Breast Cancer Jump Scare Team 5

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Agenda

This study aims to create models to determine overall survivability and predict clinical data for clinicians

About the Data	Data sources, time range, size, and end to end data flow from source to consumption with components and connectors describing how data evolves through the flow to generate end visuals and for modelling
Exploratory Analysis 1	Modeling Experiment, Evaluation
Exploratory Analysis 2	Modeling Experiment, Evaluation
Exploratory Analysis 3	Modeling Experiment, Evaluation
Considerations & Key Learning	Discussing our key learnings, future implementations, and our key learnings

About the Data

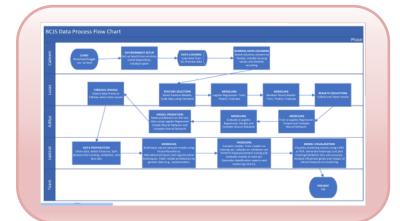


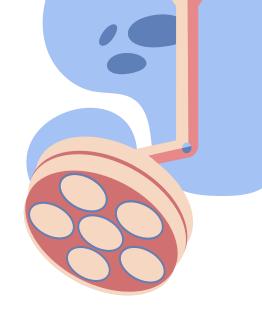
Data Description

Sourced from <u>Kaggle</u>. Data is 1904 Rows x 676 Columns. Data is taken between 2016 to 2020



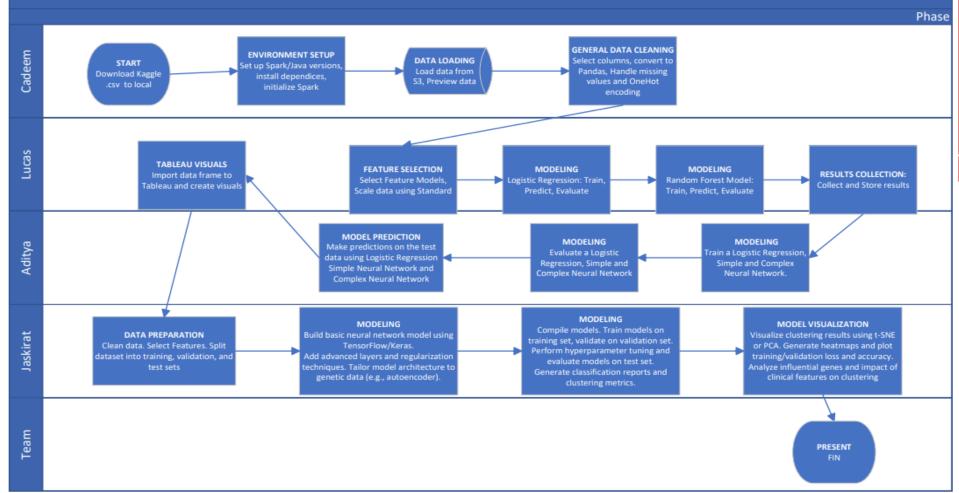
Data Process







BCJS Data Process Flow Chart



Key Insights on Target Variables

Target variable 'overall_survival' is binary. Indicates whether a patient survived over a period of time.

Modeling Experiment Design

Baseline model: Logistic Regression Model.

Random Survival Forest (RSF) Model employed to handle survivability over time (time-to-event data)

Model Evaluation

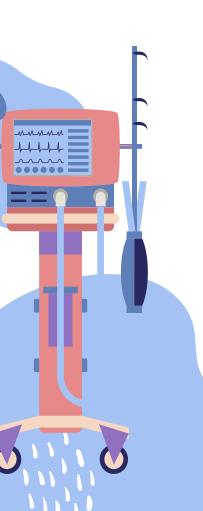
Metrics Used

- Accuracy: Primary metric in Logistic Regression
- Survival Probability Metric: RSF Model different time points (ex. 365 days)

Model Comparison

The RSF model of survival data shows better generalization and performance compared to the baseline logistic regression model.





Logistic Regression Model Accuracy

```
print("Classification Report")
    print(classification_report(y_test, y_pred))

→ Classification Report

                              recall f1-score support
                 precision
            0.0
                      0.67
                                0.65
                                          0.66
                                                    124
            1.0
                      0.56
                                0.58
                                          0.57
                                                     95
        accuracy
                                          0.62
                                                    219
       macro avg
                                          0.61
                                                    219
                      0.61
                                0.61
    weighted avg
                      0.62
                                0.62
                                          0.62
                                                    219
```



Random Survival Forest Predictions

pat	ient_id	survival probability at 180 days	survival probability at 365 days	survival probability at 545 days	survival probability at 730 days
0	0	0.998571	0.879471	0.216091	0.085850
1	1	0.987496	0.921208	0.811910	0.128509
2	2	0.999470	0.987444	0.764962	0.325275
3	3	1.000000	0.983698	0.939805	0.483045
4	4	0.967509	0.829850	0.495934	0.141528

Key Insights

Type of cancer and type of treatment, which treatment is more effective against certain types of cancers.

Model Design

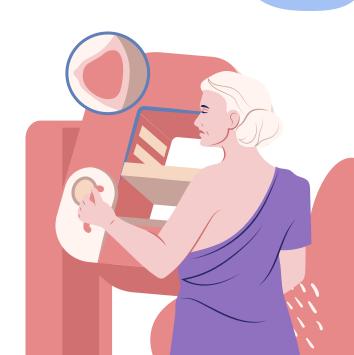
- Created Dummys for testing predictions.
- Ran Logistics Regression as the measurement is categorical

Model evaluation

- Ran with different epochs, neurons, and layers.
- To achieve 56% accuracy.



Tableau Visuals



Model 1

```
# Output layer has 8 neurons, one for each treatment type
model1 = models.Sequential([
    layers.Dense(6A, activation='relu', input_shape=(X_train.shape[1],)),
    layers.Dense(8A, activation='relu'),
    layers.Dense(8A, activation='softmax')
])
# Check the structure of the model
model1.summary()

/ /usr/local/lib/python3.10/dist-packages/keras/src/layers/core/dense.py:87: Userwarning: Do not pass an `input_shape`/`input_dim` argument to
    super().__init__(activity_regularizer=activity_regularizer, **Rwargs)
```

Layer (type)	Output Shape	Param #
dense (Dense)	(None, 64)	4,224
dense_1 (Dense)	(None, 32)	2,080
dense_2 (Dense)	(None, 8)	264

Total params: 6,568 (25.66 KB) Trainable params: 6,568 (25.66 KB) Non-trainable params: 0 (0.00 B)

Model 2

```
model2 = models.Sequential([
    layers.Dense(128, activation='relu', input_shape=(X_train.shape[1],)),
    layers.Dense(04, activation='relu'),
    layers.Dense(04, activation='relu'),
    layers.Dense(32, activation='relu'),
    layers.Dense(32, activation='relu'),
    layers.Dense(32, activation='softmax') # Output layer for 8 classes
])
# Check the structure of the model
model2.summary()
```

Model: "sequential_1"

Layer (type)	Output Shape	Param #
dense_3 (Dense)	(None, 128)	8,448
dropout (Dropout)	(None, 128)	0
dense_4 (Dense)	(None, 64)	8,256
dropout_1 (Dropout)	(None, 64)	0
dense_5 (Dense)	(None, 32)	2,080
dense_6 (Dense)	(None, 8)	264

Total params: 19,048 (74.41 KB)
Trainable params: 19,048 (74.41 KB)
Non-trainable params: 0 (0.00 B)

Training Model with 100 epoch

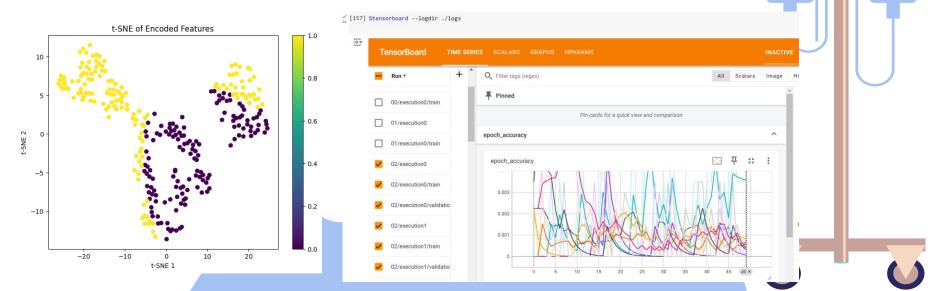
```
model2.compile(optimizer='adam', loss='sparse categorical crossentropy', metrics=['accuracy'])
# Train the model
model2.fit(X train, y train, epochs=100, batch size=64, validation data=(X test, y test))
20/20 -
                          - 0s 5ms/step - accuracy: 0.7930 - loss: 0.5112 - val accuracy: 0.5643 - val loss: 1.4376
Epoch 73/100
                          - 0s 5ms/step - accuracy: 0.8047 - loss: 0.5278 - val accuracy: 0.5486 - val loss: 1.4270
20/20 .
Epoch 74/100
20/20 -
                          - 0s 5ms/step - accuracy: 0.7848 - loss: 0.5254 - val accuracy: 0.5643 - val loss: 1.4384
Epoch 75/100
20/20 -
                          - 0s 4ms/step - accuracy: 0.7992 - loss: 0.4972 - val_accuracy: 0.5580 - val_loss: 1.4372
Epoch 76/100
                          - 0s 5ms/step - accuracy: 0.7997 - loss: 0.5350 - val accuracy: 0.5799 - val loss: 1.4075
20/20 -
Epoch 77/100
20/20 .
                          - 0s 9ms/step - accuracy: 0.7959 - loss: 0.5370 - val_accuracy: 0.5549 - val_loss: 1.4267
Epoch 78/100
20/20 -
                          — 0s 8ms/step - accuracy: 0.7809 - loss: 0.5714 - val_accuracy: 0.5674 - val_loss: 1.4606
```



Key Insights

Metrics Used: Loss and Accuracy for models and clustering metrics (silhouette score and Davies Bouldin score

Use of TensorBoard for following Model training.



Considerations & Key Lessons

Time

Horizon of the study Time constrain on the project time line

Scope

11 Models in 2 weeks

Domain Knowledge

Domain expertise in the medical industry. This model can help clinicians deal with better Prognosis for patiensts

