Project Scope Statement

POLYCYSTIC OVARY SYNDROME DATA(PCOS)

Tianqi Zhou, Zixuan Li, Yifei Wu GEORGETOWN UNIVERSITY |

Table of Contents

1 I	Project OutlinesProject Outlines	2
	Project Description	
	Date Submitted	
	Project Priority	
	1.1 Step 1. Project Deliverables	
	1.2 Step 2. List of Project Tasks	
	1.3 Step 3. Out of Scope	
	1.4 Step 4. Project Assumptions	
	1.5 Step 5. Project Constraints	
	1.6 Step 6. Updated Estimates	
2. I	ntroduction	
	Analysis of the Dataset and Trained Model	
<i>J</i> 1	3.1 Exploratory Analysis and Visualization	
	J.I LAPIOI WOLY I MILLI Y DIG WILL I I DUMINE MILLION SOCIONOS SOCIENOS SOCIONOS SOCIONOS SOCIONOS SOCIONOS SOCIONOS SOCIONOS SOCIENOS SOCIONOS SOCIENOS SOCIONOS SOC	•••¬

1 Project Outlines

Project #	Project Description	Date Submitted	Project Priority
1	Georgetown University is committed to cultivating an environment centered on personalized care tailored to the specific needs of patients. In pursuit of this goal, they are embarking on the development of an application designed to swiftly detect and diagnose PCOS (Polycystic Ovary Syndrome) as soon as a patient's laboratory results are input into the system. To support this initiative, our project group designed a machine learning application that leverages existing patient data to enhance the speed and accuracy of PCOS diagnosis, uncovering intricate details that may prove challenging for human assessment.		Priority 01

1.1 Step 1. Project Deliverables

Please list *all project deliverables* listed in the Project Charter and, if necessary, elaborate on them. *Do not list dates*. Add more rows as necessary.

Deliverable	Description			
ID#				
1	roblem Identification			
2	Method evaluation			
3	Solution by Hyperparameter Optimization			
4	Deployment Pipeline/Platform			
5	Project Demo In class			

1.2 Step 2. List of Project Tasks

Please list *all project tasks* to be completed, based on the "Deliverables" specified in the Project Charter. *Do not list dates*. Add more rows as necessary. Optional: You may substitute a work breakdown structure (WBS) or mind map in lieu of Step 2. Please attach WBS or mind-map to the document.

Task	Task to be completed	Delivery	For
ID#		Date	Deliverable #
1	Submit Project Charter	09/07/2023	1
2	Method evaluation - research on techniques, quality assurance, project approach.	09/28/2023	2
3	Solution by Hyperparameter Optimization - actual effective techniques used.	10/12/2023	3
4	Deployment Pipeline/Platform.	10/26/2023	4
5	Project Demo In class - Final report of the project in addition to the cloud deployed link.	11/16/2023	5

1.3 Step 3. Out of Scope

This project will NOT	This project will not include any analysis of other diseases of the patient
accomplish or include	and other background details.
the following:	

1.4 Step 4. Project Assumptions

Please list any project factors that will be considered to be true, real, or certain. Assumptions generally involve a certain degree of risk.

#	Assumption
1	Filling in missing data with the mean value of the feature won't our model accuracy
2	The distribution of variables won't affect the model accuracy
3	Resampling won't cause generalization issues

1.5 Step 5. Project Constraints

Project Start Date	09/07/2023
Launch/Go-Live Date	10/26/2023
Project End Date	11/16/2023
List any hard deadline(s)	09/07/2023
	09/28/2023
	10/12/2023
	10/26/2023
	11/16/2023
List other dates/descriptions of key milestones	None
Budget constraints	N/A
Enter information about project budget limitations	
Quality or performance constraints	N/A
Enter any other requirements for the functionality, performance, or quality of the	
project	
Equipment/personnel Constraints	N/A
Enter any constraints regarding equipment or people that will impact the project	
Regulatory constraints	N/A
Enter any legal, policy or other regulatory constraints	

1.6 Step 6. Updated Estimates

Estimate T&C hours	Enter total # of	If charge-back project,	Enter N/A if not
required to complete	T&C hours	list total estimated T&C	applicable.
project		cost	

2 Introduction

Polycystic Ovary Syndrome (PCOS) is a hormonal disorder that impacts women of reproductive age, potentially causing symptoms like hormonal imbalances, irregular menstrual cycles, and emotional disturbances, as recognized by the World Health Organization (WHO). It affects approximately 8–13% of women in their reproductive years, with as many as 70% of cases going unnoticed. Diagnosing PCOS

can be challenging since its symptoms can be attributed to various factors, such as heavy menstrual bleeding.

Georgetown University is deeply committed to fostering an environment focused on personalized patient care, tailored to individual needs. In pursuit of this mission, the university is embarking on the development of an application aimed at promptly identifying and diagnosing PCOS upon the entry of a patient's laboratory results into the system. The ultimate goal is to increase the rate of diagnosis.

To support this endeavor, our project group designed an application based on machine-learning approaches that harness existing patient data to enhance the speed and precision of PCOS diagnosis. The data used is from Kaggle called PCOS Diagnosis. The dataset contains 45 variables including the patient's biometric measurements, blood test, hormone data, and other symptoms. All of them are numerical and binary data. Model selection will be performed to select the most effective models for this dataset. Models for comparison include Logistic Regression, Ridge Regression, Lasso Regression, Decision Tree Classifier, Random Forest Classifier, Bagging Classifier, Gradient Boosting Classifier, and XGB Classifier. This innovative approach seeks to better evaluate the patient's situation based on the likelihood of PCOS, ultimately improving the overall management and treatment of PCOS.

3 Analysis of the Dataset and Trained Model

3.1 Exploratory Analysis and Visualization

The dataset has 541 samples with 1 missing column in marriage status and 1 missing value in AMH. The mean of the feature was used to fill in the missing value. Our project group first did the exploratory analysis to obtain the distribution of our data and the relationship between the target and feature variables.

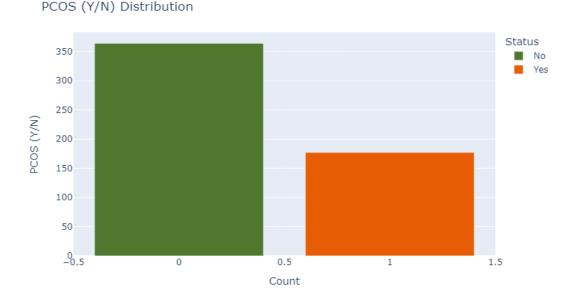


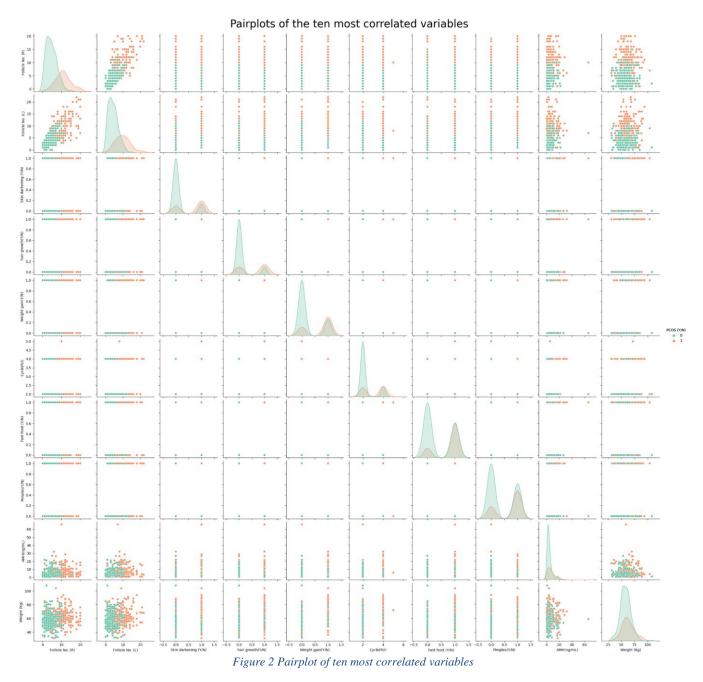
Figure 1 Distribution of the dependent variable (PCOS)

Figure 1 shows the distribution of the dependent variable, PCOS. Our dataset contains 177 positive samples and 364 negative samples. Due to the imbalanced distribution, we considered using a resampling method called SMOTE to increase the number of positive samples.

Table 1 Summary of the Numerical Variables

para m.	Age (yrs)	Weight (Kg)	Height (Cm)	BMI	Pulse rate (bpm)	RR (brths /min)	Hb(g/dl)	Cycle length (days)	Marria ge Status (Yrs)	No. of absorptions		
mean	31.43	59.64	156.48	24.32	73.25	19.24	11.16	4.94	7.68	(0.29	
std	5.41	11.03	6.03	4.05	4.43	1.69	0.87	1.49	4.8	(
min	20	31	137	12.42	13	16	8.5	0	0		0	
25%	28	52	152	21.71	72	18	10.5	4	4		0	
50%	31	59	156	24.24	72	18	11	5	7		0	
75%	35	65	160	26.64	74	20	11.7	5	10		0	
max	48	108	180	38.9	82	28	14.8	12	30		5	
para m.	I beta- HCG(m IU/mL)	II beta- HCG(m IU/mL)	FSH(m IU/mL)	LH(mIU/ mL)	FSH /LH	Hip (inch)	Waist (inch)	Waist: Hip Ratio	TSH (mIU/L)	AMH(ng/mL)		
mean	664.55	5.62	14.6	6.47	6.9	37.99	33.84	0.89	2.98	:	5.62	
std	3348.92	5.88	217.02	86.67	60.69	3.97	3.6	0.05	3.76	5.88		
min	1.3	0.1	0.21	0.02	0	26	24	0.76	0.04	0.1		
25%	1.99	2.01	3.3	1.02	1.42	36	32	0.86	1.48	2.01		
50%	20	3.7	4.85	2.3	2.17	38	34	0.89	2.26	3.7		
75%	297.21	6.9	6.41	3.68	3.96	40	36	0.93	3.57	6.9		
max	32460.9 7	66	5052	2018	1372.83	48	47	0.98	65	66		
para m.	PRL(ng/ mL)	Vit D3 (ng/mL)	PRG(n g/mL)	RBS(mg/ dl)	BP _S(mm Hg)	BP _D(m mHg)	Follicle No. (L)	Follicle No. (R)	Avg. F size (L) (mm)	Avg. F size (R) (mm)	Endometri um (mm)	
mean	24.32	49.92	0.61	99.84	114.66	76.93	6.13	6.64	15.02	15.45	8.48	
std	14.97	346.21	3.81	18.56	7.38	5.57	4.23	4.44	3.57	3.32	2.17	
min	0.4	0	0.05	60	12	8	0	0	0	0	0	
25%	14.52	20.8	0.25	92	110	70	3	3	13	13	7	
50%	21.92	25.9	0.32	100	110	80	5	6	15	16	8.5	
75%	29.89	34.5	0.45	107	120	80	9	10	18	18	9.8	
max	128.24	6014.66	85	350	140	100	22	20	24	24	18	

Table 1 presents the statistics of the numerical variables in our dataset.



Since our dataset contains 42 different features, we selected the ten most correlated variables based on the correlation coefficient and visualized their distributions. Different colors in Figure 2 indicate distinct diagnostic results. The results indicate that only weight is normally distributed, while both Follicle No. (right and left) are right-skewed and AMH gathers near 0. However, speaking of different diagnostic groups, Follicle numbers and AMH become closer to normal distributions. Moreover, from Figure 2 we can primarily assume that phenomena like skin darkening, hair growth, weight gain, and lifestyles of eating fast food have important influences on the occurrence and diagnosis of PCOS.