

**One Year Life Expectancy Post Thoracic Surgery Using Machine Learning**

[DOCUMENT TITLE]

**PROFESSIONAL TRAINING REPORT – 2**

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# **INTRODUCTION**

## **1.1.overview**

Cancer is a disease in which cells in the body grow out of control and is one of the most serious health problems in the world. Among various different types of cancer, lung cancer is one of the leading causes of death in both men and women. According to World Health Organization, it was observed that in the year 2018 2.09 million cases of lung cancer was registered and a total of 1.76 million died due to lung cancer. One of the reasons for the high death rate due to lung cancer is the late detection of the lung cancer. Also, the treatment and the prognosis depend on the type of the lung cancer, the stage and the patient's performance. Once the lung cancer is detected, possible treatments include thoracic surgery, chemotherapy and radiotherapy

## **1.2 purpose**

the United States, lung cancer claims more lives every year than colon cancer Lung cancer is the leading cause of cancer-related deaths in the world. In, prostate cancer, and breast cancer combined.

Despite the very serious prognosis (outlook) of lung cancer, some people with earlier-stage cancers are cured. More than 430,000 people alive today have been diagnosed with lung cancer at some point. The data is dedicated to classification problems related to the post-operative life expectancy in lung cancer patients: class 1 - death within one year after surgery, class 2 - survival.

We will be using classification algorithms such as Decision tree, Random forest, KNN, and xgboost. We will train and test the data with these algorithms. From this best model is selected and saved in pkl format. We will be doing flask integration and IBM deployment.

# **2.LITERATURE SURVEY**

## **2.1 Existing problem**

Disha Sharma et al. (2011), proposed an approach for the early detection of lung cancer by analyzing lungs CT images using Image Processing techniques[1]. The authors used bit-plane slicing, erosion and Weiner filter image processing techniques to extract the lung regions from the CT image. Further the extracted lung regions were segmented using Region growing segmentation algorithm and later Rule based model was used to detect the cancerous nodules. With the help of diagnostics indicator, it was observed that the proposed method achieved an overall accuracy of 80%. Hamid Bagherieh et al. (2013) gave a methodology to detect and classify the lung nodules using Image processing and Decision-Making techniques[2]. Initially, image preprocessing was carried out on a CT images by using contrast enhancement and linear filtering. Next, the filtered image was segmented using Region growing Segmentation process. Further the features like area and color was

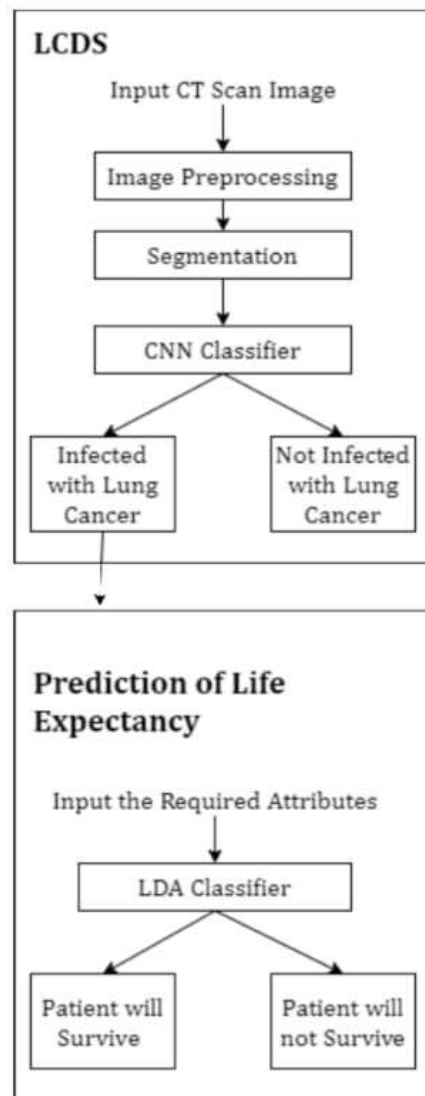
given as input to the Fuzzy system which employed fuzzy membership function to find the abnormalities. Sindhu V et al.(2014), proposed an approach where the authors aimed to classify the survival of lung cancer patients post thoracic surgery[3]. The authors used Naïve Bayes, PART, J48, OneR, Random Forest and Decision stump algorithm techniques to classify the target function. The performance measurement shows that Random Forest gave high accuracy of 95.65% compared to other ML techniques used. Prashant Naresh et al.(2014), proposed a methodology to detect the lung cancer using Image processing and Neural Techniques[4]. Initially the CT image of lung was filtered to remove Gaussian white noise and Otsu's threshold technique was used to do the segmentation of the image. The structural and features were extracted and these features were given as input to the classifier. The SVM and ANN techniques were used for the classification and it was found that SVM techniques gave a higher accuracy of 95.12%. Kwetishe Joro Danjuma (2015), proposed a methodology to predict the one-year survival of the patient post thoracic surgery[5]. Naïve Bayes, J48 and Multilayer Perceptron algorithms were used to classify the target class. The Naïve Bayes gave an accuracy of 74.4%, J48 gave an accuracy of 81.8% and MLP gave an accuracy of 82.4%.

## **2.1 proposed solution**

A system is developed which detects the lung cancer from the given input CT scanned lung images which are in DICOM (.dcm) format. In addition to this, the system also helps to predict the Life Expectancy Post Thoracic Surgery of the Lung Cancer infected patients.

### 3.THEORITICAL ANALYSIS

#### 3.1 Block diagram



**Fig-1:** Block Diagram of the System

### 3.2 Hardware/software designing

### \* Software Requirements

Processor : intel core i3

\* Operating System :windows

Speed : 1.19GHz

\* Coding Language:Python3.7

RAM : 8.0GB

Memory usage : 7.2kb

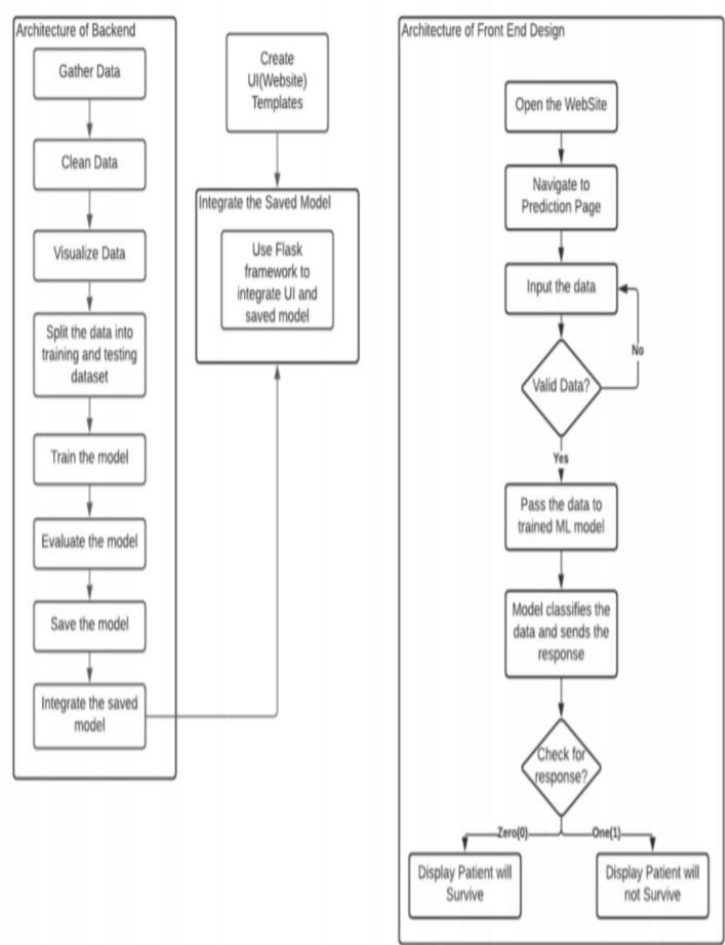
Keyboard : Standard keyboard

Monitor : 15 VGA color


## 4.EXPERIMENTAL INVESTIGATIONS

This is the second part of the system which aims at predicting the survival of lung cancer infected patient post thoracic surgery. The dataset includes 17 attributes which are specified in Table-1. Among all those attributes, Risk1Y is the target class specifying zero if the patient survives for at least one-year post thoracic surgery and one for those who died before completing one-year post surgery. The visualization of the dataset is done using the Matplotlib and Seaborn libraries of Python. Further the essential attributes are found based on the Information Gain (IG) attribute evaluation which is used to find the importance of an attribute by using the Information Gain with respect to the target class.  $IG (Class, Attribute) = E (Class) - E (Class | Attribute)$  where, E stands for Entropy After the IG Attribute Evaluation on all the 16 independent attributes in the dataset, it is found that the attributes PRE19 and PRE32 gives the IG value as zero and hence are the least useful attributes for training the model. Therefore, these two attributes are eliminated and the remaining 14 attributes are used to train the model

5. FLOWCHART



6.RESULT

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In [1]:


```
import pandas as pd
import numpy as np
import seaborn as sns
from sklearn.model_selection import train_test_split
from sklearn.linear_model import LogisticRegression
from sklearn.metrics import accuracy_score
import matplotlib.pyplot as plt
from sklearn.linear_model import LogisticRegression
from sklearn.ensemble import RandomForestClassifier, GradientBoostingClassifier
from sklearn.tree import DecisionTreeClassifier
from sklearn.neighbors import KNeighborsClassifier
from sklearn.metrics import f1_score
from sklearn.metrics import classification_report, confusion_matrix
import itertools
```

In [2]: df=pd.read\_csv("ThoracicSurgery.csv")

In [3]: df.head()

Out[3]:

	Diagnosis	FVC	FEV1	Performance	Pain	Haemoptysis	Dyspnoea	Cough	Weakness	Tumor_Size	Diabetes_Mellitus	MI_6mo	PAD	Smoking	Asthma	Age
0	2	2.88	2.16	1	0	0	0	1	1	4	0	0	0	1	0	6
1	3	3.40	1.88	0	0	0	0	0	0	2	0	0	0	1	0	8
2	3	2.76	2.08	1	0	0	0	1	0	1	0	0	0	1	0	8

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In [2]: df=pd.read\_csv("ThoracicSurgery.csv")


In [3]: df.head()

Out[3]:

	Diagnosis	FVC	FEV1	Performance	Pain	Haemoptysis	Dyspnoea	Cough	Weakness	Tumor_Size	Diabetes_Mellitus	MI_6mo	PAD	Smoking	Asthma	Age
0	2	2.88	2.16	1	0	0	0	1	1	4	0	0	0	1	0	6
1	3	3.40	1.88	0	0	0	0	0	0	2	0	0	0	1	0	8
2	3	2.76	2.08	1	0	0	0	1	0	1	0	0	0	1	0	8
3	3	3.68	3.04	0	0	0	0	0	0	1	0	0	0	0	0	8
4	3	2.44	0.96	2	0	1	0	1	1	1	0	0	0	1	0	7

In [4]: df.columns

Out[4]: Index(['Diagnosis', 'FVC', 'FEV1', 'Performance', 'Pain', 'Haemoptysis', 'Dyspnoea', 'Cough', 'Weakness', 'Tumor\_Size', 'Diabetes\_Mellitus', 'MI\_6mo', 'PAD', 'Smoking', 'Asthma', 'Age', 'Death\_1yr'], dtype='object')

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In [5]: df.describe()

Out[5]:

	Diagnosis	FVC	FEV1	Performance	Pain	Haemoptysis	Dyspnoea	Cough	Weakness	Tumor_Size	Diabetes_Mellitus	MI_6m
count	454.000000	454.000000	454.000000	454.000000	454.000000	454.000000	454.000000	454.000000	454.000000	454.000000	454.000000	454.000000
mean	3.092511	3.287952	2.51685	0.795154	0.059471	0.136564	0.055066	0.696035	0.171806	1.733480	0.074890	0.00441
std	0.715817	0.872347	0.77189	0.531459	0.236766	0.343765	0.228361	0.460475	0.377628	0.707499	0.263504	0.06628
min	1.000000	1.440000	0.96000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	1.000000	0.000000	0.00000
25%	3.000000	2.600000	1.96000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	1.000000	0.000000	0.00000
50%	3.000000	3.160000	2.36000	1.000000	0.000000	0.000000	0.000000	1.000000	0.000000	2.000000	0.000000	0.00000
75%	3.000000	3.840000	2.97750	1.000000	0.000000	0.000000	0.000000	1.000000	0.000000	2.000000	0.000000	0.00000
max	8.000000	6.300000	5.48000	2.000000	1.000000	1.000000	1.000000	1.000000	1.000000	4.000000	1.000000	1.00000

In [6]: df.shape

Out[6]: (454, 17)

In [7]: df.info()

<class 'pandas.core.frame.DataFrame'>  
RangeIndex: 454 entries, 0 to 453  
Data columns (total 17 columns):

```
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In [7]: df.info()

<class 'pandas.core.frame.DataFrame'>
RangeIndex: 454 entries, 0 to 453
Data columns (total 17 columns):
#   Column              Non-Null Count  Dtype  
---  --
0   Diagnosis            454 non-null    int64   
1   FVC                  454 non-null    float64  
2   FEV1                 454 non-null    float64  
3   Performance          454 non-null    int64   
4   Pain                 454 non-null    int64   
5   Haemoptysis          454 non-null    int64   
6   Dyspnoea             454 non-null    int64   
7   Cough                454 non-null    int64   
8   Weakness             454 non-null    int64   
9   Tumor_Size           454 non-null    int64   
10  Diabetes_Mellitus     454 non-null    int64   
11  MI_6mo               454 non-null    int64   
12  PAD                  454 non-null    int64   
13  Smoking              454 non-null    int64   
14  Asthma               454 non-null    int64   
15  Age                  454 non-null    int64   
16  Death_1yr            454 non-null    int64   
dtypes: float64(2), int64(15)
memory usage: 60.4 KB
```

```
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In [8]: df.isnull().sum()

Out[8]: Diagnosis      0
FVC      0
FEV1      0
Performance  0
Pain      0
Haemoptysis  0
Dyspnoea  0
Cough      0
Weakness   0
Tumor_Size  0
Diabetes_Mellitus  0
MI_6mo     0
PAD        0
Smoking     0
Asthma      0
Age         0
Death_1yr   0
dtype: int64

In [9]: live = df[df['Death_1yr'] == 0]
death = df[df['Death_1yr'] == 1]

cond = ['FVC', 'FEV1', 'Performance', 'Pain', 'Haemoptysis', 'Dyspnoea', 'Cough', 'Weakness', \
        'Tumor_Size', 'Diabetes_Mellitus', 'MI_6mo', 'PAD', 'Smoking', 'Asthma', 'Age']

l = [np.mean(live[c]) for c in cond]
d = [np.mean(death[c]) for c in cond]

ld = pd.DataFrame(data={'Attribute': cond, 'Live 1yr Mean': l, 'Death 1yr Mean': d})
ld = ld.set_index('Attribute')

print('Death: {d}, Live: {l}'.format(len(death), len(live)))
print("1 year death: {:.2f}% out of 454 patients".format(np.mean(df.Death_1yr)*100))
ld

Death: 69, Live: 385
1 year death: 15.20% out of 454 patients

Out[9]:
```

Attribute	Live 1yr Mean	Death 1yr Mean
FVC	1	3.195072
FEV1	1	2.383188
Performance	1	0.913043



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Out[9]:

Attribute	Live 1yr Mean	Death 1yr Mean
FVC	1	3.195072
FEV1	1	2.383168
Performance	1	0.913043
Pain	1	0.101449
Haemoptysis	1	0.202899
Dyspnoea	1	0.115942
Cough	1	0.797101
Weakness	1	0.246377
Tumor_Size	1	2.014493
Diabetes_Mellitus	1	0.144928
MI_6mo	1	0.000000
PAD	1	0.028986
Smoking	1	0.898551
Asthma	1	0.000000
Age	1	63.333333

In [10]:

```

d = np.array(d)
l = np.array(l)

p_diff = (d-l)/l*100

fig, axes = plt.subplots(2,1,figsize=(12,18))

axes[0].bar(cond, p_diff)
axes[0].set_title('Mean Difference % between Dead and Live 1yr', fontsize=18)
axes[0].set_xticks(cond)
axes[0].set_xticklabels(cond, rotation=90)
axes[0].set_ylabel('Percent', fontsize=13)

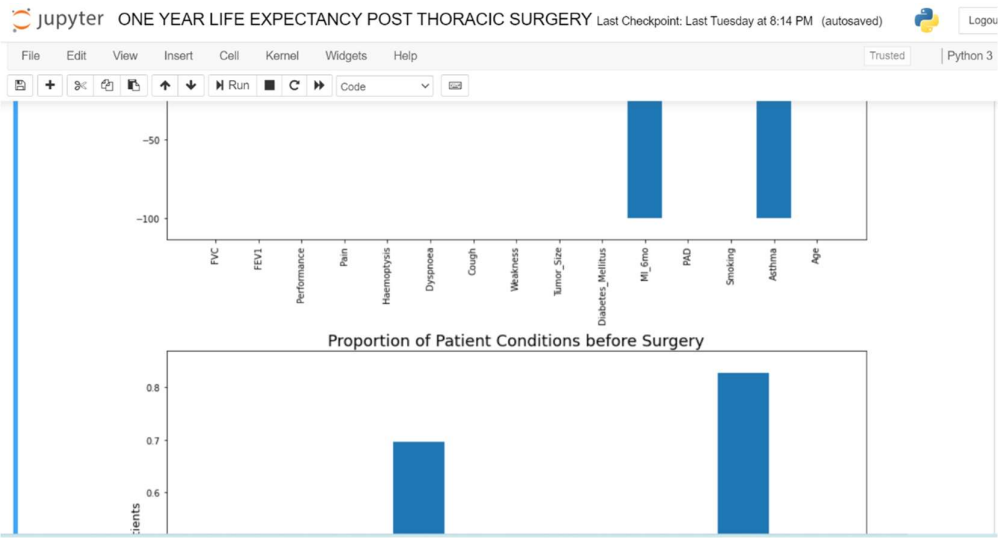
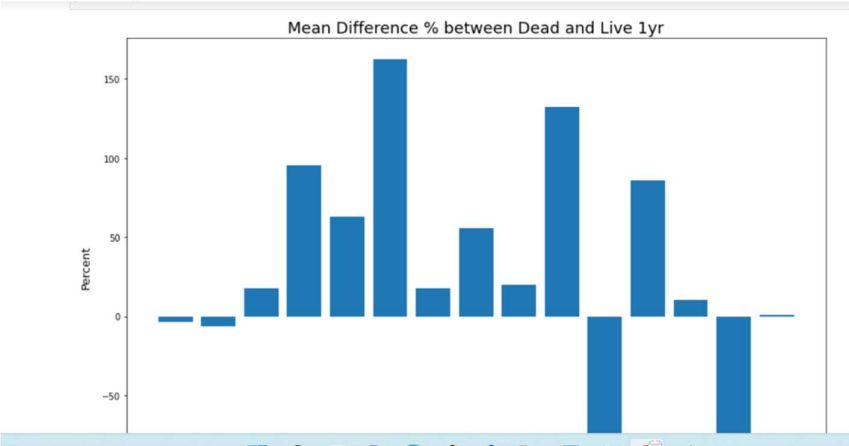
tf_col = ['Pain', 'Haemoptysis', 'Dyspnoea', 'Cough', 'Weakness', 'Diabetes_Mellitus', 'MI_6mo', 'PAD', 'Smoking', 'Asthma']
tf_sum = [df[col].sum()/454 for col in tf_col]

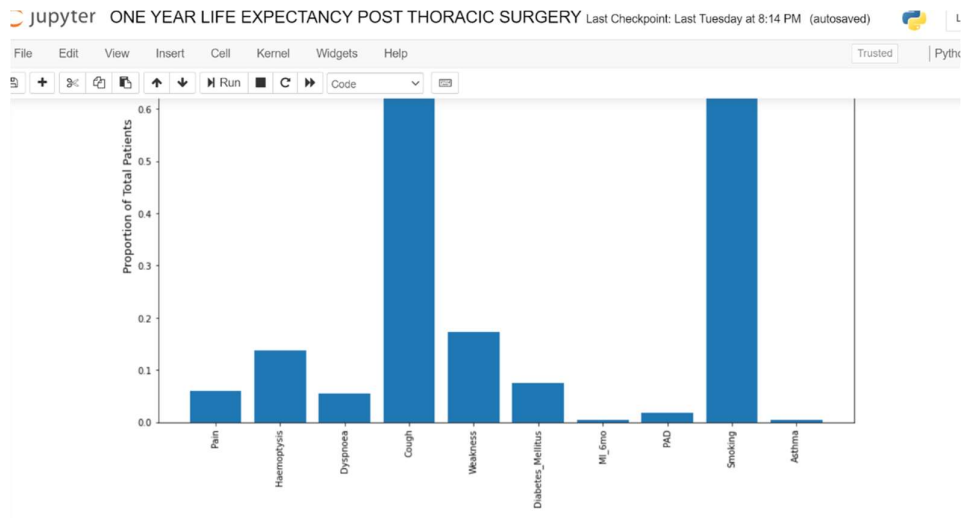
axes[1].bar(tf_col, tf_sum)
axes[1].set_xticks(tf_col)
axes[1].set_xticklabels(tf_col, rotation=90)
axes[1].set_ylabel('Proportion of Total Patients', fontsize=13)
axes[1].set_title('Proportion of Patient Conditions before Surgery', fontsize=18)

plt.tight_layout()

plt.show()

```





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```
In [12]: def permutation_sample(data1, data2):
        """Generate a permutation sample from two data sets."""
        data = np.concatenate((data1, data2))
        permuted_data = np.random.permutation(data)

        perm_sample_1 = permuted_data[:len(data1)]
        perm_sample_2 = permuted_data[len(data1):]

        return perm_sample_1, perm_sample_2

    def draw_perm_reps(data_1, data_2, func, size=1):
        """Generate multiple permutation replicates."""
        perm_replicates = np.empty(size)

        for i in range(size):
            perm_sample_1, perm_sample_2 = permutation_sample(data_1, data_2)
            perm_replicates[i] = func(perm_sample_1, perm_sample_2)

        return perm_replicates

    def diff_of_means(data_1, data_2):
        """Difference in means of two arrays."""
        diff = np.mean(data_1) - np.mean(data_2)
        return diff

In [13]: condition = ['FVC', 'FEV1', 'Performance', 'Pain', 'Haemoptysis', 'Dyspnoea', 'Cough', 'Weakness', \
                    'Tumor_Size', 'Diabetes_Mellitus', 'MI_6mo', 'PAD', 'Smoking', 'Asthma', 'Age']
        p_val = []
```

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```
In [13]: condition = ['FVC', 'FEV1', 'Performance', 'Pain', 'Haemoptysis', 'Dyspnoea', 'Cough', 'Weakness', \
                    'Tumor_Size', 'Diabetes_Mellitus', 'MI_6mo', 'PAD', 'Smoking', 'Asthma', 'Age']
        p_val = []

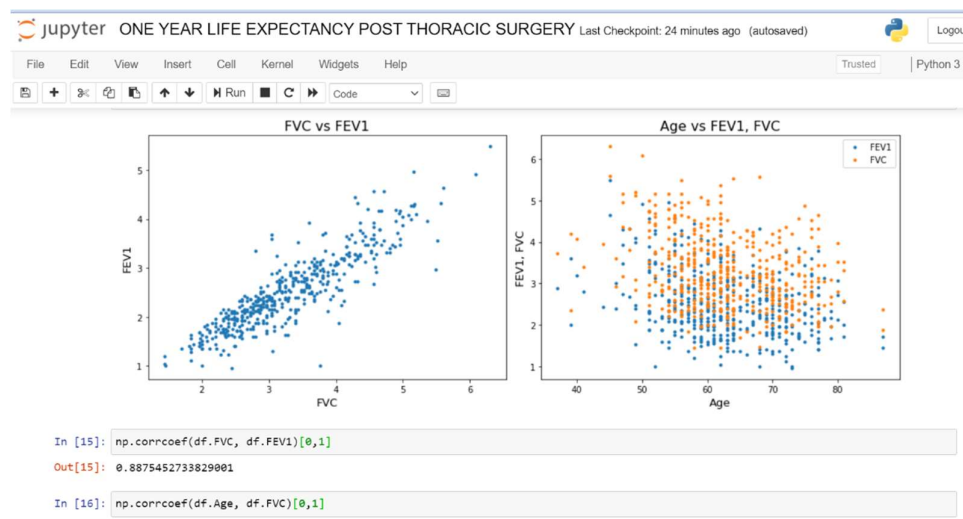
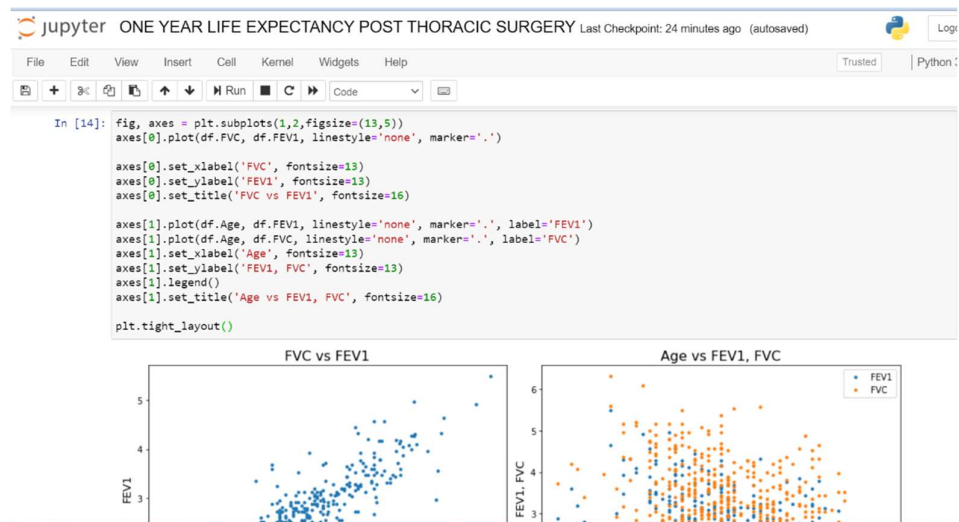
        for c in condition:
            empirical_diff_means = diff_of_means(death[c], live[c])
            perm_replicates = draw_perm_reps(death[c], live[c], diff_of_means, size=10000)
            if empirical_diff_means > 0:
                p = np.sum(perm_replicates >= empirical_diff_means) / len(perm_replicates)
                p_val.append(p)
            else:
                p = np.sum(perm_replicates <= empirical_diff_means) / len(perm_replicates)
                p_val.append(p)

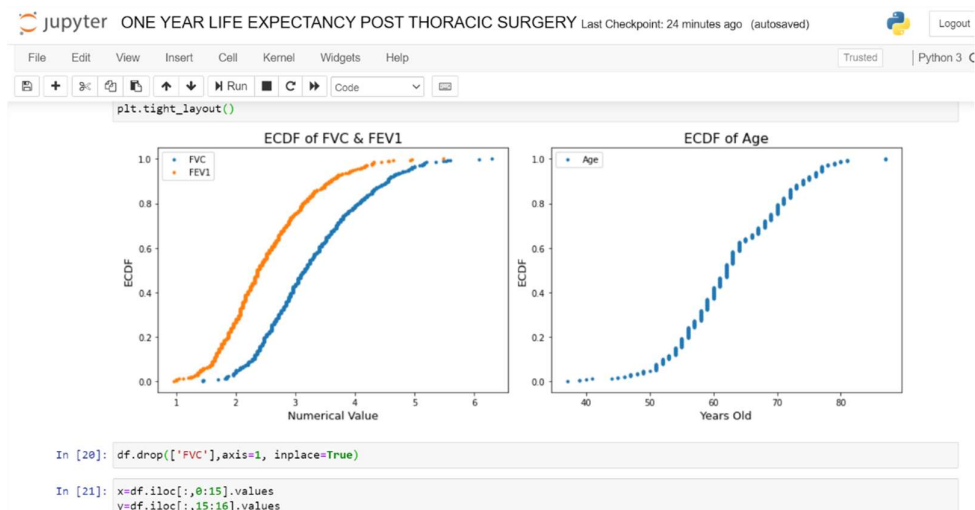
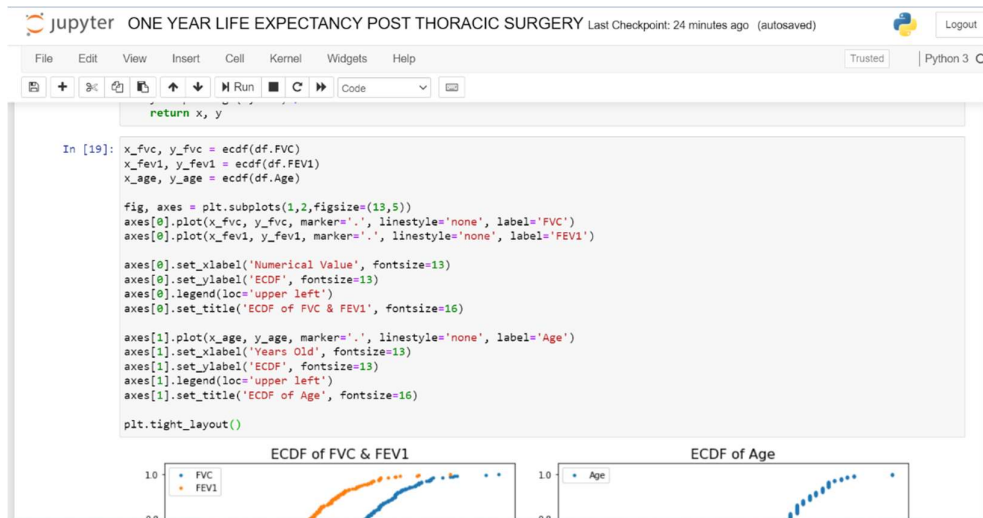
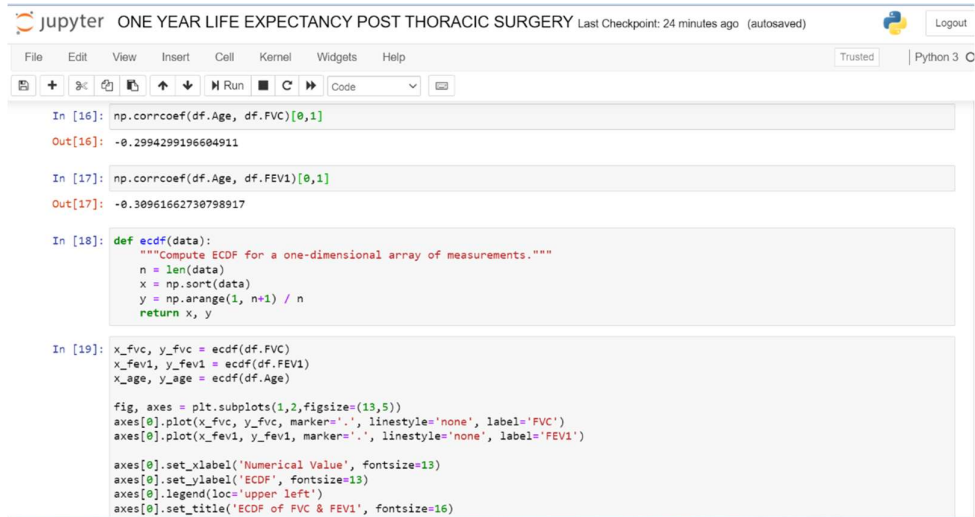
        print(list(zip(condition, p_val)))

[[('FVC', 0.1737), ('FEV1', 0.0546), ('Performance', 0.03), ('Pain', 0.0973), ('Haemoptysis', 0.0644), ('Dyspnoea', 0.0265), ('Cough', 0.0315), ('Weakness', 0.0559), ('Tumor_Size', 0.0008), ('Diabetes_Mellitus', 0.0201), ('MI_6mo', 0.7166), ('PAD', 0.3572), ('Smoking', 0.0603), ('Asthma', 0.7149), ('Age', 0.2742)]

In [14]: fig, axes = plt.subplots(1,2,figsize=(13,5))
        axes[0].plot(df.FVC, df.FEV1, linestyle='none', marker='.')

        axes[0].set_xlabel('FVC', fontsize=13)
        axes[0].set_ylabel('FEV1', fontsize=13)
        axes[0].set_title('FVC vs FEV1', fontsize=16)
```





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In [20]: `df.drop(['FVC'],axis=1, inplace=True)`

In [21]: `x=df.iloc[:,0:15].values  
y=df.iloc[:,15:16].values`

In [22]: `x_train,x_test,y_train,y_test=train_test_split(x,y,test_size=0.2,random_state=0)`

In [23]: `print('Shape of x_train {}'.format(x_train.shape))  
print('Shape of y_train {}'.format(y_train.shape))  
print('Shape of x_test {}'.format(x_test.shape))  
print('Shape of y_test {}'.format(y_test.shape))`

Shape of x\_train (363, 15)  
Shape of y\_train (363, 1)  
Shape of x\_test (91, 15)  
Shape of y\_test (91, 1)

In [24]: `def decisionTree(x_train, x_test, y_train, y_test):  
 dt=DecisionTreeClassifier()  
 dt.fit(x_train,y_train.ravel())  
 yPred = dt.predict(x_test)  
 print('***DecisionTreeClassifier***')  
 print('Confusion matrix')  
 print(confusion_matrix(y_test,yPred))  
 print('Classification report')  
 print(classification_report(y_test,yPred))`


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In [24]: `def decisionTree(x_train, x_test, y_train, y_test):  
 dt=DecisionTreeClassifier()  
 dt.fit(x_train,y_train.ravel())  
 yPred = dt.predict(x_test)  
 print('***DecisionTreeClassifier***')  
 print('Confusion matrix')  
 print(confusion_matrix(y_test,yPred))  
 print('Classification report')  
 print(classification_report(y_test,yPred))`

In [25]: `def randomForest(x_train, x_test, y_train, y_test):  
 rf = RandomForestClassifier()  
 rf.fit(x_train,y_train.ravel())  
 yPred = rf.predict(x_test)  
 print('***RandomForestClassifier***')  
 print('Confusion matrix')  
 print(confusion_matrix(y_test,yPred))  
 print('Classification report')  
 print(classification_report(y_test,yPred))`

In [26]: `def KNN(x_train, x_test, y_train, y_test):  
 knn = KNeighborsClassifier()  
 knn.fit(x_train,y_train.ravel())  
 yPred = knn.predict(x_test)  
 print('***KNeighborsClassifier***')  
 print('Confusion matrix')  
 print(confusion_matrix(y_test,yPred))`


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```
In [26]: def KNN(x_train, x_test, y_train, y_test):
        knn = KNeighborsClassifier()
        knn.fit(x_train, y_train.ravel())
        yPred = knn.predict(x_test)
        print('***KNeighborsClassifier***')
        print('Confusion matrix')
        print(confusion_matrix(y_test, yPred))
        print('Classification report')
        print(classification_report(y_test, yPred))

In [27]: def xgboost(x_train, x_test, y_train, y_test):
        xg = GradientBoostingClassifier()
        xg.fit(x_train, y_train.ravel())
        yPred = xg.predict(x_test)
        print('***GradientBoostingClassifier***')
        print('Confusion matrix')
        print(confusion_matrix(y_test, yPred))
        print('Classification report')
        print(classification_report(y_test, yPred))

In [28]: def compareModel(x_train, x_test, y_train, y_test):
        decisionTree(x_train, x_test, y_train, y_test)
        print('-'*100)
        randomForest(x_train, x_test, y_train, y_test)
        print('-'*100)
        KNN(x_train, x_test, y_train, y_test)
        print('-'*100)
        xgboost(x_train, x_test, y_train, y_test)
```

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```
In [29]: compareModel(x_train, x_test, y_train, y_test)

***DecisionTreeClassifier***
Confusion matrix
[[62 12]
 [13  4]]
Classification report
      precision    recall  f1-score   support

     0       0.83      0.84      0.83        74
     1       0.25      0.24      0.24        17

 accuracy      0.73        0.73        91
 macro avg     0.54      0.54      0.54        91
 weighted avg     0.72      0.73      0.72        91

-----
***RandomForestClassifier***
Confusion matrix
[[73  1]
 [17  0]]
Classification report
      precision    recall  f1-score   support

     0       0.81      0.99      0.89        74
     1       0.00      0.00      0.00        17

 accuracy      0.41        0.49      0.45        91
 macro avg     0.41      0.49      0.45        91
```

```
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Classification report
precision recall f1-score support
0 0.81 0.99 0.89 74
1 0.00 0.00 0.00 17
accuracy 0.80 91
macro avg 0.41 0.49 0.45 91
weighted avg 0.66 0.80 0.72 91

***KNeighborsClassifier***
Confusion matrix
[[74 0]
 [17 0]]
Classification report
precision recall f1-score support
0 0.81 1.00 0.90 74
1 0.00 0.00 0.00 17
accuracy 0.81 91
macro avg 0.41 0.50 0.45 91
weighted avg 0.66 0.81 0.73 91
```

```
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***GradientBoostingClassifier***
Confusion matrix
[[70 4]
 [17 0]]
Classification report
precision recall f1-score support
0 0.80 0.95 0.87 74
1 0.00 0.00 0.00 17
accuracy 0.77 91
macro avg 0.40 0.47 0.43 91
weighted avg 0.65 0.77 0.71 91
```



```
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In [30]: from sklearn.model_selection import cross_val_score
         rf = RandomForestClassifier()
         rf.fit(x_train,y_train.ravel())
         yPred = rf.predict(x_test)

In [31]: f1_score(yPred,y_test,average='weighted')
Out[31]: 0.8476461809795143

In [32]: cv = cross_val_score(rf,x,y.ravel(),cv=5)

In [33]: np.mean(cv)
Out[33]: 0.8414163614163614

In [34]: dt = DecisionTreeClassifier()
         dt.fit(x_train,y_train)
         yPred_dt = dt.predict(x_test)

In [35]: f1_score(yPred_dt,y_test,average='weighted')
Out[35]: 0.7358388247087242

In [36]: cv = cross_val_score(rf,x,y.ravel(),cv=5)
```

```
In [37]: np.mean(cv)
Out[37]: 0.8414163614163614

In [38]: kn = KNeighborsClassifier()
         kn.fit(x_train,y_train.ravel())
         yPred_kn = kn.predict(x_test)

In [39]: f1_score(yPred_kn,y_test,average='weighted')
Out[39]: 0.896969696969697

In [40]: cv = cross_val_score(kn,x,y.ravel(),cv=5)

In [41]: np.mean(cv)
Out[41]: 0.8326495726495727

In [42]: gb = GradientBoostingClassifier()
         gb.fit(x_train,y_train.ravel())
         yPred_gb = gb.predict(x_test)

In [43]: f1_score(yPred_gb,y_test,average='weighted')
Out[43]: 0.8313425704730053
```

```
jupyter ONE YEAR LIFE EXPECTANCY POST THORACIC SURGERY Last Checkpoint: 3 minutes ago (autosaved) Logout
File Edit View Insert Cell Kernel Widgets Help Trusted Python 3

In [44]: cv = cross_val_score(gb,x,y.ravel(),cv=5)

In [45]: np.mean(cv)
Out[45]: 0.8260073260073261

In [46]: x_test[4]
Out[46]: array([ 3. ,  3.2,  0. ,  0. ,  0. ,  0. ,  1. ,  0. ,  2. ,  0. ,  0. ,
                0. ,  1. ,  0. , 55. ])

In [47]: gb.predict([[3,4.08,0,0,0,0,1,0,2,0,0,0,1,0,55]])
Out[47]: array([0], dtype=int64)

In [48]: rf.predict([[3,4.08,0,0,0,0,1,0,2,0,0,0,1,0,55]])
Out[48]: array([0], dtype=int64)

In [49]: dt.predict([[3,4.08,0,0,0,0,1,0,2,0,0,0,1,0,55]])
Out[49]: array([0], dtype=int64)

In [50]: kn.predict([[3,4.08,0,0,0,0,1,0,2,0,0,0,1,0,55]])
Out[50]: array([0], dtype=int64)
```

```
1 import joblib
2 from flask import Flask, request, render_template
3
4 app = Flask(__name__)
5 joblib_file = "model.pkl"
6 model = joblib.load(joblib_file)
7
8
9 @app.route('/')
10 def index():
11     return render_template('index.html')
12
13 @app.route("/form", methods=['GET', 'POST'])
14 def getform():
15     if request.method == "GET":
16         return (render_template("form.html"))
17
18     if request.method == "POST":
19         if 'submit-button' in request.form:
20             diagnosis = request.form["diagnosis"]
21             fev = request.form["fev"]
22             age = request.form["age"]
23             performance = request.form["performance"]
24             tnm = request.form["tnm"]
25             hae = request.form["hae"]
26             pain = request.form["pain"]
27             dys = request.form["dys"]
28             cough = request.form["cough"]
29             weakness = request.form["weakness"]
30             dm = request.form["dm"]
31             mi = request.form["mi"]
32             pad = request.form["pad"]
33             smoking = request.form["smoking"]
34             asthma = request.form["asthma"]
```

```
Python 3.8.3 (default, Jul 2 2020, 17:30:36) [MSC v.1916 64 bit (AMD64)]
Type "copyright", "credits" or "license" for more information.

IPython 7.16.1 -- An enhanced Interactive Python.

In [1]: runfile('C:/Users/Mohansai/Downloads/5_6228888303706309736/One
Year Life expectancy post thoracic surgery using IBM watson studio/
flask/app.py', wdir='C:/Users/Mohansai/Downloads/5_6228888303706309736/
One Year Life expectancy post thoracic surgery using IBM watson studio/
flask')
* Serving Flask app "app" (lazy loading)
* Environment: production
WARNING: This is a development server. Do not use it in a production
deployment.
* Use a production WSGI server instead.
* Debug mode: off
C:/Users/Mohansai/Documents/Zoom/lib/site-packages/sklearn/base.py:329:
UserWarning: Trying to unpickle estimator DecisionTreeClassifier from
version 0.23.2 when using version 0.23.1. This might lead to breaking
code or invalid results. Use at your own risk.
warnings.warn(
C:/Users/Mohansai/Documents/Zoom/lib/site-packages/sklearn/base.py:329:
UserWarning: Trying to unpickle estimator RandomForestClassifier from
version 0.23.2 when using version 0.23.1. This might lead to breaking
code or invalid results. Use at your own risk.
warnings.warn(
* Running on http://127.0.0.1:5000/ (Press CTRL+C to quit)
```

```
24 tnm = request.form["tnm"]
25 hae = request.form["hae"]
26 pain = request.form["pain"]
27 dys = request.form["dys"]
28 cough = request.form["cough"]
29 weakness = request.form["weakness"]
30 dm = request.form["dm"]
31 mi = request.form["mi"]
32 pad = request.form["pad"]
33 smoking = request.form["smoking"]
34 asthma = request.form["asthma"]
35 total = [(diagnosis, fev, age, performance, tnm, hae, pain, dys, cough, weakness, dm,
36 #prediction = model.predict(total)
37
38 #input_variables = pd.DataFrame([[performance, dys, cough, tnm, dm]], column
39
40 prediction = model.predict(total)[0]
41
42 if int(prediction) == 1:
43     prediction = "Patient is at High Risk"
44
45 else:
46     prediction = "Patient is Not at Risk"
47
48
49 return render_template("result.html", prediction = prediction)
50
51 return render_template("result.html")
52
53
54 if __name__ == "__main__":
55     app.run(debug=False)
56
57
```

```
Python 3.8.3 (default, Jul 2 2020, 17:30:36) [MSC v.1916 64 bit (AMD64)]
Type "copyright", "credits" or "license" for more information.

IPython 7.16.1 -- An enhanced Interactive Python.

In [1]: runfile('C:/Users/Mohansai/Downloads/5_6228888303706309736/One
Year Life expectancy post thoracic surgery using IBM watson studio/
flask/app.py', wdir='C:/Users/Mohansai/Downloads/5_6228888303706309736/
One Year Life expectancy post thoracic surgery using IBM watson studio/
flask')
* Serving Flask app "app" (lazy loading)
* Environment: production
WARNING: This is a development server. Do not use it in a production
deployment.
* Use a production WSGI server instead.
* Debug mode: off
C:/Users/Mohansai/Documents/Zoom/lib/site-packages/sklearn/base.py:329:
UserWarning: Trying to unpickle estimator DecisionTreeClassifier from
version 0.23.2 when using version 0.23.1. This might lead to breaking
code or invalid results. Use at your own risk.
warnings.warn(
C:/Users/Mohansai/Documents/Zoom/lib/site-packages/sklearn/base.py:329:
UserWarning: Trying to unpickle estimator RandomForestClassifier from
version 0.23.2 when using version 0.23.1. This might lead to breaking
code or invalid results. Use at your own risk.
warnings.warn(
* Running on http://127.0.0.1:5000/ (Press CTRL+C to quit)
```



## LUNGEVITY

Lung cancer is the leading cause of cancer-related deaths in the world. In the United States, lung cancer claims more lives every year than colon cancer, prostate cancer, and breast cancer combined. Patients who receive **Thoracic Surgery for Lung Cancer** do so with the expectation that their lives will be prolonged for a sufficient amount of time after the surgery.

The problem this app tries to solve is to find whether there is a way to determine **Post-Operative Life Expectancy** from patient attributes in the dataset. If there is pattern to be recognized, this would help physicians and patients make a more educated decision on whether they should proceed forward with the surgery. Not only would this influence physicians and patients but also health insurance companies, national health organizations and clinical researchers.

[View the APP](#)[How was this made?](#)

## Will the Patient Survive Post Thoracic Surgery ?

Find Out Whether Your Patient Is High Risk Before The Surgery

DIAGNOSIS	<input type="text" value="3"/>			
FEV	<input type="text" value="3"/>			
AGE	<input type="text" value="51"/>			
PERFORMANCE	<input type="text" value="PRZ0"/>			
TNM	<input type="text" value="OCT12"/>			
PAIN	HAEMOPTYSIS	DYSPNOEA	COUGH	WEAKNESS
<input type="radio"/> Yes	<input type="radio"/> Yes	<input type="radio"/> Yes	<input type="radio"/> Yes	<input type="radio"/> Yes
<input checked="" type="radio"/> No	<input checked="" type="radio"/> No	<input checked="" type="radio"/> No	<input checked="" type="radio"/> No	<input checked="" type="radio"/> No

## Will the Patient Survive Post Thoracic Surgery ?

Patient is Not at Risk

## 7.ADVANTAGES & DISADVANTAGES

Procedure	Application	Advantages	Disadvantages
Rigid bronchoscopy	Diagnosis of carcinoma Clear airway obstruction	Removal of secretions/blood Large biopsy specimen may reveal submucosal tumor Opportunity to debulk inoperable tumors or excise benign lesions Assessment of carinal fixation	Poor view of secondary bronchial divisions
Cervical and scalene lymph node biopsy	Diagnosis of enlarged nodes	Safer technique than fine-needle aspiration (FNA) for deeper nodes Large tissue sample	Operative procedure; occasional technical difficulty
Mediastinoscopy	Diagnosis of middle mediastinal pathology (node/mass)	Direct access to mediastinal abnormality Much larger tissue sample than FNA	Operative field restricted to paratracheal and subcarinal areas
Mediastinotomy	Diagnosis of anterior mediastinal pathology (node/mass)	Reaches areas of anterior mediastinum inaccessible to mediastinoscopy Useful for subaortic node sampling May be safer to use than mediastinoscopy in patients who have superior vena cava obstruction Much larger tissue sample than in FNA	Operative field restricted to upper anterior mediastinum More complex operative procedure
Thoracoscopy	Pleural biopsy Biopsy of visceral pleura deposits	Can be effected through single puncture Visually directed biopsy	Limited to small biopsies Dependent on pleural cavity being free of adhesions
Video-assisted thorascopic/thoracic surgery (VATS)	Any ipsilateral biopsy or sampling procedure	Complementary to mediastinoscopy and mediastinotomy Excellent view and full operative intervention possible (e.g., wedge excision of pulmonary nodule, dissection of mediastinal mass)	Dependent on pleural cavity being free of adhesions Significant surgical expertise required

## **8.CONCLUSION**

Detection of lung cancer is one of the challenging problems in medical field due to structure of cancer cells, where most of the cells are overlapped to each other. Detection of lung cancer in the early stage is curable. The system contains two parts. One is Lung Cancer Detection part and the other is the Prediction of Life Expectancy Post Thoracic Surgery. Both the parts can run independently. The system is provided as a web application where anyone can upload a CT scan image of the lung in the front end and check out whether that image is infected with Lung Cancer. At the backend the image uploaded is preprocessed, segmented and predicted using the CNN model which is already trained. Also, the system provides a form requesting for the 14 attributes required to predict the survival span post Thoracic Surgery. Once the form is submitted, the form inputs are run on the LDA model and a response of whether the patient will survive or not is produced. CNN model used to detect lung cancer gave an accuracy of 95% and the LDA classifier gave the highest accuracy of 83.76% compared to other 3 algorithms used. The system considers only CT lung images as input, but further the system can be enhanced to take MRI (Magnetic Resonance Imaging) or PET (Position Emission Tomography) as input. Also, in the prediction of survival part, higher accuracies of the classifier can be achieved by considering large amount of dataset.

## **9.FUTURE SCOPE**

Thoracic surgery is living an enthusiastic era of thriving innovations. The changes in the fields of diagnostics, in the therapeutic options, the development of surgical techniques and the adoption of novel technologies has radically changed the horizons of our specialty. We must embrace the innovations, learn navigational bronchoscopy, understand the multiple targeted therapies, and learn new ways to operate on advanced cases. We will not be able to stay complacent with our current 3 ports video-assisted thoracoscopic surgery (VATS) technique with open instruments, therefore every surgeon needs to be watching for each latest development as it happens. In this brief manuscript we will summarize some of the most important development over the recent years and forecast how this will impact on the patients affected by thoracic malignancies. As thoracic surgeons we have to embrace these developments in order to redefine our specialty.

## 10.BIBLIOGRAPHY

As we need to connect to internet for opening websites and gather information about the skills required for the jobs, we can also use other programming languages like C, C++, Java but its effective to use and also have predefined functions to use in the python language since the syntax for this programming language is simple to use and is more effective comparatively. So we have used python for this project for accessing sites and recommended skills as per the present trending technologies

## APPENDIX

### A.Source code

```
import pandas as pd
import numpy as np
import seaborn as sns
from sklearn.model_selection import train_test_split
from sklearn.linear_model import LogisticRegression
from sklearn.metrics import accuracy_score
import matplotlib.pyplot as plt
from sklearn.linear_model import LogisticRegression
from sklearn.ensemble import RandomForestClassifier, GradientBoostingClassifier
from sklearn.tree import DecisionTreeClassifier
from sklearn.neighbors import KNeighborsClassifier
from sklearn.metrics import f1_score
from sklearn.metrics import classification_report, confusion_matrix
import itertools
df=pd.read_csv("ThoracicSurgery.csv")
df.head()
df.columns
df.describe()
df.shape
df.info()
df.isnull().sum()
live = df[df['Death_1yr'] == 0]
death = df[df['Death_1yr'] == 1]

cond = ['FVC', 'FEV1', 'Performance', 'Pain', 'Haemoptysis', 'Dyspnoea',
, 'Cough', 'Weakness',\
```

```

        'Tumor_Size', 'Diabetes_Mellitus', 'MI_6mo', 'PAD', 'Smoking',
        'Asthma', 'Age']

l = [np.mean(live[c]) for c in cond]
d = [np.mean(death[c]) for c in cond]

ld = pd.DataFrame(data={'Attribute': cond, 'Live 1yr Mean': l, 'Death 1
yr Mean': d})
ld = ld.set_index('Attribute')

print('Death: {:d}, Live: {:d}'.format(len(death), len(live)))
print("1 year death: {:.2f}% out of 454 patients".format(np.mean(df.Dea
th_1yr)*100))
ld
d = np.array(d)
l = np.array(l)

p_diff = (d-l)/l*100

fig, axes = plt.subplots(2,1,figsize=(12,18))

axes[0].bar(cond, p_diff)
axes[0].set_title('Mean Difference % between Dead and Live 1yr', fontsi
ze=18)
axes[0].set_xticks(cond)
axes[0].set_xticklabels(cond, rotation=90)
axes[0].set_ylabel('Percent', fontsize=13)

tf_col = ['Pain', 'Haemoptysis', 'Dyspnoea', 'Cough', 'Weakness', 'Diab
etes_Mellitus', 'MI_6mo', 'PAD', 'Smoking', 'Asthma']
tf_sum = [df[col].sum()/454 for col in tf_col]

axes[1].bar(tf_col, tf_sum)
axes[1].set_xticks(tf_col)
axes[1].set_xticklabels(tf_col, rotation=90)
axes[1].set_ylabel('Proportion of Total Patients', fontsize=13)
axes[1].set_title('Proportion of Patient Conditions before Surgery', fo
ntsize=18)

plt.tight_layout()

plt.show()
fig, axes = plt.subplots(3,1,figsize=(10,15))

sns.countplot(x='Diagnosis', hue='Death_1yr', data=df, palette='Blues_d
', ax=axes[0]).set_title('Diagnosis', fontsize=18)
sns.countplot(x='Tumor_Size', hue='Death_1yr', data=df, palette='Blues_d
', ax=axes[1]).set_title('Tumor_Size', fontsize=18)
sns.countplot(x='Performance', hue='Death_1yr', data=df, palette='Blues
_d', ax=axes[2]).set_title('Performance', fontsize=18)

plt.tight_layout()
def permutation_sample(data1, data2):
    """Generate a permutation sample from two data sets."""
    data = np.concatenate((data1, data2))
    permuted_data = np.random.permutation(data)

```



```

    perm_sample_1 = permuted_data[:len(data1)]
    perm_sample_2 = permuted_data[len(data1):]

    return perm_sample_1, perm_sample_2

def draw_perm_reps(data_1, data_2, func, size=1):
    """Generate multiple permutation replicates."""
    perm_replicates = np.empty(size)

    for i in range(size):
        perm_sample_1, perm_sample_2 = permutation_sample(data_1, data_
2)
        perm_replicates[i] = func(perm_sample_1, perm_sample_2)

    return perm_replicates

def diff_of_means(data_1, data_2):
    """Difference in means of two arrays."""
    diff = np.mean(data_1) - np.mean(data_2)
    return diff

condition = ['FVC', 'FEV1', 'Performance', 'Pain', 'Haemoptysis', 'Dysp
noea', 'Cough', 'Weakness', \
            'Tumor_Size', 'Diabetes_Mellitus', 'MI_6mo', 'PAD', 'Smoki
ng', 'Asthma', 'Age']
p_val = []

for c in condition:
    empirical_diff_means = diff_of_means(death[c], live[c])
    perm_replicates = draw_perm_reps(death[c], live[c], diff_of_means,
size=10000)
    if empirical_diff_means > 0:
        p = np.sum(perm_replicates >= empirical_diff_means) / len(perm_
replicates)
        p_val.append(p)
    else:
        p = np.sum(perm_replicates <= empirical_diff_means) / len(perm_
replicates)
        p_val.append(p)

print(list(zip(condition, p_val)))
fig, axes = plt.subplots(1,2,figsize=(13,5))
axes[0].plot(df.FVC, df.FEV1, linestyle='none', marker='.')

axes[0].set_xlabel('FVC', fontsize=13)
axes[0].set_ylabel('FEV1', fontsize=13)
axes[0].set_title('FVC vs FEV1', fontsize=16)

axes[1].plot(df.Age, df.FEV1, linestyle='none', marker='.', label='FEV1
')
axes[1].plot(df.Age, df.FVC, linestyle='none', marker='.', label='FVC')
axes[1].set_xlabel('Age', fontsize=13)
axes[1].set_ylabel('FEV1, FVC', fontsize=13)
axes[1].legend()
axes[1].set_title('Age vs FEV1, FVC', fontsize=16)

plt.tight_layout()
bnp.corrcoef(df.FVC, df.FEV1)[0,1]
np.corrcoef(df.Age, df.FVC)[0,1]

```



```

np.corrcoef(df.Age, df.FEV1)[0,1]
def ecdf(data):
    """Compute ECDF for a one-dimensional array of measurements."""
    n = len(data)
    x = np.sort(data)
    y = np.arange(1, n+1) / n
    return x, y
x_fvc, y_fvc = ecdf(df.FVC)
x_fev1, y_fev1 = ecdf(df.FEV1)
x_age, y_age = ecdf(df.Age)

fig, axes = plt.subplots(1,2,figsize=(13,5))
axes[0].plot(x_fvc, y_fvc, marker='.', linestyle='none', label='FVC')
axes[0].plot(x_fev1, y_fev1, marker='.', linestyle='none', label='FEV1'
)

axes[0].set_xlabel('Numerical Value', fontsize=13)
axes[0].set_ylabel('ECDF', fontsize=13)
axes[0].legend(loc='upper left')
axes[0].set_title('ECDF of FVC & FEV1', fontsize=16)

axes[1].plot(x_age, y_age, marker='.', linestyle='none', label='Age')
axes[1].set_xlabel('Years Old', fontsize=13)
axes[1].set_ylabel('ECDF', fontsize=13)
axes[1].legend(loc='upper left')
axes[1].set_title('ECDF of Age', fontsize=16)

plt.tight_layout()
df.drop(['FVC'],axis=1, inplace=True)
x=df.iloc[:,0:15].values
y=df.iloc[:,15:16].values
x_train,x_test,y_train,y_test=train_test_split(x,y,test_size=0.2,random
_state=0)
print('Shape of x_train {}'.format(x_train.shape))
print('Shape of y_train {}'.format(y_train.shape))
print('Shape of x_test {}'.format(x_test.shape))s
print('Shape of y_test {}'.format(y_test.shape))
def decisionTree(x_train, x_test, y_train, y_test):
    dt=DecisionTreeClassifier()
    dt.fit(x_train,y_train.ravel())
    yPred = dt.predict(x_test)
    print('***DecisionTreeClassifier***')
    print('Confusion matrix')
    print(confusion_matrix(y_test,yPred))
    print('Classification report')
    print(classification_report(y_test,yPred))
def randomForest(x_train, x_test, y_train, y_test):
    rf = RandomForestClassifier()
    rf.fit(x_train,y_train.ravel())
    yPred = rf.predict(x_test)
    print('***RandomForestClassifier***')
    print('Confusion matrix')
    print(confusion_matrix(y_test,yPred))
    print('Classification report')
    print(classification_report(y_test,yPred))
def KNN(x_train, x_test, y_train, y_test):
    knn = KNeighborsClassifier()
    knn.fit(x_train,y_train.ravel())

```

```

yPred = knn.predict(x_test)
print('***KNeighborsClassifier***')
print('Confusion matrix')
print(confusion_matrix(y_test,yPred))
print('Classification report')
print(classification_report(y_test,yPred))
def xgboost(x_train, x_test, y_train, y_test):
    xg = GradientBoostingClassifier()
    xg.fit(x_train,y_train.ravel())
    yPred = xg.predict(x_test)
    print('***GrandientBoostingClassifier***')
    print('Confusion matrix')
    print(confusion_matrix(y_test,yPred))
    print('Classification report')
    print(classification_report(y_test,yPred))
def compareModel(x_train, x_test, y_train, y_test):
    decisionTree(x_train, x_test, y_train, y_test)
    print('-'*100)
    randomForest(x_train, x_test, y_train, y_test)
    print('-'*100)
    KNN(x_train, x_test, y_train, y_test)
    print('-'*100)
    xgboost(x_train, x_test, y_train, y_test)
compareModel(x_train, x_test, y_train, y_test)
from sklearn.model_selection import cross_val_score
rf= RandomForestClassifier()
rf.fit(x_train,y_train.ravel())
yPred = rf.predict(x_test)
f1_score(yPred,y_test,average='weighted')
cv = cross_val_score(rf,x,y.ravel(),cv=5)
np.mean(cv)
import pickle
pickle.dump(rf,open('model.pkl','wb'))

```