**Early Detection of Alzheimer’s using Blood Gene Expression Data and Machine Learning Techniques**

J. Hariharan, R. Jothi

*School of Computer Science and Engineering, Vellore Institute of Technology, Chennai, India*

**Abstract:**

**Alzheimer's disease (AD), a type of neurodegenerative disorder, has seen an increase in cases over the past decade, necessitating the construction of a comprehensive early detection method. Existing methods are typically invasive and costly, so our research concentrates on blood gene expression as a possible biomarker. The high dimensionality of the gene expression data and the small sample size complicate blood gene expression data analysis. Our novel approach attempts to address these issues by identifying a suitable feature selection, a feature extraction method to reduce the dimension size, and synthetic data modelling to address the issue of a small sample size. The classification of the resulting dataset using DNN yielded an accuracy of 90.6%.  Feature selection along with synthetic data modelling significantly enhanced the early detection of Alzheimer's disease using blood gene expression.**

***Keywords: Blood Gene Expression, Feature Selection, Feature Extraction, Synthetic Data Modelling.***

I. INTRODUCTION:

Alzheimer's disease (AD) is a progressive neurodegenerative disorder characterized by the gradual loss of cognitive function and memory. It is the most common cause of dementia among older adults. Especially in India the cases are expected to grow to 11,422,692 by 2050 from 3,848,118 measured in 2019 according to Lancet report as of July 2022 [1]. The disease is caused by the accumulation of amyloid plaques and tau tangles in the brain, leading to the death of nerve cells and the disruption of communication between brain cells. As the disease progresses, individuals may have trouble with everyday tasks, behavioral changes, and eventually, complete dependence on caregivers. Despite intense research efforts, there is currently no cure for AD and available treatments only offer temporary symptom relief. Early detection and diagnosis of AD is crucial for the planning of appropriate care and support for individuals and their families, as well as for the development of disease-modifying therapies. However, current diagnostic methods for AD often involve invasive and expensive procedures, such as brain imaging or lumbar punctures. In recent years, there has been increasing interest in the use of blood-based biomarkers, such as gene expression patterns, as a less invasive and more cost-effective approach for the early detection of AD. The identification of specific gene expression patterns in the blood that are associated with AD may enable the development of simple and reliable diagnostic tools that can be used in a clinical setting.

Gene expression refers to the process by which the genetic information contained in DNA is used to synthesize the various proteins and other molecules that perform specific functions within cells. This process is regulated by a complex network of signaling pathways that control which genes are turned on or off in each cell at a given time. The measurement of gene expression, or transcriptomics, allows scientists to understand how cells respond to different stimuli and how they differ from one another. By analyzing gene expression data, researchers can gain insights into the underlying mechanisms of biological processes and diseases, such as cancer or Alzheimer's disease. Gene expression data can be obtained from a variety of sources, including tissues, cells, and biofluids such as blood. The use of blood-based gene expression data has the advantage of being non-invasive and easily accessible, making it a promising tool for the diagnosis and monitoring of diseases. In recent years, there has been growing interest in the use of gene expression data for the early detection and treatment of a wide range of conditions, including cancer, cardiovascular disease, and neurological disorders. The major problem that we must address while use blood gene expression data is the High Dimensionality of the dataset, since blood tissue can be used to extract around 10,000-30,000 genes on average and each of these genes might have 1-3 gene probes. DNA probes are usually single-stranded DNA molecules that are labeled with a detectable molecule, such as a fluorescent dye or a radioactive isotope. They are designed to bind to a complementary DNA sequence and are often used to detect the presence of specific genes or to analyze DNA modifications, such as methylation. RNA probes are like DNA probes, but they are designed to bind to complementary RNA sequences. They are often used to detect the presence and abundance of specific RNA molecules, such as mRNA or non-coding RNA. Protein probes are molecules that are designed to specifically bind to and detect the presence of a particular protein. They can be antibodies, small molecules, or other types of protein-binding molecules and are often used to analyze protein expression, localization, and function.

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| **Study** | **Dataset Used** | **Feature Selection Alg.** | **No. of Genes** | **Classification Alg.** | **Performance** |
| Lee, T et al. | GSE3060  GSE3061 | SAM | 697 | SVM | AUC: 87.4% |
| H. M. AL-Bermany et al. | GSE3060  GSE3061 | ANOVA + k-means | 2500 | CNN | ACC: 92.9% |
| Kalkan H et al. | GSE63060  GSE63061  GSE140829 | LASSO | 488 | CNN | ACC: 84.2%  AUC: 87.5% |
| El-Gawady, A et al. | GSE33000  GSE44770  GSE44768  GSE44771 | χ2, ANOVA, MI | 30 | SVM | ACC: 97.5%  AUC: 97.2% |
| S. Perera et al. | GSE5281 | PCA, RF, ETC | 14 | SVM | ACC: 93.9% |
| **Table 1:** Summary of studies conducted on detection of AD using various feature selection and classification techniques. | | | | | |

The Table 1 summarizes several studies on the detection of Alzheimer's disease (AD) using different feature selection and classification techniques. These studies utilized various datasets, including GSE3060, GSE3061, GSE63060, GSE63061, GSE140829, GSE33000, GSE44770, GSE44768, GSE44771, and GSE5281. Different feature selection algorithms such as Significance Analysis of Microarrays (SAM), Analysis of Variance (ANOVA), k-means, LASSO, χ2 (Chi-square), and Mutual Information (MI) were employed to identify informative genes. Classification algorithms like Support Vector Machine (SVM), Convolutional Neural Network (CNN), and others were used to build predictive models. The performance metrics reported included Accuracy (ACC) and Area Under the Curve (AUC), with the achieved accuracies ranging from 84.2% to 97.5% and AUCs ranging from 87.4% to 97.2%. These studies highlight the diverse strategies employed to detect AD and showcase varying levels of accuracy in distinguishing between AD and non-AD cases.

II. METHODS:

A. Dataset:

GSE63060 and GSE63061 gene expression samples were collected from GEO gene expression omnibus repository as SOFT formatted family files. GSE63060 contained samples collected from 329 individuals out of which 145 were AD samples, 104 were healthy samples (CTL), and 80 were samples collected from people with Mild Cognitive Impairment (MCI). Each samples contained expression values of 38323 probes, which were mapped to their respective gene symbols using python GEOParse annotation package. If a gene had multiple probe values, Median of the values are taken as the expression value for the gene based on the study done by Lee, T et al. [2]. A total of 29958 unique gene expression values where this extracted and combined with other attributes such as Age, Ethnicity, Gender. Gene expression value where then normalized using Min-Max normalization method and MCI sample were ignored to avoid noise in the dataset.

B. Feature Selection:

1. Chi-square (χ2): χ2 is a statistical method used to determine where there is a statistically significant relation between observed frequency and expected frequency of a particular event. If the difference between the observed and expected values are differ by a large value, then we can reject the null hypothesis and state that the variables are related.
2. Analysis of Variance (ANOVA): ANOVA is a statistical method used to test if two groups of variables related by checking if there is a statistical difference between the mean s of the groups. This is a powerful tool which is now widely used in many fields including biology and psychology.

C. Synthetic Data Modelling:

Synthetic data modeling involves generating artificial data that resembles real-world data to simulate various scenarios for research, testing, and analysis purposes. It allows researchers and data scientists to create controlled environments, explore hypothetical situations, and evaluate the performance of algorithms or models without the need for sensitive or limited data. Synthetic data modeling can aid in data privacy protection, algorithm validation, and developing robust machine learning models that generalize well to real data.

D. Feature Extraction:

1. PCA: PCA is a statistical method that involves linear transformation of the given dataset into a new system where the sample can be represented with few dimensions. This technique aims to find the set principal components that exhibit maximum variance. Due to this PCA is used widely in genetic studies to find important gene sets from the given set of gene in the sample.
2. Linear Discriminant Analysis (LDA): Linear Discriminant Analysis, is a statistical technique used for dimensionality reduction and classification tasks. It aims to find a linear combination of features that maximizes the separation between classes while minimizing the within-class scatter.

E. Classification Technique:

1. Support Vector Machine (SVM): SVM is the most popular and widely used linear classifier technique. This involves classification based on supervised learning approach.
2. Ensemble Classifiers:
   1. AdaBoost: Adaboost technique is an Ensemble method using decision trees. These uses set of weak learners for fast implementation and to converge faster into the results and doesn’t require any prior domain knowledge in creating the weak learners. The goal of the weak leaners is to identify the weak hypothesis.
   2. Random Forest (RF): RF is a most popular ensemble technique which is used for task such as classification and regression. From the give sample of dataset random sub samples are generated for each of which a classifier is modelled. The classifiers are then ensembled together to make the final classification. This technique can be used to extract the import feature by using a metric called as feature importance. For a feature this metric is found by identifying how much role it plays in its respective decisions trees classification accuracy.
3. Deep Learning Classifiers:
   1. DNN: DNNs with output layers having SoftMax activation functions are generally used for designing classification models. The number of layers, nodes at each layer, activation and loss functions are each layer depend on the domain over which classification is applied. DNN models with classification done using gene expression values with large dimensions require model to be complex with large number of hidden layers. If number of unique gene are fewer complex models lead to overfitting.
   2. CNN: A common deep learning model predominantly used for classification tasks, such as image classification, image segmentation and object detection. Has special layers called convolution layers which perform mathematical operation called convolution on the input data based on a kernel with static size. CNN models also use polling layers to down sample or to reduce dimension using min, max or average of the values over a small region. Though CNN are mainly designed for image classification this can be extended to classification of AD using gene expression as biomarker.

RESULTS AND DISCUSSION:

|  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- |
| **Feature Selection** | **Feature Extraction** | **No of Genes Selected** | **Classification** | **Modelled Synthetic Data** | **Accuracy (Training Set)** | **Accuracy (Testing Set)** |
| Chi Square | - | 10814 | SVM Gaussian Kernel | No | 99.67% | 77% |
| Chi Square | - | 10814 | DNN (64, 128, 128, 1) | No | 97.13% | 77.33% |
| Chi Square | - | 10814 | DNN (6, 4, 4, 1) | No | 96.68% | 89.33% |
| Chi Square | PCA | 30 | RF | No | 100% | 75% |
| ANOVA | PCA | 30 | RF | No | 100% | 82% |
| ANOVA | - | 514 | AdaBoost | No | 100% | 90.20% |
| ANOVA | - | 514 | RF | No | 100% | 86.45% |
| ANOVA | - | 514 | SVM Gaussian Kernel | No | 99.83% | 87.42% |
| ANOVA | - | 514 | DNN (6, 5, 5, 5, 1) | Yes | 93.77% | 90.60% |
| **Table 2:** Summary of results observed while classifying AD from CTL samples. | | | | | | |

|  |  |
| --- | --- |
|  |  |
| **Fig 1:** Loss and Accuracy variation chart over 100 epochs in Training and Testing set. | |

CONCLUSION:

REFERENCES:

1. Nandi, A et al, “Global and regional projections of the economic burden of Alzheimer's disease and related dementias from 2019 to 2050: A value of statistical life approach”, EClinicalMedicine - The Lancet Discovery Science, Volume 51, 101580, 2022. https://doi.org/10.1016/j.eclinm.2022.101580.
2. Lee, T., Lee, H. Prediction of Alzheimer’s disease using blood gene expression data. Sci Rep 10, 3485, 2020. https://doi.org/10.1038/s41598-020-60595-1.
3. M. S. Kamal, A. Northcote, L. Chowdhury, N. Dey, R. G. Crespo and E. Herrera-Viedma, "Alzheimer’s Patient Analysis Using Image and Gene Expression Data and Explainable-AI to Present Associated Genes," in IEEE Transactions on Instrumentation and Measurement, vol. 70, pp. 1-7, 2021, Art no. 2513107, doi: 10.1109/TIM.2021.3107056.
4. Mahendran N, P M DRV. A deep learning framework with an embedded-based feature selection approach for the early detection of the Alzheimer's disease. Comput Biol Med. 2022 Feb; 141:105056. doi: 10.1016/j.compbiomed.2021.105056. Epub 2021 Nov 22. PMID: 34839903.
5. S. Khanal, J. Chen, N. Jacobs, and A. -L. Lin, "Alzheimer’s Disease Classification Using Genetic Data," 2021 IEEE International Conference on Bioinformatics and Biomedicine (BIBM), 2021, pp. 2245-2252, doi: 10.1109/BIBM52615.2021.9669730.
6. El-Gawady, A.; Makhlouf, M.A.; Tawfik, B.S.; Nassar, H. Machine Learning Framework for the Prediction of Alzheimer’s Disease Using Gene Expression Data Based on Efficient Gene Selection. Symmetry 2022, 14, 491. https://doi.org/10.3390/sym14030491.
7. S. Perera, K. Hewage, C. Gunarathne, R. Navarathna, D. Herath and R. G. Ragel, "Detection of Novel Biomarker Genes of Alzheimer’s Disease Using Gene Expression Data," 2020 Moratuwa Engineering Research Conference (MERCon), 2020, pp. 1-6, doi: 10.1109/MERCon50084.2020.9185336.
8. C. Park, J. Ha, and S. Park, "Prediction of Alzheimer's disease based on deep neural network by integrating gene expression and DNA methylation dataset," Expert Systems with Applications, vol. 140, pp. 112873, 2020, doi: 10.1016/j.eswa.2019.112873.
9. H. M. AL-Bermany and S. Z. AL-Rashid, "Microarray Gene Expression Data for Detection Alzheimer’s Disease Using k-means and Deep Learning," 2021 7th International Engineering Conference “Research & Innovation amid Global Pandemic" (IEC), 2021, pp. 13-19, doi: 10.1109/IEC52205.2021.9476128.
10. A. Sharma, P. Dey, "A Machine Learning Approach to Unmask Novel Gene Signatures and Prediction of Alzheimer's Disease Within Different Brain Regions," Genomics, vol. 113, no. 4, pp. 1778-1789, Apr. 2021, doi: 10.1016/j.ygeno.2021.04.028.
11. Kalkan H, Akkaya UM, Inal-Gültekin G, Sanchez-Perez AM. Prediction of Alzheimer's Disease by a Novel Image-Based Representation of Gene Expression. Genes (Basel). 2022 Aug 8;13(8):1406. doi: 10.3390/genes13081406. PMID: 36011317; PMCID: PMC9407775.
12. D. Sun, H. Peng and Z. Wu, "Establishment and Analysis of a Combined Diagnostic Model of Alzheimer's Disease with Random Forest and Artificial Neural Network," Frontiers in Aging Neuroscience, 2022. Available: https://www.proquest.com/scholarly-journals/establishment-analysis-combined-diagnostic-model/docview/2682564611/se-2. DOI: https://doi.org/10.3389/fnagi.2022.921906.
13. Yuen, S.C., Liang, X., Zhu, H., Jia, Y., and Leung, S.W. "Prediction of differentially expressed microRNAs in blood as potential biomarkers for Alzheimer's disease by meta-analysis and adaptive boosting ensemble learning." Alzheimer's Research & Therapy, vol. 13, no. 1, 2021, p. 126. doi: 10.1186/s13195-021-00862-z.
14. S. Pavalarajan, B. A. Kumar, S. S. Hammed, K. Haripriya, C. Preethi and T. Mohanraj, "Detection of Alzheimer's disease at Early Stage using Machine Learning," 2022 International Conference on Advanced Computing Technologies and Applications (ICACTA), Coimbatore, India, 2022, pp. 1-5, doi: 10.1109/ICACTA54488.2022.9752827.
15. S. S. Rajeswari and M. Nair, "A Transfer Learning Approach for Predicting Alzheimer's Disease," 2021 4th Biennial International Conference on Nascent Technologies in Engineering (ICNTE), NaviMumbai, India, 2021, pp. 1-5, doi: 10.1109/ICNTE51185.2021.9487746.
16. J. Li, Y. Wei, C. Wang, Q. Hu, Y. Liu and L. Xu, "3-D CNN-Based Multichannel Contrastive Learning for Alzheimer’s Disease Automatic Diagnosis," in IEEE Transactions on Instrumentation and Measurement, vol. 71, pp. 1-11, 2022, Art no. 5008411, doi: 10.1109/TIM.2022.3162265.
17. S. Basheer, S. Bhatia and S. B. Sakri, "Computational Modeling of Dementia Prediction Using Deep Neural Network: Analysis on OASIS Dataset," in IEEE Access, vol. 9, pp. 42449-42462, 2021, doi: 10.1109/ACCESS.2021.3066213.
18. G. Chutani, H. Bohra, D. Diwan and N. Garg, "Improved Alzheimer Detection using Image Enhancement Techniques and Transfer Learning," 2022 3rd International Conference for Emerging Technology (INCET), Belgaum, India, 2022, pp. 1-6, doi: 10.1109/INCET54531.2022.9824008.
19. D. Chaihtra and S. Vijaya Shetty, "Alzheimer’s Disease Detection from Brain MRI Data using Deep Learning Techniques," 2021 2nd Global Conference for Advancement in Technology (GCAT), Bangalore, India, 2021, pp. 1-5, doi: 10.1109/GCAT52182.2021.9587756.
20. U. R. K, S. S. S, U. M. G and V. B. C, "Binary Classification of Alzheimer's disease using MRI images and Support Vector Machine," 2021 IEEE Mysore Sub Section International Conference (MysuruCon), Hassan, India, 2021, pp. 423-426, doi: 10.1109/MysuruCon52639.2021.9641661.
21. S. Buyrukoğlu, "Improvement of Machine Learning Models’ Performances based on Ensemble Learning for the detection of Alzheimer Disease," 2021 6th International Conference on Computer Science and Engineering (UBMK), Ankara, Turkey, 2021, pp. 102-106, doi: 10.1109/UBMK52708.2021.9558994.
22. Dubey, S. (2019, December 26). Alzheimer’s Dataset (4 class of Images), Kaggledatase. https://www.kaggle.com/tourist55/alzheimers-dataset-4-class-of-images?select=Alzheimer\_s%2Bt.