Supervised_learning_final_project

December 9, 2024

0.1 Supervised Learning Final Project

This is a fairly simple and straight forward project focusing on genetics data. The question we start with is given a set of prostate cancer patients, can we classify them into tumor or normal tissue based on the genes that are involved with prostate cancer?

For this we will start with the data downloading it from the Cancer Genome Atlas (https://portal.gdc.cancer.gov/). This is a bit of a task to get the data downloaded correctly and put in the right place. Mainly, it is a bunch of UI manipulation to get the data downloaded the way that you want. There are numerous tutorials out there that can show you how to get this information and restrict it to the proper fields.

TCGA doesn't exactly have normal tissue samples within it's database. Rather, it has surrounding tissue that has been tested to use in the place of missing normal tissue. This can complicate some analyses, but for this purpose we don't care.

0.2 Exploratory Data Analysis

```
[141]: # First step is to get some basic information

print("Normal DataFrame Shape:", normal_df.shape)

print("Tumor DataFrame Shape:", tumor_df.shape)

# Check for missing values

print("\nMissing Values in Normal DataFrame:", normal_df.isnull().sum().sum())

print("Missing Values in Tumor DataFrame:", tumor_df.isnull().sum().sum())

# Check max and average values

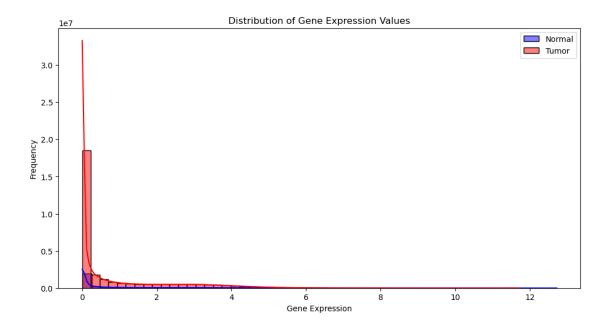
print("\nMax Values in Normal DataFrame:", normal_df.max())

print("Max Values in Tumor DataFrame:", tumor_df.max())
```

```
# Check real max value
print("\nSorted Max Values in Normal DataFrame:", normal_df.max().
  ⇔sort_values(ascending=False))
print("Sorted Max Values in Tumor DataFrame:", tumor_df.max().
  ⇔sort_values(ascending=False))
Normal DataFrame Shape: (52, 60660)
Tumor DataFrame Shape: (497, 60660)
Missing Values in Normal DataFrame: 0
Missing Values in Tumor DataFrame: 0
Max Values in Normal DataFrame: GeneSymbol
TSPAN6
              111.8758
TNMD
               52.5824
DPM1
              162.8129
SCYL3
               17.3148
Clorf112
                4.1748
AC008763.4
                0.0249
AL592295.6
               24.3267
AC006486.3
                0.0000
AL391628.1
                0.1797
AP006621.6
                1.8073
Length: 60660, dtype: float64
Max Values in Tumor DataFrame: GeneSymbol
TSPAN6
              136.6281
TNMD
                6.1874
DPM1
              174.0162
SCYL3
               22.6370
Clorf112
                8.9708
AC008763.4
                0.0581
AL592295.6
               22.7874
AC006486.3
                0.0000
AL391628.1
                0.8144
AP006621.6
                6.2973
Length: 60660, dtype: float64
Sorted Max Values in Normal DataFrame: GeneSymbol
SEMG1
              336058.1856
SEMG2
              135728.9928
MT-ND4
               63965.1804
MSMB
               58958.3913
NPY
               56558.0827
                   0.0000
OR6C76
```

```
ZNF593
                         0.0000
      Y_RNA
                         0.0000
      RNU6-449P
                         0.0000
      AL353997.3
                         0.0000
      Length: 60660, dtype: float64
      Sorted Max Values in Tumor DataFrame: GeneSymbol
      MT-RNR2
                    120956.7202
                    102051.7722
      IGKC
      MT-ND4
                    92665.8190
      MT-CO3
                     91600.2711
      MT-CO2
                     86275.2984
      AL133410.3
                         0.0000
                         0.0000
      AC106886.5
                         0.0000
      AC008581.1
      AC109486.3
                         0.0000
      AC098484.1
                         0.0000
      Length: 60660, dtype: float64
[142]: # Gene expression can have wide range of values. 120956 is too big so let's
        ⇔try to log transform this.
       import numpy as np
       normal_df = np.log(normal_df + 1)
       tumor_df = np.log(tumor_df + 1)
       # Check real max value
       print("\nSorted Log Values in Normal DataFrame:", normal_df.max().
        ⇒sort_values(ascending=False))
       print("Sorted Log Values in Tumor DataFrame:", tumor_df.max().
        ⇔sort_values(ascending=False))
      Sorted Log Values in Normal DataFrame: GeneSymbol
      SEMG1
                    12.725043
      SEMG2
                    11.818423
      MT-ND4
                    11.066110
      MSMB
                    10.984604
      NPY
                    10.943041
      OR6C76
                     0.000000
      ZNF593
                     0.000000
      Y RNA
                     0.00000
      RNU6-449P
                     0.000000
      AL353997.3
                     0.000000
      Length: 60660, dtype: float64
```

```
Sorted Log Values in Tumor DataFrame: GeneSymbol
      MT-RNR2
                    11.703196
      IGKC
                    11.533245
      MT-ND4
                    11.436766
      MT-CO3
                    11.425200
      MT-CO2
                    11.365310
      AL133410.3
                     0.000000
      AC106886.5
                     0.00000
      AC008581.1
                     0.000000
      AC109486.3
                     0.000000
      AC098484.1
                     0.000000
      Length: 60660, dtype: float64
[143]: # Check the distribution of Gene Expression Values
       # Let's subset. 60K values takes a long time on a simple machine. Let's look_{\sqcup}
        ⇔at the first 1000 genes
       normal_df_1K = normal_df.iloc[:1000]
       tumor_df_1K = tumor_df.iloc[:1000]
       plt.figure(figsize=(12, 6))
       sns.histplot(normal_df_1K.values.flatten(), bins=50, color='blue',_
        ⇔label='Normal', kde=True)
       sns.histplot(tumor_df_1K.values.flatten(), bins=50, color='red', label='Tumor',_
        →kde=True)
       plt.title('Distribution of Gene Expression Values')
       plt.xlabel('Gene Expression')
       plt.ylabel('Frequency')
       plt.legend()
      plt.show()
      C:\Users\dvanbooven\AppData\Local\anaconda3\lib\site-
      packages\seaborn\_oldcore.py:1119: FutureWarning: use_inf_as_na option is
      deprecated and will be removed in a future version. Convert inf values to NaN
      before operating instead.
        with pd.option_context('mode.use_inf_as_na', True):
      C:\Users\dvanbooven\AppData\Local\anaconda3\lib\site-
      packages\seaborn\ oldcore.py:1119: FutureWarning: use inf as na option is
      deprecated and will be removed in a future version. Convert inf values to NaN
      before operating instead.
        with pd.option_context('mode.use_inf_as_na', True):
```



```
[144]: # That's an awful lot of O's. Let's remove them.

# Add the label column to indicate normal (0) and tumor (1) status
normal_df['Label'] = 0 # Normal samples are labeled as 0
tumor_df['Label'] = 1 # Tumor samples are labeled as 1

# Combine the two DataFrames (normal and tumor)
combined_df = pd.concat([normal_df, tumor_df], axis=0)

filtered_df = combined_df.loc[:, combined_df.sum() > 100]

print("Combined DataFrame Shape:", filtered_df.shape)
```

Combined DataFrame Shape: (549, 25677)

```
[145]: # Let's check a few prostate cancer important genes expression ...

columns_to_keep = ['AR', 'KLK3', 'PTEN', 'TP53', 'ERG']

filtered_df_2 = filtered_df[columns_to_keep]

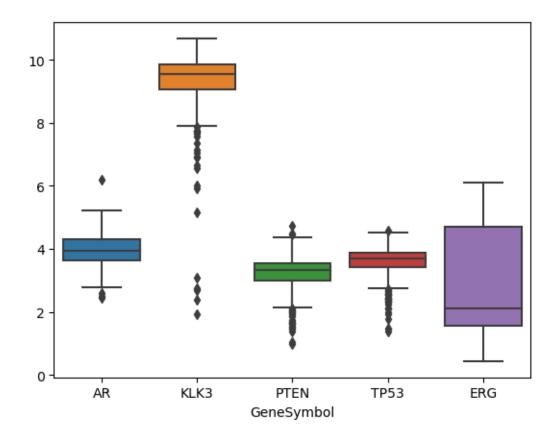
print(filtered_df_2.describe())

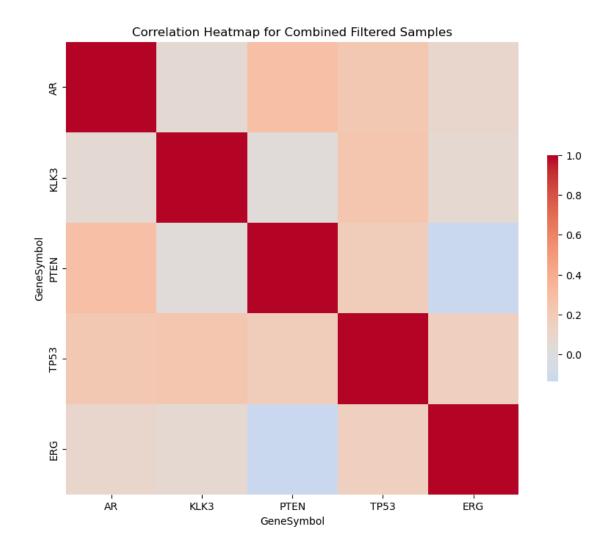
sns.boxplot(data=filtered_df_2)
```

GeneSymbol AR KLK3 PTEN TP53 ERG count 549.000000 549.000000 549.000000 549.000000

mean	3.967581	9.316864	3.222247	3.609323	2.900050
std	0.488126	0.990835	0.527577	0.471165	1.626492
min	2.452401	1.928590	0.994288	1.369275	0.437739
25%	3.632254	9.051101	2.983123	3.427836	1.573023
50%	3.954383	9.536852	3.322785	3.683558	2.123219
75%	4.313389	9.840564	3.554465	3.882431	4.697529
max	6.204963	10.678389	4.718860	4.582873	6.102934

[145]: <Axes: xlabel='GeneSymbol'>





0.3 Model Generation

For this project I've selected to use a logistic regression model. This fits in perfectly with the categorical yes/no output, but in this case it's an evaluation of tumor vs normal sample.

```
# Standardize the feature values (important for logistic regression)
scaler = StandardScaler()
X_train_scaled = scaler.fit_transform(X_train)
X_test_scaled = scaler.transform(X_test)

# Initialize the logistic regression model
logreg = LogisticRegression(max_iter=10000)

# Train the model
logreg.fit(X_train_scaled, y_train)

# Predict on the test set
y_pred = logreg.predict(X_test_scaled)
```

0.4 Model Evaluation

```
[148]: # Evaluate the model
    accuracy = accuracy_score(y_test, y_pred)
    conf_matrix = confusion_matrix(y_test, y_pred)
    class_report = classification_report(y_test, y_pred)

# Results
    print(f"Accuracy: {accuracy}")
    print(f"Confusion Matrix:\n{conf_matrix}")
    print(f"Classification Report:\n{class_report}")

cm = confusion_matrix(y_test, y_pred)
    sns.heatmap(cm, annot=True, fmt="d", cmap="Blues", xticklabels=["Pred: 0", u"Pred: 1"], yticklabels=["True: 0", "True: 1"])
    plt.title("Confusion Matrix")
    plt.show()
```

Accuracy: 0.94545454545454

Confusion Matrix:

[[7 3] [3 97]]

Classification Report:

	precision	recall	f1-score	support
0	0.70	0.70	0.70	10
1	0.97	0.97	0.97	100
accuracy			0.95	110
macro avg	0.83	0.83	0.83	110
weighted avg	0.95	0.95	0.95	110

