# Personalized cancer diagnosis

# c) Logistic regression with CountVectorizer Features, including both unigrams and bigrams

# 1. Business Problem

# 1.1. Description

Source: https://www.kaggle.com/c/msk-redefining-cancer-treatment/

Data: Memorial Sloan Kettering Cancer Center (MSKCC)

Download training\_variants.zip and training\_text.zip from Kaggle.

#### Context:

Source: https://www.kaggle.com/c/msk-redefining-cancer-treatment/discussion/35336#198462

#### Problem statement:

Classify the given genetic variations/mutations based on evidence from text-based clinical literature.

# 1.2. Source/Useful Links

Some articles and reference blogs about the problem statement

- 1. <a href="https://www.forbes.com/sites/matthewherper/2017/06/03/a-new-cancer-drug-helped-almost-everyone-who-took-it-almost-heres-what-it-teaches-us/#2a44ee2f6b25">https://www.forbes.com/sites/matthewherper/2017/06/03/a-new-cancer-drug-helped-almost-everyone-who-took-it-almost-heres-what-it-teaches-us/#2a44ee2f6b25</a>
- 2. https://www.youtube.com/watch?v=UwbuW7oK8rk
- 3. <a href="https://www.youtube.com/watch?v=qxXRKVompl8">https://www.youtube.com/watch?v=qxXRKVompl8</a>

# 1.3. Real-world/Business objectives and constraints.

- · No low-latency requirement.
- Interpretability is important.
- Errors can be very costly.
- · Probability of a data-point belonging to each class is needed.

# 2. Machine Learning Problem Formulation

# 2.1. Data

# 2.1.1. Data Overview

- Source: https://www.kaggle.com/c/msk-redefining-cancer-treatment/data
- We have two data files: one conatins the information about the genetic mutations and the other contains the clinical evidence (text) that human experts/pathologists use to classify the genetic mutations.
- Both these data files are have a common column called ID
- Data file's information:
  - training variants (ID, Gene, Variations, Class)
  - training\_text (ID, Text)

# 2.1.2. Example Data Point

#### training\_variants

ID,Gene,Variation,Class 0,FAM58A,Truncating Mutations,1 1,CBL,W802\*,2 2,CBL,Q249E,2

...

### training\_text

#### ID,Text

0||Cyclin-dependent kinases (CDKs) regulate a variety of fundamental cellular processes. CDK10 stands out as one of the last orphan CDKs for which no activating cyclin has been identified and no kinase activity revealed. Previous work has shown that CDK10 silencing increases ETS2 (v-ets erythroblastosis virus E26 oncogene homolog 2)-driven activation of the MAPK pathway, which confers tamoxifen resistance to breast cancer cells. The precise mechanisms by which CDK10 modulates ETS2 activity, and more generally the functions of CDK10, remain elusive. Here we demonstrate that CDK10 is a cyclin-dependent kinase by identifying cyclin M as an activating cyclin. Cyclin M, an orphan cyclin, is the product of FAM58A, whose mutations cause STAR syndrome, a human developmental anomaly whose features include toe syndactyly, telecanthus, and anogenital and renal malformations. We show that STAR syndrome-associated cyclin M mutants are unable to interact with CDK10. Cyclin M silencing phenocopies CDK10 silencing in increasing c-Raf and in conferring tamoxifen resistance to breast cancer cells. CDK10/cyclin M phosphorylates ETS2 in vitro, and in cells it positively controls ETS2 degradation by the proteasome. ETS2 protein levels are increased in cells derived from a STAR patient, and this increase is attributable to decreased cyclin M levels. Altogether, our results reveal an additional regulatory mechanism for ETS2, which plays key roles in cancer and development. They also shed light on the molecular mechanisms underlying STAR syndrome. Cyclin-dependent kinases (CDKs) play a pivotal role in the control of a number of fundamental cellular processes (1). The human genome contains 21 genes encoding proteins that can be considered as members of the CDK family owing to their sequence similarity with bona fide CDKs, those known to be activated by cyclins (2). Although discovered almost 20 y ago (3, 4), CDK10 remains one of the two CDKs without an identified cyclin partner. This knowledge gap has largely impeded the exploration of its biological functions. CDK10 can act as a positive cell cycle regulator in some cells (5, 6) or as a tumor suppressor in others (7, 8). CDK10 interacts with the ETS2 (v-ets erythroblastosis virus E26 oncogene homolog 2) transcription factor and inhibits its transcriptional activity through an unknown mechanism (9). CDK10 knockdown derepresses ETS2, which increases the expression of the c-Raf protein kinase, activates the MAPK pathway, and induces resistance of MCF7 cells to tamoxifen (6). ...

# 2.2. Mapping the real-world problem to an ML problem

# 2.2.1. Type of Machine Learning Problem

There are nine different classes a genetic mutation can be classified into => Multi class classification problem

#### 2.2.2. Performance Metric

Source: https://www.kaggle.com/c/msk-redefining-cancer-treatment#evaluation

Metric(s):

- · Multi class log-loss
- Confusion matrix

# 2.2.3. Machine Learing Objectives and Constraints

Objective: Predict the probability of each data-point belonging to each of the nine classes.

Constraints:

- Interpretability
- Class probabilities are needed.
- Penalize the errors in class probabilites => Metric is Log-loss.
- · No Latency constraints.

# 2.3. Train, CV and Test Datasets

Split the dataset randomly into three parts train, cross validation and test with 64%,16%, 20% of data respectively

# 3. Exploratory Data Analysis

```
In [0]:
```

```
import pandas as pd
import matplotlib.pyplot as plt
%matplotlib inline
import re
import time
import warnings
import numpy as np
from nltk.corpus import stopwords
from sklearn.decomposition import TruncatedSVD
from sklearn.preprocessing import normalize
from sklearn.feature extraction.text import CountVectorizer
from sklearn.manifold import TSNE
import seaborn as sns
from sklearn.neighbors import KNeighborsClassifier
from sklearn.metrics import confusion matrix
from sklearn.metrics.classification import accuracy score, log loss
from sklearn.feature extraction.text import TfidfVectorizer
from sklearn.linear model import SGDClassifier
from imblearn.over_sampling import SMOTE
from collections import Counter
from scipy.sparse import hstack
from sklearn.multiclass import OneVsRestClassifier
from sklearn.svm import SVC
from sklearn.model_selection import StratifiedKFold
from collections import Counter, defaultdict
from sklearn.calibration import CalibratedClassifierCV
from sklearn.naive bayes import MultinomialNB
from sklearn.naive_bayes import GaussianNB
from sklearn.model selection import train test split
from sklearn.model selection import GridSearchCV
from sklearn.metrics import normalized_mutual_info_score
from sklearn.ensemble import RandomForestClassifier
warnings.filterwarnings("ignore")
from mlxtend.classifier import StackingClassifier
from sklearn import model selection
from sklearn.linear model import LogisticRegression
```

# 3.1. Reading Data

```
In [0]:
```

```
from google.colab import drive
drive.mount('/content/drive')
```

Drive already mounted at /content/drive; to attempt to forcibly remount, call drive.mount("/content/drive", force\_remount=True).

## 3.1.1. Reading Gene and Variation Data

```
data = pd.read_csv('/content/drive/My Drive/cd/training/training_variants')
print('Number of data points : ', data.shape[0])
print('Number of features : ', data.shape[1])
print('Features : ', data.columns.values)
data.head()

Number of data points : 3321
Number of features : 4
Features : ['ID' 'Gene' 'Variation' 'Class']
```

#### Out[0]:

	ID	Gene	Variation	Class
0	0	FAM58A	Truncating Mutations	1
1	1	CBL	W802*	2
2	2	CBL	Q249E	2
3	3	CBL	N454D	3
4	4	CBL	L399V	4

training/training\_variants is a comma separated file containing the description of the genetic mutations used for training. Fields are

- ID: the id of the row used to link the mutation to the clinical evidence
- Gene: the gene where this genetic mutation is located
- Variation: the aminoacid change for this mutations
- Class: 1-9 the class this genetic mutation has been classified on

# 3.1.2. Reading Text Data

#### In [0]:

```
# note the seprator in this file
data_text =pd.read_csv("/content/drive/My Drive/cd/training/training_text", sep="\\\", engine="python", n
ames=["ID", "TEXT"], skiprows=1)
print('Number of data points : ', data_text.shape[0])
print('Number of features : ', data_text.shape[1])
print('Features : ', data_text.columns.values)
data_text.head()

Number of data points : 3321
Number of features : 2
Features : ['ID' 'TEXT']
```

# Out[0]:

	ID	TEXT
0	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	4	Oncogenic mutations in the monomeric Casitas B

# 3.1.3. Preprocessing of text

```
In [0]:
import nltk
nltk.download('stopwords')

[nltk_data] Downloading package stopwords to /root/nltk_data...
[nltk_data] Package stopwords is already up-to-date!

Out[0]:
True
```

```
# loading stop words from nltk library
stop words = set(stopwords.words('english'))
def nlp preprocessing(total text, index, column):
   if type(total_text) is not int:
       string = ""
        # replace every special char with space
       total text = re.sub('[^a-zA-z0-9]', '', total text)
        # replace multiple spaces with single space
       total text = re.sub('\s+',' ', total text)
        # converting all the chars into lower-case.
       total text = total text.lower()
       for word in total text.split():
        # if the word is a not a stop word then retain that word from the data
           if not word in stop words:
               string += word + " "
       data_text[column][index] = string
```

# In [0]:

```
#text processing stage.
start_time = time.clock()
for index, row in data_text.iterrows():
    if type(row['TEXT']) is str:
        nlp_preprocessing(row['TEXT'], index, 'TEXT')
    else:
        print("there is no text description for id:",index)
print('Time took for preprocessing the text :',time.clock() - start_time, "seconds")

there is no text description for id: 1109
there is no text description for id: 1277
there is no text description for id: 1407
there is no text description for id: 1639
there is no text description for id: 2755
Time took for preprocessing the text : 24.682213 seconds
```

#### In [0]:

```
#merging both gene_variations and text data based on ID
result = pd.merge(data, data_text,on='ID', how='left')
result.head()
```

### Out[0]:

	ID	Gene	Variation	Class	TEXT
0	0	FAM58A	Truncating Mutations	1	cyclin dependent kinases cdks regulate variety
1	1	CBL	W802*	2	abstract background non small cell lung cancer
2	2	CBL	Q249E	2	abstract background non small cell lung cancer
3	3	CBL	N454D	3	recent evidence demonstrated acquired uniparen
4	4	CBL	L399V	4	oncogenic mutations monomeric casitas b lineag

```
result[result.isnull().any(axis=1)]
```

#### Out[0]:

	ID	Gene	Variation	Class	TEXT
1109	1109	FANCA	S1088F	1	NaN
1277	1277	ARID5B	Truncating Mutations	1	NaN
1407	1407	FGFR3	K508M	6	NaN
1639	1639	FLT1	Amplification	6	NaN
2755	2755	BRAF	G596C	7	NaN

#### In [0]:

```
result.loc[result['TEXT'].isnull(),'TEXT'] = result['Gene'] +' '+result['Variation']
```

#### In [0]:

```
result[result['ID']==1109]
```

#### Out[0]:

	ID	Gene	Variation	Class	TEXT
1109	1109	FANCA	S1088F	1	FANCA S1088F

# 3.1.4. Test, Train and Cross Validation Split

#### 3.1.4.1. Splitting data into train, test and cross validation (64:20:16)

# In [0]:

```
y_true = result['Class'].values
            = result.Gene.str.replace('\s+', '')
result.Gene
result.Variation = result.Variation.str.replace('\s+', '')
# split the data into test and train by maintaining same distribution of output varaible 'y true' [stra
tify=y true]
X_train, test_df, y_train, y_test = train_test_split(result, y_true, stratify=y_true, test_size=0.2)
# split the train data into train and cross validation by maintaining same distribution of output varai
ble 'y train' [stratify=y train]
train df, cv df, y train, y cv = train test split(X train, y train, stratify=y train, test size=0.2)
```

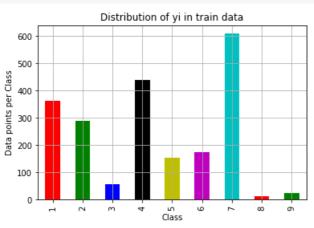
We split the data into train, test and cross validation data sets, preserving the ratio of class distribution in the original data set

# In [0]:

```
print('Number of data points in train data:', train df.shape[0])
print('Number of data points in test data:', test_df.shape[0])
print('Number of data points in cross validation data:', cv df.shape[0])
Number of data points in train data: 2124
Number of data points in test data: 665
Number of data points in cross validation data: 532
```

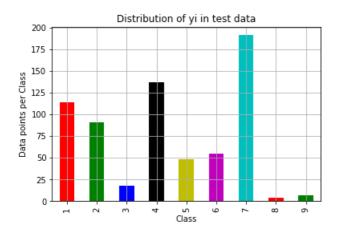
#### 3.1.4.2. Distribution of y\_i's in Train, Test and Cross Validation datasets

```
# it returns a dict, keys as class labels and values as the number of data points in that class
train class distribution = train df['Class'].value counts().sort index()
test class distribution = test df['Class'].value counts().sort index()
cv class distribution = cv df['Class'].value counts().sort index()
my colors = ['r','g','b','k','y','m','c']
train class distribution.plot(kind='bar',color=my colors)
plt.xlabel('Class')
plt.ylabel('Data points per Class')
plt.title('Distribution of yi in train data')
plt.grid()
plt.show()
{\it \# ref: argsort https://docs.scipy.org/doc/numpy/reference/generated/numpy.argsort.html}
# -(train class distribution.values): the minus sign will give us in decreasing order
sorted yi = np.argsort(-train class distribution.values)
for i in sorted yi:
   print('Number of data points in class', i+1, ':', train class distribution.values[i], '(', np.round(
(train class distribution.values[i]/train df.shape[0]*100), 3), '%)')
print('-'*80)
#my colors = 'rgbkymc'
test_class_distribution.plot(kind='bar',color=my_colors)
plt.xlabel('Class')
plt.ylabel('Data points per Class')
plt.title('Distribution of yi in test data')
plt.grid()
plt.show()
# ref: argsort https://docs.scipy.org/doc/numpy/reference/generated/numpy.argsort.html
# -(train class distribution.values): the minus sign will give us in decreasing order
sorted yi = np.argsort(-test class distribution.values)
for i in sorted_yi:
   print('Number of data points in class', i+1, ':', test class distribution.values[i], '(', np.round((
test class distribution.values[i]/test df.shape[0]*100), 3), '%)')
print('-'*80)
#my colors = 'rgbkymc'
cv_class_distribution.plot(kind='bar',color=my_colors)
plt.xlabel('Class')
plt.ylabel('Data points per Class')
plt.title('Distribution of yi in cross validation data')
plt.grid()
plt.show()
# ref: argsort https://docs.scipy.org/doc/numpy/reference/generated/numpy.argsort.html
# -(train class distribution.values): the minus sign will give us in decreasing order
sorted yi = np.argsort(-train class distribution.values)
for i in sorted yi:
   print('Number of data points in class', i+1, ':', cv class distribution.values[i], '(', np.round((cv
class distribution.values[i]/cv df.shape[0]*100), 3), '%)')
```



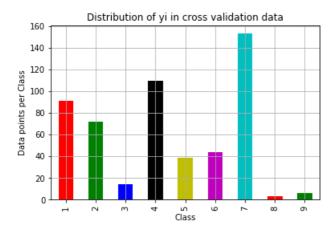
```
Number of data points in class 4 : 439 ( 20.669 \%) Number of data points in class 1 : 363 ( 17.09 \%) Number of data points in class 2 : 289 \% ( 13.606 \%) Number of data points in class 6 : 176 \% ( 8.286 \%) Number of data points in class 5 : 155 \% ( 7.298 \%) Number of data points in class 3 : 57 \% ( 2.684 \%) Number of data points in class 9 : 24 \% ( 1.13 \%) Number of data points in class 8 : 12 \%
```

\_\_\_\_\_\_



```
Number of data points in class 7 : 191 ( 28.722 \%) Number of data points in class 4 : 137 ( 20.602 \%) Number of data points in class 1 : 114 ( 17.143 \%) Number of data points in class 2 : 91 ( 13.684 \%) Number of data points in class 6 : 55 ( 8.271 \%) Number of data points in class 5 : 48 ( 7.218 \%) Number of data points in class 3 : 18 ( 2.707 \%) Number of data points in class 9 : 7 ( 1.053 \%) Number of data points in class 8 : 4 ( 0.602 \%)
```

\_\_\_\_\_



```
Number of data points in class 7 : 153 ( 28.759 \%) Number of data points in class 4 : 110 ( 20.677 \%) Number of data points in class 1 : 91 ( 17.105 \%) Number of data points in class 2 : 72 ( 13.534 \%) Number of data points in class 6 : 44 ( 8.271 \%) Number of data points in class 5 : 39 ( 7.331 \%) Number of data points in class 3 : 14 ( 2.632 \%) Number of data points in class 9 : 6 ( 1.128 \%) Number of data points in class 8 : 3 ( 0.564 \%)
```

# 3.2 Prediction using a 'Random' Model

In a 'Random' Model, we generate the NINE class probabilites randomly such that they sum to 1.

```
# This function plots the confusion matrices given y i, y i hat.
def plot confusion matrix(test y, predict y):
   C = confusion_matrix(test_y, predict_y)
    # C = 9,9 matrix, each cell (i,j) represents number of points of class i are predicted class j
   A = (((C.T) / (C.sum(axis=1))).T)
    #divid each element of the confusion matrix with the sum of elements in that column
    \# C = [[1, 2],
         [3, 4]]
   # C.T = [[1, 3],
            [2, 4]]
   # C.sum(axis = 1) axis=0 corresonds to columns and axis=1 corresponds to rows in two diamensional
arrav
   \# C.sum(axix = 1) = [[3, 7]]
   \# ((C.T)/(C.sum(axis=1))) = [[1/3, 3/7]
    \# ((C.T)/(C.sum(axis=1))).T = [[1/3, 2/3]
                                [3/7, 4/7]]
    # sum of row elements = 1
   B = (C/C.sum(axis=0))
   #divid each element of the confusion matrix with the sum of elements in that row
    \# C = [[1, 2],
         [3, 4]]
   # C.sum(axis = 0) axis=0 corresponds to columns and axis=1 corresponds to rows in two diamensional
array
   \# C.sum(axix = 0) = [[4, 6]]
   \# (C/C.sum(axis=0)) = [[1/4, 2/6],
                           [3/4, 4/6]]
   labels = [1,2,3,4,5,6,7,8,9]
    # representing A in heatmap format
   print("-"*20, "Confusion matrix", "-"*20)
   plt.figure(figsize=(20,7))
   sns.heatmap(C, annot=True, cmap="YlGnBu", fmt=".3f", xticklabels=labels, yticklabels=labels)
   plt.xlabel('Predicted Class')
   plt.ylabel('Original Class')
   plt.show()
   print("-"*20, "Precision matrix (Column Sum=1)", "-"*20)
   plt.figure(figsize=(20,7))
   sns.heatmap(B, annot=True, cmap="YlGnBu", fmt=".3f", xticklabels=labels, yticklabels=labels)
   plt.xlabel('Predicted Class')
   plt.ylabel('Original Class')
   plt.show()
    # representing B in heatmap format
   print("-"*20, "Recall matrix (Row sum=1)", "-"*20)
   plt.figure(figsize=(20,7))
   sns.heatmap(A, annot=True, cmap="YlGnBu", fmt=".3f", xticklabels=labels, yticklabels=labels)
   plt.xlabel('Predicted Class')
   plt.ylabel('Original Class')
   plt.show()
```

```
# we need to generate 9 numbers and the sum of numbers should be 1
# one solution is to genarate 9 numbers and divide each of the numbers by their sum
# ref: https://stackoverflow.com/a/18662466/4084039
test_data_len = test_df.shape[0]
cv_data_len = cv_df.shape[0]

# we create a output array that has exactly same size as the CV data
cv_predicted_y = np.zeros((cv_data_len,9))
for i in range(cv_data_len):
    rand_probs = np.random.rand(1,9)
    cv_predicted_y[i] = ((rand_probs/sum(sum(rand_probs)))[0])
print("Log_loss_on_Cross_Validation_Data_using_Random_Model",log_loss(y_cv,cv_predicted_y, eps=le-15))

# Test-Set_error.
#we create a output array that has exactly same as the test_data
test_predicted_y = np.zeros((test_data_len,9))
for i in range(test_data_len):
```

```
rand_probs = np.random.rand(1,9)
  test_predicted_y[i] = ((rand_probs/sum(sum(rand_probs)))[0])
print("Log_loss on Test_Data_using Random Model",log_loss(y_test,test_predicted_y, eps=1e-15))
predicted_y =np.argmax(test_predicted_y, axis=1)
plot_confusion_matrix(y_test, predicted_y+1)
```

- 20

- 15

- 10

- 5

0.35

- 0.30

- 0.25

- 0.20

-0.15

-0.10

- 0.05

- 0.00

0.5

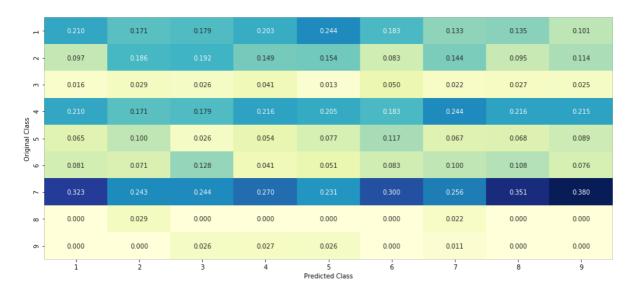
- 0.4

- 0.3

Log loss on Cross Validation Data using Random Model 2.4066319305249415 Log loss on Test Data using Random Model 2.4942359949277337 ------ Confusion matrix ------

	13.000	12.000	14.000		19.000	11.000	12.000	10.000	8.000
- 2	6.000	13.000		11.000	12.000	5.000	13.000	7.000	9.000
m -	1.000	2.000	2.000	3.000	1.000	3.000	2.000	2.000	2.000
SS 4 -	13.000	12.000	14.000		16.000	11.000	22.000	16.000	17.000
Original Class 5	4.000	7.000	2.000	4.000	6.000	7.000	6.000	5.000	7.000
oric	5.000	5.000	10.000	3.000	4.000	5.000	9.000	8.000	6.000
۲.	20.000		19.000	20.000	18.000	18.000	23.000	26.000	30.000
ω -	0.000	2.000	0.000	0.000	0.000	0.000	2.000	0.000	0.000
თ -	0.000	0.000	2.000	2.000	2.000	0.000	1.000	0.000	0.000
	i	2	3	4	5 Predicted Class	6	7	8	9

----- Precision matrix (Columm Sum=1) -----



----- Recall matrix (Row sum=1) -----

r-1	0.114	0.105	0.123	0.132	0.167	0.096	0.105	0.088	0.070
- 5	0.066	0.143	0.165	0.121	0.132	0.055	0.143	0.077	0.099
m -	0.056	0.111	0.111	0.167	0.056	0.167	0.111	0.111	0.111
- 4 -	0.095	0.088	0.102	0.117	0.117	0.080	0.161	0.117	0.124
Original Class 5	0.083	0.146	0.042	0.083	0.125	0.146	0.125	0.104	0.146
Oric	0.091	0.091	0.182	0.055	0.073	0.091	0.164	0.145	0.109
۲-	0.105	0.089	0.099	0.105	0.094	0.094	0.120	0.136	0.157
∞ -	0.000	0.500	0.000	0.000	0.000	0.000	0.500	0.000	0.000

# 3.3 Univariate Analysis

```
In [0]:
```

```
# code for response coding with Laplace smoothing.
# alpha: used for laplace smoothing
# feature: ['gene', 'variation']
# df: ['train_df', 'test_df', 'cv_df']
# algorithm
# Consider all unique values and the number of occurances of given feature in train data dataframe
# build a vector (1*9) , the first element = (number of times it occured in class1 + 10*alpha / number
of time it occurred in total data+90*alpha)
# gv_dict is like a look up table, for every gene it store a (1*9) representation of it
# for a value of feature in df:
# if it is in train data:
# we add the vector that was stored in 'gv dict' look up table to 'gv fea'
# if it is not there is train:
# we add [1/9, 1/9, 1/9, 1/9, 1/9, 1/9, 1/9] to 'gv fea'
# return 'gv fea'
# get gv fea dict: Get Gene varaition Feature Dict
def get gv fea dict(alpha, feature, df):
    # value count: it contains a dict like
    # print(train_df['Gene'].value_counts())
    # output:
             {BRCA1
                         174
              TP5.3
                         106
              EGFR
                          75
              BRCA2
                          69
              PTEN
              KIT
                          60
              ERBB2
                          47
              PDGFRA
                          46
              ...}
    # print(train df['Variation'].value counts())
    # output:
    # Truncating Mutations
                                                63
                                                4.3
    # Deletion
    # Amplification
                                                43
    # Fusions
    # Overexpression
                                                 3
    # E17K
                                                 3
    # Q61L
                                                 3
                                                 2
    # S222E
    # P130S
    # }
    value count = train df[feature].value counts()
    # gv dict : Gene Variation Dict, which contains the probability array for each gene/variation
    gv dict = dict()
    # denominator will contain the number of time that particular feature occured in whole data
    for i, denominator in value count.items():
        # vec will contain (p(yi==1/Gi) probability of gene/variation belongs to perticular class
        # vec is 9 diamensional vector
        vec = []
        for k in range(1,10):
            # print(train_df.loc[(train_df['Class']==1) & (train_df['Gene']=='BRCA1')])
                      ID
                          Gene
                                             Variation Class
            # 2470 2470 BRCA1
                                               S1715C
            # 2486 2486 BRCA1
                                                S1841R
                                                            7
            # 2614 2614 BRCA1
                                                   M1R
            # 2432 2432 BRCA1
                                                L1657P
                                                            7
            # 2567 2567 BRCA1
                                                T1685A
```

```
# 2583 2583 BRCA1
                                              E1660G
            # 2634 2634 BRCA1
                                               W1718L
            # cls cnt.shape[0] will return the number of rows
            cls cnt = train df.loc[(train df['Class']==k) & (train df[feature]==i)]
            # cls cnt.shape[0](numerator) will contain the number of time that particular feature occur
ed in whole data
           vec.append((cls cnt.shape[0] + alpha*10)/ (denominator + 90*alpha))
        # we are adding the gene/variation to the dict as key and vec as value
       gv dict[i]=vec
   return gv dict
# Get Gene variation feature
def get gv feature (alpha, feature, df):
    # print(gv dict)
        {'BRCA1': [0.2007575757575757575, 0.037878787878788, 0.0681818181818177, 0.1363636363636363
5, 0.25, 0.1931818181818181818, 0.03787878787878788, 0.03787878787878, 0.037878787878787878),
         'TP53': [0.32142857142857145, 0.061224489795918366, 0.061224489795918366, 0.2704081632653061
5, 0.061224489795918366, 0.066326530612244902, 0.051020408163265307, 0.051020408163265307, 0.0561224489
79591837],
          'EGFR': [0.056818181818181816, 0.21590909090909091, 0.0625, 0.068181818181818177, 0.06818181
   #
818181877, 0.0625, 0.34659090909090912, 0.0625, 0.056818181818181816],
          'BRCA2': [0.13333333333333333, 0.0606060606060608, 0.0606060606060608, 0.0787878787878
782, 0.13939393939394, 0.34545454545454546, 0.0606060606060608, 0.06060606060608, 0.06060606060
6060608],
           'PTEN': [0.069182389937106917, 0.062893081761006289, 0.069182389937106917, 0.465408805031446
55, 0.075471698113207544, 0.062893081761006289, 0.069182389937106917, 0.062893081761006289, 0.062893081
761006289],
# 'KIT': [0.066225165562913912, 0.25165562913907286, 0.072847682119205295, 0.07284768211920529
# 0.066225165562913912, 0.06622516556
5, 0.066225165562913912, 0.066225165562913912, 0.27152317880794702, 0.066225165562913912, 0.06622516556
           'BRAF': [0.066666666666666666, 0.17999999999999, 0.073333333333334, 0.07333333333333
   #
34, 0.09333333333333338, 0.08000000000000000, 0.2999999999999, 0.06666666666666666, 0.0666666666
666666661,
   #
   #
   gv dict = get gv fea dict(alpha, feature, df)
   # value count is similar in get gv fea dict
   value count = train df[feature].value counts()
    # gv fea: Gene variation feature, it will contain the feature for each feature value in the data
   gv fea = []
    # for every feature values in the given data frame we will check if it is there in the train data t
hen we will add the feature to gv fea
    # if not we will add [1/9,1/9,1/9,1/9,1/9,1/9,1/9,1/9] to gv fea
   for index, row in df.iterrows():
       if row[feature] in dict(value count).keys():
           gv_fea.append(gv_dict[row[feature]])
           gv fea.append([1/9,1/9,1/9,1/9,1/9,1/9,1/9,1/9])
             gv fea.append([-1,-1,-1,-1,-1,-1,-1,-1])
   return gv fea
```

when we caculate the probability of a feature belongs to any particular class, we apply laplace smoothing

• (numerator + 10\\*alpha) / (denominator + 90\\*alpha)

#### 3.2.1 Univariate Analysis on Gene Feature

Q1. Gene, What type of feature it is?

Ans. Gene is a categorical variable

Q2. How many categories are there and How they are distributed?

```
In [0]:
```

```
unique_genes = train_df['Gene'].value_counts()
print('Number of Unique Genes :', unique_genes.shape[0])
# the top 10 genes that occured most
print(unique_genes.head(10))
```

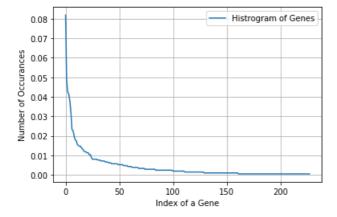
```
Number of Unique Genes : 228
          174
BRCA1
TP53
          103
           90
EGFR
BRCA2
           88
PTEN
           80
           68
KIT
ERBB2
           49
BRAF
           48
           42
ALK
CDKN2A
           38
Name: Gene, dtype: int64
```

```
print("Ans: There are", unique_genes.shape[0] ,"different categories of genes in the train data, and th
ey are distibuted as follows",)
```

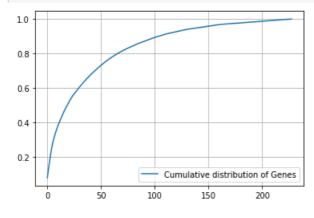
Ans: There are 228 different categories of genes in the train data, and they are distibuted as follows

## In [0]:

```
s = sum(unique_genes.values);
h = unique_genes.values/s;
plt.plot(h, label="Histrogram of Genes")
plt.xlabel('Index of a Gene')
plt.ylabel('Number of Occurances')
plt.legend()
plt.grid()
plt.show()
```



```
c = np.cumsum(h)
plt.plot(c,label='Cumulative distribution of Genes')
plt.grid()
plt.legend()
plt.show()
```



#### Q3. How to featurize this Gene feature?

Ans.there are two ways we can featurize this variable check out this video:

https://www.appliedaicourse.com/course/applied-ai-course-online/lessons/handling-categorical-and-numerical-features/

- 1. One hot Encoding
- 2. Response coding

We will choose the appropriate featurization based on the ML model we use. For this problem of multi-class classification with categorical features, one-hot encoding is better for Logistic regression while response coding is better for Random Forests.

```
#response-coding of the Gene feature
# alpha is used for laplace smoothing
alpha = 1
# train gene feature
train gene feature responseCoding = np.array(get gv feature(alpha, "Gene", train df))
# test gene feature
test_gene_feature_responseCoding = np.array(get_gv_feature(alpha, "Gene", test_df))
# cross validation gene feature
cv gene feature responseCoding = np.array(get gv feature(alpha, "Gene", cv df))
```

#### In [0]:

```
print ("train gene feature responseCoding is converted feature using respone coding method. The shape of
gene feature:", train gene feature responseCoding.shape)
```

train gene feature responseCoding is converted feature using respone coding method. The shape of gene f eature: (2124, 9)

#### In [0]:

```
# one-hot encoding of Gene feature.
gene vectorizer = CountVectorizer()
train gene feature onehotCoding = gene vectorizer.fit transform(train df['Gene'])
test_gene_feature_onehotCoding = gene_vectorizer.transform(test_df['Gene'])
cv_gene_feature_onehotCoding = gene_vectorizer.transform(cv df['Gene'])
```

#### In [0]:

```
train df['Gene'].head()
```

# Out[0]:

```
1546
       ALK
     BRCA1
2649
1302
       MLH1
      SMAD3
559
2496
      BRCA1
Name: Gene, dtype: object
```

```
gene vectorizer.get feature names()
```

```
Out[0]:
```

```
['abl1',
 'acvr1',
 'ago2',
 'akt1',
'akt2',
'akt3',
 'alk',
 'apc',
 'ar',
 'araf',
 'aridla',
```

```
'aridlb',
'arid2',
'arid5b',
'asxl2',
'atm',
'aurka',
'aurkb',
'axl',
'b2m',
'bap1',
'bard1',
'bcl2111',
'bcor',
'braf',
'brcal',
'brca2',
'brd4',
'brip1',
'btk',
'card11',
'carm1',
'casp8',
'cbl',
'ccnd1',
'ccnd2',
'ccnd3',
'cdh1',
'cdk12',
'cdk6',
'cdk8',
'cdknla',
'cdkn1b',
'cdkn2a',
'cdkn2b',
'cdkn2c',
'cebpa',
'chek2',
'cic',
'crebbp',
'ctcf',
'ctla4',
'ctnnb1',
'ddr2',
'dicer1',
'dnmt3a',
'dnmt3b',
'dusp4',
'egfr',
'eiflax',
'elf3',
'ep300',
'epas1',
'epcam',
'erbb2',
'erbb3',
'erbb4',
'ercc2',
'ercc3',
'ercc4',
'erg',
'esr1',
'etv1',
'etv6',
'ewsr1',
'ezh2',
'fam58a',
'fanca',
'fat1',
'fbxw7',
'fgf3',
'fgfr1',
'fgfr2',
'fgfr3',
'flt1',
'flt3',
'foxal'.
```

```
'foxp1',
'gata3',
'glil',
'gna11',
'gnaq',
'gnas',
'h3f3a',
'hist1h1c',
'hla',
'hnfla',
'hras',
'idh1',
'idh2',
'igf1r',
'ikbke',
'il7r',
'inpp4b',
'jak1',
'jak2',
'kdm5a',
'kdm5c',
'kdm6a',
'kdr',
'keap1',
'kit',
'kmt2a',
'kmt2b',
'kmt2c',
'kmt2d',
'knstrn',
'kras',
'lats1',
'lats2',
'map2k1',
'map2k2',
'map2k4',
'map3k1',
'mapk1',
'mdm2',
'med12',
'mef2b',
'met',
'mlh1',
'mpl',
'msh2',
'msh6',
'mtor',
'myc',
'mycn',
'myd88',
'ncor1',
'nf1',
'nf2',
'nfe212',
'nfkbia',
'nkx2',
'notch1',
'notch2',
'npm1',
'nras',
'nsd1',
'ntrk1',
'ntrk2',
'ntrk3',
'nup93',
'pbrml',
'pdgfra',
'pdgfrb',
'pik3ca',
'pik3cb',
'pik3cd',
'pik3r1',
'pik3r2',
'pik3r3',
'pim1',
'pms1',
'nms2'.
```

```
'pole',
 'ppp2r1a',
 'ppp6c',
 'prdm1',
 'pten',
 'ptpn11',
 'ptprd',
 'ptprt',
 'rab35',
 'rac1',
 'rad21',
 'rad50',
 'rad51b',
 'rad51c',
 'raf1',
 'rasal',
 'rb1',
 'rbm10',
 'ret',
 'rheb',
 'rhoa',
 'rictor',
 'rit1',
 'rnf43',
 'ros1',
 'runx1',
 'rxra',
 'rybp',
 'sdhb',
 'setd2',
 'sf3b1',
 'smad2',
 'smad3',
 'smad4',
 'smarca4',
 'smarcb1',
 'smo',
'sos1',
 'sox9',
 'spop',
 'src',
 'srsf2',
 'stag2',
 'stat3',
 'stk11',
 'tcf3',
 'tcf712',
 'tert',
 'tet1',
 'tet2',
 'tgfbr1',
 'tgfbr2',
 'tmprss2',
 'tp53',
 'tp53bp1',
 'tsc1',
 'tsc2',
 'u2af1',
 'vhl',
 'whsc111',
 'xpol',
 'xrcc2',
 'yap1']
In [0]:
```

print("train\_gene\_feature\_onehotCoding is converted feature using one-hot encoding method. The shape of gene feature:", train\_gene\_feature\_onehotCoding.shape)

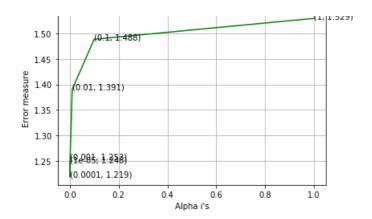
train\_gene\_feature\_onehotCoding is converted feature using one-hot encoding method. The shape of gene f eature: (2124, 228)

# **Q4.** How good is this gene feature in predicting y i?

There are many ways to estimate how good a feature is, in predicting y\_i. One of the good methods is to build a proper ML model using just this feature. In this case, we will build a logistic regression model using only Gene feature (one hot encoded) to predict y\_i.

```
In [0]:
```

```
alpha = [10 ** x for x in range(-5, 1)] # hyperparam for SGD classifier.
# read more about SGDClassifier() at http://scikit-learn.org/stable/modules/generated/sklearn.linear mo
del.SGDClassifier.html
# default parameters
# SGDClassifier(loss='hinge', penalty='12', alpha=0.0001, 11 ratio=0.15, fit intercept=True, max iter=N
one, tol=None,
# shuffle=True, verbose=0, epsilon=0.1, n jobs=1, random state=None, learning rate='optimal', eta0=0.0,
power t=0.5.
# class weight=None, warm start=False, average=False, n iter=None)
# some of methods
# fit(X, y[, coef init, intercept init, ...]) Fit linear model with Stochastic Gradient Descent.
\# predict(X) Predict class labels for samples in X.
# video link:
cv log error array=[]
for i in alpha:
   clf = SGDClassifier(alpha=i, penalty='12', loss='log', random state=42)
    clf.fit(train_gene_feature_onehotCoding, y_train)
    sig clf = CalibratedClassifierCV(clf, method="sigmoid")
    sig clf.fit(train gene feature onehotCoding, y train)
   predict y = sig clf.predict proba(cv gene feature onehotCoding)
    cv log error array.append(log loss(y cv, predict y, labels=clf.classes , eps=1e-15))
   print ('For values of alpha = ', i, "The log loss is:", log_loss(y_cv, predict_y, labels=clf.classes
, eps=1e-15))
fig, ax = plt.subplots()
ax.plot(alpha, cv log error array, c='g')
for i, txt in enumerate(np.round(cv_log_error_array,3)):
   ax.annotate((alpha[i],np.round(txt,3)), (alpha[i],cv_log_error_array[i]))
plt.grid()
plt.title("Cross Validation Error for each alpha")
plt.xlabel("Alpha i's")
plt.ylabel("Error measure")
plt.show()
best alpha = np.argmin(cv log error array)
clf = SGDClassifier(alpha=alpha[best alpha], penalty='12', loss='log', random state=42)
clf.fit(train gene feature onehotCoding, y train)
sig_clf = CalibratedClassifierCV(clf, method="sigmoid")
sig_clf.fit(train_gene_feature_onehotCoding, y_train)
predict y = sig clf.predict proba(train gene feature onehotCoding)
print('For values of best alpha = ', alpha[best alpha], "The train log loss is:", log loss (y train, pred
ict_y, labels=clf.classes_, eps=1e-15))
predict_y = sig_clf.predict_proba(cv_gene_feature_onehotCoding)
print('For values of best alpha = ', alpha[best_alpha], "The cross validation log loss is:",log_loss(y_
cv, predict_y, labels=clf.classes_, eps=1e-15))
predict y = sig clf.predict proba(test gene feature onehotCoding)
print('For values of best alpha = ', alpha[best alpha], "The test log loss is:", log loss(y test, predic
t_y, labels=clf.classes_, eps=1e-15))
For values of alpha = 1e-05 The log loss is: 1.2479139797072707
For values of alpha = 0.0001 The log loss is: 1.2189525488162916
For values of alpha = 0.001 The log loss is: 1.253279979892459
For values of alpha = 0.01 The log loss is: 1.3906735591825905
For values of alpha = 0.1 The log loss is: 1.4883831234882048
For values of alpha = 1 The log loss is: 1.5292272790268189
```



```
For values of best alpha = 0.0001 The train log loss is: 0.9728991501977299

For values of best alpha = 0.0001 The cross validation log loss is: 1.2189525488162916

For values of best alpha = 0.0001 The test log loss is: 1.2316177101876011
```

# Q5. Is the Gene feature stable across all the data sets (Test, Train, Cross validation)?

Ans. Yes, it is. Otherwise, the CV and Test errors would be significantly more than train error.

#### In [0]:

```
print("Q6. How many data points in Test and CV datasets are covered by the ", unique_genes.shape[0], "
genes in train dataset?")

test_coverage=test_df[test_df['Gene'].isin(list(set(train_df['Gene'])))].shape[0]

cv_coverage=cv_df[cv_df['Gene'].isin(list(set(train_df['Gene'])))].shape[0]

print('Ans\n1. In test data', test_coverage, 'out of', test_df.shape[0], ":", (test_coverage/test_df.shape
[0])*100)

print('2. In cross validation data', cv_coverage, 'out of ', cv_df.shape[0], ":", (cv_coverage/cv_df.shape
[0])*100)
```

Q6. How many data points in Test and CV datasets are covered by the 228 genes in train dataset? Ans

- 1. In test data 637 out of 665 : 95.78947368421052
- 2. In cross validation data 515 out of 532: 96.80451127819549

## 3.2.2 Univariate Analysis on Variation Feature

Q7. Variation, What type of feature is it?

Ans. Variation is a categorical variable

**Q8.** How many categories are there?

```
unique_variations = train_df['Variation'].value_counts()
print('Number of Unique Variations :', unique_variations.shape[0])
# the top 10 variations that occured most
print(unique_variations.head(10))
```

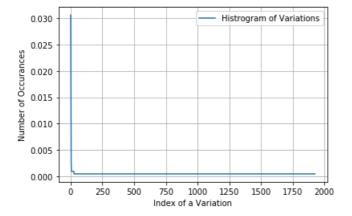
```
Number of Unique Variations: 1925
Truncating Mutations
Amplification
                              50
                              45
Deletion
Fusions
                              17
Overexpression
G12V
                               3
C618R
G13V
ETV6-NTRK3 Fusion
Promoter Hypermethylation
Name: Variation, dtype: int64
```

```
print("Ans: There are", unique_variations.shape[0] ,"different categories of variations in the train da ta, and they are distibuted as follows",)
```

Ans: There are 1925 different categories of variations in the train data, and they are distibuted as fo llows

#### In [0]:

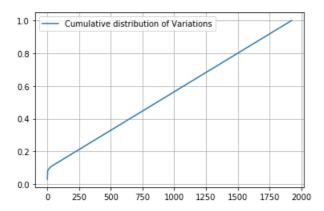
```
s = sum(unique_variations.values);
h = unique_variations.values/s;
plt.plot(h, label="Histrogram of Variations")
plt.xlabel('Index of a Variation')
plt.ylabel('Number of Occurances')
plt.legend()
plt.grid()
plt.show()
```



# In [0]:

```
c = np.cumsum(h)
print(c)
plt.plot(c,label='Cumulative distribution of Variations')
plt.grid()
plt.legend()
plt.show()
```

```
[0.03060264 0.05414313 0.07532957 ... 0.99905838 0.99952919 1.
```



#### **Q9.** How to featurize this Variation feature?

**Ans.**There are two ways we can featurize this variable check out this video: https://www.appliedaicourse.com/course/applied-ai-course-online/lessons/handling-categorical-and-numerical-features/

- 1. One hot Encoding
- 2. Response coding

```
# alpha is used for laplace smoothing
alpha = 1
# train gene feature
train_variation_feature_responseCoding = np.array(get_gv_feature(alpha, "Variation", train_df))
# test gene feature
test_variation_feature_responseCoding = np.array(get_gv_feature(alpha, "Variation", test_df))
# cross validation gene feature
cv_variation_feature_responseCoding = np.array(get_gv_feature(alpha, "Variation", cv_df))
```

## In [0]:

```
print("train_variation_feature_responseCoding is a converted feature using the response coding method.
The shape of Variation feature:", train_variation_feature_responseCoding.shape)
```

train\_variation\_feature\_responseCoding is a converted feature using the response coding method. The sha pe of Variation feature: (2124, 9)

#### In [0]:

```
# one-hot encoding of variation feature.
variation_vectorizer = CountVectorizer()
train_variation_feature_onehotCoding = variation_vectorizer.fit_transform(train_df['Variation'])
test_variation_feature_onehotCoding = variation_vectorizer.transform(test_df['Variation'])
cv_variation_feature_onehotCoding = variation_vectorizer.transform(cv_df['Variation'])
```

#### In [0]:

```
print("train_variation_feature_onehotEncoded is converted feature using the onne-hot encoding method. T
he shape of Variation feature:", train_variation_feature_onehotCoding.shape)
```

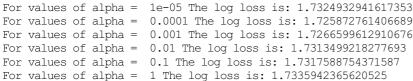
train\_variation\_feature\_onehotEncoded is converted feature using the onne-hot encoding method. The shap e of Variation feature: (2124, 1954)

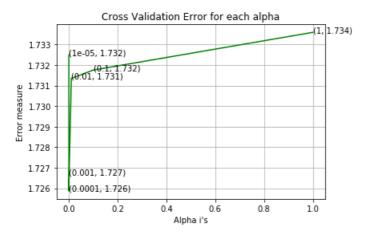
## **Q10.** How good is this Variation feature in predicting y i?

Let's build a model just like the earlier!

```
alpha = [10 ** x for x in range(-5, 1)]
# read more about SGDClassifier() at http://scikit-learn.org/stable/modules/generated/sklearn.linear mo
del.SGDClassifier.html
# default parameters
# SGDClassifier(loss='hinge', penalty='12', alpha=0.0001, 11 ratio=0.15, fit intercept=True, max iter=N
one, tol=None,
# shuffle=True, verbose=0, epsilon=0.1, n jobs=1, random state=None, learning rate='optimal', eta0=0.0,
power t=0.5,
# class weight=None, warm start=False, average=False, n iter=None)
# some of methods
# fit(X, y[, coef init, intercept init, ...]) Fit linear model with Stochastic Gradient Descent.
# predict(X) Predict class labels for samples in X.
# video link:
cv log error array=[]
for i in alpha:
   clf = SGDClassifier(alpha=i, penalty='12', loss='log', random state=42)
   clf.fit(train variation feature onehotCoding, y train)
```

```
sig clf = CalibratedClassifierCV(clf, method="sigmoid")
    sig clf.fit(train variation feature onehotCoding, y train)
    predict_y = sig_clf.predict_proba(cv_variation_feature_onehotCoding)
    cv_log_error_array.append(log_loss(y_cv, predict_y, labels=clf.classes , eps=1e-15))
    print('For values of alpha = ', i, "The log loss is:", log loss(y cv, predict y, labels=clf.classes
, eps=1e-15))
fig, ax = plt.subplots()
ax.plot(alpha, cv log error array, c='g')
for i, txt in enumerate(np.round(cv_log_error_array,3)):
   ax.annotate((alpha[i],np.round(txt,3)), (alpha[i],cv log error array[i]))
plt.grid()
plt.title("Cross Validation Error for each alpha")
plt.xlabel("Alpha i's")
plt.ylabel("Error measure")
plt.show()
best alpha = np.argmin(cv log error array)
clf = SGDClassifier(alpha=alpha[best_alpha], penalty='12', loss='log', random state=42)
clf.fit(train variation feature onehotCoding, y train)
sig_clf = CalibratedClassifierCV(clf, method="sigmoid")
sig_clf.fit(train_variation_feature_onehotCoding, y_train)
predict_y = sig_clf.predict_proba(train_variation_feature_onehotCoding)
print('For values of best alpha = ', alpha[best_alpha], "The train log loss is:",log_loss(y_train, pred
ict_y, labels=clf.classes_, eps=1e-15))
predict_y = sig_clf.predict_proba(cv_variation_feature_onehotCoding)
print('For values of best alpha = ', alpha[best alpha], "The cross validation log loss is:", log loss(y
cv, predict_y, labels=clf.classes_, eps=1e-15))
predict_y = sig_clf.predict_proba(test_variation_feature_onehotCoding)
print('For values of best alpha = ', alpha[best_alpha], "The test log loss is:",log loss(y test, predic
t y, labels=clf.classes , eps=1e-15))
For values of alpha = 1e-05 The log loss is: 1.7324932941617353
For values of alpha = 0.0001 The log loss is: 1.7258727614066895
For values of alpha = 0.001 The log loss is: 1.7266599612910676
```





```
For values of best alpha = 0.0001 The train log loss is: 0.6591569225625009
For values of best alpha = 0.0001 The cross validation log loss is: 1.7258727614066895
For values of best alpha = 0.0001 The test log loss is: 1.7052434375675716
```

## **Q11.** Is the Variation feature stable across all the data sets (Test, Train, Cross validation)?

**Ans.** Not sure! But lets be very sure using the below analysis.

```
In [0]:
```

```
print("Q12. How many data points are covered by total ", unique variations.shape[0], " genes in test an
d cross validation data sets?")
test_coverage=test_df[test_df['Variation'].isin(list(set(train_df['Variation'])))].shape[0]
cv coverage=cv df[cv df['Variation'].isin(list(set(train df['Variation'])))].shape[0]
```

```
print('Ans\n1. In test data',test_coverage, 'out of',test_df.shape[0], ":",(test_coverage/test_df.shape
[0])*100)
print('2. In cross validation data',cv_coverage, 'out of ',cv_df.shape[0],":",(cv_coverage/cv_df.shape
[0])*100)
```

Q12. How many data points are covered by total 1925 genes in test and cross validation data sets?

- 1. In test data 72 out of 665 : 10.827067669172932
- 2. In cross validation data 50 out of 532: 9.398496240601503

# 3.2.3 Univariate Analysis on Text Feature

- 1. How many unique words are present in train data?
- 2. How are word frequencies distributed?
- 3. How to featurize text field?
- 4. Is the text feature useful in predicitng y i?
- 5. Is the text feature stable across train, test and CV datasets?

#### In [0]:

#### In [0]:

```
import math
#https://stackoverflow.com/a/1602964
def get_text_responsecoding(df):
    text_feature_responseCoding = np.zeros((df.shape[0],9))
    for i in range(0,9):
        row_index = 0
        for index, row in df.iterrows():
            sum_prob = 0
            for word in row['TEXT'].split():
                  sum_prob += math.log(((dict_list[i].get(word,0)+10 )/(total_dict.get(word,0)+90)))
            text_feature_responseCoding[row_index][i] = math.exp(sum_prob/len(row['TEXT'].split()))
            row_index += 1
    return text_feature_responseCoding
```

#### In [0]:

```
# building a CountVectorizer with all the words that occured minimum 3 times in train data
text_vectorizer = CountVectorizer(ngram_range=(1,2),min_df=3)
train_text_feature_onehotCoding = text_vectorizer.fit_transform(train_df['TEXT'])
# getting all the feature names (words)
train_text_features= text_vectorizer.get_feature_names()

# train_text_feature_onehotCoding.sum(axis=0).Al will sum every row and returns (1*number of features)
vector
train_text_fea_counts = train_text_feature_onehotCoding.sum(axis=0).Al

# zip(list(text_features),text_fea_counts) will zip a word with its number of times it occured
text_fea_dict = dict(zip(list(train_text_features),train_text_fea_counts))

print("Total number of unique words in train data :", len(train_text_features))
```

Total number of unique words in train data : 781887

```
In [0]:
```

```
dict list = []
# dict list =[] contains 9 dictoinaries each corresponds to a class
for i in range (1,10):
   cls text = train df[train df['Class']==i]
    # build a word dict based on the words in that class
   dict list.append(extract dictionary paddle(cls text))
   # append it to dict_list
# dict list[i] is build on i'th class text data
# total dict is buid on whole training text data
total dict = extract dictionary paddle(train df)
confuse array = []
for i in train text features:
   ratios = []
   \max \text{ val} = -1
   for j in range (0,9):
      ratios.append((dict list[j][i]+10 )/(total dict[i]+90))
   confuse_array.append(ratios)
confuse array = np.array(confuse array)
```

```
#response coding of text features
train_text_feature_responseCoding = get_text_responsecoding(train_df)
test_text_feature_responseCoding = get_text_responsecoding(test_df)
cv_text_feature_responseCoding = get_text_responsecoding(cv_df)
```

#### In [0]:

```
# https://stackoverflow.com/a/16202486
# we convert each row values such that they sum to 1
train_text_feature_responseCoding = (train_text_feature_responseCoding.T/train_text_feature_responseCod
ing.sum(axis=1)).T
test_text_feature_responseCoding = (test_text_feature_responseCoding.T/test_text_feature_responseCoding
.sum(axis=1)).T
cv_text_feature_responseCoding = (cv_text_feature_responseCoding.T/cv_text_feature_responseCoding.sum(axis=1)).T
```

## In [0]:

```
# don't forget to normalize every feature
train_text_feature_onehotCoding = normalize(train_text_feature_onehotCoding, axis=0)

# we use the same vectorizer that was trained on train data
test_text_feature_onehotCoding = text_vectorizer.transform(test_df['TEXT'])
# don't forget to normalize every feature
test_text_feature_onehotCoding = normalize(test_text_feature_onehotCoding, axis=0)

# we use the same vectorizer that was trained on train data
cv_text_feature_onehotCoding = text_vectorizer.transform(cv_df['TEXT'])
# don't forget to normalize every feature
cv_text_feature_onehotCoding = normalize(cv_text_feature_onehotCoding, axis=0)
```

# In [0]:

```
#https://stackoverflow.com/a/2258273/4084039
sorted_text_fea_dict = dict(sorted(text_fea_dict.items(), key=lambda x: x[1] , reverse=True))
sorted_text_occur = np.array(list(sorted_text_fea_dict.values()))
```

```
# Number of words for a given frequency.
#print(Counter(sorted_text_occur))
```

```
# Train a Logistic regression+Calibration model using text features whicha re on-hot encoded
alpha = [10 ** x for x in range(-5, 1)]
# read more about SGDClassifier() at http://scikit-learn.org/stable/modules/generated/sklearn.linear mo
del.SGDClassifier.html
# default parameters
# SGDClassifier(loss='hinge', penalty='12', alpha=0.0001, 11 ratio=0.15, fit intercept=True, max iter=N
# shuffle=True, verbose=0, epsilon=0.1, n jobs=1, random state=None, learning rate='optimal', eta0=0.0,
power t=0.5,
# class weight=None, warm start=False, average=False, n iter=None)
# some of methods
# fit(X, y[, coef init, intercept init, ...]) Fit linear model with Stochastic Gradient Descent.
# predict(X) Predict class labels for samples in X.
# video link:
cv_log_error_array=[]
for i in alpha:
    clf = SGDClassifier(alpha=i, penalty='12', loss='log', random_state=42)
    clf.fit(train text feature onehotCoding, y train)
    sig clf = CalibratedClassifierCV(clf, method="sigmoid")
    sig clf.fit(train_text_feature_onehotCoding, y_train)
    predict y = sig clf.predict proba(cv text feature onehotCoding)
    cv log error array.append(log loss(y cv, predict y, labels=clf.classes , eps=1e-15))
   print ('For values of alpha = ', i, "The log loss is:", log loss (y cv, predict y, labels=clf.classes
, eps=1e-15))
fig, ax = plt.subplots()
ax.plot(alpha, cv log error array, c='g')
for i, txt in enumerate(np.round(cv_log_error_array,3)):
   ax.annotate((alpha[i],np.round(txt,3)), (alpha[i],cv log error array[i]))
plt.grid()
plt.title("Cross Validation Error for each alpha")
plt.xlabel("Alpha i's")
plt.ylabel("Error measure")
plt.show()
best_alpha = np.argmin(cv_log_error_array)
clf = SGDClassifier(alpha=alpha[best alpha], penalty='12', loss='log', random state=42)
clf.fit(train text feature onehotCoding, y train)
sig clf = CalibratedClassifierCV(clf, method="sigmoid")
sig_clf.fit(train_text_feature_onehotCoding, y_train)
predict y = sig clf.predict proba(train text feature onehotCoding)
print('For values of best alpha = ', alpha[best_alpha], "The train log loss is:",log_loss(y_train, pred
ict_y, labels=clf.classes_, eps=1e-15))
predict y = sig clf.predict proba(cv text feature onehotCoding)
print('For values of best alpha = ', alpha[best_alpha], "The cross validation log loss is:",log_loss(y_
cv, predict_y, labels=clf.classes_, eps=1e-15))
predict y = sig clf.predict proba(test text feature onehotCoding)
print('For values of best alpha = ', alpha[best alpha], "The test log loss is:",log loss(y test, predic
t_y, labels=clf.classes_, eps=1e-15))
For values of alpha = 1e-05 The log loss is: 1.4579396920648464
For values of alpha = 0.0001 The log loss is: 1.4184348029795248
For values of alpha = 0.001 The log loss is: 1.25553492519071
For values of alpha = 0.01 The log loss is: 1.2368617612994484
For values of alpha = 0.1 The log loss is: 1.2662170433876874
For values of alpha = 1 The log loss is: 1.3283015648597083
```

1.45 (1e-05, 1.458) (0.0001, 1.418)

```
130

125

(0.001, 1.256)

(0.001, 1.237)

0.0 0.2 0.4 0.6 0.8 1.0

Alpha i's
```

```
For values of best alpha = 0.01 The train log loss is: 0.6723933204672076

For values of best alpha = 0.01 The cross validation log loss is: 1.2368617612994484

For values of best alpha = 0.01 The test log loss is: 1.2049870534511515
```

# Q. Is the Text feature stable across all the data sets (Test, Train, Cross validation)?

Ans. Yes, it seems like!

```
In [0]:
```

```
def get_intersec_text(df):
    df_text_vec = CountVectorizer(min_df=3)
    df_text_fea = df_text_vec.fit_transform(df['TEXT'])
    df_text_features = df_text_vec.get_feature_names()

df_text_fea_counts = df_text_fea.sum(axis=0).A1
    df_text_fea_dict = dict(zip(list(df_text_features),df_text_fea_counts))
    len1 = len(set(df_text_features))
    len2 = len(set(train_text_features) & set(df_text_features))
    return len1,len2
```

#### In [0]:

```
len1,len2 = get_intersec_text(test_df)
print(np.round((len2/len1)*100, 3), "% of word of test data appeared in train data")
len1,len2 = get_intersec_text(cv_df)
print(np.round((len2/len1)*100, 3), "% of word of Cross Validation appeared in train data")
```

96.956 % of word of test data appeared in train data 98.83 % of word of Cross Validation appeared in train data

# 4. Machine Learning Models

#### In [0]:

```
#Data preparation for ML models.

#Misc. functionns for ML models

def predict_and_plot_confusion_matrix(train_x, train_y, test_x, test_y, clf):
    clf.fit(train_x, train_y)
    sig_clf = CalibratedClassifierCV(clf, method="sigmoid")
    sig_clf.fit(train_x, train_y)
    pred_y = sig_clf.predict(test_x)

# for calculating log_loss we will provide the array of probabilities belongs to each class
    print("Log loss:",log_loss(test_y, sig_clf.predict_proba(test_x)))
    # calculating the number of data points that are misclassified
    print("Number of mis-classified points:", np.count_nonzero((pred_y- test_y))/test_y.shape[0])
    plot_confusion_matrix(test_y, pred_y)
```

```
def report_log_loss(train_x, train_y, test_x, test_y, clf):
    clf.fit(train_x, train_y)
    sig_clf = CalibratedClassifierCV(clf, method="sigmoid")
    sig_clf.fit(train_x, train_y)
```

```
sig_clf_probs = sig_clf.predict_proba(test_x)
return log_loss(test_y, sig_clf_probs, eps=1e-15)
```

```
In [0]:
```

```
# this function will be used just for naive bayes
# for the given indices, we will print the name of the features
# and we will check whether the feature present in the test point text or not
def get_impfeature_names(indices, text, gene, var, no_features):
   gene_count_vec = CountVectorizer(ngram_range=(1,2))
   var_count_vec = CountVectorizer(ngram_range=(1,2))
   text count vec = CountVectorizer(ngram range=(1,2),min df=3)
   gene vec = gene count vec.fit(train df['Gene'])
   var_vec = var_count_vec.fit(train_df['Variation'])
   text vec = text count vec.fit(train df['TEXT'])
    fea1 len = len(gene vec.get feature names())
   fea2 len = len(var count vec.get feature names())
   word_present = 0
   for i,v in enumerate(indices):
       if (v < feal len):</pre>
           word = gene vec.get feature names()[v]
            yes no = True if word == gene else False
           if yes no:
               word present += 1
               print(i, "Gene feature [{}] present in test data point [{}]".format(word, yes no))
       elif (v < fea1_len+fea2_len):</pre>
           word = var_vec.get_feature_names()[v-(fea1_len)]
            yes_no = True if word == var else False
            if yes no:
                word present += 1
                print(i, "variation feature [{}] present in test data point [{}]".format(word,yes no))
       else:
           word = text vec.get feature names()[v-(fea1 len+fea2 len)]
           yes no = True if word in text.split() else False
           if yes no:
               word present += 1
                print(i, "Text feature [{}] present in test data point [{}]".format(word, yes no))
   print("Out of the top ", no features," features ", word present, "are present in query point")
```

# Stacking the three types of features

```
# merging gene, variance and text features
# building train, test and cross validation data sets
# a = [[1, 2],
      [3, 4]]
#b = [[4, 5],
      [6, 7]]
# hstack(a, b) = [[1, 2, 4, 5],
                 [ 3, 4, 6, 711
train gene var onehotCoding = hstack((train gene feature onehotCoding, train variation feature onehotCod
test gene var onehotCoding = hstack((test gene feature onehotCoding, test variation feature onehotCoding
cv gene var onehotCoding = hstack((cv gene feature onehotCoding,cv variation feature onehotCoding))
\label{train_x_onehotCoding} \ = \ hstack((train\_gene\_var\_onehotCoding, \ train\_text\_feature\_onehotCoding)).tocsr()
train y = np.array(list(train df['Class']))
test x onehotCoding = hstack((test gene var onehotCoding, test text feature onehotCoding)).tocsr()
test y = np.array(list(test df['Class']))
cv x onehotCoding = hstack((cv gene var onehotCoding, cv text feature onehotCoding)).tocsr()
cv y = np.array(list(cv df['Class']))
```

```
train gene var responseCoding = np.hstack((train gene feature responseCoding, train variation feature re
sponseCoding))
test gene var responseCoding = np.hstack((test gene feature responseCoding,test variation feature respo
cv gene var responseCoding = np.hstack((cv gene feature responseCoding,cv variation feature responseCod
train_x_responseCoding = np.hstack((train_gene_var_responseCoding, train_text_feature_responseCoding))
test x responseCoding = np.hstack((test gene var responseCoding, test text feature responseCoding))
cv x responseCoding = np.hstack((cv gene var responseCoding, cv text feature responseCoding))
In [0]:
print("One hot encoding features :")
print("(number of data points * number of features) in train data = ", train x onehotCoding.shape)
print("(number of data points * number of features) in test data = ", test x onehotCoding.shape)
print("(number of data points * number of features) in cross validation data =", cv x onehotCoding.shap
e)
One hot encoding features :
(number of data points * number of features) in train data = (2124, 784069)
(number of data points * number of features) in test data = (665, 784069)
(number of data points * number of features) in cross validation data = (532, 784069)
In [0]:
print(" Response encoding features :")
```

```
print("(number of data points * number of features) in train data = ", train_x_responseCoding.shape)
print("(number of data points * number of features) in test data = ", test_x_responseCoding.shape)
print("(number of data points * number of features) in cross validation data = ", cv_x_responseCoding.sh
ape)
```

```
Response encoding features:
(number of data points * number of features) in train data = (2124, 27)
(number of data points * number of features) in test data = (665, 27)
(number of data points * number of features) in cross validation data = (532, 27)
```

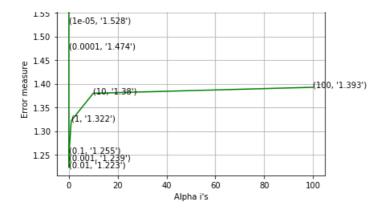
# 4.3. Logistic Regression

### 4.3.1. With Class balancing

### 4.3.1.1. Hyper paramter tuning

```
# default paramters
# sklearn.calibration.CalibratedClassifierCV(base estimator=None, method='sigmoid', cv=3)
# some of the methods of CalibratedClassifierCV()
# fit(X, y[, sample weight]) Fit the calibrated model
# get params([deep]) Get parameters for this estimator.
# predict(X) Predict the target of new samples.
# predict proba(X) Posterior probabilities of classification
# video link:
alpha = [10 ** x for x in range(-6, 3)]
cv log error array = []
for i in alpha:
    print("for alpha =", i)
    clf = SGDClassifier(class_weight='balanced', alpha=i, penalty='12', loss='log', random_state=42)
    clf.fit(train_x_onehotCoding, train_y)
    sig clf = CalibratedClassifierCV(clf, method="sigmoid")
    sig_clf.fit(train_x_onehotCoding, train_y)
    sig_clf_probs = sig_clf.predict_proba(cv_x_onehotCoding)
    cv_log_error_array.append(log_loss(cv_y, sig_clf_probs, labels=clf.classes_, eps=1e-15))
    # to avoid rounding error while multiplying probabilites we use log-probability estimates
    print("Log Loss :",log_loss(cv_y, sig_clf_probs))
fig, ax = plt.subplots()
ax.plot(alpha, cv_log_error_array,c='g')
for i, txt in enumerate(np.round(cv log error array,3)):
   ax.annotate((alpha[i],str(txt)), (alpha[i],cv log error array[i]))
plt.grid()
plt.title("Cross Validation Error for each alpha")
plt.xlabel("Alpha i's")
plt.ylabel("Error measure")
plt.show()
best_alpha = np.argmin(cv_log_error_array)
clf = SGDClassifier(class weight='balanced', alpha=alpha[best alpha], penalty='12', loss='log', random
clf.fit(train x onehotCoding, train y)
sig clf = CalibratedClassifierCV(clf, method="sigmoid")
sig clf.fit(train x onehotCoding, train y)
predict y = sig clf.predict proba(train x onehotCoding)
print('For values of best alpha = ', alpha[best alpha], "The train log loss is:", log loss (y train, pred
ict_y, labels=clf.classes_, eps=1e-15))
predict y = sig clf.predict proba(cv x onehotCoding)
print('For values of best alpha = ', alpha[best_alpha], "The cross validation log loss is:",log_loss(y_
cv, predict_y, labels=clf.classes_, eps=1e-15))
predict y = sig clf.predict proba(test x onehotCoding)
print('For values of best alpha = ', alpha[best_alpha], "The test log loss is:",log_loss(y_test, predic
t_y, labels=clf.classes_, eps=1e-15))
for alpha = 1e-06
Log Loss: 1.558141916589646
for alpha = 1e-05
Log Loss: 1.527881308282556
for alpha = 0.0001
Log Loss: 1.4740906779331717
for alpha = 0.001
Log Loss: 1.2392503761304874
for alpha = 0.01
Log Loss: 1.223295676590564
for alpha = 0.1
Log Loss: 1.254611523310852
for alpha = 1
Log Loss: 1.3220206892100543
for alpha = 10
Log Loss: 1.3799078458889427
for alpha = 100
Log Loss: 1.3925458236832124
```

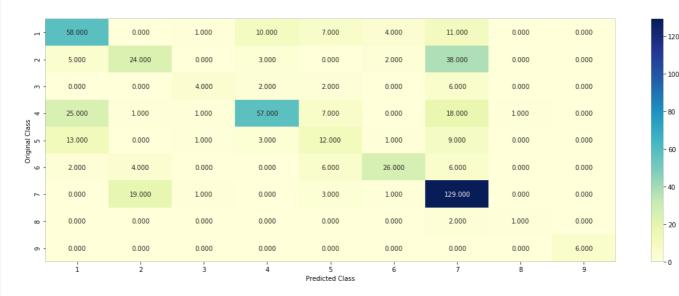
III. CAIIDIA CIOII. CAIIDIA CCACIASSIIICI CV. IIGIII



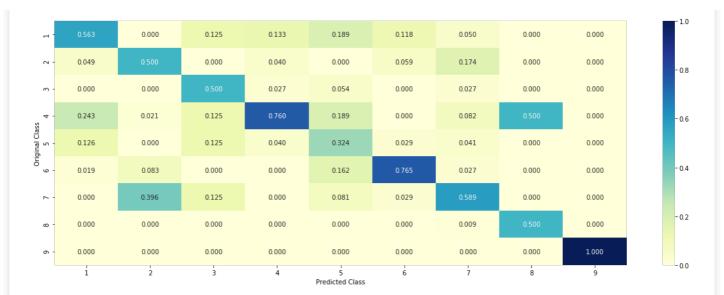
For values of best alpha = 0.01 The train log loss is: 0.6535361214259147 For values of best alpha = 0.01 The cross validation log loss is: 1.223295676590564 For values of best alpha = 0.01 The test log loss is: 1.195179296880062

#### 4.3.1.2. Testing the model with best hyper paramters

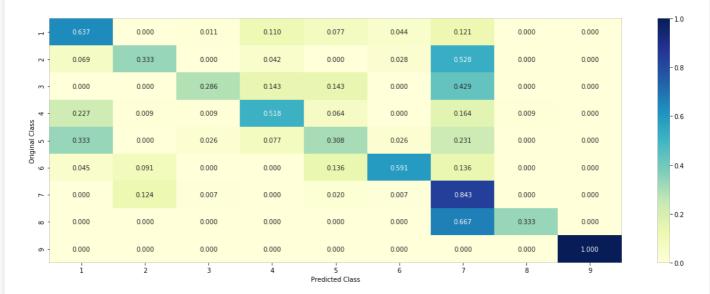
#### In [0]:



------ Precision matrix (Columm Sum=1) ------



----- Recall matrix (Row sum=1) -----



#### 4.3.1.3. Feature Importance

```
def get_imp_feature_names(text, indices, removed_ind = []):
    word present = 0
    tabulte list = []
    incresingorder_ind = 0
    for i in indices:
        if i < train gene feature onehotCoding.shape[1]:</pre>
            tabulte_list.append([incresingorder_ind, "Gene", "Yes"])
        elif i< 18:
            tabulte_list.append([incresingorder_ind,"Variation", "Yes"])
        if ((i > 17) & (i not in removed ind)):
            word = train_text_features[i]
            yes_no = True if word in text.split() else False
            if yes_no:
                word present += 1
            tabulte_list.append([incresingorder_ind,train_text_features[i], yes_no])
        incresingorder_ind += 1
    print (word present, "most importent features are present in our query point")
    print("-"*50)
    \label{lem:print} \mbox{print("The features that are most importent of the ",predicted\_cls[0]," class:")}
    print (tabulate(tabulte list, headers=["Index", 'Feature name', 'Present or Not']))
```

```
In [0]:
```

```
# from tabulate import tabulate
clf = SGDClassifier(class weight='balanced', alpha=alpha[best alpha], penalty='12', loss='log', random
clf.fit(train x onehotCoding, train y)
 test point index = 1
 no feature = 500
 predicted cls = sig clf.predict(test x onehotCoding[test point index])
 print("Predicted Class:", predicted cls[0])
print("Predicted Class Probabilities:", np.round(sig clf.predict proba(test x onehotCoding[test point i
 print("Actual Class:", test_y[test_point_index])
 indices = np.argsort(-1*abs(clf.coef))[predicted cls-1][:,:no feature]
 print("-"*50)
\texttt{get impfeature\_names(indices[0], test\_df['\texttt{TEXT'}].iloc[test\_point\_index], test\_df['\texttt{Gene'}].iloc[test\_point\_index], test\_df['\texttt{Gene'}].iloc['\texttt{Gene
  index], test df['Variation'].iloc[test point index], no feature)
Predicted Class: 6
Predicted Class Probabilities: [[0.0066 0.0069 0.0011 0.0039 0.048 0.9219 0.0035 0.0064 0.0017]]
Actual Class: 6
Out of the top 500 features 0 are present in query point
```

#### 4.3.1.3.2. Incorrectly Classified point

```
In [0]:
```

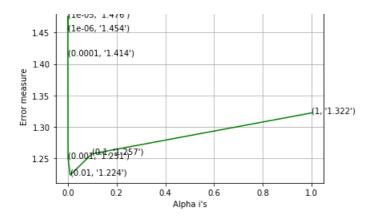
```
test point index = 100
no_feature = 500
predicted_cls = sig_clf.predict(test_x_onehotCoding[test_point_index])
print("Predicted Class :", predicted_cls[0])
print("Predicted Class Probabilities:", np.round(sig_clf.predict_proba(test_x_onehotCoding[test_point_i
ndex1),4))
print("Actual Class :", test_y[test_point_index])
indices = np.argsort(-1*abs(clf.coef ))[predicted cls-1][:,:no feature]
print("-"*50)
get impfeature names(indices[0], test df['TEXT'].iloc[test point index], test df['Gene'].iloc[test point
index], test df['Variation'].iloc[test point index], no feature)
Predicted Class: 2
Predicted Class Probabilities: [[0.0717 0.4325 0.0183 0.0574 0.0321 0.0267 0.3472 0.0071 0.007 ]]
Actual Class: 7
382 Text feature [cys1156] present in test data point [True]
Out of the top 500 features 1 are present in query point
```

## 4.3.2. Without Class balancing

#### 4.3.2.1. Hyper paramter tuning

```
# find more about CalibratedClassifierCV here at http://scikit-learn.org/stable/modules/generated/sklea
rn.calibration.CalibratedClassifierCV.html
# default paramters
# sklearn.calibration.CalibratedClassifierCV(base estimator=None, method='sigmoid', cv=3)
# some of the methods of CalibratedClassifierCV()
\# fit(X, y[, sample weight]) Fit the calibrated model
# get_params([deep]) Get parameters for this estimator.
# predict(X) Predict the target of new samples.
# predict proba(X) Posterior probabilities of classification
# video link:
alpha = [10 ** x for x in range(-6, 1)]
cv log error array = []
for i in alpha:
   print("for alpha =", i)
    clf = SGDClassifier(alpha=i, penalty='12', loss='log', random state=42)
   clf.fit(train_x_onehotCoding, train_y)
    sig clf = CalibratedClassifierCV(clf, method="sigmoid")
    sig clf.fit(train x onehotCoding, train y)
    sig_clf_probs = sig_clf.predict_proba(cv_x_onehotCoding)
    cv_log_error_array.append(log_loss(cv_y, sig_clf_probs, labels=clf.classes_, eps=1e-15))
    print("Log Loss :",log loss(cv y, sig clf probs))
fig, ax = plt.subplots()
ax.plot(alpha, cv log error array, c='g')
for i, txt in enumerate(np.round(cv log error array,3)):
   ax.annotate((alpha[i],str(txt)), (alpha[i],cv log error array[i]))
plt.grid()
plt.title("Cross Validation Error for each alpha")
plt.xlabel("Alpha i's")
plt.ylabel("Error measure")
plt.show()
best_alpha = np.argmin(cv_log_error_array)
clf = SGDClassifier(alpha=alpha[best alpha], penalty='12', loss='log', random state=42)
clf.fit(train_x_onehotCoding, train_y)
sig clf = CalibratedClassifierCV(clf, method="sigmoid")
sig clf.fit(train x onehotCoding, train y)
predict y = sig clf.predict proba(train x onehotCoding)
print('For values of best alpha = ', alpha[best_alpha], "The train log loss is:",log_loss(y train, pred
ict_y, labels=clf.classes_, eps=1e-15))
predict y = sig clf.predict proba(cv x onehotCoding)
print('For values of best alpha = ', alpha[best alpha], "The cross validation log loss is:",log loss(y
cv, predict_y, labels=clf.classes_, eps=1e-15))
predict y = sig clf.predict proba(test x onehotCoding)
print('For values of best alpha = ', alpha[best_alpha], "The test log loss is:",log_loss(y_test, predic
t_y, labels=clf.classes_, eps=1e-15))
for alpha = 1e-06
Log Loss: 1.4538556392957696
for alpha = 1e-05
Log Loss: 1.475576127170645
for alpha = 0.0001
Log Loss: 1.413765961776697
for alpha = 0.001
Log Loss : 1.2509724087932241
for alpha = 0.01
Log Loss: 1.2237825948063414
for alpha = 0.1
Log Loss: 1.2574156264144203
for alpha = 1
Log Loss: 1.3223089085194708
```

on-1/



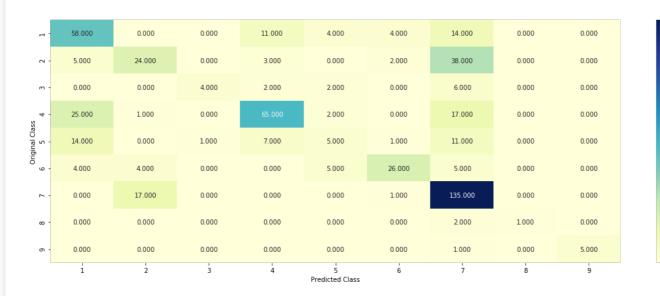
For values of best alpha = 0.01 The train log loss is: 0.6423459018626693

For values of best alpha = 0.01 The cross validation log loss is: 1.2237825948063414

For values of best alpha = 0.01 The test log loss is: 1.1925278039958513

# 4.3.2.2. Testing model with best hyper parameters

#### In [0]:



- 120

- 100

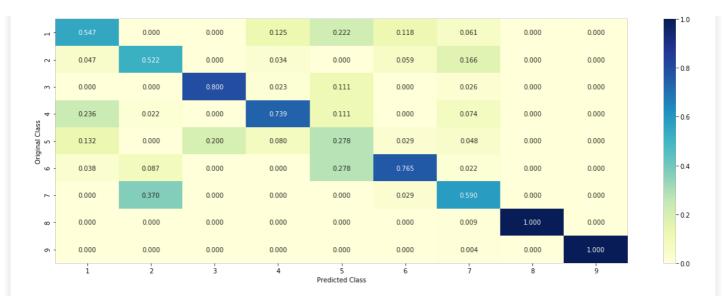
- 80

60

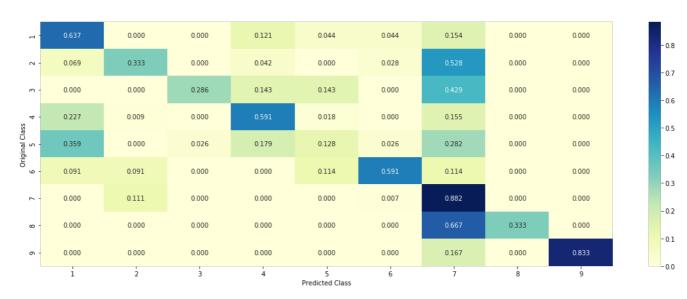
- 40

- 20

----- Precision matrix (Columm Sum=1)



----- Recall matrix (Row sum=1) -----



#### 4.3.2.3. Feature Importance, Correctly Classified point

```
In [0]:
clf = SGDClassifier(alpha=alpha[best alpha], penalty='12', loss='log', random state=42)
clf.fit(train x onehotCoding,train y)
test point index = 1
no feature = 500
predicted cls = sig clf.predict(test x onehotCoding[test point index])
print("Predicted Class :", predicted cls[0])
print("Predicted Class Probabilities:", np.round(sig_clf.predict_proba(test_x_onehotCoding[test_point_i
ndex]),4))
print("Actual Class:", test_y[test_point_index])
indices = np.argsort(-1*abs(clf.coef_))[predicted_cls-1][:,:no_feature]
print("-"*50)
get_impfeature_names(indices[0], test_df['TEXT'].iloc[test_point_index],test_df['Gene'].iloc[test_point
_index], test_df['Variation'].iloc[test_point_index], no_feature)
Predicted Class: 6
Predicted Class Probabilities: [[9.30e-03 7.70e-03 1.50e-03 5.90e-03 3.54e-02 9.32e-01 3.90e-03 4.00e-0
  2.00e-04]]
Actual Class : 6
```

#### 4.3.2.4. Feature Importance, Inorrectly Classified point

Out of the top 500 features 0 are present in query point

• • •

```
In [0]:
test point index = 100
no feature = 500
predicted cls = sig clf.predict(test x onehotCoding[test point index])
print("Predicted Class :", predicted_cls[0])
print("Predicted Class Probabilities:", np.round(sig clf.predict proba(test x onehotCoding[test point i
ndex]),4))
print("Actual Class :", test_y[test_point_index])
indices = np.argsort(-1*abs(clf.coef))[predicted cls-1][:,:no feature]
print("-"*50)
_index],test_df['Variation'].iloc[test_point_index], no_feature)
Predicted Class: 2
Predicted Class Probabilities: [[0.0725 0.4559 0.0213 0.0553 0.035 0.0288 0.3191 0.0072 0.0049]]
Actual Class: 7
Out of the top 500 features 0 are present in query point
```