Natural Scene Statistics Account for the Representation of Scene Categories in Human Visual Cortex

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SUMMARY

During natural vision, humans categorize the scenes they encounter: an office, the beach, and so on. These categories are informed by knowledge of the way that objects co-occur in natural scenes. How does the human brain aggregate information about objects to represent scene categories? To explore this issue, we used statistical learning methods to learn categories that objectively capture the co-occurrence statistics of objects in a large collection of natural scenes. Using the learned categories, we modeled fMRI brain signals evoked in human subjects when viewing images of scenes. We find that evoked activity across much of anterior visual cortex is explained by the learned categories. Furthermore, a decoder based on these scene categories accurately predicts the categories and objects comprising novel scenes from brain activity evoked by those scenes. These results suggest that the human brain represents scene categories that capture the co-occurrence statistics of objects in the world.

INTRODUCTION

During natural vision, humans categorize the scenes that they encounter. A scene category can often be inferred from the objects present in the scene. For example, a person can infer that she is at the beach by seeing water, sand, and sunbathers. Inferences can also be made in the opposite direction: the category “beach” is sufficient to elicit the recall of these objects plus many others such as towels, umbrellas, sandcastles, and so on. These objects are very different from those that would be recalled for another scene category such as an office. These observations suggest that humans use knowledge about how objects co-occur in the natural world to categorize natural scenes.

There is substantial behavioral evidence to show that humans exploit the co-occurrence statistics of objects during natural vision. For example, object recognition is faster when objects in a scene are contextually consistent (Biederman, 1972; Biederman et al., 1973; Palmer, 1975). When a scene contains objects that are contextually inconsistent, then scene categorization is more difficult (Potter, 1975; Davenport and Potter, 2004; Joubert et al., 2007). Despite the likely importance of object co-occurrence statistics for visual scene perception, few fMRI studies have investigated this issue systematically. Most previous fMRI studies have investigated isolated and decontextualized objects (Kanwisher et al., 1997; Downing et al., 2001) or a few, very broad scene categories (Epstein and Kanwisher, 1998; Peelen et al., 2009). However, two recent fMRI studies have investigated isolated and decontextualized objects (Walther et al., 2009; MacEvoy and Epstein, 2011) provide some evidence that the human visual system represents information about individual objects during scene perception.

Here we test the hypothesis that the human visual system represents scene categories that capture the statistical relationships between objects in the natural world. To investigate this issue, we used a statistical learning algorithm originally developed to model large text corpora to learn scene categories that capture the co-occurrence statistics of objects found in a large collection of natural scenes. We then used fMRI to record blood oxygenation level-dependent (BOLD) activity evoked in the human brain when viewing natural scenes. Finally, we used the learned scene categories to model the tuning of individual voxels and we compared predictions of these models to alternative models based on object co-occurrence statistics that lack the statistical structure inherent in natural scenes.

We report three main results that are consistent with our hypothesis. First, much of anterior visual cortex represents scene categories that reflect the co-occurrence statistics of objects in natural scenes. Second, voxels located within and beyond the boundaries of many well-established functional ROIs in anterior visual cortex are tuned to mixtures of these scene categories. Third, scene categories and the specific objects that occur in novel scenes can be accurately decoded from evoked brain activity alone. Taken together, these results suggest that scene categories represented in the human brain...
Learning the statistical relationships between objects in the natural world.

RESULTS

Learning Natural Scene Categories

To test whether the brain represents scene categories that reflect the co-occurrence statistics of objects in natural scenes, we first had to obtain such a set of categories. We used statistical learning methods to solve this problem (Figures 1A and 1B). First, we created a learning database by labeling the individual objects in a large collection of natural scenes (Figure 1A). The frequency counts of the objects that appeared in each scene were then used as input to the Latent Dirichlet Allocation (LDA) learning algorithm (Blei et al., 2003). LDA was originally developed to learn underlying topics in a collection of documents based on the co-occurrence statistics of the words in the documents. When applied to the frequency counts of the objects in the learning database, the LDA algorithm learns an underlying set of scene categories that capture the co-occurrence statistics of the objects in the database.

LDA defines each scene category as a list of probabilities that are assigned to each of the object labels within an available vocabulary. Each probability reflects the likelihood that a specific
object occurs in a scene that belongs to that category (Figure 1B). LDA learns the probabilities that define each scene category without supervision. However, the number of distinct categories the algorithm learns and the object label vocabulary must be specified by the experimenter. The vocabulary used for our study consisted of the most frequent objects in the learning database. For each category are shown. Format is the same as in Figure 1B. Ten examples taken from the 20 best scene categories identified across subjects (corresponding to the red dot in A). The seven most probable objects (left column) is aptly named “Roadway” because it is most likely to contain the objects “car,” “vehicle,” “highway,” “crash barrier,” and “street lamp.” The other examples shown in Figure 1B can also be assigned intuitive names that describe typical natural scenes. Once a set of scene categories has been learned, the LDA algorithm also offers a probabilistic inference procedure that can be used to estimate the probability that a new scene belongs to each of the learned categories, conditioned on the objects in the new scene.

**Voxelwise Encoding Models Based on Learned Scene Categories**

To determine whether the brain represents the scene categories learned by LDA, we recorded BOLD brain activity evoked when human subjects viewed 1,260 individual natural scene images. We used the LDA probabilistic inference procedure to estimate the probability that each of the presented stimulus scenes belonged to each of a learned set of categories. For instance, if a scene contained the objects “plate,” “table,” “fish,” and “beverage,” LDA would assign the scene a high probability of belonging to the “Dining” category in Figure 1B, a lower probability to the “Aquatic” category, and near zero probability to the remaining categories (Figure 1C, green oval).

The category probabilities inferred for each stimulus scene were used to construct voxelwise encoding models. The encoding model for each voxel consisted of a set of weights that best mapped the inferred category probabilities of the stimulus scenes onto the BOLD responses evoked by the scenes (Figure 1C, green hexagon). Model weights were estimated using regularized linear regression applied independently for each subject and voxel. The prediction accuracy for each voxelwise encoding model was defined to be the correlation coefficient (Pearson’s r score) between the responses evoked by a novel set of stimulus scenes and the responses to those scenes predicted by the model.

Introspection suggests that humans can conceive of a vast number of distinct objects and scene categories. However, because the spatial and temporal resolution of fMRI data are fairly coarse (Buxton, 2002), it is unlikely that all these objects or scene categories can be recovered from BOLD signals. BOLD signal-to-noise ratios (SNRs) also vary dramatically across individuals, so the amount of information that can be recovered from individual fMRI data also varies. Therefore, before proceeding with further analysis of the voxelwise models, we first identified the single set of scene categories that provided the best predictions of brain activity recorded from all subjects. To do so, we examined how the amount of accurately predicted cortical territory across subjects varied with specific settings of the number of individual categories and object vocabulary size assumed by the LDA algorithm during category learning. Specifically, we incremented the number of individual scene categories and object vocabulary size by the LDA algorithm during category learning. The number of individual scene categories learned using LDA, for example, ranged from 2 to 40 while also varying the size of the object label vocabulary from the 25 most frequent to 950 most frequent objects in the learning database (see Experimental Procedures for further details). Figure 2A shows the relative amount of accurately predicted cortical territory across subjects based on each setting. Accurate predictions are stable across a wide range of settings.

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**Table 1:**

<table>
<thead>
<tr>
<th>Category</th>
<th>Number of Individual Categories</th>
<th>Vocabulary Size</th>
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<tbody>
<tr>
<td>Plants</td>
<td></td>
<td></td>
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<tr>
<td>Urban Street</td>
<td></td>
<td></td>
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<tr>
<td>Bath Bedroom</td>
<td></td>
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<tr>
<td>Living Room</td>
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<tr>
<td>People Moving</td>
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<tr>
<td>Water Animals</td>
<td></td>
<td></td>
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<tr>
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<td></td>
<td></td>
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<tr>
<td>Desk</td>
<td></td>
<td></td>
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<tr>
<td>Boatway</td>
<td></td>
<td></td>
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<tr>
<td>Signs/Text</td>
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**Figure 2:** Identifying the Best Scene Categories for Modeling Data across Subjects

(A) Encoding model performance across a range of settings for the specified number of distinct categories learned using LDA (y axis) and vocabulary size (x axis). Each pixel corresponds to one of the candidate scene categories learned by LDA when applied to the learning database. The color of each pixel represents the relative amount of cortical territory across subjects that is accurately predicted by encoding models based on a specific setting for the number of individual categories and vocabulary size. The number of individual categories was incremented from 2 to 40. The object vocabulary was varied from the 25 most frequent to the 950 most frequent objects in the learning database. The red dot identifies the number of individual categories and vocabulary size that produce accurate predictions for the largest amount of cortical territory across subjects. For individual results, see Figure 1B. (B) Ten examples taken from the 20 best scene categories identified across subjects (corresponding to the red dot in A). The seven most probable objects for each category are shown. Format is the same as in Figure 1B. See Figures S4 and S5 for interpretation of all 20 categories.
Across subjects, the encoding models perform best when based on 20 individual categories and composed of a vocabulary of 850 objects (Figure 2A, indicated by red dot; for individual subject results, see Figure S3 available online). Examples of these categories are displayed in Figure 2B (for an interpretation of all 20 categories, see Figures S4 and S5). To the best of our knowledge, previous fMRI studies have only used two to eight distinct categories and 2–200 individual objects (see Walther et al., 2009; MacEvoy and Epstein, 2011). Thus, our results show there is more information in BOLD signals related to encoding scene categories than has been previously appreciated.

We next tested whether natural scene categories were necessary to accurately model the measured fMRI data. We derived a set of null scene categories by training LDA on artificial scenes. The artificial scenes were created by scrambling the objects in the learning database across scenes, thus removing the natural statistical structure of object co-occurrences inherent in the original learning database. If the brain incorporates information about the co-occurrence statistics of objects in natural scenes, then the prediction accuracy of encoding models based upon these null scene categories should be much poorer than encoding models based on scene categories learned from natural scenes.

Indeed, we find that encoding models based on the categories learned from natural scenes provide significantly better predictions of brain activity than do encoding models based on the null categories and for all subjects (p < 1 × 10^−15 for all subjects, Wilcoxon rank-sum test for differences in median prediction accuracy across all cortical voxels and candidate scene category settings; subject S1: W(15,025,164) = 9.96 × 10^13; subject S2: W(24,440,399) = 3.04 × 10^14; subject S3: W(15,778,360) = 9.93 × 10^13; subject S4: W(14,705,625) = 1.09 × 10^14). In a set of supplemental analyses, we also compared the LDA-based models to several other plausible models of scene category representation. We find that the LDA-based models provide superior prediction accuracy to all these alternative models (see Figures S12–S15). These results support our central hypothesis that the human brain encodes categories that reflect the co-occurrence statistics of objects in natural scenes.

Categories Learned From Natural Scenes Explain Selectivity in Many Anterior Visual ROIs

Previous fMRI studies have identified functional regions of interest (ROIs) tuned to very broad scene categories, such as places (Epstein and Kanwisher, 1998), as well as to narrow object categories such as faces (Kanwisher et al., 1997) or body parts (Downing et al., 2001). Can selectivity in these regions be explained in terms of the categories learned from natural scene object statistics?

We evaluated scene category tuning for voxels located within the boundaries of several conventional functional ROIs: the fusiform face area (FFA; Kanwisher et al., 1997), the occipital face area (OFA; Gauthier et al., 2000), the extrastriate body area (EBA; Downing et al., 2001), the parahippocampal place area (PPA; Epstein and Kanwisher, 1998), the transverse occipital sulcus (TOS; Nakamura et al., 2000; Grill-Spector, 2003; Hasson et al., 2003), the retrosplenial cortex (RSC; Maguire, 2001), and lateral occipital cortex (LO; Malach et al., 1995).

Figure 3A shows the boundaries of these ROIs, identified using separate functional localizer experiments, and projected on the cortical flat map of one representative subject. The color of each location on the cortical map indicates the prediction accuracy of the corresponding encoding model. All encoding models were based on the 20 best scene categories identified across subjects. These data show that the encoding models accurately predict responses of voxels located in many ROIs within anterior visual cortex. To quantify this effect, we calculated the proportion of response variance explained by the encoding models, averaged across all voxels within each ROI. We find that the average proportion of variance explained to be significantly greater than chance for every anterior visual cortex ROI and for all subjects (p < 0.01; see Experimental Procedures for details). Thus, selectivity in many previously identified ROIs can be explained in terms of tuning to scene categories learned from natural scene statistics.

To determine whether scene category tuning is consistent with tuning reported in earlier localizer studies, we visualized the weights of encoding models fit to voxels within each ROI. Figure 3C shows encoding model weights averaged across all voxels located within each function ROI. Scene category selectivity is broadly consistent with the results of previous functional localizer experiments. For example, previous studies have suggested that PPA is selective for presence of buildings (Epstein and Kanwisher, 1998). The LDA algorithm suggests that images containing buildings are most likely to belong to the “Urban/Street” category (see Figure 2B), and we find that voxels within PPA have large weights for the “Urban/Street” category (see Figures S4 and S5). To take another example, previous studies have suggested that OFA is selective for the presence of human faces (Gauthier et al., 2000). Under the trained LDA model, images containing faces are most likely to belong to the “Portrait” category (see Figures S4 and S5), and we find that voxels within OFA have large weights for the “Portrait” category.

Although category tuning within functional ROIs is generally consistent with previous reports, Figure 3C demonstrates that tuning is clearly more complicated than assumed previously. In particular, many functional ROIs are tuned for more than one scene category. For example, both FFA and OFA are thought to be selective for human faces, but voxels in both these areas also have large weights for the “Plants” category. Additionally, area TOS, an ROI generally associated with encoding information important for navigation, has relatively large weights for the “Portrait” and “People Moving” categories. Thus, our results suggest that tuning in conventional ROIs may be more diverse than generally believed (for additional evidence, see Huth et al., 2012 and Naselaris et al., 2012).

Decoding Natural Scene Categories from Evoked Brain Activity

The results presented thus far suggest that information about natural scene categories is encoded in the activity of many voxels located in anterior visual cortex. It should therefore be possible to decode these scene categories from brain activity evoked by viewing a scene. To investigate this possibility, we constructed a decoder for each subject that uses voxel activity evoked in anterior visual cortex to predict the probability that a
viewed scene belongs to each of 20 best scene categories identified across subjects. To maximize performance, the decoder used only those voxels for which the encoding models produced accurate predictions on a held-out portion of the model estimation data (for details, see Experimental Procedures).

We used the decoder to predict the 20 category probabilities for 126 novel scenes that had not been used to construct the decoder. Figure 4A shows several examples of the category probabilities predicted by the decoder. The scene in the upper right of Figure 4A depicts a harbor in front of a city skyline. The predicted category probabilities indicate that the scene is most likely a mixture of the categories “Urban” and “Boatway,” which is an accurate description of the scene. Inspection of the other examples in the figure suggests that the predicted scene category probabilities accurately describe many different types of natural scenes.

To quantify the accuracy of each decoder, we calculated the correlation (Pearson’s $r$) between the scene category probabilities predicted by the decoder and the probabilities inferred using the LDA algorithm (conditioned on the labeled objects in each scene). Figure 4B shows the distribution of decoding accuracies across all decoded scenes, for each subject. The median accuracies and 95% confidence interval (CI) on median estimates are indicated by the black cross-hairs. Most of the novel scenes are decoded significantly for all subjects. Prediction accuracy was calculated on responses from a separate validation set of stimuli not used to estimate the model. ROI Abbreviations: V1–V4, retinotopic visual areas 1–4; PPA, parahippocampal place area; FFA, fusiform face area; EBA, extrastriate body area; OFA, occipital face area; RSC, retrosplenial cortex; TOS, transverse occipital sulcus. Center key: A, anterior; P, posterior; S, superior; I, inferior. For remaining subjects’ data, see Figure S6.

(B) Each bar indicates the average proportion of voxel response variance in an ROI that is explained by voxelwise encoding models estimated for a single subject. Bar colors distinguish individual subjects. Error bars represent SEM. For all anterior visual ROIs and for all subjects, encoding models based on scene categories learned from natural scenes explain a significant proportion of voxel response variance ($p < 0.01$, indicated by red lines).

(C) The average encoding model weights for voxels within distinct functional ROIs. Averages are calculated across all voxels located within the boundaries of an ROI and across subjects. Each row displays the average weights for the scene category listed on the left margin. Each column distinguishes average weights for individual ROIs. The color of each pixel represents the positive (red) or negative (blue) average ROI weight for the corresponding category. The size of each pixel is inversely proportional to the magnitude of the SEM estimate; larger pixels indicate selectivity estimates with greater confidence. SE scaling is according to the data within an ROI (column). ROI tuning is generally consistent with previous findings. However, tuning also appears to be more complex than indicated by conventional ROI-based analyses. For individual subjects’ data, see Figure S7; see also Figures S8–S15.
across all scenes exhibited systematically greater-than-chance performance for all subjects \( (p < 0.02 \text{ for all subjects, Wilcoxon rank-sum test; subject S1: } W(126) = 18,585; \text{ subject S2: } W(126) = 17,274; \text{ subject S3: } W(126) = 17,018; \text{ subject S4: } W(126) = 19,214. \) The voxels selected for the decoding analysis summarized in Figure 4 were located throughout the visual cortex. However, we also find that accurate decoding can be obtained using the responses of subsets of voxels located within specific ROIs (see Figures S16–S19).

### Predicting the Objects that Occur in Decoded Natural Scenes

Our results suggest that the visual system represents scene categories that capture the co-occurrence statistics of objects in the natural world. This suggests that we should be able to predict accurately the likely objects in a scene based on the scene category probabilities decoded from evoked brain activity.

To investigate this issue, we estimated the probability that each of the 850 objects in the vocabulary for the single best set of scene categories identified across subjects occurred in each of the 126 decoded validation set scenes. The probabilities were estimated by combining the decoded category probabilities, to the likelihood of the labeled objects in each scene if all were selected with equal probability (chance). A likelihood ratio greater than one (red line) indicates that the objects in a scene are better predicted by the decoded object probabilities than by selecting objects randomly. Each plot shows the (horizontally mirrored) histogram of likelihood ratios for a single subject. Median likelihood ratios and 95% CI are represented by the black cross-hairs. For subjects S1–S4, the median likelihood ratio was 1.67 (CI: [1.57, 1.76]), 1.66 (CI: [1.52, 1.72]), 1.62 (CI: [1.45, 1.78]), and 1.66 (CI: [1.56, 1.78]) for subjects S1–S4, respectively.
Scene to calculate the likelihood of the labeled objects in the scene. We then calculated the likelihood of the labeled objects from a naive distribution that assumes all 850 objects are equally likely to occur. The ratio of these likelihoods provides a measure of accuracy for the estimated object probabilities. Likelihood ratios greater than one indicate that the estimated object probabilities better predict the labeled objects in the scene than by picking objects at random (see Experimental Procedures for details).

Figure 4C shows the distribution of likelihood ratios for each subject, calculated for all 126 decoded scenes. The medians and 95% confidence intervals of the median estimates are indicated by the black cross-hairs. Object prediction accuracy across all scenes indicates systematically greater-than-chance performance for all subjects ($p < 1 \times 10^{-15}$ for all subjects, Wilcoxon rank-sum test; subject S1: \(W(126) = 9,983\); subject S2: \(W(126) = 11,375\); subject S3: \(W(126) = 11,103\); subject S4: \(W(126) = 10,715\)).

The estimated object probabilities and the likelihood ratio analysis both show that the objects that are likely to occur in a scene can be predicted probabilistically from natural scene categories that are encoded in human brain activity. This suggests that humans might use a probabilistic strategy to help infer the likely objects in a scene from fragmentary information available at any point in time.

**DISCUSSION**

This study provides compelling evidence that the human visual system encodes scene categories that reflect the co-occurrence statistics of objects in the natural world. First, categories that capture co-occurrence statistics are consistent with our intuitive interpretations of natural scenes. Second, voxelwise encoding models based on these categories accurately predict visually evoked BOLD activity across much of anterior visual cortex, including within several conventional functional ROIs. Finally, the category of a scene and its constituent objects can be decoded from BOLD activity evoked by viewing the scene.

Previous studies of scene representation in the human brain used subjective categories that were selected by the experimenters. In contrast, our study used a data-driven, statistical algorithm (LDA) to learn the intrinsic categorical structure of natural scenes from object labels. These learned, intrinsic scene categories provide a more objective foundation for scene perception research than is possible using subjective categories.

One previous computer vision study used a similar statistical learning approach to investigate the intrinsic category structure of natural scenes (Fei-Fei and Perona, 2005). In that study, the input to the learning algorithm was visual features of intermediate spatial complexity. Because our goal was to determine whether the brain represents the object co-occurrence statistics of natural scenes, we used object labels of natural scenes as input to the learning algorithm rather than intermediate visual features.

The voxelwise modeling and decoding framework employed here (Kay et al., 2008b; Mitchell et al., 2008; Naselaris et al., 2009, 2012; Nishimoto et al., 2011; Thirion et al., 2006) provides a powerful alternative to conventional methods based on statistical parametric mapping (Friston et al., 1996) or multivariate pattern analysis (MVPA; Norman et al., 2006). Studies based on statistical mapping or MVPA do not aim to produce explicit predictive models of voxel tuning, so it is difficult to generalize their results beyond the specific stimuli or task conditions used in each study. In contrast, the goal of voxelwise modeling is to produce models that can accurately predict responses to arbitrary, novel stimuli or task conditions. A key strategy for developing theoretical models of natural systems has been to validate model predictions under novel conditions (Hastie et al., 2008). We believe that this strategy is also critically important for developing theories of representation in the human brain.

Our results generally corroborate the many previous reports of object selectivity in anterior visual cortex. However, we find that tuning properties in this part of visual cortex are more complex than reported in previous studies (see Figures S7, S8–S11, and S16–S19 for supporting results). This difference probably reflects the sensitivity afforded by the voxelwise modeling and decoding framework. Still, much work remains before we can claim a complete understanding of what and how information is represented in anterior visual cortex (Huth et al., 2012; Naselaris et al., 2012).

Several recent studies (Kim and Biederman, 2011; MacEvoy and Epstein, 2011; Peelen et al., 2009) have suggested that the lateral occipital complex (LO) represents, in part, the identity of scene categories based on the objects therein. Taken together, these studies suggest that some subregions within LO should be accurately predicted by models that link objects with scene categories. Our study employs one such model. We find that the encoding models based on natural scene categories provide accurate predictions of activity in anterior portions of LO (Figures 3A and 3B). Note, however, that our results do not necessarily imply that LO represents scene categories explicitly (see Figures S16–S19 for further analyses).

fMRI provides only a coarse proxy of neural activity and has a low SNR. In order to correctly interpret the results of fMRI experiments, it is important to quantify how much information can be recovered from these data. Here we addressed this problem by testing many candidate models in order to determine a single set of scene categories that can be recovered reliably from the BOLD activity measured across all of our subjects (Figure 2A). This test places a clear empirical limit on the number of scene categories and objects that can be recovered from our data. These numbers are larger than what has typically been assumed in previous fMRI studies of scene perception (Epstein and Kanwisher, 1998; Peelen et al., 2009; Walther et al., 2009; MacEvoy and Epstein, 2011), but they are still far smaller than the likely representational capacity of the human visual system.

Theoreticians have argued that the simple statistical properties of natural scenes explain selectivity to low-level features in peripheral sensory areas (Oshhausen and Field, 1996; Smith and Lewicki, 2006). Behavioral data suggest that low-level natural scene statistics also influence the perception of scene categories (Oliva and Torralba, 2001; Torralba and Oliva, 2003). Though several qualitative theories have been proposed that link the object statistics of natural scenes with human scene perception (Biederman, 1981; Palmer, 1975), none have provided an objective, quantitative framework to support this link. The current study provides such a framework. Our data-driven,
model-based approach shows that scene categories encoded in the human brain can be derived from the co-occurrence statistics of objects in natural scenes. This further suggests that the brain exploits natural scene statistics at multiple levels of abstraction. If this is true, then natural scene statistics might be used as a principled means to develop quantitative models of representation throughout the visual hierarchy.

The work reported here could be extended in several ways. For example, although the spatial distribution of objects within a scene appears to influence the representation of the scene (Biederman et al., 1982; Green and Hummel, 2006; Kim and Biederman 2011), the modeling framework used here makes no assumptions about the spatial distribution of objects within scenes. More sophisticated models that incorporate spatial statistics or other mediating factors such as attention may provide further information about the representation of scenes and scene categories in the human brain.

**EXPERIMENTAL PROCEDURES**

**fMRI Data Acquisition**

The experimental protocol used was approved by the UC Berkeley Committee for the Protection of Human Subjects. All fMRI data were collected at the UC Berkeley Brain Imaging Center using a 3 Tesla Siemens Tim Trio MR scanner (Siemens, Germany). For subjects S1, S3, and S4, a gradient-echo echo planar imaging sequence, combined with a custom fat saturation RF pulse, was used for functional data collection. Twenty-five axial slices covered occipital, occipitoparietal, and occipitotemporal cortex. Each slice had a 234 × 234 mm² field of view, 2.60 mm slice thickness, and 0.39 mm slice gap (matrix size = 104 × 104; TR = 2,009.9 ms; TE = 35 ms; flip angle = 74°; voxel size = 2.25 × 2.25 × 2.99 mm³).

For subject S2 only, a gradient-echo echo planar imaging sequence, combined with a custom water-specific excitation (fat-shunting) RF pulse was used for functional data collection. In this case, 31 axial slices covered the entire brain, and each slice had a 224 × 224 mm² field of view, 3.50 mm slice thickness, and 0.63 mm slice gap (matrix size = 100 × 100; TR = 2,004.5 ms; TE = 33 ms; flip angle = 74°; voxel size = 2.24 × 2.24 × 4.13 mm³).

Subject S1 experienced severe visual occlusion of the stimuli when the whole head coil was used. Therefore, for subject S1 the back portion (20 channels) of the Siemens 32 channel quadrature receive head coil was used as a surface coil. The full 32 channel head coil was used for subjects S2, S3, and S4.

**Stimuli**

All stimuli consisted of color images selected from a large database of natural scenes collected from various sources. Each image was presented on an iso-luminant gray background and subtended the central 20° × 20° square of the visual field. Images were presented in successive 4 s trials. On each trial, a photo was flashed for 1 s at 5 Hz, followed by a 3 s period in which only the gray background was present. A central fixation square was superimposed at the center of the display, subtending 0.2° × 0.2° of the visual field. To facilitate fixation, we randomly permuted the fixation square in color (red, green, blue, white) at a rate of 3 Hz. No eye tracking was performed during stimulus presentation. However, all subjects in the study were highly trained psychophysical observers having extensive experience with fixation tasks, and preliminary data collected during an identical visual task showed that the subject cohort maintained stable fixation. Note also that the visual stimuli contained no object labels.

**Experimental Design**

fMRI experiments consisted of interleaved runs that contained images from separate model estimation and validation sets. Data were collected over six sessions for subjects S1 and S4, and seven sessions for subjects S2 and S3. Each of the 35 estimation set runs was 5.23 min in duration and consisted of 36 distinct images presented two times each. Evoked responses to these 1,260 images were used during model estimation. Each of 21 5.23-min-long validation set runs consisted of six distinct images presented 12 times each. The evoked responses to these 126 images were used during model validation. All images were randomly selected for each run with no repeated images across runs.

**fMRI Data Processing**

The SPM8 package (University College, London, UK) was used to perform motion correction, coregistration, and reslicing of functional images. All other preprocessing of functional data was performed using custom software (MATLAB, R2010a, MathWorks). Preprocessing was conducted across all sessions for each subject, using the first run of the first session as the reference. For each voxel, the preprocessed time series was used to estimate the hemodynamic response function (Kay et al., 2008a). Deconvolving each voxel time course from the stimulus design matrix produced an estimate of the response amplitude—a single value—evoked by each image, for each voxel. These response amplitude values were used in both model estimation and validation stages of data analysis. Retinotopic visual cortex was identified in separate scan sessions using conventional methods (Hansen et al., 2007). Standard functional localizers (Spiridon et al., 2006) were also collected in separate scan sessions and were used to identify the anatomical boundaries of conventional ROIs.

**Learning Database and Stimulus Data Sets**

Natural scene categories were learned using Latent Dirichlet Allocation (Blei et al., 2003; see Figure S1 for more details). The LDA algorithm was applied to the object labels of a learning database of 4,116 natural scenes compiled from two image data sets. The first image data set (Lotus Hill; Yao et al., 2007) provided learning data of 903 (71%) of the learning database scenes. The remaining scenes were sampled from an image data set that was created in house. In both data sets, all objects within the visible area of each image were outlined and labeled. Each in-house image was labeled by one of 15 naive labelers. Since each image was labeled by a single labeler, no labels were combined when compiling the databases. In a supplemental analysis, we verify that scene context created negligible bias in the statistics of the object labels (Figure S2). Ambiguous labels, misspelled labels, and rare labels having synonyms within the learning database were edited accordingly (see Supplemental Experimental Procedure 1). Note that the 1,260 stimulus scenes in the estimation set were sampled from the learning database. The validation set consisted of an independent set of 126 natural scenes labeled in house.

**Voxelwise Encoding Modeling Analysis**

Encoding models were estimated separately for each voxel using 80% of the responses to the estimation set stimuli selected at random. The model weights were estimated using regularized linear regression in order to best map the scene category probabilities for a stimulus scene onto the voxel responses evoked when viewing that scene. The category probabilities for a stimulus scene were calculated from the posterior distribution of the LDA inference procedure, conditioned on the labeled objects in the scene (see Supplemental Experimental Procedure 6 for details). Half of the remaining 20% of the estimation data was used to determine model regularization parameters and the other half of the estimation data was used to estimate model prediction accuracy (see Supplemental Experimental Procedure 7 for more details on encoding model parameter estimation).

Prediction accuracy estimates were used to determine the single best set of categories across subjects. For each of the 760 different scene category settings (defining the number of distinct categories and vocabulary size assumed by LDA during learning), we calculated the number of voxels with prediction accuracy above a statistical significance threshold (correlation coefficient > 0.21; p < 0.01; see Supplemental Experimental Procedure 8 for details on defining statistically significant prediction accuracy). This resulted in a vector of 760 values for each subject, where each entry in the vector provided an estimate of the amount of cortical territory that was accurately predicted by encoding models based on each category setting. To combine the cortical territory estimates across subjects, we normalized the vector for each subject to sum to 1 (normalization was done to control for differences in brain size and...
signal-to-noise ratios across subjects) and the Hadamard (element-wise) product of the normalized vectors was calculated. This resulted in a combined distribution of 760 values (see Figure 2A). The peak of the combined distribution gave the single best set of categories across subjects. For more details on this issue, see Supplemental Experimental Procedure 9.

When calculating the proportion of response variance explained in each ROI by the encoding models, statistical significance was determined by permutation. Specifically, the proportion of variance explained was estimated using the responses to the validation set for each voxelwise encoding model. These explained variance estimates were then permuted across all cortical locations and the average was estimated within each functional ROI. Thus, each permutation produced a random sample of average explained variance within the boundaries of each functional ROI. Statistical significance was defined as the upper 99th percentile of the distribution of average explained variance estimates calculated within each ROI after 1,000 voxel permutations. For more details on this procedure, see Supplemental Experimental Procedure 10.

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SUPPLEMENTAL INFORMATION
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