MACHINE LEARNING LABORATORY

ASSIGNMENT 1

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Github: github.com/atmikgoswami/ML-Lab

DATASETS:

1. Iris Plants Dataset

• Features: Sepal Length, Sepal Width, Petal Length, Petal Width

• Classes: Setosa, Versicolor, Virginica

• Total Samples: 150

Sample Data:

	sepal length (cm)	sepal width (cm)	petal length (cm)	petal width (cm)	species
0	5.1	3.5	1.4	0.2	setosa
1	4.9	3.0	1.4	0.2	setosa
2	4.7	3.2	1.3	0.2	setosa
3	4.6	3.1	1.5	0.2	setosa
4	5.0	3.6	1.4	0.2	setosa

2. Breast Cancer Wisconsin (Diagnostic) Dataset

• Features: 30 numeric features (e.g., radius, texture, area, etc.)

• Classes: Malignant (M), Benign (B)

• Total Samples: 569

Sample Data:

	mean radius	mean texture	mean perimeter	mean area	mean smoothness	mean compactness	mean concavity	mean concave points	mean symmetry	mean fractal dimension	 worst perimeter	worst area	worst smoothness	worst compactness	worst concavity	worst concave points	w symm
0	17.99	10.38	122.80	1001.0	0.11840	0.27760	0.30010	0.14710	0.2419	0.07871	184.60	2019.0	0.16220	0.66560	0.7119	0.2654	0.
1	20.57	17.77	132.90	1326.0	0.08474	0.07864	0.08690	0.07017	0.1812	0.05667	 158.80	1956.0	0.12380	0.18660	0.2416	0.1860	0.:
2	19.69	21.25	130.00	1203.0	0.10960	0.15990	0.19740	0.12790	0.2069	0.05999	152.50	1709.0	0.14440	0.42450	0.4504	0.2430	0.:
3	11.42	20.38	77.58	386.1	0.14250	0.28390	0.24140	0.10520	0.2597	0.09744	 98.87	567.7	0.20980	0.86630	0.6869	0.2575	0.0
4	20.29	14.34	135.10	1297.0	0.10030	0.13280	0.19800	0.10430	0.1809	0.05883	152.20	1575.0	0.13740	0.20500	0.4000	0.1625	0.:
564	21.56	22.39	142.00	1479.0	0.11100	0.11590	0.24390	0.13890	0.1726	0.05623	166.10	2027.0	0.14100	0.21130	0.4107	0.2216	0.:
565	20.13	28.25	131.20	1261.0	0.09780	0.10340	0.14400	0.09791	0.1752	0.05533	 155.00	1731.0	0.11660	0.19220	0.3215	0.1628	0.:
566	16.60	28.08	108.30	858.1	0.08455	0.10230	0.09251	0.05302	0.1590	0.05648	126.70	1124.0	0.11390	0.30940	0.3403	0.1418	0.:
567	20.60	29.33	140.10	1265.0	0.11780	0.27700	0.35140	0.15200	0.2397	0.07016	 184.60	1821.0	0.16500	0.86810	0.9387	0.2650	0.
568	7.76	24.54	47.92	181.0	0.05263	0.04362	0.00000	0.00000	0.1587	0.05884	59.16	268.6	0.08996	0.06444	0.0000	0.0000	0.:
569 rd	ws × 32 c	columns															

1. Employ Naive Bayes (Gaussian, Multinomial & Bernoulli) classifier and show classification results (Accuracy, Precision, Recall, F-score, confusion matrix).

Code:

```
import pandas as pd
import seaborn as sns
import matplotlib.pyplot as plt
from sklearn.model selection import train test split
from sklearn.metrics import classification report, confusion matrix
from sklearn.datasets import load iris
iris = load iris()
df = pd.DataFrame(data=iris.data, columns=iris.feature_names)
df['species'] = iris.target
df['species'] = df['species'].map({0: 'setosa', 1: 'versicolor', 2:
'virginica'})
X = df.drop(['species'], axis=1)
y = df['species']
X_train, X_test, y_train, y_test = train_test_split(X, y,
test size=0.2)
from sklearn.naive bayes import MultinomialNB
classifier = MultinomialNB(alpha=1.5, fit prior=False)
classifier.fit(X train, y train)
y pred = classifier.predict(X test)
print("Confusion Matrix")
print(confusion matrix(y test, y pred))
print("-----
print("Classification Report")
print(classification_report(y_test, y_pred))
from sklearn.naive bayes import GaussianNB
```

```
classifier = GaussianNB()
classifier.fit(X_train, y_train)
y pred = classifier.predict(X test)
print("Confusion Matrix")
print(confusion_matrix(y_test, y_pred))
print("-----")
print("Classification Report")
print(classification_report(y_test, y_pred))
from sklearn.naive_bayes import BernoulliNB
classifier = BernoulliNB(binarize=1.5)
classifier.fit(X train, y train)
y pred = classifier.predict(X test)
print("Confusion Matrix")
print(confusion_matrix(y_test, y_pred))
print("-----
print("Classification Report")
print(classification report(y test, y pred))
from sklearn.datasets import load breast cancer
data = load breast cancer()
df = pd.DataFrame(data.data, columns=data.feature_names)
df['target'] = data.target
df['target_label'] = df['target'].map({0: 'malignant', 1: 'benign'})
X = df.drop(['target', 'target_label'], axis=1)
y = df['target']
X train, X test, y train, y test = train test split(X, y,
test size=0.2)
from sklearn.naive bayes import MultinomialNB
classifier = MultinomialNB()
```

```
classifier.fit(X_train, y_train)
y pred = classifier.predict(X test)
print("Confusion Matrix")
print(confusion matrix(y test, y pred))
print("----")
print("Classification Report")
print(classification_report(y_test, y_pred))
from sklearn.naive_bayes import GaussianNB
classifier = GaussianNB(var smoothing=1e-09)
classifier.fit(X_train, y_train)
y pred = classifier.predict(X test)
print("Confusion Matrix")
print(confusion_matrix(y_test, y_pred))
print("----")
print("Classification Report")
print(classification_report(y_test, y_pred))
from sklearn.naive bayes import BernoulliNB
classifier = BernoulliNB(binarize=1.5, fit prior=False,
class prior=[0.8, 0.2])
classifier.fit(X train, y train)
y pred = classifier.predict(X test)
print("Confusion Matrix")
print(confusion_matrix(y_test, y_pred))
print("----")
print("Classification Report")
print(classification_report(y_test, y_pred))
```

Results and Discussion:

Iris Dataset:

Multinomial Naive Bayes:

Confusion Matrix [[7 0 0] [0 11 0] [0 1 11]]

-----Classification Report precision recall f1-score support

 setosa
 1.00
 1.00
 1.00
 7

 versicolor
 0.92
 1.00
 0.96
 11

 virginica
 1.00
 0.92
 0.96
 12

 accuracy
 0.97
 30

 macro avg
 0.97
 0.97
 0.97
 30

 weighted avg
 0.97
 0.97
 0.97
 30

Gaussian Naive Bayes:

Confusion Matrix [[11 0 0] [0 9 0] [0 1 9]]

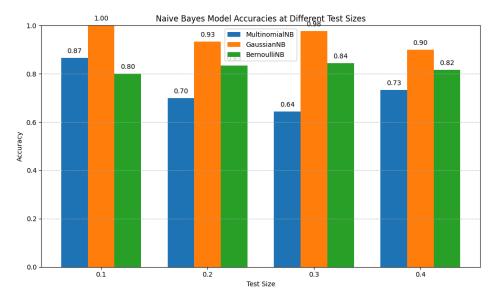
Classificatio	n Report			
	precision	recall	f1-score	support
setosa	1.00	1.00	1.00	11
versicolor	0.90	1.00	0.95	9
virginica	1.00	0.90	0.95	10
accuracy			0.97	30
macro avg	0.97	0.97	0.96	30
weighted avg	0.97	0.97	0.97	30

Bernoulli Naive Bayes:

Confusion Matrix [[9 2 0] [0 9 0] [0 0 10]]

Classificatio	n Report						
	precision	recall	f1-score	support			
	4 00						
setosa	1.00	0.82	0.90	11			
versicolor	0.82	1.00	0.90	9			
virginica	1.00	1.00	1.00	10			
accuracy			0.93	30			
macro avg	0.94	0.94	0.93	30			
weighted avg	0.95	0.93	0.93	30			

Comparison of accuracies for different test sizes:



Wisconsin Breast Cancer Dataset

Multinomial Naive Bayes:

Confusion Matrix [[27 7] [1 79]]

Classification Report

Classificati	ion Report			
	precision	recall	f1-score	support
(0.96	0.79	0.87	34
1	0.92	0.99	0.95	80
accuracy	/		0.93	114
macro av	g 0.94	0.89	0.91	114
weighted av	g 0.93	0.93	0.93	114

Gaussian Naive Bayes:

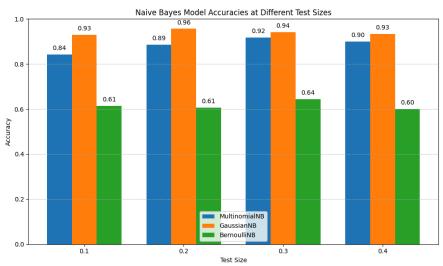
Confusion Matrix [[31 3] [0 80]]

[0 00]]								
Classification Report precision recall f1-score support								
	precision	recarr	11-2001.6	Support				
0	1.00	0.91	0.95	34				
1	0.96	1.00	0.98	80				
accuracy			0.97	114				
macro avg	0.98	0.96	0.97	114				
weighted avg	0.97	0.97	0.97	114				

Bernoulli Naive Bayes:

Classific	atio	n Report precision	recall	f1-score	support
	0 1	1.00 0.69	0.35 1.00	0.52 0.82	46 68
accur macro weighted	avg	0.85 0.82	0.67 0.74	0.74 0.67 0.70	114 114 114

Comparison of accuracies for different test sizes:



Discussion:

Naive Bayes classifiers assume feature independence and use probability distributions to classify data. For the **Iris dataset**, Gaussian Naive Bayes generally performed best, as the features (lengths and widths) are continuous and approximately normally distributed. Multinomial Naive Bayes was less effective since it is designed for count-based features, and Bernoulli Naive Bayes was the least suitable because binarization of continuous features led to information loss. This highlights how model assumptions directly impact classification performance. Accuracy and F-scores were higher for Gaussian NB, showing that modeling continuous distributions is appropriate for this dataset.

For the **Breast Cancer dataset**, Gaussian Naive Bayes again outperformed the other variants due to the continuous, real-valued features like radius, texture, and area. Multinomial Naive Bayes, while usable, is not ideal for such data and gave comparatively lower results. Bernoulli Naive Bayes struggled most because binarization discards crucial variance across 30 features. Overall, Gaussian NB provided robust results with good precision and recall for both malignant and benign classes. The confusion matrices showed very few misclassifications, reinforcing its effectiveness.

Thus, the key observation is that Gaussian NB consistently works well for continuous feature datasets, while Multinomial and Bernoulli are more suited for text or count data.

2. Use Decision Tree classifier for all the two datasets and show classification results (Accuracy, Precision, Recall, F-score, confusion matrix). Generate the decision tree images for all cases highlighting information like Gini and Entropy.

Code:

```
import pandas as pd
import seaborn as sns
import matplotlib.pyplot as plt
from sklearn.model selection import train test split
from sklearn.metrics import classification report, confusion matrix
from sklearn.tree import DecisionTreeClassifier
from sklearn.model selection import GridSearchCV
from sklearn.datasets import load iris
iris = load iris()
df = pd.DataFrame(data=iris.data, columns=iris.feature names)
df['species'] = iris.target
df['species'] = df['species'].map({0: 'setosa', 1: 'versicolor', 2:
'virginica'})
X = df.drop(['species'], axis=1)
y = df['species']
X_train, X_test, y_train, y_test = train_test_split(X, y,
test size=0.2, stratify=y)
classifier = DecisionTreeClassifier(random state=1)
param grid = {
    'criterion': ['gini', 'entropy'],
    'splitter': ['best', 'random'],
    'max depth': [2, 3, 4, 5, 6],
    'min samples split': [2, 3, 4, 5],
    'min samples leaf': [1, 2, 3, 4, 5],
    'max features': [None, 'sqrt', 'log2']
# Grid search
```

```
grid = GridSearchCV(classifier, param grid, cv=5, scoring='accuracy')
grid.fit(X train, y train)
# Best model
best model = grid.best estimator
print("Best Parameters:", grid.best params )
# Evaluate on test set
y pred = best model.predict(X test)
print("\nConfusion Matrix")
print(confusion_matrix(y_test, y_pred))
print("-----
print("Classification Report")
print(classification_report(y_test, y_pred,
target names=iris.target names))
# Plotting of decision tree
from IPython.display import Image
from sklearn.tree import export graphviz
!pip install pydotplus
import pydotplus
features = X.columns
dot data = export graphviz(best model, out file=None,
feature names=iris.feature names)
graph = pydotplus.graph_from_dot data(dot data)
Image(graph.create png())
from sklearn.datasets import load breast cancer
data = load breast cancer()
df = pd.DataFrame(data.data, columns=data.feature names)
df['target'] = data.target
df['target label'] = df['target'].map({0: 'malignant', 1: 'benign'})
X = df.drop(['target', 'target label'], axis=1)
y = df['target label']
```

```
X_train, X_test, y_train, y_test = train_test_split(X, y,
test size=0.2, stratify=y)
classifier = DecisionTreeClassifier(random state=1)
param grid = {
    'criterion': ['gini', 'entropy'],
   'splitter': ['best', 'random'],
   'max depth': [2, 3, 4, 5, 6],
   'min_samples_split': [2, 3, 4, 5],
   'min samples leaf': [1, 2, 3, 4, 5],
   'max_features': [None, 'sqrt', 'log2']
# Grid search
grid = GridSearchCV(classifier, param grid, cv=5, scoring='accuracy')
grid.fit(X train, y train)
# Best model
best model = grid.best estimator
print("Best Parameters:", grid.best params )
# Evaluate on test set
y pred = best model.predict(X test)
print("\nConfusion Matrix")
print(confusion matrix(y test, y pred))
print("----")
print("Classification Report")
print(classification report(y test, y pred,
target names=data.target names))
# Plotting of decision tree
from IPython.display import Image
from sklearn.tree import export graphviz
!pip install pydotplus
import pydotplus
features = X.columns
dot data = export graphviz(best model, out file=None,
feature names=data.feature names)
```

```
graph = pydotplus.graph_from_dot_data(dot_data)
Image(graph.create_png())
```

Results and Discussion:

Iris Dataset:

Confusion Matrix [[10 0 0] [0 10 0] [0 0 10]]

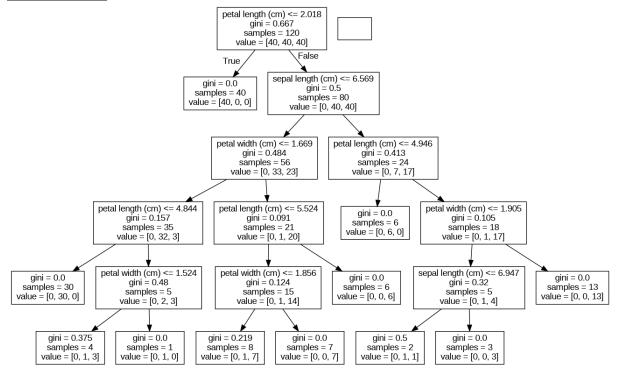
Classification Report

nocall fi scone support

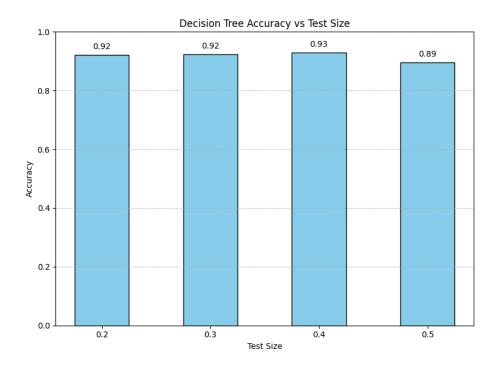
Classification	n keport
	precisio

	precision	recarr	T1-Score	support
setosa	1.00	1.00	1.00	10
versicolor	1.00	1.00	1.00	10
virginica	1.00	1.00	1.00	10
accuracy			1.00	30
macro avg	1.00	1.00	1.00	30
weighted avg	1.00	1.00	1.00	30

Decision Tree:



Comparison for accuracies for different test sizes:



Wisconsin Breast Cancer Dataset

Confusion Matrix

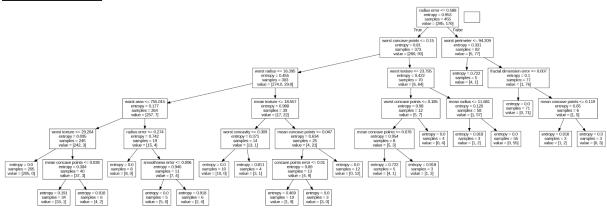
[[72 0]

[3 39]]

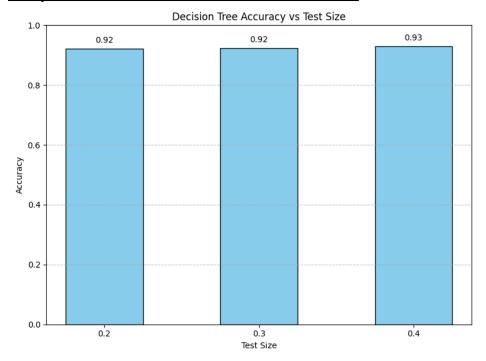
Classification Report

	precision	recall	f1-score	support
malignant benign	0.96 1.00	1.00 0.93	0.98 0.96	72 42
J	1.00	0.93		
accuracy macro avg	0.98	0.96	0.97 0.97	114 114
weighted avg	0.97	0.97	0.97	114

Decision Tree:



Comparison for accuracies for different test sizes:



Discussion:

Decision Trees learn feature-based rules and partition the feature space using criteria like Gini index or Entropy. For the Iris dataset, the Decision Tree achieved very high classification accuracy, often close to perfect, because the dataset is small, well-balanced, and highly separable (e.g., petal length and width easily distinguish Setosa). The visualization of the tree made the classification process interpretable, showing clear splits based on petal dimensions. Overfitting was controlled using hyperparameters like max_depth and min_samples_split.

For the Breast Cancer dataset, the Decision Tree also performed well, with strong precision and recall, especially for the benign class. However, because this dataset has higher dimensionality (30 features), trees can easily overfit. Grid search tuning helped in selecting optimal depth and splitting criteria to balance accuracy with generalization. The resulting trees highlighted key medical features (such as mean radius and texture) that strongly influence the classification.

In comparison with Naive Bayes, Decision Trees provided better interpretability and comparable or higher accuracy, especially when tuned. However, they are more prone to overfitting, while Naive Bayes is simpler and more robust for small datasets.