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Review on Classification of Amyotrophic Lateral Sclerosis Using Ensemble Classifiers †

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Abstract: Amyotrophic Lateral Sclerosis is a neurodegenerative disorder that has a complex differential diagnosis that has not been elucidated due to the variety of clinical manifestations or specific biological markers. This review aims at studying the application of ensemble machine learning classifiers for enhancing the classification of ALS based on several models like Random Forest, Extra-Trees, XG Boost, Light GBM, Cat Boost, AdaBoost, Voting and stacking classifiers. Ensemble models offer enhanced performance in diagnosing ALS by utilizing diverse classification techniques with appropriate feature selection method. This study finds that while the Voting classifier produces comparatively inferior results, the Extra-Trees and Cat Boost models perform better with adequate precision and recall and specificity. These studies on improvements in ensembled learning methods have the potential to greatly improve disease diagnosis and early identification, which will support individualized care plans for patient with ALS.

Keywords: Amyotrophic Lateral Sclerosis (ALS); machine learning; classification; ensemble methods

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1. Introduction

Neurological disorders such as sporadic ALS and familial ALS are two main categories of amyotrophic lateral sclerosis. Sporadic ALS is more frequent, and it occurs in roughly 90–95% of patients, meaning they can't trace the disease back to a genetic source in their families. Such changes in vocal characteristics manifested themselves surprised patients at first of the presence of diseases on average at the age of 55–65 years. It develops by attacking the motor neurons that leads to muscle wastage and weakness in the muscles of speech, swallowing and muscle, and paralysis. One of this gene can develop the symptom of ALS and it can be caused by mutations. Familial ALS is responsible for 5–10% of cases and the pattern of inheritance is autosomal dominant. Familial ALS generally progresses more rapidly than the sporadic type, and is usually seen in a younger population.

Clinical examination together with elimination of other possible diseases forms part of the diagnostic procedure in ALS. For assessing the muscle and nerve involvement and the characteristics of denervation and reinnervation that are particular to ALS electroneuromyography and nerve conduction studies are obligatory. For instance, blood and spinal fluid test will help eliminate other cause of neurological issues while magnetic resonance imaging curtails structural issues of the brain and spine. Genetic testing is essential when diagnosing Familial ALS since, in most of the cases, the patient will have previously seen other family members suffering from the same disease. There are not many biomarkers detectable in ALS patients therefore early detection of the diseases is challenging given the symptoms of the disease mentioned above. By examining patient histories, genetic

profiles, and EMG results to find patterns for an early diagnosis, machine learning applied to clinical data has improved the detection and management of ALS in recent years. Additionally, wearable technology, voice, and movement data can be used by ML models to follow the evolution of diseases, enabling individualized care. This methodology improves early identification, forecasts results, and could provide fresh perspectives on ALS and potential therapeutic options.

2. Literature Review

ALS is a type of neurological disorder that usually leads to muscle atrophy, muscle weakness, and eventual paralysis. Most other conditions may pose challenges to the physician attempting to make a diagnosis at early stages when muscular strength is slightly affected. Earlier, patients were diagnosed based on clinical examinations, electrodiagnostic techniques, and imaging studies using magnetic resonance. Studies are being done that implement deep learning and machine learning in relation to diagnosis and prognosis of patients with amyotrophic lateral sclerosis. We developed a Deep Ensemble Forward Neural Network aimed at measuring cognitive performance of patients suffering from ALS. The network simulated has a target accuracy rate of 95 percent. There is one neural network that demonstrates the potential power of machine learning to enhance diagnostic capability by integrating imaging and clinical parameters. This network consists of multiple types. Erdaş et al. (2021) focused on improving the diagnosis of neurodegenerative diseases by 3-D CNN and Convolutional LSTM models. These include Huntington's disease, Parkinson's disease and Amyotrophic lateral sclerosis.

These models demonstrate some approaches that deep learning and the machine learning employed may operate on different sorts of motor function deficits. In the course of their research, Ladanza et al. employed gait test data for their model training. Leão et al. (2021) describe a vision transformer model as an efficient means in the diagnosis proven in the analysis of patients with amyotrophic lateral sclerosis. This model analyses images of MRI and carries out the extraction of spatial as well as frequency information. These snapshots illustrate the likely integration of various patient data into deep learning approaches that could assist in the more challenging aspect of developing and implementing diagnostic and treatment options for ALS. Longitudinal studies are necessary for the assessment of ALS regarding the further patients' potentialities. In their paper, Migliorelli et al. (2023) developed a CNN named DDK-AID for assessment of ALS language functions. This CNN was created based on studies of neurological audio recordings. The recent developments in technology have made it easy to keep track of a disease in real time with more focus on early detection of language-related problems. Studies have been done on sEMG as a method among the accessible non-invasive approaches to diagnostics. Chang et al. (2014) devised three diagnostic criteria designed to differentiate between healthy individuals and patients with ALS including the clustering index, the kurtosis of the EMG amplitude histogram, and the kurtosis of the EMG crossing rate expansion. At the same time, Behler et al. (2023) studied whether it is plausible to obtain the biological markers of ALS using Machine Learning and diffusion tensor imaging (DTI). Without going into too much detail, it was found that with the help of machine learning algorithms, abnormalities in large DTI datasets caused by degenerative processes of white matter motor neurons in ALS could be detected. If this is how, it could help in the development of concepts in new diagnostic as well as predictive interventions for ALS. Ahangaran et al. (2022) utilized causal graph analysis to examine the connection between genetics and the progression of ALS. There is also a huge potential in this domain for applying machine learning and deep learning techniques. The model examined three levels and rates of progression in those affected by ALS it was slow, moderate, and rapid progression. The AALSFRS-R score was fed into the program. In the end, comparing their approaches to more traditional methods, such as linear regression or support vector machine regression, they came to the conclusion that the causal graph approach brought the best results concerning the development of the disease. To some extent, and more recently, the genetic components are being

integrated into the medical diagnosis. In research involving 99 caregivers and individuals, with sclerosis (ALS) it was discovered that the burden on caregivers was greatly influenced by their emotional well-being and overall quality of life highlighting the importance of utilizing technology to identify the need for tailored therapy solutions for caregivers facing challenges in caregiving situations, for ALS patients. Burgh et al. (2017) study examined how clinical information and MRI scans could help predict the life expectancy of patients, with ALS using learning techniques. The combination of data and MRI images advanced the development of deep learning algorithms. The results demonstrated an 84. Percent improvement in accuracy when utilizing this approach suggesting that MRI data could play a role in predicting the survival outlook for ALS patients. Bean et al. (2020), on the other hand utilized machine learning to discover genes associated with ALS.

This was achieved by an independent study. An ontology-driven system for mining gene annotations and protein-protein interaction data discovered ALS and other significant biological processes. This study is going to be about how machine learning can be applied to help discover new genetic targets for ALS research and treatment. Despite the great advance, ALS pathogenesis and diagnosis are still vague. The unpredictable nature of ALS throws a barrier towards the development of effective diagnostic and prognostic tools. Onset of ALS cannot be predicted due to the inherent unpredictability of the disease in such a manner that diagnostic tests would work at the time individuals develop symptoms. A large dataset, spanning genomic, biomarker, and clinical information is needed for a more detailed mapping of the genetic basis of ALS.

Forecasting of the disease might be facilitated by an ALS-Net, the deep learning architecture designed by Yin et al. (2019). While increasing detection rates could be an objective, benefits of such strategies have not been ruled out. Deep learning and machine learning studies have expanded our understanding of ALS significantly and diagnosis mechanisms related to it. Thus, diagnostics have received a big improvement due to the integration of the imaging, genetic, and clinical data. In addition, recent strategies for offering treatment for ALS are emerging based on machine learning-based computational algorithms used to decide on the stress factors of caregivers and survival prospects. The intrinsic diversity of ALS and the need for bigger, more comprehensive data sets are two ongoing problems. More research is needed to develop expandable and non-invasive monitoring technologies, improve genetic testing procedures, and create personalized diagnostic models. If these barriers can be removed, the quality of life for those suffering from ALS and their caregivers might improve dramatically. It is equally important to promote early diagnosis and individualized treatment. Tabe 1 offers a brief review on existing works for ALS classification.

Table 1. Reviews on existing works for ALS classification.

Author	Feature Extraction Method	Classification Models	Result	Observations
Alqahtani et al., 2024	Neuroimaging, Clinical data	Deep Resemble Forward Neural Network	95% accuracy in classifying cognitive function	Highly effective in combining neuroimaging and clinical data
ErdaÅŸ et al., 2021	Gait analysis data	Convolutional LSTM, 3D CNN	Detects neurodegenerative diseases using gait analysis	Demonstrates potential for ML in multiple neurodegenerative diseases
Iadanza et al., 2022	Gene expression data	Pathway Analysis using ML	Identified immune response and inflammation as key biological processes in ALS progression	Offers valuable insights into ALS mechanisms and potential therapeutic targets

LeÃ£o et al., 2021	Time-dependent and independent patient data	Dynamic Bayesian Networks	Predicts ALS progression over time	Addresses ALS progression, aiding in prognosis
Kew et al., 2024	Brain signals from BCIs	ML applied to Brain-Computer Interfaces (BCI)	ML applied to BCIs can decode complex brain signals for ALS patients	Synergy between ML and BCIs holds promise for ALS patient independence
Alzahrani et al., 2024	Dual neural network approach (UNET and GAN)	UNET and GAN architectures	Early ALS detection with promising results (details unspecified)	Uses innovative dual architecture for early ALS detection
Ã‡elik et al., 2024	Chromatin accessibility outliers	EpiOut Algorithm	Identifies genetic factors contributing to ALS through chromatin accessibility outliers	Significant in understanding genetic contributions to ALS
Alemasov et al., 2022	Hydrogen bond stability in SOD1 mutants	Accelerated Molecular Dynamics	Hydrogen bond stability correlated with patient survival times	Protein stability plays a role in ALS progression
Ahangaran et al., 2022	Causal graph analysis, ALSFRS-R score	Causal Graph Method, SVM Regression	Causal graph method showed greater precision in predicting ALS progression	Causal graph analysis improves precision in progression prediction
Migliorelli et al., 2023	Audio recordings (Speech performance)	CNN (DDK-AID)	Remote monitoring and speech assessment tool for ALS progression	Enables early detection and monitoring of speech performance
Burgh et al., 2017	MRI and clinical data	Deep Learning	84.4% accuracy in predicting ALS survival	Combining clinical and MRI data improves survival prediction
Pancotti et al., 2022	Pooled ALS clinical trial data	Deep Learning	Accurate prediction of ALS progression using clinical data	Deep learning shows promise in predicting ALS progression
Zhang et al., 2014	Surface Electromyography (sEMG) data	Linear Discriminant Analysis	90% sensitivity, 100% specificity in differentiating ALS patients using sEMG	Non-invasive sEMG method useful in ALS diagnosis
Gordon & Lerner, 2019	Variety of ALS-related data sources	Machine Learning	ML provides insights into ALS heterogeneity	ML models need further validation for heterogeneous ALS
Yin et al., 2019	Genome data (genotype prediction)	Deep Learning (ALS-Net)	ALS-Net outperforms other methods in recall	ALS-Net offers strong recall and precision for genotype-based ALS prediction
Antoniadi et al., 2020	Caregiver data (Quality of life, psychological distress)	Random Forests	ML can predict caregiver burden and quality of life	ML tools provide actionable insights for caregivers
Behler et al., 2023	Diffusion Tensor Imaging (DTI)	Machine Learning (for biomarkers)	DTI and ML show promise in detecting ALS biomarkers	Combining ML with neuroimaging can improve ALS understanding
Bean et al., 2020	Protein-protein interaction data, gene annotations	Knowledge-Based ML	ML identified new candidate genes for ALS	ML offers a pathway for discovering new ALS-related genes

3. Methodology

Several ensemble learning models are used in this work to improve prediction robustness and accuracy for ALS classification. Developed from numerous decision trees, the Random Forest Classifier is a basic model that reduces overfitting and enhances generalization. In a comparable way, the Extra Trees Classifier expands on this idea by employing random splits, which frequently produce different trees with quicker training periods. Using a distinct methodology, the Gradient Boosting Classifier builds models one after the other, correcting the mistakes of the earlier models to produce higher accuracy. XGBoost is an optimized version of gradient boosting which is prominent for its efficiency making it suitable for large datasets due to its ability to handle missing values and implement regularization techniques.

LightGBM, another important model examined in this paper, has been optimized to be highly efficient and has the ability to process categorical information directly, which minimizes the need for preprocessing. With its unique ordered boosting technique, the CatBoost model reduces overfitting and performs exceptionally well with categorical data. While the Bagging Classifier trains models on various subsets of the data to efficiently minimize variance and prevent overfitting, AdaBoost combines many weak classifiers to boost accuracy by modifying the weights of instances based on prior misclassifications. Together with these models, this work constructs a Stacking Classifier, which makes advantage of the capabilities of each model by leveraging the information from many base classifiers to inform a higher-level meta-classifier. To improve accuracy, the Voting Classifier integrates predictions from multiple models using both hard and soft voting techniques. Last but not least, an ensemble neural network that leverages neural networks' capacity to identify intricate patterns in data combines two Multi-layer Perceptron classifiers through soft voting. In medical contexts, where precise diagnoses can have a substantial impact on patient outcomes, it is imperative that ensemble approaches such as this one together to improve the prediction performance in ALS classification.

4. Experiments and Results

In order to improve model performance for the categorization of ALS, feature selection is essential similar to any domain where classification involved features. To get ready for model training, the dataset first goes through preprocessing which involves encoding categorical features and standardizing numerical values. To systematically select and maintain the most significant features, the Random Forest Classifier is used as the base estimator in the Recursive Feature Elimination approach. By using an incremental approach in which eliminating the features with the lowest significance each time, the final four features are determined to have incredible potential of enhancing the predictive value of the models. The pair plot of these features is provided in the Figure 1 below. Due to these extracted features, the subsequent ensemble models are able to work more effectively and be computationally more explosive by eliminating dimensionality. The experiments carried out include setting up and assessing multiple kinds of ensemble classifiers, such as Random Forest, XGBoost, LightGBM, and voting and stacking techniques. Through each model, various methods of performance analysis are examined and include accuracy, recall, specificity, F1 score, and AUC-ROC for a comprehensive understanding of how well each model predicts data. This systematic approach to feature selection and model experimentation ensures that the classification framework is both robust and interpretable that ultimately facilitate more accurate predictions in ALS diagnosis. Figure 2 provides Performance of classification of ALS using different ensemble machine leaning classifiers.

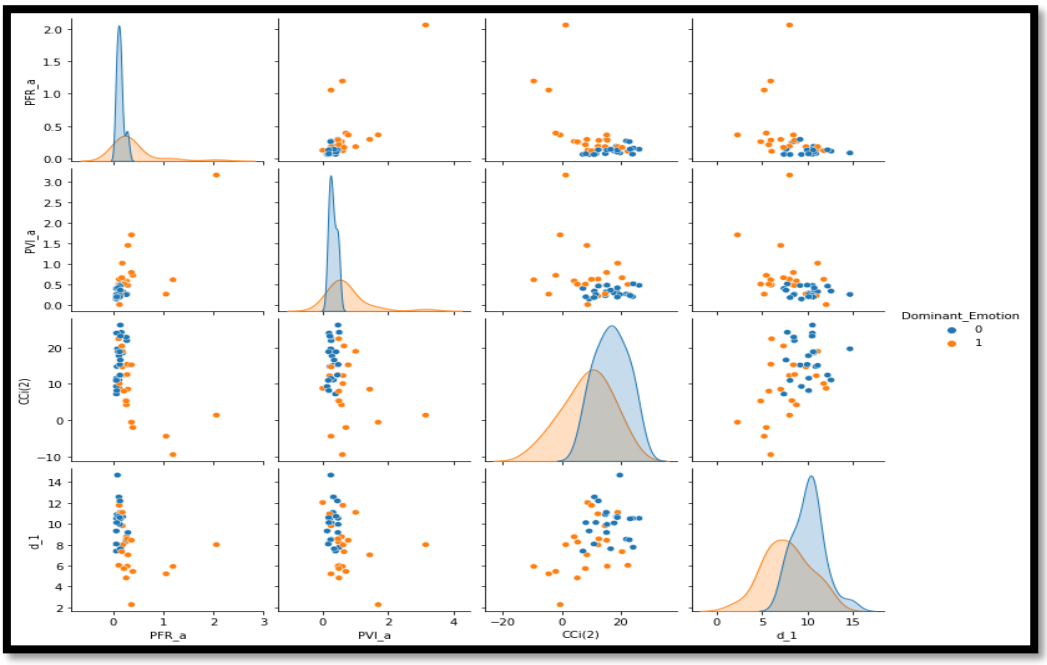

Figure 1. Pair plot for prominent features selected through RFE method.

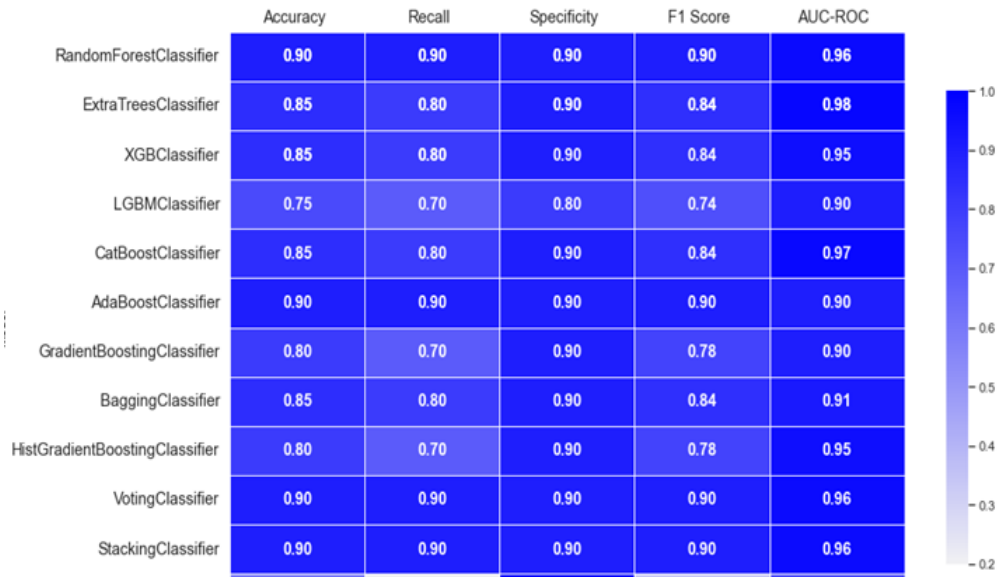

Figure 2. Performance of classification of ALS using different ensemble machine leaning classifiers.

The Random Forest Classifier and AdaBoost Classifier show strong performance across all metrics, with 0.90 for Accuracy, Recall, Specificity, and F1 Score, and AUC-ROC values above 0.95. Extra Trees, XG Boost, and Cat Boost also perform well with slight differences in Recall and Specificity. LGBM Classifier lags slightly, especially in Recall and F1 Score. Figure 3 offers accuracy comparison plot for classification of ALS using different ensemble machine leaning classifiers. The Accuracy Comparison chart shows that AdaBoost Classifier and Stacking Classifier have the highest accuracy slightly above 0.9. LGBM Classifier has the lowest accuracy, around 0.75 while the other models, including Random Forest, XG Boost, and Extra Trees, perform similarly score close to 0.85.

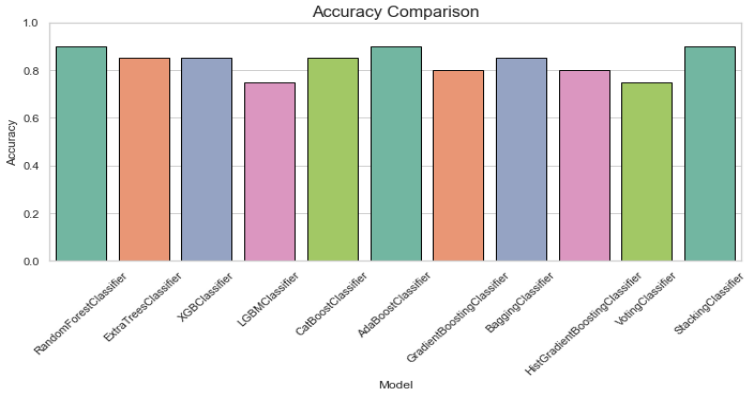

Figure 3. Accuracy comparison plot for classification of ALS using different ensemble machine leaning classifiers.

Figure 4 displays specificity comparision plot for classification of ALS using diffrent ensemble machine leaning classifiers. The Specificity Comparison chart indicates that all classifiers perform well, with most achieving scores above 0.8. The VotingClassifier stands out with a perfect specificity of 1.0, while the other models, including RandomForest and ExtraTrees, maintain strong performance with scores close to 0.9. Figure 5 provided F1 score comparision plot for ALS classification using diffrent ensemble machine leaning classifiers. The chart shows that most models achieve F1 scores above 0.8, with StackingClassifier performing the best followed by GradientBoostingClassifier and AdaBoostClassifier. The VotingClassifier has the lowest F1 score, suggesting it underperforms compared to the other ensemble methods.

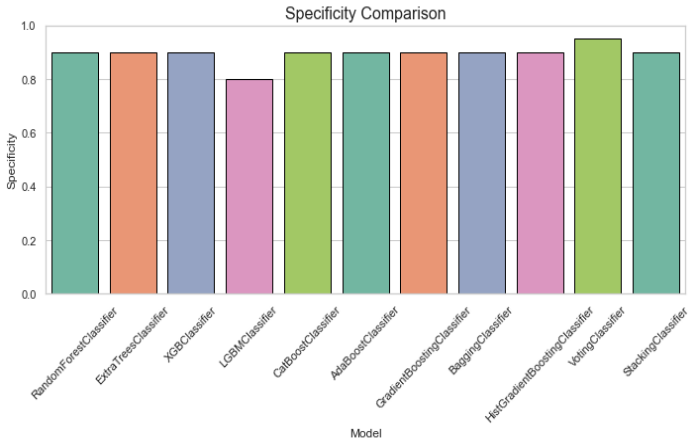

Figure 4. Specificity comparison plot for classification of ALS using different ensemble machine leaning classifiers.

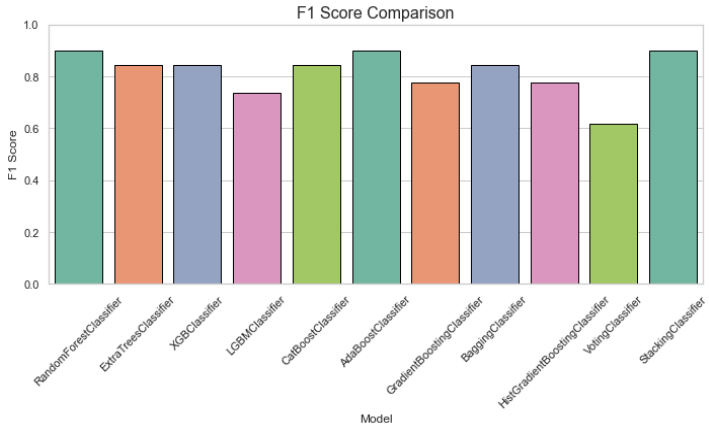

Figure 5. F1 score comparision plot for classification of using diffrent ensemble machine leaning classifiers.

Figure 6 displays AUC-ROC comparision plot for ALS classification. The AUC-ROC comparison chart shows that most models perform well with AUC-ROC values close to 0.9. StackingClassifier and RandomForestClassifier achieve the highest AUC-ROC, while VotingClassifier performs the worst, slightly below the others.

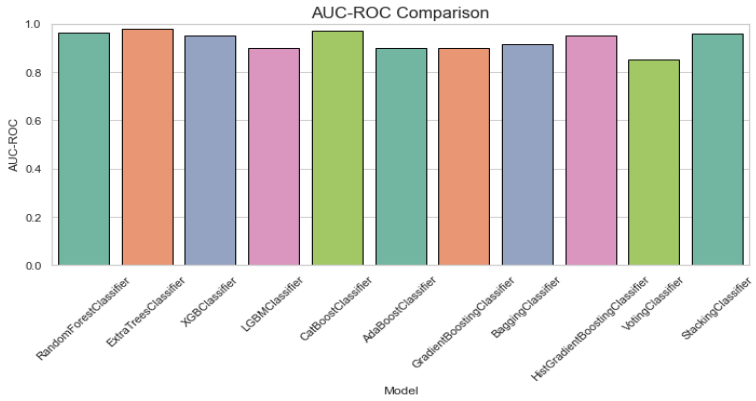

Figure 6. Specificity comparision plot for classification of ALS using diffrent ensemble machine leaning classifiers.

Figure 7 displays the precision recall curves for classification of ALS using diffrent ensemble machine leaning classifiers. The Precision-Recall curve shows that ExtraTreesClassifier and CatBoostClassifier have the highest average precision i.e., 0.98 and 0.97 respectively. This indicate that superior precision and recall performance. VotingClassifier performs the worst i.e., 0.78, suggesting weaker precision-recall trade-offs compared to other ensemble models like RandomForestClassifier and StackingClassifier with 0.96 score.

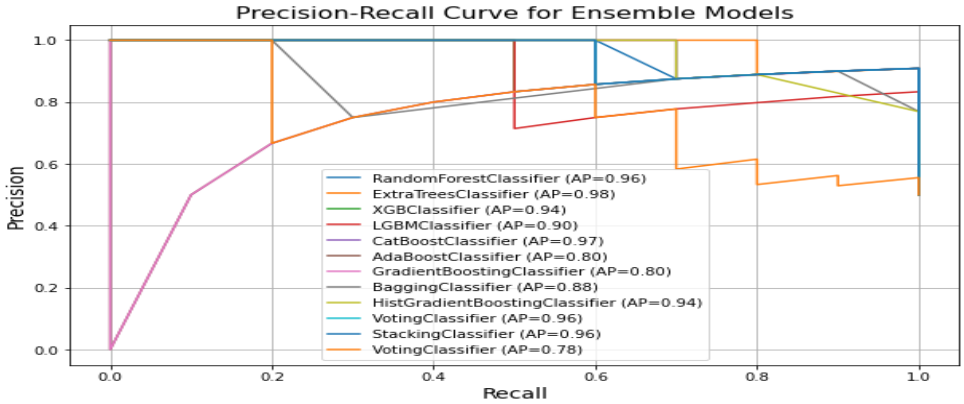

Figure 7. Specificity comparision plot for classification of ALS using diffrent ensemble machine leaning classifiers.

5. Conclusions

The predicted accuracy and resilience of ALS classification models are greatly improved by the application of ensemble machine learning classifiers. The models with the highest precision and recall are the Extra-Trees Classifier and Cat Boost Classifier, whereas the Voting Classifier performs poorly. Through the reduction of dimensionality and enhancement of computing efficiency, feature selection techniques such as RFE aid in the optimization of model performance. Despite these advancements few challenges remain unsolved such as including the need for larger datasets and more comprehensive diagnostic biomarkers. Continued refinement of machine learning models and the integration of novel data sources hold great promise for improving early diagnosis, treatment outcomes, and the overall quality of life for ALS patients.

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