Cervical Cancer Detection Using Cytology Data This notebook demonstrates the process of detecting cervical cancer using cytology data. It includes data preprocessing, handling missing values, feature scaling, addressing class imbalance, model training, evaluation, and saving the trained model for future use. 1. Import Necessary Libraries In [1]: import pandas as pd import numpy as np import matplotlib.pyplot as plt import seaborn as sns from sklearn.model_selection import train_test_split from sklearn.impute import KNNImputer from sklearn.preprocessing import StandardScaler from sklearn.ensemble import RandomForestClassifier from sklearn.metrics import classification report from imblearn.over sampling import SMOTE import joblib import warnings warnings.filterwarnings("ignore") # Set seaborn style for better aesthetics sns.set(style='whitegrid', palette='muted', font_scale=1.1) plt.rcParams['figure.figsize'] = (10, 6) 2: Load and Inspect the Dataset Load the dataset and perform initial inspection to understand its structure and identify any missing values. In [2]: # Load the dataset data = pd.read_csv('cervical_cancer_dataset.csv') In [3]: # Display the shape of the dataset print(f"Dataset Shape: {data.shape}") Dataset Shape: (858, 36) In [4]: # Show the first five rows of the dataset data.head() Out[4]: STDs: STDs: Number Time Hormonal of First sexual **Smokes Smokes** Time Num of Hormonal Dx:Cancer Dx:CIN Dx:HPV Dx Hinselmann Schiller Citolo **Smokes** Contraceptives IUD ... since Age (years) (packs/year) Contraceptives sexual intercourse pregnancies since last first (years) diagnosis partners diagnosis 0.0 0.0 ... 0 0 18 4.0 15.0 1.0 0.0 0.0 0.0 0.0 ? ? 0 0 0 0 0 0.0 ... 0 0 15 14.0 0.0 0.0 0.0 0.0 ? ? 0 0 0 1.0 1.0 0.0 0 0.0 0.0 0.0 ... ? 0 0 34 1.0 ? 1.0 0.0 0.0 ? 0 0 0 0 0.0 0 52 16.0 37.0 37.0 1.0 0.0 ... ? ? 1 1 0 0 5.0 4.0 1.0 3.0 0 46 0.0 15.0 0.0 ... ? 0 0 ? 0 0 0 0 3.0 21.0 0.0 0.0 1.0 4.0 5 rows × 36 columns In [5]: # Check for missing values represented by '?' missing_values = data.isin(['?']).sum() print("Missing values in each column:") print(missing values[missing values > 0]) Missing values in each column: Number of sexual partners 26 7 First sexual intercourse 56 Num of pregnancies Smokes 13 13 Smokes (years) Smokes (packs/year) 13 108 Hormonal Contraceptives Hormonal Contraceptives (years) 108 117 IUD (years) 117 STDs 105 105 STDs (number) STDs:condylomatosis 105 STDs:cervical condylomatosis 105 STDs:vaginal condylomatosis 105 STDs:vulvo-perineal condylomatosis 105 105 STDs:syphilis STDs:pelvic inflammatory disease 105 STDs:genital herpes 105 STDs:molluscum contagiosum 105 STDs:AIDS 105 STDs:HIV 105 105 STDs:Hepatitis B 105 STDs:HPV 787 STDs: Time since first diagnosis STDs: Time since last diagnosis 787 dtype: int64 3: Data Preprocessing Replace placeholder missing values, convert data types, and select relevant features for analysis. In [6]: # Replace '?' with NaN and convert all data to numeric types data.replace('?', np.nan, inplace=True) data = data.apply(pd.to_numeric, errors='coerce') In [7]: # Define feature columns and target variable user_features = ['Age', 'Number of sexual partners', 'First sexual intercourse', 'Num of pregnancies', 'Smokes', 'Smokes (years)', 'Smokes (packs/year)', 'STDs', 'STDs (number)', 'Hormonal Contraceptives', 'Hormonal Contraceptives (years)', 'IUD', 'IUD (years)', target = 'Dx:Cancer' In [8]: # Exclude irrelevant or redundant features features_to_exclude = ['STDs: Time since first diagnosis', 'STDs: Time since last diagnosis', data.drop(columns=features_to_exclude, inplace=True, errors='ignore') In [9]: # Separate features and target variable X = data[user_features] y = data[target] 4: Handle Missing Values Identify and impute missing values using K-Nearest Neighbors imputation. In [10]: print("Number of missing values per column:") print(X.isnull().sum()) Number of missing values per column: Age 26 Number of sexual partners First sexual intercourse Num of pregnancies Smokes Smokes (years) 13 Smokes (packs/year) 13 STDs 105 STDs (number) 105 Hormonal Contraceptives 108 Hormonal Contraceptives (years) 108 IUD 117 IUD (years) 117 dtype: int64 In [11]: # Initialize KNN Imputer with 5 neighbors imputer = KNNImputer(n_neighbors=5) # Impute missing values X_imputed = pd.DataFrame(imputer.fit_transform(X), columns=X.columns) 5: Feature Scaling Standardize numerical features to ensure they contribute equally to the model training. In [12]: # Define numerical features for scaling numerical_features = ['Age', 'Number of sexual partners', 'First sexual intercourse', 'Num of pregnancies', 'Smokes (years)', 'Smokes (packs/year)', 'Hormonal Contraceptives (years)', 'IUD (years)', 'STDs (number)', # Initialize StandardScaler scaler = StandardScaler() # Apply scaling to numerical features X_imputed[numerical_features] = scaler.fit_transform(X_imputed[numerical_features]) 6: Address Class Imbalance Visualize class distribution and apply SMOTE to balance the classes. In [13]: # Visualize class distribution before applying SMOTE plt.figure(figsize=(6, 4)) sns.countplot(x=y) plt.title('Class Distribution Before SMOTE') plt.xlabel('Cancer Diagnosis') plt.ylabel('Count') plt.show() Class Distribution Before SMOTE 800 700 600 500 400 300 200 100 0 0 Cancer Diagnosis In [14]: print("Class distribution before SMOTE:") print(y.value counts()) Class distribution before SMOTE: Dx:Cancer 840 Name: count, dtype: int64 In [15]: # Initialize SMOTE with a fixed random state for reproducibility sm = SMOTE(random_state=42) # Apply SMOTE to balance the classes X_resampled, y_resampled = sm.fit_resample(X_imputed, y) In [16]: # Visualize class distribution after applying SMOTE plt.figure(figsize=(6, 4)) sns.countplot(x=y resampled) plt.title('Class Distribution After SMOTE') plt.xlabel('Cancer Diagnosis') plt.ylabel('Count') plt.show() Class Distribution After SMOTE 800 700 600 500 400 300 200 100 0 0 **Cancer Diagnosis** In [17]: print("Class distribution after SMOTE:") print(y_resampled.value_counts()) Class distribution after SMOTE: Dx:Cancer 840 840 Name: count, dtype: int64 7: Split the Data into Training and Testing Sets Divide the dataset into training and testing subsets to evaluate model performance. In [18]: X_train, X_test, y_train, y_test = train_test_split(X_resampled, y_resampled, test_size=0.2, random_state=42, stratify=y_resampled 8: Train the Random Forest Classifier Initialize and train the Random Forest model with specified hyperparameters. In [19]: # Initialize the Random Forest Classifier with balanced class weights model = RandomForestClassifier(n_estimators=200, max_depth=6, min_samples_split=2, min_samples_leaf=1, class_weight='balanced', random_state=42 In [20]: # Train the model on the training data model.fit(X_train, y_train) Out[20]: 🔻 RandomForestClassifier RandomForestClassifier(class_weight='balanced', max_depth=6, n_estimators=200, random_state=42) 9: Evaluate the Model Assess the trained model's performance using classification metrics. In [21]: # Make predictions on the test set y_pred = model.predict(X_test) # Obtain prediction probabilities for the positive class y_proba = model.predict_proba(X_test)[:, 1] In [22]: # Display the classification report print("Classification Report:") print(classification_report(y_test, y_pred)) Classification Report: precision recall f1-score support 0.94 0.93 0.94 168 1 0.93 0.94 0.94 168 336 0.94 accuracy 0.94 0.94 0.94 336 macro avg weighted avg 0.94 0.94 0.94 336 In [23]: # Visualize feature importance importances = model.feature_importances_ indices = np.argsort(importances)[::-1] features = X_train.columns plt.figure(figsize=(12, 8)) sns.barplot(x=importances[indices], y=features[indices], palette='viridis') plt.title('Feature Importances') plt.xlabel('Importance') plt.ylabel('Feature') plt.show() /var/folders/ff/sds78knx6lv3kyl_xckqx8cc0000gn/T/ipykernel_8023/1891037928.py:7: FutureWarning: Passing `palette` without assigning `hue` is deprecated and will be removed in v0.14.0. Assign the `y` variable to `hue` and set `legend=False` for the same effect. sns.barplot(x=importances[indices], y=features[indices], palette='viridis') Feature Importances First sexual intercourse Num of pregnancies Age Number of sexual partners IUD (years) Hormonal Contraceptives (years) Feature IUD **Hormonal Contraceptives** Smokes STDs (number) Smokes (packs/year) STDs Smokes (years) 0.00 0.05 0.10 0.15 0.20 0.25 Importance 10. Correlation Matrix Generate and visualize the correlation matrix to understand relationships between features. In [24]: # Compute the correlation matrix corr_matrix = data[user_features + [target]].corr() # Plot the heatmap plt.figure(figsize=(14, 12)) sns.heatmap(corr_matrix, annot=True, fmt=".2f", cmap='coolwarm', linewidths=0.5, annot_kws={"size": 10} plt.title('Correlation Matrix of Features', fontsize=16) plt.xticks(rotation=45, ha='right') plt.yticks(rotation=0) plt.tight_layout() plt.show() **Correlation Matrix of Features** 0.09 0.06 0.22 0.13 -0.02 0.07 0.28 0.22 Age 1.00 0.37 0.55 0.01 0.29 Number of sexual partners -0.15 0.19 0.09 1.00 0.08 0.25 0.18 0.06 0.04 0.01 0.02 0.03 0.00 0.02 - 0.8 First sexual intercourse -0.15 -0.12-0.06 -0.06 -0.01 0.37 1.00 -0.060.01 0.02 0.01 -0.02-0.030.07 Num of pregnancies 0.08 0.55 0.08 -0.06 0.18 0.10 0.05 0.00 0.16 0.22 0.22 0.15 0.04 1.00 0.06 -0.120.72 0.13 0.12 0.05 -0.06 -0.04Smokes 0.25 0.08 1.00 0.49 -0.00 -0.01 - 0.6 0.04 Smokes (years) 0.22 -0.06 0.18 0.72 0.72 0.10 -0.01 0.05 0.03 0.06 0.19 1.00 Smokes (packs/year) -0.06 0.72 0.01 0.04 0.02 0.01 0.13 0.18 0.49 1.00 0.03 0.03 - 0.4 STDs 0.01 0.06 0.05 0.13 0.10 0.92 -0.03 0.00 0.06 0.02 0.00 -0.01 0.03 1.00 STDs (number) -0.02 0.04 0.01 0.00 0.12 0.03 -0.04 -0.01 0.05 0.02 -0.02 0.92 1.00 **Hormonal Contraceptives** -0.03 0.07 0.01 0.02 0.16 -0.00-0.01 0.01 -0.03-0.040.45 0.03 0.03 1.00 - 0.2 Hormonal Contraceptives (years) 0.02 0.22 0.05 0.05 0.04 0.00 -0.01 1.00 0.10 0.00 0.05 0.29 0.01 0.45 IUD 0.03 0.22 -0.06 0.03 0.06 0.05 0.03 0.28 0.01 -0.021.00 - 0.0 IUD (years) -0.04 0.04 1.00 0.00 -0.030.15 0.02 0.02 0.02 -0.03 0.00 0.10 0.75 0.22 Dx:Cancer 0.02 0.07 0.04 -0.01 0.06 0.00 -0.02 0.03 0.05 1.00 STDS (humber)
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STDS (humber) 11: Define the Prediction Function Create a function to predict the risk of cervical cancer based on user input. def predict_risk(input_data): Predicts the risk probability of cervical cancer based on input features. Parameters: input_data (dict): A dictionary containing feature values. Returns: dict: A dictionary with the risk probability. # Convert input data to DataFrame input_df = pd.DataFrame([input_data]) # Ensure all required features are present for feature in X.columns: if feature not in input_df.columns: input_df[feature] = np.nan # Impute missing values input_df_imputed = pd.DataFrame(imputer.transform(input_df), columns=input_df.columns) # Scale numerical features input_df_imputed[numerical_features] = scaler.transform(input_df_imputed[numerical_features]) # Predict risk probability risk_proba = model.predict_proba(input_df_imputed)[0][1] return {'Risk Probability': risk_proba} 12: Example User Input and Prediction Provide an example of how to use the prediction function with sample user data. In [26]: # Example user input user_input = { 'Age': 25, 'Number of sexual partners': 0, 'First sexual intercourse': 0, 'Num of pregnancies': 0, 'Smokes': 0, 'Smokes (years)': 0, 'Smokes (packs/year)': 0, 'STDs': 0, 'STDs (number)': 0, 'Hormonal Contraceptives': 0, 'Hormonal Contraceptives (years)': 0, 'IUD': 0, 'IUD (years)': 0, # Make a prediction prediction = predict_risk(user_input) # Display the prediction result print("\nRisk Prediction for the User:") print(f"Risk Probability: {prediction['Risk Probability']*100:.2f}%") Risk Prediction for the User: Risk Probability: 1.87% 13: Save the Model for Future Use Persist the trained model to disk for later deployment or inference. In [27]: # Save the trained model to a file joblib.dump(model, "ML_model.pkl") print("Model saved as 'ML_model.pkl'") Model saved as 'ML_model.pkl' Summary This notebook provides a complete workflow for detecting cervical cancer based on cytology data using traditional machine learning techniques. It encompasses data loading and inspection, preprocessing steps such as handling missing values and feature scaling, and addressing class imbalance with SMOTE. A Random Forest Classifier is trained and evaluated with detailed classification metrics. The notebook includes insightful visualizations like feature importance and a correlation matrix to understand feature relationships. Additionally, it offers a user-friendly prediction function for assessing individual risk and demonstrates how to save the trained model for future use. The well-structured and commented code ensures ease of understanding and adaptability for similar datasets and classification tasks.