## Week 13 Project Deliverable

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## **Problem Statement**

One of the challenge for all Pharmaceutical companies is to understand the persistency of drug as per the physician prescription. To solve this problem ABC pharma company approached an analytics company to automate this process of identification.

The main objective is to build a classification model to predict the NTM drug persistence of patients based on several factors.

# Project Schedule

Week	Date	Goal			
07	06/19/2023	Problem Statement, Data Preview			
08	06/26/2023	Data Preprocessing			
09	07/02/2023	Data Prepare and cleaning			
10	07/09/2023	EDA			
11	07/16/2023	Recommendation and Model			
		Suggestion			
12	07/23/2023	Model Building and Evaluation			

13	07/30/2023	Presentation

# Data Understanding

## **Predictor Variables**

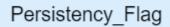
### Demographic features

Age	Age of the patient during their therapy
Race	Race of the patient from the patient table
Region	Region of the patient from the patient table
Ethnicity	Ethnicity of the patient from the patient table
Gender	Gender of the patient from the patient table



IDN Indicator	Flag indicating patients mapped to IDN		
NTM - Physician Specialty	Specialty of the HCP that prescribed the NTM $\ensuremath{Rx}$		
NTM - T-Score	T Score of the patient at the time of the NTM $\dots$		
Change in T Score	Change in Tscore before starting with any ther		
NTM - Risk Segment	Risk Segment of the patient at the time of the		
Change in Risk Segment	Change in Risk Segment before starting with an		
NTM - Multiple Risk Factors	Flag indicating if patient falls under multip		
NTM - Dexa Scan Frequency	Number of DEXA scans taken prior to the first		
NTM - Dexa Scan Recency	Flag indicating the presence of Dexa Scan befo		
Dexa During Therapy	Flag indicating if the patient had a Dexa Scan		
NTM - Fragility Fracture Recency	Flag indicating if the patient had a recent fr		
Fragility Fracture During Therapy	Flag indicating if the patient had fragility f		
NTM - Glucocorticoid Recency	Flag indicating usage of Glucocorticoids (>=7		
Glucocorticoid Usage During Therapy	Flag indicating if the patient had a Glucocort		
NTM - Injectable Experience	Flag indicating any injectable drug usage in t		
NTM - Risk Factors	Risk Factors that the patient is falling into		
NTM - Comorbidity	Comorbidities are divided into two main catego		
NTM - Concomitancy	Concomitant drugs recorded prior to starting w		
Adherence	Adherence for the therapies		

## Target Variable:



Flag indicating if a patient was persistent or...

Boolean Predictor Variable List: (All the variables in the list have two values of Yes or No)

```
['Gluco Record Prior Ntm',
 'Gluco_Record_During_Rx',
 'Dexa_During_Rx',
 'Frag_Frac_Prior_Ntm',
 'Frag_Frac_During_Rx',
 'Idn_Indicator',
'Injectable_Experience_During_Rx',
 'Comorb_Encounter_For_Screening_For_Malignant_Neoplasms',
 'Comorb_Encounter_For_Immunization',
 'Comorb_Encntr_For_General_Exam_W_O_Complaint,_Susp_Or_Reprtd_Dx',
 'Comorb_Vitamin_D_Deficiency',
 'Comorb Other Joint Disorder Not Elsewhere Classified',
 'Comorb_Encntr_For_Oth_Sp_Exam_W_O_Complaint_Suspected_Or_Reprtd_Dx',
 'Comorb_Long_Term_Current_Drug_Therapy',
 'Comorb_Dorsalgia',
 'Comorb_Personal_History_Of_Other_Diseases_And_Conditions',
 'Comorb_Other_Disorders_Of_Bone_Density_And_Structure',
 'Comorb_Disorders_of_lipoprotein_metabolism_and_other_lipidemias',
 'Comorb_Osteoporosis_without_current_pathological_fracture',
 'Comorb_Personal_history_of_malignant_neoplasm',
 'Comorb Gastro esophageal reflux disease'.
 'Concom_Cholesterol_And_Triglyceride_Regulating_Preparations',
 'Concom_Narcotics',
 'Concom Systemic Corticosteroids Plain'
 'Concom_Anti_Depressants_And_Mood_Stabilisers',
 'Concom_Fluoroquinolones',
 'Concom Cephalosporins',
 'Concom_Macrolides_And_Similar_Types',
 'Concom_Broad_Spectrum_Penicillins',
 'Concom Anaesthetics General',
 'Concom_Viral_Vaccines',
 'Risk_Type_1_Insulin_Dependent_Diabetes',
 'Risk Osteogenesis Imperfecta',
 'Risk_Rheumatoid_Arthritis',
 'Risk_Untreated_Chronic_Hyperthyroidism',
 'Risk Untreated Chronic Hypogonadism',
 'Risk_Untreated_Early_Menopause',
 'Risk_Patient_Parent_Fractured_Their_Hip',
 'Risk_Smoking_Tobacco',
 'Risk_Chronic_Malnutrition_Or_Malabsorption',
 'Risk_Chronic_Liver_Disease',
 'Risk Family History Of Osteoporosis',
 'Risk_Low_Calcium_Intake',
 'Risk_Vitamin_D_Insufficiency',
 'Risk_Poor_Health_Frailty',
 'Risk_Excessive_Thinness',
 'Risk_Hysterectomy_Oophorectomy',
 'Risk_Estrogen_Deficiency',
 'Risk Immobilization'.
 'Risk_Recurring_Falls']
```

All these data will be dummy coded with 0 and 1 values for future classification model

Columns with quantitative data:

	Dexa_Freq_During_Rx	Count_Of_Risks
0	0	0
1	0	0
2	0	2
3	0	1
4	0	1
3419	0	1
3420	0	0
3421	7	1
3422	0	0
3423	0	1

	Dexa_Freq_During_Rx	Count_Of_Risks
count	3424.000000	3424.000000
mean	3.016063	1.239486
std	8.136545	1.094914
min	0.000000	0.000000
25%	0.000000	0.000000
50%	0.000000	1.000000
75%	3.000000	2.000000
max	146.000000	7.000000

No outliers and missing values exist in the dataset

Data Cleaning and Transformation

Since there is no missing data in the dataset, the only values that need to be cleaned was features that contains 'Unknown'

Values:

```
# impute unknown value
df.loc[df['Change_Risk_Segment'] == 'Unknown', 'Change_Risk_Segment'] = 'No change'
df.loc[df['Change_T_Score'] == 'Unknown', 'Change_T_Score'] = 'No change'
```

For unknown changes, I grouped them all into unchanged group which is the majority of the variable

```
def transform_speciality(value):
    transform medical speciality
    if 'MEDICINE' in value.split(' '):
        return 'MEDICINE'
    elif 'SURGERY' in value.split(' '):
        return 'SURGERY'
    elif df['Ntm_Speciality'].value_counts()[value] < 10 or value == 'Unknown':
        return 'OTHER'
    return value</pre>
```

Cleaned and transform medical specialty to decrease the variable numbers.

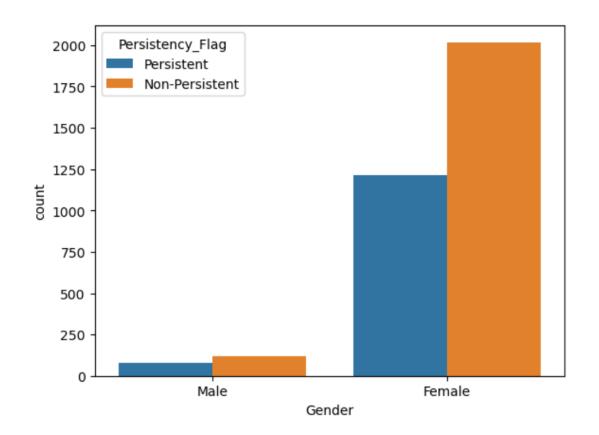
Use label encoding and one hot encoding to transform different types of categorical features

	Ethnicity_Hispanic	Ethnicity_Not Hispanic	Ethnicity_Unknown	Age_Bucket_55- 65	Age_Bucket_65- 75	Age_Bucket_<55	Age_Bucket_>75	Ntm_Speciality_Bucket_Endo/Onc/U
0	0.0	1.0	0.0	0.0	0.0	0.0	1.0	(
1	0.0	1.0	0.0	1.0	0.0	0.0	0.0	(
2	1.0	0.0	0.0	0.0	1.0	0.0	0.0	(
3	0.0	1.0	0.0	0.0	0.0	0.0	1.0	(
4	0.0	1.0	0.0	0.0	0.0	0.0	1.0	(

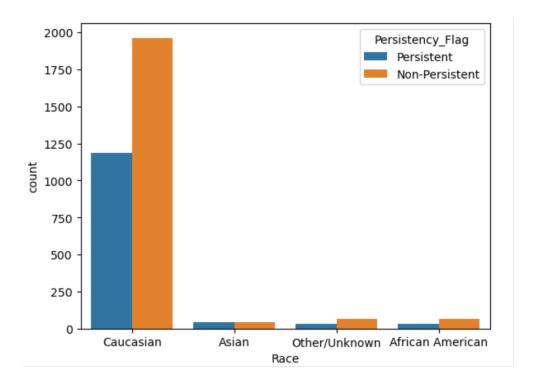
Code target variables

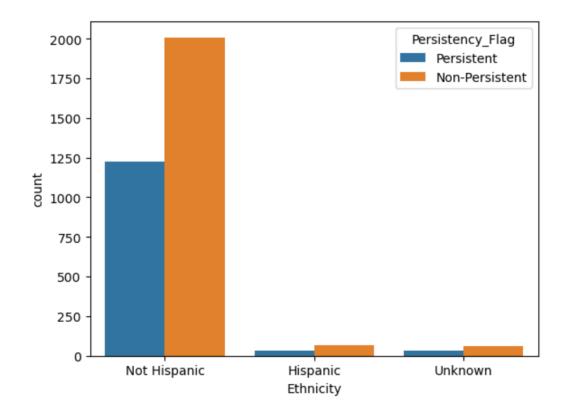
```
# label encode other features
for col in bool_cols:
   label(df, col)
# code target variables
df['target'] = np.where(df['Persistency_Flag'] == 'Persistent', 1, 0)
df['target']
       1
2
4
3419
      1
3420
3421
      1
3422
3423
Name: target, Length: 3424, dtype: int32
```

# Exploratory Data Analysis and Recommendations



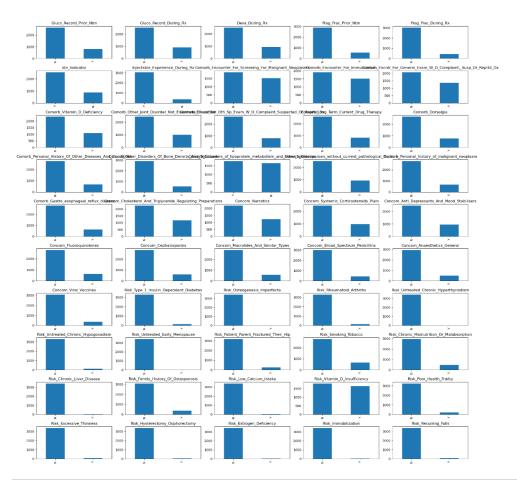
There is a huge distinction between the number of Male and Female patients, we will recommend not use this feature or weight it less.





Again, Race and Ethnicity groups are not balanced in the dataset, one way is that we can group other groups together into a new bigger group.

All the variables above have very unbalanced data distribution, as a result we recommend to group other values together into a new group for later model feature selection.



The healthcare dataset has many Boolean columns that contains YES or NO values. Again many of the unbalanced features will not be recommended for later use.

## **Model Selection and Building**

Through the model-building process, I evaluated two Linear Classification models (Logistic Regression, Support Vector Machine), one Ensemble model (Random Forest), and one Boosting model (Extreme Gradient Boosting).

First, the dataset was split into train and test sets for model evaluation. Then, we use the training set to train fit the model, then take the test set to predict target variables. Finally, use different evaluation metrics to evaluate performance of models.

#### **Logistic Regression:**

```
X_train, X_test, y_train, y_test = train_test_split(X, y, test_size=0.2, random_state=1)

clr = LogisticRegressionCV(cv=5, scoring='accuracy')

clr.fit(X_train, y_train)

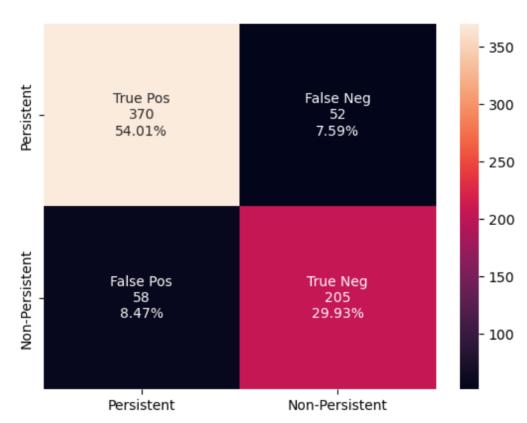
LogisticRegressionCV(cv=5, scoring='accuracy')

pred = clr.predict(X_test)

acc_log = accuracy_score(y_test, pred)
    print("Accuracy Score:", acc_log)
    pre_log = precision_score(y_test, pred)
    print('Precision Score:', pre_log)
    recall_log = recall_score(y_test, pred)
    print('Recall Score:', recall_log)
    fl_log = fl_score(y_test, pred)
    print('F1 Score:', f1_log)
    roc_log = roc_auc_score(y_test, pred)
    print('ROC Area:', roc_log)
```

Accuracy Score: 0.8394160583941606 Precision Score: 0.8148148148148

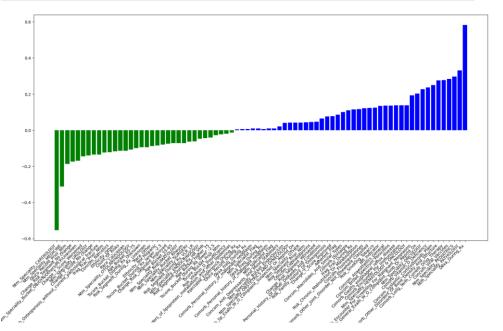
Recall Score: 0.752851711026616 F1 Score: 0.7826086956521738 ROC Area: 0.8231083199682843



#### Linear SVC:

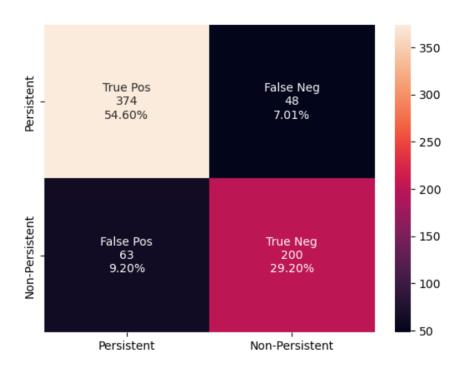
After fitting Linear SVC Model, I visualized all features by their importance level and remove the unimportant features:

```
# visualize feature importance
num_features = 39
coef = svc.coef_.ravel()
top_positive_coefficients = np.argsort(coef)[-num_features:]
top_negative_coefficients = np.argsort(coef)[:num_features]
top_coefficients = np.hstack([top_negative_coefficients, top_positive_coefficients])
plt.figure(figsize=(20,10))
colors = ['green' if c < 0 else 'blue' for c in coef[top_coefficients]]
plt.bar(np.arange(2 * num_features), coef[top_coefficients], color=colors)
feature_names = np.array(X.columns.values)
plt.xticks(np.arange(2 * num_features), feature_names[top_coefficients], rotation=45, ha='right')
plt.show()</pre>
```



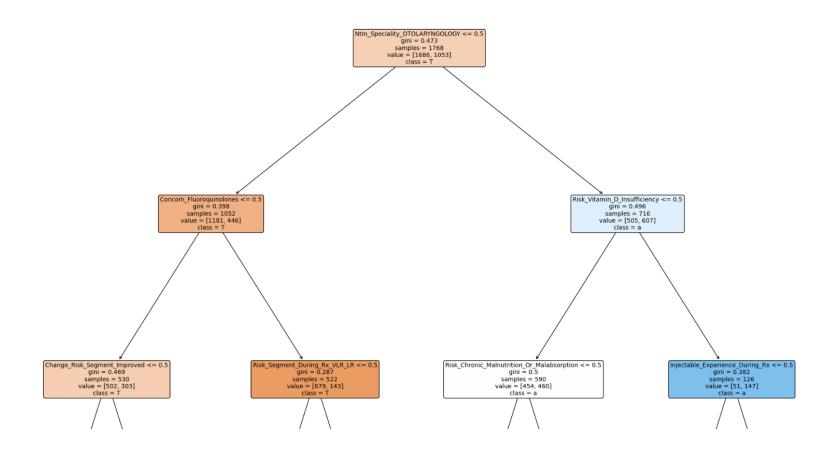
Accuracy Score: 0.8379562043795621 Precision Score: 0.8064516129032258 Recall Score: 0.7604562737642585

F1 Score: 0.7827788649706457 ROC Area: 0.8233560989674372



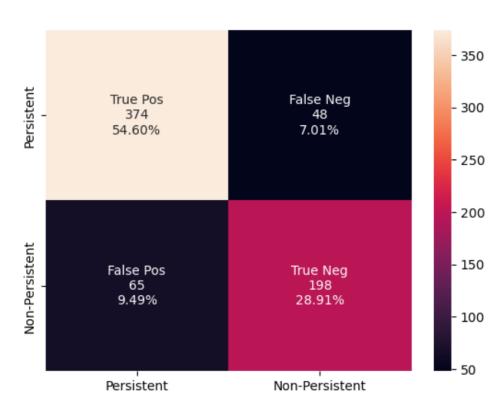
#### **Random Forest:**

Fit the data to Ensemble model Random Forest and visualized the top nodes of some of the decision trees:



Accuracy Score: 0.8350364963503649 Precision Score: 0.8048780487804879 Recall Score: 0.752851711026616

F1 Score: 0.7779960707269156 ROC Area: 0.8195538175986161



### **XGboost:**

For Boosting model, I selected XGboost to fit the data. I used Bayesian optimization to tune the hyperparameters:

```
params_gbm ={
    'max_depth':(1, 10),
    'learning_rate':(0.01, 1),
    'n_estimators':(80, 500),
    'subsample': (0.8, 1),
}
gbm_bo = BayesianOptimization(gbm_cl_bo, params_gbm, random_state=1)
gbm_bo.maximize(init_points=20, n_iter=4)
```

iter	target	learni	max_depth	n_esti	subsample
1	0.793	0.4229	7.483	80.05	0.8605
2	0.8109	0.1553	1.831	158.2	0.8691
3	0.7824	0.4028	5.849	256.1	0.937
4	0.7904	0.2124	8.903	91.5	0.9341
5	0.7857	0.4231	6.028	139.0	0.8396
6	0.789	0.8027	9.714	211.6	0.9385
7	0.782	0.8776	9.051	115.7	0.8078
8	0.797	0.1781	8.903	121.3	0.8842
9	0.7715	0.9583	5.798	370.6	0.8631
10	0.7839	0.6896	8.512	87.68	0.95
11	0.7802	0.989	7.733	197.8	0.9579
12	0.7915	0.1122	5.031	461.6	0.8587
13	0.8109	0.2949	2.17	88.13	0.9358
14	0.8003	0.2195	3.39	286.5	0.8107
15	0.7901	0.5784	2.321	327.5	0.94
16	0.7923	0.1113	4.727	371.6	0.8828
17	0.7967	0.05945	5.823	358.8	0.903
18	0.7882	0.9451	6.279	459.4	0.8275
19	0.7981	0.1479	8.267	247.0	0.8331
20	0.778	0.9282	4.13	395.3	0.9452
21	0.8076	0.01764	2.457	377.9	0.9529
22	0.7864	0.4481	4.721	310.1	0.9488
23	0.7956	0.3617	8.191	247.1	0.9915
24	0.8098	0.09787	1.855	159.1	0.8381
========			=========		

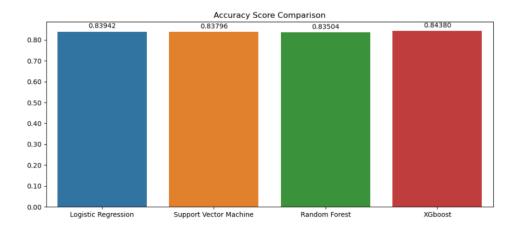
Accuracy Score: 0.8437956204379562 Precision Score: 0.8023255813953488

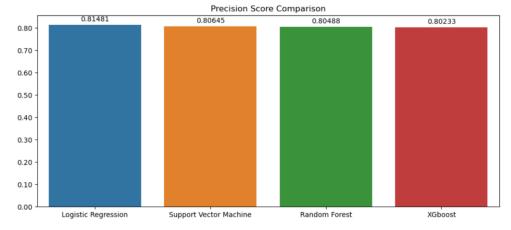
Recall Score: 0.7870722433460076

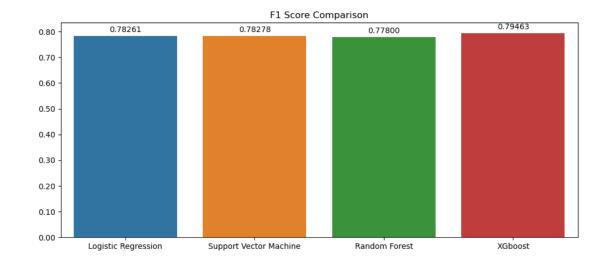
F1 Score: 0.7946257197696737 ROC Area: 0.8331095813886435

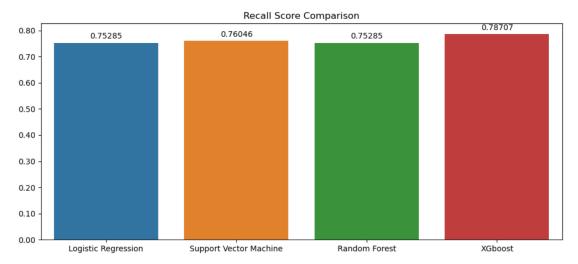


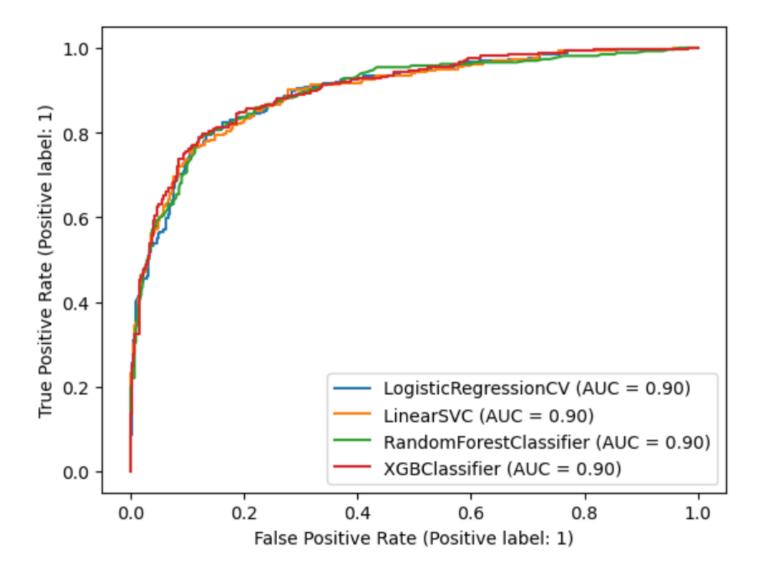
### I then grouped all the performances together into barplots to compare the difference:











Overall, every model has pretty similar performance regarding different evaluation metrics. Among them, XGboost has the best average performance. So it is recommended to use the XGboost model to make predictions. The model is save as the best model file for later use.

```
import pickle

# save the best model
filename = 'best_model.sav'
pickle.dump(clf, open(filename, 'wb'))

loaded_model = pickle.load(open(filename, 'rb'))
loaded_model.score(X_test, y_test)

0.8437956204379562
```