

# **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.

# **Table of contents**

- 1. Load the Cancer data
- 2. Modeling
- 3. Evaluation
- 4. Practice

### 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]. 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				

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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#### In [2]:

```
#Click here and press Shift+Enter
!wget -0 cell samples.csv https://s3-api.us-geo.objectstorage.softlayer.net/cf-courses-data/Cognit
iveClass/ML0101ENv3/labs/cell samples.csv
--2020-06-16 06:54:05-- 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)...
67.228.254.196
Connecting to s3-api.us-geo.objectstorage.softlayer.net (s3-api.us-
geo.objectstorage.softlayer.net)|67.228.254.196|:443... connected.
HTTP request sent, awaiting response... 200 OK
Length: 20675 (20K) [text/csv]
Saving to: 'cell samples.csv'
                  in 0.02s
cell samples.csv
2020-06-16 06:54:05 (971 KB/s) - 'cell samples.csv' saved [20675/20675]
```

#### **Load Data From CSV File**

```
In [3]:
```

```
cell df = pd.read csv("cell samples.csv")
cell df.head()
```

# Out[3]:

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

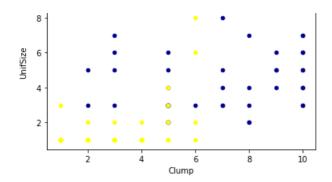
The ID field contains the patient identifiers. The characteristics of the cell samples from each patient are contained in fields Clump to Mit. The values are graded from 1 to 10, with 1 being the closest to benign.

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:

```
In [4]:
```

```
ax = cell df[cell df['Class'] == 4][0:50].plot(kind='scatter', x='Clump', y='UnifSize', color='Dark
Blue', 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
             int64
int64
UnifSize
UnifShape
             int64
MargAdh
SingEpiSize
             int64
BareNuc
           object
            int64
BlandChrom
NormNucl
             int64
Mit
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]:
ID
              int64
             int64
Clump
UnifSize
UnifShape int64
MargAdh int64
SingEpiSize int64
BareNuc int64
BlandChrom int64
NormNucl
              int64
Mit
               int64
Class
                int64
dtype: object
In [7]:
feature_df = cell_df[['Clump', 'UnifSize', 'UnifShape', 'MargAdh', 'SingEpiSize', 'BareNuc', 'Bland
Chrom', 'NormNucl', 'Mit']]
X = np.asarray(feature df)
X[0:5]
Out[7]:
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]])
```

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_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:

```
    Linear
    Polynomial
    Radial basis function (RBF)
    Sigmoid
```

shrinking=True, tol=0.001, verbose=False)

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]:
```

```
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 features. Set gamma explicitly to 'auto' or 'scale' to avoid this warning.
   "avoid this warning.", FutureWarning)

Out[10]:
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,
```

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")
    else:
       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]:

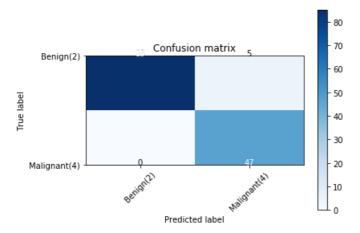
2 1.00 0.94 0.97	ort
4 0.90 1.00 0.95 4	90 47

```
        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 [17]:

```
# write your code here
```

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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