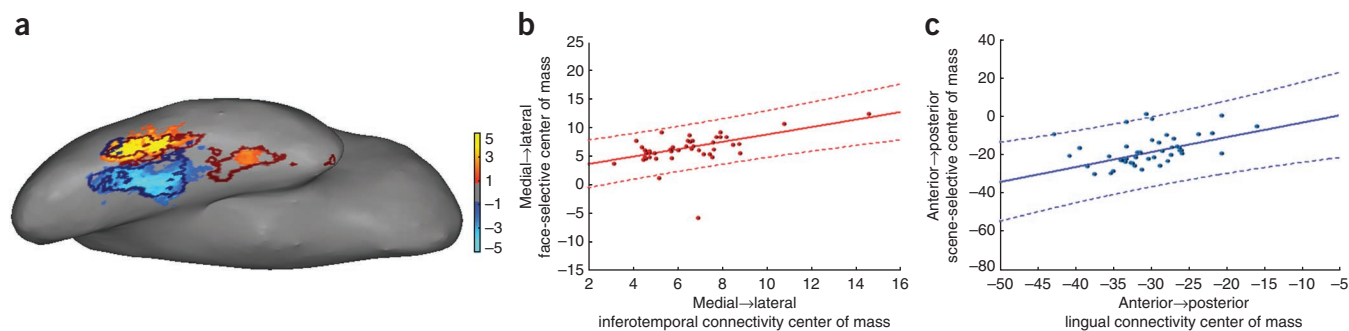


Figure 5 Spatial relationship of function with connection strength to the highest predictors. (a) Functional activation for faces > scenes of an example participant, with the thresholded boundaries of inferotemporal connectivity overlaid in dark red and boundaries of lingual connectivity overlaid in dark blue. Units are t -statistic. (b) Each participant's center of mass of connectivity to the inferotemporal region plotted against that participant's center of mass of positively responding voxels along the medial–lateral dimension, along which each participant's connectivity varied with face selectivity. (c) Centroids of lingual connectivity plotted against centroids of negatively responding voxels along the anterior–posterior dimension. Data in **b, c** are in millimeters relative to each participants' fusiform centroid. Solid lines are the least-square fits of these data; dashed lines, 99% confidence intervals.

To better establish how individual subject variability in connectivity profiles can be sensitive to individual subject variability in functional responses, we tested whether connectivity patterns of one participant can better predict that participant's functional activation than another participant's connectivity patterns. Unlike any of the analyses above, this relied on identifying the same voxel spatially across participants, so data from each participant in group 2 were registered to Montreal Neurological Institute (MNI) template space and subsequently onto each other participant's native anatomical space. We then made functional predictions for each participant based on each other participant's connectivity pattern. A participant's own connectivity values were better at predicting their own functional activation than other participants' connectivity values ($t_{419} = 11.67$, paired t -test, $P = 0$). Thus, the connectivity model is detecting relationships between functional responses and connectivity patterns that capture individual variation.

DISCUSSION

The present study provides evidence of a direct relationship between structural connectivity and function in the human brain. Specifically, we demonstrate that the responses to faces within an individual's right fusiform gyrus can be predicted from that individual's patterns of structural connectivity alone. This approach further reveals which targets are most influential in predicting function. Voxels with higher responses to faces had characteristic patterns of connectivity to other brain regions that distinguished them from neighboring voxels with lower responses to faces, or higher responses to scenes.

The connectivity model outperformed the random permutation control, indicating that there exists a strong relationship between connectivity and function. Moreover, it outperformed the distance control, suggesting that spatial information alone is insufficient for predicting functional activity and that connectivity offers information above and beyond the topographic information inherently embedded in it (owing to the posited small-world organization of cortical connectivity^{24,25}). The relationship between function and spatial information was highly variable across participants, whereas the connectivity data were consistent across participants in relationship to the functional responses. When compared to the group-average benchmark, a standard method of defining face-selective ROIs in fMRI studies, connectivity was a better predictor of the individual's actual activation pattern in over 70% of the participants. One reason that the group average did not successfully predict the activation pattern could be the high variability of activation loci, relative to the standard template (for example, ref. 26).

Although we have treated spatial metrics as potential confounds and controlled for them by using distance and group activation models as controls, future studies may build other geometric models that do predict intersubject variability in functional activation. For example, detailed models of cortical folding patterns²⁷, myelination²⁸ and/or cortical thickness²⁹ may be detectable with MRI and predictive of functional regions. Connectivity can provide a complementary source of evidence in some cases, whereas in others it may be the only gross morphological marker available.

Despite spatial variability in functional responses, the connectivity model was highly accurate across participants. We found that the spatial distribution of face and scene selectivity varied in tandem with connection strength to their most predictive targets. A direct analysis of subject-to-subject variability revealed that although each participant's connectivity profile did well at predicting that person's own functional response, it predicts another participant's functional responses relatively poorly. Overall, the connectivity patterns seemed highly sensitive to individual variation in function.

Although the results from group 1 were noteworthy, they could have been specific to one data set²². The findings from group 2 demonstrate that this is not the case: the connectivity model's predictions from group 1 were much more accurate than both the distance and group-average models in more than 70% of the new group of participants. This result was especially noteworthy because the participants in group 2 had been scanned while performing a different functional task. The two tasks differed in the type of stimuli presented (1-s static images versus 3-s movie clips), type of design (event-related versus block), number of runs (1 versus 3) and scan parameters (also see Online Methods for other differences). Further, the structural connectivity measures in this second group were acquired using a DWI sequence with half as many gradient directions (30 versus 60), indicating the generalizability of the connectivity model across functional tasks and diffusion sequences.

This analysis also revealed the target brain regions for which connectivity with the fusiform was most predictive of face- or scene-selective activity in the fusiform. Face-selective fusiform voxels were predicted by connectivity with regions that have been previously reported to function in face processing, such as the inferior and superior temporal cortices (for example, refs. 30,31). Scene-selective voxels, by contrast, were best predicted by their connectivity to key brain areas involved in scene recognition, such as the isthmuscingulate (containing the retrosplenial cortex) and the parahippocampal cortex^{10,32,33}. Unlike functional connectivity, structural connectivity

models are naive to the functional responses of the target regions. Therefore, a region need not be category selective to be connected to (and predictive of) selective voxels in the fusiform. For example, we also discovered unexpected predictors of face selectivity, such as the cerebellar cortices. Even though the cerebellum is not commonly considered as part of the “core” or “extended” face processing network^{3,30,34}, tracer^{35–37} studies have revealed disynaptic connections with extrastriate visual cortices by way of the pons, which tractography is able to reconstruct (see **Supplementary Figs. 1 and 2**) and which is corroborated by functional connectivity³⁸. Future studies may explore these relationships to further expand on the role of functional responses in components of a structural network. New structure–function relationships could be investigated in macaques with functional and connectivity data, and subsequently validated more directly through more invasive techniques involving tracer injections (for example, refs. 39,40).

The final connectivity model also provides a framework with which to evaluate the impact of the most predictive targets and their spatial distribution. The model built from only the significantly predictive targets resulted in more accurate predictions than the model based on all of the target regions. Although some of the best predictors from this model were nearby regions, most of them were distant to the fusiform; further analyses excluding the fusiform’s neighbors (**Supplementary Results and Discussion**) revealed that although proximal targets are part of the fusiform’s network, they do not fully account for the connectivity model’s performance. Altogether, a distributed network of brain regions characterizes category-specific visual processing in the fusiform gyrus.

The connectivity fingerprint has practical applications, both for defining ROIs independently of a task and also for exploring group differences in structural connectivity signatures. Researchers or clinicians can apply the relationships discovered here to predict functional activation at the single-subject level in populations who do not or cannot have a functional localizer, and they should expect that this will be a more accurate prediction than group-based methods. The connectivity model provided here can also be directly compared to a connectivity model built from study participants with specific conditions or lesions. For instance, compromised structural connectivity in individuals with congenital prosopagnosia has previously been suggested to influence their deficits in face recognition, in light of their surprisingly normal functional activation in the fusiform⁴¹. This type of analysis can shed light on which components (if any) of the fusiform connective fingerprint are altered or compromised in individuals with congenital prosopagnosia. A similar analysis can be used to explore possible substrates of face-processing differences in autism, normal development and aging.

Future studies can also extend the present methods to other brain regions and contrasts that are commonly used as functional localizers, such as retinotopy in visual cortices, scene selectivity in the parahippocampal place area¹⁰ or expression specificity in the superior temporal sulcus. In some cases, more complex or non-linear approaches might better capture the relationship of connectivity and function. We implemented a linear fit to provide more parsimonious interpretations and to establish the feasibility of modeling structure–function relationships. Since these relationships are probably not strictly linear in a complex system such as the brain (**Supplementary Fig. 3**), future work can expand these findings, creating better models and elucidating a more detailed relationship between connectivity and function. Also, voxel-to-voxel tractography may help to more finely characterize the structure–function relationships identified here.

These findings open a window into the coupling between structural and functional organization in the brain. The operations of a brain region are determined by both its intrinsic properties (such as cytoarchitecture) that likely determine the operations that it can perform and by the extrinsic connectivity that defines the input–output relations of that brain region. Neuroimaging can relate localized functions (determined by means of fMRI) to input–output patterns of cortical connectivity (determined by means of probabilistic tractography) in an individual. The present findings demonstrate that brain structure–function relations can be defined for category-selective functional activation.

METHODS

Methods and any associated references are available in the online version of the paper at <http://www.nature.com/natureneuroscience/>.

Note: Supplementary information is available on the Nature Neuroscience website.

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AUTHOR CONTRIBUTIONS

Z.M.S. and D.E.O. designed and performed experiments, analyzed data and wrote the manuscript. K.K. designed and performed experiments. G.R. performed experiments and analyzed data. J.D.E.G. and R.R.S. designed experiments and helped write the manuscript.

COMPETING FINANCIAL INTERESTS

The authors declare no competing financial interests.

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ONLINE METHODS

Participants. For group 1, 23 participants between the ages of 19 and 42 (mean age = 27.9 ± 1.06 , 12 female) were recruited from the greater Boston area. Group 2 included 21 participants between the ages of 19 and 44 (mean age = 26.9 ± 1.45 , 13 female) who were similarly recruited. Both groups of participants were screened for history of mental illness, gave written informed consent and were compensated at \$30 per hour. The studies were approved by the Massachusetts Institute of Technology and Massachusetts General Hospital ethics committees.

Acquisition parameters for group 1. DWI data were acquired using echo planar imaging (64 slices, voxel size $2 \times 2 \times 2$ mm, 128×128 base resolution, diffusion weighting isotropically distributed along 60 directions, b -value 700 s mm^{-2}) on a 3-T Siemens scanner with a 32-channel head-coil⁴². A high-resolution (1 mm^3) three-dimensional magnetization-prepared rapid acquisition with gradient echo (MPRAGE) scan was acquired on these participants.

We acquired event-related fMRI data (gradient echo sequence 2,000 ms TR, 30 ms TE, 90° flip, 324 volumes, $3.1 \times 3.1 \times 4.0$ mm voxel size) while the same participants viewed color images of faces or scenes while indicating any immediate repetition of an image with a button press. Face stimuli (IASLab, <http://www.affective-science.org/>) consisted of neutral and emotional faces (angry, disgusted and happy). Scene stimuli were all neutral outdoor and indoor scenes⁴³ (<http://cvcl.mit.edu/database.htm>). Face and scene stimuli were ordered using optseq2 (<http://surfer.nmr.mgh.harvard.edu/optseq/>)⁴⁴, an optimization program for jittering trials in event-related experiments.

Acquisition parameters for group 2. DWI acquisition parameters for group 2 were different, with 30 directions of diffusion, 64 slices, voxel size $2 \times 2 \times 2$ mm, 128×128 base resolution, b -value 700 s mm^{-2} , but were acquired on the same scanner with the same 32-channel head-coil as group 1. A high-resolution (1 mm^3) three-dimensional MPRAGE scan was also acquired on these participants.

Stimuli for the fMRI consisted of 3-s movie clips of faces, bodies, scenes, objects and scrambled objects. Movies of faces and bodies were filmed against a black background and framed to reveal just the faces or bodies of seven children, shown one at a time. Scenes consisted primarily of pastoral scenes filmed through a car window while driving slowly through the countryside or suburb. Objects were selected specifically to minimize any suggestion of animacy of the object itself or of an invisible actor pushing the object. Scrambled object clips were constructed by dividing each object movie clip into a 15×15 box grid and spatially rearranging the location of each of the resulting boxes. Pilot testing indicated that a contrast of the response for moving faces versus moving objects identified the same FFA as that identified in a standard static localizer. Further studies in adults show that the FFA responds similarly to movies of faces as to static snapshots of faces⁴⁵.

Functional data were acquired over four block-design functional runs (gradient echo sequence 2,000 ms TR, 30 ms TE, 90° flip, 234 volumes, $3 \times 3 \times 3$ mm voxel size). Each functional run contained three 18-s fixation blocks at the beginning, middle and end of the run. During these blocks, a series of six uniform color fields were presented for 3 s each. Each run also contained two sets of five consecutive stimulus blocks (faces, bodies, scenes, objects or scrambled objects) sandwiched between these rest blocks, resulting in two blocks per stimulus category per run. Each block lasted 18 s and contained six 3-s movie clips from each of the five stimulus categories. The order of stimulus category blocks in each run was palindromic, and specific movie clips were chosen randomly to be presented during the block. Participants were asked to passively view the stimuli.

Functional magnetic resonance imaging analysis. For group 1, functional neuroimaging data were analyzed using Statistical Parametric Mapping software (SPM8, Wellcome Department of Cognitive Neurology, London). Preprocessing included slice timing correction, motion correction and linear trend, and temporal filtering with a 128s cutoff. The images were not spatially normalized. Statistical parametric maps of blood oxygenation level-dependent (BOLD) activation were created using a multiple regression analysis, with regressors defined for the five stimulus categories (neutral, angry, disgusted and happy faces, and scenes). Boxcar functions for each trial type were convolved with a standard double-gamma hemodynamic function (SPM8, <http://www.fil.ion.ucl.ac.uk/spm/>) to generate each regressor. The resulting maps were spatially smoothed

with a 6-mm (full width at half maximum) Gaussian kernel, and the t -statistic image was generated per participant for the contrast of faces > scenes.

Group 2's data were analyzed with FSL software (<http://www.fmrib.ox.ac.uk/fsl/>). Image preprocessing was similar to group 1: images were motion corrected, smoothed (5 mm Gaussian kernel, full width at half maximum) and detrended, and were fit using a standard gamma function ($\delta = 2.25$ and $\tau = 1.25$). Data were not spatially normalized. Statistical modeling was then performed using a general linear model on the preprocessed functional images. Next, t -maps corresponding to the contrast of interest for faces > scenes were overlaid on each participant's high-resolution anatomical image.

For both groups, each participant's functional image for the faces > scenes contrast was registered to his or her diffusion-weighted image. Because we were interested in predicting relative activation values that were independent of task-specific parameters such as the degrees of freedom, we standardized the t -statistic values (x) across the fusiform gyrus per participant. This detrending was performed for each participant j , such that the mean value in the fusiform was subtracted from each voxel's fMRI value (x_{ij}) and divided by the s.d. The standardized value per fusiform voxel (x_{zij}) of participant j was then used for the subsequent regression models.

Tractography. Automated cortical and subcortical parcellation was performed with FreeSurfer^{46,47} to define specific cortical and subcortical regions in each individual's T1 scan, using a published atlas⁴⁸. Automated segmentation results were reviewed for quality control and were then registered to each individual's diffusion images and used as the seed and target regions for fiber tracking. The resulting cortical and subcortical targets were then checked and corrected for automatic parcellation or segmentation errors if necessary. There was one seed region per participant, and the 85 target regions were defined as all other automatic parcels, not including the seed. The principal diffusion directions were calculated per voxel, and probabilistic diffusion tractography was carried out using FSL-FDT^{17,49} with 25,000 streamline samples in each seed voxel to create a connectivity distribution to each of the target regions, while avoiding a mask consisting of the ventricles.

Regressions. All analyses were performed on subject-specific anatomy, rather than extrapolation from a template brain, except for the group-average models. For the regression models, each observation was an individual voxel in native space, and there was no identifying or matching of spatial location of voxels across participants. Further, the model was blind to the participant each voxel belonged to.

On group 1, we built a regression model using LOOCV: the model was trained to predict the standardized fMRI value for each native-space fusiform voxel based on connectivity data concatenated across 22 of 23 participants and was tested using the remaining participant's data (Fig. 1a). This was performed iteratively for all participants. For group 2, the analyses were performed in a similar manner, except that the regressions were performed on all the participants in group 1 (23 of 23) and simply applied to each participant in group 2's connectivity data to produce an fMRI image of predicted activation. This was then compared to the participants' own observed fMRI images, and MAEs were calculated.

Using the same LOOCV method, we trained a regression model to predict t -values of fusiform voxels based on each voxel's physical Euclidian distance to each other target region's center of mass, rather than each voxel's connection probability to each target region. In this way, the connectivity and distance models had the same number of dimensions and were generated identically except for the information present in each model. We also considered other 85-dimensional spatial metrics, such as distance to the nearest voxel of each target, and found that these measures were highly similar to the present one. We applied the regression coefficients from the distance model generated from all group 1 participants to each participant in group 2, as described for the connectivity model.

We created random distributions by training models using the observed fMRI images and connection probabilities but randomizing the voxel data. We permuted across 5,000 random combinations of connection probability to fMRI activation values per participant and thus obtained a distribution of random MAE per participant. We then performed a one-tailed t -test to determine whether the mean of the participant's random distribution was significantly greater than the same participant's MAE for connectivity-based predictions.

Each participant's functional data were spatially normalized into MNI space with FSL and FreeSurfer, checked and corrected for registration errors, and superimposed to create composite maps. For group 1 cross-validation, we performed LOOCV: a random-effects test on whole-brain fMRI data was performed with SPM8 on the contrast images for faces > scenes from all but one participant. The resulting *t*-statistic image, which was based on all the other participants in normalized space, was applied to the participant left out of the group analysis and was registered back into his or her native space. We analyzed only the right fusiform gyrus in comparing the group average prediction to that participant's actual fMRI image using measures of MAE (Fig. 1b).

For group 2, we created the group average fMRI image using the same method above, but from all group 1 participants' observed (actual) fMRI images. This fMRI image was mapped onto each participant in group 2's native-space coordinates and compared to that participant's observed fMRI pattern.

Accuracy and benchmark comparisons. As a measure of accuracy, we measured the absolute error per voxel (reported in standardized units) per participant, by calculating the absolute difference between the predicted and actual values. To statistically compare the performance of the connectivity model to the random and benchmark models, we performed a pairwise *t*-test per participant across all their fusiform voxels. A criterion threshold of $P < 0.001$ was used to report the number of participants whose activation pattern was better predicted by one model versus another. Mean absolute error (MAE) was also calculated per participant for each model by averaging the absolute error across the fusiform voxels. A two-tailed Student's *t*-test of the MAEs per participant was then used to compare models, with the same threshold ($P < 0.001$) to decide which model's predictions were significantly better.

Spatial relationship of function and connectivity. We registered the connectivity data for the right inferotemporal and lingual targets to the native-space anatomical image of each participant in groups 1 and 2 and projected these data to each participant's native surface vertices using FreeSurfer. The functional data were similarly projected to the surface. We calculated the center of mass for the targets with respect to a reference frame fixed at the center of mass for each participant's fusiform gyrus (also on the surface). After partitioning the functionals into positive and negative values, we similarly calculated their centers

of mass with respect to the fusiform. We observed more subject variability in the medial-lateral dimension for the positive functionals and more anterior-posterior variability for the negative functionals, and we therefore calculated correlations between functional values and connectivity strengths along those dimensions respectively. Since both functional and connectivity centers of mass were calculated with respect to the subject's own fusiform, the correlations were not biased by cross-subject variability in the boundaries between the seed region and the predictive regions.

For the direct analyses of individual subject variation, we registered each group 2 participant's connectivity data to MNI space, and subsequently onto each other participant's brain, using FreeSurfer and FSL registration tools. We then applied the final model designed from group 1 to both the original participant's and registered participant's connectivity values. This was done for all combinations of participant pairs (420). We then compared the MAEs from predictions built from each participant's own connectivity with those built from another participant's connectivity across all participants in group 2. All of the above predictions were restricted to those voxels that overlapped between the original and registered participants.

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