priyadarshini-sm-hw4

March 4, 2024

1 Support Vector Machines (SVM)

2 HW 4

2.0.1 Set up

```
[3]: # Python 3.5 is required
import sys
assert sys.version_info >= (3, 5)

# Scikit-Learn 0.20 is required
import sklearn
assert sklearn.__version__ >= "0.20"

# Common imports
import numpy as np
import pandas as pd
import os

np.random.seed(42)

%matplotlib inline
import matplotlib as mpl
import matplotlib.pyplot as plt
```

2.0.2 utility functions

```
[4]: def plot_class_regions_for_classifier_subplot(clf, X, y, X_test, y_test, title, u_subplot, target_names = None, plot_decision_regions = True):

numClasses = np.amax(y) + 1
color_list_light = ['#FFFFAA', '#EFEFEF', '#AAFFAA', '#AAAAFF']
color_list_bold = ['#EEEE00', '#000000', '#0000000', '#000000C']
cmap_light = ListedColormap(color_list_light[0:numClasses])
cmap_bold = ListedColormap(color_list_bold[0:numClasses])
h = 0.03
```

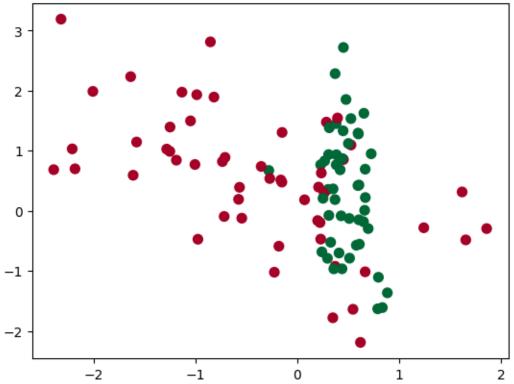
```
k = 0.5
    x_plot_adjust = 0.1
    y_plot_adjust = 0.1
    plot_symbol_size = 50
    x_min = X[:, 0].min()
    x_max = X[:, 0].max()
    y_min = X[:, 1].min()
    y_max = X[:, 1].max()
    x2, y2 = np.meshgrid(np.arange(x_min-k, x_max+k, h), np.arange(y_min-k,_
 \rightarrowy max+k, h))
    P = clf.predict(np.c_[x2.ravel(), y2.ravel()])
    P = P.reshape(x2.shape)
    if plot_decision_regions:
        subplot.contourf(x2, y2, P, cmap=cmap_light, alpha = 0.8)
    subplot.scatter(X[:, 0], X[:, 1], c=y, cmap=cmap_bold, s=plot_symbol_size,_
 ⇔edgecolor = 'black')
    subplot.set_xlim(x_min - x_plot_adjust, x_max + x_plot_adjust)
    subplot.set_ylim(y_min - y_plot_adjust, y_max + y_plot_adjust)
    if (X_test is not None):
        subplot.scatter(X_test[:, 0], X_test[:, 1], c=y_test, cmap=cmap_bold,_
 s=plot_symbol_size, marker='^', edgecolor = 'black')
        train score = clf.score(X, y)
        test_score = clf.score(X_test, y_test)
        title = title + "\nTrain score = {:.2f}, Test score = {:.2f}".
 →format(train_score, test_score)
    subplot.set_title(title)
    if (target_names is not None):
        legend handles = []
        for i in range(0, len(target_names)):
            patch = mpatches.Patch(color=color_list_bold[i],__
 →label=target_names[i])
            legend_handles.append(patch)
        subplot.legend(loc=0, handles=legend_handles)
def plot_class_regions_for_classifier(clf, X, y, X_test=None, y_test=None, u
 stitle=None, target_names = None, plot_decision_regions = True):
    numClasses = np.amax(y) + 1
```

```
color_list_light = ['#FFFFAA', '#EFEFEF', '#AAFFAA', '#AAAAFF']
  color_list_bold = ['#EEEE00', '#000000', '#000000', '#000000']
  cmap_light = ListedColormap(color_list_light[0:numClasses])
  cmap_bold = ListedColormap(color_list_bold[0:numClasses])
  h = 0.03
  k = 0.5
  x_plot_adjust = 0.1
  y_plot_adjust = 0.1
  plot_symbol_size = 50
  x_min = X[:, 0].min()
  x_max = X[:, 0].max()
  y_min = X[:, 1].min()
  y_max = X[:, 1].max()
  x2, y2 = np.meshgrid(np.arange(x_min-k, x_max+k, h), np.arange(y_min-k,_
\rightarrowy_max+k, h))
  P = clf.predict(np.c_[x2.ravel(), y2.ravel()])
  P = P.reshape(x2.shape)
  plt.figure()
  if plot_decision_regions:
      plt.contourf(x2, y2, P, cmap=cmap_light, alpha = 0.8)
  plt.scatter(X[:, 0], X[:, 1], c=y, cmap=cmap_bold, s=plot_symbol_size,__
⇔edgecolor = 'black')
  plt.xlim(x_min - x_plot_adjust, x_max + x_plot_adjust)
  plt.ylim(y_min - y_plot_adjust, y_max + y_plot_adjust)
  if (X test is not None):
      plt.scatter(X_test[:, 0], X_test[:, 1], c=y_test, cmap=cmap_bold,__
⇔s=plot_symbol_size, marker='^', edgecolor = 'black')
      train_score = clf.score(X, y)
      test_score = clf.score(X_test, y_test)
      title = title + "\nTrain score = {:.2f}, Test score = {:.2f}".
→format(train_score, test_score)
  if (target_names is not None):
      legend_handles = []
      for i in range(0, len(target_names)):
          patch = mpatches.Patch(color=color_list_bold[i],__
⇔label=target_names[i])
          legend_handles.append(patch)
      plt.legend(loc=0, handles=legend_handles)
  if (title is not None):
      plt.title(title)
```

```
plt.show()
```

2.0.3 Synthetic dataset

Sample binary classification problem with two informative features



2.0.4 Linear Support Vector Machine

```
[6]: from sklearn.svm import SVC
from sklearn.model_selection import train_test_split
from sklearn.metrics import classification_report

X_train, X_test, y_train, y_test = train_test_split(X, y, random_state = 0)

clf = SVC(kernel = 'linear', C=1.0).fit(X_train, y_train)

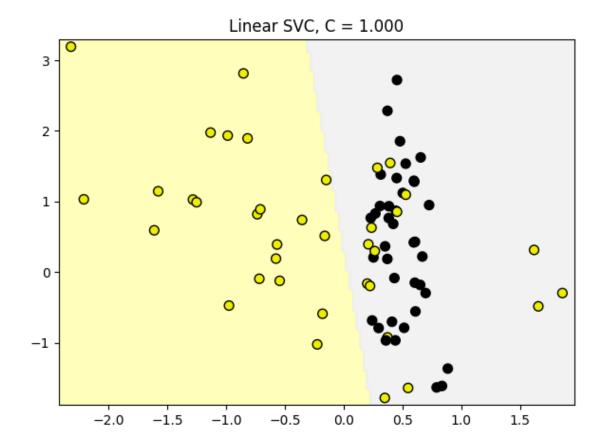
y_pred = clf.predict(X_test)

result_metrics = classification_report(y_test, y_pred)
print(result_metrics)

fig, subaxes = plt.subplots(1, 1, figsize=(7, 5))

title = 'Linear SVC, C = {:.3f}'.format(1.0)
plot_class_regions_for_classifier_subplot(clf, X_train, y_train, None, None, u_stitle, subaxes)
```

	precision	recall	f1-score	support
0	0.91	0.67	0.77	15
1	0.64	0.90	0.75	10
accuracy			0.76	25
macro avg	0.78	0.78	0.76	25
weighted avg	0.80	0.76	0.76	25



2.0.5 Linear Support Vector Machine: C parameter

• C is regularization parameter. The strength of the regularization is inversely proportional to C. Must be strictly positive. The penalty is a squared 12 penalty.

```
[7]: from sklearn.svm import LinearSVC

X_train, X_test, y_train, y_test = train_test_split(X, y, random_state = 0)
this_C = 1.0

fig, subaxes = plt.subplots(1, 2, figsize=(8, 4))

for this_C, subplot in zip([0.00001, 100], subaxes):
    clf = LinearSVC(C=this_C, max_iter=1000).fit(X_train, y_train)

    y_pred = clf.predict(X_test)
    result_metrics = classification_report(y_test, y_pred)
    print(result_metrics)

title = 'Linear SVC, C = {:.5f}'.format(this_C)
```

C:\Users\priya\AppData\Roaming\Python\Python312\site-packages\sklearn\svm_classes.py:31: FutureWarning: The default value of `dual` will change from `True` to `'auto'` in 1.5. Set the value of `dual` explicitly to suppress the warning.

warnings.warn(

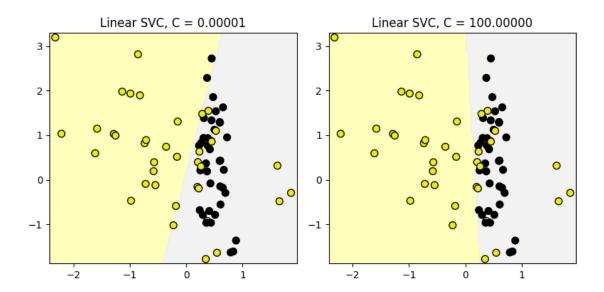
C:\Users\priya\AppData\Roaming\Python\Python312\site-packages\sklearn\svm_classes.py:31: FutureWarning: The default value of `dual` will change from `True` to `'auto'` in 1.5. Set the value of `dual` explicitly to suppress the warning.

warnings.warn(

C:\Users\priya\AppData\Roaming\Python\Python312\site-packages\sklearn\svm_base.py:1237: ConvergenceWarning: Liblinear failed to converge, increase the number of iterations.

warnings.warn(

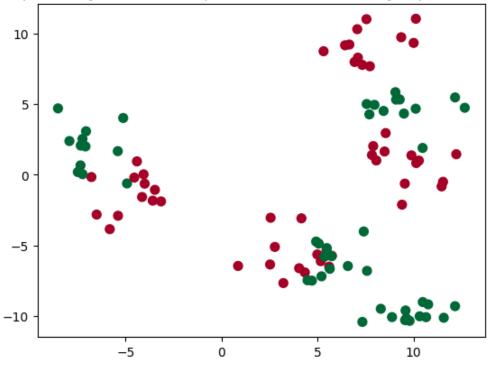
	precision	recall	f1-score	support
0	0.91	0.67	0.77	15
1	0.64	0.90	0.75	10
accuracy			0.76	25
macro avg	0.78	0.78	0.76	25
weighted avg	0.80	0.76	0.76	25
	precision	recall	f1-score	support
0	precision 0.92	recall	f1-score 0.81	support
0 1	•			••
1	0.92	0.73	0.81 0.78	15
	0.92	0.73	0.81	15 10



2.0.6 Kernelized Support Vector Machines

• More complex synthetic dataset

Sample binary classification problem with non-linearly separable classes

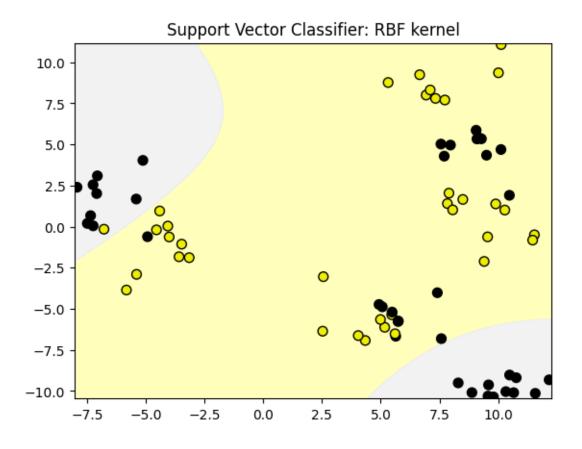


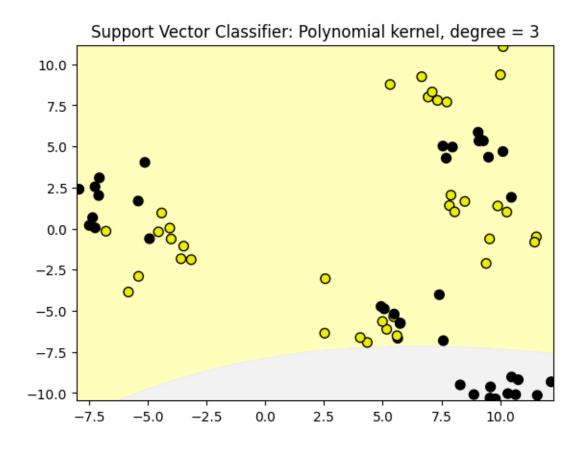
2.0.7 Classification using kernels

RBF kernel (Gaussian kernel)

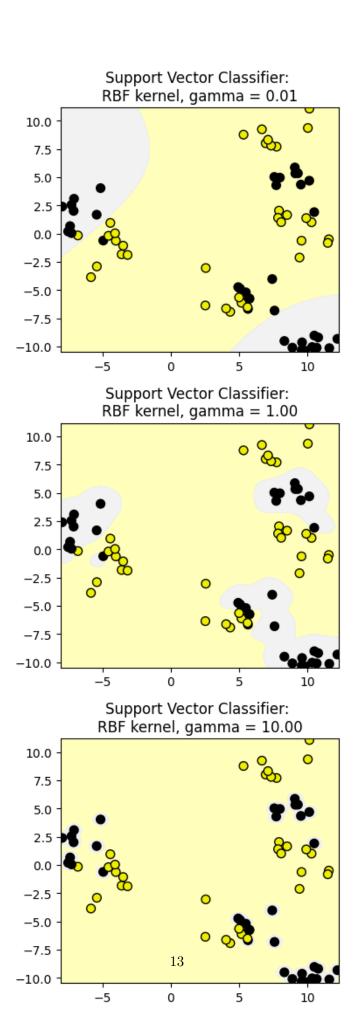
Polynomial kernel

RBF kernel (Ga	aussian) resu	1+e		
TIDI NOTHOL (GC	precision		f1-score	support
0	0.62	1.00	0.76	13
1	1.00	0.33	0.50	12
accuracy			0.68	25
macro avg	0.81	0.67	0.63	25
weighted avg	0.80	0.68	0.64	25
Polynomial ker	nel results			
	precision	recall	f1-score	support
0	0.63	0.92	0.75	13
1	0.83	0.42	0.56	12
accuracy			0.68	25
macro avg	0.73	0.67	0.65	25
weighted avg	0.73	0.68	0.66	25

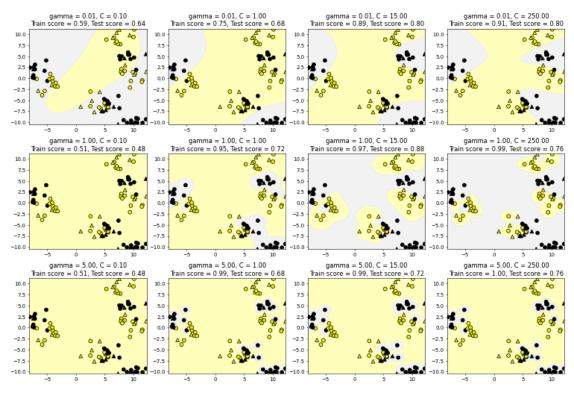




2.0.8 Support Vector Machine with RBF kernel: gamma parameter



2.0.9 Support Vector Machine with RBF kernel: using both C and gamma parameter



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Part 1:

4 Breast cancer dataset for classification

Apply SVM linear kernel (basically no kernel) and print the performance metrics.

Apply SVM RBF kernel (Gaussian kernel) and print the performance metrics with non-normalized dataset.

Apply SVM RBF kernel with normalized dataset.

Apply SVM RBF kernel using varying C and gamma parameter values. Use C= 0.1, 1, 15, 250. Use gamma= 0.01, 1, 5. Hence, 12 subplots, similar to the above example, should be drawn.

Part 2:

Write a short comparisons of SVM linear kernel and RBF kernel.

Write a short summary of how C and gamma parameters play in SVM RBF kernel.

5 PART 1

6 Breast cancer dataset for classification

```
[12]: from sklearn.datasets import load_breast_cancer

# Breast cancer dataset for classification
cancer = load_breast_cancer()
#(X_cancer, y_cancer) = load_breast_cancer(return_X_y = True)
```

```
Breast Cancer Dataset Information:
```

Number of samples: 569
Number of features: 30
Target classes: [0 1]

```
[14]: cancer
```

```
[14]: {'data': array([[1.799e+01, 1.038e+01, 1.228e+02, ..., 2.654e-01, 4.601e-01,
             1.189e-01],
             [2.057e+01, 1.777e+01, 1.329e+02, ..., 1.860e-01, 2.750e-01,
             8.902e-02],
             [1.969e+01, 2.125e+01, 1.300e+02, ..., 2.430e-01, 3.613e-01,
             8.758e-02],
             [1.660e+01, 2.808e+01, 1.083e+02, ..., 1.418e-01, 2.218e-01,
             7.820e-02],
             [2.060e+01, 2.933e+01, 1.401e+02, ..., 2.650e-01, 4.087e-01,
             1.240e-01],
             [7.760e+00, 2.454e+01, 4.792e+01, ..., 0.000e+00, 2.871e-01,
             7.039e-02]]),
      1,
             0, 0, 1, 0, 1, 1, 1, 1, 0, 0, 1, 0, 0, 1, 1, 1, 1, 0, 1, 0, 0,
             1, 1, 1, 1, 0, 1, 0, 0, 1, 0, 1, 0, 0, 1, 1, 1, 0, 0, 1, 0, 0, 0,
             1, 1, 1, 0, 1, 1, 0, 0, 1, 1, 1, 0, 0, 1, 1, 1, 1, 0, 1, 1, 0, 1,
             1, 1, 1, 1, 1, 1, 1, 0, 0, 0, 1, 0, 0, 1, 1, 1, 0, 0, 1, 0, 1, 0,
             0, 1, 0, 0, 1, 1, 0, 1, 1, 0, 1, 1, 1, 1, 0, 1, 1, 1, 1, 1, 1, 1, 1,
             1, 1, 0, 1, 1, 1, 1, 0, 0, 1, 0, 1, 1, 0, 0, 1, 1, 0, 0, 1, 1, 1,
             1, 0, 1, 1, 0, 0, 0, 1, 0, 1, 0, 1, 1, 1, 0, 1, 1, 0, 0, 1, 0, 0,
             0, 0, 1, 0, 0, 0, 1, 0, 1, 0, 1, 1, 0, 1, 0, 0, 0, 0, 1, 1, 0, 0,
             1, 1, 1, 0, 1, 1, 1, 1, 1, 0, 0, 1, 1, 0, 1, 1, 0, 0, 1, 0, 1, 1,
             1, 1, 0, 1, 1, 1, 1, 1, 0, 1, 0, 0, 0, 0, 0, 0, 0, 0, 0, 0, 0, 0,
             0, 0, 1, 1, 1, 1, 1, 0, 1, 0, 1, 1, 0, 1, 1, 0, 1, 0, 0, 1, 1,
             1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 0, 1, 1, 0, 1, 0, 1, 1, 1, 1, 1,
             1, 1, 1, 1, 1, 1, 1, 1, 0, 1, 1, 0, 1, 0, 1, 1, 1, 1, 1, 0, 0,
             0, 1, 1, 1, 1, 0, 1, 0, 1, 1, 1, 1, 0, 1, 1, 1, 1, 1, 1, 1, 0,
             0, 0, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 0, 0, 1, 0, 0, 1, 0, 0,
             1, 1, 1, 1, 1, 0, 1, 1, 1, 1, 1, 0, 1, 1, 0, 1, 1, 0, 0, 1, 1,
             1, 1, 1, 1, 0, 1, 1, 1, 1, 1, 1, 0, 1, 1, 1, 1, 1, 0, 1, 1, 0,
             1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 0, 1, 0, 0, 1, 0, 1, 1, 1, 1,
             1, 0, 1, 1, 0, 1, 0, 1, 1, 0, 1, 0, 1, 1, 1, 1, 1, 1, 1, 1, 1, 0, 0,
             1, 1, 1, 1, 1, 1, 0, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 0, 1, 1, 1, 1,
             1, 1, 1, 0, 1, 0, 1, 1, 0, 1, 1, 1, 1, 1, 0, 0, 1, 0, 1, 0, 1, 1,
             1, 1, 1, 0, 1, 1, 0, 1, 0, 1, 0, 0, 1, 1, 1, 0, 1, 1, 1, 1, 1, 1,
             1, 1, 1, 1, 1, 0, 1, 0, 0, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1,
             1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 0, 0, 0, 0, 0, 0, 1]),
      'frame': None,
      'target_names': array(['malignant', 'benign'], dtype='<U9'),
      'DESCR': '.. breast_cancer_dataset:\n\nBreast_cancer_wisconsin (diagnostic)
     dataset\n-----\n\n**Data Set
     Characteristics:**\n\n:Number of Instances: 569\n\n:Number of Attributes: 30
     numeric, predictive attributes and the class\n\n:Attribute Information:\n
     radius (mean of distances from center to points on the perimeter)\n
                                                                      - texture
```

```
(standard deviation of gray-scale values)\n - perimeter\n - area\n
smoothness (local variation in radius lengths)\n - compactness (perimeter^2 /
               - concavity (severity of concave portions of the contour)\n
- concave points (number of concave portions of the contour)\n
- fractal dimension ("coastline approximation" - 1)\n\n
                                                     The mean, standard
error, and "worst" or largest (mean of the three\n worst/largest values) of
these features were computed for each image,\n resulting in 30 features. For
instance, field 0 is Mean Radius, field\n
                                         10 is Radius SE, field 20 is Worst
Radius.\n\n
              - class:\n
                                  - WDBC-Malignant\n
Max\n=======\nradius (mean):
6.981 28.11\ntexture (mean):
                                                 9.71
                                                       39.28\nperimeter
(mean):
                          43.79 188.5\narea (mean):
143.5 2501.0\nsmoothness (mean):
                                                  0.053 0.163\ncompactness
                        0.019 \quad 0.345 \setminus \text{mean}:
0.0
      0.427\nconcave points (mean):
                                                 0.0
                                                       0.201\nsymmetry
                           0.106 0.304\nfractal dimension (mean):
(mean):
     0.097\nradius (standard error):
                                                 0.112 2.873\ntexture
(standard error):
                            0.36
                                  4.885\nperimeter (standard error):
0.757 21.98\narea (standard error):
                                                 6.802 542.2 \times 10^{-2}
                         0.002 0.031\ncompactness (standard error):
(standard error):
0.002 0.135\nconcavity (standard error):
                                                 0.0
                                                       0.396\nconcave points
(standard error):
                     0.0
                           0.053\nsymmetry (standard error):
0.079\nfractal dimension (standard error): 0.001 0.03\nradius (worst):
      36.04\ntexture (worst):
                                                 12.02 49.54\nperimeter
(worst):
                          50.41 251.2\narea (worst):
185.2 4254.0\nsmoothness (worst):
                                                 0.071 0.223 \setminus ncompactness
                        0.027 1.058\nconcavity (worst):
(worst):
      1.252\nconcave points (worst):
                                                 0.0
                                                       0.291\nsymmetry
(worst):
                           0.156 0.664\nfractal dimension (worst):
0.055 0.208\n========\n\n:Missing
Attribute Values: None\n\n:Class Distribution: 212 - Malignant, 357 -
Benign\n\n:Creator: Dr. William H. Wolberg, W. Nick Street, Olvi L.
Mangasarian\n\n:Donor: Nick Street\n\n:Date: November, 1995\n\nThis is a copy of
UCI ML Breast Cancer Wisconsin (Diagnostic)
datasets.\nhttps://goo.gl/U2Uwz2\n\nFeatures are computed from a digitized image
of a fine needle\naspirate (FNA) of a breast mass. They
describe\ncharacteristics of the cell nuclei present in the image.\n\nSeparating
plane described above was obtained using\nMultisurface Method-Tree (MSM-T) [K.
P. Bennett, "Decision Tree\nConstruction Via Linear Programming." Proceedings of
the 4th\nMidwest Artificial Intelligence and Cognitive Science Society,\npp.
97-101, 1992], a classification method which uses linear\nprogramming to
construct a decision tree. Relevant features\nwere selected using an exhaustive
search in the space of 1-4\nfeatures and 1-3 separating planes.\n\nThe actual
linear program used to obtain the separating plane\nin the 3-dimensional space
is that described in:\n[K. P. Bennett and O. L. Mangasarian: "Robust
```

```
Linear\nProgramming Discrimination of Two Linearly Inseparable
Sets",\nOptimization Methods and Software 1, 1992, 23-34].\n\nThis database is
also available through the UW CS ftp server:\n\nftp ftp.cs.wisc.edu\ncd math-
prog/cpo-dataset/machine-learn/WDBC/\n\n|details-
start|\n**References**\n|details-split|\n\n- W.N. Street, W.H. Wolberg and O.L.
Mangasarian. Nuclear feature extraction\n for breast tumor diagnosis. IS&T/SPIE
1993 International Symposium on \n Electronic Imaging: Science and Technology,
volume 1905, pages 861-870,\n San Jose, CA, 1993.\n- O.L. Mangasarian, W.N.
Street and W.H. Wolberg. Breast cancer diagnosis and \n prognosis via linear
programming. Operations Research, 43(4), pages 570-577,\n July-August 1995.\n-
W.H. Wolberg, W.N. Street, and O.L. Mangasarian. Machine learning techniques\n
to diagnose breast cancer from fine-needle aspirates. Cancer Letters 77 (1994)\n
163-171.\n\details-end\n',
 'feature_names': array(['mean radius', 'mean texture', 'mean perimeter', 'mean
area',
        'mean smoothness', 'mean compactness', 'mean concavity',
        'mean concave points', 'mean symmetry', 'mean fractal dimension',
        'radius error', 'texture error', 'perimeter error', 'area error',
        'smoothness error', 'compactness error', 'concavity error',
        'concave points error', 'symmetry error',
        'fractal dimension error', 'worst radius', 'worst texture',
        'worst perimeter', 'worst area', 'worst smoothness',
        'worst compactness', 'worst concavity', 'worst concave points',
        'worst symmetry', 'worst fractal dimension'], dtype='<U23'),
 'filename': 'breast_cancer.csv',
 'data module': 'sklearn.datasets.data'}
```

Synthetic dataset

```
[15]: import matplotlib.colors import ListedColormap
  from matplotlib.colors import ListedColormap
  from sklearn.preprocessing import StandardScaler
  from sklearn.decomposition import PCA

# Load breast cancer dataset
#cancer = load_breast_cancer()
X_cancer, y_cancer = cancer.data, cancer.target

# Standardize the features
scaler = StandardScaler()
X_cancer_scaled = scaler.fit_transform(X_cancer)

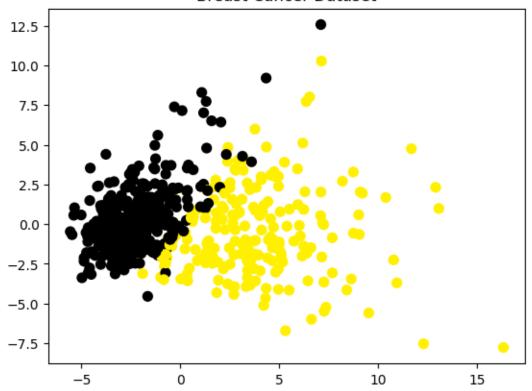
# Apply PCA for dimensionality reduction to 2 dimensions
pca = PCA(n_components=2)
X_cancer_pca = pca.fit_transform(X_cancer_scaled)

# Define a custom colormap
```

```
cmap_bold = ListedColormap(['#FFF000', '#000000'])

# Plot the breast cancer dataset
plt.figure()
plt.title('Breast Cancer Dataset')
plt.scatter(X_cancer_pca[:, 0], X_cancer_pca[:, 1], c=y_cancer, marker='o', \( \text{s} \)
$\times = 50$, cmap=cmap_bold)
plt.show()
```

Breast Cancer Dataset



7 Apply SVM Linear (No kernel) and Print the performance metrics

```
cmap_bold = ListedColormap(color_list_bold[0:numClasses])
  h = 0.03
  k = 0.5
  x_plot_adjust = 0.1
  y_plot_adjust = 0.1
  plot_symbol_size = 50
  x \min = X[:, 0].min()
  x_max = X[:, 0].max()
  y_min = X[:, 1].min()
  y_max = X[:, 1].max()
  x2, y2 = np.meshgrid(np.arange(x_min - k, x_max + k, h), np.arange(y_min -__
\hookrightarrow k, y_max + k, h))
  P = clf.predict(np.c_[x2.ravel(), y2.ravel()])
  P = P.reshape(x2.shape)
  if plot_decision_regions:
       subplot.contourf(x2, y2, P, cmap=cmap_light, alpha=0.8)
  subplot.scatter(X[:, 0], X[:, 1], c=y, cmap=cmap_bold, s=plot_symbol_size,_
⇔edgecolor='black')
  subplot.set_xlim(x_min - x_plot_adjust, x_max + x_plot_adjust)
  subplot.set_ylim(y_min - y_plot_adjust, y_max + y_plot_adjust)
  if X test is not None:
       subplot.scatter(X_test[:, 0], X_test[:, 1], c=y_test, cmap=cmap_bold,__
⇒s=plot_symbol_size, marker='^', edgecolor='black')
       train_score = clf.score(X, y)
       test_score = clf.score(X_test, y_test)
      title = title + "\nTrain score = {:.2f}, Test score = {:.2f}".
→format(train_score, test_score)
  subplot.set_title(title)
  if target_names is not None:
       legend handles = []
       for i in range(0, len(target_names)):
           patch = mpatches.Patch(color=color_list_bold[i],__
→label=target_names[i])
           legend_handles.append(patch)
       subplot.legend(loc=0, handles=legend_handles)
```

```
[17]: from sklearn.model_selection import train_test_split from sklearn.svm import SVC from sklearn.metrics import classification_report, accuracy_score
```

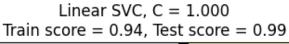
```
# Split the data into training and testing sets
X_train, X_test, y_train, y_test = train_test_split(X_cancer_pca, y_cancer,_u

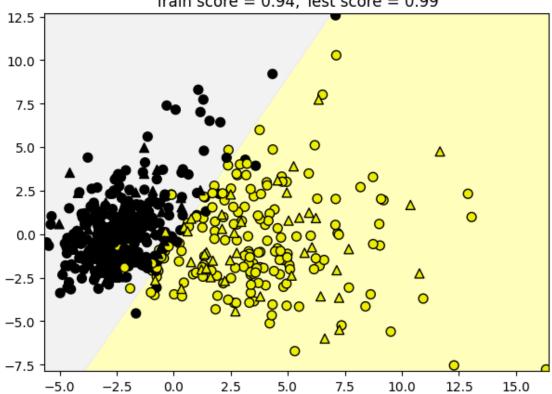
→test_size=0.2, random_state=42)
# Create SVM classifier with linear kernel
svm_linear = SVC(kernel='linear',C=1.0)
# Train the classifier
svm_linear.fit(X_train, y_train)
# Predict on the test set
y_pred = svm_linear.predict(X_test)
# Print performance metrics
print("Accuracy:", accuracy_score(y_test, y_pred))
print("\nClassification Report:\n", classification_report(y_test, y_pred))
fig, subaxes = plt.subplots(1, 1, figsize=(7, 5))
title = 'Linear SVC, C = {:.3f}'.format(1.0)
plot_for_classifier_subplot(svm_linear, X_train, y_train, X_test, y_test,_u
 →title, subaxes)
```

Accuracy: 0.9912280701754386

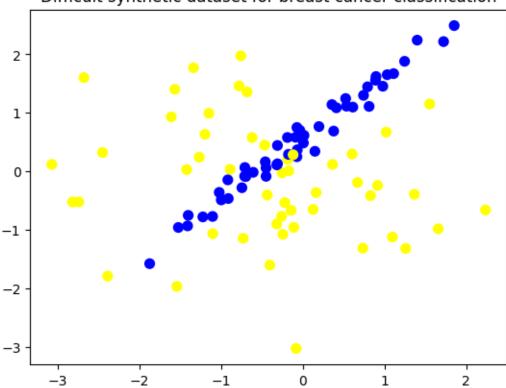
Classification Report:

	precision	recall	f1-score	support
0	1.00	0.98	0.99	43
1	0.99	1.00	0.99	71
accuracy			0.99	114
macro avg	0.99	0.99	0.99	114
weighted avg	0.99	0.99	0.99	114





Difficult synthetic dataset for breast cancer classification

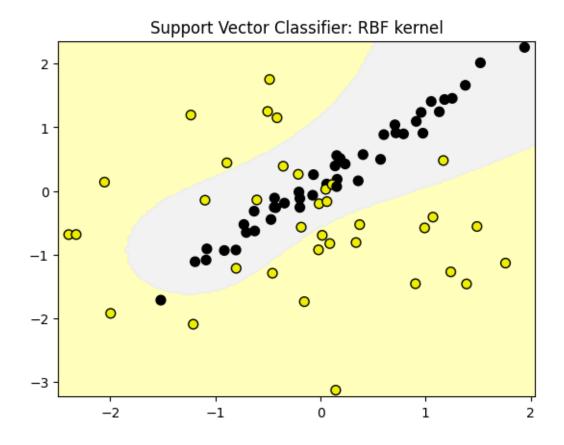


Apply SVM RBF kernel (Gaussian kernel) and print the performance metrics with non-normalized dataset.

Accuracy: 0.8

Classification Report:

	precision	recall	f1-score	support
0	1.00	0.71	0.83	14
1	0.60	1.00	0.75	6
accuracy			0.80	20
macro avg	0.80	0.86	0.79	20
weighted avg	0.88	0.80	0.81	20



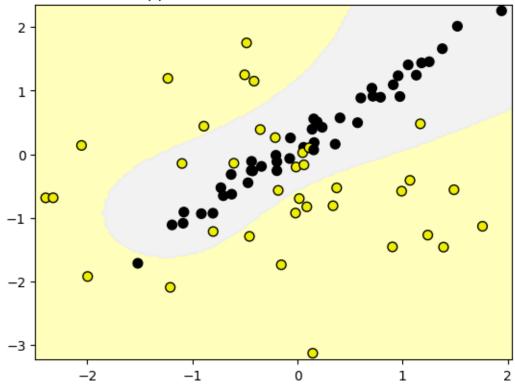
8 Apply SVM RBF kernel with normalized dataset.

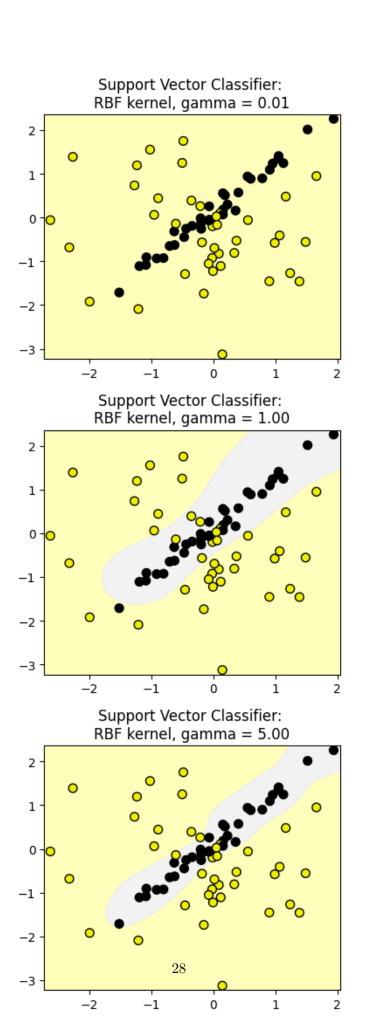
Accuracy: 0.8

Classification Report:

	precision	recall	f1-score	support
0	1.00	0.71	0.83	14
1	0.60	1.00	0.75	6
accuracy			0.80	20
macro avg	0.80	0.86	0.79	20
weighted avg	0.88	0.80	0.81	20

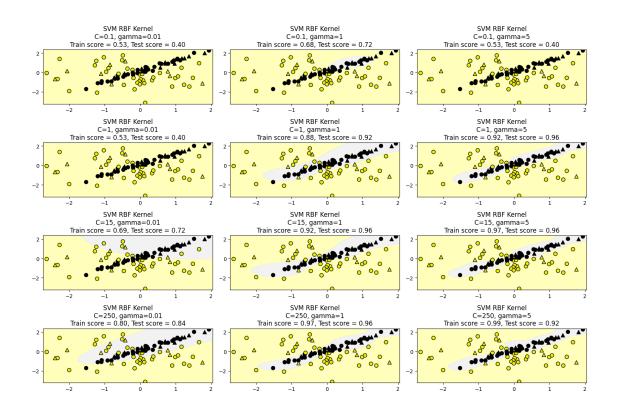
Support Vector Classifier: RBF kernel





Apply SVM RBF kernel using varying C and gamma parameter values. Use C= 0.1, 1, 15, 250. Use gamma= 0.01, 1, 5. Hence, 12 subplots, similar to the above example, should be drawn.

```
[22]: from sklearn.svm import SVC
      import matplotlib.pyplot as plt
      # Define the range of C and gamma values
      C_{values} = [0.1, 1, 15, 250]
      gamma_values = [0.01, 1, 5]
      # Create subplots
      fig, subaxes = plt.subplots(len(C_values), len(gamma_values), figsize=(15, 10))
      # Iterate over each combination of C and gamma values
      for i, C in enumerate(C_values):
          for j, gamma in enumerate(gamma_values):
              # Train the SVM classifier with RBF kernel
              clf = SVC(kernel='rbf', C=C, gamma=gamma).fit(X_train, y_train)
              # Generate title for the subplot
              title = f'SVM RBF Kernel\nC={C}, gamma={gamma}'
              # Plot decision regions for the classifier
              plot_class_regions_for_classifier_subplot(clf, X_train, y_train,
                                                        X_test, y_test, title,
                                                         subaxes[i, j])
      # Adjust layout
      plt.tight_layout()
      # Show the plots
      plt.show()
```



9 PART 2

9.1 Write a short comparisons of SVM linear kernel and RBF kernel.

Linear Kernel: - The linear kernel computes a decision boundary as a straight line, making it suitable for datasets with linear separability. - It is optimal for scenarios where classes can be cleanly separated by a single hyperplane. - Due to its simplicity, the linear kernel is computationally efficient, making it preferable for large-scale datasets. - It performs well when the number of features is high relative to the number of samples.

RBF (Radial Basis Function) Kernel: - The RBF kernel computes a non-linear decision boundary, allowing for the capture of complex relationships between features and target classes. - It is effective for datasets with non-linear separability, where classes are not easily separated by a linear boundary. - The RBF kernel's flexibility comes with two important hyperparameters: C and gamma. C balances margin maximization and classification error minimization, while gamma controls the influence of individual training examples. - It is capable of handling intricate datasets by capturing fine-grained details in the data.

The choice between linear and RBF kernels depends on the dataset's nature and the desired complexity of the decision boundary. Linear kernel is straightforward and computationally efficient, suitable for simpler datasets with linear separability. In contrast, the RBF kernel offers more flexibility and can handle complex relationships, making it preferable for datasets with non-linear separability.

9.2 Write a short summary of how C and gamma parameters play in SVM RBF kernel.

The C parameter in SVM with RBF kernel plays a pivotal role in controlling the trade-off between achieving a smooth decision boundary and accurately classifying training data points. A smaller value of C leads to a softer margin, allowing more misclassifications but potentially increasing the model's generalization ability. A larger value of C results in a narrower margin, aiming to correctly classify more training samples, which may lead to overfitting if set too high. Tuning C allows balancing between bias and variance in the model, with smaller C values promoting a more flexible decision boundary and larger C values promoting a more strict boundary.

In the context of the breast cancer dataset, smaller C values allow for a more flexible decision boundary, potentially capturing subtle patterns in the data. However, this flexibility might lead to misclassification of some training examples, particularly outliers or noisy data points. Conversely, larger C values impose a stricter penalty for misclassification, resulting in a decision boundary that aims to correctly classify as many training examples as possible. However, this could lead to a narrower margin between classes and potential overfitting, especially if the dataset contains noise or outliers.

On the other hand, the gamma parameter determines the influence of individual training examples on the shape of the decision boundary. In the context of the breast cancer dataset, smaller gamma values result in a smoother decision boundary, which could be advantageous for datasets with a relatively simple structure or when overfitting is a concern. Conversely, larger gamma values lead to a more complex decision boundary that closely fits the training data, potentially resulting in overfitting, particularly if the dataset is noisy or contains outliers. A smaller value of gamma results in a smoother decision boundary, with points further away from the decision boundary having less influence. Tuning gamma allows controlling the model's complexity, with smaller gamma values promoting smoother decision boundaries and larger gamma values promoting more complex boundaries.

Overall, effectively tuning the C and gamma parameters in SVM with RBF kernel for the breast cancer dataset is crucial for achieving a balance between model complexity and generalization ability. This often requires careful experimentation and validation, using techniques such as cross-validation, to find the optimal values that yield the best classification performance and robustness on unseen data.

10 Part 3

Reflection

Working with Support Vector Machines (SVMs) offered valuable insights into the fundamental concepts of machine learning algorithms, particularly the capacity of SVMs to discern optimal decision boundaries. Exploring hyperparameters such as C and gamma underscored the significance of hyperparameter tuning in machine learning, demonstrating how different parameter configurations influence model performance and generalization capabilities.

Visualizing decision boundaries provided a tangible understanding of how hyperparameters impact model behavior, offering a visual representation of their effects on the model's decision-making process. However, fine-tuning hyperparameters like C and gamma posed challenges, requiring iterative experimentation and validation, especially for larger datasets, which can be time-consuming and computationally demanding.

Assessing model performance and interpreting the results presented essential yet intricate tasks. It involved evaluating multiple performance metrics, comprehending their implications, and making informed decisions aligned with the specific problem domain. This process underscored the importance of continual learning and exploration in machine learning, highlighting the necessity of staying abreast of new techniques, tools, and best practices to effectively address complex modeling tasks.

Conclusion

The experimentation with different C and gamma values in SVM models employing the RBF kernel has yielded insightful conclusions. Firstly, the C parameter significantly influences the decision boundary's smoothness and margin width. Smaller C values yield smoother decision boundaries with wider margins, fostering higher bias and lower variance. Conversely, larger C values result in tighter boundaries, aiming to accurately classify more training examples, potentially enhancing performance but increasing the risk of overfitting.

Secondly, the gamma parameter plays a pivotal role in shaping the decision boundary's complexity. Smaller gamma values produce smoother boundaries with reduced influence from individual data points, suitable for simpler datasets or scenarios with numerous samples. On the other hand, larger gamma values lead to more intricate boundaries closely aligned with the training data, beneficial for capturing intricate patterns but posing a higher risk of overfitting.

The interplay between C and gamma underscores the delicate balance between bias and variance, necessitating careful parameter tuning. Achieving optimal performance often entails iterative experimentation and validation, leveraging techniques like grid search or randomized search to explore various parameter combinations effectively.

In summary, the choice of C and gamma values profoundly impacts the SVM model's complexity, bias-variance trade-off, and generalization capability. Understanding their interaction and discerning their effects on the decision boundary are paramount for developing robust classifiers across diverse machine learning tasks.