MIMIC-IV Mortality Prediction Tutorial

A step-by-step guide using clinical data and machine learning

Introduction

- Predictive Analytics in Healthcare
 - Leveraging large-scale EHR data to predict patient outcomes
 - MIMIC-IV: Publicly available ICU dataset with structured and unstructured data

Dataset Overview

- Key MIMIC-IV Tables:
 - ADMISSIONS
 - PATIENTS
 - **ICUSTAYS**
 - DIAGNOSES_ICD

Goal

 Predict in-hospital mortality at the time of ICU admission using EHR data

Import Libraries

- pandas
- numpy
- scikit-learn
- tensorflow / PyTorch
- transformers (ClinicalBERT)

Data Loading

Load data from MIMIC IV BigQuery into

DataFrames

Admissions Data

Diagnoses code categories

```
# Encode categorical variables

model_df = admissions_df.copy)

# Calculate length of stay in days

model_df('idschtime'l) - pd.to_datetime(model_df('dschtime'l) - pd.to_datetime(model_df('admittime'l)).dt.days

# Merge diagnosis category dumines from ding_dumines (created in next cell)

# (This merge will be performed after diag_dumines is created)

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# (This merge will be performed after install().astype(int)

# Display the first fav rows and basic statistics of the prepared dataset print(model_df, head())

# print(model_df, head())

# admissions.df('gender'.lamp(('N': 8, 'F': 11))

# The diagnosis category columes will be added after merging below.

# Final feature set will be updated after merging diagnosis dumines.

## Obs.

## Display.diagnosis category columes will be added after merging below.

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```

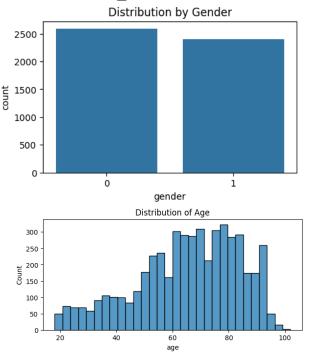
Length of Stay, Mortality and Gender

```
mimic_diag_query = """
SELECT hadm id. icd code
FROM `physionet-data.mimiciv_3_1_hosp.diagnoses_icd`
diagnoses_df = client.query(mimic_diag_query).to_dataframe()
def icd to category(icd code):
        code = str(icd_code)
        if code[0].isalpha():
           return 'misc
        code int = int(code[:3])
        if 1 <= code_int < 140: return 'infectious'
        if 140 <= code_int < 240: return 'neoplasms'
        if 240 <= code_int < 280: return 'endocrine'
        if 280 <= code_int < 290: return 'blood'
        if 290 <= code_int < 320: return 'mental'
        if 320 <= code int < 390; return 'nervous'
        if 390 <= code_int < 460: return 'circulatory
        if 460 <= code_int < 520: return 'respiratory
        if 520 <= code_int < 580: return 'digestive
        if 580 <= code_int < 630: return 'genitourinary'
        if 630 <= code_int < 680: return 'pregnancy'
        if 680 <= code int < 710: return 'skin'
        if 710 <= code_int < 740: return 'muscular'
        if 760 <= code_int < 780: return 'prenatal'
        if 780 <= code int < 800: return 'misc'
        if 800 <= code int < 1000: return 'injury
        return 'misc'
```

Feature Engineering

- Demographics: age, gender, ethnicity
- Admission info: type, source
- Clinical measurements: lab results, vitals
- Diagnosis codes grouped by ICD-9 categories

```
# Drop datetime and 'gender' columns for modeling
datetime_cols = ['admittime', 'dischtime', 'deathtime']
drop_cols = ['subject_id', 'hadm_id'] + datetime_cols
diag categories = [
for cat in diag_categories:
   x_col = f'{cat}_x'
    if x_col in X_train.columns and y_col in X_train.columns:
        X_train[f'{cat}'] = X_train[x_col].fillna(0) + X_train[y_col].fillna(0)
        X_test[f'{cat}'] = X_test[x_col].fillna(0) + X_test[y_col].fillna(0)
diag_x_{cols} = [f'_{cat}_x' \text{ for cat in diag_categories if } f'_{cat}_x' \text{ in } X_{train.columns}]
diag_y_cols = [f'{cat}_y' for cat in diag_categories if f'{cat}_y' in X_train.columns]
X_train_clean = X_train.drop(columns=drop_cols + diag_x_cols + diag_y_cols + ['gender'])
X_test_clean = X_test.drop(columns=['gender'] + diag_x_cols + diag_y_cols + drop_cols)
imputer = SimpleImputer(strategy='mean')
# Ensure dense output in case imputer returns a sparse matrix
if hasattr(X_train_imputed, "toarray") (function) toarray: Unknown
    X_train_imputed = X_train_imputed.toarray()
X_train_clean = pd.DataFrame(X_train_imputed, columns=X_train_clean.columns, index=X_train_clean.index)
X_test_imputed = imputer.transform(X_test_clean)
if hasattr(X_test_imputed, "toarray"):
```



Modeling

- Train classifiers: Logistic Regression, Random Forest, XGBoost and SVM
- Fine-tune ClinicalBERT for text classification
- Split data: 80% train, 20% test

```
# Fine-tuned models with parameters
models = {
    'Logistic Regression': LogisticRegression(max_iter=1000, solver='lbfgs'),
    'Random Forest': RandomForestClassifier(n_estimators=100, max_depth=7),
    'XGBoost': XGBClassifier(use_label_encoder=False, eval_metric='logloss', max_depth=7, learning_rate=0.1),
    'SVM': SVC(probability=True, kernel='rbf', C=1.0)
}

results = {}
for name, model in models.items():
    model.fit(X_train_clean, y_train)
    y_pred = model.predict(X_test_clean)
    results[name] = {'model': model, 'y_pred': y_pred}
    print(f'{name} training and prediction done.')
```

Evaluation

- R^2 and RMSE Score Comparison for various models
- Confusion matrix analysis
- Feature Importance

```
# Evaluate models using R<sup>2</sup> and RMSE
for name, result in results.items():
   y_pred = result['y_pred']
    r2 = r2_score(y_test, y_pred)
    rmse = np.sqrt(mean_squared_error(y_test, y_pred))
    print(f'{name}: R² = {r2:.3f}, RMSE = {rmse:.3f}')
    results[name]['r2'] = r2
    results[name]['rmse'] = rmse
# Plot comparison of models for R2 and RMSE
model names = list(results.keys())
r2_scores = [results[m]['r2'] for m in model_names]
rmse_scores = [results[m]['rmse'] for m in model_names]
plt.figure(figsize=(10,4))
plt.subplot(1,2,1)
sns.barplot(x=model_names, y=r2_scores)
plt.title('R2 Comparison')
plt.ylabel('R2')
plt.subplot(1,2,2)
sns.barplot(x=model_names, y=rmse_scores)
plt.title('RMSE Comparison')
plt.ylabel('RMSE')
plt.tight_layout()
plt.show()
```

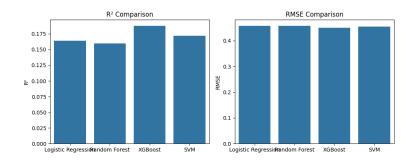
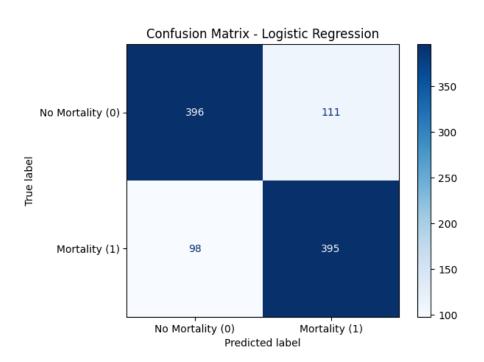


Table 1

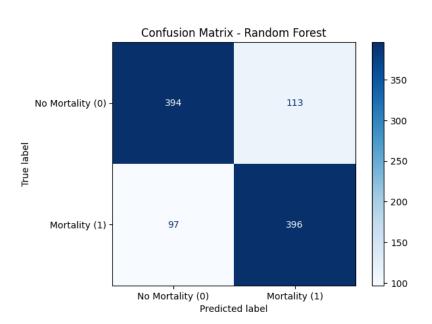
Logistic Regression: R ² = 0.164	RMSE = 0.457
Random Forest: R ² = 0.160	RMSE = 0.458
XGBoost: R ² = 0.188	RMSE = 0.451
SVM: R ² = 0.172	RMSE = 0.455

Evaluation - Logistics Regression



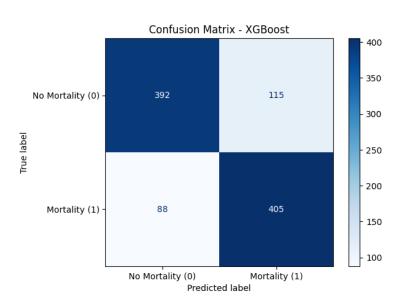
Logistic Regression: Accuracy=0.791, Precision=0.781, Recall=0.801, F1=0.791

Evaluation - Random Forest



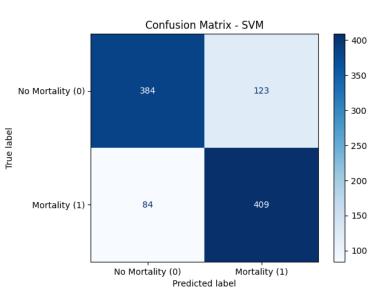
Random Forest: Accuracy=0.790, Precision=0.778, Recall=0.803, F1=0.790

Evaluation - XGBoost



XGBoost: Accuracy=0.797, Precision=0.779, Recall=0.822, F1=0.800

Evaluation - SVM



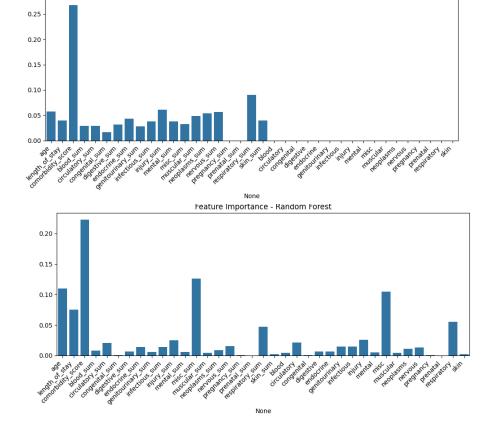
SVM: Accuracy=0.793, Precision=0.769, Recall=0.830, F1=0.798

Feature Importance

Identify top predictive features influencing

mortality risk

```
# Feature importance for tree-based models
for name in ['Random Forest', 'XGBoost']:
   model = results[name]['model']
    importances = model.feature_importances_
    feature_names = X_train_clean.columns
   plt.figure(figsize=(10, 5))
   sns.barplot(x=feature names, y=importances)
   plt.title(f'Feature Importance - {name}')
   plt.xticks(rotation=45, ha='right')
   plt.tight_layout()
   plt.show()
# Coefficients for linear model
lr model = results['Logistic Regression']['model']
if hasattr(lr_model, 'coef_'):
   coef = lr_model.coef_[0]
    feature_names = X_train_clean.columns
    plt.figure(figsize=(8, 5))
    sns.barplot(x=coef, y=feature_names, orient='h')
   plt.title('Feature Coefficients - Logistic Regression')
   plt.xlabel('Coefficient Value')
   plt.ylabel('Feature')
   plt.tight_layout()
    plt.show()
```

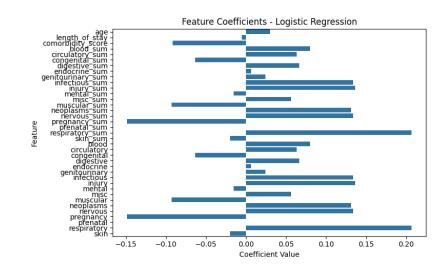


Feature Importance - XGBoost

Feature Importance

 Identify top predictive features influencing mortality risk

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   plt.xlabel('Coefficient Value')
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   plt.tight_layout()
    plt.show()
```



Conclusion & Next Steps

- With R^2 score of 0.172 and RMSE of 0.455, SVM model provide the best recall (True Positive) for prediction
- Integrate real-time EHR streaming data
- Deploy model in clinical decision support systems