

Fraud Detection in Healthcare Insurance Claims

Ashpak Rakeeb
B00913796

Shravya Reddy
B00911193

Sri Ramya
B00900307

Venkata Vijaya
B00912916

Meagan Sinclair
B00737317

Abstract—In terms of the value of our healthcare system and in terms of money, fraudulent claims come with a very high price. Healthcare spending in the United States totals at 3.6 trillion dollars in 2018, which equates to billions in insurance claims. [1]. There is no denying that some of these claims are fraudulent. [1]. In a huge set of Medicare data (more than 550,000 claims), we developed a process to predict potential fraud in medical insurance claims where the doctor files the paperwork and submits the claim on the patient’s behalf. We experimented with four distinct approaches: Decision Tree Classifier, XGBoost Classifier, Bagging Classifier, and Stacked Classifier. According to our experiments, the XGBoost classifier had the best generalization and a 78% accuracy rate on the reserved test set. The other models’ performances were likewise satisfactory, with the lowest-scoring model, the Stacked Classifier, having an accuracy of 72%.

I. INTRODUCTION

Every year, healthcare fraud costs the industry billions of dollars as fraudsters find the healthcare industry an attractive target [1]. Common healthcare frauds include beneficiaries claiming medical conditions for which they were not treated and submitting multiple claims for the same service. Claiming more than the actual monetary cost spent on treatment is also a common fraud technique [2].

If insurance companies don’t have procedures in place to identify and stop fraudulent practices, they are at risk to lose significant profit. Further, if fraud detection techniques are unreliable, healthcare providers could be incorrectly subjected to an inquiry that could harm their reputation and revenue, as well as damaging the reputation and wasting resources of the insurance provider. This can raise health insurance premiums for customers. Insurance companies prefer to make payments without conducting investigations since doing so is expensive and time-consuming. Additionally, incorrectly detecting fraudulent claims could delay reimbursement, which is undesirable for honest customers and degrades the public opinion of the insurance provider [3].

To solve this business problem, insurance providers now use predictive analytical methods to prevent false medical bills before healthcare providers receive payments. The goal of this project is to create a model which can predict whether claim is potentially fraudulent or not based on the claim information. This would

allow the insurance providers to assess claim details and determine if further fraud investigation is required. This is a supervised classification problem as we have a specific binary target, potential fraud.

II. LITERATURE REVIEW

Several researchers have tackled the problem of fraud detection in healthcare insurance claims. Multiple clustering methods have been employed to identify suspicious claims within databases and provide insight into groups within the data [4]. This is a useful basis for further work toward classification but does not provide conclusive fraud detection. Bayesian co-clustering methods have also been employed [5]. This method is beneficial for the problem as co-clustering allows for modeling relationships between entities and potentially better results. There is little work done on ensemble models or boosted learning algorithms [6].

The accompanying notebook on Kaggle.com utilized feature selection methods and showed testing of various models [7]. Model types trialed included logistic regression, Random Forest classifier, and auto-encoders with good results, around 90% accuracy on test data. The author noted that increasing the database size, vectoring medical codes, and utilizing further ensemble models could lead to better results. The author of the notebook also noted the hyperparameter tuning was not performed on the models, so careful tuning will also be a focus of the project. While this previous work attempted to predict fraud for each provider from the information of all claims belonging to the provider, this project will attempt to predict fraud based on a single claim. This is more useful for the business problem as fraud could be identified immediately after 1 claim is submitted, as opposed to requiring multiple claims information, which is less timely.

This project is valuable as various new model types will be applied to the problem and evaluated. Ensembled and boosted models will also be tested to determine if performance can be further improved. This work extends current knowledge of fraudulent claim detection methods.

III. METHODOLOGY

A. Data Exploration

The data set chosen for the project contains relevant information on fraudulent and non-fraudulent health-care claims [8]. This data is open source and is hosted on Kaggle.com. The raw data is broken down into 4 sections: inpatient data, outpatient data, beneficiary details data, list of possibly fraudulent providers.

Inpatient data contains information about claims made by beneficiaries who were admitted into the hospital. The data describes the corresponding beneficiary (the person who submitted the claim), claim date range, relevant monetary amounts, provider and physician codes, diagnoses information and procedure information. These values are summarized below in Figure 1a. Outpatient data describes claims made by patients who were not admitted. The data contains the same fields as inpatient data, excluding the admission and discharge dates.

Beneficiary data includes information describing the person who submitted the claim. This includes date of birth and death, gender, race, residential location, the status of various chronic conditions, length of time covered by insurance, and relevant annual monetary values such as deductibles and reimbursements. These values are summarized below in Figure 1b.

Finally, the last data set contains a list of provider codes marked as either potentially fraudulent or not. The domain concepts graph of the entire feature set is shown below in Figure 2. The corresponding Analytic Base Table with all raw features and their descriptions is shown in Figure 3.

Since the data is split in separate subsets, merging the data into one comprehensive set was the first step. The inpatient and outpatient data sets were merged and the corresponding potential fraud indicator based on the provider value was added to each claim. Next, separate categorical and continuous feature reports were created for the combined data set, these are shown in Appendix A and B.

B. Data Pre-Processing

Several steps were required to prepare the data for model training. First, the binary features such as disease indicators, chronic conditions, and potential fraud indicator were transformed into binary values 0 and 1. Since the claim codes contained alphanumeric strings to indicate the diagnostic claim, these were re-encoded into numerals. The procedure codes were strictly numeric so these were left.

Next, derived features were selected and created. The date of birth and date of death were transformed into an age and a binary feature indicating if the beneficiary was living or not. Time in hospital was derived from the admission and discharge dates and length of claim

#	Column	Non-Null Count	Dtype
0	BeneID	40474 non-null	object
1	ClaimID	40474 non-null	object
2	ClaimStartDt	40474 non-null	object
3	ClaimEndDt	40474 non-null	object
4	Provider	40474 non-null	object
5	InscClaimAmtReimbursed	40474 non-null	int64
6	AttendingPhysician	40362 non-null	object
7	OperatingPhysician	23830 non-null	object
8	OtherPhysician	4690 non-null	object
9	AdmissionDt	40474 non-null	object
10	ClmAdmitDiagnosisCode	40474 non-null	object
11	DeductibleAmtPaid	39575 non-null	float64
12	DischargeDt	40474 non-null	object
13	DiagnosisGroupCode	40474 non-null	object
14	ClmDiagnosisCode_1	40474 non-null	object
15	ClmDiagnosisCode_2	40248 non-null	object
16	ClmDiagnosisCode_3	39798 non-null	object
17	ClmDiagnosisCode_4	38940 non-null	object
18	ClmDiagnosisCode_5	37580 non-null	object
19	ClmDiagnosisCode_6	35636 non-null	object
20	ClmDiagnosisCode_7	33216 non-null	object
21	ClmDiagnosisCode_8	30532 non-null	object
22	ClmDiagnosisCode_9	26977 non-null	object
23	ClmDiagnosisCode_10	3927 non-null	object
24	ClmProcedureCode_1	23148 non-null	float64
25	ClmProcedureCode_2	5454 non-null	float64
26	ClmProcedureCode_3	965 non-null	float64
27	ClmProcedureCode_4	116 non-null	float64
28	ClmProcedureCode_5	9 non-null	float64
29	ClmProcedureCode_6	0 non-null	float64

(a) Inpatient Data Summary

#	Column	Non-Null Count	Dtype
0	BeneID	138556 non-null	object
1	DOB	138556 non-null	object
2	DOD	1421 non-null	object
3	Gender	138556 non-null	int64
4	Race	138556 non-null	int64
5	RenalDiseaseIndicator	138556 non-null	object
6	State	138556 non-null	int64
7	County	138556 non-null	int64
8	NoOfMonths_PartACov	138556 non-null	int64
9	NoOfMonths_PartBCov	138556 non-null	int64
10	ChronicCond_Alzheimer	138556 non-null	int64
11	ChronicCond_HeartFailure	138556 non-null	int64
12	ChronicCond_KidneyDisease	138556 non-null	int64
13	ChronicCond_Cancer	138556 non-null	int64
14	ChronicCond_ObstrPulmonary	138556 non-null	int64
15	ChronicCond_Depression	138556 non-null	int64
16	ChronicCond_Diabetes	138556 non-null	int64
17	ChronicCond_IschemicHeart	138556 non-null	int64
18	ChronicCond_Osteoporosis	138556 non-null	int64
19	ChronicCond_rheumatoidarthritis	138556 non-null	int64
20	ChronicCond_stroke	138556 non-null	int64
21	IPAnnualReimbursementAmt	138556 non-null	int64
22	IPAnnualDeductibleAmt	138556 non-null	int64
23	OPAnnualReimbursementAmt	138556 non-null	int64
24	OPAnnualDeductibleAmt	138556 non-null	int64

(b) Beneficiary Data Summary

Fig. 1: Raw Data Summaries

was derived from the claim start date and claim end date. The features which were used for derivation were dropped.

The features were then assessed for quality. The percent of values missing, the cardinality, and the presence of outliers were determined for each feature. From large portions of missing values as observed in the feature reports, the claim diagnoses codes 8, 9, and 10 were dropped along with all claim procedure codes excluding the first one. While claim procedure code 1 also had a high percentage of missing values (96%) it was decided to retain this feature's information for the time being. The remaining diagnosis codes also had high missing values percentages and we chose the cut-off for retention conservatively at less than 90% missing values. Again, this was to retain more infor-

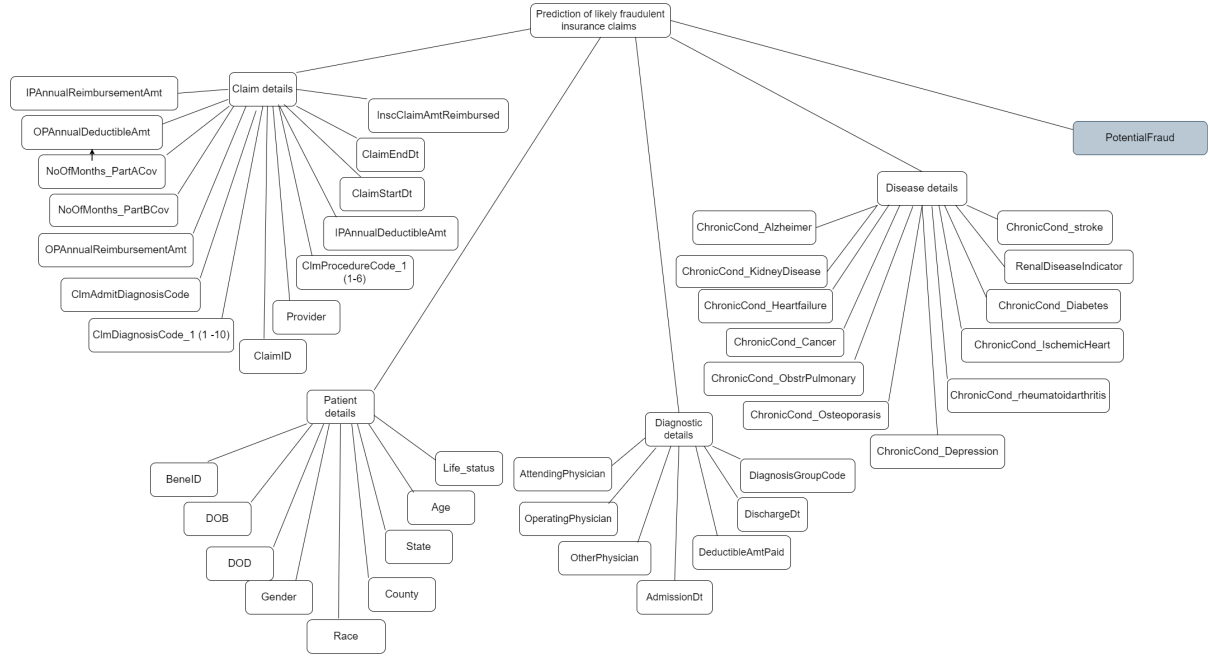


Fig. 2: Domain Concepts

Feature Name	Domain Concept	Feature Description	Feature Type	Data Type
ChronicCond_Alzheimer	Disease Details	Do patient has this particular disease ?	Categorical	Int
ChronicCond_Heartfailure	Disease Details	Do patient has this particular disease ?	Categorical	Int
ChronicCond_KidneyDisease	Disease Details	Do patient has this particular disease ?	Categorical	Int
ChronicCond_Cancer	Disease Details	Do patient has this particular disease ?	Categorical	Int
ChronicCond_ObstPulmonary	Disease Details	Do patient has this particular disease ?	Categorical	Int
ChronicCond_Depression	Disease Details	Do patient has this particular disease ?	Categorical	Int
ChronicCond_Diabetes	Disease Details	Do patient has this particular disease ?	Categorical	Int
ChronicCond_IschemicHeart	Disease Details	Do patient has this particular disease ?	Categorical	Int
ChronicCond_Osteoporosis	Disease Details	Do patient has this particular disease ?	Categorical	Int
ChronicCond_rheumatoidarthritis	Disease Details	Do patient has this particular disease ?	Categorical	Int
ChronicCond_stroke	Disease Details	Do patient has this particular disease ?	Categorical	Int
DOB	Patient Details	Date of birth of the patient	Categorical	datetime
DOD	Patient Details	Date of death of the patient	Categorical	datetime
Gender	Patient Details	Gender of the patient	Categorical	Int
Race	Patient Details	Race of the patient	Categorical	Int
State	Patient Details	State of the patient	Categorical	Int
BenefitID	Patient Details	Beneficiary id of the patient	Continuous	String
Life_status	Patient Details	Is the patient alive?	Categorical	Int
County	Patient Details	Origin of the patient	Categorical	Int
Provider	Claim Details	Insurance provider	Categorical	String
ClaimID	Claim Details	Insurance claimid	Continuous	String
ClaimStartDt	Claim Details	Insurance claim start date	Continuous	datetime
ClaimEndDt	Claim Details	Insurance claim end date	Continuous	datetime
InscClaimAmtReimbursed	Claim Details	Insurance amount reimbursed	Continuous	Int
CimDiagnosisCode_1	Claim Details	Patient claim diagnosis code	Continuous	Int
CimDiagnosisCode_2	Claim Details	Patient claim diagnosis code	Continuous	Int
CimDiagnosisCode_3	Claim Details	Patient claim diagnosis code	Continuous	Int
CimDiagnosisCode_4	Claim Details	Patient claim diagnosis code	Continuous	Int
CimDiagnosisCode_5	Claim Details	Patient claim diagnosis code	Continuous	Int
CimDiagnosisCode_6	Claim Details	Patient claim diagnosis code	Continuous	Int
CimDiagnosisCode_7	Claim Details	Patient claim diagnosis code	Continuous	Int
CimDiagnosisCode_8	Claim Details	Patient claim diagnosis code	Continuous	Int
CimDiagnosisCode_9	Claim Details	Patient claim diagnosis code	Continuous	Int
CimDiagnosisCode_10	Claim Details	Patient claim diagnosis code	Continuous	Int
CimProcedureCode_1	Claim Details	Patient claim procedure code	Continuous	Float
CimProcedureCode_2	Claim Details	Patient claim procedure code	Continuous	Float
CimProcedureCode_3	Claim Details	Patient claim procedure code	Continuous	Float
CimProcedureCode_4	Claim Details	Patient claim procedure code	Continuous	Float
CimProcedureCode_5	Claim Details	Patient claim procedure code	Continuous	Float
CimProcedureCode_6	Claim Details	Patient claim procedure code	Continuous	Float
IPAnnualReimbursementAmt	Claim Details	Insurance annual reimbursement amount	Continuous	Int
OPAnnualDeductibleAmt	Claim Details	Out patient deductible annual amount	Continuous	Int
OPAnnualReimbursementAmt	Claim Details	Out patient reimbursement annual amount	Continuous	Int
AttendingPhysician	Diagnostic Details	Attending physician for the patient	Categorical	String
OperatingPhysician	Diagnostic Details	Operating physician for the patient	Categorical	String
OtherPhysician	Diagnostic Details	Other physician for the patient	Categorical	String
AdmissionDt	Diagnostic Details	Admission date of the patient into the hospital	Continuous	datetime
DischargeDt	Diagnostic Details	Discharge date of the patient into the hospital	Continuous	datetime
DeductibleAmtPaid	Diagnostic Details	Total cost for the procedure	Continuous	Int
DiagnosisGroupCode	Diagnostic Details	Code for the diagnosis provided to the patient	Continuous	Int

Fig. 3: Analytical Base Table

mation at this stage. The claim admission diagnosis code, diagnosis group code, operating physician, and other physicians were also dropped for missing values. Attending physician was also dropped as the cardinality was too high to be useful for a categorical feature (~82,000). No issues with outliers were detected. The

claim and beneficiary IDs were dropped next as they contain identification codes and not useful data.

Finally, the provider code was also dropped. Since the original data set tagged potential fraud based on provider code, but the desired model would predict fraud based on one specific claim, this would potentially cause the model to simply learn which providers are fraudulent over the claims. This would not allow the model to predict on new data. The final feature set is summarized below in Figure 4. The final number of instances is 558211.

C. Feature Selection

The sklearn library tools SelectKBest and mutual_info_classif was utilized to determine the most useful 15 features to use for model training [9], [10]. mutual_info_classif estimates the mutual information to score each feature for dependency concerning the target feature. It functions based on entropy calculations using k-nearest neighbors' distance. SelectKBest selects the top K features based on the scores from mutual_info_classif. The top 15 features and the corresponding mutual information score are shown below in Figure 5.

D. Model Building, Parameter Tuning, and Training

This section describes the models chosen for testing, the justification of choice, and the hyperparameter tuning performed on each before the final training. Section 4 will describe the chosen evaluation plan and the results of training and evaluation will be shown in Section 5. Hyperparameter tuning was performed with 5 cross-fold shuffle split validation with 20% test size

#	Column	Non-Null Count	Dtype
0	LengthofClaim	558211 non-null	int64
1	Provider	558211 non-null	string
2	InscClaimAmtReimbursed	558211 non-null	int64
3	TimeInHosp	558211 non-null	float64
4	DeductibleAmtPaid	558211 non-null	int64
5	ClmDiagnosisCode_1	558211 non-null	int64
6	ClmDiagnosisCode_2	558211 non-null	int64
7	ClmDiagnosisCode_3	558211 non-null	int64
8	ClmDiagnosisCode_4	558211 non-null	int64
9	ClmDiagnosisCode_5	558211 non-null	int64
10	ClmDiagnosisCode_6	558211 non-null	int64
11	ClmDiagnosisCode_7	558211 non-null	int64
12	ClmProcedureCode_1	558211 non-null	int64
13	Gender	558211 non-null	int64
14	Race	558211 non-null	int64
15	RenalDiseaseIndicator	558211 non-null	int64
16	State	558211 non-null	int64
17	County	558211 non-null	int64
18	NoOfMonths_PartACov	558211 non-null	int64
19	NoOfMonths_PartBCov	558211 non-null	int64
20	ChronicCond_Alzheimer	558211 non-null	int64
21	ChronicCond_HeartFailure	558211 non-null	int64
22	ChronicCond_KidneyDisease	558211 non-null	int64
23	ChronicCond_Cancer	558211 non-null	int64
24	ChronicCond_ObstrPulmonary	558211 non-null	int64
25	ChronicCond_Depression	558211 non-null	int64
26	ChronicCond_Diabetes	558211 non-null	int64
27	ChronicCond_IschemicHeart	558211 non-null	int64
28	ChronicCond_Osteoporosis	558211 non-null	int64
29	ChronicCond_rheumatoidarthritis	558211 non-null	int64
30	ChronicCond_stroke	558211 non-null	int64
31	IPAnnualReimbursementAmt	558211 non-null	int64
32	IPAnnualDeductibleAmt	558211 non-null	int64
33	OPAnnualReimbursementAmt	558211 non-null	int64
34	OPAnnualDeductibleAmt	558211 non-null	int64
35	Life_status	558211 non-null	int64
36	Age	558211 non-null	int64
37	PotentialFraud	558211 non-null	bool

Fig. 4: Final Data Set

NoOfMonths_PartBCov	64.998847
NoOfMonths_PartACov	64.869859
State	44.014857
Race	43.493692
ChronicCond_IschemicHeart	42.577699
County	42.417695
Gender	38.611772
ChronicCond_Diabetes	36.021233
ChronicCond_HeartFailure	24.956785
ChronicCond_Depression	13.519686
DeductibleAmtPaid	13.272918
ChronicCond_KidneyDisease	12.551674
IPAnnualDeductibleAmt	12.318085
ChronicCond_Alzheimer	10.114031
InscClaimAmtReimbursed	8.246388

Fig. 5: Feature Selection Results

using sklearn's GridSearchCV [11]. This parameter selection function was used as it performs an exhaustive search for the given parameter ranges for the ideal combination. All other model input parameters which were not tuned were left as the default values. At this stage, randomly sampled 33% of the data was reserved for final testing.

1) *Decision Tree Classifier*: The decision tree classifier was chosen as the baseline model as this is a powerful but simple model to implement, and the logic within the model can be investigated. The implementation of DecisionTreeClassifier from sklearn was used [12]. The parameters max_depth and max_features were tuned with the discussed method. The max_depth parameter was selected as 18 and the max_features selected as 15.

2) *XGBoost Classifier*: Next, XGBClassifier was used from the implementation provided by the XGBoost Python library [13]. This classifier uses boosting on multiple decision tree classifiers. This model was chosen because testing ensemble models was a focus of

this project and boosting can potentially reduce bias in the model results. The max_depth of the model was tuned to be 15.

3) *Bagging Classifier*: The BaggingClassifier implemented in sklearn's ensemble package was used next [14]. A bootstrap aggregating (boosting) model based on decision tree classifiers is a good option as boosting can reduce variance and avoid over fitting our model. The number of estimators, n_estimators, parameter was tuned to be 10.

4) *Stacked Classifier*: Finally, a stacked classifier was tested using StackingClassifier as implemented by sklearn [15]. The stacked model consisted of a Logistic Regression model, K-Neighbours Classifier, Decision Tree Classifier, and Gaussian Naive Bayes, with the final estimator set to an addition Logistic Regression model. These base models are all implemented by sklearn [16]. This model type was chosen as using multiple types of models and combining the results could give more diverse analysis of the data and utilize the strengths of each. Hyperparameter tuning is not applicable for this model type.

E. Evaluation

We employed a variety of evaluation criteria to compare and analyze the output quality from our models, including the following:

1) *Confusion Matrix*: Confusion matrices give you the ability to see the various predictions, errors you might make. By comparing the characteristics of correctly and incorrectly classified data, you can learn more about how to apply machine learning more effectively. The confusion matrix format is shown below in Figure 6 and defines True Positive, False Positive, True Negative, and False Negative values.

		Actual Values	
		Positive (1)	Negative (0)
Predicted Values	Positive (1)	TP	FP
	Negative (0)	FN	TN

Fig. 6: Confusion matrix

2) *Accuracy*: One parameter for assessing classification models is accuracy. The percentage of predictions that our model correctly predicted is known as accuracy.

$$Accuracy = \frac{\text{Number of correct predictions}}{\text{Total number of predictions}}$$

Accuracy can also be determined in terms of positives and negatives for binary classification, as seen below:

$$Accuracy = \frac{TP + TN}{TP + TN + FP + FN}$$

3) *Precision*: Precision speaks about a model's ability to make a correct estimate. When calculating precision, one divides the total number of positive predictions by the proportion of actual positives.

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

4) *Recall*: The recall is determined as the proportion of Positive samples that were correctly identified as Positive to all Positive samples. The recall measures how well the model can identify positive samples. The more positive samples that are identified, the higher the recall.

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

5) *F1 score*: Instead of evaluating a model's overall performance like accuracy does, F1score focuses on how well it performs in each class to determine how predictive it is. The precision and recall scores of a model are combined into one statistic called the F1 score.

$$F1\ Score = 2 \times \frac{Precision \times Recall}{Precision + Recall}$$

6) *Learning Curve*: An improvement in a process over time as a result of learning and growing skill is represented visually by a learning curve. According to the learning curve idea, tasks will demand less time and resources as they are carried out more frequently due to the proficiency that is acquired when the procedure is learned. Training was performed with 5-fold cross validation.

IV. EXPERIMENTS AND RESULTS

The evaluation was performed on the 33% reserved data. Here we have the confusion matrix, and classification report with accuracy, precision, recall, and f1-score. We also have the learning curve, scalability, and performance of each model. The confusion matrix is giving the number of TP, TN, FP and FN for each model. The classification report gives the values of precision, recall, f1-score, support, and accuracy. Overall, our models have shown acceptable accuracy. Although all the models have shown almost similar results, the highest recorded was for XGBoost at 78% and the lowest for Stacked Classifier at 72%. Close to XGBoost, we have the Decision Tree classification where the accuracy has been recorded at 76%.

From our learning curves of all our models, as the training set increases, the training score curve, and the cross-validation curve are about to converge at some point. In the Decision tree classification and the XGBoost classification, as the instances are increasing,

there is a significant change in the error which explains that the model has a proper learning rate.

The confusion matrix for each model are shown below in Figure 8. The classification report of each model are shown below in Figure 7. The learning curves for each classifier are shown in Figures 9 - 12. The means of the 5-fold cross validation are plotted with the standard deviation shaded.

Classification Report for Decision Tree			
Provider Class	Precision	Recall	f1-Score
Non-Fraudulent	0.79	0.84	0.81
Fraudulent	0.7	0.63	0.67
Accuracy	76%		

(a) Decision Tree

Classification Report for XGBoost			
Provider Class	Precision	Recall	f1-Score
Non-Fraudulent	0.8	0.86	0.83
Fraudulent	0.73	0.65	0.69
Accuracy	78%		

(b) XGBoost Classifier

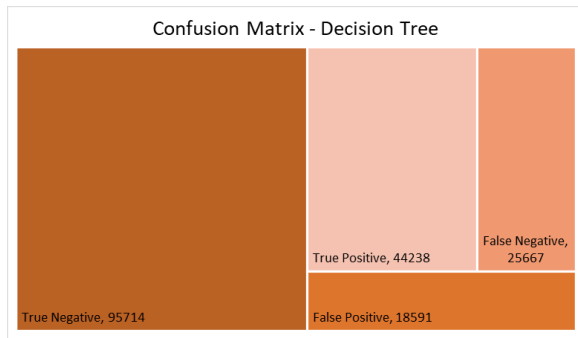
Classification Report for Bagging Classifier			
Provider Class	Precision	Recall	f1-Score
Non-Fraudulent	0.77	0.81	0.79
Fraudulent	0.66	0.61	0.63
Accuracy	73%		

(c) Bagging Classifier

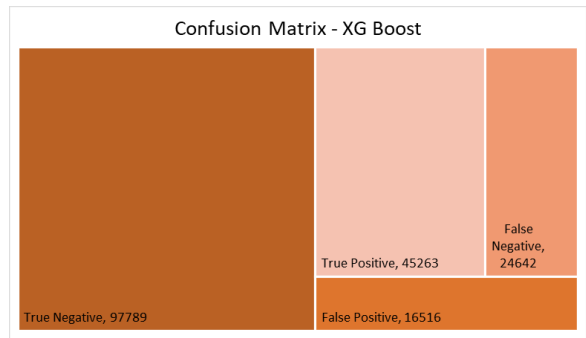
Classification Report for Stacked Classifier			
Provider Class	Precision	Recall	f1-Score
Non-Fraudulent	0.75	0.82	0.78
Fraudulent	0.65	0.55	0.6
Accuracy	72%		

(d) Stacked Classifier

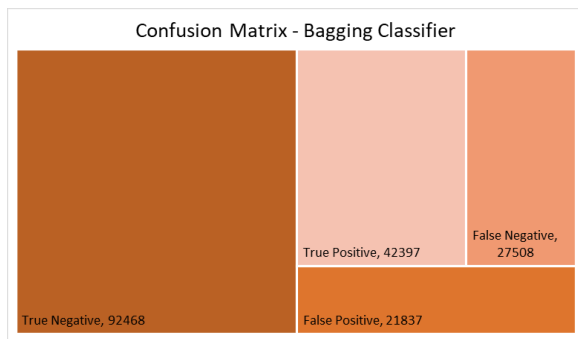
Fig. 7: Classification Reports



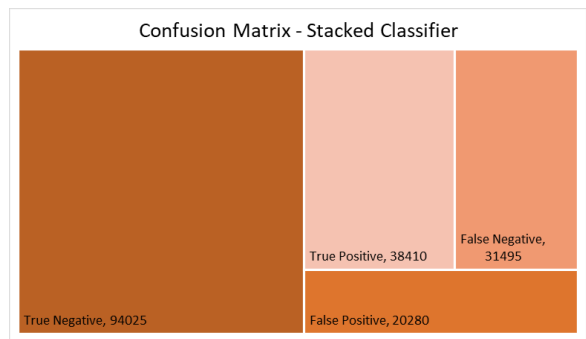
(a) Decision Tree



(b) XG Boost Classifier



(c) Bagging Classifier



(d) Stacked Classifier

Fig. 8: Confusion Matrices

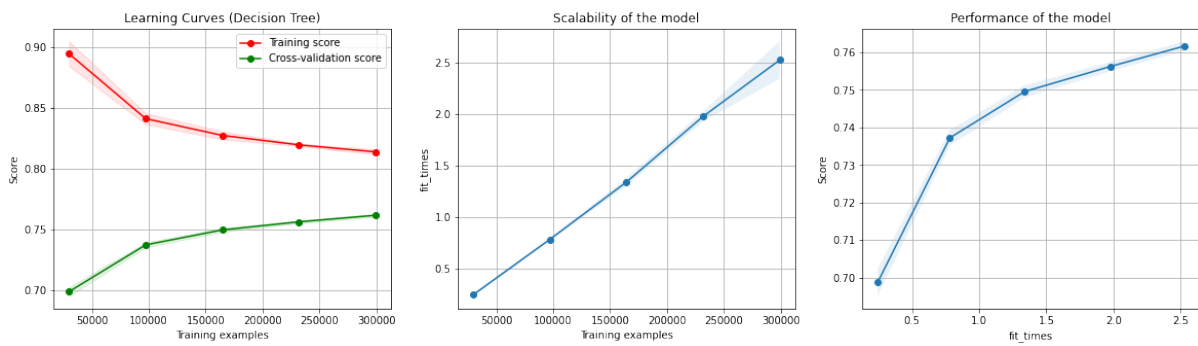


Fig. 9: Decision Tree Classifier Learning Curves

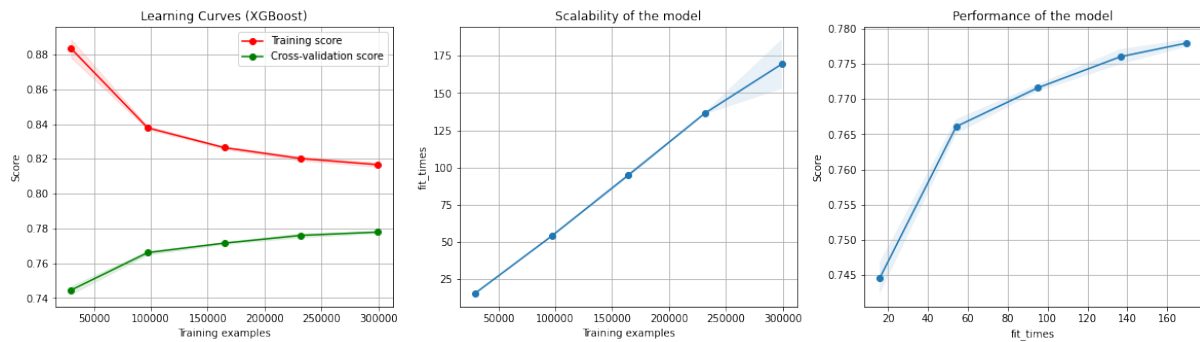


Fig. 10: XGBoost Classifier Learning Curves

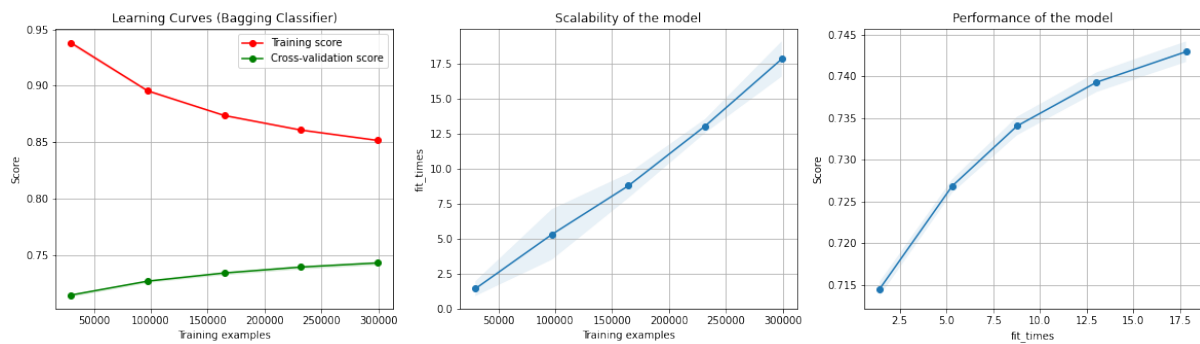


Fig. 11: Bagging Classifier Learning Curves

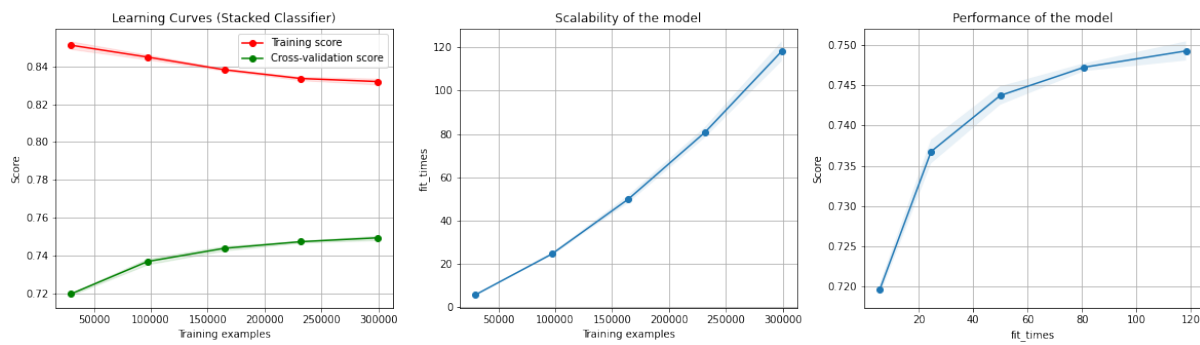


Fig. 12: Stacked Classifier Learning Curves

Our results show that fraudulent claims can predicted with moderate accuracy. While improvements to accuracy would likely be needed before final deployment, the model output solves the business problem. In deployment, the submitted claim details would be automatically input to the fraud detection model and the output potential fraud prediction would flag suspicious claims for further review by the insurance company.

V. CONCLUSION

From a vast data set of both potentially fraudulent and non-fraudulent claims, we identified the potentially fraudulent claims in this paper, employing the Decision Tree Classifier, XGBoost Classifier, Bagging Classifier, and Stacked Classifier supervised methods. In order to approach our solution, we used the CRISP-DM paradigm, beginning with business understanding, where we attempted to comprehend fraudulent claims and identify the problem statement. We derived new features and identified key features that are useful in spotting possibly fraudulent providers' actions. Our experiments revealed that each algorithm can produce predictions with a good degree of accuracy and usability. Despite the similar accuracies of all models, the stacking classifier model under performed than other models, with a performance of 72% accuracy. According to our investigation, XGBoost was the best-performing model, with an accuracy of 78%, while Decision Tree Classifier and Bagging Classifier were at 76% and 73% respectively.

In future work, we aim to investigate how well unsupervised models perform with the data we have. We would also analyze the K-nearest neighbor algorithm in combination with a genetic algorithm in order to find the ideal weighting of the features. These insights could assist with increasing performance of the prediction models. Gathering more data for training would also be attempted to improve model results.

REFERENCES

- [1] T. L. Leap, "The major health care fraud laws," in *Phantom Billing, Fake Prescriptions, and the High Cost of Medicine: Health Care Fraud and What to Do about It*, T. L. Leap, Ed. Cornell University Press, p. 0. [Online]. Available: <https://doi.org/10.7591/cornell/9780801449796.003.0002>
- [2] <https://revcycleintelligence.com/features/how-providers-can-detect-prevent-healthcare-fraud-and-abuse>, accessed: 2022-12-20.
- [3] G. Brooks, M. Button, and J. Gee, "The scale of health-care fraud: A global evaluation," vol. 25, no. 1, pp. 76–87. [Online]. Available: <https://doi.org/10.1057/sj.2011.7>
- [4] Y. Peng, G. Kou, A. Sabatka, Z. Chen, D. Khazanchi, and Y. Shi, "Application of clustering methods to health insurance fraud detection," in *2006 International Conference on Service Systems and Service Management*, vol. 1, pp. 116–120, ISSN: 2161-1904.
- [5] T. Ekin, "Application of bayesian methods in detection of healthcare fraud," vol. 33.
- [6] R. Bauder, T. M. Khoshgoftaar, and N. Seliya, "A survey on the state of healthcare upcoding fraud analysis and detection," vol. 17, no. 1, pp. 31–55. [Online]. Available: <https://doi.org/10.1007/s10742-016-0154-8>
- [7] R. Anand Gupta. Medical provider fraud detection. [Online]. Available: <https://kaggle.com/code/rohitrox/medical-provider-fraud-detection>
- [8] ——. Healthcare provider fraud detection analysis. [Online]. Available: <https://www.kaggle.com/datasets/rohitrox/healthcare-provider-fraud-detection-analysis>
- [9] sklearn.feature_selection.SelectKBest. [Online]. Available: https://scikit-learn/stable/modules/generated/sklearn.feature_selection.SelectKBest.html
- [10] sklearn.feature_selection.mutual_info_classif. [Online]. Available: https://scikit-learn/stable/modules/generated/sklearn.feature_selection.mutual_info_classif.html
- [11] sklearn.model_selection.GridSearchCV. [Online]. Available: https://scikit-learn/stable/modules/generated/sklearn.model_selection.GridSearchCV.html
- [12] sklearn.tree.DecisionTreeClassifier. [Online]. Available: <https://scikit-learn/stable/modules/generated/sklearn.tree.DecisionTreeClassifier.html>
- [13] Python API reference — xgboost 1.7.2 documentation. [Online]. Available: https://xgboost.readthedocs.io/en/stable/python/python_api.html
- [14] sklearn.ensemble.BaggingClassifier. [Online]. Available: <https://scikit-learn/stable/modules/generated/sklearn.ensemble.BaggingClassifier.html>
- [15] sklearn.ensemble.StackingClassifier. [Online]. Available: <https://scikit-learn/stable/modules/generated/sklearn.ensemble.StackingClassifier.html>
- [16] scikit-learn: machine learning in python — scikit-learn 1.2.0 documentation. [Online]. Available: <https://scikit-learn.org/stable/>

APPENDIX A: CATEGORICAL FEATURE REPORT

	Count	Miss %	Card.		Mode	Mode Freq	Mode %		2nd Mode	2nd Mode Freq	2nd Mode %
BenetID	558211	0.000000	138556	[BENE118316, BENE42721, BENE59303]	87	0.015586	[BENE36330, BENE44241, BENE80977]	84	0.015048		
ClaimID	558211	0.000000	558211	[CLM110011, CLM110012, CLM110013, CLM110014, C...	558211	100.000000				0	0.000000
ClaimStartDt	558211	0.000000	398	[2009-01-31]	1709	0.306157		[2009-03-03]	1706	0.305619	
ClaimEndDt	558211	0.000000	366	[2009-03-03]	1707	0.305798		[2009-02-11]	1682	0.301320	
Provider	558211	0.000000	5410	[PRV51459]	8240	1.476144		[PRV53797]	4739	0.848962	
AttendingPhysician	558211	0.270149	82063	[PHY330576]	2534	0.453950		[PHY350277]	1628	0.291646	
OperatingPhysician	558211	79.497538	35315	[PHY330576]	424	0.075957		[PHY424897]	293	0.052489	
OtherPhysician	558211	64.218548	46457	[PHY412132]	1247	0.223392		[PHY341578]	1098	0.196700	
AdmissionDt	558211	92.749337	398	[2009-02-10]	144	0.025797		[2009-01-31, 2009-02-26]	286	0.051235	
ClinAdmitDiagnosisCode	558211	73.863109	4098	[V7612]	4074	0.729832		[42731]	3634	0.651008	
DischargeDt	558211	92.749337	365	[2009-02-11]	153	0.027409		[2009-01-10]	147	0.026334	
DiagnosisGroupCode	558211	92.749337	736	[882]	179	0.032067		[884]	174	0.031171	
ClinDiagnosisCode_1	558211	1.872589	10450	[4019]	13886	2.487590		[4011]	12512	2.241446	
ClinDiagnosisCode_2	558211	35.041588	5300	[4019]	22378	4.008878		[25000]	11744	2.103864	
ClinDiagnosisCode_3	558211	56.458221	4756	[4019]	14408	2.581103		[25000]	7946	1.423476	
ClinDiagnosisCode_4	558211	70.524407	4359	[4019]	9188	1.645973		[25000]	5250	0.940505	
ClinDiagnosisCode_5	558211	79.949517	3970	[4019]	6005	1.075758		[25000]	3451	0.618225	
ClinDiagnosisCode_6	558211	84.881702	3607	[4019]	4170	0.747029		[25000]	2506	0.448934	
ClinDiagnosisCode_7	558211	88.144805	3388	[4019]	3014	0.539939		[25000]	1822	0.326400	
ClinDiagnosisCode_8	558211	90.425843	3070	[4019]	2257	0.404327		[25000]	1399	0.250622	
ClinDiagnosisCode_9	558211	92.509105	2774	[4019]	1581	0.283226		[25000]	1100	0.197058	
ClinDiagnosisCode_10	558211	99.102490	1158	[4019]	169	0.030275		[25000]	125	0.022393	
DOB	558211	0.000000	900	[1943-12-01 00:00:00]	2072	0.371186		[1939-03-01 00:00:00]	2030	0.363662	
DOD	558211	99.259957	11	[2009-12-01 00:00:00]	710	0.127192		[2009-10-01 00:00:00]	572	0.102470	
RenalDiseaseIndicator	558211	0.000000	2	[0]	448363	80.321420		[1]	109848	19.678580	
PotentialFraud	558211	0.000000	2	[No]	345415	61.878931		[Yes]	212796	38.121069	

APPENDIX B: CONTINUOUS FEATURE REPORT

	Count	Miss %	Card.	Min	1st Qrt.	Mean	Median	3rd Qrt	Max	Std. Dev.
InscClaimAmtReimbursed	558211	0.000000	438	0	40.0	997.012133	80.0	300.0	125000	3821.534891
DeductibleAmtPaid	558211	0.161050	17	0	0.0	78.421085	0.0	0.0	1068	274.016812
ClmProcedureCode_1	558211	95.824160	1117	11.0	3848.0	5896.154612	5363.0	8669.0	9999.0	3050.489933
ClmProcedureCode_2	558211	99.016501	300	42.0	2724.0	4106.358106	4019.0	4439.0	9999.0	2031.640878
ClmProcedureCode_3	558211	99.826410	154	42.0	2724.0	4221.123839	4019.0	5185.0	9999.0	2281.849885
ClmProcedureCode_4	558211	99.978861	48	42.0	2754.25	4070.262712	4019.0	4439.0	9986.0	2037.62699
ClmProcedureCode_5	558211	99.998388	6	2724	4139.0	5269.444444	4139.0	5185.0	9982	2780.071632
ClmProcedureCode_6	558211	100.000000	0	<NA>	<NA>	<NA>	<NA>	<NA>	<NA>	<NA>
Gender	558211	0.000000	2	1	1.0	1.578838	2.0	2.0	2	0.493746
Race	558211	0.000000	4	1	1.0	1.255011	1.0	1.0	5	0.717437
State	558211	0.000000	52	1	11.0	25.446969	24.0	38.0	54	15.192784
County	558211	0.000000	314	0	150.0	378.588195	350.0	570.0	999	265.215531
NoOfMonths_PartACov	558211	0.000000	13	0	12.0	11.931472	12.0	12.0	12	0.889712
NoOfMonths_PartBCov	558211	0.000000	13	0	12.0	11.93877	12.0	12.0	12	0.7859
ChronicCond_Alzheimer	558211	0.000000	2	0	0.0	0.401868	0.0	1.0	1	0.490276
ChronicCond_Heartfailure	558211	0.000000	2	0	0.0	0.590427	1.0	1.0	1	0.491755
ChronicCond_KidneyDisease	558211	0.000000	2	0	0.0	0.412002	0.0	1.0	1	0.492196
ChronicCond_Cancer	558211	0.000000	2	0	0.0	0.151385	0.0	0.0	1	0.358424
ChronicCond_ObstrPulmonary	558211	0.000000	2	0	0.0	0.31293	0.0	1.0	1	0.463687
ChronicCond_Depression	558211	0.000000	2	0	0.0	0.434807	0.0	1.0	1	0.495732
ChronicCond_Diabetes	558211	0.000000	2	0	0.0	0.705395	1.0	1.0	1	0.455866
ChronicCond_IschemicHeart	558211	0.000000	2	0	1.0	0.759265	1.0	1.0	1	0.42753
ChronicCond_Osteoporosis	558211	0.000000	2	0	0.0	0.317647	0.0	1.0	1	0.465562
ChronicCond_rheumatoidarthritis	558211	0.000000	2	0	0.0	0.311171	0.0	1.0	1	0.462973
ChronicCond_stroke	558211	0.000000	2	0	0.0	0.10172	0.0	0.0	1	0.302279
IPAnnualReimbursementAmt	558211	0.000000	3004	-8000	0.0	5227.971466	0.0	6000.0	161470	11786.274732
IPAnnualDeductibleAmt	558211	0.000000	147	0	0.0	568.756807	0.0	1068.0	38272	1179.172616
OPAnnualReimbursementAmt	558211	0.000000	2078	-70	460.0	2278.225348	1170.0	2590.0	102960	3881.846386
OPAnnualDeductibleAmt	558211	0.000000	789	0	120.0	649.698745	340.0	790.0	13840	1002.020811
Life_status	558211	0.000000	2	0	0.0	0.0074	0.0	0.0	1	0.085707
Age	558211	0.000000	76	26	68.0	73.852368	75.0	83.0	101	13.020485