

Using Support Vector Machine and 3D Convolutional Neural Networks to Predict Brain States in Task-Based Functional Magnetic Resonance Imaging (fMRI)

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You can find the work we did in our Repository on Github at: [Team Brainiacs](#)

The landing page gives a taste of what we did in this project: [Team Brainiac Landing Page](#)

INTRODUCTION

Real-time functional magnetic resonance imaging neurofeedback (rtfMRI-NFB) is a technique used in functional magnetic resonance imaging (fMRI) to measure an individual's control over target brain activity. Brain computer interfaces are used to provide subjects with neurofeedback during a scan in the magnetic resonance imaging (MRI) machine in order to help subjects learn to modulate their neural activity during a given task[1]. Researchers use these tools to help study brain states and regions of interest (ROI) in the brain that may be involved in the given task. Traditionally, research using these tools has been analyzed using a univariate approach. Researchers have been studying the application of multivariate analyses[2] on rtfMRI-NFB using machine learning models, such as Support Vector Machines (SVM), and deep learning models, such as convolutional neural networks (CNN). The machine learning and deep learning methods hold promise in clinical and exploratory research.

Real-time fMRI has potential to aid mental health experts in helping people who have substance use disorders learn to better regulate their reward system leading to rehabilitation and recovery. Research in rtfMRI for substance use disorders typically uses an a priori knowledge of predefined ROI centered around the left and right Nucleus Accumbens (NAcc) to analyze the output from images. Subjects studied are provided feedback using a brain computer interface based on percent signal change in voxel activity from the previous timepoints in these regions. However, little is known whether more regions exist that could be involved in reward circuitry. It may also be the case that there are individual differences in the mental strategies applied to the increase or decrease of neural activation.

We are interested in studying reward circuitry and regions of interest in task-based rtfMRI using these brain images to conduct a data-driven analysis. Through the use of SVM, we hope to discover other brain regions that may be active during the up and down regulation during rtfMRI. We are also interested in how a deep learning approach would compare against our predictions with SVM models.

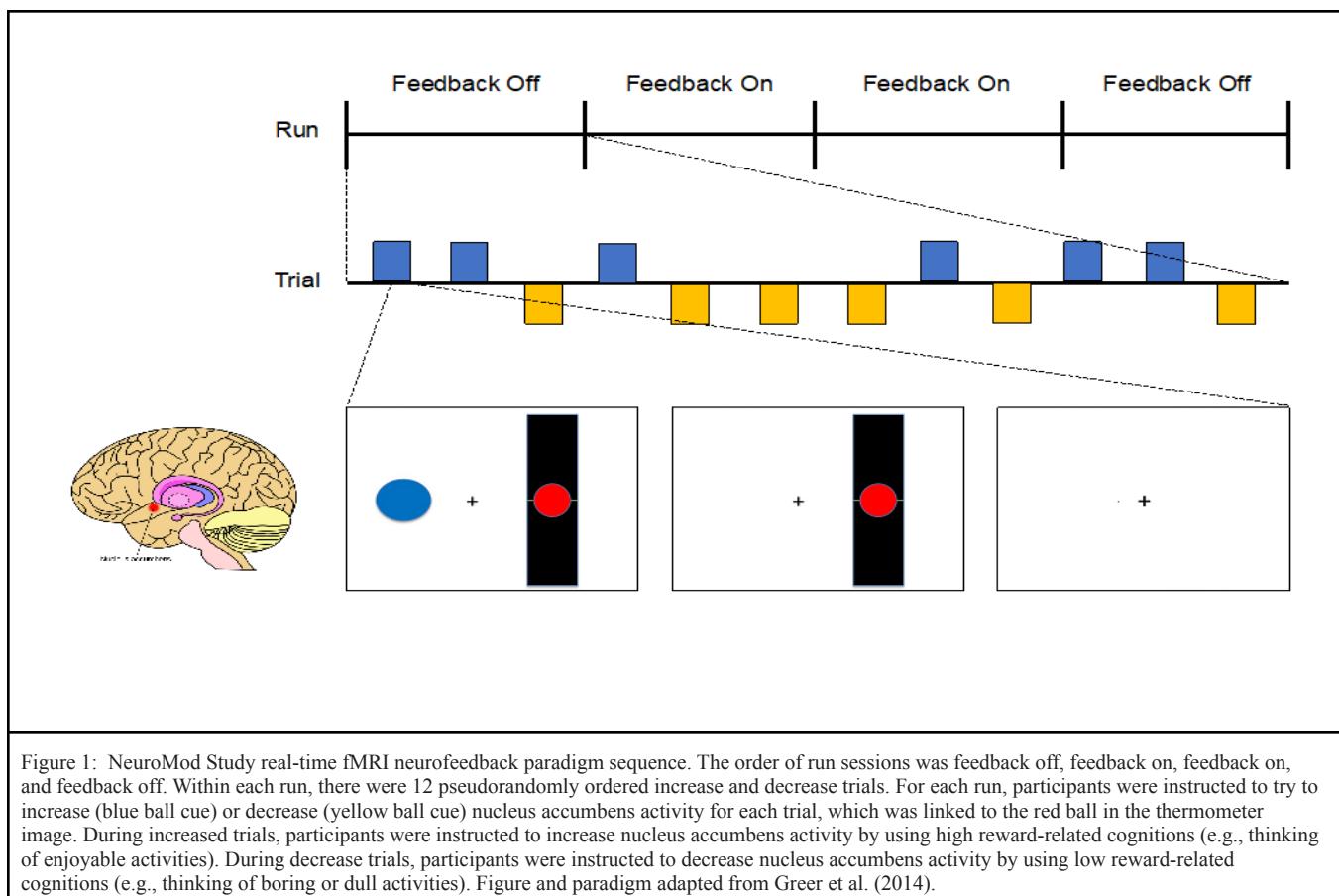
MOTIVATION

We are collaborating with the University of Michigan Medicine Psychology Research Department under the research of Principal Investigator Dr. Meghan Martz, PhD. This opportunity was presented to us by one of our authors, Mary Soules, who works as an Application Programming Analyst Senior in the UM research lab processing and analyzing brain images captured through Dr. Martz study of real-time fMRI. We have three approaches to looking at the fMRI data captured through this study. The first approach is using SVM on single subjects to look at individual differences in brain activation and metrics associated within a single subject. This approach would allow us to use trained models at the individual level to help in personalizing treatment for addiction and help individuals up or down-regulate areas in the brain that could be playing a role in their addiction. The second approach is using SVM at the group-level to study whether we can predict brain states between adolescents and young adults and apply it across subjects. The third approach is using a deep-learning model at the group level to see if we can provide better predictions to brain-state data.

The brain data collected uses a task that captures percent signal change in the Nucleus Accumbens (NAcc) since research shows that the NAcc is a key area of interest in the brain associated with the reward circuitry and may play a vital role in drug and alcohol addiction[3]. This is an a priori approach to an area we think may play an active role in addiction. However, we are also interested in whether other regions are involved in addiction, and whether individual differences exist at the group and individual level. Additionally, we want to investigate

whether there are key differences in brain states between adolescents and young adults. Since young adults are better able to control inhibition and impulse through their reward circuitry than adolescents, they are less vulnerable to risk-taking behaviors and are less likely to show substance use behavior. As adolescents mature, these differences could be heightened or diminished. Those that show potential for substance abuse could, through maturity, learn to control it while those that do not could go on to develop substance abuse disorders. The product from these analyses would be to develop classifiers that are able to predict brain states from the fMRI data between groups of adolescents and young adults, and within individuals, and deploy these models in production to be used in real-time. This would improve the feedback for up and down-regulation training to participants in the scanner and help researchers uncover potential ways to use this as therapy for addiction.

TASK DESIGN



The data we use are rt-fMRI images of temporal brain states that are captured during tasks that specifically target the NAcc. (Figure 1) Subjects are taught to increase or decrease their brain activity in the reward system through a real-time NAcc task that calculates the percent signal change of the NAcc during the task. The task consists of four runs. In run 1, subjects either see a blue or yellow dot on the brain computer interface to up-regulate or down-regulate the NAcc but do not receive visual feedback from the red dot in the thermometer on how well they are able to up or down-regulate, nor do they receive feedback from researchers after the scan is completed. In run 2, visual feedback from the red dot in the thermometer is provided on the brain computer interface to subjects based on the percent signal change of the NAcc while they are up or down-regulating, and they also receive

feedback by the researchers on how well they did once the scan is completed. In run 3, subjects receive the same feedback stimuli as in run 2 from the brain computer interface as well as from researchers. Lastly, in run 4, no feedback is provided by the red dot in the thermometer on the brain computer interface nor by researchers once the scan is completed. In this run, subjects are expected to up or down-regulate without these cues and are expected to use their own strategies learned from prior training.

DATA

The fMRI brain data we used for these analyses was obtained from 52 voluntary, healthy subjects, 33 of which are adolescents ranging from 14 -16 years old (50% female) and 19 of which are young adults ranging from 25 - 27 years old (50% female), from the University of Michigan Medicine Research lab. The data is proprietary and every measure has been taken to ensure privacy of all subjects. The data structure is temporal, such that a single subject MR run contains 144 volumes, also known as time points, of whole brain image scans. Each scan represents a 3D brain image where the pixels are 3D voxels. These pixels represent the feature space for our machine and deep learning models. There are 79 voxels in the x direction, 95 voxels in the y direction and 79 voxels in the z direction. The data is gathered and preprocessed in SPM 12 then flattened so that each brain volume of a given run is flattened to 592,895 voxels per time point. Each subject participates in four runs of equal duration, and 84 time points in a given run are dedicated to the subject attempting to down-regulate or up-regulate based on the task given in the MR machine. From the 84 time points, there are 42 labeled time points for down-regulating and 42 labeled time points for up-regulating. The remaining 60 time points are rest periods in the scanner. Therefore, for a given subject there are 576 total brain volumes, 336 of which are used in our analyses and are labeled for regulation. Our total dataset includes 17,472 brain volumes of 592,895 voxels each. We used other data in this study, such as data with masked out regions and data of regions of interest. We will explain these data further in the Masking section. See Appendix I for a visual representation of our data.

QUALITY CONTROL AND PRE-PROCESSING

All data captured for this analysis went through a stringent quality control and preprocessing pipeline before being uploaded to AWS for further analysis. All pre-processing steps were employed using SPM12, which was first released 1st October 2014 and last updated 13th January 2020. In this time, substantial theoretical, algorithmic, structural and interface enhancements were made in the software. The steps for preprocessing include manually inspecting images in raw form to look for any deformities or anomalies that would make the images unusable for analysis. If a run did not pass the quality control criteria, it was excluded from analysis and the subject was excluded from the dataset. All subjects and runs that passed the quality control criteria were put through a pre-processing pipeline using SPM12. (Table 1) Individual runs were inspected to make sure co-registration and warping were consistent across structural and functional images. Since individuals in the scanner may move and introduce noise into the images, we want to exclude runs that have a lot of motion in them. This is done by creating a threshold and looking at motion from one brain image to the next and if a run exceeds the threshold, we exclude it from analysis. Because we only wanted individuals included that had four quality runs, only individuals that passed all of the quality measures were included in the dataset. Lastly, subject runs were flattened into a 2-D array and stored in a mat file to be uploaded to AWS for analysis.

Pre-Processing Steps	
Slice Timing Correction	Unlike a photograph that is captured in a single instance, a full brain image is captured over time. In order to analyze the brain, we need to apply slice-timing correction to be able to analyze the voxels of the brain as though they were captured simultaneously.
Realignment	If a person moves a lot during their scan, they could introduce noise to the data and one image to the next could be shifted. In order to correct this, we need to shift everyone's individual images in each run and over runs to the same space.
Co-registration	In order to put the fMRI images into a common space, we need to have their high-resolution structural images aligned with their functional images and this is achieved in this step.
Warping	In order to analyze between subjects, these data need to be in a common space. In our lab, we use the MNI152 brain to change the subject's functional data to a common space.

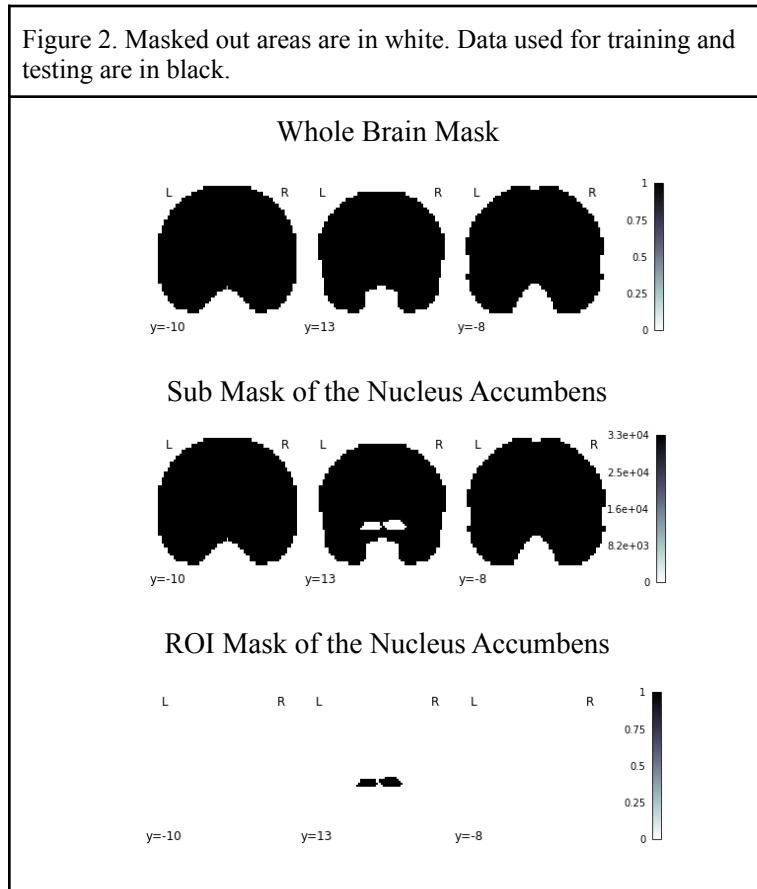
Table 1. Pre-processing steps. It should be noted SPM runs under MATLAB which requires a license. All of the pre-processing was done outside of this analysis. For more information about SPM and pre-processing: <https://www.fil.ion.ucl.ac.uk/spm/doc/>

MASKING

Masking fMRI data is an important preprocessing step where regions of the brain are removed to allow a focus on other regions of interest. Our mask data is 3D with the shape of (79,95,79) and was flattened before application to our brain data. We applied various masks to the original data and utilized a fourier transformation to ensure that proper masking coordinates in Python's numpy reshape package matched MATLAB's masking application technique. We used a whole brain mask in our first application for model training, which removes areas of the image that are not considered a part of the brain. We also studied other masking strategies, which remove certain areas of the brain. Such areas that we studied and removed for separate analyses include a submask of the Anterior Cingulate Cortex, Anterior Insula (Right-side), Nucleus Accumbens, and the Medial Prefrontal Cortex. Refer to Table 2. to better understand the significance of these regions. The other masks we studied include removing all brain data except for the region of interest. The same regions removed in the submasks are the same regions kept in the ROI masks. (Figure 2)

All brain masks were created outside of the repository using SPM 12. The whole brain mask was created using a single subject T1 brain image supplied by SPM. We used IMCALC, a function in SPM12, to set dimensions outside of the brain to values of 0, and masks within the brain to a value of 1. Individual ROI masks were created using the AAL toolbox for SPM12. Once masks were created, we had to run the SPM reslice tool to get masks into our correct dimensions (79,95,79) which are different from default dimensions (91,109,91). Once all the masks were in the correct dimension, we used IMCALC to subtract the individual ROIs to give us the submasks. These submasks contain the whole brain without the region of interest. The ROI masks were made in a similar fashion. We put all mask data into a .mat file along with the labels corresponding to time points and uploaded these with the brain data to AWS.

Regions of Interest	Table 2. Functions of the Regions of Interest
Anterior Cingulate Cortex	Known for error detection and monitoring; alerts other areas of the brain of this conflict to activate a resolution.[4] It is also an area of the brain involved in memory, motivation, decision-making, and cognitive inhibition. These functions are important in understanding substance use disorders.[5]
Anterior Insula (Right Hemisphere)	Involved in various brain disorders and serves as an area of the brain that creates subjective feelings. This is important for understanding how one prioritizes stimuli and salient information, thus, impacting motivation. With substance abuse, users seek hedonic experiences, influencing this motivational salience. Also linked to the ACC.[6]
Medial Prefrontal Cortex	Research has shown the Medial Prefrontal Cortex to be inhibited in subjects with substance use disorders.[7] It is an important part of self-regulation, inhibition, decision-making and conflict resolution/monitoring.[8]
Nucleus Accumbens	The Nucleus Accumbens regulates depression and addiction and is the reward center of the brain that activates dopamine and pleasure. This area of the brain has been studied as one that is activated in people with substance use disorders.



DATA STORAGE AND ENVIRONMENT

We used various types of data storage in this project. Once the lab preprocessing was completed, we uploaded the data to an AWS S3 bucket where each subject's data, label data and mask data are stored in their own folders. We would access the data from AWS for further processing like applying masks to each image as well as filtering the time points of the image by labels. Once these processes were completed we uploaded these files back to AWS. Eventually, it would make more sense to run these processing steps in real-time since we had to make various changes to the processing pipeline and it was not feasible to upload back to AWS after each iteration. In some cases, we were able to upload model and metric data to AWS. Due to pickle file formatting issues, we opted to save models locally, and some metric data back to the cloud.

We set up our work environment initially by connecting to a Docker container in Pycharm Student Edition, working locally and within Jupyter Notebooks, using Github for version control. Once we realized that the size of our data required RAM greater than what was capable locally, we opted to build our pipelines using a virtual environment provided to us with a subscription to Google Colab Pro+. All data training and analyses require enabling 'High-RAM' in the runtime settings in Google Colab, which provides 51GB of memory. Each step of the single subject, group-level and deep learning data pipelines is organized in python files and imported into Google Colab for execution and demonstration. The python files are organized by project analysis type and task, as are the notebooks that run these modules.

NORMALIZATION

Before we began training with machine learning models, we researched the most commonly used strategies for fMRI data normalization and consulted with domain experts. We decided to perform an exploratory analysis on the different normalization strategies applied to fMRI data. (Figure 3)

Within these analyses, we plotted voxel distributions on subjects split by age groups and looked at distributions across MR scan runs. We also performed cross validation on each different normalization strategy and compared the accuracy results. These processes were repeated twice as we later realized from our research on fMRI analyzes that the temporal aspect of fMRI data requires detrending as the data drifts over time points. We looked at the voxel distributions on non-detrended unnormalized data, detrended unnormalized data, detrended percent signal change normalized and detrended Z-score normalized data. The non-detrended, unnormalized data was not normally distributed, and that adding a normalization technique to the data would improve the distribution. Detrending unnormalized data surprisingly improved the variance of the data, as did applying percent signal change to detrended data. However, the data in these graphs are still noisy, so we lastly applied Z-score normalization to the detrended data and found that this distribution was the least noisy with lower variance. (Figure 4, Appendix II)

Accuracy Scores of Normalization Strategies

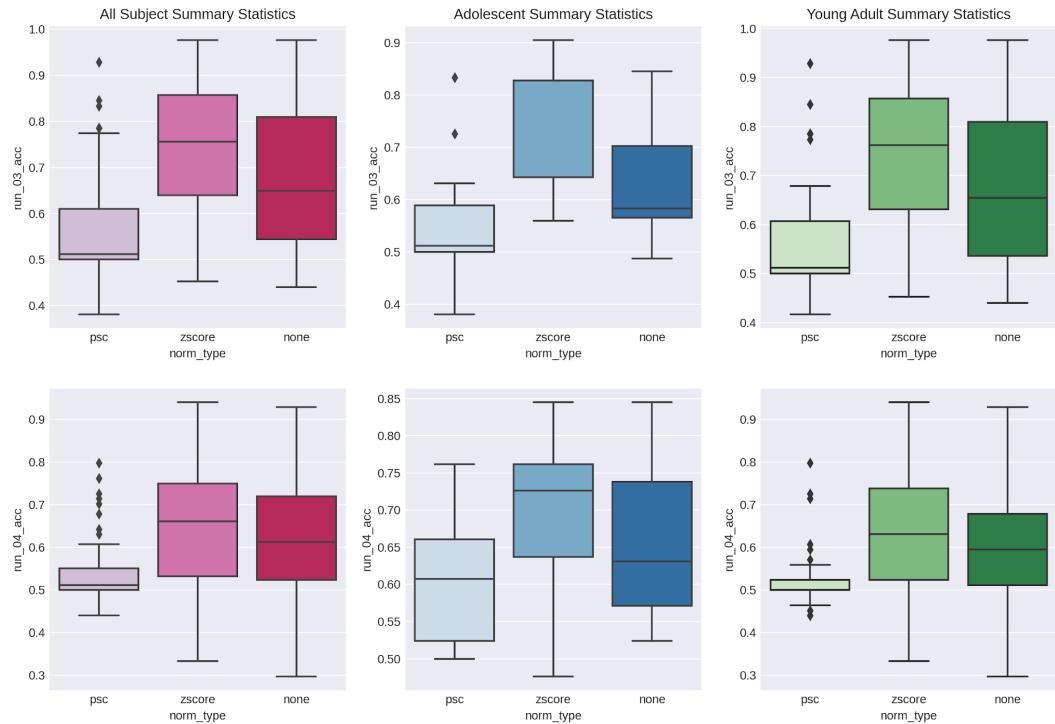
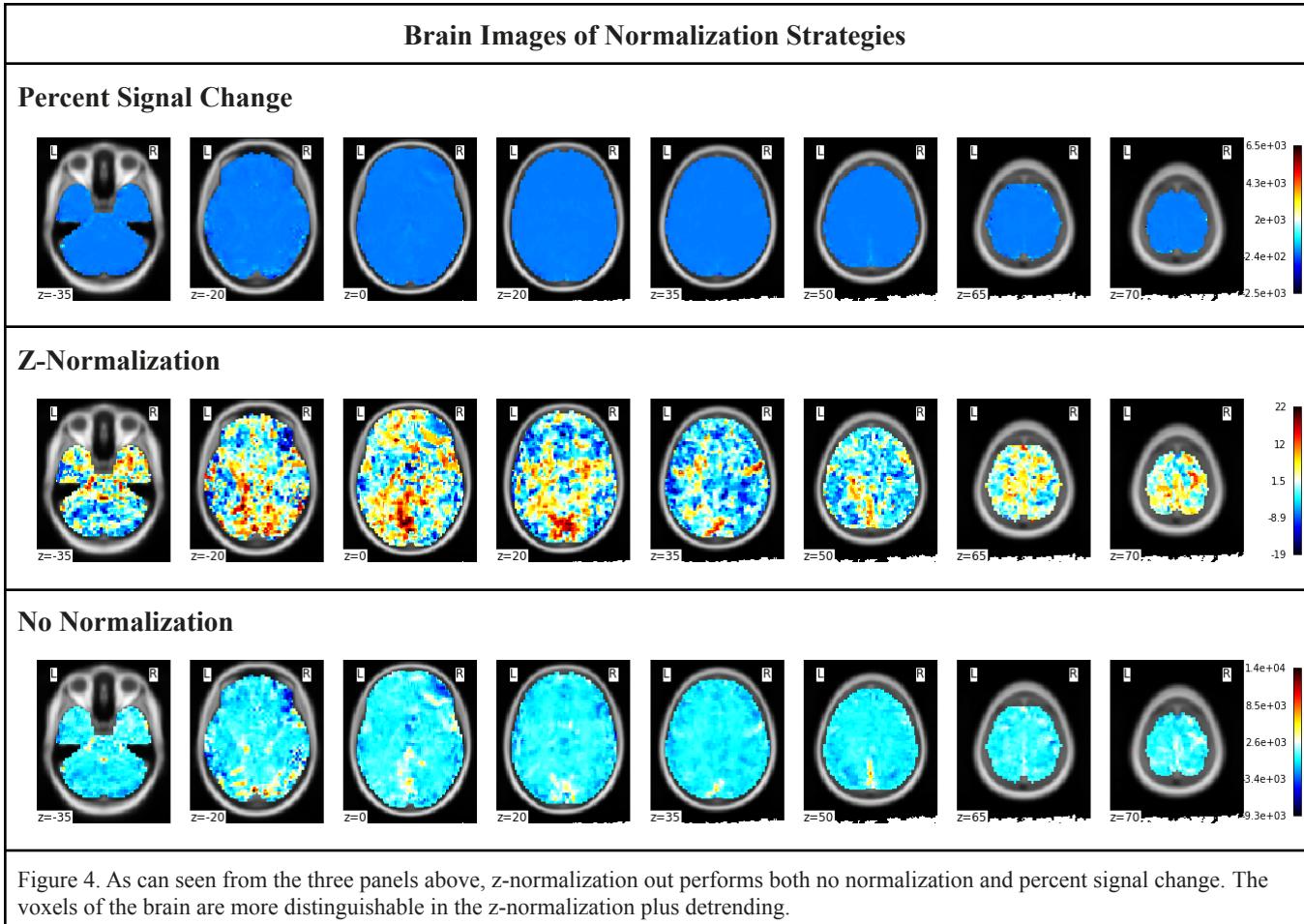


Figure 3. As seen above, we can see that z-normalization on detrended data out-performs percent signal change and no normalization.



SINGLE SUBJECT SVM

Methods

Data Processing and Model Building

The single-subject level pipeline reads in each subject's data, the masking data and the label data from AWS using a dictionary of key paths. We then applied the mask and label data to each subject's run data separately, which resulted in each run containing 84 timepoints and 237,979 features. Once we got our final masked data per run, we detrended and z-score normalized the time series voxels using Nilearn's signal clean package. The detrending and normalization was done per subject per run.

Once the data was normalized, we built our models training on run 2 using Scikit-Learn's `svm.SVC` classifier. The parameters we chose were found during the cross-validation phase of the single-subject data exploration. The final customized model parameters we chose were `C=5`, `gamma='auto'`, and `kernel='rbf'`.

Cross-Validation Results

To ensure we were getting the best parameters for our models, we chose to use Scikit-Learn's GridSearchCV. Data for the cross validation grid search was processed the same way as for model building. We chose to only look at whole brain masking in the cross-validation and apply to all masked models. Each individual's run 2 went through a 5-fold cross validation and the data was summarized across all individuals to find the best parameters for use across all subjects. (Figure 5, Figure 6)

We also chose to summarize the data and not customize the parameters per subject which could be an area for further research.

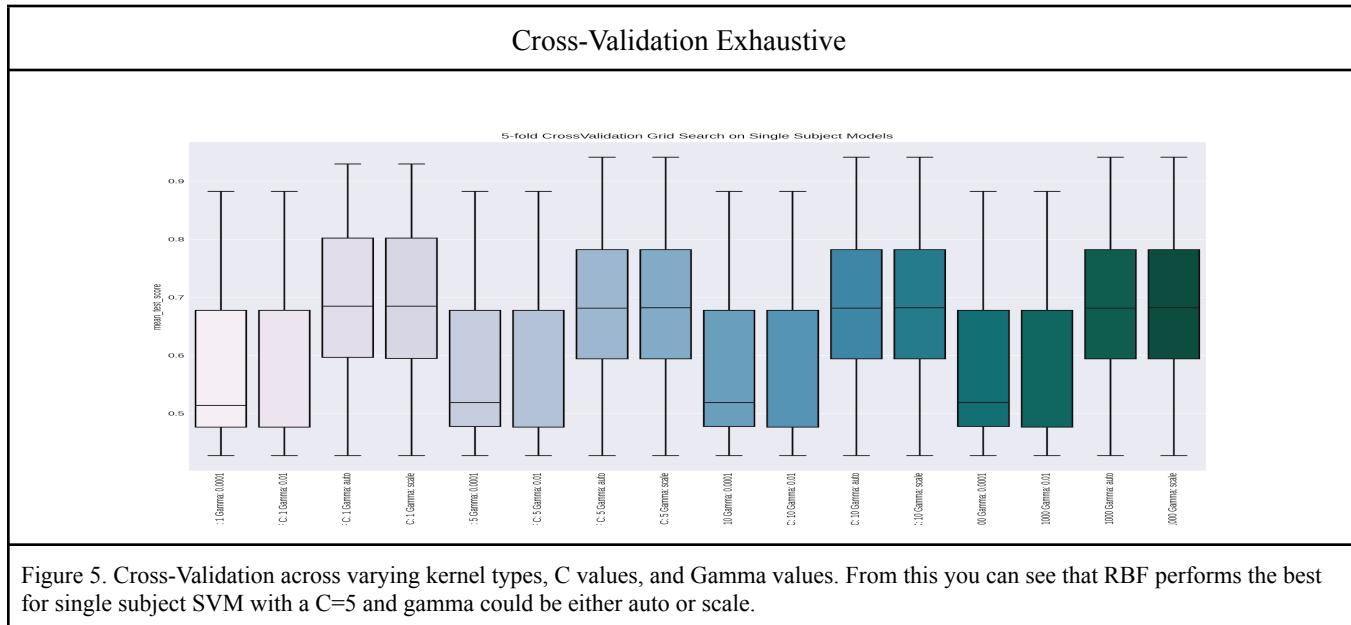


Figure 5. Cross-Validation across varying kernel types, C values, and Gamma values. From this you can see that RBF performs the best for single subject SVM with a C=5 and gamma could be either auto or scale.

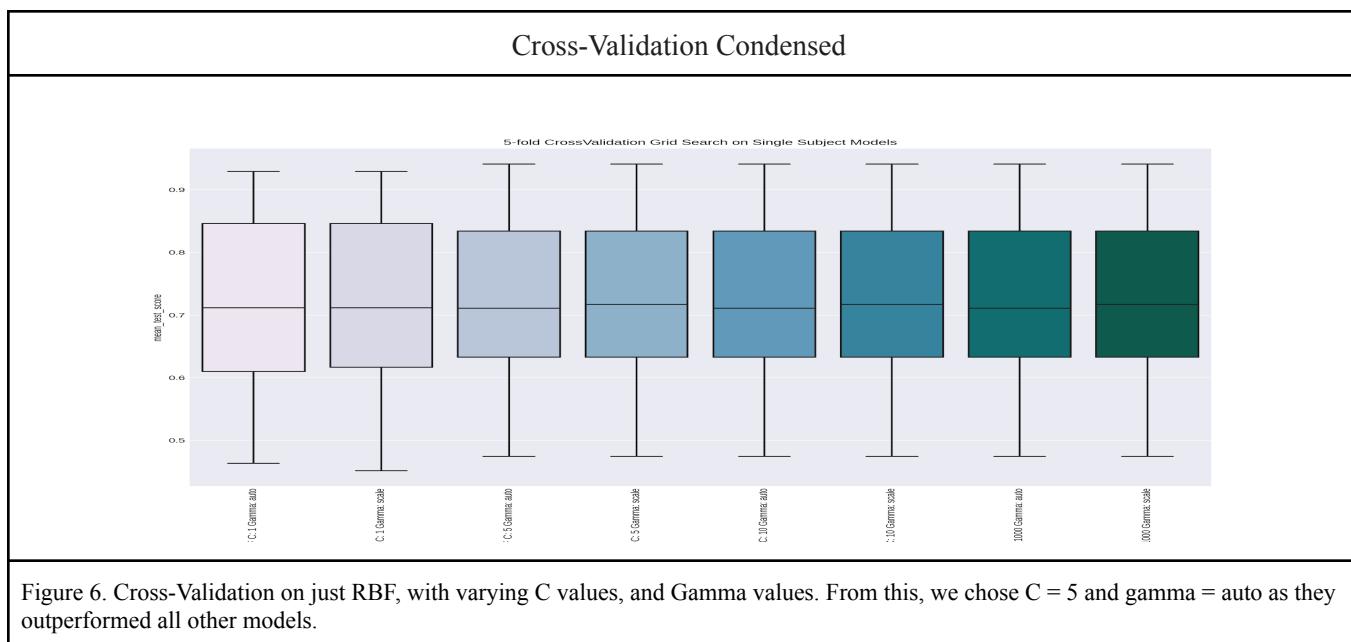


Figure 6. Cross-Validation on just RBF, with varying C values, and Gamma values. From this, we chose C = 5 and gamma = auto as they outperformed all other models.

Single Subject SVM Results

Being able to train SVM models on individuals that may utilize different brain regions for up and down-regulation is an important data-driven approach to studying addiction. Through this we may be able to customize therapies for individuals to help aid in recovery from substance abuse disorders. In these analyses, we were able to explore whether we could predict brain states on the individual level. As seen in Figure 1, there are four distinct runs in the task design. For this analysis, we chose to use run 2 data for training our classifier since this run is the first to provide subjects with feedback on their ability to up or down-regulate the NAcc while in the scanner. We also chose to test separately on run 3 (another feed back run) and run 4 (a no feedback run). This would give us insight into whether there were differences in individuals' abilities to maintain what they had learned in the feedback runs. Further analysis was done looking at the different masks, which either removed ROIs from brain data or only trained on ROIs to see if areas that have been studied and thought to be important in substance abuse disorders had an effect on the model's ability to predict brain states in individual subjects.

As shown in Figure 7, we see a lot of variation in model performance across individuals. Some of the individual models performed very well at predicting brain states while other individual model performances were at or below chance. This could be due to the fact that there are individual differences in how well an individual is at regulating their brain states and may need further training on how to regulate. We investigated the models performance looking at sensitivity and specificity, which looks at how well the model predicts the positive and negative class, respectively. In the single subject models, the mean sensitivity and specificity were similar, which suggests the model was good at distinguishing the two classes when it was able to predict well.

Mean Metrics for Single Subject SVM for Run 3 Feedback Run						
Mask Type	Run	Accuracy	Sensitivity	Specificity	Precision	AUC
mask	run_03	0.74	0.741	0.739	0.746	0.809
masksubACC	run_03	0.741	0.74	0.742	0.748	0.808
masksubAI	run_03	0.741	0.742	0.739	0.746	0.808
masksubNAcc	run_03	0.74	0.74	0.739	0.746	0.809
masksubmPFC	run_03	0.74	0.741	0.739	0.746	0.808
acc_aal	run_03	0.68	0.695	0.664	0.678	0.737
anterior_insula_aal	run_03	0.613	0.632	0.595	0.611	0.644
mPFC	run_03	0.612	0.634	0.588	0.61	0.655
nacc_aal	run_03	0.636	0.641	0.631	0.636	0.681

Mean Metrics for Single Subject SVM for Run 4 No Feedback Run						
Mask Type	Run	Accuracy	Sensitivity	Specificity	Precision	AUC
mask	run_04	0.663	0.665	0.662	0.665	0.712
masksubACC	run_04	0.665	0.667	0.664	0.667	0.712
masksubAI	run_04	0.664	0.665	0.663	0.666	0.712
masksubNAcc	run_04	0.663	0.665	0.663	0.666	0.712
masksubmPFC	run_04	0.664	0.665	0.663	0.666	0.712
acc_aal	run_04	0.602	0.623	0.58	0.599	0.647
anterior_insula_aal	run_04	0.557	0.563	0.554	0.557	0.578
mPFC	run_04	0.57	0.588	0.552	0.567	0.592
nacc_aal	run_04	0.569	0.577	0.561	0.572	0.592

Table 3: Shows mean metrics measures for feedback vs. no feedback runs used to test the model. Across all metrics and different masking strategies, we can see that the model was better able to predict on the feedback run.

Single Subject Model SVM Boxplots Showing Variation of Metrics of Individuals

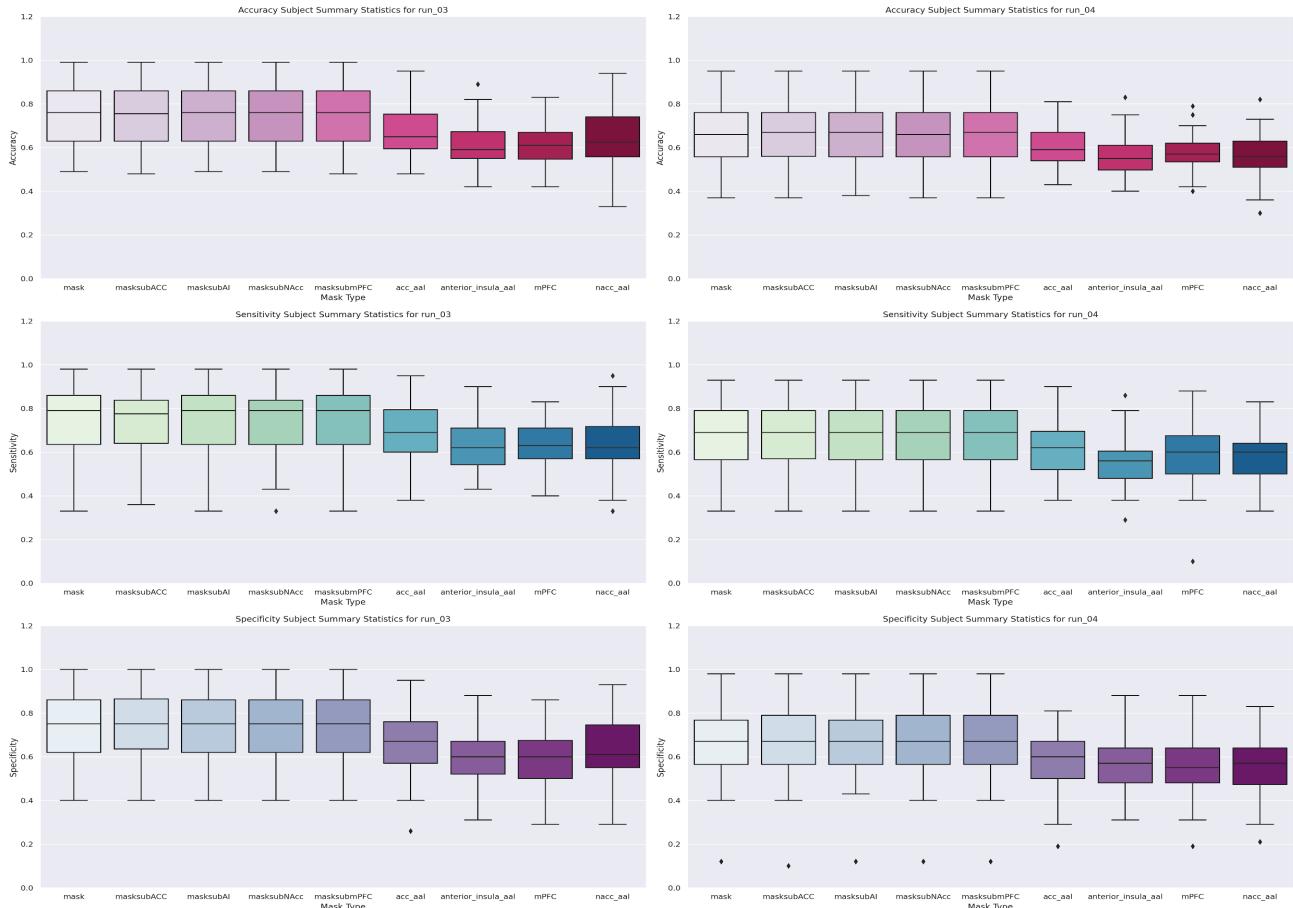


Figure 7. Boxplots showing variation in the predictions for single subject models. Top Panel is showing accuracy scores, the middle panel is showing sensitivity scores, and the bottom panel is showing specificity scores. The left panel shows the metrics on feedback run and the right panel shows metrics on the no feedback run.

We wanted to also look at how removing different ROIs from the brain affected prediction. The subtraction of ROIs didn't affect model performance suggesting that there may be individual differences in areas of the brain a person uses to control their brain states. (Table 3, Figure 7) Furthermore, individual ROIs showed a decrease in all metrics which may further suggest that there may be more regions than just that interconnect when a person tries to regulate their brain state.

In investigating the ability to maintain up or down-regulation of brain states, we analyzed performance of the predictions over the feedback vs no feedback runs. Individual models were better able to predict on the feedback run versus the no feedback run in all our metrics (Table 3). This could suggest that the individuals may need more training to maintain reward regulation when not being given feedback. The ROC curves and AUC scores show that in the no feedback runs, the model performance did not evaluate as well as in the feedback run. (Table 3, Figure 8)

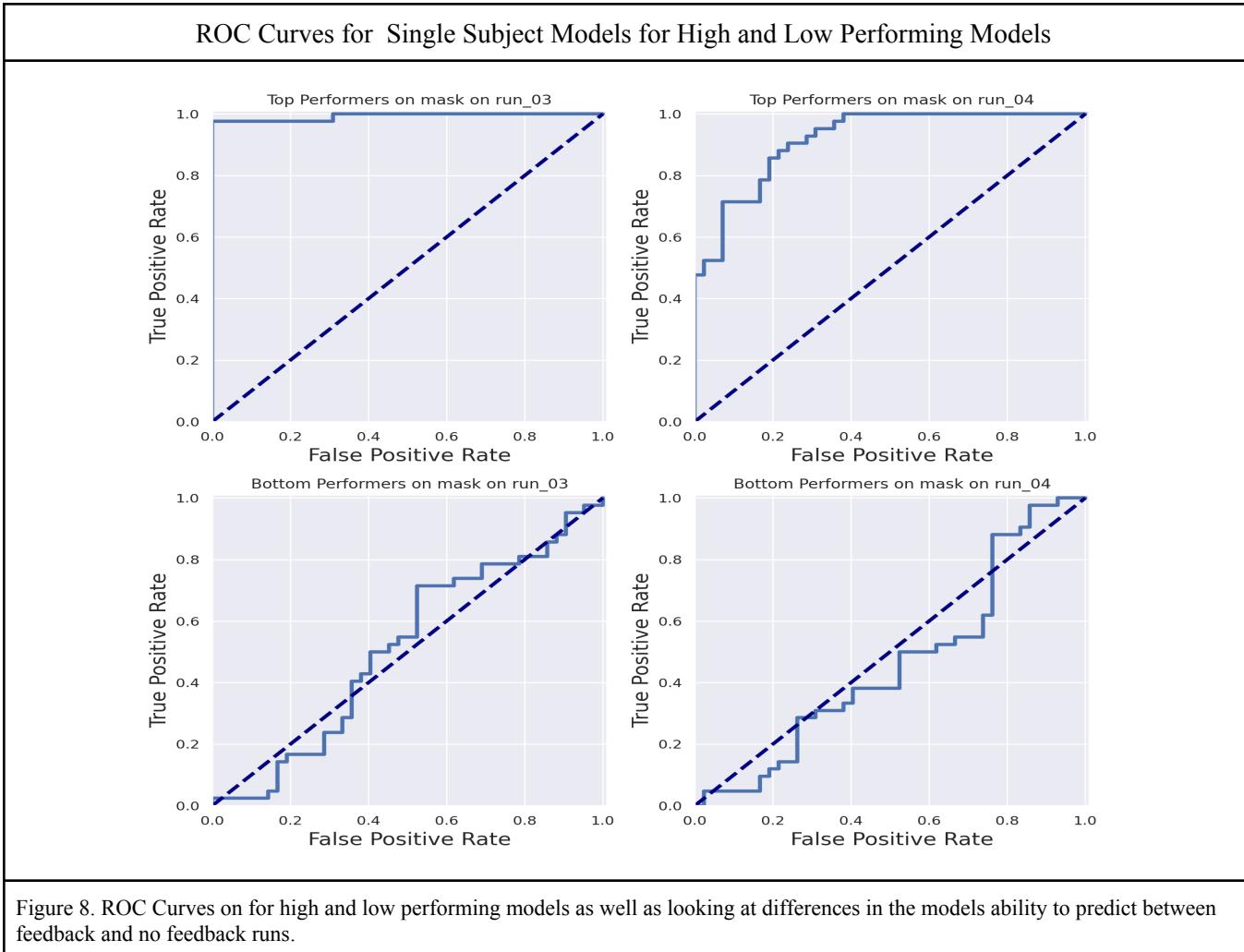
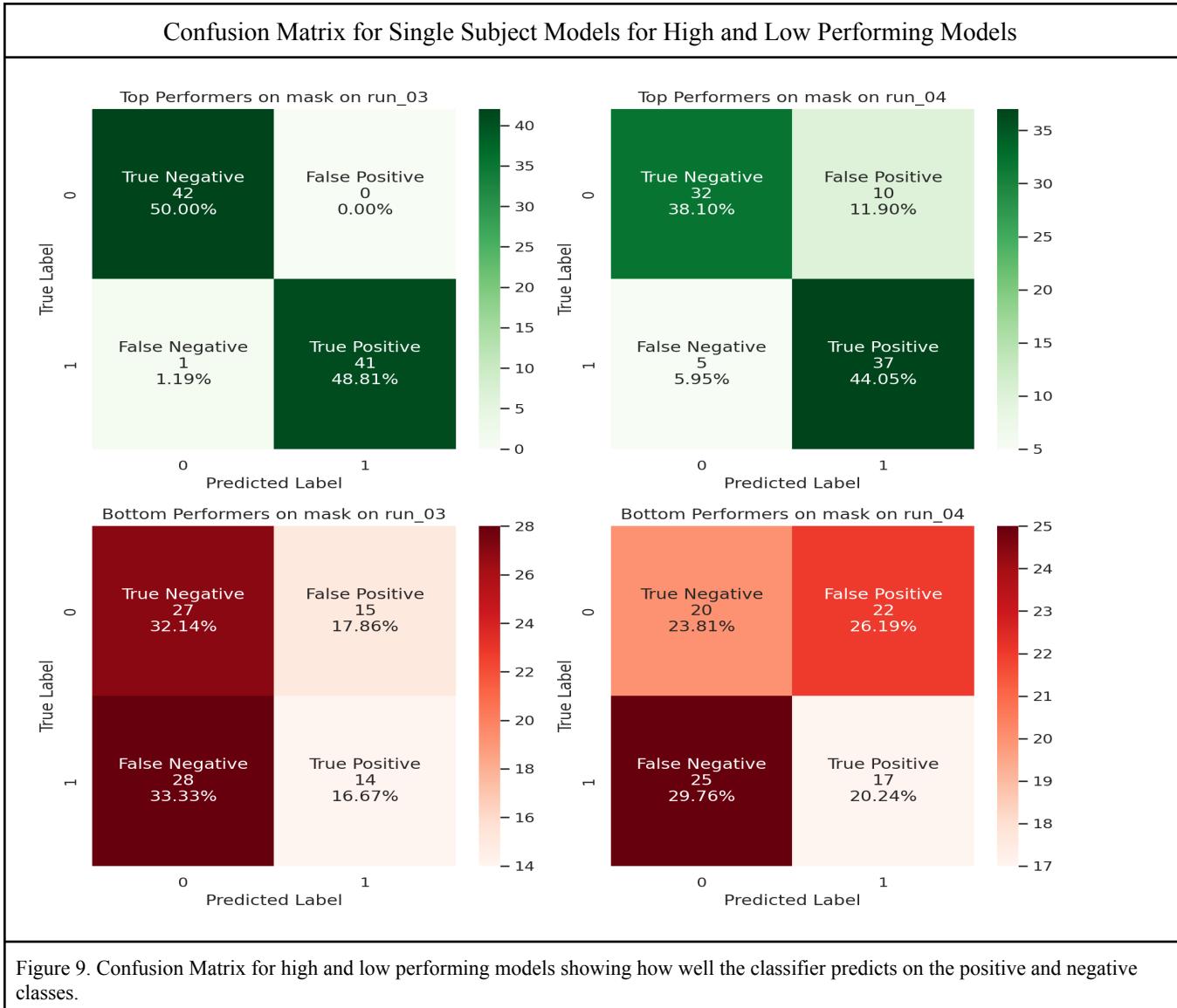


Figure 8. ROC Curves on for high and low performing models as well as looking at differences in the models ability to predict between feedback and no feedback runs.

In order to understand how individual models performed differently, we explored data from individuals with good model performance and individuals with poor model performance. We used different visualization and metric techniques to explore these differences. We looked at model ROC curves and AUC scores to compare model performance between these individuals for the whole brain masked models. ROC curves showed that the model was unable to distinguish well between the two brain states for models that perform poorly.

To understand how the individual models were differing, we looked at how the model predicted labels for the up and down-regulation classes. In the models that were able to predict well, the model was able to evenly classify both up and down regulation brain states. The poor performance model had trouble classifying both up and down, but had a harder time distinguishing the positive class. This could suggest that this person was not able to properly regulate their brain state. We also show that the model was able to more accurately predict both the positive and negative class in the feedback run versus the no feedback run. Again, this suggests that individuals are having a harder time carrying over from feedback runs to no feedback runs. (Figure 9)



In order to understand how the classifier was making its decisions on the two individual models, we explored the difference in decision function scores, which are measurements of the distances of the significant voxels from separating hyperplanes the classifier was using to make its predictions. We plotted these against the true labels over the time course. You can see that the low performing models had a harder time distinguishing between the two classes (shown in the bottom panel of Figure 10). We also can see that it had a harder time distinguishing the no feedback run (seen in the right side of Figure 10). Which may suggest that individuals are not able to carry over their training from the feedback runs. You can also see how different the brain maps which are used to show how significantly different regions of the brain map on to up and down-regulation states. (Figure 11)

Decision Function Scores for High and Low Performing Models

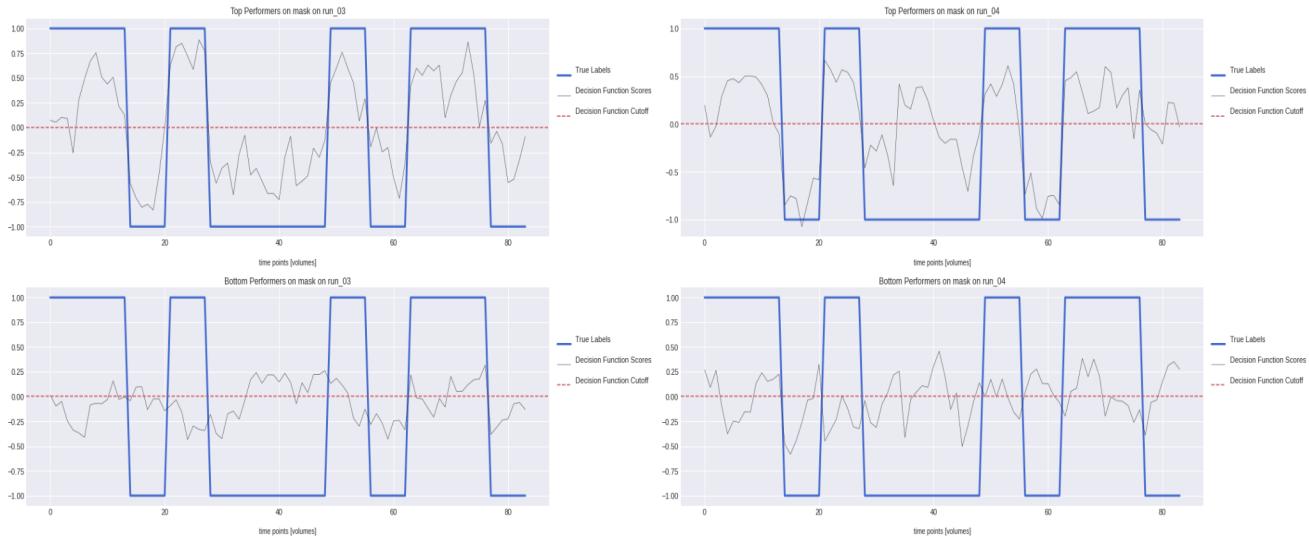


Figure 10. Decision Function Scores for top and low performing models. Top panel shows decision scores on a top performer and the bottom panel shows decision scores on low performers. The left panel shows feedback run and the right panel shows no feedback run.

Visualize Voxel Patterns for Whole Brain Masks for High and Low Performance Models

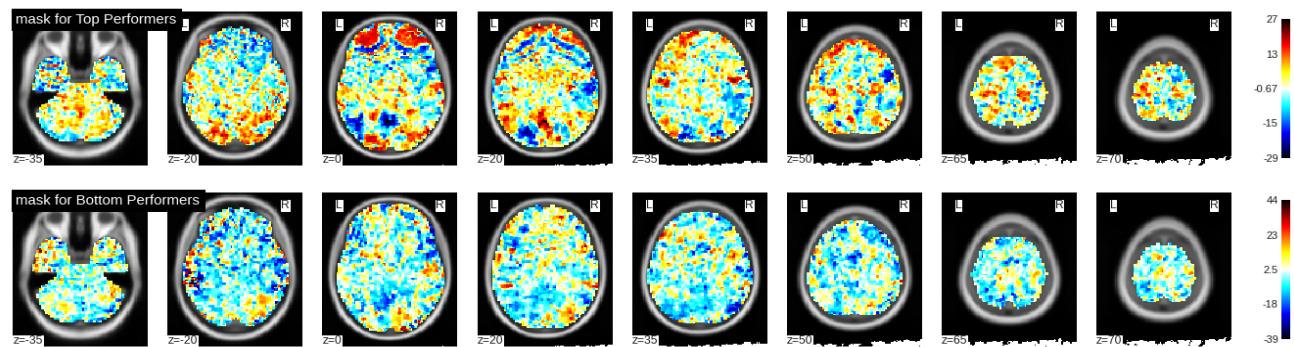


Figure 11. Showing voxel distributions across the whole brain. The top panel shows distributions for the top performing model and the bottom panel shows voxel distributions for the low performing model. The high performer has more clustered voxels than the low performing model. This suggests that the individual had greater control in up and down-regulating their brain states.

GROUP LEVEL METHODS AND RESULTS

Data Processing, Preparation and Model Building

The group level pipeline is set differently than the single subject pipeline. Some tasks are similar in that we read in subject, mask and label data from AWS using a dictionary of key paths and then apply the specified mask to the data and filter out the resting time points per subject per run. The data for group analyses is split between adolescents and young adults. Adolescent subject IDs begin with a ‘1’ and young adult IDs begin with a ‘3’. Once groups are separated, we perform an 80-20 split for training and test data and do this using subject IDs to ensure that each dataset contains unique IDs to prevent data leakage. Adolescent training is performed on 26 subjects and tested on 7 subjects for runs 2 and 3 in the scanner. Young adult training is performed on 15 subjects and tested on 4 subjects on runs 2 and 3.

Once the data is split, masked and labeled filtered, it is then detrended and scaled using z-score normalization using Nilearn’s signal.clean package by subject by run over the time points. The training and test subject timepoints are concatenated separately. This provides a data shape of (4368, 237979) and (1176, 237979) for adolescent training and test subjects, respectively. Young adult data shapes after concatenation are (2520, 237979) and (672, 237979) for training and test sets, respectively. We chose to use Scikit-Learn’s svm.SVC classifier for model training with parameters for class balance, max_iter of 1000, probabilities as True. The regularization C parameter of 10 was chosen for both young adult and adolescent and gamma was set to ‘scale’ and ‘auto’ for adolescent and young adult, respectively. The data metrics are saved as a dictionary in a pickle file and uploaded to AWS. The models are saved locally to Google Drive.

Time Series Cross-Validation with Halving Gridsearch

Preparing the data for cross-validation was performed differently than for model training. We wanted to ensure that a held out test set for model training would not be included in the cross-validation study. We separated the data on subject IDs between adolescent and young adult subjects and split the data by subject IDs into 80% training for cross-validation and 20% for held-out test sets for each group. In the adolescent group, we further split the training set by 80% training for a total of 20 subject IDs and approximately 20% validation set of 6 subject IDs. For young adults, we split the original set of 19 subjects into 80% training and 20% held out test data. Of the 80% training, 12 subjects were assigned the training set, and 3 subjects assigned for the validation set. Each split throughout was on unique subject IDs to prevent data leakage. The data was then processed for detrending, z-score normalization, and all the subjects were concatenated for the training as well as for validation sets.

Time series cross-validation was the method used at the group level since the same subject data is not included in both the training and test sets. We used Scikit-Learn’s TimeSeriesSplit¹ package to perform the 5-fold split of the data. This method consists of splitting a data set into a training set of size n (with the max size defined prior to calling the function) for observations that occurred prior to the time points for the validation. This method preserves the time series aspect of the data and shuffling is not performed. This prevents the model from observing future time points during training, providing a more accurate prediction on the validation set. The data was then fed into Scikit-learn’s HalvingGridSearch² package for hyperparameter tuning.

¹ https://scikit-learn.org/stable/modules/generated/sklearn.model_selection.TimeSeriesSplit.html

² https://scikit-learn.org/stable/modules/generated/sklearn.model_selection.HalvingGridSearchCV.html

Cross-Validation Results

The parameters were chosen in the cross-validation grid search for both adolescent and young adults using Scikit-Learn's `svm.SVC` API.³ Since we iterated this process more than once due to fundamental changes in the data that we needed to make in regards to detrending and normalization, and where the time to complete the cross validation averaged 6 hours per group, we limited the number of parameters tested and did not fine-tune as much as we originally expected. We did observe through these iterations that a radial basis function kernel consistently outperformed the other optional kernels. The regularization C parameter of 10.0 for both groups had overall better accuracy scores, though the spread was larger, likely due differences between subjects. The gamma parameter 'scale' was chosen for the adolescent model. The C parameter in the young adult cross-validation study beyond C = 5.0 statistically appeared the same and therefore, choosing the regularization value was more decided on trial and error testing. We found that the gamma parameter 'auto' was the best choice for our young adult model. Overall, we did not observe great parameter results for either young adult and adolescent cross validations and given more time, we would have been able to explore other cross-validation parameters and methods to produce better results. (Figure 12)

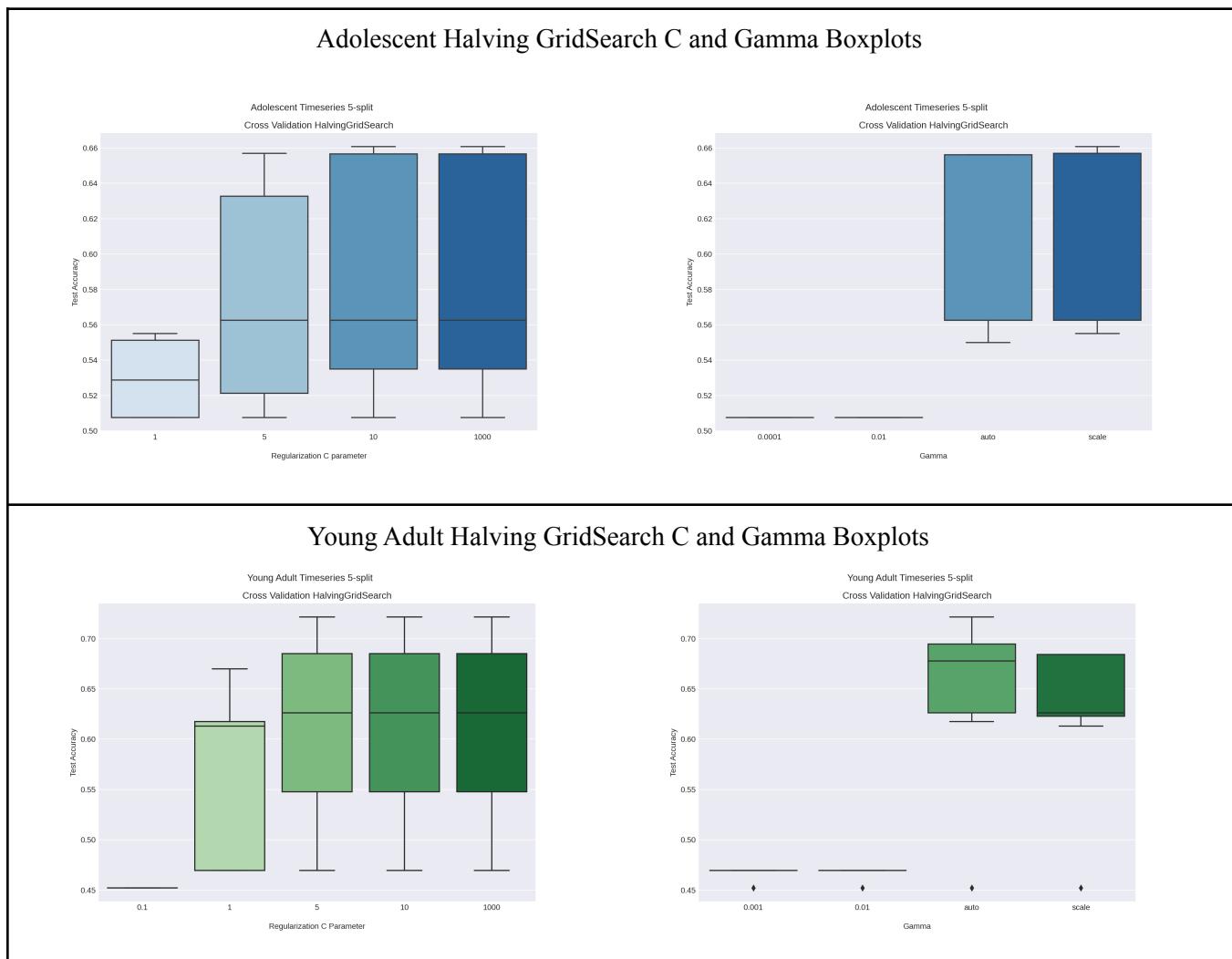


Figure 12. Adolescent C parameters have large spreads beyond 1, though the medians are higher. We use C = 10 for both Adolescent and Young Adults. Gamma 'auto' for Young Adult and 'scale' for Adolescent.

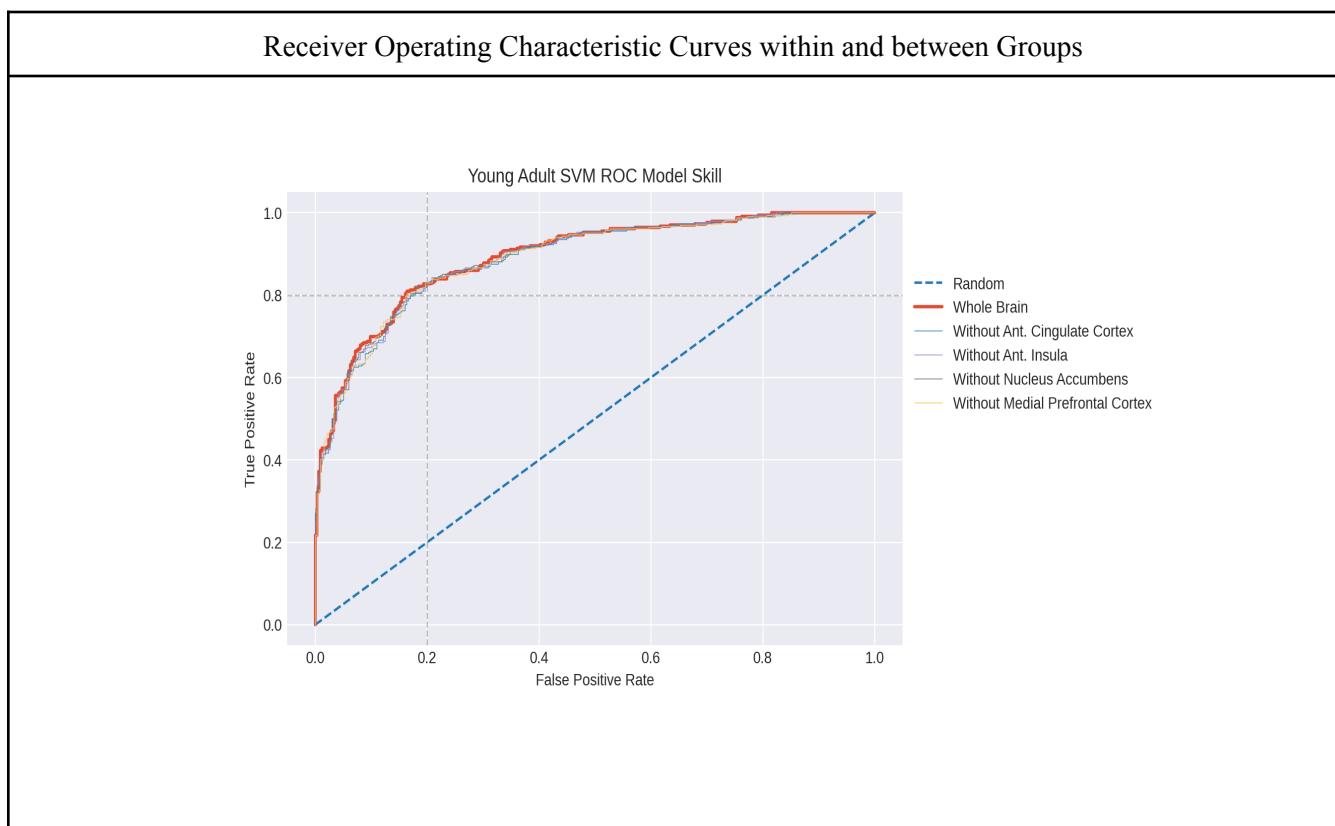
³ <https://scikit-learn.org/stable/modules/generated/sklearn.svm.SVC.html>

Adolescent and Young Adult Results

We chose to train our group level models on brain scan runs 2 and 3. These runs contain data where subjects receive feedback for up and down-regulating the reward centers while in the scanner using a brain computer interface, as well as feedback from researchers at the end of the scan. Run 2 is the first time when subjects are given feedback. Run 3 is when subjects are provided with feedback again, though this time it is thought they have already been able to learn or practice strategies since this is their second exposure. Further analyses at the group level could look at training and testing on other runs to determine if subjects were able to carry over their learning strategies using the neurofeedback from previous runs. We were able to achieve this first step in our analyses.

We looked at model ROC curves and AUC scores to compare model skill between the whole brain mask and all of the submasks for young adults and adolescents as well as comparing whole brain mask model skill between the two groups. Young adults score better than adolescents in the whole brain model with an average score of 89% and 82%, respectively. This is not surprising since young adults are thought to be better at regulation. The whole brain model underperformed slightly compared to the submasks in the adolescent group and all models performed similarly in young adults. (Figure 13)

We use a confusion matrix to compare label predictions between adolescent and young adults in the whole brain model. Both group model sensitivity of the positive class were similar with 82% of subjects up-regulating when expected. The specificity of young adults was 13% better than adolescents where 81% were down-regulating when expected versus 69% for adolescents. The young adult group performed better in sensitivity and specificity overall compared to the adolescent group. (Figure 14)



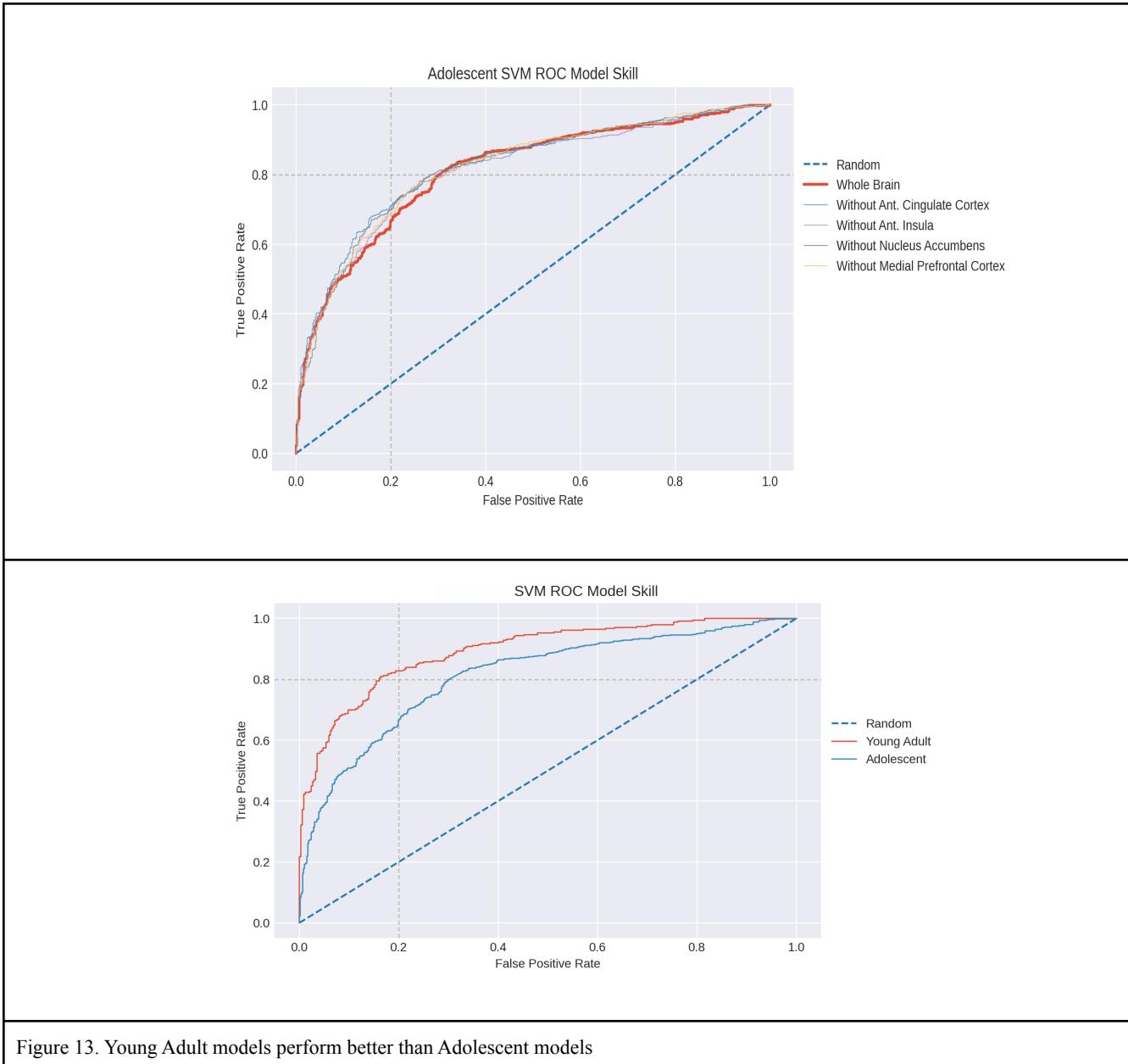


Figure 13. Young Adult models perform better than Adolescent models

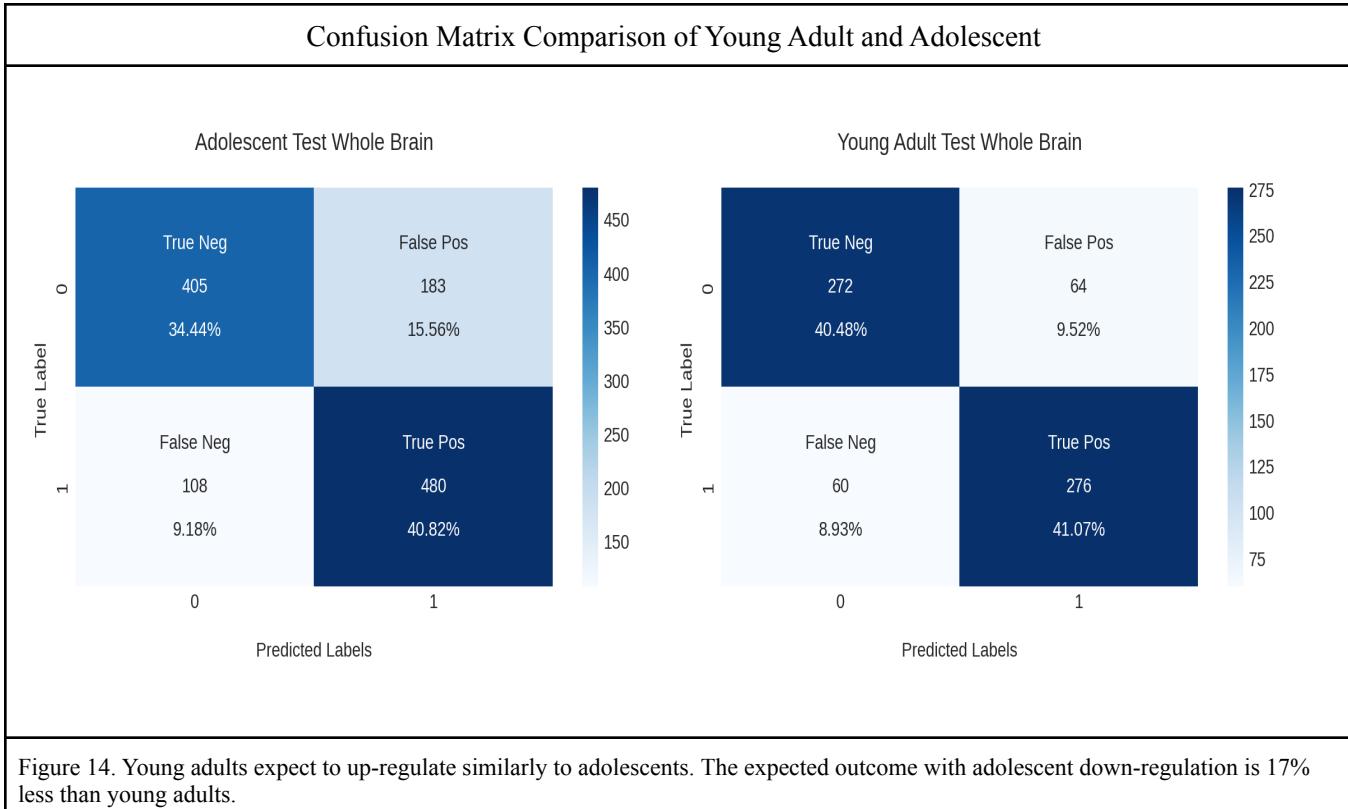
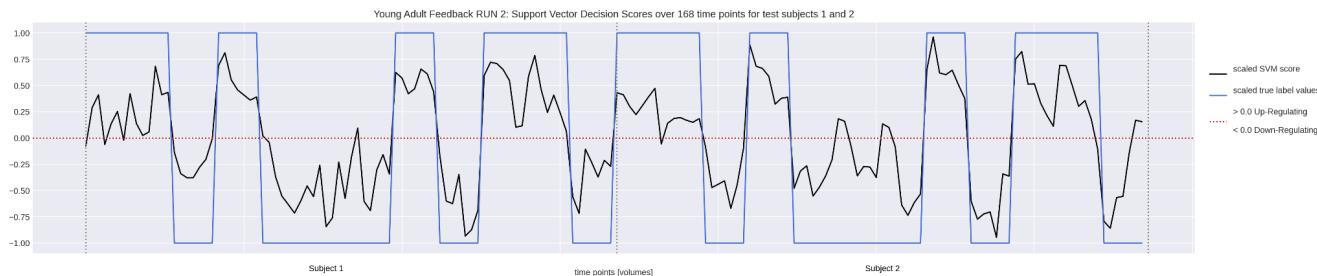


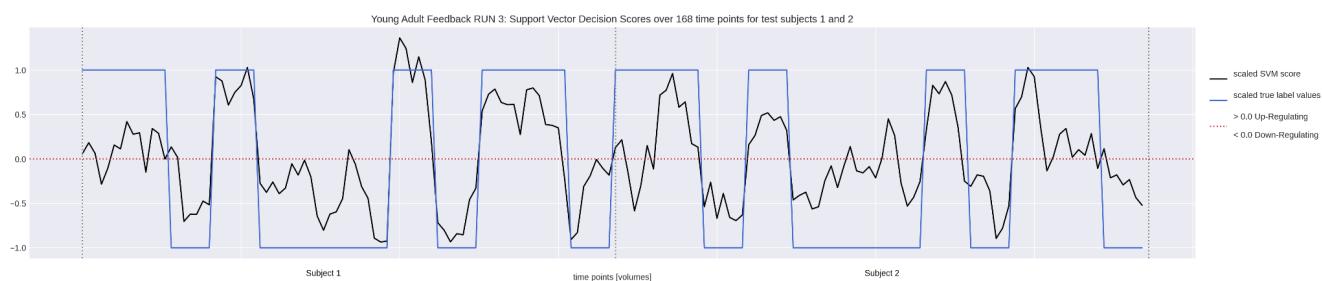
Figure 14. Young adults expect to up-regulate similarly to adolescents. The expected outcome with adolescent down-regulation is 17% less than young adults.

Since our data feature space is highly dimensional, we are not able to visualize the hyperplane and decision boundaries of the SVM classifier. In order to appreciate the decisions made by the classifier, we looked at decision function scores, which are measurements of the distances of the significant voxels from separating hyperplanes. Figure 15 shows examples of young adult and adolescent decision scores plotted against time and true labels. We chose to show an example of how two subjects might differ from one another as well as differ between themselves from run 2 to run 3 where feedback from the task might be carried over. By looking at how well the decision score line follows the curves of the true labels we can tell that young adults are overall better able to regulate than adolescents. We can also see better regulation between adolescent subjects where subject 4 regulates better than subject 5. Interestingly, we see an improvement in both of these adolescent subjects from run 2 to run 3, which may suggest some level of carryover from one feedback run to the next. (Figure 15)

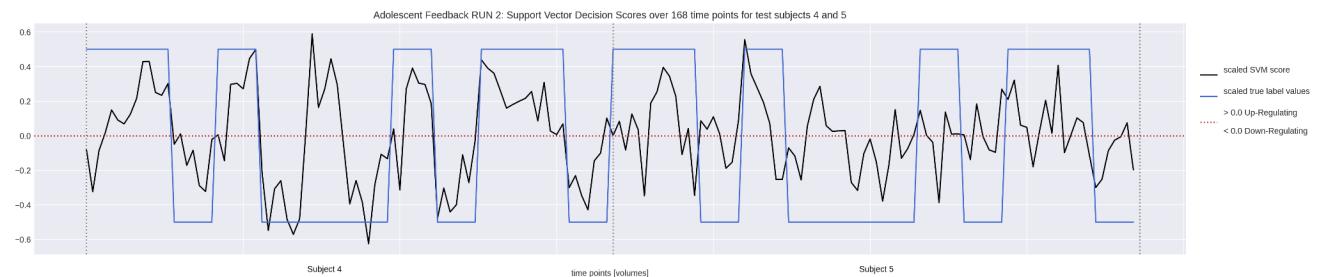
Run 2: Young Adult Support Vector Decision Scores for subjects 1 and 2



Run 3: Young Adult Support Vector Decision Scores for subjects 1 and 2



Run 2: Adolescent Support Vector Decision Scores for subjects 4 and 5



Run 3: Adolescent Support Vector Decision Scores for subjects 4 and 5

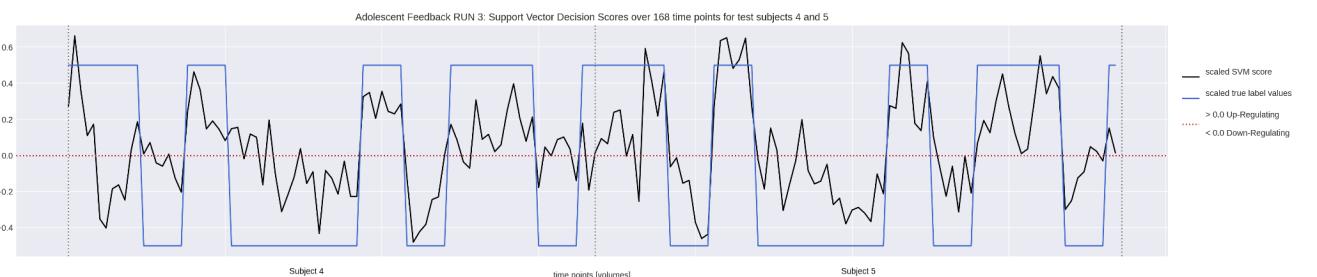
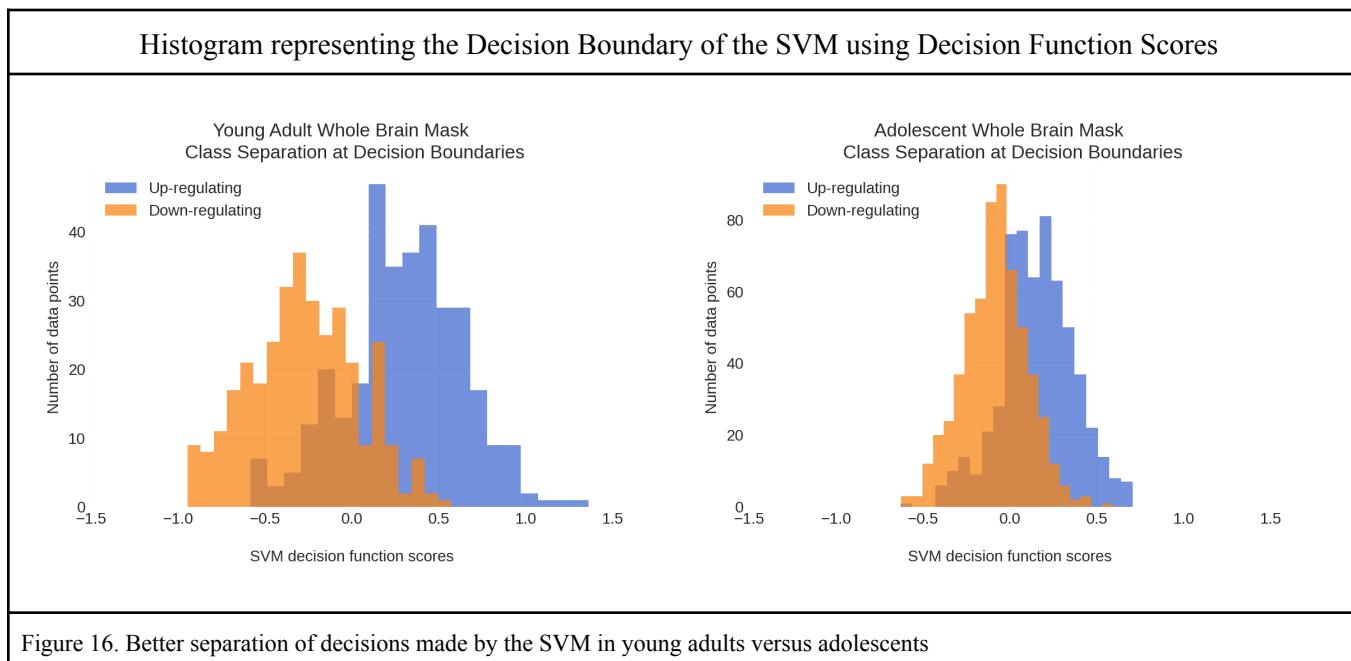


Figure 15. You can see how the adolescent subjects 4 and 5 appear to learn to regulate and carry over this knowledge from runs 2 and 3.

We can summarize all of our observations together in a histogram of the decision scores to represent the boundaries that the classifier created to make predictions. True-positive predictions sit to the right of the center of the x-axis, true-negative predictions sit to the left, while areas that overlap between -0.5 to 0.5 have been misclassified, (brown shaded region between down-regulating and up-regulating). Overall, you can see how the SVM is better at separating the decision boundaries for young adults versus adolescents. The histogram for young adults overlaps less frequently between classes, where the overlap's spread appears similar to adolescents. (Figure 16)



Analysis beyond metric comparisons at the group level looked at statistical tests of the decision function scores and beta maps between adolescents and young adults as well as within groups on the whole brain mask, submasks and regions of interest. The beta maps represent the applied support vector voxel weights to the data at the voxel indices that correspond to the support vectors (voxels that are most significant and sit near the decision boundaries). What is most notable from these tests are the differences between young adult and adolescent decision scores or beta maps for the Medial Prefrontal Cortex, the Anterior Cingulate Cortex, the whole brain mask without the Anterior Cingulate Cortex and the Anterior Insula. It does not seem clear to compare the significant scores we obtained from tests comparing the whole brain to the regions of interest since we discovered model predictions on regions of interest to be 30% lower on average than prediction scores on the whole brain. Further analysis is needed to understand how these regions of interest are different from each group and within groups. (Figure 17)

Statistical tests between groups				
Two Tailed T-Test Between Young Adult and Adolescent Decision Scores				
Region	Test Statistic	p-value	alpha	Hypothesis
Medial Prefrontal Cortex decision score	3.32825	0.000891	0.05	Reject Null H0
Anterior Insula (Right side)	2.309826	0.021008	0.05	Reject Null H0
Whole Brain without the Anterior Cingulate Cortex	-1.997647	0.045901	0.05	Reject Null H0
Two Tailed T-Test Between Beta Maps for Young Adults and Adolescents				
Region	Test Statistic	p-value	alpha	Hypothesis
Whole Brain Beta Maps	62.00712	0	0.05	Reject Null H0
Medial Prefrontal Cortex from Whole Brain Beta Map	-4.1317	3.6011e-5	0.05	Reject Null H0
Nucleus Accumbens from Whole Brain Beta Map	-1.44904	0.14733	0.05	Fail to Reject Null H0
Anterior Cingulate Cortex from Whole Brain Beta Map	16.0073	1.15223e-57	0.05	Reject Null H0
Anterior Insula (Right) from Whole Brain Beta Map	0.26111	0.79401	0.05	Fail to Reject Null H0

Figure 17. Further analyses is needed to investigate how the AI differs between groups

DEEP LEARNING METHODS AND RESULTS

Data and Preprocessing

Convolutional neural networks (CNNs) are the most widely used deep learning models in computer vision and imaging tasks. We used Pytorch, a useful library to build CNNs for 3D imaging tasks.

We built the model on runs 2 and 3 for all subjects. These runs were set aside because in these runs, patients received real-time feedback in the scanner of their ability to up and down-regulate. This creates a more consistent model premise to train, validate, and test the CNN. CNNs also need a lot of data to train and can pick up complex features in images. To increase the number of images given to the CNN, we decided not to create separate models for young adults and adolescents and instead take runs 2 and 3 for all subjects, regardless of their age group. This was a reasonable choice because CNNs have the ability to pick up features in the data, like age, that are too complex for non-deep-learning models.

The preprocessing pipeline for deep learning was slightly different than single subject and group model preprocessing regarding masking and preparing images for modeling. Researchers have found success in previous deep learning real-time fMRI imaging projects[9] using full brain masks, and we decided to format our images similarly. This involved the same normalization and detrending seen in single and group models with the addition of demasking and casting images back into their original 3D space.

We split 52 subjects into training, test, and validation sets. The training, validation, and test sets contained 36, 5, and 11 subjects respectively as a 70%, 10%, and 20% split. We split the data by subject Id to avoid data leakage. Only 36 subjects and 72 runs in the training set is not enough data to train by run. Instead, we decided to increase the training data size and train by individual image and label. This allowed us to have 6048 total labels and images in our final training dataset. With the images and labels separated, the data was model ready and saved in AWS S3.

CNN Training

The CNN parameters that we chose were based on another fMRI CNN research paper.[9] The structure of the CNN can be seen in the image below.

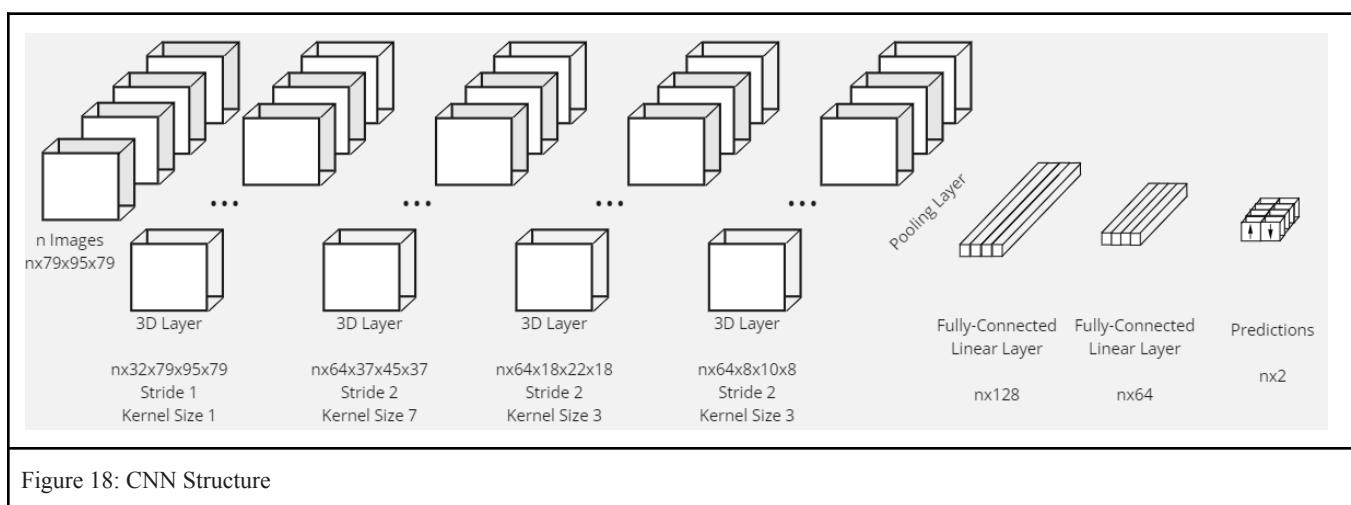
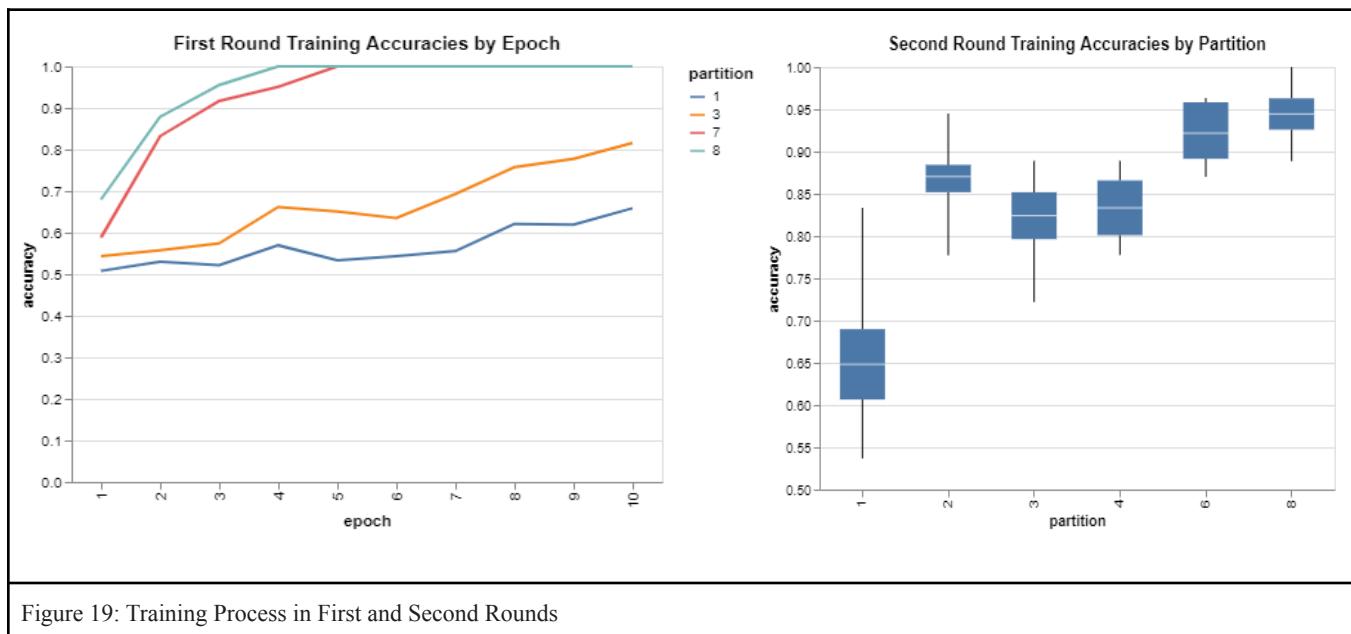


Figure 18: CNN Structure

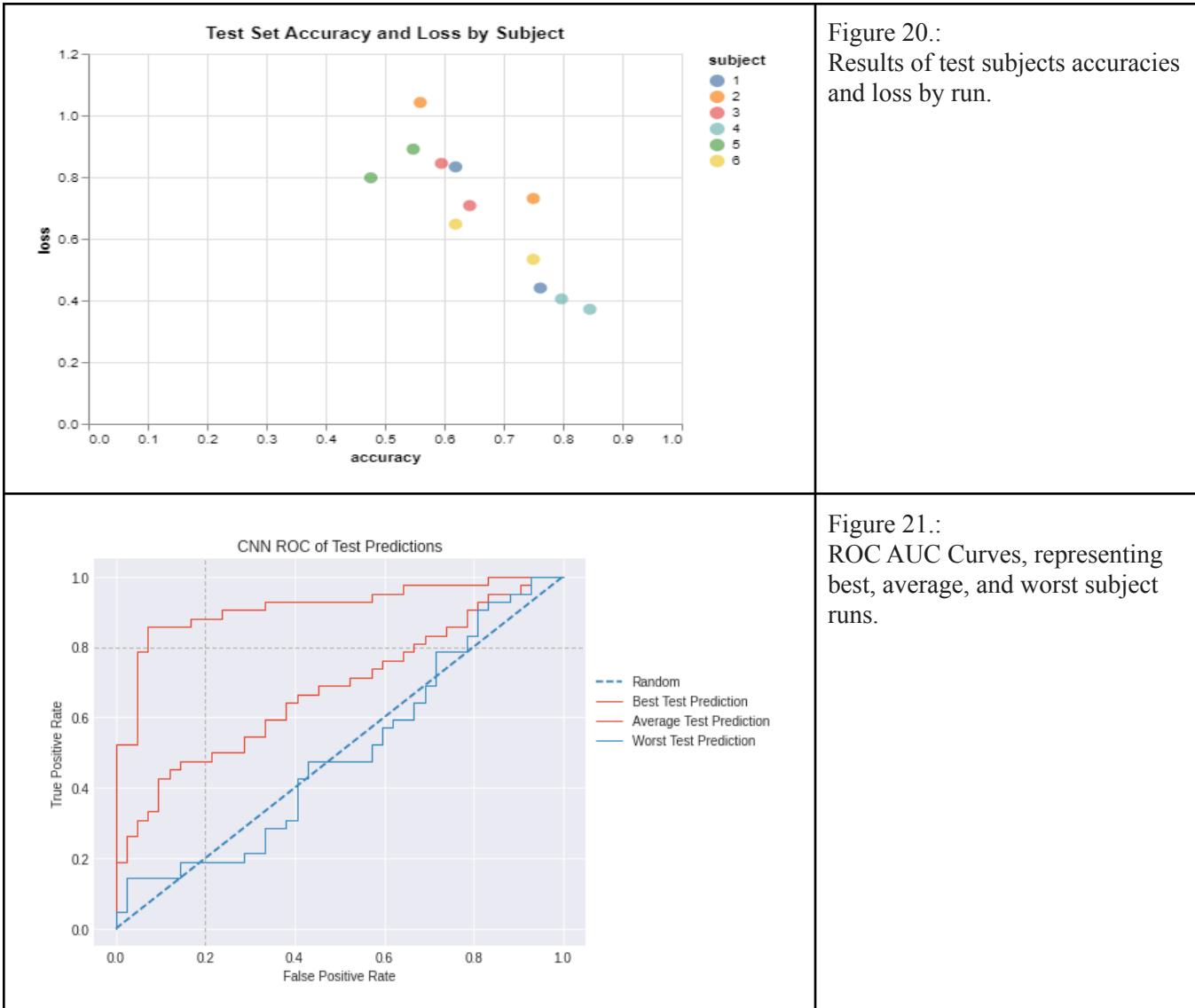
To start, we trained the data in 8 partitions of 756 images with 10 epochs for each partition. Due to the large size of our dataset, the complexity of the CNN, and the limit of 51 GB of RAM in Google Colab Pro+, we decided to train the data in 8 partitions. This involved random shuffling of individual images and labels and loading them into a Pytorch dataloader object. You can see the training results in Figure 19. As a note, some of the partition metrics were deprecated, returning only metrics on partitions 1, 3, 7, and 8. This was one of the many struggles that we had with technology. Although each partition's first epoch's accuracy was under 70%, the initial partitions of data trained much slower than the later partitions. We implemented early stopping when any batch of images returned perfect predictions in hopes to avoid model overfitting.

With the first round of 8 partitions and 10 epochs complete, it was clear that the model was overfit to the later partitions. To generalize the model, we trained the model a second time on each partition for a single epoch. The accuracy and spread of the predictions decrease from early to late partitions. This could either be because the model is able to generalize better on new data or the model remembered the later partitions better than the earlier partitions. When we ran the validation set through the model, we saw similar accuracies to partition 1. This confirmed that the CNN continued to overfit the later partitions of the data.



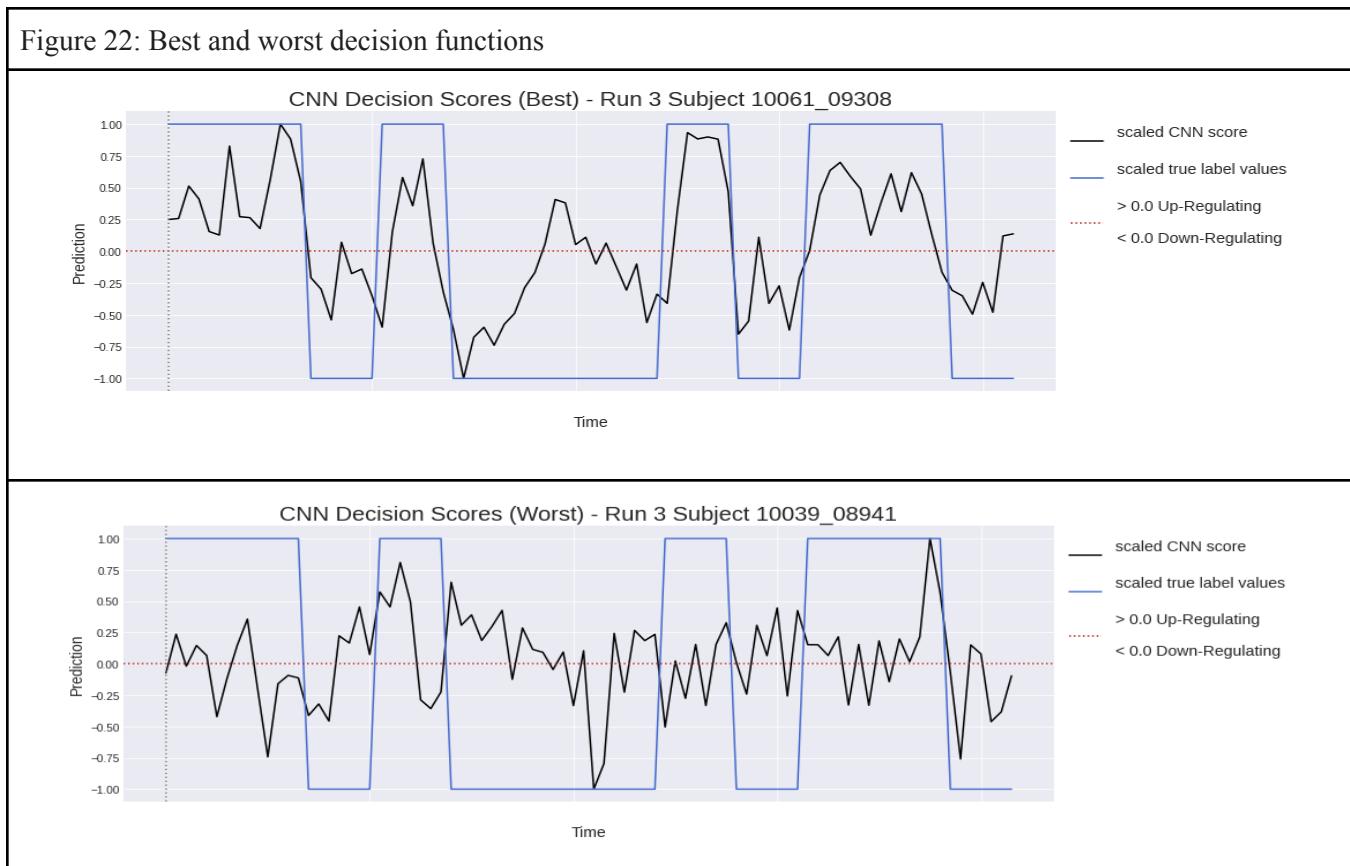
Test Results

We then took the 11 subjects in the test set and predicted each run individually through the trained CNN. Six of the subject runs 2 and 3 are shown in Figure 20, while the metrics from the other 5 were deprecated. We see here that the trained CNN had inconsistent accuracies among subjects, ranging from below random prediction accuracies to 85% accuracy. The losses were equally inconsistent. Despite these inconsistencies, each subject generally had comparable losses and accuracies between runs.



Finally, to better understand the results and the models ability to predict a subject's regulation in the scanner, we selected the best and worst subject runs by accuracy. In the ROC AUC curve in Figure 21, we see that the best prediction has an even curve, indicating that true and false positives are best distributed with around an 85% accuracy, which is very promising. In the worst test prediction, we see that the outcome follows the random line, indicating that there is little to no model indication of this subject's ability to up and down-regulate. In Figure 22, we see two decision function graphs that compare model prediction with a subject's ability to up and down regulate. In the best test prediction, the model clearly understands the subject's ability to up and down-regulate except for a small down-regulation in the middle of the run. The worst prediction does the opposite, and is predicting inconsistently with low prediction values. This is an indication that our CNN model makes volatile predictions on unseen data.

Figure 22: Best and worst decision functions



CONCLUSION

We have only begun to understand how single subject models can be trained using SVM and used in real-time to predict brain states for individuals trying to up and down-regulate the reward system. There are a lot of potential avenues to take when looking at these types of real-time classifiers. Our analyses failed to show ROIs and ROI sub-masks that may potentially play a role in the reward circuitry as being important predictors of up and down-regulation brain states. This is likely to do with the complexity of the brain and how different regions of the brain work together to control our emotional states and drive our reward system. Although we had true labels of up and down-regulation, we are not certain that all individuals will actually perform the task the way we would expect. Individuals are highly unique and the approaches for controlling the reward system could be different between individuals. We do know they are getting feedback based on percent signal change in the NAcc and future work could analyze subject prediction scores against their percent signal change neurofeedback data. Future work in single subject SVM model training would include using the program SearchLight supplied by Nilearn to create F-statistics and probability maps based on the classifier built by SVM to see if individuals that are successful at up and down-regulation have specific regions they are employing that differ between individuals.

Like in the single subject analyses, we have determined that we are able to take a machine learning approach using a SVM classifier to study at the group level. We were able to observe differences in model training between young adults and adolescents as well as differences in the ability to up and down-regulate. We also noted significant differences in statistical scores between adolescents and young adults at the group level when comparing decision scores as well as beta maps for the Medial Prefrontal Cortex, Anterior Cingulate Cortex and the Anterior Insula. These findings should be further analyzed in SearchLight as previously mentioned. Although

we focused mainly on whole brain comparisons of the two groups, we also looked at masking out regions of interest in the brain as well as looking at only the regions of interest. Further model development and analyses for the other mask models is needed to form robust conclusions between young adults and adolescents in this study to better understand regions of interest and influence in substance use disorders.

The convolutional neural network we built in this situation underperformed when compared to the support vector machines models. It had lower prediction accuracies overall and greater volatility. This comes as no great surprise because CNNs need a lot of training data. In many CNN projects, there are thousands or millions of images, and likely six thousand training images is not enough to build a consistent classifier. It is also difficult to avoid overfitting the model with such a small set of training images.

In conclusion, CNNs have potential for better accuracy in the future. As the size of training data increases, better models will likely emerge. To increase the size of data, it may be possible to use data augmentation techniques. Similarly, using transfer learning techniques on previously trained models in fMRI and then parameter tuning with our data may also increase the accuracy of a CNN and would involve a much more complex data pipeline.

STATEMENT OF WORK

Stacey Rivet Beck set up the repository and Docker container, storage bucket and group access in AWS, wrote various functions for accessing/loading and uploading data to and from AWS, created the group-level analyses pipeline from accessing the data in AWS, cross-validation, normalization exploration, model training, brain visualization and metrics analyses. She also set-up and added to the landing page, attempted to rework a pre-trained deep learning model to run with the project data, researched and built out the deep learning model architecture, participated in each general aspect of the write-up as well as full group-level write up.

Ben Merrill created the deep learning portion of the study. He created a modified pipeline to turn data into a Pytorch compatible format in AWS, dealt with scalability issues related to training the CNN and computing hardware, and created a framework for deep learning metrics and interpretation. He participated in group aspects of the project including peer review, project process and workflow, and creation of many video presentation slides.

Mary Soules contributed domain knowledge of real-time NAcc task design and fMRI domain knowledge. She supplied knowledge on how to translate SVM outputs to brain visualizations and explored how to visualize these outputs in python. She did all the quality control and preprocessing of the data to get ready for analysis. Once she created the final dataset to deploy for the project, she wrote a script to flatten the 4-D images to a 2-D matrix for each subject run and saved these data in a file that could be read from python and uploaded all data to AWS. She created all masks, sub-masks, and ROIs and uploaded those to AWS. She created the single-subject pipelines for accessing data in AWS, cross-validation, normalization exploration, model training, brain visualizations, and metric analysis. She also participated in the general aspects of the write-up as well as the full single subject SVM write up.

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Dr. Meghan Martz, Ph.D., Principal Investigator of the Neuromodulation Study

Dr. Katherine McCurry, Ph.D., Post Doctoral Fellow

Dr. Scott Peltier, Ph.D., University of Michigan fMRI Director

Dr. Qiaozhu Mei, Ph.D., University of Michigan Founding Director of MADS

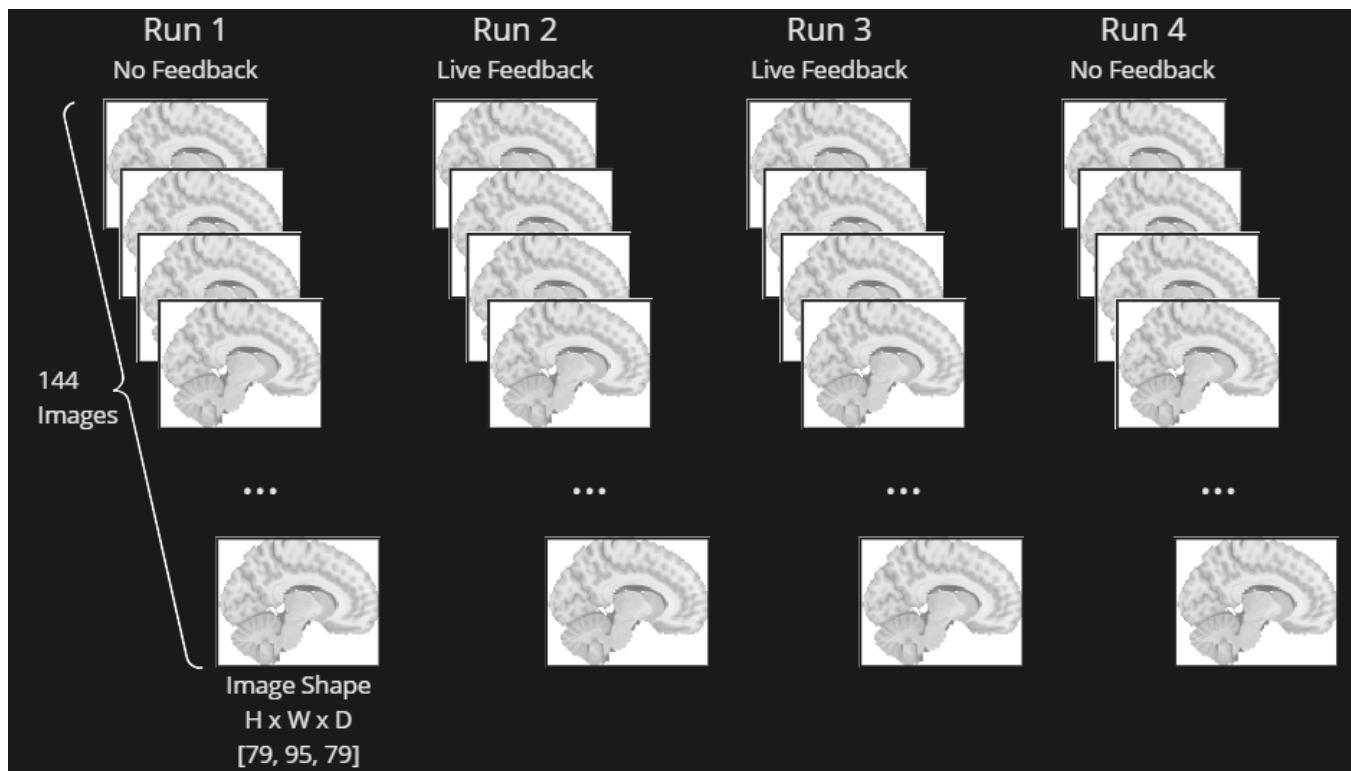
Michelle LeBlanc, Data Scientist at Blue Cross Blue Shield of Illinois, Intermittent Lecturer at University of Michigan

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APPENDIX I

Data representation of one single subject prior to masking and filtering of time points



APPENDIX II

