Predicting Alzheimer's in Patients

Final Project

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DATA 824

Purpose

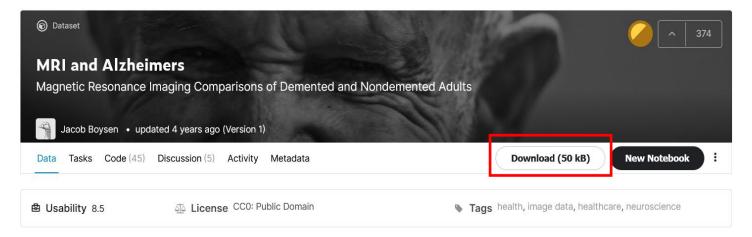
- Conduct an exploratory data analysis (EDA)
 on a dataset containing MRI and Alzheimer's
 patient data
- Conduct an analysis to predict (classify)
 whether a patient has dementia or not
- Look at available features in the data to determine importance in predicting whether patient has dementia

Overview: OASIS

- Open Access Series of Imaging Studies (OASIS)
 - Website: https://www.oasis-brains.org/
 - A project that focuses on providing neuroimaging datasets for scientific use
 - Project responsible for making dataset freely available
 - Data is generated by the Knight ADRC and its affiliated studies
 - Website: https://knightadrc.wustl.edu/

Data Acquisition

- Source: <u>https://www.kaggle.com/jboysen/mri-and-alzheimers</u>
 - Note: Data is made available and credited to the OASIS project
- How to Download via Kaggle:



- File Type: ZIP file
 - Contains 2 CSV files
 - File 1: oasis_cross-sectional.csv
 - File 2: oasis_longitudinal.csv

Data: Features

• Number of Features: 12 (see table)

Feature	Description	Туре
ID	Identification (Patient)	Character
M.F	Gender (Male or Female)	Character/Nominal
Hand	Dominant Hand	Character
Age	Age in Years	Numeric
Educ	Education Level	Numeric/Ordinal
SES	Socioeconomic Status	Numeric/Ordinal
MMSE	Mini Mental State Examination	Numeric
CDR	Clinical Dementia Rating	Numeric/Ordinal
eTIV	Estimated Total Intracranial Volume	Numeric
nWBV	Normalize Whole Brain Volume	Numeric
ASF	Atlas Scaling Factor	Numeric
Delay	Time (Feature Unknown)	Numeric

Data: Missingness

- Table shows the missingness by number of observations
- 5 variables have significant amount of missingness:
 - Educ, SES, MMSE, CDR and Delay
- What can be done?
 - Go back and try to get values from the study
 - Omit missing records/observations
 - Imputation

Feature	Missingness (# of obs)
ID	0
M.F	0
Hand	0
Age	0
Educ	201
SES	220
MMSE	201
CDR	201
eTIV	0
nWBV	0
ASF	0
Delay	416

Data: Missingness

- What was done?
 - Trying to retrieve missing data from the previous study was not possible. Therefore, the option chosen was to omit records/observations with significant missingness.
 - This is likely not the best option
 - The option for imputation was not utilized but could be used in the future

Data: Preparation

- Based on the features there are several that will be dropped for the analysis:
 - ID Is a unique identifier and is unnecessary (not helpful) in prediction
 - Hand There is only one option in the dataset and that's R (right-handed)
 - Delay Missingness in the data is approximately 95%
- Based on the features there are several that are ordinal and need to be converted to a factor:
 - M.F, Educ, SES, CDR
 - CDR will be converted to a two-level factor (dementia/Yes or no dementia/No) for prediction, i.e. binary variable

Data: Privacy & Security

- The data has been redacted such that information that can identify the patient is unavailable in this dataset
- Since this is actual patient information as part of the OASIS project it is important to follow the data agreement when using the data
 - Visit website: https://www.oasis-brains.org/ for details

Exploratory Data Analysis: Visualization

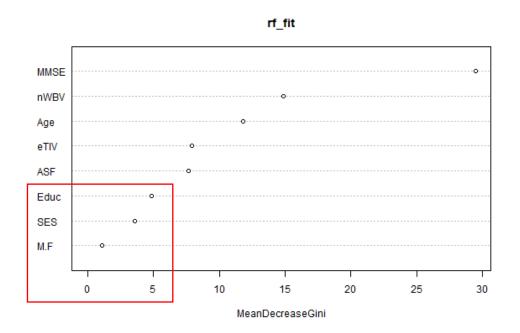
- Several data visualizations (plots/charts) were created:
 - Density plot showing Age vs. Dementia
 - Correlation Plot for numerical variables
 - Histogram for MMSE
 - Bar plots for categorical variables
 - Educ (Education Score)
 - SES (Socioeconomic Status Score)
 - M.F (Gender)
 - Boxplots for numerical variables
 - ASF
 - MMSE
 - eTIV
 - nWBV
 - Correlation plot for numerical variables (interactive)

Exploratory Data Analysis: Takeaways

- Patients with dementia have ages between 60 and 90 years of age
- Patients with a high MMSE score (approximately 25) tend to not have dementia
 - MMSE seems to be an indicator of dementia
- Patients with higher nMBV values tend to not have dementia
- Factors such as Education and Socioeconomic Scores don't seem to indicate dementia
- Correlated variables include:
 - eTIV and ASF: -0.99 correlation
 - nWBV and Age: -0.74 correlation
 - MMSE and nWBV: 0.48 correlation

Model Building: Feature Selection

- There were 8 features in the dataset, but reduced to 5 features after conducting a Random Forest feature selection method
 - The criteria of a mean decrease in Gini value of 5 or less was used arbitrarily
 - Educ, SES and M.F were dropped from the model based on this criteria



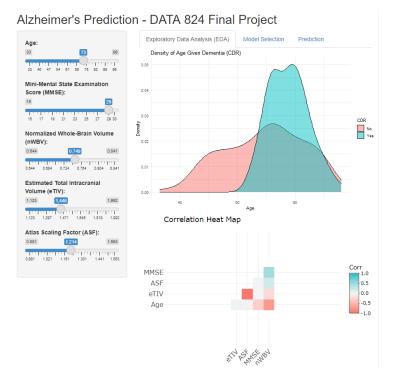
Model Building: Final Model

- Data was split into test and validation datasets and cross-validation was utilized to determine final model
 - The metric for assessment was the F1 Score
- Models selected for comparison were:
 - Logistic Regression
 - Single Decision Tree
 - KNN
 - Flexible Discriminant Analysis (FDA)
 - Support Vector Machine (SVM) using a radial kernel
- The "best" model i.e. final model was the SVM model with an F1 Score of 0.91

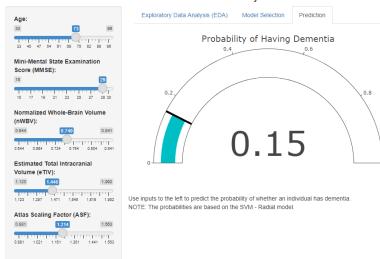
Shiny App: Description & Location

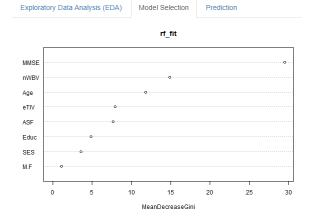
- The Alzheimer's Prediction app was developed using the R package, Shiny
- The layout is structured using 1 side panel for inputs and 3 tabs where information is displayed
 - Tab 1 shows visualizations as part of the exploratory data analysis (EDA)
 - Tab 2 shows visualizations as part of the model selection process
 - Tab 3 shows a gauge visualization and utilizes the slider inputs to predict the probability that a patient has dementia
- The code can be located at the following GitHub repository
 - https://github.com/wacorda/MSASADS-DATA-824-Final-Project

Shiny App: Screenshots



Alzheimer's Prediction - DATA 824 Final Project





NOTE: Based on the MeanDecreaseGini, any feature that is less than 5 was removed from the model. Three features were removed: Educ, SES and M.F.

Models	F1 Score
Logistic Regression	0.89
Single Decision Tree	0.88
KNN	0.77
Flexible DA	0.89
SVM - Radial	0.91

NOTE: The final model selected was the SVM - Radial as it had the highest F1 Score (0.91).

Conclusion

- I hope you enjoyed my Shiny app that was used to predict the probability that a patient has dementia
- If you have any questions, please feel free to contact me
 - E-mail: wacorda@kumc.edu