Interpretable and Lightweight Machine Learning Approach for Autism Classification Using Biomarkers Derived from **Multi-trial Resting EEG**

BOSTON

Michelle Su^{1,2}, Deeksha M. Shama^{2,3}, Archana Venkataraman^{2,3}

Lincoln-Sudbury Regional High School, 390 Lincoln Rd, Sudbury, MA¹; Department of Electrical and Computer Engineering, Boston University, 8 St. Mary's St, Boston, MA²; Department of Electrical and Computer Engineering, Johns Hopkins University, 3400 North Charles Street, Baltimore, MD³

INTRODUCTION

- Autism Spectrum Disorder (ASD): heterogenous, hard to identify biomarkers for diagnosis
- Electroencephalography (EEG): valuable tool for biomarker identification—non-invasive, high temporal resolution, affordable

1) DATA¹ Age (years) Male/Female Size Group Autism (ASD) 55/39 12.5 ± 2.9 94 Control (CON) 96 12.9 ± 2.8 49/47

3) FEATURE EXTRACTION

Electrodes

frequency band

Within-band features

 $128 \times 5 \times 4 = 2560$ features

Frequency bands

Get Power Spectral Density (PSD) at each electrode and frequency

2) PREPROCESSING

- Low pass (<100 Hz) and Notch (60Hz) filters</p>
- Wavelet threshold and bad segment rejection
- ICA and bad channel rejection¹

4) TRAINING AND TESTING

Four classifiers trained on the two



Subset of EEG signals (Figure from Biopac)

EEG head map (Template from Neuhaus *et. al.*¹)

Coccipital

GOALS

- Predict ASD from EEG using machine learning tools
- Explore distributional features across EEG trials
- Identify important features for ASD diagnosis

band using Welch's estimate {hamming, 512 point}

Two Feature Aggregation Types

Mean-EEG		Across-trial
PSD of Mean EEG (conventional)		PSD of EEG for each
Mean, std. deviation, skew, & kurtosis of PSD in each		Mean, std. deviation, & kurtosis of PSD in e

skew, & kurtosis of PSD in each frequency band

trial

METHODS

Mean and std. deviation of band features <u>across trials</u>

2 x 2560 = 5210 features

standardized feature types separately

- 1. Logistic regression (LR)
- 2. Random forest without bootstrap (RF)
- 3. Kernel-support vector machine (SVM)
- 4. An artificial neural network (ANN)
- Stratified 5-fold cross-validation
 - Repeated 5x
- Evaluation: accuracy, F1 score, precision, recall, specificity, AUROC, AUPRC metrics
- Feature importance: determined by magnitude of mean feature weights in cross-validation across-trial LR models

RESULTS

REPEATED 5-FOLD CROSS-VALIDATION MODEL PERFORMANCE

Model	Features	Accuracy	F1 score	Precision	Recall	Specificity	AUROC	AUPRC
LR	Across-trial	0.678±0.083	0.678±0.083	0.674±0.101	0.693±0.108	0.664±0.123	0.748±0.073	0.751±0.086

REPEATED 5-FOLD CROSS-VALIDATION ACCURACY



	Mean-EEG	0.608 ± 0.060	0.597±0.088	0.601±0.082	0.604±0.117	0.61±0.085	0.648 ± 0.070	0.678±0.084
RF	Across-trial	0.660±0.085	0.646±0.102	0.665±0.112	0.649±0.147	0.681±0.121	0.742±0.082	0.744±0.089
	Mean-EEG	0.614±0.068	0.589±0.076	0.626±0.095	0.571±0.109	0.662±0.115	0.672±0.092	0.693±0.088
SVM	Across-trial	0.659±0.079	0.656±0.089	0.659±0.101	0.679±0.146	0.654±0.106	0.734±0.087	0.743±0.093
	Mean-EEG	0.577±0.064	0.532±0.087	0.608±0.136	0.509±0.161	0.669±0.146	0.608±0.146	0.619±0.102
ANN	Across-trial	0.654±0.088	0.682±0.099	0.650±0.108	0.751±0.156	0.590±0.160	0.721±0.095	0.709±0.084
	Mean-EEG	0.622±0.079	0.616±0.112	0.600±0.112	0.661±0.160	0.564±0.158	0.651±0.108	0.683±0.098

Table 1: Metrics from all experiments. Significant differences (p<0.05) from two-sample t-tests between the two feature types are highlighted in green. (No significant differences between model choices found.)

Fig 1: Accuracy of all models using the two feature types. Across-trial feature models show higher accuracy.



DISTRIBUTION OF TOP 5% MOST IMPORTANT FEATURES

SIGNIFICANTLY DIFFERENT BRAIN REGIONS BETWEEN GROUPS



P-value = 6.38e-14P-value = 5.94e-06

Fig 2: Frequency band distribution (left) and across-trial feature distribution (right) of the top 5% most important features. P-values from chi-square goodness-of-fit tests with the null hypothesis of a uniform distribution.

-log10(P-value) p>0.05 p<0.05 2 3 4

Fig 3: Topological head plots of significant regions based on two-sample t-tests for the difference between ASD and CON in the mean of summed across-trial features adjusted by LR model weights.

DISCUSSION AND CONCLUSION

Features recording variability across trials are more important

- Trial-to-trial variability differs between ASD and CON groups²
- The choice of machine learning model is not significant
- Beta frequency: highest diagnostic value in all brain regions
- Occipital and frontal temporal regions: higher diagnostic value across multiple frequency bands

Future work:

- Incorporate features from task-EEG data
- Use deep learning for feature extraction

ACKNOWLEDGEMENTS

We thank the Autism Center For Excellence at the University of Virginia and Seattle Children's Research Institute for acquiring the data. Thank you to Prof. Archana Venkataraman and Deeksha M. Shama for providing unwavering support and guidance. Thank you to BU RISE for this incredible opportunity.

REFERENCES

(1) Neuhaus, E.; Lowry, S. J.; Santhosh, M.; Kresse, A.; Edwards, L. A.; Keller, J.; Libsack, E. J.; Kang, V. Y.; Naples, A.; Jack, A.; Jeste, S.; McPartland, J. C.; Aylward, E.; Bernier, R.; Bookheimer, S.; Dapretto, M.; Van Horn, J. D.; Pelphrey, K.; Webb, S. J. Resting State EEG in Youth with ASD: Age, Sex, and Relation to Phenotype. Journal of Neurodevelopmental Disorders 2021, 13 (1). https://doi.org/10.1186/s11689-021-09390-1.

(2) Milne, E. Increased Intra-Participant Variability in Children with Autistic Spectrum Disorders: Evidence from Single-Trial Analysis of Evoked EEG. Frontiers in Psychology 2011, 2. https://doi.org/10.3389/fpsyg.2011.00051.