



Predicting Hospital Readmissions

Data606 phase 3

Team C:

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Files

main

Go to file

PHASE 0 CREATING TEAMS.txt

PHASE 1 PROJECT PITCH.txt

PHASE 2 EDA & Model Construct...

PHASE 3 Execution and Interpret...

Capstone-Project / PHASE 2 EDA & Model Construction.txt



Brett-Duvall Update PHASE 2 EDA & Model Construction.txt

35d9bec · last month

History

Code

Blame

Raw



```
1 In the second part, groups will
2 • complete their data exploration stage (the dataset should be completely ready after appropriate cleansing and trans-
3   all the members are expected to be familiar with all the major patterns and trends in dataset)
4 • construct their model (i.e. if it is a regression problem, then groups should have their codes ready that are compa-
5   if it is a neural network implementation, then students should complete at least one successful training. etc., A1C, gluco-
6   (Especially, provide the details on how you do the splitting of your data into training, validation, and test sets. etc., A1C, gluco-
7   folds (e.g. 40/30/30? 10-fold? leave-one-out/LOO?).)
8 • Again, each group will make a presentation (P2) to their classmates about their findings.
```

- Size: 101,766 encounters, 50 variables, 18 MB.
- Categories: demographics, diagnoses, labs (A1C, glucose), medication use, utilization.
- Target: 30 day readmission.

TABLE 1: List of features and their descriptions in the initial dataset (the dataset is also available at the website of Data Mining and Biomedical Informatics Lab at VCU (<http://www.cioslab.vcu.edu/>)).

| Feature name | Type | Description and values | % missing |
|-----------------------------|---------|--|-----------|
| Encounter ID | Numeric | Unique identifier of an encounter | 0% |
| Patient number | Numeric | Unique identifier of a patient | 0% |
| Race | Nominal | Values: Caucasian, Asian, African American, Hispanic, and other | 2% |
| Gender | Nominal | Values: male, female, and unknown/invalid | 0% |
| Age | Nominal | Grouped in 10-year intervals: [0, 10), [10, 20), ..., [90, 100) | 0% |
| Weight | Numeric | Weight in pounds. | 97% |
| Admission type | Nominal | Integer identifier corresponding to 9 distinct values, for example, emergency, urgent, elective, newborn, and not available | 0% |
| Discharge disposition | Nominal | Integer identifier corresponding to 29 distinct values, for example, discharged to home, expired, and not available | 0% |
| Admission source | Nominal | Integer identifier corresponding to 21 distinct values, for example, physician referral, emergency room, and transfer from a hospital | 0% |
| Time in hospital | Numeric | Integer number of days between admission and discharge | 0% |
| Payer code | Nominal | Integer identifier corresponding to 23 distinct values, for example, Blue Cross\Blue Shield, Medicare, and self-pay | 52% |
| Medical specialty | Nominal | Integer identifier of a specialty of the admitting physician, corresponding to 84 distinct values, for example, cardiology, internal medicine, family/general practice, and surgeon | 53% |
| Number of lab procedures | Numeric | Number of lab tests performed during the encounter | 0% |
| Number of procedures | Numeric | Number of procedures (other than lab tests) performed during the encounter | 0% |
| Number of medications | Numeric | Number of distinct generic names administered during the encounter | 0% |
| Number of outpatient visits | Numeric | Number of outpatient visits of the patient in the year preceding the encounter | 0% |
| Number of emergency visits | Numeric | Number of emergency visits of the patient in the year preceding the encounter | 0% |
| Number of inpatient visits | Numeric | Number of inpatient visits of the patient in the year preceding the encounter | 0% |
| Diagnosis 1 | Nominal | The primary diagnosis (coded as first three digits of ICD9); 848 distinct values | 0% |
| Diagnosis 2 | Nominal | Secondary diagnosis (coded as first three digits of ICD9); 923 distinct values | 0% |
| Diagnosis 3 | Nominal | Additional secondary diagnosis (coded as first three digits of ICD9); 954 distinct values | 1% |
| Number of diagnoses | Numeric | Number of diagnoses entered to the system | 0% |
| Glucose serum test result | Nominal | Indicates the range of the result or if the test was not taken. Values: ">200," ">300," "normal," and "none" if not measured | 0% |
| Alc test result | Nominal | Indicates the range of the result or if the test was not taken. Values: ">8" if the result was greater than 8%, ">7" if the result was greater than 7% but less than 8%, "normal" if the result was less than 7%, and "none" if not measured. | 0% |
| Change of medications | Nominal | Indicates if there was a change in diabetic medications (either dosage or generic name). Values: "change" and "no change" | 0% |
| Diabetes medications | Nominal | Indicates if there was any diabetic medication prescribed. Values: "yes" and "no" | 0% |
| 24 features for medications | Nominal | For the generic names: metformin, repaglinide, nateglinide, chlorpropamide, glimepiride, acetohexamide, glipizide, glyburide, tolbutamide, pioglitazone, rosiglitazone, acarbose, miglitol, troglitazone, tolazamide, examide, sitagliptin, insulin, glyburide-metformin, glipizide-metformin, glimepiride-pioglitazone, metformin-rosiglitazone, and metformin-pioglitazone, the feature indicates whether the drug was prescribed or there was a change in the dosage. Values: "up" if the dosage was increased during the encounter, "down" if the dosage was decreased, "steady" if the dosage did not change, and "no" if the drug was not prescribed | 0% |
| Readmitted | Nominal | Days to inpatient readmission. Values: "<30" if the patient was readmitted in less than 30 days, ">30" if the patient was readmitted in more than 30 days, and "No" for no record of readmission. | 0% |

Machine Learning – Classification

Target ('readmitted') has 3 categories:

- <30 (the focus of our classification)
- >30
- NO

Features kept even though missing values:

- A1C test result
- Glucose serum test result

Unnecessary features (administrative or missing values):

- Encounter ID
- patient number
- weight
- admission type
- discharge disposition
- admission source
- payer code
- medical specialty (of admitting physician)

Dataset:

<https://drive.google.com/file/d/1fMbjB-5I0Suifh4QrpH1ewtXvkAtayd/view?usp=sharing>

Literature Review

- **Ashfaq et al. (2019)** implemented deep learning models such as convolutional and recurrent neural networks for hospital readmission prediction using electronic health records. Their study demonstrated improved recall and AUC over traditional models, motivating our comparison between ensemble and neural approaches for clinical prediction tasks.
- **Emi-Johnson (2025)** showed the potential of ML algorithms, particularly XGBoost, in enhancing the prediction of 30-day hospital readmissions using structured EHR. By comparing four different models, they demonstrated that ensemble-based methods such as XGBoost not only provide superior predictive performance but also robust interpretability via SHAP analysis.
- **Shukla and Tripathi (2020)** proposed an embedding-based model (EmbPred30) for predicting 30-day readmissions for diabetic patients using the same UCI dataset. Their results showed that categorical embeddings and gradient boosting improved model accuracy compared with traditional one-hot encoding. This supports our focus on feature engineering to enhance model performance.
- **Strack et al. (2014)** analyzed 70,000 clinical records in the UCI Diabetes Readmission dataset and found that conducting HbA1c testing during hospitalization was strongly associated with reduced 30-day readmission rates. Their study introduced the dataset that serves as the foundation for many subsequent projects, including ours.
- **Wang and Zhu (2021)** reviewed challenges and solutions in predictive modeling of hospital readmissions using machine learning. They compared several algorithms, including Random Forest, Support Vector Machines, and Logistic Regression, reporting AUC values typically between 0.62 and 0.70. Their work emphasizes issues such as class imbalance and preprocessing, which our project addresses using class weighting and consistent encoding.

Literature Review Metrics

Ashfaq et al. (2019)

- AUC 0.73

Emi-Johnson (2025)

- AUC 0.58 - 0.67

Shukla and Tripathi (2020)

- AUC 0.71

Strack et al. (2014)

- AUC 0.61 – 0.63

Wang and Zhu (2021)

- AUC 0.62 – 0.70

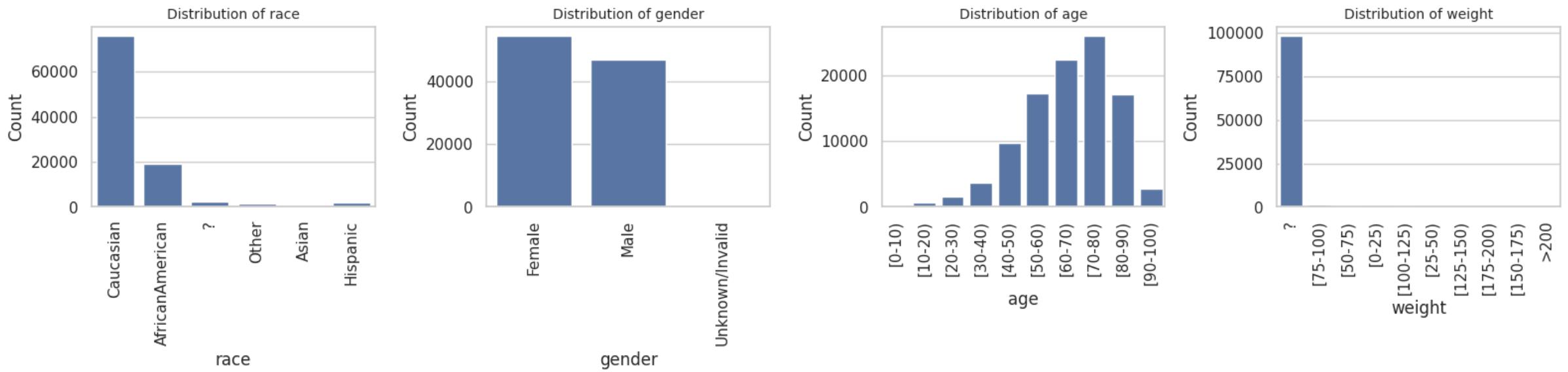
Additional Literature Review

- Combining Patient Visits –
Burrill (2025)
 - Adding created features
- A1C Test Reasoning -
Strack et al. (2014)
 - 3 main admittance values
 - Diabetes
 - Circulatory
 - Respiratory

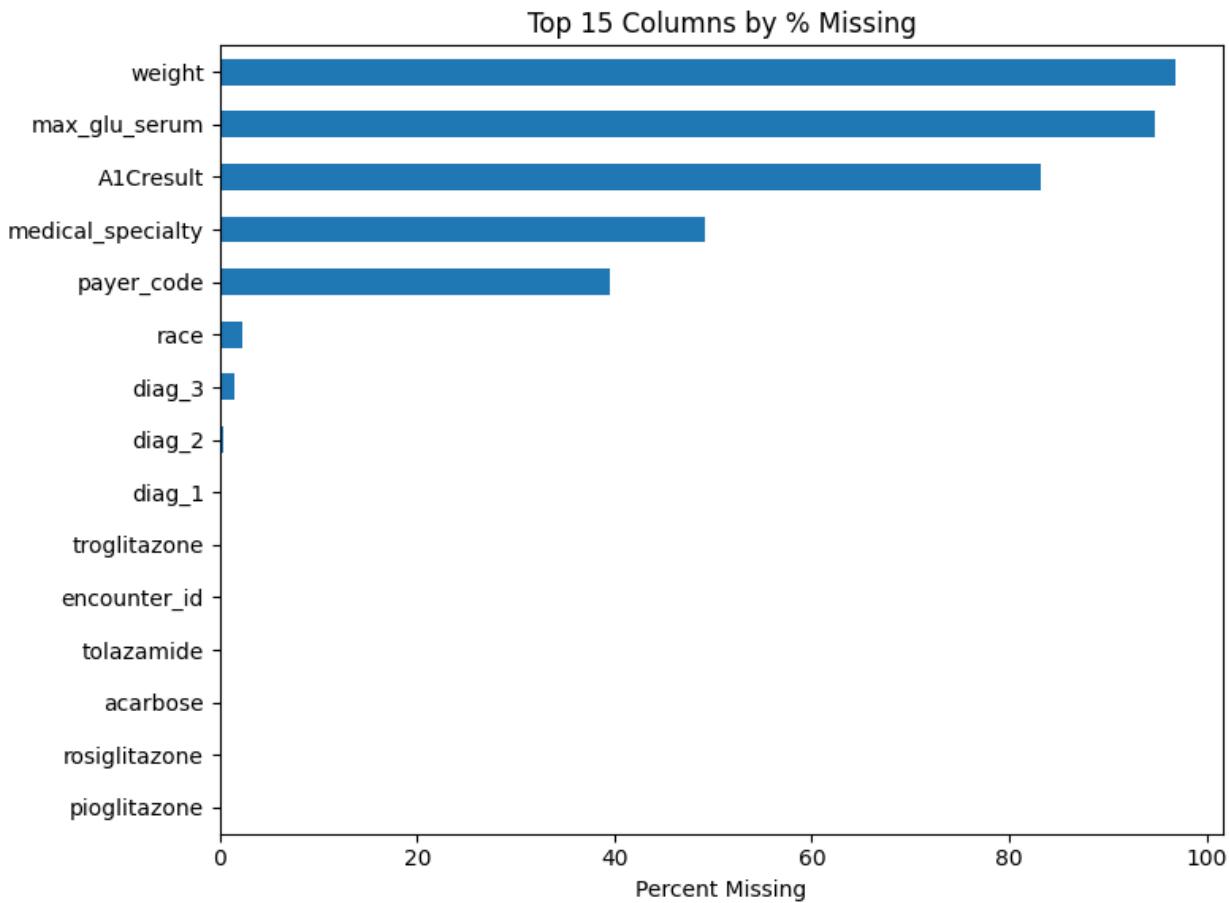
Data Exploration



Data Distribution



Missing values



- Serum glucose is a test that measures the amount of glucose (sugar) in a patient's blood.
- A1C test measures the average amount of sugar in your blood over the past few months

| | Missing Values | Percentage |
|---------------|----------------|------------|
| max_glu_serum | 96420 | 94.746772 |
| A1Cresult | 84748 | 83.277322 |

Dealing with missing values

Why keeping two of the highest features with missing values?

```
# Fill 'missing' for selected categoricals with moderate-to-high missingness
categorical_missing_cols = ["max_glu_serum", "A1Cresult", "medical_specialty", "payer_code", "race"]
for c in categorical_missing_cols:
    if c in X.columns:
        X[c] = X[c].fillna("missing")

# Quick check: confirm no remaining NaNs in categorical columns
print("Remaining NaNs in categorical features:")
print(X[categorical_missing_cols].isna().sum())

# Define numeric and categorical feature sets
numeric_cols = X.select_dtypes(include=["number"]).columns.tolist()
categorical_cols = [c for c in X.columns if c not in numeric_cols]

# Define preprocessing pipeline
from sklearn.compose import ColumnTransformer
from sklearn.preprocessing import StandardScaler, OneHotEncoder

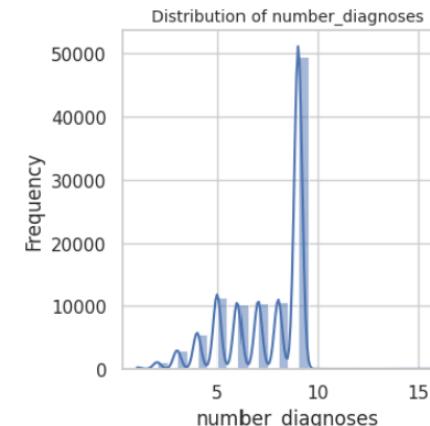
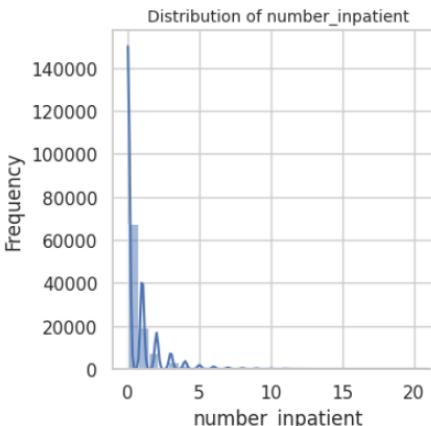
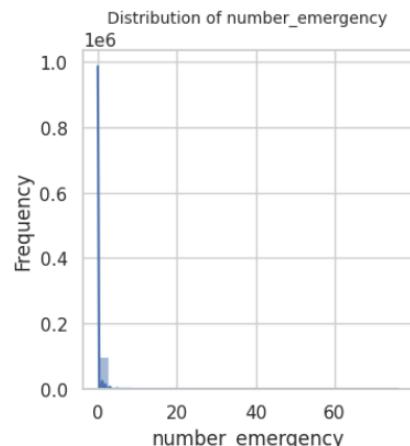
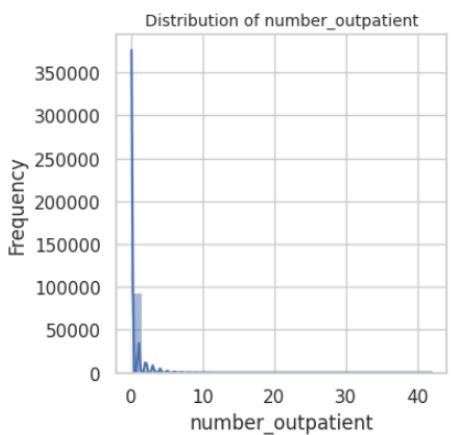
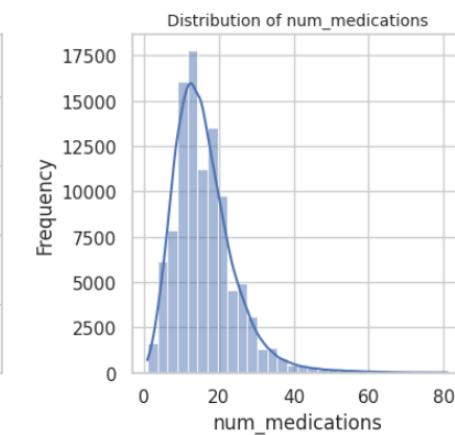
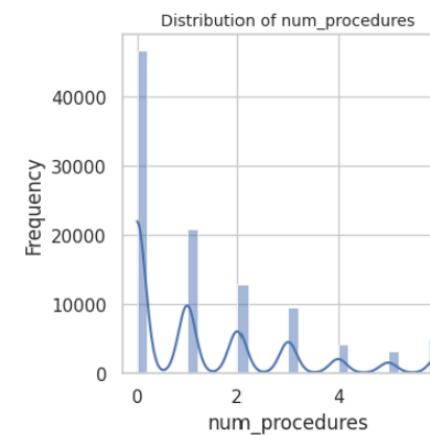
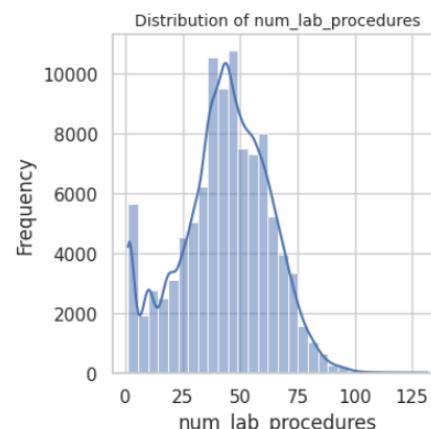
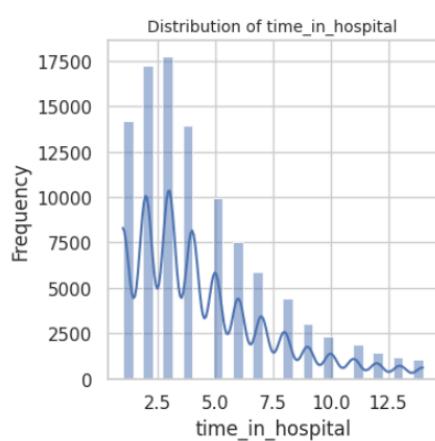
preprocess = ColumnTransformer(
    transformers=[
        ("num", StandardScaler(with_mean=False), numeric_cols),
        ("cat", OneHotEncoder(handle_unknown='ignore', min_frequency=50), categorical_cols)
    ],
    remainder="drop", sparse_threshold=0.3
)

print(f"Preprocessor ready with {len(numeric_cols)} numeric and {len(categorical_cols)} categorical features")
```

Remaining NaNs in categorical features:

| | |
|-------------------|---|
| max_glu_serum | 0 |
| A1Cresult | 0 |
| medical_specialty | 0 |
| payer_code | 0 |
| race | 0 |

Distribution of Numeric Features



Correlation Among Numeric Features

- Moderate positive correlations between longer hospital stays and more medications, also medications and procedures(other than lab tests).
- Most variables have weak relationships.

Correlation Heatmap - Numeric Features



Model Construction



Model Construction and Validation

Goal: Predict 30-day hospital readmission among diabetic patients using the UCI Diabetes Readmission dataset (101,766 records, 8 numeric, 38 categorical features).

Task Type: Binary classification

Target variable: *readmitted <30 days (1) vs ≥30 days or no readmission (0)*

Imbalance: ~11% positive (readmitted within 30 days)

Data Split:

- Training: 70%
- Validation: 10%
- Testing: 20%

Cross-Validation:

10-fold Stratified CV to preserve class proportions

Evaluation Metrics:

- ROC-AUC (overall ranking ability)
- PR-AUC (performance on rare positives)
- Precision@Top-10% (clinical targeting usefulness)
- F1 score (balance of precision and recall)

Model Portfolio

Model Comparison

- Logistic Regression – interpretable linear baseline.
- Random Forest – ensemble of decision trees capturing non-linear patterns.
- Gradient Boosting – sequential tree model emphasizing hard cases.
- XGBoost – optimized gradient boosting; best for structured data.
- SVM – margin-based classifier for comparison.

Design Choices:

- Missing values handled; categorical variables encoded; numeric features scaled.
- Class imbalance addressed with `class_weight='balanced'`.
- Consistent preprocessing pipeline applied to all models.

Model Performance (Base vs Cross-Validation)

Base Model Performance

| Model | ROC-AUC | PR-AUC | Precision@10 % |
|---------------------|---------|--------|----------------|
| Logistic Regression | 0.678 | 0.226 | 0.260 |
| Random Forest | 0.679 | 0.234 | 0.277 |
| Gradient Boosting | 0.687 | 0.241 | 0.285 |
| XGBoost | 0.688 | 0.241 | 0.280 |

Cross-Validation (10-Fold Mean ± SD)

| Model | ROC-AUC | PR-AUC |
|---------------------|-------------------|-------------------|
| Logistic Regression | 0.669 ± 0.007 | 0.218 ± 0.009 |
| Random Forest | 0.676 ± 0.007 | 0.230 ± 0.007 |
| Gradient Boosting | 0.680 ± 0.005 | 0.232 ± 0.009 |
| XGBoost | 0.681 ± 0.005 | 0.231 ± 0.008 |

Choosing the Right Metric for Hospital Readmission

Why Accuracy Is Misleading

Only 11 percent of patients are readmitted, so a model can appear “accurate” just by predicting “no readmission.”

ROC-AUC (Overall Ranking Ability)

Shows how well the model separates readmitted from non-readmitted patients across all thresholds.

Good global indicator but less sensitive to class imbalance.

PR-AUC (Focus on Rare Positives)

Captures the trade-off between precision and recall for the *readmitted* class.

Best single metric for imbalanced clinical data—tells how well the model identifies true readmissions among many non-readmissions.

Precision @ Top 10 Percent (Actionability)

Measures how many of the top 10 percent highest-risk patients truly return within 30 days.

Directly reflects how well hospitals can target limited follow-up resources.

Key Insights

- Baseline PR-AUC ≈ 0.11 (random) → our best models ≈ 0.24 ($>2\times$ improvement).
- Precision @ Top 10 % ≈ 0.28 → roughly 28 of 100 flagged patients actually readmit.
- **XGBoost** shows the best overall ranking (ROC-AUC ≈ 0.69).
- **Gradient Boosting** achieves slightly higher precision in the actionable top 10 percent.

Takeaway

Focus on PR-AUC and Precision @ Top 10 % for evaluation—these best represent clinical usefulness in predicting rare readmissions.

Benchmarking Against Prior Studies

How Our Results Compare

- *Strack et al., 2014 (UCI baseline)* – AUC 0.61 – 0.63 → our XGBoost 0.69 = major improvement over the original logistic model.
- *Wang & Zhu, 2021* – AUC 0.62 – 0.70 → our ensemble 0.68 – 0.69 = comparable to modern ML benchmarks.
- *Shukla & Tripathi, 2020* – AUC 0.71 (embedding model) → our XGBoost 0.69 = slightly lower but close.
- *Ashfaq et al., 2019 (deep learning)* – AUC 0.73 → our XGBoost 0.69 = near deep-learning performance with simpler, interpretable methods.

Interpretation

Our ensemble models (XGBoost, Gradient Boosting) perform on par with or better than most published traditional ML approaches.

They achieve strong, stable AUC and PR-AUC scores while remaining easier to interpret and deploy.

Key Takeaways

- **Best Model:** XGBoost (ROC-AUC ≈ 0.69, PR-AUC ≈ 0.24)
- **Clinical Targeting:** Gradient Boosting slightly higher precision @ 10 %.
- **Consistency:** Results align with the 0.62–0.73 AUC range reported across the literature.

Next Step: Apply SHAP analysis to explain which factors (insulin use, HbA1c level, utilization) drive readmission risk.

Gradio Demonstration

- Model will classify whether patient is at risk for <30 days readmittance
- Client can input top features
 - This is for clean demo purposes
 - Final product can allow for all features
- Client can choose to adjust patient's treatment

Hospital Readmission Predictor

Predicts the probability of readmission within 30 days based on patient history and discharge details.

Age Group

Time in Hospital (Days)

Number of Procedures

Number of Medications

Inpatient Visits (Prior Year)

Total Visit Count (History)

Primary Diagnosis

Discharge Destination

Clear

Submit

Flag

output

Thank You. Any Questions?

REFERENCES

- Ashfaq, A., Sant'Anna, A., Lingman, M., & Nowaczyk, S. (2019). Readmission prediction using deep learning on electronic health records. *Journal of Biomedical Informatics*, 97, 103271. <https://doi.org/10.1016/j.jbi.2019.103271>
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