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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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In [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 (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
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.

Did you know? When it comes to Machine Learning, you will likely be working with large datasets. As a business, where can you host your data? IBM is offering a unique opportunity for businesses, with 10 Tb of IBM Cloud Object Storage: Sign up now for free (http://cocl.us/ML0101EN-IBM-Offer-CC)

In [2]:

```
ses-data/CognitiveClass/ML0101ENv3/labs/cell samples.csv
--2019-10-12 06:49:09-- 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.obje
ctstorage.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'
cell samples.csv
                    100%[========>]
                                                20.19K --.-KB/s
                                                                    in
0.02s
2019-10-12 06:49:09 (1.01 MB/s) - 'cell_samples.csv' saved [20675/2067
5 ]
```

!wget -0 cell samples.csv https://s3-api.us-geo.objectstorage.softlayer.net/cf-cour

Load Data From CSV File

#Click here and press Shift+Enter

In [3]:

```
cell_df = pd.read_csv("cell_samples.csv")
cell_df.head()
```

Out[3]:

	ID	Clump	UnifSize	UnifShape	MargAdh	SingEpiSize	BareNuc	BlandChrom	NormNu
0	1000025	5	1	1	1	2	1	3	
1	1002945	5	4	4	5	7	10	3	
2	1015425	3	1	1	1	2	2	3	
3	1016277	6	8	8	1	3	4	3	
4	1017023	4	1	1	3	2	1	3	

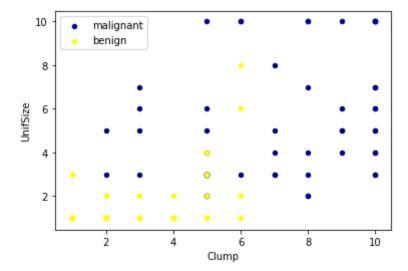
The ID field contains the patient identifiers. The characteristics of the cell samples from each patient are contained in fields Clump to Mit. The values are graded from 1 to 10, with 1 being the closest to benign.

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:

```
In [4]:
```

```
ax = cell_df[cell_df['Class'] == 4][0:50].plot(kind='scatter', x='Clump', y='UnifSi
ze', color='DarkBlue', label='malignant');
cell_df[cell_df['Class'] == 2][0:50].plot(kind='scatter', x='Clump', y='UnifSize',
color='Yellow', label='benign', ax=ax);
plt.show()
```



Data pre-processing and selection

Lets first look at columns data types:

```
In [5]:
```

```
cell_df.dtypes
```

Out[5]:

```
int64
ID
                 int64
Clump
UnifSize
                 int64
UnifShape
                 int64
MargAdh
                 int64
                 int64
SingEpiSize
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:

```
In [6]:
cell df = cell_df[pd.to_numeric(cell_df['BareNuc'], errors='coerce').notnull()]
cell df['BareNuc'] = cell df['BareNuc'].astype('int')
cell_df.dtypes
Out[6]:
                int64
ID
Clump
                int64
UnifSize
                int64
UnifShape
                int64
MargAdh
                int64
                int64
SingEpiSize
BareNuc
                int64
BlandChrom
                int64
NormNucl
                int64
Mit
                int64
Class
                int64
dtype: object
In [7]:
feature_df = cell_df[['Clump', 'UnifSize', 'UnifShape', 'MargAdh', 'SingEpiSize',
'BareNuc', 'BlandChrom', 'NormNucl', 'Mit']]
X = np.asarray(feature_df)
X[0:5]
Out[7]:
array([[ 5,
              1,
                  1,
                      1,
                           2, 1,
                                   3,
                                        1,
                                            1],
                           7, 10,
                                        2,
       [5,
              4,
                  4,
                       5,
                                    3,
                                            1],
                                    3,
                                        1,
                                            1],
                       1,
                           2,
                               2,
       [ 3,
              1,
                  1,
              8,
                  8,
                      1,
                           3,
                               4,
                                    3,
                                        7,
                                            1],
       [6,
                           2,
                                   3,
       [4,
              1,
                  1,
                       3,
                               1,
                                        1,
                                            1]])
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.
```

```
In [8]:

cell_df['Class'] = cell_df['Class'].astype('int')
y = np.asarray(cell_df['Class'])
y [0:5]

Out[8]:
array([2, 2, 2, 2, 2])
```

Train/Test dataset

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

```
In [9]:
```

```
X_train, X_test, y_train, y_test = train_test_split( X, y, test_size=0.2, random_st
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.

```
In [10]:
```

Out[10]:

```
from sklearn import svm
clf = svm.SVC(kernel='rbf')
clf.fit(X train, y train)
```

/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 f eatures. Set gamma explicitly to 'auto' or 'scale' to avoid this warnin "avoid this warning.", FutureWarning)

```
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:

```
In [11]:
    yhat = clf.predict(X_test)
    yhat [0:5]
Out[11]:
    array([2, 4, 2, 4, 2])
```

Evaluation

```
In [12]:
```

```
from sklearn.metrics import classification_report, confusion_matrix
import itertools
```

In [13]:

```
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')
```

In [14]:

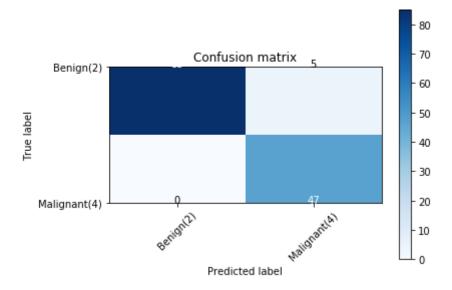
```
# Compute confusion matrix
cnf_matrix = confusion_matrix(y_test, yhat, labels=[2,4])
np.set_printoptions(precision=2)

print (classification_report(y_test, yhat))

# Plot non-normalized confusion matrix
plt.figure()
plot_confusion_matrix(cnf_matrix, classes=['Benign(2)','Malignant(4)'],normalize= F
alse, title='Confusion matrix')
```

		precision	recall	f1-score	support
	2	1.00	0.94	0.97	90
	4	0.90	1.00	0.95	47
micro	avg	0.96	0.96	0.96	137
macro	avg	0.95	0.97	0.96	137
weighted	avg	0.97	0.96	0.96	137

Confusion matrix, without normalization
[[85 5]
 [0 47]]



You can also easily use the **f1_score** from sklearn library:

In [15]:

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

Out[15]:

0.9639038982104676

Lets try jaccard index for accuracy:

```
In [16]:
```

```
from sklearn.metrics import jaccard_similarity_score
jaccard_similarity_score(y_test, yhat)
```

Out[16]:

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 svm. How the accuracy changes with the new kernel function?

In [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

In [21]:

```
jaccard_similarity_score(y_test, yhat2)
```

Out[21]:

0.9635036496350365

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 (http://cocl.us/ML0101EN-SPSSModeler)

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Thanks for completing this lesson!

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