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# Predicting Alzheimer's Disease at Low Cost Using Machine Learning

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Abstract— Modeling and predicting Alzheimer's disease (AD) is a significant field of study for many years. However, it is a luxury to diagnose and treat AD for poor people in many developing and underdeveloped countries. Many countries lack the proper tools. Many people do not have the capability of bearing the cost of MRI tests. In this paper, we analyzed a way of identifying AD as cheaply as possible 8 Ve investigated the difference in efficiency between the Machine learning models using Magnetic Resonance Imaging (MRI) data versus models without using MRI data. We had around a 3% difference between the best models of these tests. We used the OASIS dataset. We used patients' longitudinal lifestyle data like age, gender, education, income, MMSC score, and other features for Non-MRI models. For MRI models, we also analyzed MRI data along with the lifestyle data. We trained different Machine learning models like Random Forest Classifier, GaussianNB, LinearSVC, Logistic Regression, KNeighbors Classifier, Adaboost Classifier, and several other models. After that, we combined our best models and created a hybrid model. Our best result was 96.07% accuracy with MRI data and 93.37% accuracy without MRI data. The non-MRI result may not be as efficient as the models with the MRI dataset and not crucially inefficient. Our result can be summarized as the Non-MRI data can be used for starting the diagnosis for those poor people.

Keywords—Machine learning, Alzheimer's disease. MRI data, Non-MRI data, cheap.

#### I. INTRODUCTION

Alzheimer's disease is a progressive neurologic disorder, it 32 ses the brain to shrink and brain cells to die. Every four seconds, someone is diagnosed with Alzheimer's disease. Alzheimer's disease is one of the most common forms of dementia affecting millions of older people worldwide.

Dementia refers to diseases that are characterized by a loss of memory or other cognitive impairments. It is caused by damage to nerve cells in the brain. It becomes more severe over time, and it is not curable.

Every country has elderly with dementia. However, the number is alarmingly increasing in the least developing and developing nations. Research [1] has shown that by 2040, 71% of dementia patients will be from developing nations, and the number will double every 20 years. There is low awareness about dementia in those countries. Many people in those countries think the loss of memory is a typical aging problem. The lack of diagnostic assessment [2], cost of diagnosis deters people from diagnosis of dementia. The average cost of MRI in the USA is 1100 USD, the MMSE is 1.23 USD [3]. In Bangladesh, the price of a brain MRI is about 46 USD. In India, it is about 136 USD. So, Brain MRI is significantly more expensive than MMSE, which many families cannot bear.

There are several factors in dementia. Age and gender are also essential factors. Females are more like to be affected by AD than Male [4]. Low education level [5] is also a crucial factor. Low social class, income, depression, head trauma, epilepsy, diabetes, and stroke are relevant factors [6]. All these factors can be used with Mini-Mental State Exam (MMSE) to early diagnosis of AD. The lower cost of diagnosis will open doors for many poor people and people living in the rural area for AD diagnosis.

Early confirmation of Alzheimer's disease may be helpful to start proper treatment. If the disease is predicted earlier, the progression of the symptoms of the disease can be slow down and save lives.

Several Machine learning models were tested for MRI data and nor-MRI data to measure the gap between those two approaches. Analyzing MRI data gives more accurate results but is expassive. Analyzing non-MRI data is less efficient but is cheap. In this research, we utilize the effectiveness of the popular Machine learning algorithms to prove that analyzing non-47 I data is a viable option for starting the AD treatment. Our research can be summarized as,

- We investigated the existing Machine learning algorithms to produce the best result possible for both of our tests, with and without MRI data. Our research is the first research to perform this benchmark.
- We experimented with several Machine learning algorithms and found AdaBoost gives the best result in MRI data analysis, and our Hybrid model gives the best cauself for non-MRI data analysis.

The rest of the paper is organized as follows. In section II, we conducted a literature review. In section III, we described the detailed description of our study, finally, in section IV, the conclusion of our research.

#### II. LITERATURE REVIEW

Predicting Alzheimer's disease is a popular research field for researchers. A significant amount of work has been done in this field. MRI and PET data analysis is the dominant subfield in predicting Alzheimer's disease. Biomarkers analysis and longitudinal data analysis for classification and progression are also much popular. A few of them are,

7 Gopi Battineni et al. [7] also used MRI data to develop different ML models to predict dementia in the elderly. Subjects going through brain changes, like leukoaraiosis, mild atrophy, and regular dementia cass of AD, were included in this study. They trained four Machine Learning models. The combination of all four models with selective features enhanced the Accuracy of dementia prediction. The combination of all four models with selective features increased the Accuracy to 98%.

Siqi Liu et al. [8] designed a deep learning architecture of stacked auto-encoders and a SoftMax output layer and applied them on MRI images from ADNI. They could do it with less minimal domain prio 42 nowledge and labelled training samples. They produced overall Accuracy (87.76%) in the binary classification of AD.

Yanyan Lin et al. [9] studied to develop a longitudinal structural magnetic resonance imaging-based prediction system for MCI progression. They ellected longitudinal data from 164 MCI patients. They used a discriminative dictionary learning framework to distinguish MCI patches, avoiding the segmentation of regions of interest. They achieved an accuracy of 97%.

Courtney Cocherane et al. [10] analyzed different preprocessing methods, machine learning models, and feature selection techniques. Instead of using MRI data, they used longitudinal lifestyle interventions. They achieved more than 90% accuracy and recall in predicting Alzheimer's disease. They produced a "lean" diagnostic protocol that can predict AD development in someone with only three tests and four clinical visits with 87% accuracy and 70% recall.

XIN HONG et al. [11] focused on <sup>28</sup> tiffy time rela <sup>28</sup> biomarkers associated with disease status. They found that the Cortical Thickness Average (TA) is significant in predicting A heimer's disease progression. They propose a predicting model bas <sup>28</sup> on Long short-term memory (LSTM), which might connect previous information to the present task.

Garam Lee et al. [12] developed a framework that combines cross-sectional neuroimaging biomarkers at baseline, longitudinal cerebrospinal fluid (CSF), and cognitive performance biom 4 ers obtained from ADNI. They took advantage of the longitudinal and multi-modal nature of available data for discovering nonlinear patterns associated with MCI progression. The proposed framework integrates 4 ngitudinal multi-domain data. The most significant advantage of their approach is that irregular longitudi 4 data can be used. Their model achieved 81% accuracy when incorporating longitudinal multi-domain data.

Subramoniam et al. [13] proposed the classification of Alzheimer's disease based on a deep neural network using Magnetic Resonance Images (MRI) as input for the classification task. They have proved that among the VGG architectures, the VGG-16 performed better than VGG-19. Among the residual neural network architectures, Resnet-18 was more accurate than Resnet-101.

We have reviewed many papers, but none of them focused on the differences in efficiency between MRI and non-MRI data analysis. In this paper, we did precisely that analysis to determine the efficiency difference.

#### III. METHODOLOGY

#### A. Dataset used

For this 5 dy, we used the OASIS [14] longitudinal data.

OASIS is the Open Access Series of Imaging Studies (OASIS). This project aims to make neuroimaging data sets of the brain freely available to the scientific community.

OASIS Dataset provides several independent variables including sociodemographic and clinical variables. Sociodemographic variables used in this experiment are:

- Gender: Male = 150, Female = 204.
- Age: Min = 60, Max = 98, Mean = 77.03.
- Years of Education (EDUC): Min = 6, Max =23, Mean
   14.7.
- Socioeconomic Status (SES): Lower = 88, Lower Middle
   = 103, Middle = 82, Upper Middle = 74, Upper = 7.

#### Clin 29 Variables are:

- Mini-Mental State Examination score (MMSE): The MMSE is a 30-point questionnaire used broadly in clinical and research environments to measure cognitive impairment [15]. However, MMSE's alone cannot be 25 d for the classification of a disease.
- Clinical Dementia Rating (CDR): The CDR is a 5-point scale used to characterizing and tracking a patient's dementia level. CDR is estimated based on a semi-structured interview of the subject and the caregiver (informant) and the clinician's clinical judgme 48
- Estimated total intracranial volume (eTIV): The eTIV variable estimates total intracranial brain volume.
- 7 prmalize Whole Brain Volume (nWVB): The nWVB variable measures the 5 ume of the whole brain.
- Atlas Scaling Factor (ASF): Atlas Scaling Factor is a computed scaling factor that transforms native-space brain and skull to the atlas target (i.e., the determinant of the transform matrix). [16]

39 In our test, we used Gender, Age, Years of education, Socioeconomic Status, Mini-Mental State Examination score, and Clinical Dementia Rating for the non-MRI test. In our Test with MRI data, we used all of these variables.

#### B. Data processing

First, we removed all the rows with empty data. Initially, we had 373 subjects, but after modifying our dataset, we had 354 subjects. Then we used the One-hot encoder to encoded all categ 44 all variables and performed feature engineering. Finally, we split 41 data into two parts, the train set and the test set. We used 80% of the data for training our models and 20% for testing purpose.

We conducted our experiment in two steps. In the first step, we included all the MRI data and longitudinal lifestyle data like age, gender, income, education, and other lifestyle data and neuropsychological scores like MMSC. In the second step, we excluded all the MRI-related data and did the same experiment again. Our dataset has multiple classes like demented, non-demented, and converters.

#### C. Machine learning algorithms.

After p 26 essing our data, we moved onto selecting an efficient Machine learning. Selecting the best machine learning algorithm is very important. So, we experimented 26 h several machine learning algorithms. We used both classification and ensemble 26 arner algorithms. Ensemble learners can sometimes outperform the classification algorithms by combining the predictions of multiple base estimators.

For our test with MRI data, we started with several Machine learning models. We first run all the models without hyperparameter tuning. We only selected the models that had more than 50% accuracy and recall. Some models performed very poorly. For example, KNeighbours Classifier trained with MRI data only got an accuracy score of 45.07% and recall 0.45. So, we did not invest time in tunning hyper-parameters of these models.

We got five models that performed well: Random Forest Classifier, Gaussian NB, Linear SVC, Logistic Regression, and Ada boo 6 classifier. So, we moved on to tunning these models. The supervised learning module performs a stratified ten-fold cross-validation and grid search over selected features for each model. 38r example, for the Random Forest method, we tunned n\_estimators, max\_depth, max\_features, min samples split, min samples leaf, and bootstrap for tunning the model. We created a list of different values for each of them and searched for the best parameters using grid search. After finding the best parameters the first time, for the numerical value parameters like n estimators, we changed the list of values, this time, we took values close to the previous best and ran again to get the maximum accuracy recall possible. We repeated this process until the Accuracy was not increasing anymore. Then using the parameters that maximized the Accuracy and Recall, we predicted the results for our test dataset

We repeated the same procedure for our experiment without MRI data.

After that, we used our tunned model to create a hybrid model for both tests. We used soft voting classifier for building the hybrid models. We combined Adaboost classifier, Random Forest, Linear SVC, and Logistic Regression with soft voting classifier for building this hybrid model for MRI data. For non-MRI data, We combined five models. They are Random Forest Classifier, GaussianNB, LinearSVC, Logistic Regression, and AdaBoost Classifier.

#### D. Comparison

We compared the result of the two experiments. We compared the result in different scales like Accuracy, F1 score, Recall, Precision. Accuracy is the percentage of correct predictions for the test data. The equations are of Accuracy is.

$$Accuracy = \frac{TP + TN}{11 + TN + FP + FN}$$
Here, TP stands for True Positive, TN denotes True

Here, TP stands for True Positive, TN denotes True Negative, FP denotes False Positive, and FN denotes False Negative. However, Accuracy is not alway 43 to best scale to measure a model's performance. Precision tells us how many of the predicted positive results are positive. While Recall tells us how many of the positives are captured by our model. The equation of Recall and Precision are,

$$Recall = \frac{TP}{TP + FN}$$
 
$$Precision = \frac{TP}{TP + FP}$$

F1 score is used when we need to balance between Precision and Recall. The equation for measuring the F1 score is,

$$F1\:score = 2*\frac{Precision*Recall}{Precision + Recall}$$

We analyzed the Area Under the Receiver Operating Characteristics to determine which model best distinguishes classes. The model with the highest ROC – AUC value has the best separability. F1 score also tells us which model has the optimal false positive and true positive rates.

#### E. Results

a) MRI data results: We got the best Accuracy and a recall score from AdaBoost Classifier. It had an Accuracy score of 96.07%, a Recall of 0.96, a Precision of 0.96, and an F1 score of 0.96. The 23 curacy score of Logistic Regression was 94.37%, a Recall score of 0.94, a Precision score of 0.94, and an F1 score of 0.94.

We got the best AUC score from Logistic Regress 18, 0.97, AdaBoost Classifier had an AUC score of 0.95. The Area Under the Curve (AUC) is equal to the probability that a classifier will rank a randomly chosen positive instance higher than a randomly chosen negative example. So the logistic Regression model has the best separability between positive and negative results.

The results of our hybrid model were not as good as the individual models. Our hybrid model \$23 ed an Accuracy score of 92.55%, a Recall score of 0.93, a Precision score of 0.93, an F1 score of 0.93, and an AUC score of 0.95.

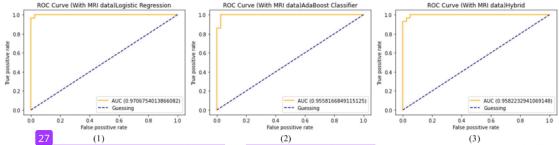


Fig 1. The Receiver Operating Characteristic curve (ROC) and with Area Under the Curve (AUC) score of Logistic Regression (1), AdaBoost Classifier (2), and Our Hybrid model that was created by combining Adaboost classifier, Random Forest, Linear SVC and Logistic Regression with soft voting classifier in the experiment using MRI data.

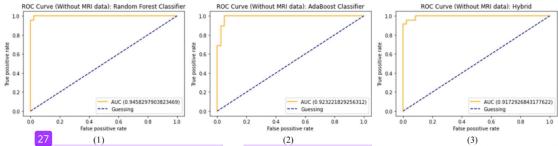


Fig 2. The Receiver Operating Characteristic curve (ROC) and with Area Under the Curve (AUC) score of Random Forest Classifier (1), AdaBoost Classifier (2), and Our Hybrid model that was created by Random Forest Classifier, GaussianNB, LinearSVC, Logistic Regression, and AdaBoost Classifier with soft voting classifier in the experiment without using MRI data.

TABLE I. This table shows the accuracy, recall, precision and f1 score of all the models trained using MRI data.

	49			
Models	Accuracy	recall	precision	F1 Score
Random Forest	<mark>93</mark> .14%	0.93	0.93	0.93
GaussianNB	86.00%	0.86	0.86	0.86
Logistic Regression	94.37%	0.94	0.94	0.94
AdaBoost Classifier	96.07%	0.96	0.96	0.96
Linear SVC	90.14%	0.90	0.90	0.90
Hybrid	92.55%	0.92	0.92	0.92

a) Non-MRI data results: When we trained 24 models without MRI data, Our best performing model was the Random forest classifier. This model scored an Accuracy score of 90.14%, a Recall of 0.90, a Precision of 0.90, and an F1 score of 0.90. The second-best performing models were LinearSVC, and the AdaBoost classifier. Both models achieved an Accuracy score of 89.14%, a Recall of 0.89, a Precision of 0.89, and an F1 score of 0.89. The AUC score for these Random forest Classifier, AdaBoost classifier and LinearSVC were 0.945 and 0.923, 0.92 respectively.

In our training with MRI data KNeighbors Classifier performed very poorly. However, in this case, when trained

TABLE II. This table shows the accuracy, recall, precision and f1 score of all the models trained without MRI data.

Model	Accuracy	recall	precision	F1 Score
Random Forest	90.14%	0.90	0.90	0.90
GaussianNB	88.73%	0.89	0.89	0.89
Logistic Regression	88.73%	0.88	0.88	0.88
AdaBoost Classifier	89.14%	0.89	0.89	0.89
Linear SVC	89.14%	0.89	0.89	0.89
KNeighbors	83.10%	0.83	0.83	0.83
Hybrid	93.37%	0.93	0.93	0.93

without MRI data, this model performed better than before. It is a perfect example that modifying a dataset can improve the performance of a model.

We got our best result when we combined all the models. Our Hybrid model scored an Accuracy score of 93.37%, a Recall of 0.93, a Precision of 0.93, and an F1 score of 0.93. The AUC score for this hybrid model was 0.917.

If we compare this result with our previous results, we can see that GaussianNB and Hybrid model also performed better when trained without MRI data, like KNeighbors Classifier. Linear SVC performed the same as before.

However, our overall Accuracy decreased. Previously our best model had 96.07% Accuracy, but when we removed all the MRI data, our Accuracy was 93.37%. So, we got an Accuracy drop of 2.7%. When it comes to detecting a patient, we always want to avoid False Negatives. So, we can consider the recall score for selecting our optimal model. The Recall decreased by 0.03, from 0.96 to 0.93. Which is not a significant drop. The drop in AUC score was 0.03. It dropped from 0.97 to 0.94.

#### IV. CONCLUSION & FUTURE SCOPE

From our result, we can see that there was no considerable gap in efficiency between our two experiments. We got about a 3% difference. MRI tests have significant importance for monitoring the disease and treatment. Some papers have found better accuracy with MRI data than ours. Fulton et al. [17] found 99.34% correct classification rate using MRI images and 91.3% accuracy in predicting CDR using non-MRI data. So, MRI classification is nearly perfect. So non-MRI methods actually have about 7% gap, which is significant. However, the main objective of our research was not to find an optimal way of predicting Alzheimer's disease but to show that the non-MRI methods are not too inefficient comparing to MRI data analysis methods.

Though the Non-MRI method has good results, it is not perfect. Some Demented patients may not be detected with this approach. It is recommended to do an MRI test if possible for better detection and avoid the risk of falsenegative prediction. False-negative predictions can be dangerous sometimes when it comes to detecting a disease. However, the non-MRI methods can provide reliable results when there is a lack of diagnostic assessment or cannot afford an MRI test. This will reduce their initial cost by a margin. The non-MRI method can be used to raise awareness 25 leveloping nations, as most of them are unaware of Alzheimer's disease.

These results can be further improved by counting more factors related to AD, like depression, head trauma, epilepsy, diabetes, and stroke and similar factors. We firmly believe our research will pave the way to raise awareness and cheap diagnosis of AD with excellent efficiency.

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