Personalized cancer diagnosis

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

- 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
- 2. https://www.youtube.com/watch?v=UwbuW7oK8rk
- 3. https://www.youtube.com/watch?v=qxXRKVompl8

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.

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 [1]:
```

```
import pandas as pd
import matplotlib.pyplot as plt
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.cross validation 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
import math
# 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
from tqdm import tqdm notebook
```

3.1. Reading Data

3.1.1. Reading Gene and Variation Data

```
In [2]:
```

```
data = pd.read_csv('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[2]:

	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 [3]:

```
# note the seprator in this file
data_text =pd.read_csv("training_text",sep="\|\\",engine="python",names=["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[3]:

	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 [4]:

```
# 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\n]', ' ', 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 [5]:

```
#text processing stage.
start_time = time.clock()
for index, row in tqdm_notebook(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 : 154.3329649 seconds

In [6]:

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

Out[6]:

	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

In [7]:

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

Out[7]:

	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 [8]:

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

In [9]:

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

Out[9]:

	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 [10]:

```
y_true = result['Class'].values
result.Gene = result.Gene.str.replace('\s+', '_')
result.Variation = result.Variation.str.replace('\s+', '_')

# split the data into test and train by maintaining same distribution of output varaible 'y_true' [stratify=y_true]
X_train, test_df, y_train, y_test = train_test_split(result, y_true, stratify=y_true, test_size=0.2, random_state=1)

# split the train data into train and cross validation by maintaining same distribution of output varaible '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, random_state=1)
```

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

In [11]:

```
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 cross validation data: 532

Number of data points in test data: 665

3.1.4.2. Distribution of y_i's in Train, Test and Cross Validation datasets

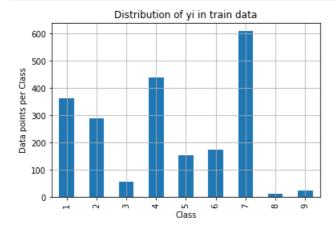
In [12]:

```
# 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()

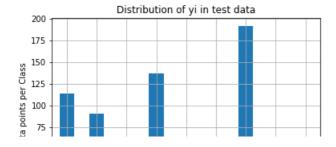
my_colors = 'rgbkymc'
train_class_distribution.plot(kind='bar')
plt.xlabel('Class')
plt.ylabel('Distribution of yi in train data')
plt.grid()
plt.grid()
plt.grid()
plt.show()

# 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:
```

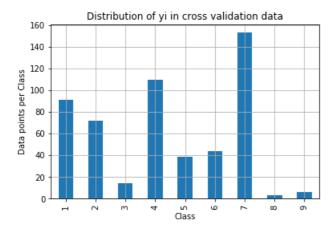
```
print('Number of data points in class', i+1, ':', train class distribution.values[i], '(', np.round(
(train_class_distribution.values[i]/train_df.shape[0]*100), 3), '%)')
print('-'*80)
my colors = 'rgbkymc'
test class distribution.plot(kind='bar')
plt.xlabel('Class')
plt.ylabel('Data points per Class')
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')
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 7 : 609 ( 28.672 \%) 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 \% ( 0.565 \%)
```



```
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.

In [13]:

```
# 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 array
    # C.sum(axix = 1) = [[3, 7]]
    # ((C.T)/(C.sum(axis=1))) = [[1/3, 3/7]
    # [2/3, 4/7]]

# ((C.T)/(C.sum(axis=1))).T = [[1/3, 2/3]
```

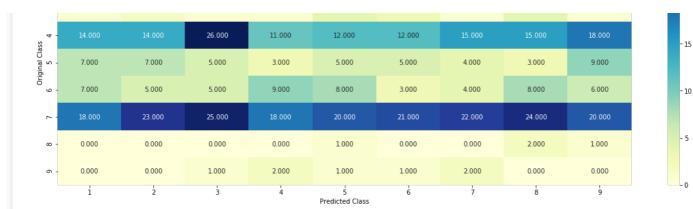
```
[3/7, 4/7]]
    # sum of row elements = 1
   B = (C/C.sum(axis=0))
   #divid each element of the confusion matrix with the sum of elements in that row
    \# C = [[1, 2],
         [3, 4]]
   # C.sum(axis = 0) axis=0 corresonds to columns and axis=1 corresponds to rows in two diamensional
arrav
    \# 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()
```

In [14]:

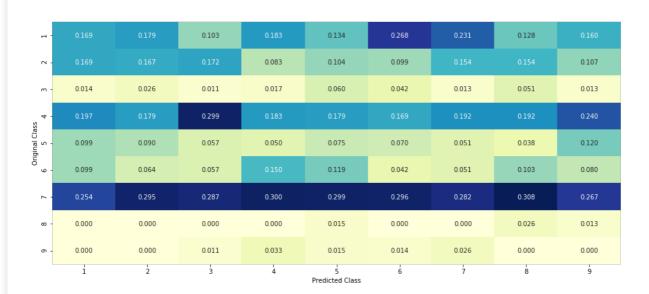
```
# we need to generate 9 numbers and the sum of numbers should be 1
# one solution is to genarate 9 numbers and divide each of the numbers by their sum
# ref: https://stackoverflow.com/a/18662466/4084039
test data len = test df.shape[0]
cv data len = cv df.shape[0]
# we create a output array that 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=1e-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)
```

Log loss on Cross Validation Data using Random Model 2.5104111490430268 Log loss on Test Data using Random Model 2.5158449593068393 ------ Confusion matrix ------

н -	12.000	14.000	9.000	11.000	9.000	19.000	18.000	10.000	12.000
- 5	12.000		15.000	5.000	7.000	7.000	12.000	12.000	8.000
m -	1.000	2.000	1.000	1.000	4.000	3.000	1.000	4.000	1.000



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



- 0.24

-0.18

- 0.06

- 0.00

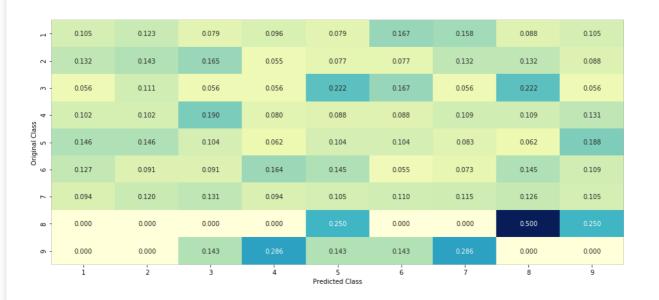
- 0.4

- 0.3

- 0.2

- 0.1

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



3.3 Univariate Analysis

```
In [15]:
```

```
# 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, 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:
                      174
            {BRCA1
             TP53
                       106
             EGFR
             BRCA2
                        75
             PTEN
                         69
    #
             KIT
                         61
             BRAF
                         60
             ERBB2
             PDGFRA
                        46
             . . . }
   # print(train df['Variation'].value counts())
   # output:
   # {
    # Truncating Mutations
                                              63
   # Deletion
                                              43
   # Amplification
                                              43
   # Fusions
                                              22
                                               .3
   # Overexpression
   # E17K
    # 061L
   # S222D
    # 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 = n  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
# 2614 2614 BRCA1
                                             S1841R
                                                M1R
           # 2432 2432 BRCA1
                                             L1657P
           # 2567 2567 BRCA1
                                             T1685A
                                            E1660G
           # 2583 2583 BRCA1
           # 2634 2634 BRCA1
                                             W1718T
            # 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(av dict)
```

```
{'BRCA1': [0.200757575757575, 0.037878787878788, 0.068181818181818177, 0.1363636363636363
5, 0.25, 0.193181818181818181, 0.037878787878787878, 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, 0.2159090909090901, 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, 0.1393939393934, 0.34545454545454546, 0.0606060606060608, 0.06060606060608, 0.06060606060
60606081.
           '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
2913912],
           'BRAF': [0.06666666666666666, 0.17999999999999, 0.0733333333333334, 0.07333333333333
34, 0.09333333333333338, 0.08000000000000000, 0.2999999999999, 0.06666666666666666, 0.0666666666
66666666],
   #
   #
   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 [16]:
```

```
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: 235
BRCA1
         168
TP53
         111
          86
EGFR
BRCA2
          80
BRAF
          65
          60
KTT
ALK
          44
          39
ERBB2
PDGFRA
          38
Name: Gene, dtype: int64
```

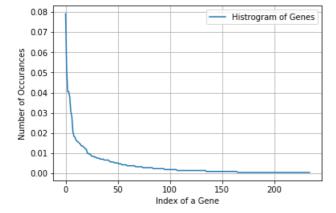
In [17]:

```
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 235 different categories of genes in the train data, and they are distibuted as follows

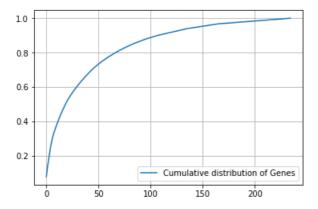
In [18]:

```
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()
```



In [19]:

```
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.

In [20]:

```
# 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 [21]:

```
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 [22]:

```
# 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 [23]:

```
gene_vectorizer.get_feature_names()
```

Out[23]:

```
['abl1',
 'acvr1'
'ago2',
'akt1',
'akt2',
'akt3',
 'alk',
 'apc',
'ar',
'araf',
'aridla',
 'arid1b',
 'arid2',
 'arid5b'.
'asxl1',
'asxl2',
 'atm',
 'atr',
 'atrx',
'aurka',
'axin1',
'axl',
 'b2m',
 'bap1',
 'bard1',
'bcl10',
'bcl2111',
 'bcor',
 'braf',
 'brcal',
'brca2',
'brd4',
'brip1',
 'btk',
 'card11',
```

'carm1'.

```
'casp8',
'cbl',
'ccnd1',
'ccnd2',
'ccnd3',
'ccne1',
'cdh1',
'cdk12',
'cdk4',
'cdk6',
'cdknla',
'cdkn1b',
'cdkn2a',
'cdkn2b',
'cdkn2c',
'cebpa',
'chek2',
'cic',
'crebbp',
'ctcf',
'ctla4',
'ctnnb1',
'ddr2',
'dicer1',
'dnmt3a',
'dnmt3b',
'egfr',
'elf3',
'ep300',
'epas1',
'epcam',
'erbb2',
'erbb3',
'erbb4',
'ercc2',
'ercc3',
'ercc4',
'erg',
'errfil',
'esr1',
'etv1',
'etv6',
'ewsr1',
'ezh2',
'fanca',
'fat1',
'fbxw7',
'fgf4',
'fgfr1',
'fgfr2',
'fgfr3',
'fgfr4',
'flt1',
'flt3',
'foxal',
'foxl2',
'foxp1',
'fubp1',
'gata3',
'glil',
'gnaq',
'gnas',
'h3f3a',
'hist1h1c',
'hla',
'hras',
'idh1',
'idh2',
'igf1r',
'ikbke',
'ikzf1',
'jak1',
'jak2',
'jun',
'kdm5c',
'kdm6a',
'kdr'
```

```
nur,
'keap1',
'kit',
'kmt2a',
'kmt2b',
'kmt2c',
'kmt2d',
'knstrn',
'kras',
'lats2',
'map2k1',
'map2k2',
'map2k4',
'map3k1',
'mapk1',
'mdm2',
'mdm4',
'med12',
'mef2b',
'men1',
'met',
'mlh1',
'mpl',
'msh2',
'msh6',
'mtor',
'myc',
'mycn',
'myd88',
'ncor1',
'nf1',
'nf2',
'nfe212',
'nfkbia',
'nkx2',
'notch1',
'notch2',
'nras',
'ntrk1',
'ntrk2',
'ntrk3',
'nup93',
'pak1',
'pax8',
'pbrm1',
'pdgfra',
'pdgfrb',
'pik3ca',
'pik3cb',
'pik3cd',
'pik3r1',
'pik3r2',
'pik3r3',
'pim1',
'pms1',
'pms2',
'pole',
'ppmld',
'ppp2r1a',
'ppp6c',
'prdm1',
'ptch1',
'pten',
'ptpn11',
'ptprd',
'ptprt',
'rab35',
'rac1',
'rad21',
'rad50',
'rad51b',
'rad51d',
'raf1',
'rara',
'rasa1',
'rb1',
'rbm10',
1 20+ 1
```

```
TEC ,
'rheb',
'rhoa',
'rictor',
'rit1',
'rnf43',
'ros1',
'rras2',
'runx1',
'rybp',
'sdhb',
'sdhc',
'setd2',
'sf3b1',
'shoc2',
'smad2',
'smad3',
'smad4',
'smarca4'
'smarcb1'
'smo',
'sox9'
'spop',
'src',
'stag2',
'stat3',
'stk11',
'tcf712',
'tert',
'tet1',
'tet2',
'tgfbr1',
'tgfbr2',
'tmprss2',
'tp53',
'tp53bp1',
'tsc1',
'tsc2',
'u2af1',
'vegfa',
'vhl',
'whsc1'
'whsc1l1'
'xpol',
'xrcc2'
'yap1']
```

In [24]:

```
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, 235)

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

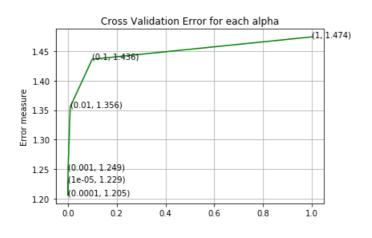
There are many ways to estimate how good a feature is, in predicting y_i. One of the good methods is to build a proper ML model using just this feature. In this case, we will build a logistic regression model using only Gene feature (one hot encoded) to predict y_i.

In [25]:

```
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.arid()
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: 1.2285570397154246
For values of alpha = 0.0001 The log loss is: 1.205397678433504
For values of alpha = 0.001 The log loss is: 1.2491650009150361
For values of alpha = 0.01 The log loss is: 1.3556362010078462
For values of alpha = 0.1 The log loss is: 1.4361429435939435
For values of alpha = 1 The log loss is: 1.4740301432719325
```



```
For values of best alpha = 0.0001 The train log loss is: 1.004691587658098 For values of best alpha = 0.0001 The cross validation log loss is: 1.205397678433504 For values of best alpha = 0.0001 The test log loss is: 1.1962093819056512
```

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 [26]:

```
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 $\,$ 235 $\,$ genes in train dataset? Ans

- 1. In test data 644 out of 665 : 96.84210526315789
- 2. In cross validation data 511 out of 532: 96.05263157894737

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?

In [27]:

```
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: 1920
Truncating Mutations 58
Amplification
                       51
Deletion
                       45
Fusions
                       2.5
G12V
Overexpression
                        3
Y64A
C618R
Q61R
                        2
061T
Name: Variation, dtype: int64
```

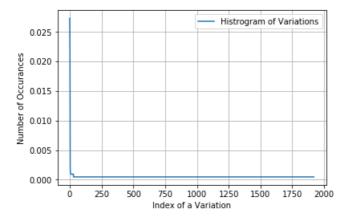
In [28]:

```
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 1920 different categories of variations in the train data, and they are distibuted as fo llows

In [29]:

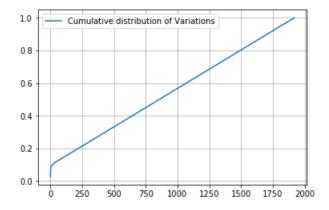
```
s = sum(unique_variations.values);
h = unique_variations.values/s;
plt.plot(h, label="Histrogram of Variations")
plt.xlabel('Index of a Variation')
plt.ylabel('Number of Occurances')
plt.legend()
plt.grid()
plt.show()
```



In [30]:

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

[0.02730697 0.05131827 0.07250471 ... 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

We will be using both these methods to featurize the Variation Feature

In [31]:

```
# Response coding on 'Variation'
# alpha is used for laplace smoothing
alpha = 1
# train cone 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 [32]:

```
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 [33]:

```
# 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 [34]:

```
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, 1948)

Q10. How good is this Variation feature in predicting y_i?

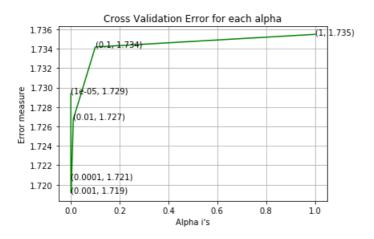
Let's build a model just like the earlier!

In [35]:

```
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: 1.7293899332490221 For values of alpha = 0.0001 The log loss is: 1.7205800598416228 For values of alpha = 0.001 The log loss is: 1.719154795433691 For values of alpha = 0.01 The log loss is: 1.7267443626734562 For values of alpha = 0.1 The log loss is: 1.7341851235193924 For values of alpha = 1 The log loss is: 1.7354863846993358
```



```
For values of best alpha = 0.001 The train log loss is: 1.0984554907261728

For values of best alpha = 0.001 The cross validation log loss is: 1.719154795433691

For values of best alpha = 0.001 The test log loss is: 1.7066671376163376
```

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 [36]:

```
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 1920 genes in test and cross validation data sets? Ans
```

- 1. In test data 61 out of 665 : 9.172932330827068
- 2. In cross validation data 58 out of 532 : 10.902255639097744

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 [37]:

In [38]:

In [39]:

```
# building a CountVectorizer with all the words that occured minimum 3 times in train data
text_vectorizer = CountVectorizer(min_df=3, ngram_range=(1,2))
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: 780378

```
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 = train df.groupby('Class').count().to dict().get('ID')
confuse array = []
for i in train text features:
   ratios = []
   \max val = -1
   for j in range (0,9):
       ratios.append((dict_list[j][i]+10 )/(total_dict[j+1]+90))
   confuse array.append(ratios)
confuse_array = np.array(confuse_array)
```

In [41]:

```
#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 [42]:

```
# 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 [43]:

```
# 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 [44]:

```
#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()))
```

In [45]:

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

Counter({3: 150307, 4: 102413, 5: 64028, 6: 62907, 7: 48553, 8: 44824, 9: 38974, 12: 24979, 10: 21951, 11: 16250, 13: 13621, 14: 13301, 16: 13030, 15: 11945, 18: 9064, 17: 9024, 20: 6431, 19: 6367, 28: 5094 . 21: 5036. 22: 4997. 24: 4658. 30: 4317. 26: 4207. 23: 3710. 41: 3372. 25: 3300. 27: 3093. 42: 2894. 2

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In [46]:

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# 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
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 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.arid()
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=le-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=le-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, predict_y, labels=clf.classes_, eps=le-15))
```

```
For values of alpha = 1e-05 The log loss is: 1.5167231920253488

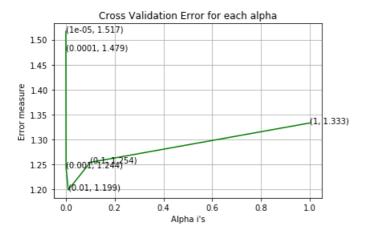
For values of alpha = 0.0001 The log loss is: 1.4794519724011637

For values of alpha = 0.001 The log loss is: 1.2444734717983619

For values of alpha = 0.01 The log loss is: 1.1988087536395524

For values of alpha = 0.1 The log loss is: 1.2541335677728367

For values of alpha = 1 The log loss is: 1.3331017378409953
```



```
For values of best alpha = 0.01 The train log loss is: 0.7301697590831281

For values of best alpha = 0.01 The cross validation log loss is: 1.1988087536395524

For values of best alpha = 0.01 The test log loss is: 1.1950351201142333
```

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

Ans. Yes, it seems like!

```
In [59]:
```

```
def get_intersec_text(df):
    df_text_vec = CountVectorizer(min_df=3, ngram_range=(1,2))
    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 [60]:

```
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")
```

```
92.847~\% of word of test data appeared in train data 95.593~\% of word of Cross Validation appeared in train data
```

4. Machine Learning Models

In [50]:

```
#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)
```

In [51]:

```
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 [78]:

```
# generate ngram from sentence
# Ref: http://www.albertauyeung.com/post/generating-ngrams-python/
def generate_ngrams(s, n):
    # Convert to lowercases
    s = s.lower()

# Replace all none alphanumeric characters with spaces
s = re.sub(r'[^a-zA-Z0-9\s]', ' ', s)

# Break sentence in the token, remove empty tokens
tokens = [token for token in s.split(" ") if token != ""]

# Use the zip function to help us generate n-grams
# Concatentate the tokens into ngrams and return
ngrams = zip(*[tokens[i:] for i in range(n)])
return [" ".join(ngram) for ngram in ngrams]
```

In [95]:

```
# 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()
   var count vec = CountVectorizer()
   text_count_vec = CountVectorizer(min_df=3, ngram_range=(1,2))
   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'])
   all features = gene count vec.get feature names() + var count vec.get feature names() + text count
vec.get feature names()
   feal 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 = all features[v]
           yes no = True if word == gene else False
            if was no
```

```
TT ACD 110.
            word present += 1
            print(i, "Gene feature [{}] present in test data point [{}]".format(word,yes_no))
    elif (v < feal len+fea2 len):</pre>
       word = all features[v]
        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 = all features[v]
         # list of text contain unigram
        text 1 = generate ngrams(text,1)
        # list of text contain bigram
        text 2 = generate ngrams (text, 2)
        yes no 1 = True if word in text 1 else False
        yes no 2 = True if word in text 2 else False
        if yes no 1:
            word present += 1
            print(i, "Text feature [{}] present in test data point [{}]".format(word, yes no 1))
        if yes no 2:
            word present += 1
            print(i, "Text feature [{}] present in test data point [{}]".format(word, yes no 2))
print("Out of the top ", no features," features ", word present, "are present in query point")
```

In [93]:

```
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)

IOPub data rate exceeded.
The notebook server will temporarily stop sending output to the client in order to avoid crashing it.
To change this limit, set the config variable
`--NotebookApp.iopub_data_rate_limit`.

Current values:
NotebookApp.iopub_data_rate_limit=1000000.0 (bytes/sec)
NotebookApp.rate_limit_window=3.0 (secs)

147 Text feature [greater ability] present in test data point [True]
247 Text feature [likely sufficient] present in test data point [True]
971 Text feature [inhibitors determine] present in test data point [True]
```

Stacking the three types of features

Out of the top 1000 features 4 are present in query point

In [53]:

```
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
nseCoding))
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 [54]:
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, 782561)
(number of data points * number of features) in test data = (665, 782561)
(number of data points * number of features) in cross validation data = (532, 782561)
In [55]:
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. Logistic Regression

4.1 With Class balancing

4.1.1. Hyper paramter tuning

In [56]:

```
# 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,
# 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: https://www.appliedaicourse.com/course/applied-ai-course-online/lessons/geometric-intuiti
on-1/
# 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, 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.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: 1.5394824675008616
for alpha = 1e-05
Log Loss: 1.5139569947164189
for alpha = 0.0001
Log Loss: 1.4616013682227202
for alpha = 0.001
Log Loss: 1.2141890807075546
for alpha = 0.01
Log Loss : 1.1792090537315691
for alpha = 0.1
```

Log Loss: 1.243222167047315

for alpha = 1

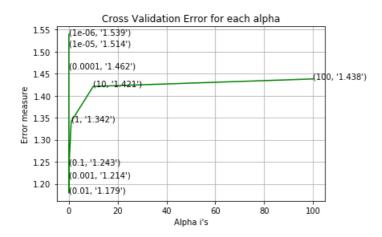
Log Loss: 1.3419419370775192

for alpha = 10

Log Loss: 1.4211980261907622

for alpha = 100

Log Loss: 1.438172305924132



For values of best alpha = 0.01 The train log loss is: 0.7070545649046233

For values of best alpha = 0.01 The cross validation log loss is: 1.1792090537315691 For values of best alpha = 0.01 The test log loss is: 1.1776257465537177

4.1.2. Testing the model with best hyper paramters

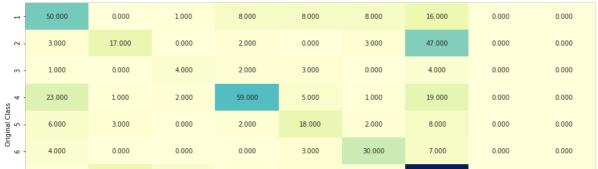
In [57]:

```
# 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: https://www.appliedaicourse.com/course/applied-ai-course-online/lessons/geometric-intuiti
clf = SGDClassifier(class weight='balanced', alpha=alpha[best alpha], penalty='12', loss='log', random
predict and plot confusion matrix(train x onehotCoding, train y, cv x onehotCoding, cv y, clf)
```

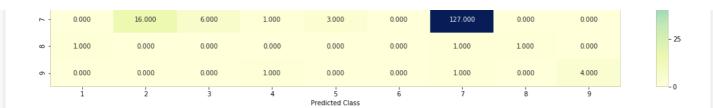
Log loss: 1.1792090537315691

Number of mis-classified points : 0.41729323308270677 ----- Confusion matrix -----

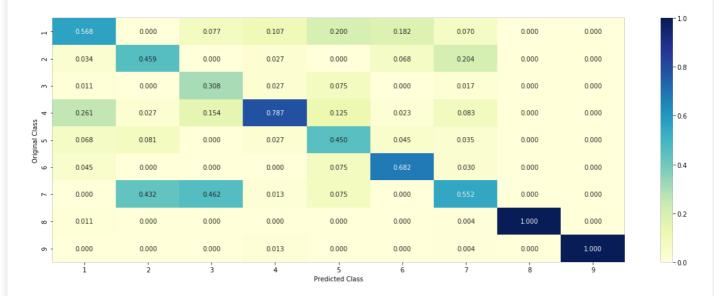
50.000 0.000 1.000



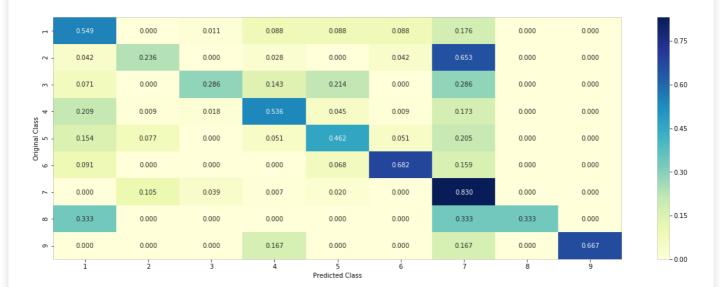
- 100 75



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



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



4.1.3. Feature Importance

4.1.3.1. Correctly Classified point

In [98]:

```
# from tabulate import tabulate
clf = SGDClassifier(class_weight='balanced', alpha=alpha[best_alpha], penalty='12', loss='log', random_
state=42)
clf.fit(train_x_onehotCoding,train_y)
test_point_index = 1
no_feature = 1500
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_index])
```

4.1.3.2. Incorrectly Classified point

```
In [97]:
```

```
test point index = 2
no feature = 1000
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(-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: 7
Predicted Class Probabilities: [[0.1203 0.1052 0.0144 0.1138 0.0458 0.0308 0.5566 0.0063 0.0068]]
Actual Class: 6
147 Text feature [greater ability] present in test data point [True]
247 Text feature [likely sufficient] present in test data point [True]
506 Text feature [thus appear] present in test data point [True]
971 Text feature [inhibitors determine] present in test data point [True]
Out of the top 1000 features 4 are present in query point
```

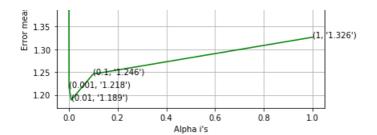
4.2. Without Class balancing

4.2.1. Hyper paramter tuning

In [99]:

```
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: 1.536591020907485
for alpha = 1e-05
Log Loss: 1.5274646873956272
for alpha = 0.0001
Log Loss: 1.4838997135067387
for alpha = 0.001
Log Loss: 1.2175398616768762
for alpha = 0.01
Log Loss: 1.1888815510143445
for alpha = 0.1
Log Loss: 1.2455676860536395
for alpha = 1
Log Loss: 1.3264610661900667
             Cross Validation Error for each alpha
```

	Cross Validation Error for each alpha						
1.55	(18:89; 11:537)						
1.50 -	(0.0001, '1.484')						
1.45 -							
일 1.40 -							



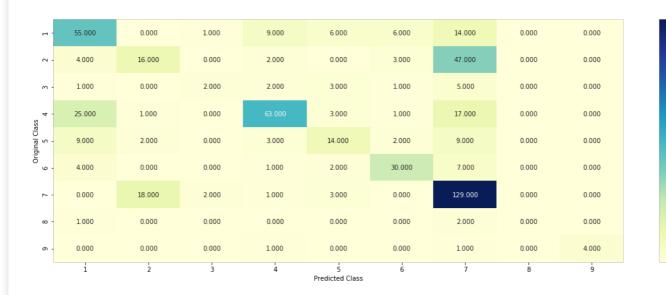
For values of best alpha = 0.01 The train log loss is: 0.7046715086493371

For values of best alpha = 0.01 The cross validation log loss is: 1.1888815510143445

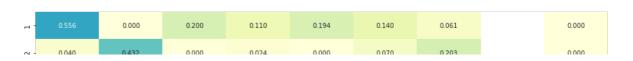
For values of best alpha = 0.01 The test log loss is: 1.1815596595987967

4.3.2.2. Testing model with best hyper parameters

In [100]:



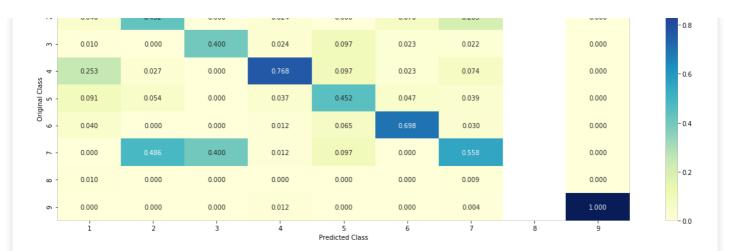
------ Precision matrix (Columm Sum=1) ------



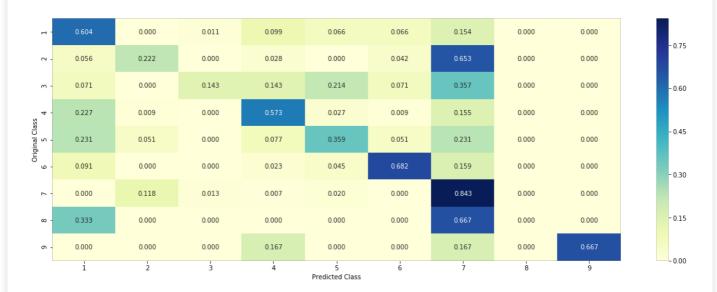
75

50

- 25



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



4.3.2.3. Feature Importance, Correctly Classified point

```
In [103]:
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 = 2000
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(-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: [[1.310e-02 1.130e-02 1.100e-03 1.020e-02 5.200e-02 9.008e-01 7.900e-03
```

1169 Text feature [african] present in test data point [True]
1367 Text feature [given family] present in test data point [True]
1790 Text feature [20c] present in test data point [True]
Out of the top 2000 features 3 are present in query point

2.900e-03 6.000e-04]]

Actual Class: 6

In [104]: test_point_index = 2 no_feature = 2000 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_index]),4)) print("Actual Class :", test y[test_point_index])

```
indices = np.argsort(-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: 7
Predicted Class Probabilities: [[0.1149 0.103 0.0173 0.1032 0.047 0.0303 0.5727 0.0053 0.0064]]
Actual Class: 6
344 Text feature [likely sufficient] present in test data point [True]
449 Text feature [greater ability] present in test data point [True]
499 Text feature [transforming] present in test data point [True]
670 Text feature [downstream signaling] present in test data point [True]
771 Text feature [p85 heterodimer] present in test data point [True]
781 Text feature [within helical] present in test data point [True]
1011 Text feature [levels phosphorylated] present in test data point [True]
1012 Text feature [5a left] present in test data point [True]
1071 Text feature [effects downstream] present in test data point [True]
1464 Text feature [genetic aberrations] present in test data point [True]
1510 Text feature [thus appear] present in test data point [True]
1570 Text feature [activation event] present in test data point [True]
1632 Text feature [cancer lineage] present in test data point [True]
1645 Text feature [inhibitors determine] present in test data point [True]
1862 Text feature [present 30] present in test data point [True]
1879 Text feature [determine effects] present in test data point [True]
1977 Text feature [studies 26] present in test data point [True]
Out of the top 2000 features 17 are present in query point
```

Conclusion

- 1. EDA
 - · Read training data 'Gene and Variation'

- · Read training data 'Text'
- Preprocessing 'Text' data
- Handling missing 'Text' data and merge all features
- 2. Split data into 3 parts in ratio 64:20:16
- 3. See the distribution of each class in train, cv and test data
- 4. Train the Random Model (worst/dump) to find the upperbound logloss
- 5. Univariant analysis on Gene and Variant
 - What type of feature is?
 - How many categories are present
 - How they are distributed
 - · How to featurize the feature: responsecoding with Laplace smoothing and onehotencoding (CountVectorizer)
 - How good is this feature in prediction y?
- 6. Univariant Analysis on Text data
 - Extract the number of words from each class
 - Featurized the text with onhotencoding(CountVectorizer) witn unigram and bigram and responsecoding with Laplace smoothing
 - How good is this feature in predicting y?
- 7. Stacked all features
- 8. Finally, train model and plot and observe confusion matrix, precision and recall
 - · Logistic Regr (LR) with class balance and get feature importance
 - Logistic Regr (LR) without class balance and get feature importance

In [3]:

```
x = PrettyTable()
x.field names = ["Feature", "Hyperparameter ", "Model", "Train logloss", "CV logloss", "Test logloss", "
# of Misclassified pts"]
print('OHE -> OneHotEncoding\nRC-> ResponseCoding with Laplace smoothing')
feature = ['OHE','OHE']
param_ = [0.01,0.01]
model_ = ['LR (class balance)','LR (no class balance)']
train_loss = [0.70, 0.70]
cv_loss = [1.18, 1.19]
test loss = [1.18, 1.18]
mis_{class} = [0.47, 0.412]
for i in range (0,2):
x.add row([feature[i],param [i],model [i],train loss[i],cv loss[i],test loss[i],mis class[i]])
print(x)
print('\n1. From the observation above, LR(no class balance) are perform better than others.')
OHE -> OneHotEncoding
RC-> ResponseCoding with Laplace smoothing
| Feature | Hyperparameter | Model | Train logloss | CV logloss | Test logloss | # of
Misclassified pts |
+----
                     | OHE | 0.01 | LR (class balance) | 0.7 | 1.18 | 1.18 | 0.47 |
OHE | 0.01
                      | LR (no class balance) | 0.7
                                                         | 1.19 | 1.18
                                                                                  1. From the observation above, LR(no class balance) are perform better than others.
In [ ]:
```