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1. INTRODUCTION

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Diabetes is a metabolic disease impacting 37.3 million Americans. Those affected by the disease have complications producing insulin, a chemical messenger that our body uses to store energy. Although it is uncommon, diabetics can be hospitalized for having critically low or high blood glucose levels. These hospitalizations can be life threatening and should be minimized at all costs.

In this case study, we will use a diabetes data set procured by Dr. Slater, to identify what factors most significantly result in diabetics getting readmitted to hospitals. To accomplish this, we will build a Logistic Regression model and extract its respective feature importances. It is our hope that this research can be leveraged by medical professionals to help treat hospitalized diabetics and to ensure that these patients are not readmitted in the future.

DATA UNDERSTANDING:

2. METHODS

Data used in this case study was a diabetes.csv provided by Dr. Slater. Our diabetes.csv contained data related to hospitalized diabetic patients including columns such as: readmitted, patient_nbr, insulin, and time in hospital, Upon reviewing the contents of our data set, we saved the data into a data frame named "diabetes_data" and began pre-processing.

The first step we performed in pre-processing was reviewing our full data set. Immediately, we recognized that missing values existed in the columns of: 1. race

DATA PREPROCESSING:

2. weight 3. payer_code

4. medical_specialty 5. diag_1

6. diag_2

7. diag_3

35000

DATA IMPUTATION:

data set.

felt this represented our data the best.

RE-CODING CATEGORICAL COLUMNS:

Given that machine learning models do not handle missing values well, we imputed them using appropriate statistical methods. Full details on how these columns were imputed can be found in the sub-header of this case study titled, "Data Imputation."

Our data frame contained 101,766 rows. Our data frame contained 50 columns. No null values existed in our data frame contained 13 numeric columns. Our data frame contained 37 categorical columns.

After identifying that missing values existed in our data set, we ran the command, diabetes_data_info() and noted the following details of our data frame:

From this output we recognized that one hot encoding (OHE), would need to be performed on our categorical columns. For additional details on our OHE process, please see the sub-header of this case study titled, One Hot Encoding.

The next step performed in pre-processing was running the command diabetes_data.describe() to view the summary statistics of our data frame. Output from this command showed that several columns contained outliers. This was something we remained cognizant of throughout our analysis. Finally, to view the distributions of our categorical columns with missing values, we created count plots. Visualizing these columns was important, as it helped us determine what data imputation method was most

type, we noted that imputing these columns with either the mean or median value would be appropriate. 40000

appropriate for our data. Output from our count plots showed that all seven of our columns with missing data contained non-normal distributions (Exhibit 1.0). Since all seven columns were of the categorical data

30000 25000 count 20000 -15000 10000 5000 payer_code Exhibit 1.0: Count Plot of Payer_Code The last step performed in pre-processing was calculating the percentage of missing values in our categorical columns. Exhibit 1.1 details our findings: **Percentage of Missing Values Column Name**

0.02% diag_1 0.35% diag_2

Exhibit 1.1: Percent Missing Values

Upon reviewing our full data set and calculating the percentage of missing values that existed in our categorical columns, we proceeded to impute our missing values.

All the missing values in our data set were denoted by: ? . Since computers cannot impute data with special characters, we converted the question marks to "NaN". Once

2.23%

96.85%

39.55%

diag_3 1.39%

race

weight

payer_code

this was complete, we re-calculated the sum of missing values in our columns to validate that no data loss had occurred in our conversion process.

No

Yes

0.1

0.01

0.9

0.9

0.9

0.1

0.09

metformin

repalglinide

nateglinide

glimepiride

glipizide

glyburide

chloropropamide

diabetesMed

medical_specialty 49.08%

When we considered imputing the columns: race, payer_code, medical_specialty, diag_1, diag_2, and diag_3, we tried two different approaches. One approach was imputing these columns with the mode of each column, and the second approach was leaving the columns as is with missing values. We fit our Logistic Regression model on both approaches and found that our performance results were negligible. Consequently, we decided to leave the columns with missing values, as we

Next, we imputed the columns: diabetesMed, change, and readmitted with values of 0 and 1. This was done to simplify OHE as these columns had a maximum of three classes. Please note that although the column: readmitted contains three classes, we chose to convert it to a binary variable as we are only concerned with whether a patient has been readmitted or not. Exhibit 1.2 details our conversion process of these columns: Column Name Original Classes Data Dictionary for Converted Classes

0=No

1=Yes

Exhibit 1.2: Imputation process for the columns: "diabetesMed", "change", and" readmitted"

The first column we chose to impute was our weight column. Given that 96% of the data in our weight column were missing, we chose to drop the column from our

Ch 0=No Change No 1=Yes NO 0=No <30 readmitted 1=Yes >30

result, we decided to reduce the classes in each categorical variable by specifying a threshold for infrequent observations. Exhibit 1.3 details the thresholds that were chosen for each variable, as well as explanations as to why thresholds were chosen. **Column Name Selected Threshold Explanation** ~90% of our data falling into the top 7 classes 0.02 payer_code medical_specialty 0.03 ~85% of our data falling into the top 5 classes max_glu_serum ~96% of our data falling into the top 2 classes 0.02 A1Cresult 0.08 ~91% of our data falling into the top 2 classes

~98% of our data falling into the top 2 classes

~99% of our data falling into the top 2 classes

~99% of our data falling into the top class

~99% of our data falling into the top class

~95% of our data falling into the top class

~97 of our data falling into the top 2 classes

~98 of our data falling into the top 2 classes

When viewing the shape of our data set, we recognized that if we one hot encoded all 37 of our categorical variables, that our data set would be extremely wide. As a

pioglitazone 0.06 ~98 of our data falling into the top 2 classes rosiglitazone 0.05 ~98% of our data falling into the top 2 classes acarbose 0.9 ~99% of our data falling into the top class 0.9 ~99% of our data falling into the top class miglitol 0.0004 tolazamide ~99% of our data falls into the top class ~99% of our data falling into the class "No" glyburide_metformin 0.9 0.0075 many of the columns contained values <= 0.000010 diag_1 diag_2 0.0075 many of the columns contained values <= 0.000010 many of the columns contained values <= 0.000010 diag_3 0.0075 Exhibit 1.3: Detailed Recoding Threshold Once we imputed our missing values and re-coded our categorical columns we separated our diabetes data set into two variables, one containing all our numeric columns and the other containing all our categorical columns. Next, using Pandas' get dummies function, we one hot encoded our categorical columns and joined our one hot encoded data to our numeric columns to arrive at our final full data set. Please note that for modeling, we scaled our non-hot encoded data to ensure that our full data set was on the same scale. For additional modeling details, please see our For our exploratory data analysis (EDA), we began by viewing histograms and pair plots of our data (Exhibits 1.4 & 1.5). Two takeaways from these visualizations included:

2. Our variables were not on the same scale.

modeling assumptions.

3. Lack of outliers.

"num_medications"

1.0

0.8

0.4

0.2

0.0

encounter_id

metformin_pioglitazone_No

0.8

0.6

0.4

0.2

0.0

assumption was met.

1.0

0.8

0.2

0.0

encounter_id

0.4

0.2

0.0

12.5

10.0

5.0

Model HalvingRandomSearchCV:

readmitted

readmitted

2. Absence of multicollinearity.

ONE HOT ENCODING:

sub-header below titled "Modeling."

EXPLORATORY DATA ANALYSIS:

5

encounter_id

3

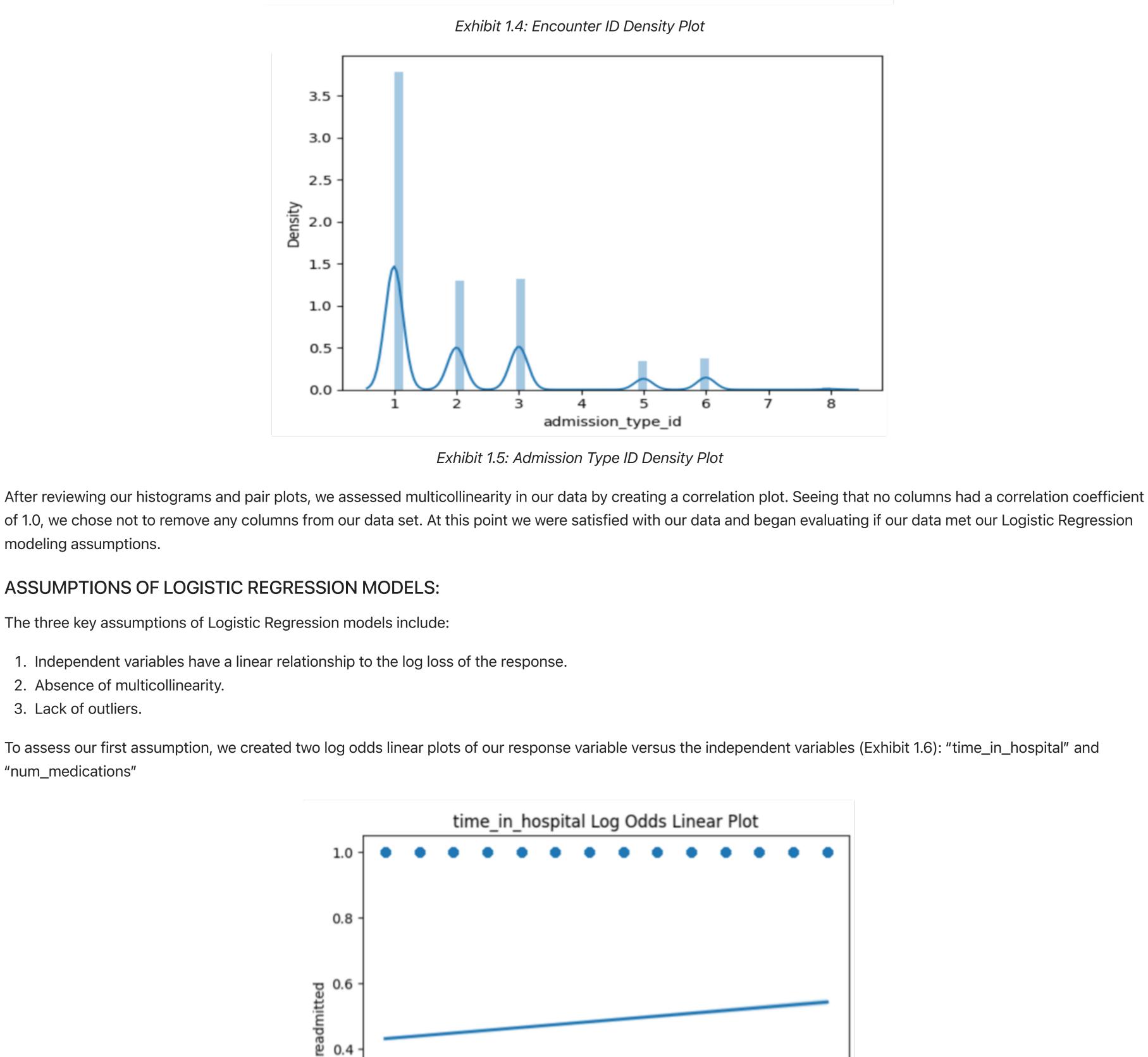
1e8

Density

2

1. Many of our numeric columns exhibited non-normal distributions.

1e-9



0.2 0.0

10

8

time_in_hospital

num_medications Log Odds Linear Plot

Exhibit 1.6: Log Odds for Time in Hospital

1.0

0.6

12

14

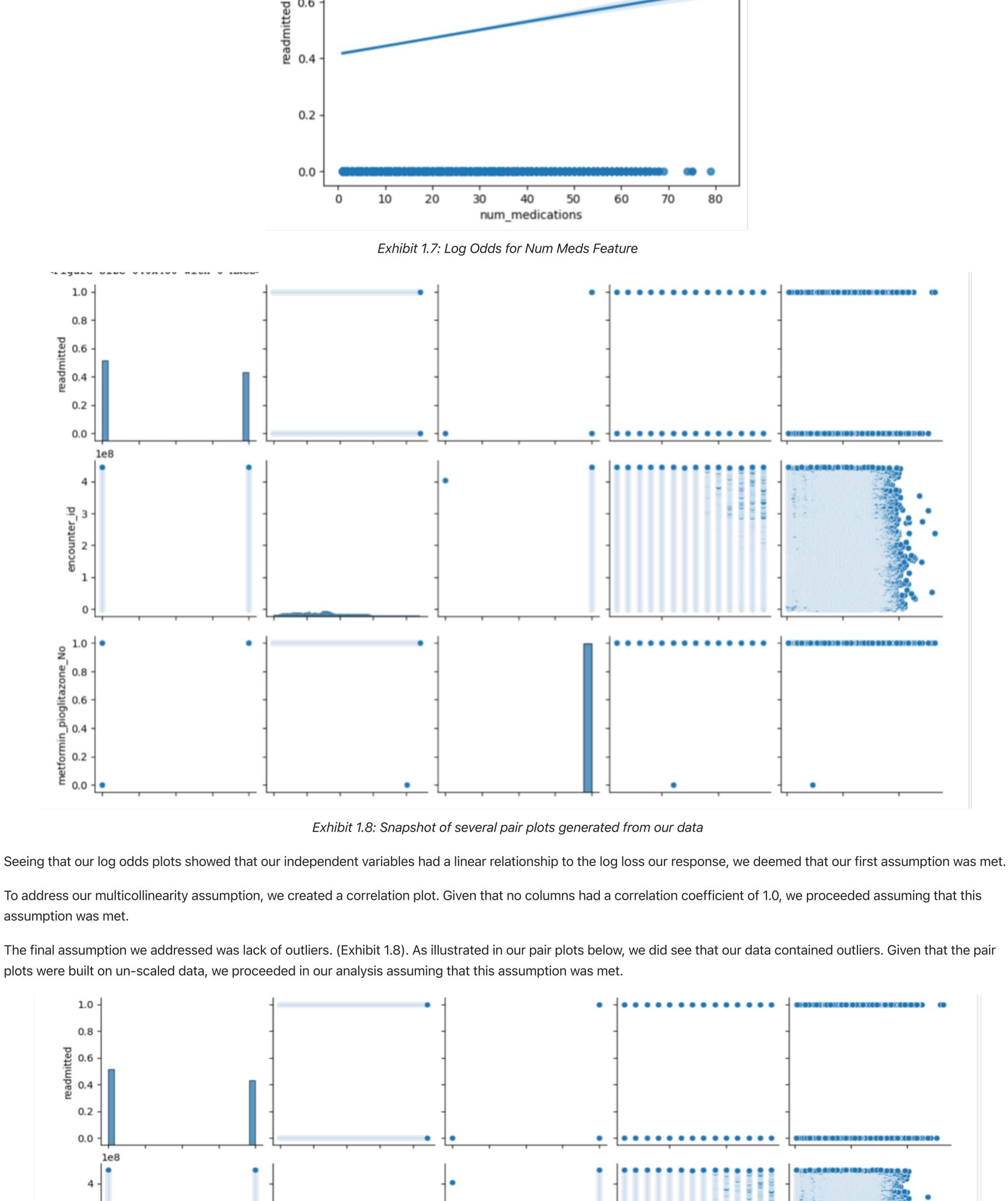


Exhibit 1.9: Addressing our Logistic Regression outlier assumption with pair plots

After preprocessing, EDA, and scaling the data, modeling was able to begin. To determine the best hyperparameters that we should use, we needed to iterate through

several of sklearn's modules. We began with utilizing GridSearchCV, but due to the shape of our data and inability to scale our CPU, GPU, and Memory for the

needs of this project, GridSearchCV was unable to complete and we needed to try other methods of tuning hyperparameters. With the use of Skelearn's

experimental and model_selection packages we were able to utilize HalvingRandomSearchCV to obtain good, but potentially not the best hyperparameters for this model. HalvingRandomSearchCV combines the idea of HalvingSearchCV and RandomizedSearchCV. HalvingSearchCV works by modeliong all potential candidates with less data and selects half of the best performing models to add additional resources and data until a "best" model is output. RandomizedSearchCV randomly picks candidate modles from the grid to model.

3. MODEL BUILDING & RESULTS

The use of Sklearn's LogisticRegression was used to model the data for this case study.

Halving Random Search CV Parameters: "C": np.logspace(-3,3,7),"l1_ratio": np.arange(0.0,1.0,0.1), 'solver': ['saga'],

We passed the below parameters into HalvingRandomSearchCV and the best model outputs were the following:

Models use: 10-fold Cross validation (Kfold), random_seed = 0, and max_iter = 50000, and scoring metric of F1

"solver": 'saga' "tol": 0.001 **ElasticNetCV with GridSearchCV Tuned Parameters:**

> 1 3.08698 0.00623727 LogisticRegression(C=100.0, I1_ratio=0.8, n_jobs=-1, penalty='elasticnet',random_state=0, solver='saga', tol=0.001) 0.56834 0.568627 3.93618 0.00589871 LogisticRegression(C=100.0, I1_ratio=0.8, n_jobs=-1, penalty='elasticnet',random_state=0, solver='saga', tol=0.001) 0.567456 0.568824 2 3 0.566277 0.56902 3.40509 0.00616121 LogisticRegression(C=100.0, I1_ratio=0.8, n_jobs=-1, penalty='elasticnet',random_state=0, solver='saga', tol=0.001) 4 2.62886 0.00576496 LogisticRegression(C=100.0, I1_ratio=0.8, n_jobs=-1, penalty='elasticnet',random_state=0, solver='saga', tol=0.001) 0.56726 0.568922 5 0.00782394 LogisticRegression(C=100.0, I1_ratio=0.8, n_jobs=-1, penalty='elasticnet',random_state=0, solver='saga', tol=0.001) 0.572271 0.568409 3.54918 0.00629807 LogisticRegression(C=100.0, I1_ratio=0.8, n_jobs=-1, penalty='elasticnet',random_state=0, solver='saga', tol=0.001) 0.574194 0.567999 2.96987 0.00986385 LogisticRegression(C=100.0, I1_ratio=0.8, n_jobs=-1, penalty='elasticnet',random_state=0, solver='saga', tol=0.001) 0.573015 0.568162 LogisticRegression(C=100.0, I1_ratio=0.8, n_jobs=-1, penalty='elasticnet',random_state=0, solver='saga', tol=0.001) 0.561812 0.56956 LogisticRegression(C=100.0, I1_ratio=0.8, n_jobs=-1, penalty='elasticnet',random_state=0, solver='saga', tol=0.001) 0.568101 0.568796 MEAN 3.18602 0.00685496 0.568746 0.568688 Exhibit 2.0: 10 Fold Cross Validation Model Results

In conclusion, after significant updates to thresholds and hyperparameters, we have determined that logistic regression does not properly model this data due to inablilty for the coefficient's to converge. Potential models that would be better for this data set would include decision trees or any sort of gradient boosting.

5. CODE:

Attached in file CS2_CODE.ipynb

4. CONCLUSION

['elasticnet'], 'penalty': "tol": [1e-9, 1e-8, 1e-7, 1e-6, 1e-5, 1e-4, 1e-3, 1e-2, 1e-1]**Best Model Output:** "C": 100.0 "l1_ratio": 0.8 "n_jobs": "penalty": 'elasticnet' After performing HalvingRandomSearchCV to tune the model parameters, Sklearn's cross_validate was used to validate the model and determine final performance. The results of all 10 folds are below with a mean F1 score of .568746. Model results for the 10 fold Cross Validation are below in Exhibit 2.0. fit_time score_time estimator test_score train_score 0 2.79806 0.00683784 LogisticRegression(C=100.0, I1_ratio=0.8, n_jobs=-1, penalty='elasticnet',random_state=0, solver='saga', tol=0.001) 0.568733 0.568562