

Project Report

December 7, 2018

1 Preparation

We first load the data into memory and split data into training data and validation data as following:

```
In [2]: import numpy as np
        np.random.seed(0)
        import mltools as ml
        import matplotlib.pyplot as plt
        import pandas as pd
        from sklearn.model_selection import train_test_split
        CSV_FILE_PATH = 'data/diabetic_data.csv'
        data = pd.read_csv(CSV_FILE_PATH)

        print(data.shape)
        print(data.info())
```

```
(101766, 50)
<class 'pandas.core.frame.DataFrame'>
RangeIndex: 101766 entries, 0 to 101765
Data columns (total 50 columns):
encounter_id      101766 non-null int64
patient_nbr       101766 non-null int64
race              101766 non-null object
gender            101766 non-null object
age              101766 non-null object
weight           101766 non-null object
admission_type_id 101766 non-null int64
discharge_disposition_id 101766 non-null int64
admission_source_id 101766 non-null int64
time_in_hospital  101766 non-null int64
payer_code        101766 non-null object
medical_specialty 101766 non-null object
num_lab_procedures 101766 non-null int64
num_procedures    101766 non-null int64
num_medications   101766 non-null int64
number_outpatient 101766 non-null int64
```

```

number_emergency          101766 non-null int64
number_inpatient          101766 non-null int64
diag_1                    101766 non-null object
diag_2                    101766 non-null object
diag_3                    101766 non-null object
number_diagnoses          101766 non-null int64
max_glu_serum             101766 non-null object
A1Cresult                 101766 non-null object
metformin                 101766 non-null object
repaglinide               101766 non-null object
nateglinide               101766 non-null object
chlorpropamide            101766 non-null object
glimepiride               101766 non-null object
acetohexamide             101766 non-null object
glipizide                 101766 non-null object
glyburide                 101766 non-null object
tolbutamide               101766 non-null object
pioglitazone              101766 non-null object
rosiglitazone             101766 non-null object
acarbose                  101766 non-null object
miglitol                  101766 non-null object
troglitazone              101766 non-null object
tolazamide                101766 non-null object
examide                   101766 non-null object
citoglipton               101766 non-null object
insulin                   101766 non-null object
glyburide-metformin       101766 non-null object
glipizide-metformin       101766 non-null object
glimepiride-pioglitazone  101766 non-null object
metformin-rosiglitazone   101766 non-null object
metformin-pioglitazone    101766 non-null object
change                    101766 non-null object
diabetesMed                101766 non-null object
readmitted                101766 non-null object
dtypes: int64(13), object(37)
memory usage: 38.8+ MB
None

```

Because not all features are useful, we could do a feature selection before training our model. Our features include much personal information about patients such as age, weight, race and many other medical treatment results.

We can see there are 101766 data and 49 features in the original data. And Y is the class label of each specific data point. For some very obvious features (like "encounter_id", "patient_nbr"), we can say they are just identities and are not related to our prediction (readmitted info) so we just ignore them. For feature examide and citoglipton, they only have 1 value and cannot be used in prediction. Further, among our data there are many missing value as mentioned in the essay. And we can just drop those features with missing rate higher than certain value (eg. 40%).

```
In [3]: for col in data.columns:
        if data[col].dtype == object:
            count = data[col][data[col] == '?'].count()
            if count != 0:
                print(col, count)
        print('gender', data['gender'][data['gender'] == 'Unknown/Invalid'].count())

('race', 2273)
('weight', 98569)
('payer_code', 40256)
('medical_specialty', 49949)
('diag_1', 21)
('diag_2', 358)
('diag_3', 1423)
('gender', 3)
```

```
In [4]: # identities
        data = data.drop('encounter_id', axis=1)
        data = data.drop('patient_nbr', axis=1)
        # only contains one value
        data = data.drop('examide', axis=1)
        data = data.drop('citoglipton', axis=1)
        # missing rate is high
        data = data.drop('weight', axis=1)
        data = data.drop('payer_code', axis=1)
        data = data.drop('medical_specialty', axis=1)
```

We should also remove some records: for those records with diag1, diag2 and diag3 all missing, we should get rid of them. And we also remove records with gender = 'Unknown/Invalid'.

```
In [5]: diag_idx = set(data[(data['diag_1'] == '?') & (data['diag_2'] == '?') & (data['diag_3'] == '?')].index)
        gender_idx = set(data['gender'][data['gender'] == 'Unknown/Invalid'].index)
        data = data.drop(data.index[list(diag_idx.union(gender_idx))])
        print(data.shape)

(101762, 43)
```

For the remaining data missing, we should replace them with np.nan.

```
In [6]: data = data.replace('?', np.nan)
```

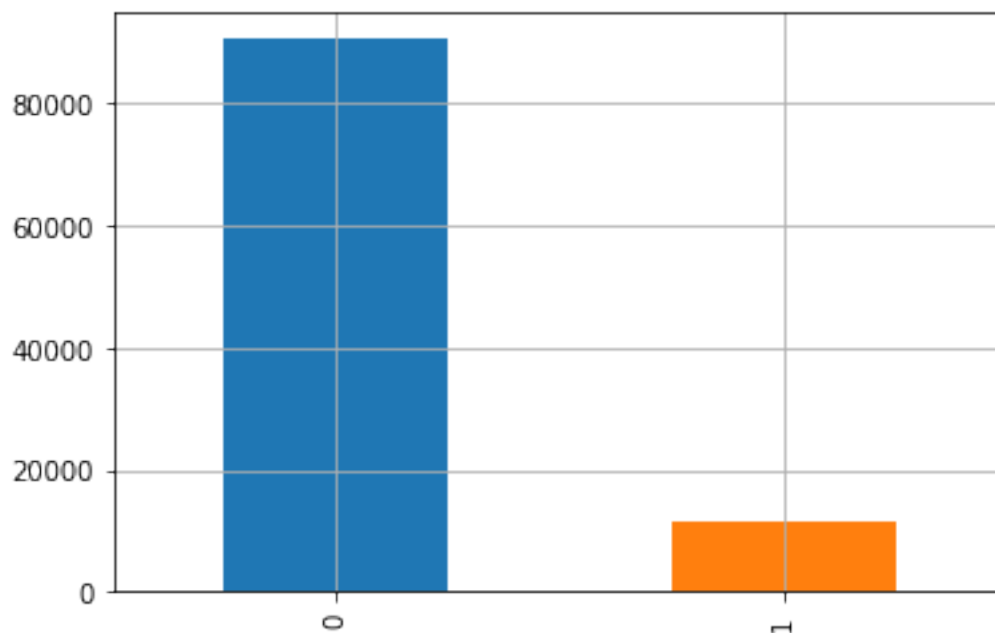
We are predicting whether a patient will occur readmission after treatment. And for patients with >30 days' readmitted, we replace them with 0. For readmitted in more than 30 days has very little correlated factor with first admitted. The distribution of our classes is like the following:

```
In [7]: data['readmitted'] = data['readmitted'].replace('>30', 0)
        data['readmitted'] = data['readmitted'].replace('<30', 1)
```

```

data['readmitted'] = data['readmitted'].replace('NO', 0)
data['readmitted'].value_counts().plot(kind='bar', grid=True)
plt.figure(figsize=(5,3))
plt.show()
print(data.groupby('readmitted').size())
print(data.head().T)

```



<Figure size 360x216 with 0 Axes>

```

readmitted
0    90405
1    11357
dtype: int64

```

	0	1	2	3 \
race	Caucasian	Caucasian	AfricanAmerican	Caucasian
gender	Female	Female	Female	Male
age	[0-10)	[10-20)	[20-30)	[30-40)
admission_type_id	6	1	1	1
discharge_disposition_id	25	1	1	1
admission_source_id	1	7	7	7
time_in_hospital	1	3	2	2
num_lab_procedures	41	59	11	44
num_procedures	0	0	5	1
num_medications	1	18	13	16
number_outpatient	0	0	2	0

number_emergency	0	0	0	0
number_inpatient	0	0	1	0
diag_1	250.83	276	648	8
diag_2	NaN	250.01	250	250.43
diag_3	NaN	255	V27	403
number_diagnoses	1	9	6	7
max_glu_serum	None	None	None	None
A1Cresult	None	None	None	None
metformin	No	No	No	No
repaglinide	No	No	No	No
nateglinide	No	No	No	No
chlorpropamide	No	No	No	No
glimepiride	No	No	No	No
acetohexamide	No	No	No	No
glipizide	No	No	Steady	No
glyburide	No	No	No	No
tolbutamide	No	No	No	No
pioglitazone	No	No	No	No
rosiglitazone	No	No	No	No
acarbose	No	No	No	No
miglitol	No	No	No	No
troglitazone	No	No	No	No
tolazamide	No	No	No	No
insulin	No	Up	No	Up
glyburide-metformin	No	No	No	No
glipizide-metformin	No	No	No	No
glimepiride-pioglitazone	No	No	No	No
metformin-rosiglitazone	No	No	No	No
metformin-pioglitazone	No	No	No	No
change	No	Ch	No	Ch
diabetesMed	No	Yes	Yes	Yes
readmitted	0	0	0	0

	4
race	Caucasian
gender	Male
age	[40-50)
admission_type_id	1
discharge_disposition_id	1
admission_source_id	7
time_in_hospital	1
num_lab_procedures	51
num_procedures	0
num_medications	8
number_outpatient	0
number_emergency	0
number_inpatient	0
diag_1	197

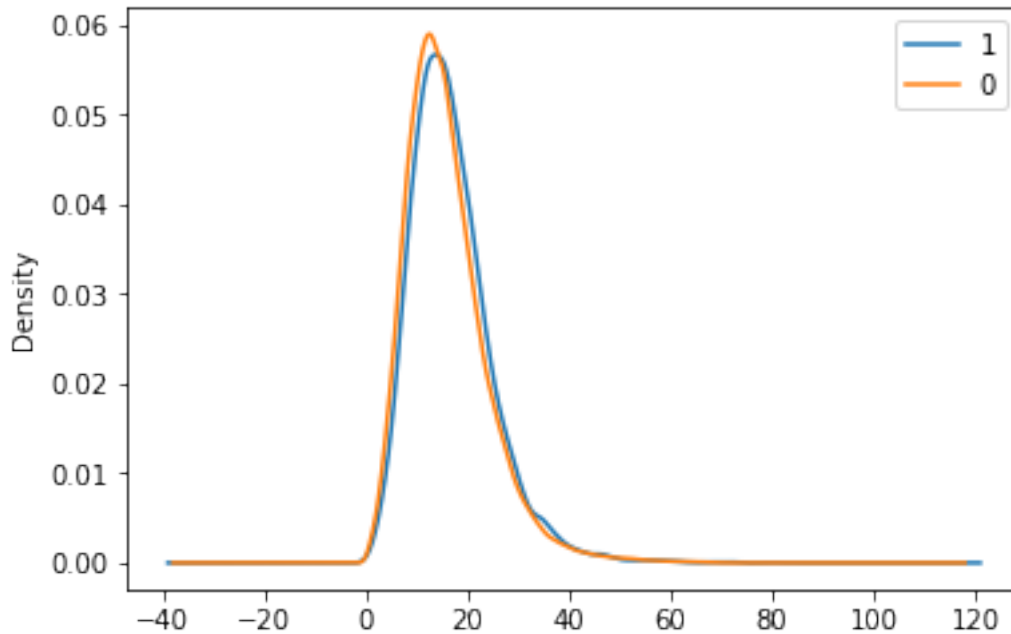
diag_2	157
diag_3	250
number_diagnoses	5
max_glu_serum	None
A1Cresult	None
metformin	No
repaglinide	No
nateglinide	No
chlorpropamide	No
glimepiride	No
acetoexamide	No
glipizide	Steady
glyburide	No
tolbutamide	No
pioglitazone	No
rosiglitazone	No
acarbose	No
miglitol	No
troglitazone	No
tolazamide	No
insulin	Steady
glyburide-metformin	No
glipizide-metformin	No
glimepiride-pioglitazone	No
metformin-rosiglitazone	No
metformin-pioglitazone	No
change	Ch
diabetesMed	Yes
readmitted	0

From bar chart above among 101,767 data, we could see that nearly 90,405 data point with "readmitted", 11,357 labeled "no readmitted". That means patients without readmitted are more frequent and patients with readmitted much smaller which is consistent with common sense. After knowing about our classes, we want to know more about our features which is vital to train our model and make corresponding prediction in the following.

2 Feature selection

For some numeric features, we could plot the density distribution of each feature and get rid of unuseful features in the following (classification density is same over different values).

```
In [8]: # for feature num_medications
data['num_medications'][data['readmitted'] == 1].plot(kind='kde')
data['num_medications'][data['readmitted'] == 0].plot(kind='kde')
plt.legend(('1', '0'),loc='best')
plt.show()
```



For this 'num_medications' feature, we find the density of 0, 1 and 2 almost overlapped. It means it nearly makes no contribution to class decision. We should get rid of it.

```
In [9]: data = data.drop('num_medications', axis=1)
```

After that, we should do some encoding for some nominal features as following:

```
In [10]: from collections import OrderedDict as odict

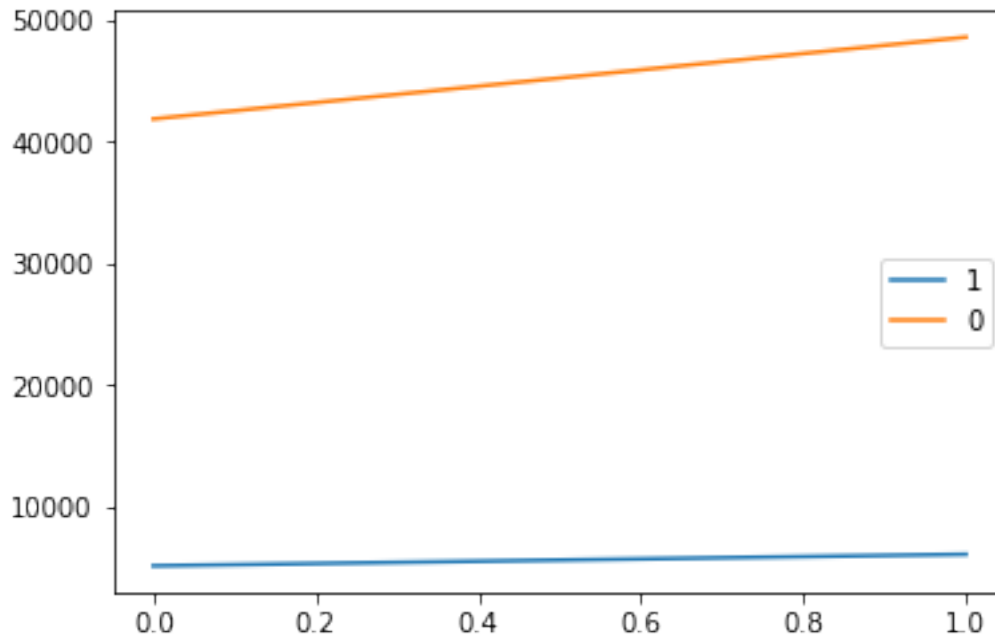
# For feature gender, we should code 'male' and 'female' using 0 and 1
data['gender'] = data['gender'].replace('Male',0)
data['gender'] = data['gender'].replace('Female',1)

feat2 = pd.concat([data.iloc[:,1], data['readmitted']], axis=1)
feat2_lt30 = feat2[feat2['readmitted'] == 1]
feat2_no = feat2[feat2['readmitted'] == 0]

feat2_1 = dict(feat2_lt30['gender'].value_counts())
feat2_1 = odict(sorted(feat2_1.items()))

feat2_2 = dict(feat2_no['gender'].value_counts())
feat2_2 = odict(sorted(feat2_2.items()))

plt.plot(list(feat2_1.keys()), list(feat2_1.values()), label="class_1")
plt.plot(list(feat2_2.keys()), list(feat2_2.values()), label="class_2")
plt.legend(('1', '0'),loc='best')
plt.show()
```



In [11]: # for feature age

```
# for age feature with '[i - i+10)' representation,
# we should replace them with the beginning number to turn this feature into numeric
data['age'] = data['age'].replace(' [0-10)',0)
data['age'] = data['age'].replace(' [10-20)',10)
data['age'] = data['age'].replace(' [20-30)',20)
data['age'] = data['age'].replace(' [30-40)',30)
data['age'] = data['age'].replace(' [40-50)',40)
data['age'] = data['age'].replace(' [50-60)',50)
data['age'] = data['age'].replace(' [60-70)',60)
data['age'] = data['age'].replace(' [70-80)',70)
data['age'] = data['age'].replace(' [80-90)',80)
data['age'] = data['age'].replace(' [90-100)',90)

feat4 = pd.concat([data.iloc[:,2], data['readmitted']], axis=1)
feat4_lt30 = feat4[feat4['readmitted'] == 1]
feat4_no = feat4[feat4['readmitted'] == 0]

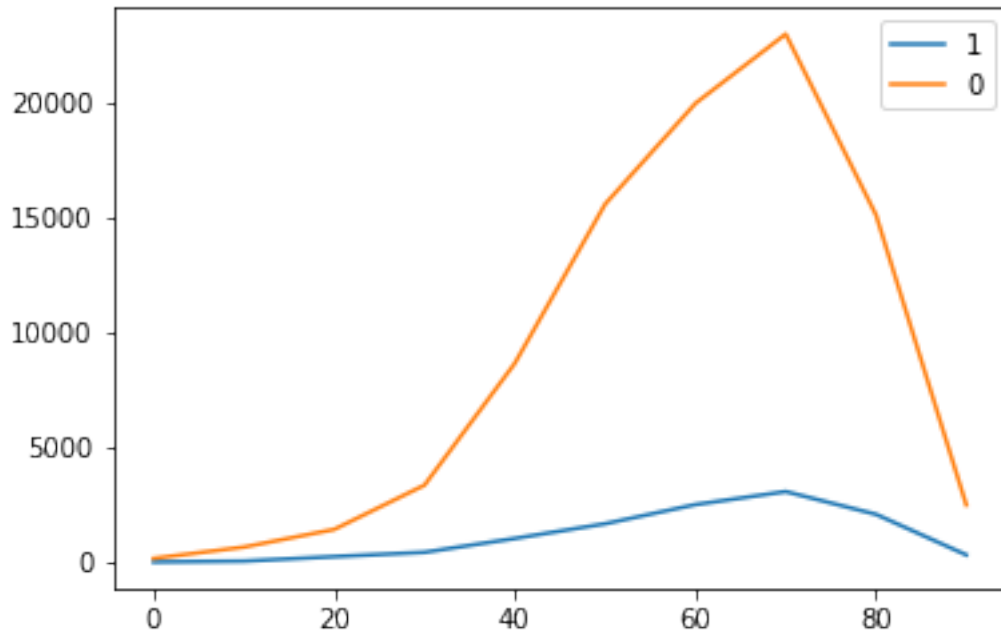
feat4_1 = dict(feat4_lt30['age'].value_counts())
feat4_1 = OrderedDict(sorted(feat4_1.items()))

feat4_2 = dict(feat4_no['age'].value_counts())
feat4_2 = OrderedDict(sorted(feat4_2.items()))

plt.plot(list(feat4_1.keys()), list(feat4_1.values()), label="class_1")
```



```
plt.plot(list(feat4_2.keys()), list(feat4_2.values()), label="class_2")
plt.legend(('1', '0'),loc='best')
plt.show()
```



```
In [12]: # For other discrete features, we should also replace them with numeric value
data['change'] = data['change'].replace('Ch', 1)
data['change'] = data['change'].replace('No', 0)

data['diabetesMed'] = data['diabetesMed'].replace('Yes', 1)
data['diabetesMed'] = data['diabetesMed'].replace('No', 0)

data['max_glu_serum'] = data['max_glu_serum'].replace('>300', 3)
data['max_glu_serum'] = data['max_glu_serum'].replace('>200', 2)
data['max_glu_serum'] = data['max_glu_serum'].replace('Normal', 1)
data['max_glu_serum'] = data['max_glu_serum'].replace('None', 0)

data['A1Cresult'] = data['A1Cresult'].replace('>8', 3)
data['A1Cresult'] = data['A1Cresult'].replace('>7', 2)
data['A1Cresult'] = data['A1Cresult'].replace('Normal', 1)
data['A1Cresult'] = data['A1Cresult'].replace('None', 0)
```

For feature race, it is a discrete Nominal feature with values: Caucasian, Asian, African American, Hispanic, and other. We would like to transform it into a numeric feature. Here we use one-hot encoding. In one-hot encoding, we create 5 additional columns for every values in 'race' and each has value 1 if record has that value or 0 otherwise.

```
In [13]: def create_dummies(data, column_name):
    dummies = pd.get_dummies(data[column_name], prefix=column_name, drop_first = True)
    print(dummies.head().T)
    data = pd.concat([data, dummies], axis=1)
    data = data.drop(column_name, axis=1)
    return data

data = create_dummies(data, 'race')
# data = create_dummies(data, 'max_glu_serum')
# data = create_dummies(data, 'A1Cresult')

      0  1  2  3  4
race_Asian      0  0  0  0  0
race_Caucasian  1  1  0  1  1
race_Hispanic   0  0  0  0  0
race_Other      0  0  0  0  0
```

There are 24 kinds of drug related features, we didn't know the details of each drugs and their efficacy. They indicate whether the dosage of each drug has been changed during treatment. "up" indicates increasing during the encounter, "down" indicates decreasing, "steady" indicates no change, and "no" indicates no prescribed (which also means no change). In order to simplify our model, we only use the number of dosage change during encounter instead of using them all which means we don't distinguish changes between drugs and will combine the changes together among all 24 changes.

```
In [14]: drugs = ['metformin', 'repaglinide', 'nateglinide', 'chlorpropamide',
    'glimepiride', 'glipizide', 'glyburide', 'pioglitazone',
    'rosiglitazone', 'acarbose', 'miglitol', 'insulin',
    'glyburide-metformin', 'tolazamide',
    'metformin-pioglitazone', 'metformin-rosiglitazone',
    'glimepiride-pioglitazone', 'glipizide-metformin',
    'troglitazone', 'tolbutamide', 'acetohexamide']

data['drug_change_num'] = 0
for drug in drugs:
    data['drug_change_num'] += data[drug].apply(lambda x: 0 if (x == 'No' or x == 'Sto
    data = data.drop(drug, axis=1)

print(data['drug_change_num'].value_counts())

0    74059
1    26272
2     1318
3      108
4        5
Name: drug_change_num, dtype: int64
```

After all data preprocessing is done, we should shuffle our data and split them into training data (80%) and testing data (20%).

```
In [15]: data = data.sample(n = len(data)).reset_index(drop = True)
        print(data.shape)
```

(101762, 25)

```
In [16]: Y = data.readmitted
        X = data.drop('readmitted', axis=1)
        print(X.shape)
        print(Y.shape)
        X = X.apply(pd.to_numeric, errors='coerce')
        Y = Y.apply(pd.to_numeric, errors='coerce')
        X.fillna(0, inplace=True)
        Y.fillna(0, inplace=True)
        X_train, X_test, y_train, y_test = train_test_split(X, Y, test_size=0.2)
```

(101762, 24)

(101762,)

3 Model build

3.1 KNN

We first choose KNN classifier model. We use our training data to train the model and make prediction for both training data and testing data and finally calculate ROC AUC and Accuracy for our prediction.

```
In [22]: from sklearn.neighbors import KNeighborsClassifier
        neigh = KNeighborsClassifier(n_neighbors=50)
        neigh.fit(X_train, y_train)
        # print(X_train.info())

        knn_y_train_predict = neigh.predict_proba(X_train)[:,-1]
        knn_y_test_predict = neigh.predict_proba(X_test)[:,-1]

        from sklearn.metrics import roc_auc_score, accuracy_score

        threshold = 0.5\

        print('Training result:')
        fpr_train, tpr_train, thresholds_train = roc_curve(y_train, knn_y_train_predict)
        print('AUC=', auc(fpr_train, tpr_train))
        print('Accuracy=', accuracy_score(y_train, (knn_y_train_predict > threshold)))

        print('Testing result:')
        fpr_test, tpr_test, thresholds_test = roc_curve(y_test, knn_y_test_predict)
        print('AUC=', auc(fpr_test, tpr_test))
        print('Accuracy=', accuracy_score(y_test, (knn_y_test_predict > threshold)))
```

```
Training result:
('AUC=', 0.652560500760044)
('Accuracy=', 0.8885012713582037)
Testing result:
('AUC=', 0.5478778285548946)
('Accuracy=', 0.8883702648258242)
```

3.2 Gradient boosting

This time we use a gradient boosting classifier model.

```
In [21]: from sklearn.ensemble import GradientBoostingClassifier

gboost = GradientBoostingClassifier(learning_rate=0.1, n_estimators=100,
                                     subsample=0.8, min_samples_split=2,
                                     min_samples_leaf=1, max_depth=3)

gboost.fit(X_train, y_train)

gboost_y_train_predict = gboost.predict_proba(X_train)[:,-1]
gboost_y_test_predict = gboost.predict_proba(X_test)[:,-1]

print('Training result:')
fpr_train, tpr_train, thresholds_train = roc_curve(y_train, gboost_y_train_predict)
print('AUC=', auc(fpr_train, tpr_train))
print('Accuracy=', accuracy_score(y_train, (gboost_y_train_predict > threshold)))

print('Testing result:')
fpr_test, tpr_test, thresholds_test = roc_curve(y_test, gboost_y_test_predict)
print('AUC=', auc(fpr_test, tpr_test))
print('Accuracy=', accuracy_score(y_test, (gboost_y_test_predict > threshold)))
```

```
Training result:
('AUC=', 0.6882818744849524)
('ROC AUC=', 0.6882818744849524)
('Accuracy=', 0.8895699492685084)
Testing result:
('AUC=', 0.6721436777626622)
('ROC AUC=', 0.6721436777626622)
('Accuracy=', 0.8887633272736206)
```

4 Random forest

```
In [26]: from sklearn.ensemble import RandomForestClassifier
         from sklearn.metrics import roc_curve, auc

         max_depth_arr = [2,4,6,8,10,12,14,16,18,20]
```

```

min_sample_split=[4,8,12,16,20,24]
tr_auc = np.zeros((len(max_depth_arr),len(min_sample_split)))
va_auc = np.zeros((len(max_depth_arr),len(min_sample_split)))
for i,d in enumerate(max_depth_arr):
    for j,s in enumerate(min_sample_split):
        clf = RandomForestClassifier(n_estimators=100, n_jobs=4,
                                    max_depth=d, min_samples_split=s)

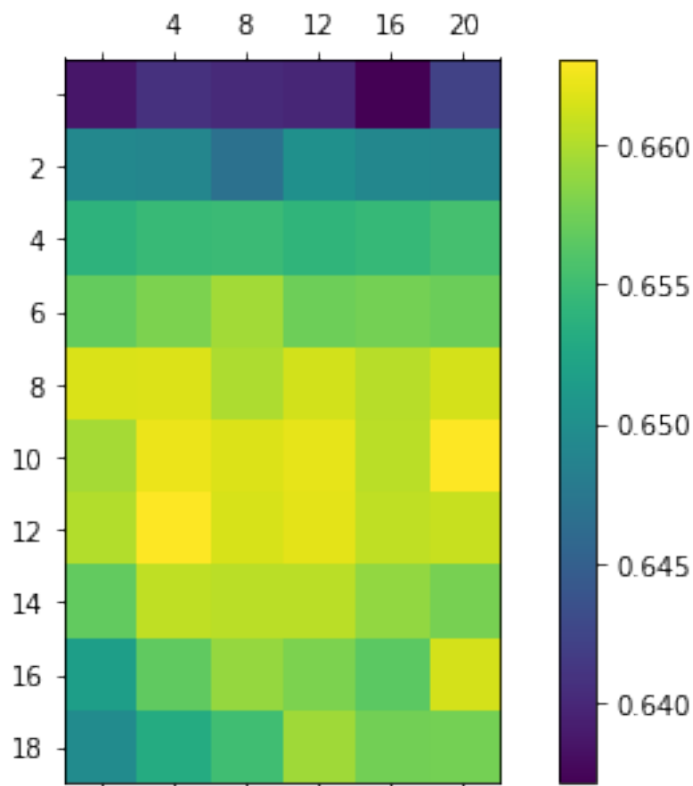
        clf.fit(X_train, y_train)
        randomforest_y_test_predict = clf.predict_proba(X_test)[:,:1]
        fpr, tpr, thresholds = roc_curve(y_test, randomforest_y_test_predict)
        va_auc[i][j]=auc(fpr, tpr)
f, ax = plt.subplots(1, 1, figsize=(8, 5))
cax = ax.matshow(va_auc, interpolation='nearest')
f.colorbar(cax)
ax.set_yticklabels(['']+list(max_depth_arr))
ax.set_xticklabels(['']+list(min_sample_split))
ax.set_yticks(np.arange(len(max_depth_arr)))
ax.set_xticks(np.arange(len(min_sample_split)))

```

```

Out[26]: [<matplotlib.axis.XTick at 0x1a158ee5d0>,
<matplotlib.axis.XTick at 0x1a158eeb10>,
<matplotlib.axis.XTick at 0x1a158ee890>,
<matplotlib.axis.XTick at 0x1030ed2d0>,
<matplotlib.axis.XTick at 0x1030ed890>,
<matplotlib.axis.XTick at 0x1030ede50>]

```



From plot above, we could know min sample split produce the best result when it equals to 12 and max depth produce best result when it equals to 10

```
In [20]: results=np.zeros(20)
print(results.shape)
x=range(0,20)
for i in range(0,20):
    clf = RandomForestClassifier(n_estimators=20*i+1, n_jobs=4, max_depth=10,
                               min_samples_split=12, min_samples_leaf=1)

    clf.fit(X_train, y_train)
    randomforest_y_test_predict1=clf.predict_proba(X_test)[:,-1]
    fpr, tpr, thresholds = roc_curve(y_test, randomforest_y_test_predict1)
    results[i]=auc(fpr, tpr)
    x[i] = 20*i+1
plt.plot(x,results)
plt.show()
```

(20,)

