# homework-4-ishagore-4

March 26, 2024

# 1 NEU502B Homework 4: Multivariate pattern analysis

Due March 27, 2024

Submission instructions: First, rename your homework notebook to include your name (e.g. homework-4-nastase.ipynb); keep your homework notebook in the homework directory of your clone of the class repository. Prior to submitting, restart the kernel and run all cells (see Kernel > Restart Kernel and Run All Cells...) to make sure your code runs and the figures render properly. Only include cells with necessary code or answers; don't include extra cells used for troubleshooting. To submit, git add, git commit, and git push your homework to your fork of the class repository, then make a pull request on GitHub to sync your homework into the class repository.

In this homework assignment, you will work through three commonly used methods in cognitive computational neuroscience: (1) neural decoding via multivariate pattern analysis (MVPA); (2) representational similarity analysis (RSA); and (3) voxelwise encoding analysis using regularized regression. Each of these problems builds on tools and ideas we've introduced in the in-class lab notebooks.

```
[152]: import numpy as np import matplotlib.pyplot as plt import seaborn as sns
```

#### 1.0.1 Problem 1: Multivariate pattern classification

First, we'll start with a simple example of classifying distributed response patterns for different object categories from Haxby et al., 2001. We'll begin by loading in the data, as well as labels for the stimuli and runs. You'll need to change data\_dir to a directory on your computer (or the server); if you've already downloaded this dataset in lab, you can set data\_dir to the existing directory to save time.

```
[153]: from nilearn import datasets
from nilearn.image import index_img
import pandas as pd

# Change this path to a directory on your computer!
data_dir = '.../../nilearn-data'

# Load the Haxby et al., 2001 data via Nilearn
haxby_dataset = datasets.fetch_haxby(data_dir=data_dir)
```

```
# Load in session metadata as pandas DataFrame
session = pd.read_csv(haxby_dataset.session_target[0], sep=" ")

# Extract stimuli and run labels for this subject
stimuli, runs = session['labels'].values, session['chunks'].values

# Create a boolean array indexing TRs containing a stimulus (non-rest)
task_trs = stimuli != 'rest'

# Get list of unique stimulus categories (excluding rest)
categories = [c for c in np.unique(stimuli) if c != 'rest']

# Extract task TRs for fMRI data and stimulus/run labels
func_task = index_img(haxby_dataset.func[0], task_trs)
stimuli_task = stimuli[task_trs]
runs_task = runs[task_trs]
```

Use NiftiMasker (with standardize=True) to create a masker for ventral temporal (VT) cortex. Use the masker to extract the NumPy array containing the functional data. (We'll analyze the data using scikit-learn rather than nilearn.)

Now, we'll set up a full SVM classification analysis using leave-one-run-out outer cross-validation with a nested leave-one-run-out inner cross-validation loop for grid search across the values of the SVM regularization parameter C. Sounds like a lot! But scikit-learn makes it pretty straightforward. First, initialize the LinearSVC estimator. Since this well-behaved dataset has the same number of samples for each stimulus category in each run, we can perform leave-one-run-out cross-validation using just KFold rather than having to specify the runs directly. Initialize an outer KFold cross-validator with 12 splits and an inner KFold cross-validator with 11 splits. We'll search over a handful of C parameters: param\_grid = {'C': [1e-2, 1e-1, 1]}. Initialize the GridSearchCV estimator with the SVM estimator, the parameter grid, and the inner cross-validator; then, submit this estimator to cross\_val\_predict with the outer cross-validator to run the full analysis. (This may take a few minutes to run!)

```
[155]: | # Suppress some warnings (e.g. SVM convergence) just to clean up output
       import warnings
       from sklearn.model_selection import cross_val_predict
       from sklearn.model_selection import (cross_val_predict,
                                            GridSearchCV,
                                            KFold)
       from sklearn.svm import LinearSVC
       warnings.filterwarnings("ignore")
       # Initialize SVM and outer/inner CVs:
       classifier = LinearSVC()
       outer cv= KFold(n splits= 12)
       inner_cv= KFold (n_splits = 11)
       # Set up parameter grid:
       param_grid = {'C': [1e-2, 1e-1, 1]}
       # Initialize GridSearchCV estimator:
       grid_search_estimator= GridSearchCV(classifier, param_grid, cv= inner_cv)
       # Generate predictions using cross_val_predict:
       y_pred = cross_val_predict(grid_search_estimator, functional_data,_
        ⇔stimuli_task, cv= outer_cv)
```

Inspect the resulting predictions. We'll evaluate our classifier's predictions in two ways. First, use accuracy\_score from sklearn.metrics to evaluate the predictions (across all test sets) against the actual labels in terms of a single classification accuracy. Procedurally, this is slightly different from computing accuracies on the test for each fold and averaging them—but the resulting value should be the same.

```
[156]: # Print accuracy score:
    from sklearn.metrics import accuracy_score
    accuracy= accuracy_score(stimuli_task, y_pred)
    print(f'The overall classification accuracy is : {accuracy}')
```

The overall classification accuracy is: 0.7233796296296297

To better understand what our classifer is doing (i.e. what it's getting right and what it's getting wrong), we'll construct a confusion matrix. Construct the confusion matrix from the actual stimulus labels and the classifer's predicted labels and plot it below. What categories does the classifier tend to misclassify?

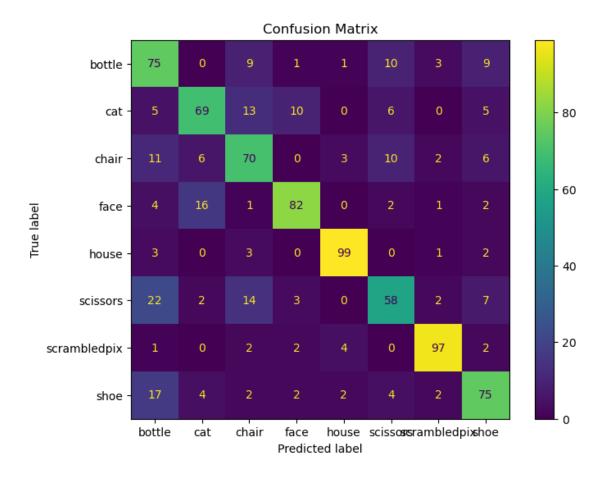
```
[157]: # Create confusion matrix from true and predicted labels:
    from sklearn.model_selection import cross_val_predict
    from sklearn.metrics import confusion_matrix, ConfusionMatrixDisplay

# Plot confusion matrix:
    cm= confusion_matrix(stimuli_task, y_pred)
    fig, ax = plt.subplots(figsize=(8, 6))
```

```
ConfusionMatrixDisplay(cm, display_labels=categories).plot(ax=ax)
plt.title('Confusion Matrix')

#The classifier tends to misclassify scissors, cats.
```

[157]: Text(0.5, 1.0, 'Confusion Matrix')



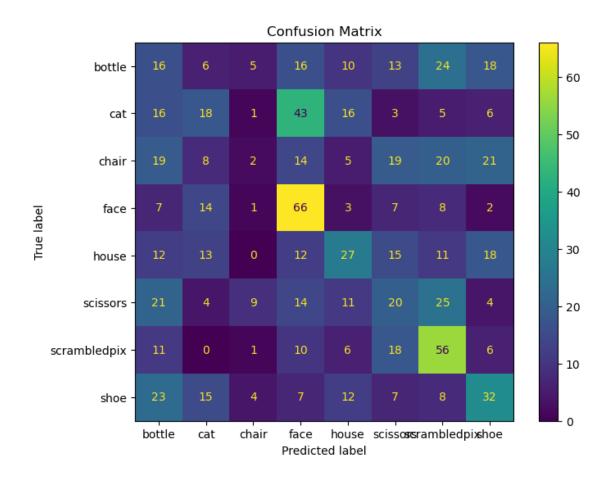
Lastly, we'll repeat the same analysis for functional regions of interest (ROIs) maximally responsive to faces (roughly FFA) and houses (roughly PPA). Use the mask\_face and mask\_house files from the dataset to create an FFA masker and a PPA masker; extract the functional data for both. Submit these datasets to the same analysis as above, and visualize the results in terms of an overall accuracy score and confusion matrix. Interpret the accuracies and confusion matrices in light of the expected chance accuracy, given what you know about these ROIs.

```
[50]: # Create masker for FFA:
    mask_face = haxby_dataset['mask_face'][0]
    masker_ffa = NiftiMasker(mask_img= mask_face, standardize=True)

# Create masker for PPA:
    mask_house = haxby_dataset['mask_house'][0]
```

```
masker_ppa = NiftiMasker(mask_img= mask_house, standardize=True)
      # Uses masker to extract numpy array for VT:
      functional_data_ffa = masker_ffa.fit_transform(func_task)
      functional_data_ppa = masker_ppa.fit_transform(func_task)
[51]: # Initialize SVM and outer/inner CVs:
      classifier_face = LinearSVC()
      outer_cv_face = KFold(n_splits= 12)
      inner_cv_face = KFold (n_splits = 11)
      # Set up parameter grid:
      param_grid = {'C': [1e-2, 1e-1, 1]}
      # Initialize GridSearchCV estimator:
      grid_search_estimator= GridSearchCV(classifier_face, param_grid, cv=_
       →inner cv face)
      # Generate predictions using cross_val_predict:
      y_pred_face = cross_val_predict(grid_search_estimator, functional_data_ffa, u
       ⇔stimuli_task, cv= outer_cv)
[36]: # Print accuracy score and plot confusion matrix:
      accuracy= accuracy_score(stimuli_task, y_pred_face)
      print(f'The classification accuracy is : {accuracy}')
      cm_face= confusion_matrix(stimuli_task, y_pred)
      fig, ax = plt.subplots(figsize=(8, 6))
      ConfusionMatrixDisplay(cm_face, display_labels=categories).plot(ax=ax)
      plt.title('Confusion Matrix')
```

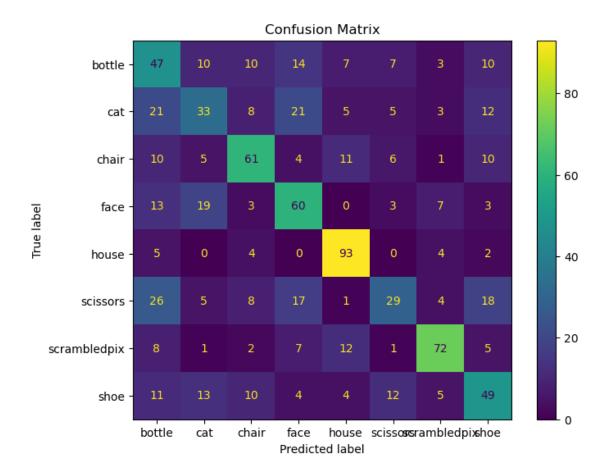
[36]: Text(0.5, 1.0, 'Confusion Matrix')



```
[161]: # Print accuracy score and plot confusion matrix:
    accuracy= accuracy_score(stimuli_task, y_pred_house)
    print(f'The classification accuracy is : {accuracy}')
```

```
cm_face= confusion_matrix(stimuli_task, y_pred_house)
fig, ax = plt.subplots(figsize=(8, 6))
ConfusionMatrixDisplay(cm_face, display_labels=categories).plot(ax=ax)
plt.title('Confusion Matrix')
```

[161]: Text(0.5, 1.0, 'Confusion Matrix')



[158]: # The FFA ROI's accuracy is low, but above chance, suggesting some level of specialization for face recognition. However, the relatively widespread misclassifications across categories indicate that this area may also respond to other stimuli

#The PPA ROI's accuracy is significantly above chance, with a strong indication of house recognition specialization.

## 1.0.2 Problem 2: Representational similarity analysis

In this problem, we'll apply representational similarity analysis (RSA) to the human fMRI dataset from Kriegeskorte et al., 2008. We'll begin by loading in the ROI data and labels.

We provide a rank\_percentile function for visualizing RDMs in a way that more closely matches the paper.

```
[163]: from scipy.stats import rankdata

def rank_percentile(a):
    return rankdata(a) / len(a) * 100
```

First, compute RDMs for the 'lffa', 'rffa', 'lppa', and 'rppa' ROIs for subject 'TI' using correlation distance. Here, we recommend z-soring each voxel across samples prior to computing the pairwise dissimilarities. Plot the RDMs for each ROI using the rank\_percentile function provided above.

```
[164]: # Plot ROI RDMs for subject TI:
    from scipy.stats import zscore
    from scipy.spatial.distance import pdist, squareform

subject= 'TI'

for roi in roi_data[subject]:
    print(f"{roi} ROI shape: {roi_data[subject][roi].shape}")

fig, axs = plt.subplots(1, len(roi_labels), figsize=(20, 5))

for ax, roi in zip(axs, roi_labels):

    roi_data_current = roi_data[subject][roi]
    zscored_data = zscore(roi_data_current, axis=0)
    rdm = pdist(zscored_data, metric='correlation')

sns.heatmap(squareform(rank_percentile(rdm)), ax=ax, square=True,u
cmap='RdBu_r',
```

```
xticklabels=False, yticklabels=False, cbar_kws={'label':___
a'Dissimilarity'})
   ax.set_title(f'RDM for {roi}')

plt.tight_layout()
plt.show()

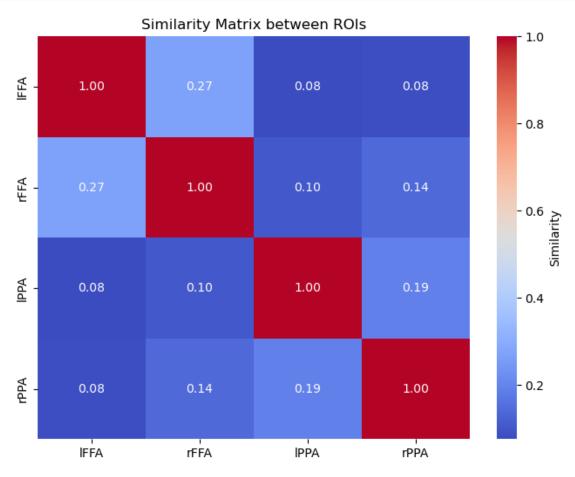
IFFA ROI shape: (96, 178)
rFFA ROI shape: (96, 407)
lPPA ROI shape: (96, 81)
rPPA ROI shape: (96, 81)
```

RSA allows us to compare the representational geometries of different ROIs. Compute the correlation between each pair of the four ROIs. Plot this similarity matrix. Which ROIs have the most similar representational geometries?

```
[165]: # Compute correlations between each pair or ROI RDMs:
    def get_upper_triangular(matrix):
        m = matrix.shape[0]
        r, c = np.triu_indices(m, k=1)
        return matrix[r, c]

similarity_matrix = np.zeros((len(roi_labels), len(roi_labels)))

similarity_matrices = {}
for roi in roi_labels:
    roi_data_current = roi_data[subject][roi]
    zscored_data = zscore(roi_data_current, axis=0)
    rdm = pdist(zscored_data, metric='correlation')
    similarity_matrices[roi] = 1 - squareform(rdm) # Convert from_u
    dissimilarity to similarity
```



Stack all four ROIs to create a single combined ROI for each subject 'SN' and 'TI'. What is the Spearman correlation between 'SN''s and 'TI''s representational geometries?

```
[166]: # Combine SN and TI ROIs into single VT ROI and compute RDMs:
    from scipy.stats import spearmanr
    combined_roi_SN = np.column_stack([roi_data['SN'][roi] for roi in roi_labels])
    combined_roi_TI = np.column_stack([roi_data['TI'][roi] for roi in roi_labels])

zscored_data_SN = zscore(combined_roi_SN, axis=0)
    zscored_data_TI = zscore(combined_roi_TI, axis=0)

rdm_SN = pdist(zscored_data_SN, metric='correlation')
    rdm_TI = pdist(zscored_data_TI, metric='correlation')

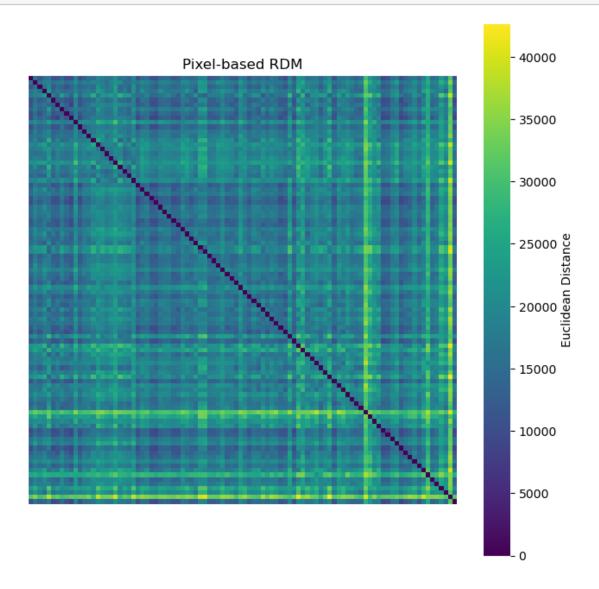
flattened_rdm_SN = get_upper_triangular(squareform(rdm_SN))
    flattened_rdm_TI = get_upper_triangular(squareform(rdm_TI))

spearman_corr, _ = spearmanr(flattened_rdm_SN, flattened_rdm_TI)

print(f'Spearman_correlation_between_SN_RDM_and_TI_RDM: {spearman_corr}')
```

Spearman correlation between SN RDM and TI RDM: 0.394897619062583

We can test different "model" RDMs according to how well they approximate a given neural. Here, for the sake of brevity, we'll construct an extremely simple RDM capturing low-level visual structure. Flatten each image file into a one-dimensional array of pixel values (across three color channels). Next, compute the pairwise Euclidean distances between these image vectors to construct an RDM capture low-level visual similarities. Plot this pixel RDM and compute it's Spearman correlation with 'TI's VT RDM?



Spearman correlation with TIs VT RDM: 0.02179628629185863

## 1.0.3 Problem 3: Voxelwise encoding analysis

In this problem, we'll return to *encoding analysis*, using regularized regression and out-of-sample prediction in individual voxels. We will use word embeddings derived from the natural language processing (NLP) model GloVe to map semantic encoding onto the brain. You can simply load the story\_transcript.txt file in a text editor to visualize the transcript for the spoken story by Carol Daniel. Each line of this file corresponds to a TR in the fMRI data. Next, we extracted word embeddings from GloVe for each word in each TR. For TRs containing multiple words, we averaged the embeddings. Finally, we horizontally stacked the embeddings at lags of 2, 3, 4, and 5 TRs (3, 4.5, 6, and 7.5 seconds relative to word onset) to account variable hemodynamic lags

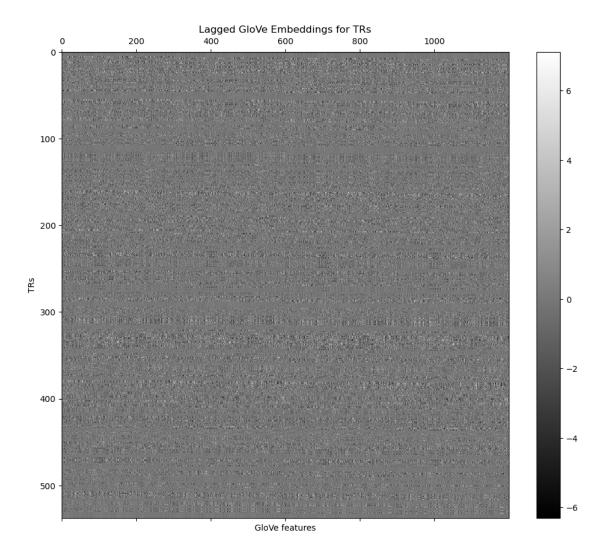
(this is effectively a finite impulse response model). Inspect and interpret the shape of the word embeddings, and visualize this matrix.

```
[168]: from os.path import join import numpy as np import matplotlib.pyplot as plt import seaborn as sns import nibabel as nib import pandas as pd import json
```

```
[169]: import numpy as np
       import json
       from scipy.stats import zscore
       import matplotlib.pyplot as plt
       transcript_fn = 'story_transcript.txt'
       with open(transcript_fn, 'r') as file:
           transcript_trs = [line.strip().split(',') for line in file]
       # Load GloVe embeddings
       with open('glove embeddings.json') as f:
           glove = json.load(f)
       # Load list of standard stop words
       stopwords = np.load('nltk_stopwords.npy').tolist()
       # Set GloVe embeddings dimension
       glove_dim = 300
       embeddings = []
       for words in transcript_trs:
           embeddings tr = [glove[word.lower()] for word in words if word.lower() not__
        →in stopwords and word.lower() in glove]
           if embeddings tr:
               embeddings_tr = np.mean(embeddings_tr, axis=0)
           else:
               embeddings_tr = np.zeros(glove_dim)
           embeddings.append(embeddings_tr)
       embeddings = np.stack(embeddings, axis=0)
       stim_trs = len(embeddings)
       # Incorporate temporal lags
       lags = [2, 3, 4, 5]
```

```
embeddings_lag = []
max_lag = max(lags)
for lag in lags:
   prepend = np.zeros((lag, glove_dim))
    # Remove the last 'lag' rows to keep the shape consistent
   lagged = embeddings[:stim_trs - lag]
   lagged = np.vstack((prepend, lagged))
    embeddings_lag.append(lagged)
min_trs = min([embed.shape[0] for embed in embeddings_lag])
embeddings_lag = [embed[:min_trs] for embed in embeddings_lag]
embeddings_lag = np.hstack(embeddings_lag)
print(embeddings_lag.shape)
embeddings_lag_zscored = zscore(embeddings_lag, axis=0)
plt.figure(figsize=(12, 10))
plt.matshow(embeddings_lag_zscored, fignum=1, aspect='auto', cmap='gray')
plt.xlabel('GloVe features')
plt.ylabel('TRs')
plt.title('Lagged GloVe Embeddings for TRs')
plt.colorbar()
plt.show()
```

(538, 1200)



We used fMRI to measure a subject's brain activity while they listened to the spoken story. Here, to reduce computational demands, we have spatially downsampled the fMRI data using an atlas containing 400 parcels. That is, for each parcel, we averaged the voxel time series within that parcel. Rather than fitting encoding models to tens of thousands of voxels, we'll fit our encoding model to each of the 400 parcels. Load in the story\_parcels.npy dataset as well as the story\_atlas.nii.gz NIfTI image from which the parcels were derived (for later visualization).

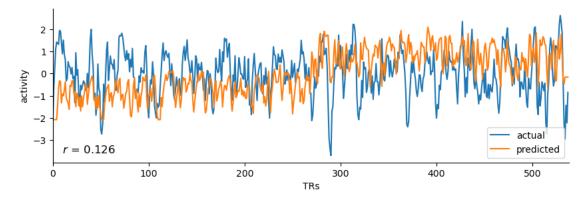
```
[170]: import nibabel as nib
from nilearn.maskers import NiftiLabelsMasker
# Load in parcel time series:
parcels_fn = 'story_parcels.npy'
story_parcels = np.load(parcels_fn)

# Load the story atlas NIfTI image
atlas_fn = 'story_atlas.nii.gz'
```

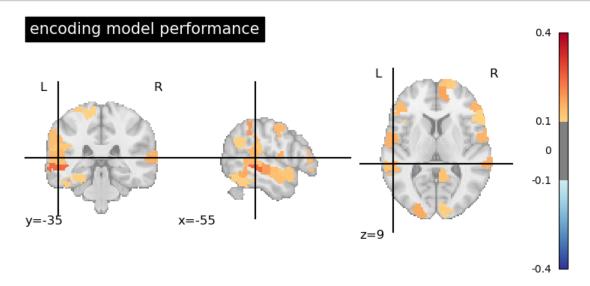
```
story_atlas = nib.load(atlas_fn)
story_img = story_atlas.get_fdata()
```

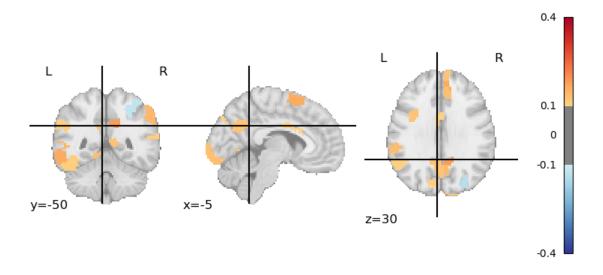
Our word embedding "model" is much wider than the number of samples, so we'll need to use regularization and out-of-sample prediction to mitigate overfitting. We'll use ridge regression to fit encoding models to predict the parcel time series from the word embeddings. First, set up an split-half outer cross-validator using KFold with n\_splits=2; next, set up an inner cross-validator using KFold with n\_splits=5 to perform grid search for the alpha hyperparameter using 5-fold cross-validation within each training set of the otuer loop. Initialize your RidgeCV estimator with the inner cross-validator and the following grid of alphas: alphas = [0.1, 1.0, 10.0, 100.0, 1000.0, 10000.0]. For each training and testing split of the other cross-validation loop, fit the ridge model on the training set of embeddings and parcel time series, and generate predicted parcel time series from the test embeddings. Compile these predicted parcel time series for model evaluation in the next step:

To evaluate our encoding model's predictions, correlate the predicted parcel time series with the actual parcel time series for each parcel.



Finally, to visualize the performance of our semantic encoding model on the brain, we need to use the atlast NIfTI image to convert from parcels back to the original brain image. You can start by creating an empty brain image (i.e. zeros) the size of the atlas image. Next, loop through each parcel and insert the prediction scores (i.e. correlations between actual and predicted parcel time series) into all voxels where the atlas correponds to that parcel label. Convert this image to a NIfTI image and visualize with plot\_stat\_map; you may want to set a particular vmax and use a threshold to exclude voxels with poor prediction performance for the sake of visualization.





[]: