# ML0101EN-Clas-SVM-cancer-py-v1

## August 12, 2019

## **SVM** (Support Vector Machines)

In this notebook, you will use SVM (Support Vector Machines) to build and train a model using human cell records, and classify cells to whether the samples are benign or malignant.

SVM works by mapping data to a high-dimensional feature space so that data points can be categorized, even when the data are not otherwise linearly separable. A separator between the categories is found, then the data is transformed in such a way that the separator could be drawn as a hyperplane. Following this, characteristics of new data can be used to predict the group to which a new record should belong.

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```
[1]: import pandas as pd
import pylab as pl
import numpy as np
import scipy.optimize as opt
from sklearn import preprocessing
from sklearn.model_selection import train_test_split
%matplotlib inline
import matplotlib.pyplot as plt
```

#### Load the Cancer data

The example is based on a dataset that is publicly available from the UCI Machine Learning Repository (Asuncion and Newman, 2007)[http://mlearn.ics.uci.edu/MLRepository.html]. The dataset consists of several hundred human cell sample records, each of which contains the values of a set of cell characteristics. The fields in each record are:

Field name	Description
ID	Clump thickness
Clump	Clump thickness
UnifSize	Uniformity of cell size
UnifShape	Uniformity of cell shape

Field name	Description
MargAdh	Marginal adhesion
SingEpiSize	Single epithelial cell size
BareNuc	Bare nuclei
BlandChrom	Bland chromatin
NormNucl	Normal nucleoli
Mit	Mitoses
Class	Benign or malignant

For the purposes of this example, we're using a dataset that has a relatively small number of predictors in each record. To download the data, we will use !wget to download it from IBM Object Storage.

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```
[2]: #Click here and press Shift+Enter
!wget -O cell_samples.csv https://s3-api.us-geo.objectstorage.softlayer.net/cf-courses-data/

CognitiveClass/ML0101ENv3/labs/cell_samples.csv
```

```
--2019-08-12 08:58:51-- https://s3-api.us-geo.objectstorage.softlayer.net/cf-courses-data/CognitiveClass/ML0101ENv3/labs/cell_samples.csv Resolving s3-api.us-geo.objectstorage.softlayer.net (s3-api.us-geo.objectstorage.softlayer.net)... 67.228.254.193 Connecting to s3-api.us-geo.objectstorage.softlayer.net (s3-api.us-geo.objectstorage.softlayer.net)|67.228.254.193|:443... connected. HTTP request sent, awaiting response... 200 OK Length: 20675 (20K) [text/csv] Saving to: cell_samples.csv 100\%[========================] 20.19K --.-KB/s \text{ in } 0.02s 2019-08-12 \ 08:58:51 \ (816 \ KB/s) - \text{cell samples.csv saved } [20675/20675]
```

#### 0.0.1 Load Data From CSV File

```
[3]: |\text{cell df} = \text{pd.read csv("cell samples.csv")}|
    cell df.head()
[3]:
           ID Clump UnifSize UnifShape MargAdh SingEpiSize BareNuc \
    0 1000025
                    5
                            1
                                      1
                                             1
                                                        2
                                                              1
                                                        7
                                                              10
    1 1002945
                    5
                            4
                                     4
                                             5
    2 1015425
                    3
                            1
                                     1
                                             1
                                                        2
                                                              2
    3 1016277
                    6
                            8
                                     8
                                             1
                                                        3
                                                              4
                                                        2
    4 1017023
                    4
                            1
                                     1
                                             3
                                                              1
```

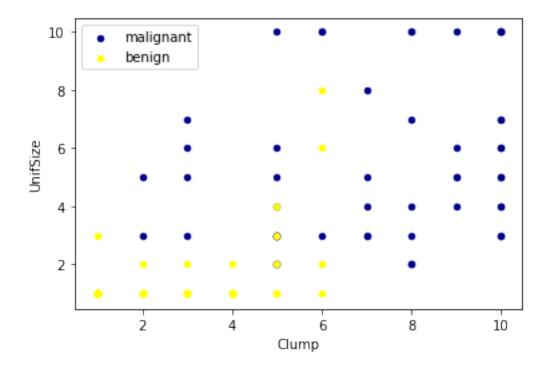
BlandChrom NormNucl Mit Class

```
0
                                2
           3
                     1
                          1
1
           3
                     2
                                 2
2
           3
                                2
                     1
                          1
3
           3
                     7
                                2
                          1
4
           3
                     1
                          1
                                 2
```

The ID field contains the patient identifiers. The characteristics of the cell samples from each patient are contained in fields Clump to Mit. The values are graded from 1 to 10, with 1 being the closest to benign.

The Class field contains the diagnosis, as confirmed by separate medical procedures, as to whether the samples are benign (value = 2) or malignant (value = 4).

Lets look at the distribution of the classes based on Clump thickness and Uniformity of cell size:



# 0.1 Data pre-processing and selection

Lets first look at columns data types:

```
[9]: cell_df.dtypes
```

```
[9]: ID
                  int64
     Clump
                    int64
     UnifSize
                   int64
     UnifShape
                    int64
     MargAdh
                     int64
     SingEpiSize
                    int64
     BareNuc
                    object
     BlandChrom
                      int64
     NormNucl
                     int64
     Mit
                  int64
     Class
                  int64
     dtype: object
        It looks like the BareNuc column includes some values that are not numerical. We can drop
     those rows:
[10]: cell df = cell df[pd.to numeric(cell df['BareNuc'], errors='coerce').notnull()]
     cell df['BareNuc'] = cell df['BareNuc'].astype('int')
     cell df.dtypes
[10]: ID
                 int64
     Clump
                   int64
     UnifSize
                  int64
     UnifShape
                    int64
     MargAdh
                    int64
                    int64
     SingEpiSize
     BareNuc
                    int64
     BlandChrom
                     int64
     NormNucl
                    int64
     Mit
                  int64
```

```
[11]: feature_df = cell_df[['Clump', 'UnifSize', 'UnifShape', 'MargAdh', 'SingEpiSize', 'BareNuc',

→'BlandChrom', 'NormNucl', 'Mit']]

X = np.asarray(feature_df)

X[0:5]
```

```
[11]: array([[ 5, 1, 1, 1, 2, 1, 3, 1, 1], [ 5, 4, 4, 5, 7, 10, 3, 2, 1], [ 3, 1, 1, 1, 2, 2, 3, 1, 1], [ 6, 8, 8, 1, 3, 4, 3, 7, 1], [ 4, 1, 1, 3, 2, 1, 3, 1, 1]])
```

int64

Class

dtype: object

We want the model to predict the value of Class (that is, benign (=2) or malignant (=4)). As this field can have one of only two possible values, we need to change its measurement level to reflect this.

```
[12]: cell_df['Class'] = cell_df['Class'].astype('int')

y = np.asarray(cell_df['Class'])

y = [0:5]
```

```
[12]: array([2, 2, 2, 2, 2])
```

#### 0.2 Train/Test dataset

Okay, we split our dataset into train and test set:

```
[13]: X_train, X_test, y_train, y_test = train_test_split( X, y, test_size=0.2, random_state=4) print ('Train set:', X_train.shape, y_train.shape) print ('Test set:', X_test.shape, y_test.shape)
```

```
Train set: (546, 9) (546,)
Test set: (137, 9) (137,)
```

Modeling (SVM with Scikit-learn)

The SVM algorithm offers a choice of kernel functions for performing its processing. Basically, mapping data into a higher dimensional space is called kernelling. The mathematical function used for the transformation is known as the kernel function, and can be of different types, such as:

- 1.Linear
- 2.Polynomial
- 3. Radial basis function (RBF)
- 4.Sigmoid

Each of these functions has its characteristics, its pros and cons, and its equation, but as there's no easy way of knowing which function performs best with any given dataset, we usually choose different functions in turn and compare the results. Let's just use the default, RBF (Radial Basis Function) for this lab.

/home/jupyterlab/conda/envs/python/lib/python3.6/site-packages/sklearn/svm/base.py:196: FutureWarning: The default value of gamma will change from 'auto' to 'scale' in version 0.22 to account better for unscaled features. Set gamma explicitly to 'auto' or 'scale' to avoid this warning.

"avoid this warning.", FutureWarning)

```
[14]: SVC(C=1.0, cache_size=200, class_weight=None, coef0=0.0, decision_function_shape='ovr', degree=3, gamma='auto_deprecated', kernel='rbf', max_iter=-1, probability=False, random_state=None, shrinking=True, tol=0.001, verbose=False)
```

After being fitted, the model can then be used to predict new values:

```
[15]: yhat = clf.predict(X_test)

yhat [0:5]
```

[15]: array([2, 4, 2, 4, 2])

Evaluation

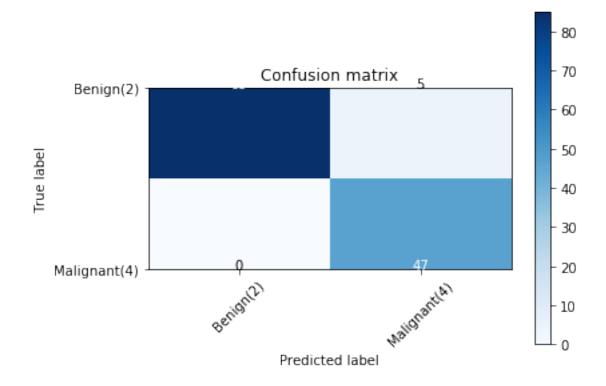
```
[16]: from sklearn.metrics import classification report, confusion matrix
      import itertools
[17]: def plot confusion matrix(cm, classes,
                         normalize=False,
                         title='Confusion matrix',
                         cmap=plt.cm.Blues):
         This function prints and plots the confusion matrix.
         Normalization can be applied by setting `normalize=True`.
         if normalize:
            cm = cm.astype('float') / cm.sum(axis=1)[:, np.newaxis]
            print("Normalized confusion matrix")
            print('Confusion matrix, without normalization')
         print(cm)
         plt.imshow(cm, interpolation='nearest', cmap=cmap)
         plt.title(title)
         plt.colorbar()
         tick marks = np.arange(len(classes))
         plt.xticks(tick marks, classes, rotation=45)
         plt.yticks(tick marks, classes)
         fmt = '.2f' if normalize else 'd'
         thresh = cm.max() / 2.
         for i, j in itertools.product(range(cm.shape[0]), range(cm.shape[1])):
            plt.text(j, i, format(cm[i, j], fmt),
                  horizontalalignment="center",
                  color="white" if cm[i, j] > thresh else "black")
         plt.tight layout()
         plt.ylabel('True label')
         plt.xlabel('Predicted label')
[18]: # Compute confusion matrix
      conf matrix = confusion matrix(y test, yhat, labels=[2,4])
      np.set printoptions(precision=2)
      print (classification report(y test, yhat))
      # Plot non-normalized confusion matrix
      plt.figure()
      plot confusion matrix(cnf matrix, classes=['Benign(2)','Malignant(4)'],normalize=False,
       →title='Confusion matrix')
```

precision recall f1-score support 2 1.00 0.940.9790 4 0.901.00 0.9547 micro avg 0.960.960.96137 macro avg 0.950.97 0.96137 weighted avg 0.970.960.96137

 $Confusion\ matrix,\ without\ normalization$ 

 $[[85 \ 5]$ 

 $[0 \ 47]]$ 



### You can also easily use the **f1\_score** from sklearn library:

```
[19]: from sklearn.metrics import f1_score f1_score(y_test, yhat, average='weighted')
```

[19]: 0.9639038982104676

# Lets try jaccard index for accuracy:

[20]: 0.9635036496350365

Practice

Can you rebuild the model, but this time with a **linear** kernel? You can use **kernel='linear'** option, when you define the sym. How the accuracy changes with the new kernel function?

```
[22]: # write your code here
clf2 = svm.SVC(kernel='linear')
clf2.fit(X_train, y_train)
yhat2 = clf2.predict(X_test)
print("Avg F1-score: %.4f" % f1_score(y_test, yhat2, average='weighted'))
print("Jaccard score: %.4f" % jaccard_similarity_score(y_test, yhat2))
```

Avg F1-score: 0.9639 Jaccard score: 0.9635

Double-click here for the solution.

Want to learn more?

IBM SPSS Modeler is a comprehensive analytics platform that has many machine learning algorithms. It has been designed to bring predictive intelligence to decisions made by individuals, by groups, by systems – by your enterprise as a whole. A free trial is available through this course, available here: SPSS Modeler

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Thanks for completing this lesson!

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