## **Cancerous Gene Classification**

- This problem statement was a part of the Kaggle competition https://www.kaggle.com/c/msk-redefining-cancer-treatment/overview
- Download the datasets 'training\_text' & 'training\_variants' from https://www.kaggle.com/c/msk-redefining-cancer-treatment/data

#### Summary of the code below:

- 1. Importing dependencies and acquiring data
- 2. Performing EDA and Data Pre-processing
- 3. Splitting the dataset into train, cross-validation and test sets
- 4. Performing hyper-parameter tuning over the cross-validation set for each model namely, Logistic Regression, Linear SVM and Random Forest.
- The metric used to evaluate and compare the results of the models was log-loss.

```
import pandas as pd
In [1]:
        import numpy as np
        import seaborn as sb
        import matplotlib.pyplot as plt
        import re
        import math
        from nltk.corpus import stopwords
        import pdb
        import warnings
        from sklearn.model selection import train test split
        from sklearn.feature extraction.text import TfidfVectorizer
        from sklearn.feature extraction.text import CountVectorizer
        from sklearn.preprocessing import normalize
        from scipy.sparse import hstack
        from sklearn.linear model import SGDClassifier
        from sklearn.svm import SVC
        from sklearn.calibration import CalibratedClassifierCV
        warnings.filterwarnings("ignore")
        import seaborn as sns
        from sklearn.metrics import plot confusion matrix
        from sklearn.metrics.classification import accuracy score, log loss
        from sklearn.feature extraction.text import TfidfVectorizer
        from sklearn.svm import SVC
        from sklearn.ensemble import RandomForestClassifier
        from sklearn.linear model import LogisticRegression
```

### Data Acquisition

1

CBL

1

W802\*

2

```
      2
      2
      CBL
      Q249E
      2

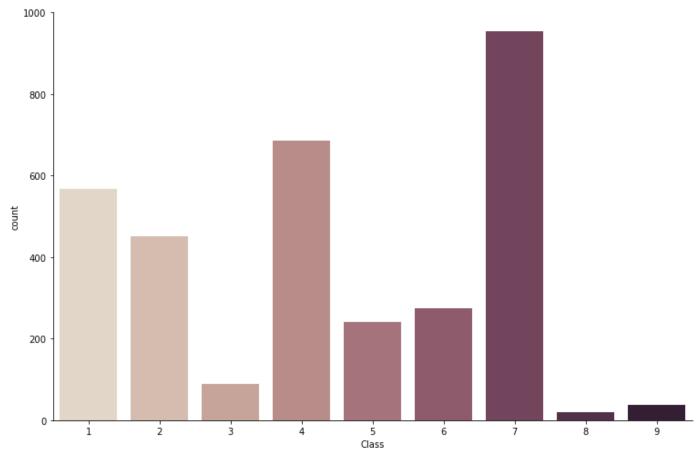
      3
      3
      CBL
      N454D
      3

      4
      4
      CBL
      L399V
      4
```

```
In [3]: print('Number of classes: ',data.Class.unique())
Number of classes: [1 2 3 4 5 6 7 8 9]
```

Thus, the gene-variation pairs will get classified into 9 different classes

```
In [4]: sb.catplot(x="Class", kind="count", palette="ch:.25", data=data, height = 7, aspect = 1.
Out[4]: <seaborn.axisgrid.FacetGrid at 0x21f4e030cc0>
```



Since the distribution of classes is skewed, we have to maintain this disb. throughout the train, cv and test sets

(3321, 2)

```
Out[5]: ID Text

O 0 Cyclin-dependent kinases (CDKs) regulate a var...

1 1 Abstract Background Non-small cell lung canc...

2 2 Abstract Background Non-small cell lung canc...

3 3 Recent evidence has demonstrated that acquired...

4 Oncogenic mutations in the monomeric Casitas B...
```

### Data Pre-processing

Out[8]:

ID

1

0 0

```
# loading stop words from nltk library
In [6]:
        stop words = set(stopwords.words('english'))
        # referenced from https://stackoverflow.com/questions/23996118/replace-special-character
        # and https://pythonexamples.org/python-replace-multiple-spaces-with-single-space-in-tex
        def clean text(corpus):
            if type(corpus) is not int:
                text = re.sub('[^a-zA-Z0-9]',' ',corpus)
                text = re.sub('\s+',' ',corpus)
                text = text.lower()
                for word in text.split():
                    if word in stop words:
                        a += ''
                    else:
                        a += word + " "
                return a
In [7]: ids = text_data['ID'].copy()
        index = 0
        for i in text data. Text:
            #pdb.set trace()
            if type(i) is str:
                x = clean text(i)
                text data['Text'][index] = x
            else:
                b = ids[index]
                print(f'Text NA for ID:{b}')
            index += 1
        Text NA for ID:1109
        Text NA for ID:1277
        Text NA for ID:1407
        Text NA for ID:1639
        Text NA for ID:2755
In [8]: | text_data.head()
```

**Text** 

cyclin-dependent kinases (cdks) regulate varie...

abstract background non-small cell lung cancer...

- **2** abstract background non-small cell lung cancer...
- **3** recent evidence demonstrated acquired uniparen...
- 4 4 oncogenic mutations monomeric casitas b-lineag...

```
In [9]: #merging both gene_variations and text data based on ID
    result = pd.merge(data, text_data, on='ID', how='left')
    result.head()
```

```
Out[9]:
              ID
                      Gene
                                        Variation Class
                                                                                                       Text
           0
               0
                   FAM58A
                            Truncating Mutations
                                                       1
                                                              cyclin-dependent kinases (cdks) regulate varie...
                                           W802*
                                                       2
           1
               1
                       CBL
                                                             abstract background non-small cell lung cancer...
           2
               2
                       CBL
                                           Q249E
                                                       2
                                                             abstract background non-small cell lung cancer...
               3
                       CBL
                                           N454D
           3
                                                           recent evidence demonstrated acquired uniparen...
               4
                       CBL
                                           L399V
                                                           oncogenic mutations monomeric casitas b-lineag...
```

```
ID
                              Variation Class Text
             Gene
1109 1109
            FANCA
                                S1088F
                                              NaN
                                           1
1277 1277 ARID5B
                    Truncating Mutations
                                              NaN
1407 1407
                                K508M
                                              NaN
             FGFR3
1639 1639
              FLT1
                           Amplification
                                              NaN
2755 2755
              BRAF
                                G596C
                                           7 NaN
```

Out[11]:

```
In [12]: pd.options.display.min_rows = 20
b = result.groupby('Gene')['Gene'].count()
b
b.sort_values(ascending = False)
```

```
Gene
Out[12]:
         BRCA1
                     264
         TP53
                    163
         EGFR
                     141
         PTEN
                    126
                    125
         BRCA2
         KIT
                     99
         BRAF
                      93
                      69
         ERBB2
         ALK
                      69
         PDGFRA
                      60
         PPM1D
                       1
         PMS1
                       1
```

```
PAX8
                     1
        ERRFI1
        PAK1
                     1
        FAM58A
                     1
        FANCC
                     1
        FGF19
                     1
        KLF4
                     1
        Name: Gene, Length: 264, dtype: int64
In [ ]:
         # referenced from https://www.geeksforgeeks.org/iterating-over-rows-and-columns-in-panda
In [13]:
         for i, j in a.iterrows():
             count = 0
             for k, v in result.iterrows():
                 #pdb.set trace()
                 if (j['Gene'] == v['Gene']) & (j['Variation'] == v['Variation']):
                     count+=1
             if count >= 1:
                print('The count of the pair of {} and {} is {}'.format(j['Gene'],j['Variation']
        The count of the pair of FANCA and S1088F is 1
        The count of the pair of ARID5B and Truncating Mutations is 1
        The count of the pair of FGFR3 and K508M is 1
        The count of the pair of FLT1 and Amplification is 1
        The count of the pair of BRAF and G596C is 1
```

PIK3R3

1

• We see that the above pairs having no supporting text are only present once throughout the dataset. If we remove these rows then we could loose their classification (especially for class 6 as they are less in count throughout the dataset)

Instead of removing these gene-variation pairs, we can just replace the NaN with the gene and variation name in the respective 'Text' field.

```
In [14]: | result.loc[result['Text'].isnull(),'Text'] = result['Gene'] +' '+result['Variation']
In [15]: result[result['ID']==2755]
                ID Gene Variation Class
Out[15]:
                                            Text
                                   7 BRAF G596C
         2755 2755 BRAF
                           G596C
In [16]:
         len(result)
         3321
Out[16]:
         1.1.1
In [17]:
         Since, we're to do onehot encoding for the features, it is better that we replace the sp
         feature with an underscore
         1.1.1
         strat disb = result['Class'].values
         result.Gene = result.Gene.str.replace('\s+', ' ')
         result. Variation = result. Variation.str.replace('\s+', ' ')
         # using stratify will help us here to maintain the same distribution of class labels in
```

```
x train, x cv, y train, y cv = train test split(X train, train y, stratify=train y, test
In [18]: # featurization of the text feature using one hot encoding (tfidf)
         # limiting the featurization to those words which occur atleast 3 times in the corpus
         text tfidf = TfidfVectorizer(min df=3)
         text tfidf.fit(x train['Text'])
         train_text = text_tfidf.transform(x train['Text'])
         train text = normalize(train text, axis=0)
         # using the same vectorizer that was trained on train data
         cv text = text tfidf.transform(x cv['Text'])
         cv text = normalize(cv text, axis=0)
         test text = text tfidf.transform(x test['Text'])
         test text = normalize(test text, axis=0)
In [19]: # featurization of the gene and variation features using one hot encoding
         gene ohe = CountVectorizer()
         train gene = gene ohe.fit transform(x train['Gene'])
         train gene = normalize(train gene, axis=0)
         cv gene = gene ohe.transform(x cv['Gene'])
         cv gene = normalize(cv gene, axis=0)
         test gene = gene ohe.transform(x test['Gene'])
         test gene = normalize(test gene, axis=0)
         variation ohe = CountVectorizer()
         train variation = variation ohe.fit transform(x train['Variation'])
         train variation = normalize(train variation, axis=0)
         cv variation = variation ohe.transform(x cv['Variation'])
         cv_variation = normalize(cv_variation, axis=0)
         test variation = variation ohe.transform(x test['Variation'])
         test variation = normalize(test variation, axis=0)
In [20]: print(train_gene.shape)
         print(train variation.shape)
         print(train text.shape)
         (2124, 233)
         (2124, 1974)
         (2124, 56489)
In [21]: x train final = hstack((train gene, train variation, train text)).tocsr()
         y train final = np.array(list(x train['Class']))
         x cv final = hstack((cv gene, cv variation, cv text)).tocsr()
         y cv final = np.array(list(x cv['Class']))
         x test final = hstack((test gene, test variation, test text)).tocsr()
         y test final = np.array(list(x test['Class']))
In [22]: print(x_train final.shape)
         print(x test final.shape)
         print(x cv final.shape)
```

X train, x test, train y, y test = train test split(result, strat disb, stratify=strat d

```
(2124, 58696)
(665, 58696)
(532, 58696)
```

# **Logistic Regression**

Hyper-parameter tuning

```
In [23]: # Using SGD Classifier with loss argument as log loss so that it functions as a Logistic
         # Using 'balanced' class weight here to maintain the ratio of the classes
        Since we do not want a blackbox model and want to classify the mutations with utmost cer
        referenced from https://machinelearningmastery.com/calibrated-classification-model-in-sc
         alpha = [10 ** x for x in range(-5, 2)]
         list log loss = []
         for i in alpha:
            print("For alpha =", i)
            logistic = SGDClassifier(class weight='balanced', alpha=i, penalty='12', loss='log',
            logistic.fit(x train final, y train final)
            prob logistic = CalibratedClassifierCV(logistic)
            prob logistic.fit(x train final, y train final)
            prob x cv = prob logistic.predict proba(x cv final)
            list log loss.append(log loss(y cv final, prob x cv, labels=logistic.classes ))
             # to avoid rounding error while multiplying probabilites I used log-probability esti
            print("CV log Loss :",log loss(y cv final, prob x cv))
        For alpha = 1e-05
        CV log Loss: 1.2554856827616867
        For alpha = 0.0001
        CV log Loss: 1.1013905238914352
        For alpha = 0.001
        CV log Loss: 1.165879328943427
        For alpha = 0.01
        CV log Loss: 1.2518909495939732
        For alpha = 0.1
        CV log Loss: 1.4005614924980676
        For alpha = 1
        CV log Loss : 1.560547384571933
        For alpha = 10
        CV log Loss: 1.5931389097786206
```

### Modelling with the best value of alpha

For the value of best alpha = 0.0001

```
In [24]: best_alpha = np.argmin(list_log_loss)
    logistic = SGDClassifier(class_weight='balanced', alpha=alpha[best_alpha], penalty='12',
    logistic.fit(x_train_final, y_train_final)
    prob_logistic = CalibratedClassifierCV(logistic, method="sigmoid")
    prob_logistic.fit(x_train_final, y_train_final)

print('For the value of best alpha =', alpha[best_alpha])
    prob_final_train = prob_logistic.predict_proba(x_train_final)
    print('\nTrain log loss :',log_loss(y_train_final, prob_final_train, labels=logistic.cla
    prob_final_cv = prob_logistic.predict_proba(x_cv_final)
    print('Cross Validation log loss :',log_loss(y_cv_final, prob_final_cv, labels=logistic.
    prob_final_test = prob_logistic.predict_proba(x_test_final)
    print('Test log loss :',log_loss(y_test_final, prob_final_test, labels=logistic.classes_
```

```
Cross Validation log loss : 1.1013905238914352
        Test log loss: 1.069329304971553
In [25]: pred_labels = []
         for i in prob final test:
            i = list(i)
            pos = i.index(max(i))
            pred labels.append(pos+1)
        print("Misclassification %age:", np.count nonzero((pred labels - y test final))/y test
        Misclassification %age: 0.35037593984962406
In [26]: test point index = 5
        predicted class = prob logistic.predict(x test final[test point index])
        print("Predicted Class:", predicted class[0])
        print("Predicted Class Probabilities:", np.round(prob logistic.predict proba(x test fina
        print("Actual Class :", y_test_final[test point index])
        Predicted Class: 1
        Predicted Class Probabilities: [[0.89 0.03 0.01 0.01 0.02 0.01 0.02 0.01 0.01]]
        Actual Class : 1
```

## **Linear SVM**

Train log loss : 0.5428493957402157

Hyper-parameter tuning

```
In [27]: # Since we have used the loss argument as 'hinge', it will function as an **SVM Classifi
         c = [10 ** x for x in range(-5, 3)]
         list log loss = []
         for i in c:
            print("For C =", i)
            svc = SVC(C=i,kernel='linear',probability=True, class weight='balanced', random sta
            svc = SGDClassifier(class weight='balanced', alpha=i, penalty='12', loss='hinge', ra
            svc.fit(x train final, y train final)
            prob svc = CalibratedClassifierCV(svc, method="sigmoid")
            prob svc.fit(x train final, y train final)
            prob x cv = prob svc.predict proba(x cv final)
            list log loss.append(log loss(y cv final, prob x cv, labels=svc.classes , eps=1e-15)
            print("CV log Loss :",log loss(y cv final, prob x cv))
        For C = 1e-05
        CV log Loss : 1.2888112114505472
        For C = 0.0001
        CV log Loss: 1.2341552428671316
        For C = 0.001
        CV log Loss: 1.130969643092841
        For C = 0.01
        CV log Loss: 1.242584803512788
        For C = 0.1
        CV log Loss : 1.4057934721134888
        For C = 1
        CV log Loss: 1.5963262909367837
        For C = 10
        CV log Loss: 1.5961996011305433
        For C = 100
        CV log Loss: 1.5961992802457856
```

```
best_alpha = np.argmin(list_log_loss)
In [28]:
        svc = SGDClassifier(class weight='balanced', alpha=alpha[best alpha], penalty='12', loss
        svc.fit(x train final, y train final)
        prob svc = CalibratedClassifierCV(svc, method="sigmoid")
        prob svc.fit(x train final, y train final)
        print('For values of best alpha = ', alpha[best alpha])
        prob final train = prob svc.predict proba(x train final)
        print('\nTrain log loss:',log loss(y train final, prob final train, labels=svc.classes
        prob final cv = prob svc.predict proba(x cv final)
        print('Cross Validation log loss:',log loss(y cv final, prob final cv, labels=svc.class
        prob final test = prob svc.predict proba(x test final)
        print('Test log loss:',log loss(y test final, prob final test, labels=svc.classes))
        For values of best alpha = 0.001
        Train log loss: 0.6078792833811073
        Cross Validation log loss: 1.130969643092841
        Test log loss: 1.1197654783603976
In [29]: pred_labels = []
        for i in prob final test:
            i = list(i)
            pos = i.index(max(i))
            pred labels.append(pos+1)
        print("Misclassification %age :", np.count nonzero((pred labels - y test final))/y test
        Misclassification %age: 0.3518796992481203
In [30]: test point index = 5
        predicted class = prob svc.predict(x test final[test point index])
        print("Predicted Class :", predicted class[0])
        print("Predicted Class Probabilities:", np.round(prob svc.predict proba(x test final[tes
        print("Actual Class:", y_test_final[test point index])
        Predicted Class : 1
        Predicted Class Probabilities: [[0.84 0.04 0. 0.04 0.02 0. 0.05 0. 0.01]]
        Actual Class: 1
```

### **Random Forest**

Hyper-parameter tuning

```
In [32]: n_{est} = [100, 200, 500, 1000, 2000]
         list log loss = []
         for i in n est:
                 print("for n estimators =", i)
                 rf = RandomForestClassifier(n estimators=i, criterion='gini', max depth=10, rand
                 rf.fit(x train final, y train final)
                 prob rf = CalibratedClassifierCV(rf, method="sigmoid")
                 prob rf.fit(x train final, y train final)
                 prob cv rf = prob rf.predict proba(x cv final)
                list log loss.append(log loss(y cv final, prob cv rf, labels=rf.classes , eps=1e
                 print("CV log Loss :",log loss(y cv final, prob cv rf))
        for n estimators = 100
        CV log Loss: 1.239328435179232
        for n estimators = 200
        CV log Loss: 1.2296648304669864
        for n estimators = 500
        CV log Loss: 1.2221892286188227
        for n estimators = 1000
        CV log Loss: 1.2189293146494184
```

```
for n_estimators = 2000
CV log Loss : 1.2176029595437161
```

We notice that there isnt much difference in the log loss values of (n\_estimators = 500 & max\_depth = 10) and (n\_estimators = 2000 & max\_depth = 10), but there is a big difference in the time complexity of the two. Hence, we select the best pair to be n\_estimators = 500 and max\_depth = 10 as it has lower time complexity.

Modellling with the best value of estimators and max\_depth

```
In [33]: rf = RandomForestClassifier(n estimators=1000, criterion='gini', max depth=10, random st
         rf.fit(x train final, y train final)
         prob rf = CalibratedClassifierCV(rf, method="sigmoid")
         prob rf.fit(x train final, y train final)
         print('For values of best estimator = ', 1000)
         prob final train = prob rf.predict proba(x train final)
         print('\nTrain log loss:',log loss(y train final, prob final train, labels=rf.classes)
         prob final cv = prob rf.predict proba(x cv final)
         print('Cross Validation log loss:',log loss(y cv final, prob final cv, labels=rf.classe
         prob final test = prob rf.predict proba(x test final)
         print('Test log loss:',log loss(y test final, prob final test, labels=rf.classes))
         For values of best estimator = 1000
        Train log loss: 0.6342129238888813
        Cross Validation log loss : 1.2189293146494184
        Test log loss : 1.1526961241417986
        pred labels = []
In [34]:
         for i in prob final test:
            i = list(i)
            pos = i.index(max(i))
            pred labels.append(pos+1)
         print("Misclassification %age:", np.count nonzero((pred labels - y test final))/y test
        Misclassification %age: 0.37142857142857144
In [35]: test_point index = 5
         predicted class = prob rf.predict(x test final[test point index])
         print("Predicted Class :", predicted class[0])
         print("Predicted Class Probabilities:", np.round(prob rf.predict proba(x test final[test
        print("Actual Class :", y_test_final[test point index])
        Predicted Class: 1
        Predicted Class Probabilities: [[0.5 0.06 0.02 0.24 0.05 0.04 0.07 0.01 0.01]]
        Actual Class : 1
```

The best result from all the three models was from 'Logistic Regression', probably because of Logistic Regression's ablility to handle large dimensional dataset better.