



Breast Cancer Prediction

Breast Cancer Prediction is a classification task aimed at predicting the diagnosis of a breast mass as either malignant or benign. The dataset used for this prediction consists of features computed from a digitized image of a fine needle aspirate (FNA) of the breast mass. These features describe various characteristics of the cell nuclei present in the image.

The dataset contains the following information for each instance:

1. **ID number:** A unique identifier for each sample.
2. **Diagnosis:** The target variable indicating the diagnosis, where 'M' represents malignant and 'B' represents benign.

For each cell nucleus, ten real-valued features are computed, which are:

1. **Radius:** The mean distance from the center to points on the perimeter of the nucleus.
2. **Texture:** The standard deviation of gray-scale values in the nucleus.
3. **Perimeter:** The perimeter of the nucleus.
4. **Area:** The area of the nucleus.
5. **Smoothness:** A measure of local variation in radius lengths.
6. **Compactness:** Computed as the square of the perimeter divided by the area minus 1.0.
7. **Concavity:** Describes the severity of concave portions of the nucleus contour.
8. **Concave points:** Represents the number of concave portions of the nucleus contour.
9. **Symmetry:** Measures the symmetry of the nucleus.
10. **Fractal dimension:** This feature approximates the "coastline" of the nucleus, using the concept of fractal geometry.

These features provide quantitative measurements that can be used to assess the characteristics of cell nuclei and aid in distinguishing between malignant and benign breast masses. By training a machine learning model on this dataset, it is possible to develop a predictive model that can assist in the early detection and diagnosis of breast cancer.

```
In [1]: # Importing libraries
import numpy as np
import pandas as pd
import matplotlib.pyplot as plt
import seaborn as sns
```

```
In [2]: # loading Dataset
df = pd.read_csv("/content/Breast_Cancer_Dataset.csv")
```

EDA

```
In [3]: df.head()
```

```
Out[3]:
```

| | id | diagnosis | radius_mean | texture_mean | perimeter_mean | area_mean |
|---|----------|-----------|-------------|--------------|----------------|-----------|
| 0 | 842302 | M | 17.99 | 10.38 | 122.80 | 1001. |
| 1 | 842517 | M | 20.57 | 17.77 | 132.90 | 1326. |
| 2 | 84300903 | M | 19.69 | 21.25 | 130.00 | 1203. |
| 3 | 84348301 | M | 11.42 | 20.38 | 77.58 | 386. |
| 4 | 84358402 | M | 20.29 | 14.34 | 135.10 | 1297. |

5 rows × 33 columns

```
In [4]: df.shape
```

```
Out[4]: (569, 33)
```

```
In [5]: df.info()
```

```

<class 'pandas.core.frame.DataFrame'>
RangeIndex: 569 entries, 0 to 568
Data columns (total 33 columns):
#   Column                                     Non-Null Count  Dtype
---  -
0   id                                           569 non-null    int64
1   diagnosis                                    569 non-null    object
2   radius_mean                                569 non-null    float64
3   texture_mean                               569 non-null    float64
4   perimeter_mean                             569 non-null    float64
5   area_mean                                  569 non-null    float64
6   smoothness_mean                            569 non-null    float64
7   compactness_mean                           569 non-null    float64
8   concavity_mean                             569 non-null    float64
9   concave points_mean                        569 non-null    float64
10  symmetry_mean                              569 non-null    float64
11  fractal_dimension_mean                     569 non-null    float64
12  radius_se                                  569 non-null    float64
13  texture_se                                 569 non-null    float64
14  perimeter_se                               569 non-null    float64
15  area_se                                    569 non-null    float64
16  smoothness_se                              569 non-null    float64
17  compactness_se                             569 non-null    float64
18  concavity_se                               569 non-null    float64
19  concave points_se                          569 non-null    float64
20  symmetry_se                                569 non-null    float64
21  fractal_dimension_se                       569 non-null    float64
22  radius_worst                               569 non-null    float64
23  texture_worst                              569 non-null    float64
24  perimeter_worst                            569 non-null    float64
25  area_worst                                 569 non-null    float64
26  smoothness_worst                           569 non-null    float64
27  compactness_worst                          569 non-null    float64
28  concavity_worst                             569 non-null    float64
29  concave points_worst                       569 non-null    float64
30  symmetry_worst                              569 non-null    float64
31  fractal_dimension_worst                     569 non-null    float64
32  Unnamed: 32                                0 non-null      float64
dtypes: float64(31), int64(1), object(1)
memory usage: 146.8+ KB

```

```

In [6]: # Duplicate values check
df.duplicated().sum()

```

```

Out[6]: np.int64(0)

```

The Breast Cancer dataset consists of 569 observations and 33 columns with no duplicated rows, combining an identifier, a binary target variable, and multiple numerical features derived from cell nucleus measurements. As there are no missing values are present in the core features. But one column (Unnamed: 32) contains only null values and provides no information. Unnamed: 32 should be dropped during preprocessing

Target Variable Analysis

```
In [7]: # Class distribution (M vs B)
df['diagnosis'].value_counts()
```

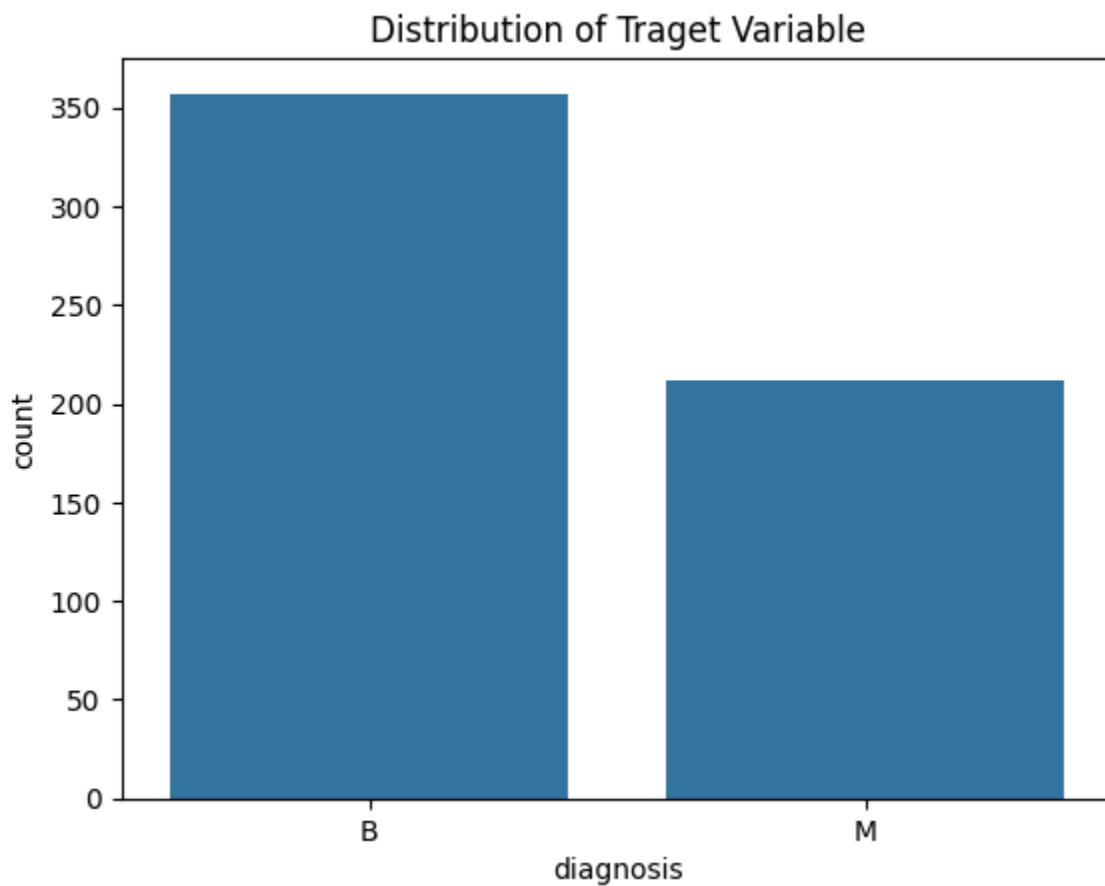
```
Out[7]:
```

| | count |
|---|-------|
| B | 357 |
| M | 212 |

| diagnosis | count |
|-----------|-------|
| B | 357 |
| M | 212 |

dtype: int64

```
In [8]: # Visualizing Class Distribution
sns.countplot(data=df, x='diagnosis', order=df['diagnosis'].value_counts().index)
plt.title('Distribution of Target Variable')
plt.show()
```



The dataset contains a moderate class imbalance, with Benign (B) cases occurring more frequently than Malignant (M) cases.

Univariate Analysis

```
In [9]: df.describe()
```

```
Out[9]:
```

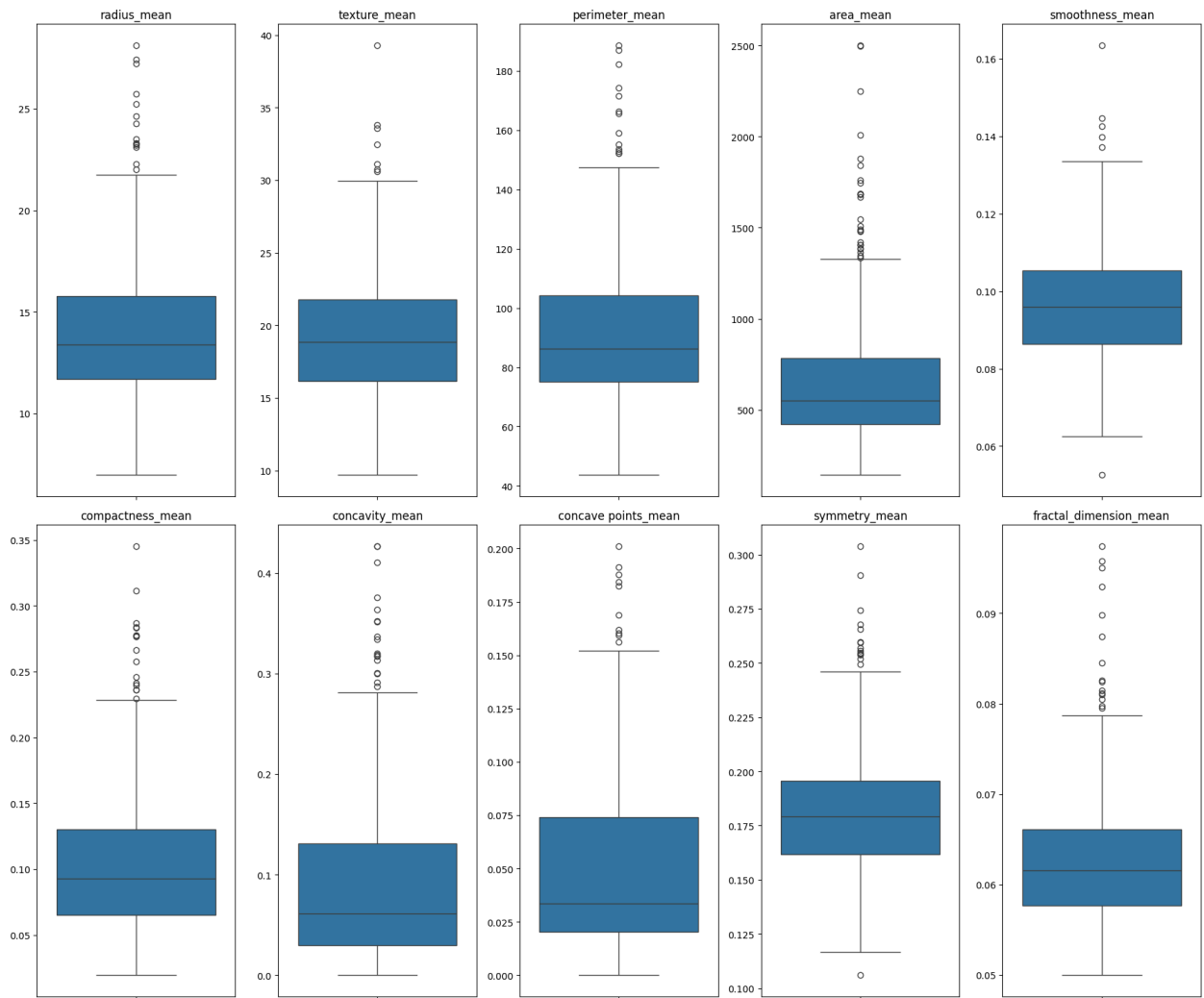
| | id | radius_mean | texture_mean | perimeter_mean | area_mean |
|--------------|--------------|-------------|--------------|----------------|-------------|
| count | 5.690000e+02 | 569.000000 | 569.000000 | 569.000000 | 569.000000 |
| mean | 3.037183e+07 | 14.127292 | 19.289649 | 91.969033 | 654.889104 |
| std | 1.250206e+08 | 3.524049 | 4.301036 | 24.298981 | 351.914129 |
| min | 8.670000e+03 | 6.981000 | 9.710000 | 43.790000 | 143.500000 |
| 25% | 8.692180e+05 | 11.700000 | 16.170000 | 75.170000 | 420.300000 |
| 50% | 9.060240e+05 | 13.370000 | 18.840000 | 86.240000 | 551.100000 |
| 75% | 8.813129e+06 | 15.780000 | 21.800000 | 104.100000 | 782.700000 |
| max | 9.113205e+08 | 28.110000 | 39.280000 | 188.500000 | 2501.000000 |

8 rows × 32 columns

```
In [10]: df.columns
```

```
Out[10]: Index(['id', 'diagnosis', 'radius_mean', 'texture_mean', 'perimeter_mean',  
                'area_mean', 'smoothness_mean', 'compactness_mean', 'concavity_mean',  
                'concave points_mean', 'symmetry_mean', 'fractal_dimension_mean',  
                'radius_se', 'texture_se', 'perimeter_se', 'area_se', 'smoothness_se',  
                'compactness_se', 'concavity_se', 'concave points_se', 'symmetry_se',  
                'fractal_dimension_se', 'radius_worst', 'texture_worst',  
                'perimeter_worst', 'area_worst', 'smoothness_worst',  
                'compactness_worst', 'concavity_worst', 'concave points_worst',  
                'symmetry_worst', 'fractal_dimension_worst', 'Unnamed: 32'],  
               dtype='object')
```

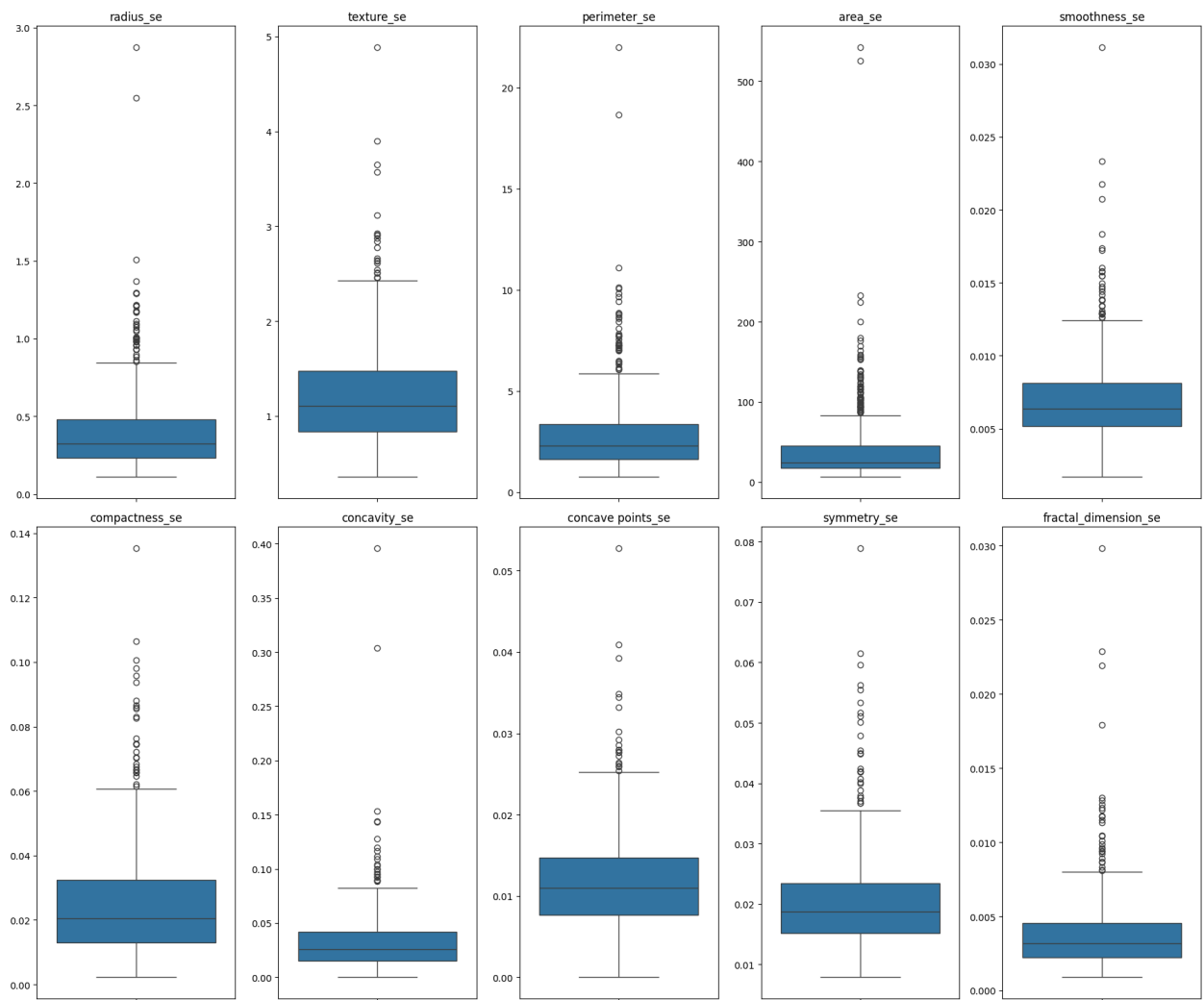
```
In [11]: # Cheking Outliers for _mean columns  
mean_cols = [c for c in df.columns if c.endswith('_mean')]  
fig, axes = plt.subplots(nrows=2, ncols=5, figsize=(18,15))  
axes = axes.flatten()  
  
for ax, col in zip(axes, mean_cols):  
    sns.boxplot(y=df[col], ax=ax)  
    ax.set_title(col)  
    ax.set_ylabel('')  
  
plt.tight_layout()  
plt.show()
```



```
In [12]: # Cheking Outliers for _se columns
se_cols = [c for c in df.columns if c.endswith('_se')]
fig, axes = plt.subplots(nrows=2, ncols=5, figsize=(18,15))
axes = axes.flatten()

for ax, col in zip(axes, se_cols):
    sns.boxplot(y=df[col], ax=ax)
    ax.set_title(col)
    ax.set_ylabel('')

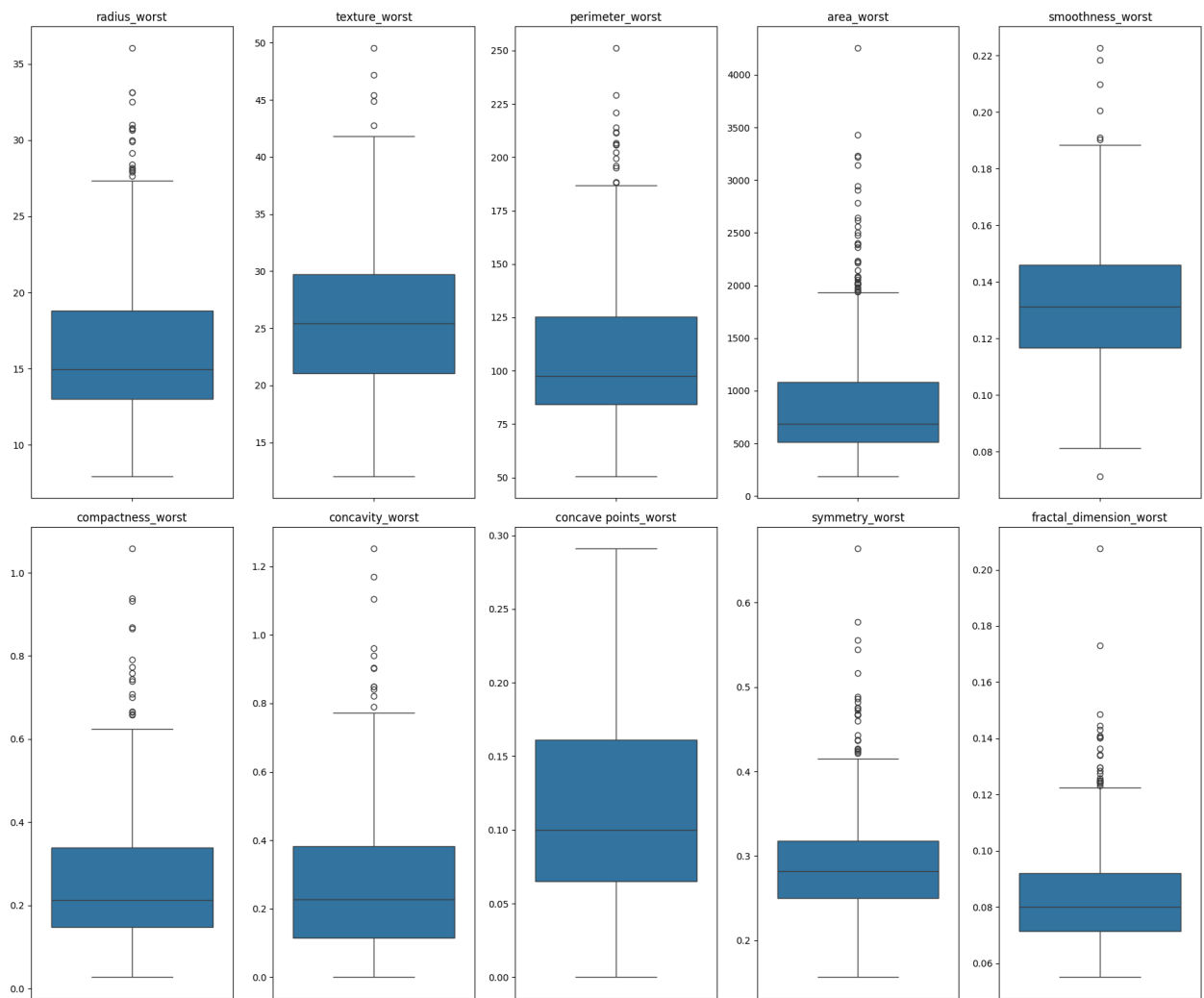
plt.tight_layout()
plt.show()
```



```
In [13]: # Cheking Outliers for _worst columns
worst_cols = [c for c in df.columns if c.endswith('_worst')]
fig, axes = plt.subplots(nrows=2, ncols=5, figsize=(18,15))
axes = axes.flatten()

for ax, col in zip(axes, worst_cols):
    sns.boxplot(y=df[col], ax=ax)
    ax.set_title(col)
    ax.set_ylabel('')

plt.tight_layout()
plt.show()
```



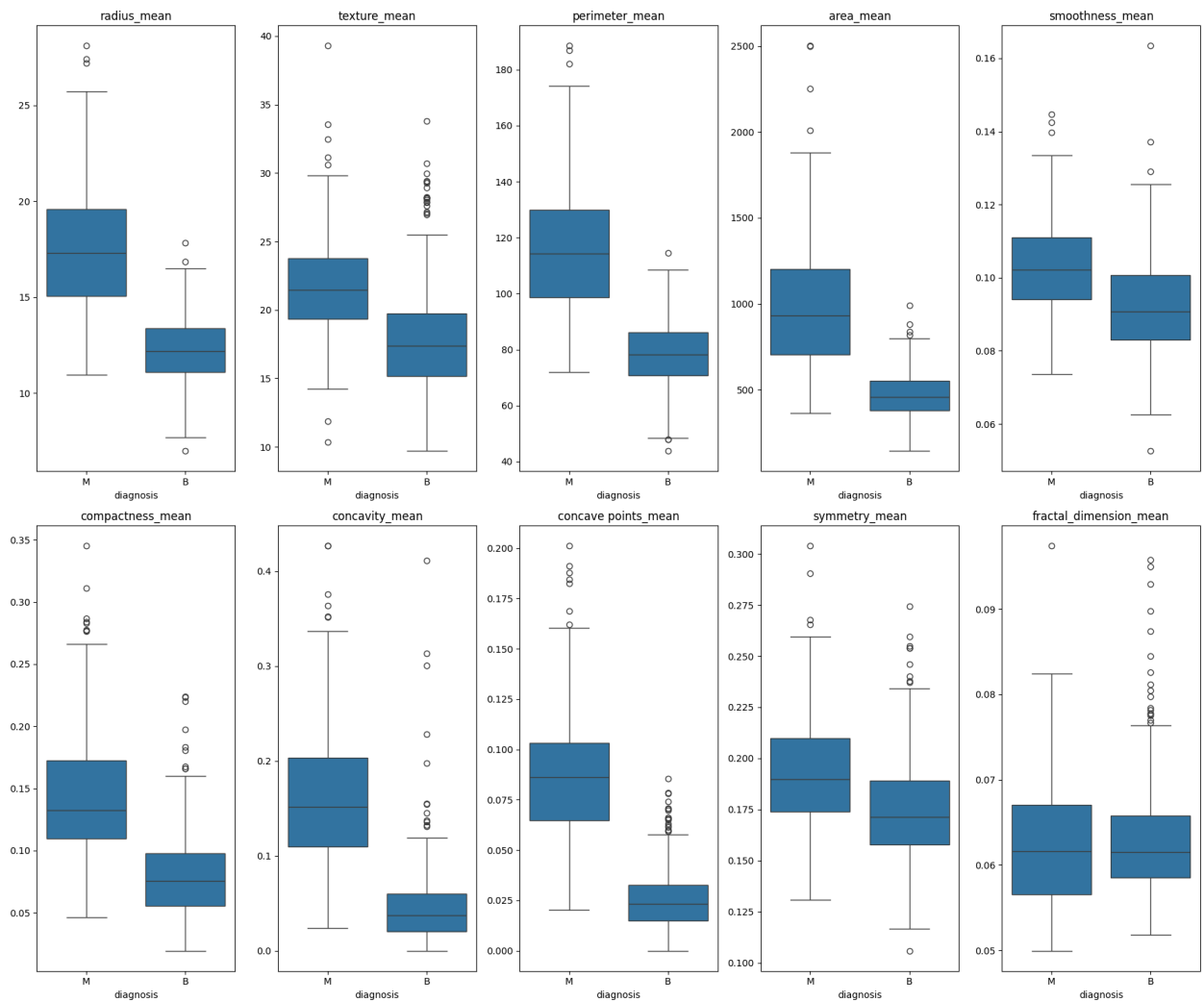
Boxplots reveal the presence of extreme values across several size and shape related features. These outliers reflect genuine biological variation rather than data quality issues. Consequently, no outlier removal was performed. Although feature scaling is required to ensure fair model training.

Feature vs Target Analysis

```
In [14]: fig, axes = plt.subplots(nrows=2, ncols=5, figsize=(18,15))
         axes = axes.flatten()

         for ax, col in zip(axes, mean_cols):
             sns.boxplot(data=df, y=col, x='diagnosis', ax=ax)
             ax.set_title(col)
             ax.set_ylabel('')
             ax.set_xlabel('')

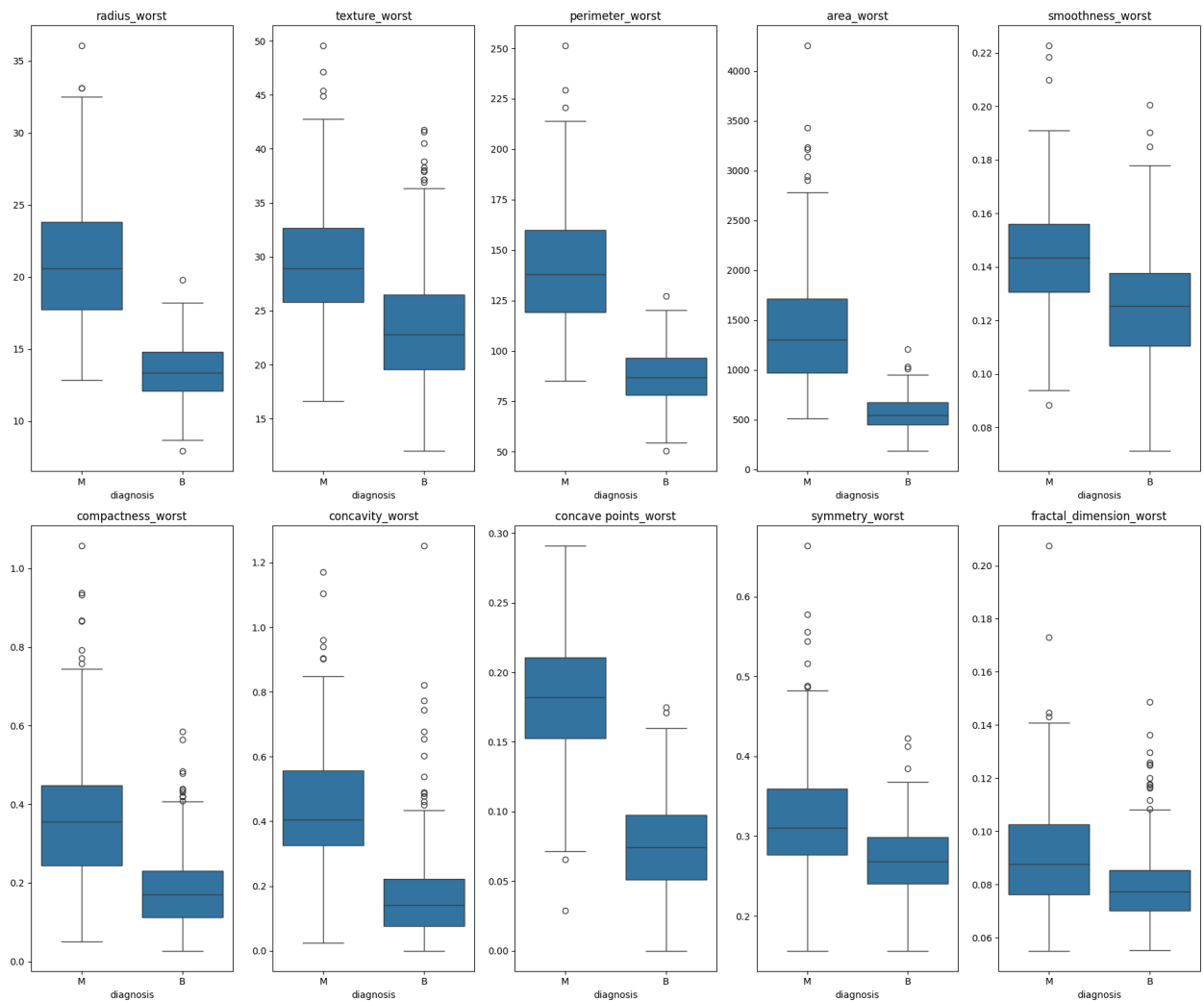
         plt.tight_layout()
         plt.show()
```

```
In [15]: fig, axes = plt.subplots(nrows=2, ncols=5, figsize=(18,15))
axes = axes.flatten()

for ax, col in zip(axes, worst_cols):
    sns.boxplot(data=df, y=col, x='diagnosis', ax=ax)
    ax.set_title(col)
    ax.set_ylabel('')
    ax.set_xlabel('')

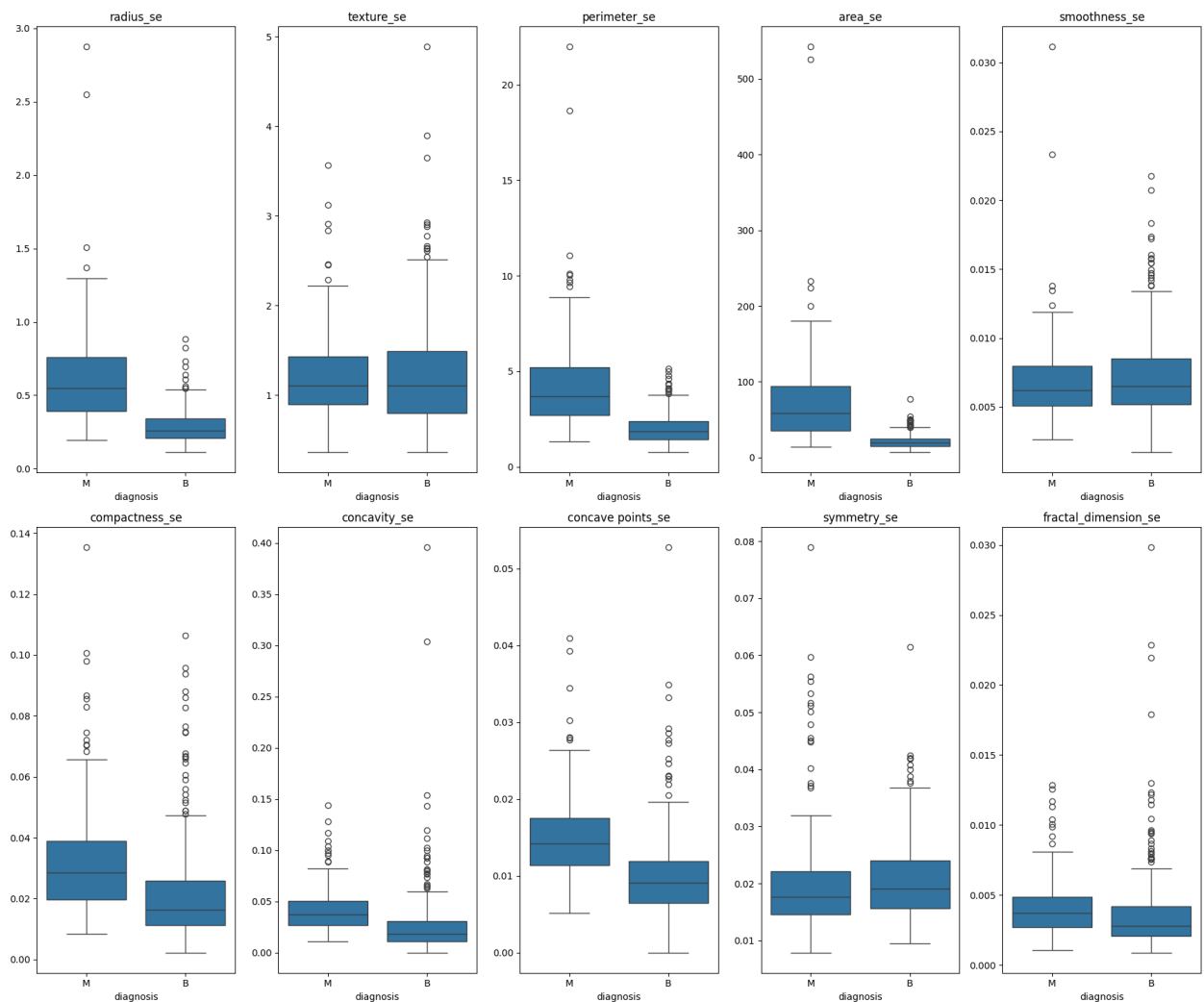
plt.tight_layout()
plt.show()
```



```
In [16]: fig, axes = plt.subplots(nrows=2, ncols=5, figsize=(18,15))
axes = axes.flatten()

for ax, col in zip(axes, se_cols):
    sns.boxplot(data=df, y=col, x='diagnosis', ax=ax)
    ax.set_title(col)
    ax.set_ylabel('')
    ax.set_xlabel('')

plt.tight_layout()
plt.show()
```



- Mean features (`*_mean`) show clear separation between classes, with malignant tumors exhibiting consistently higher values for size- and shape-related attributes (e.g., radius, perimeter, area, concavity).
- Worst-case features (`*_worst`) provide the strongest discrimination, capturing extreme tumor characteristics that are more prevalent in malignant cases.
- Standard error features (`*_se`) display substantial overlap between classes, indicating weaker standalone discriminative power and a more supportive role.

Correlation Analysis

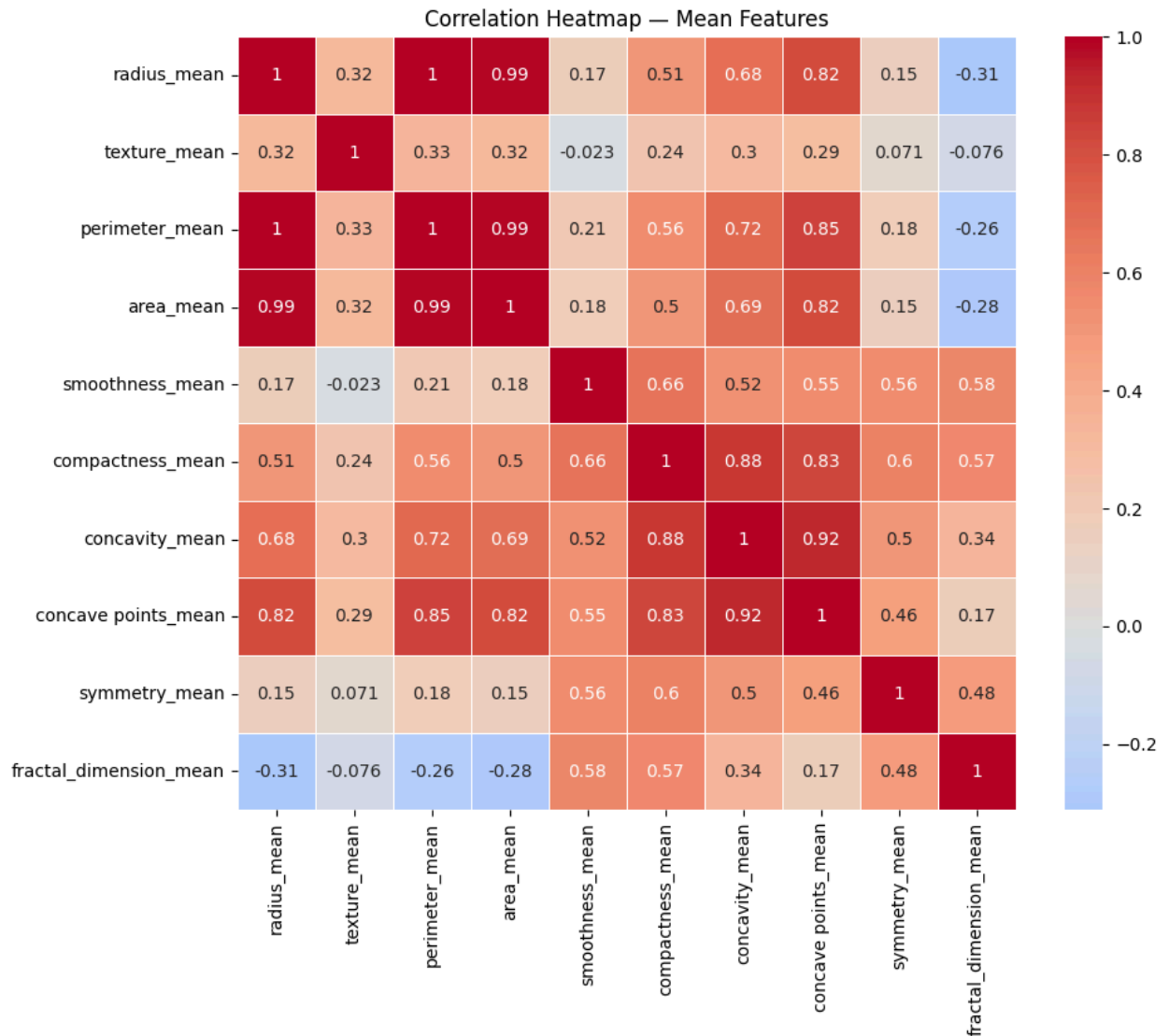
In [16]:

In [17]: `plt.figure(figsize=(10, 8))`

```

sns.heatmap(
    df[mean_cols].corr(),
    cmap='coolwarm',
    annot=True,
    center=0,
    linewidths=0.5
)
plt.title("Correlation Heatmap – Mean Features")
plt.show()

```



```

In [18]: plt.figure(figsize=(10, 8))
sns.heatmap(
    df[worst_cols].corr(),
    cmap='coolwarm',
    annot=True,
    center=0,
    linewidths=0.5
)
plt.title("Correlation Heatmap – Mean Features")
plt.show()

```



Correlation analysis revealed strong multicollinearity among size-related features such as radius, perimeter, and area, as well as among shape irregularity measures including concavity and concave points. These correlations are expected due to the physical relationships between features and justify the use of feature scaling and regularization during modeling rather than aggressive feature removal.

Pre-processing and Modeling

I will be using following models for this current project

- Logistic Regression
- Support Vector Machine (SVM)
- Random Forest
- K-Nearest Neighbours (KNN)

```
In [19]: # Importing model libraries
from sklearn.pipeline import Pipeline
from sklearn.preprocessing import StandardScaler
from sklearn.model_selection import train_test_split
from sklearn.linear_model import LogisticRegression
from sklearn.ensemble import RandomForestClassifier
from sklearn.svm import SVC
from sklearn.neighbors import KNeighborsClassifier
from sklearn.metrics import classification_report, roc_auc_score, accuracy_score
```

```
In [20]: df.columns
```

```
Out[20]: Index(['id', 'diagnosis', 'radius_mean', 'texture_mean', 'perimeter_mean',
               'area_mean', 'smoothness_mean', 'compactness_mean', 'concavity_mean',
               'concave points_mean', 'symmetry_mean', 'fractal_dimension_mean',
               'radius_se', 'texture_se', 'perimeter_se', 'area_se', 'smoothness_se',
               'compactness_se', 'concavity_se', 'concave points_se', 'symmetry_se',
               'fractal_dimension_se', 'radius_worst', 'texture_worst',
               'perimeter_worst', 'area_worst', 'smoothness_worst',
               'compactness_worst', 'concavity_worst', 'concave points_worst',
               'symmetry_worst', 'fractal_dimension_worst', 'Unnamed: 32'],
              dtype='object')
```

```
In [21]: # Features and target
X = df.drop(columns=['id', 'diagnosis', 'Unnamed: 32'])
y = df['diagnosis'].map({'B':0, 'M':1})

# Train/Test split (80%/20%)
X_train, X_test, y_train, y_test = train_test_split(X, y, test_size=0.2, stratify=y)

# Common preprocessing
preprocessor = Pipeline(steps=[
    ('scaler', StandardScaler())
])
```

Baseline Models

```
In [22]: # Logistic Regression
baseline_log_reg = Pipeline(steps=[
    ('preprocessing', preprocessor),
    ('model', LogisticRegression(max_iter=1000, class_weight='balanced', random_state=42))
])

# Support Vector Machine (RBF)
baseline_svm = Pipeline(steps=[
    ('preprocessing', preprocessor),
    ('model', SVC(kernel='rbf', probability=True, class_weight='balanced', random_state=42))
])

# Random Forest Classifier - Random Forest does not require scaling, so preprocessing is not needed
baseline_rf = Pipeline(steps=[
    ('model', RandomForestClassifier(n_estimators=200, random_state=42, class_weight='balanced'))
])
```

```

])

# KNN
baseline_knn = Pipeline(steps=[
    ('preprocessing', preprocessor),
    ('model', KNeighborsClassifier(n_neighbors=5))
])

models = {
    'Logistic Regression': baseline_log_reg,
    'SVM (RBF)': baseline_svm,
    'Random Forest Classification': baseline_svm,
    'KNN': baseline_knn
}

```

```

In [23]: # Model Evaluation
for name, pipeline in models.items():
    pipeline.fit(X_train, y_train)
    y_pred = pipeline.predict(X_test)
    y_proba = pipeline.predict_proba(X_test)[:, 1]

    print(f'\n{name}')
    print('ROC-AUC:', roc_auc_score(y_test, y_proba))
    print(classification_report(y_test, y_pred))

```

Logistic Regression

ROC-AUC: 0.9953703703703703

| | precision | recall | f1-score | support |
|--------------|-----------|--------|----------|---------|
| 0 | 0.97 | 0.99 | 0.98 | 72 |
| 1 | 0.98 | 0.95 | 0.96 | 42 |
| accuracy | | | 0.97 | 114 |
| macro avg | 0.97 | 0.97 | 0.97 | 114 |
| weighted avg | 0.97 | 0.97 | 0.97 | 114 |

SVM (RBF)

ROC-AUC: 0.9953703703703705

| | precision | recall | f1-score | support |
|--------------|-----------|--------|----------|---------|
| 0 | 0.99 | 0.99 | 0.99 | 72 |
| 1 | 0.98 | 0.98 | 0.98 | 42 |
| accuracy | | | 0.98 | 114 |
| macro avg | 0.98 | 0.98 | 0.98 | 114 |
| weighted avg | 0.98 | 0.98 | 0.98 | 114 |

Random Forest Classification

ROC-AUC: 0.9953703703703705

| | precision | recall | f1-score | support |
|--------------|-----------|--------|----------|---------|
| 0 | 0.99 | 0.99 | 0.99 | 72 |
| 1 | 0.98 | 0.98 | 0.98 | 42 |
| accuracy | | | 0.98 | 114 |
| macro avg | 0.98 | 0.98 | 0.98 | 114 |
| weighted avg | 0.98 | 0.98 | 0.98 | 114 |

KNN

ROC-AUC: 0.982308201058201

| | precision | recall | f1-score | support |
|--------------|-----------|--------|----------|---------|
| 0 | 0.95 | 0.99 | 0.97 | 72 |
| 1 | 0.97 | 0.90 | 0.94 | 42 |
| accuracy | | | 0.96 | 114 |
| macro avg | 0.96 | 0.95 | 0.95 | 114 |
| weighted avg | 0.96 | 0.96 | 0.96 | 114 |

Final Project Conclusion

Summary of Findings

- The dataset is **clean, well-structured**, and **information-rich**, with strong signal present in size- and shape-related features.
- Exploratory Data Analysis revealed:
 - Clear but overlapping separation between classes
 - Strong multicollinearity among related features
- No data quality issues requiring outlier removal
- Multiple baseline models were evaluated using a consistent preprocessing and evaluation framework.

Model Performance Overview

- **Logistic Regression** achieved strong performance with **high ROC-AUC** and **good malignant recall**, serving as a reliable and interpretable baseline.
- **Support Vector Machine (RBF)** and **Random Forest** slightly outperformed Logistic Regression, **achieving near-perfect ROC-AUC** and **higher malignant recall**.
- **KNN underperformed relative to other models**, particularly in malignant recall, and was therefore not considered a final candidate.

Across all strong models, **ROC-AUC scores** were already near saturation (**~0.995**), and differences in accuracy and recall were marginal.

Final Model Consideration

Logistic Regression remains valuable for **interpretability** and **transparency**.

SVM (RBF) and **Random Forest** provide the best balance of **performance** and **robustness**.

Given the negligible performance differences, **model choice can be guided by deployment constraints (interpretability vs flexibility)** rather than raw metrics alone.