

HOSPITAL MORTALITY



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DATASET: In Hospital Mortality Prediction

from [Kaggle.com](https://www.kaggle.com)

Our motivation: We wanted to work with data that had meaning and could impact everyday life.

It is essential to understand what features can lead to hospital mortality and what other diseases could possibly hold more weight in predictions.

VARIABLES

What are the predictors? What are we predicting?

Age, Gender, BMI, Hypertensive, Atrialfibrillation, CHD with no MI, Diabetes, Deficiencyanemias, Depression, Hyperlipemia, Renal failure, COPD, Heart Rate, Systolic blood pressure, Diastolic blood pressure, Respiratory rate, Temperature, SP O2, Urine output, Hematocrit, RBC, MCH, MCHC, MCV, RDW, Leucocyte, Platelets, Neutrophils, Basophils, Lymphocyte, INR, NT-proBNP, Creatine kinase, Creatinine, Urea nitrogen, Glucose, Blood potassium, Blood sodium, Blood calcium, Chloride, Anion gap, Magnesium ion, PH, Bicarbonate, Lactic acid, PCO2, EF





DATA CLEANING & STANDARDIZING

- Check NULL Values
- Drop them
- Z-scale to make sure all variables are on the same scale



Question 1

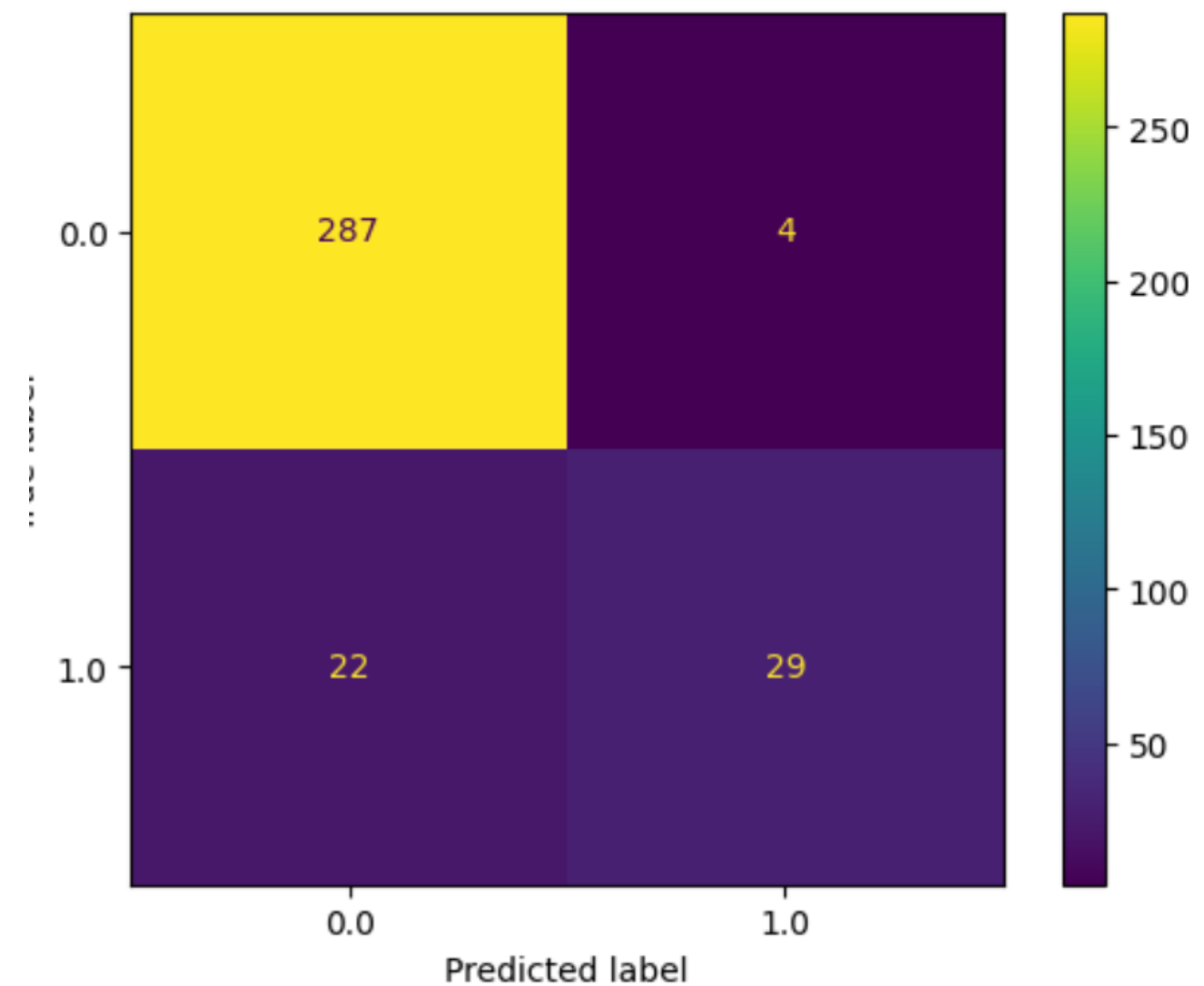
What are the most common health conditions that lead to in-hospital mortality?

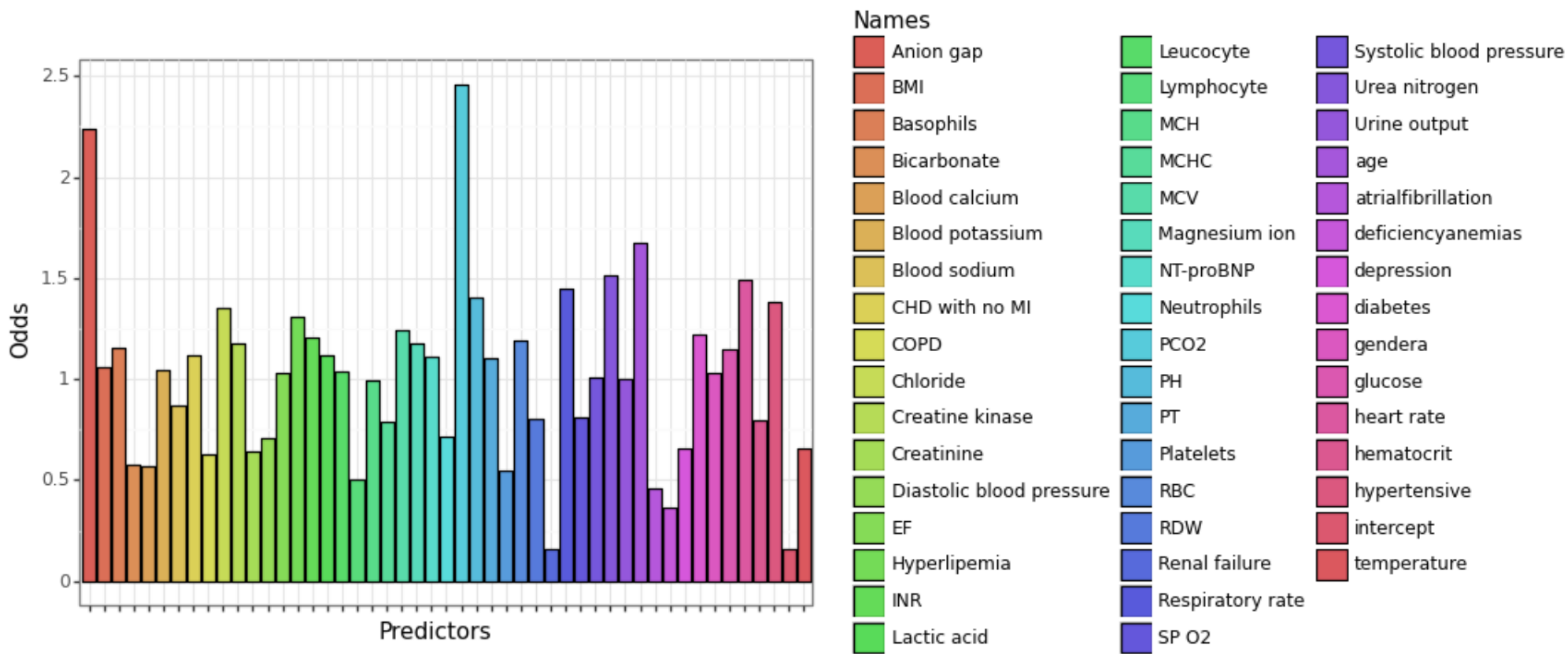
```
# metrics
print("Accuracy: ", accuracy_score(y_test, predictedVals))
print("F1 Score: ", f1_score(y_test, predictedVals))
print("Recall: ", recall_score(y_test, predictedVals))
print("Precision: ", precision_score(y_test, predictedVals))

Accuracy:  0.8953488372093024
F1 Score:  0.6086956521739131
Recall:  0.5
Precision:  0.7777777777777778

# metrics
print("Accuracy: ", accuracy_score(y_train, myLogit.predict(X_train)))
print("F1 Score: ", f1_score(y_train, myLogit.predict(X_train)))
print("Recall: ", recall_score(y_train, myLogit.predict(X_train)))
print("Precision: ", precision_score(y_train, myLogit.predict(X_train)))

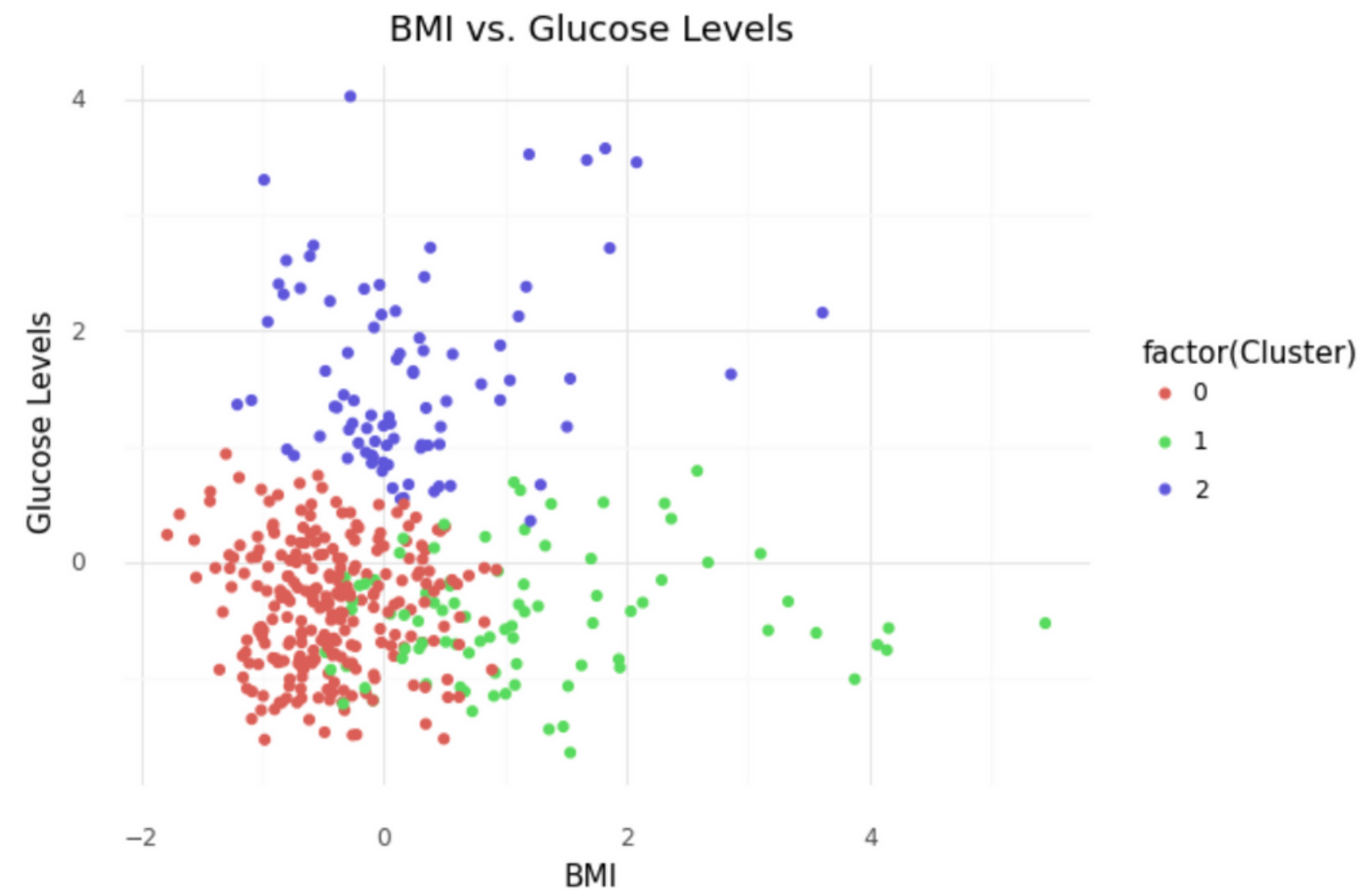
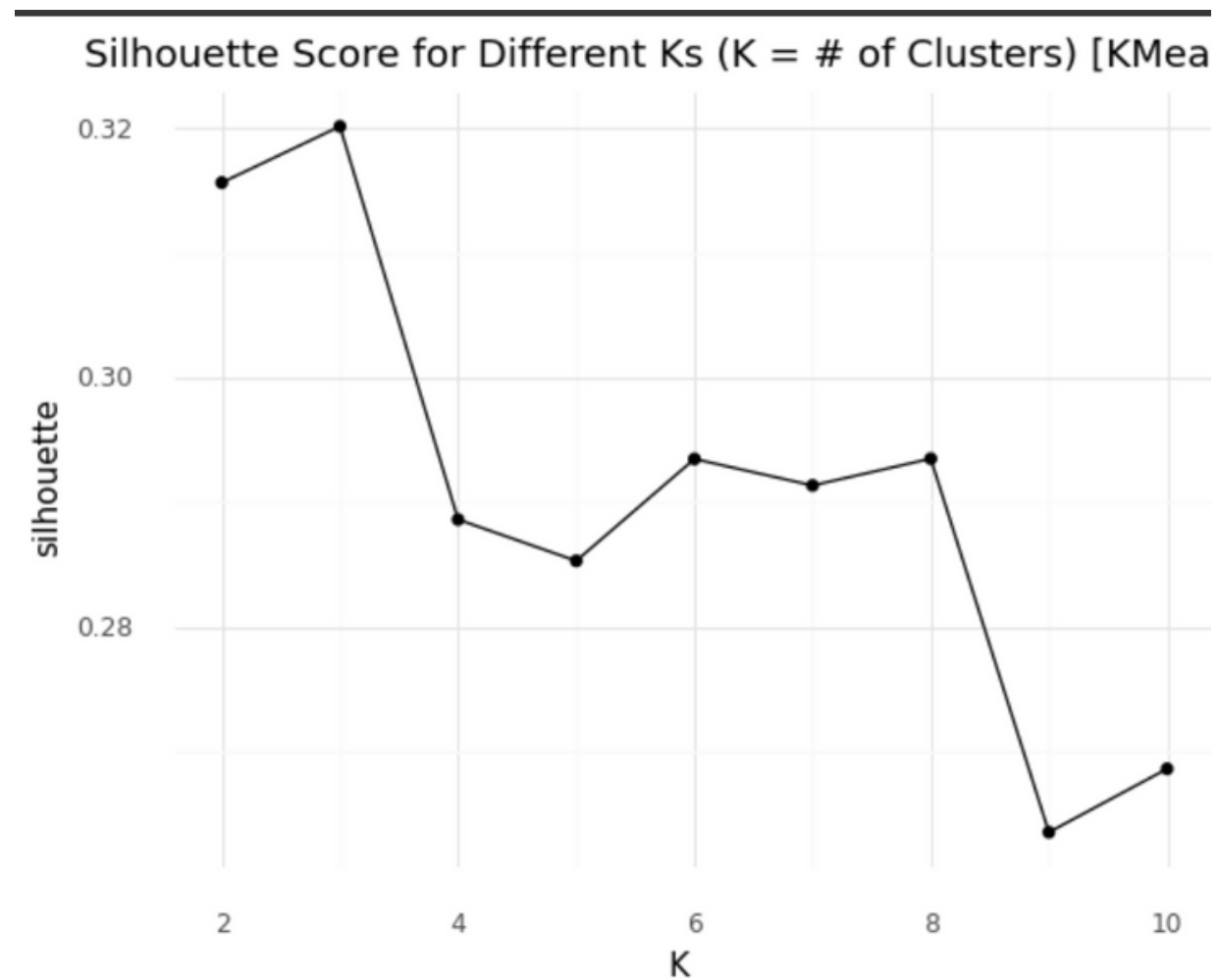
Accuracy:  0.9239766081871345
F1 Score:  0.6904761904761905
Recall:  0.5686274509803921
Precision:  0.8787878787878788
```





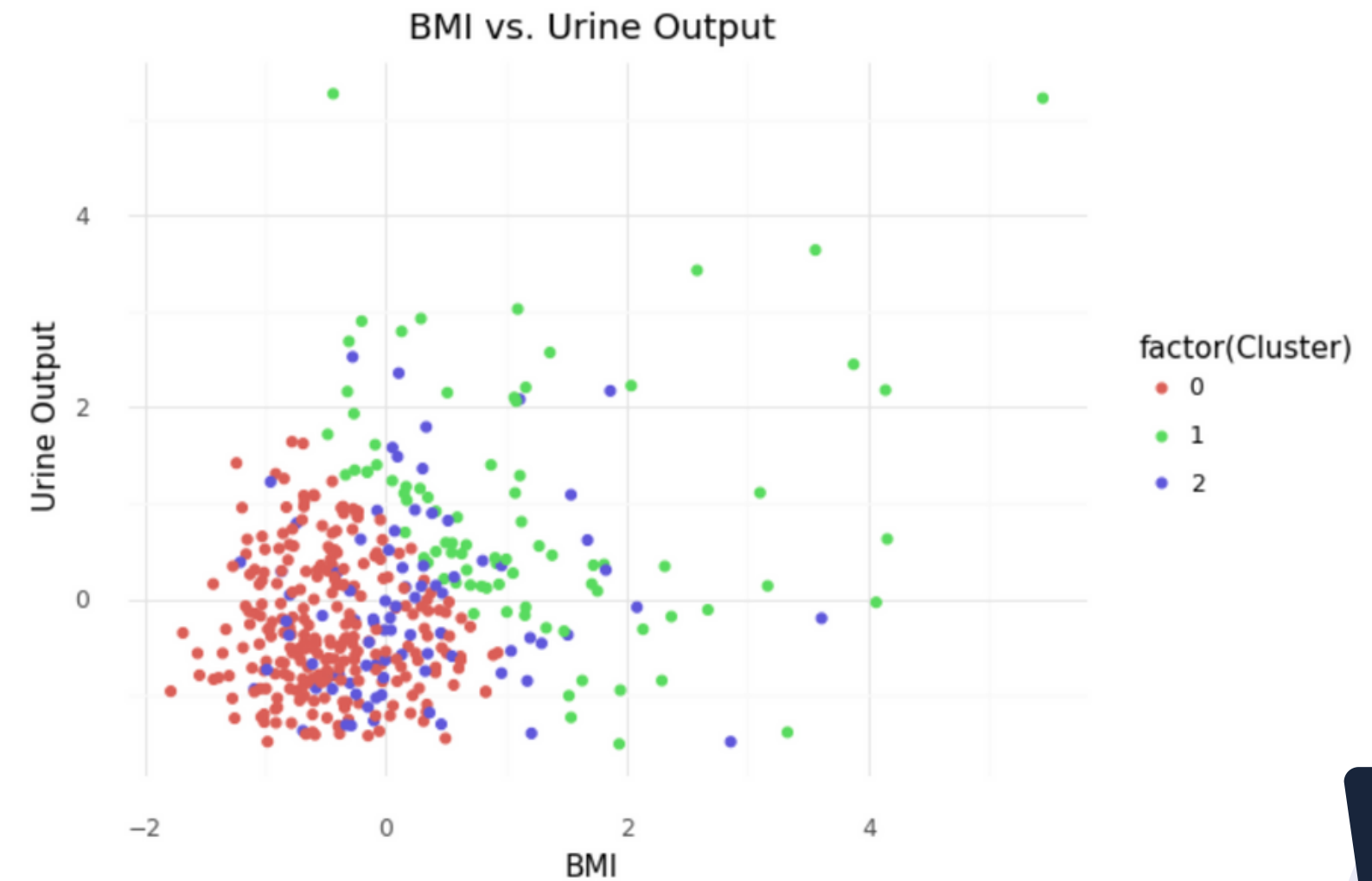
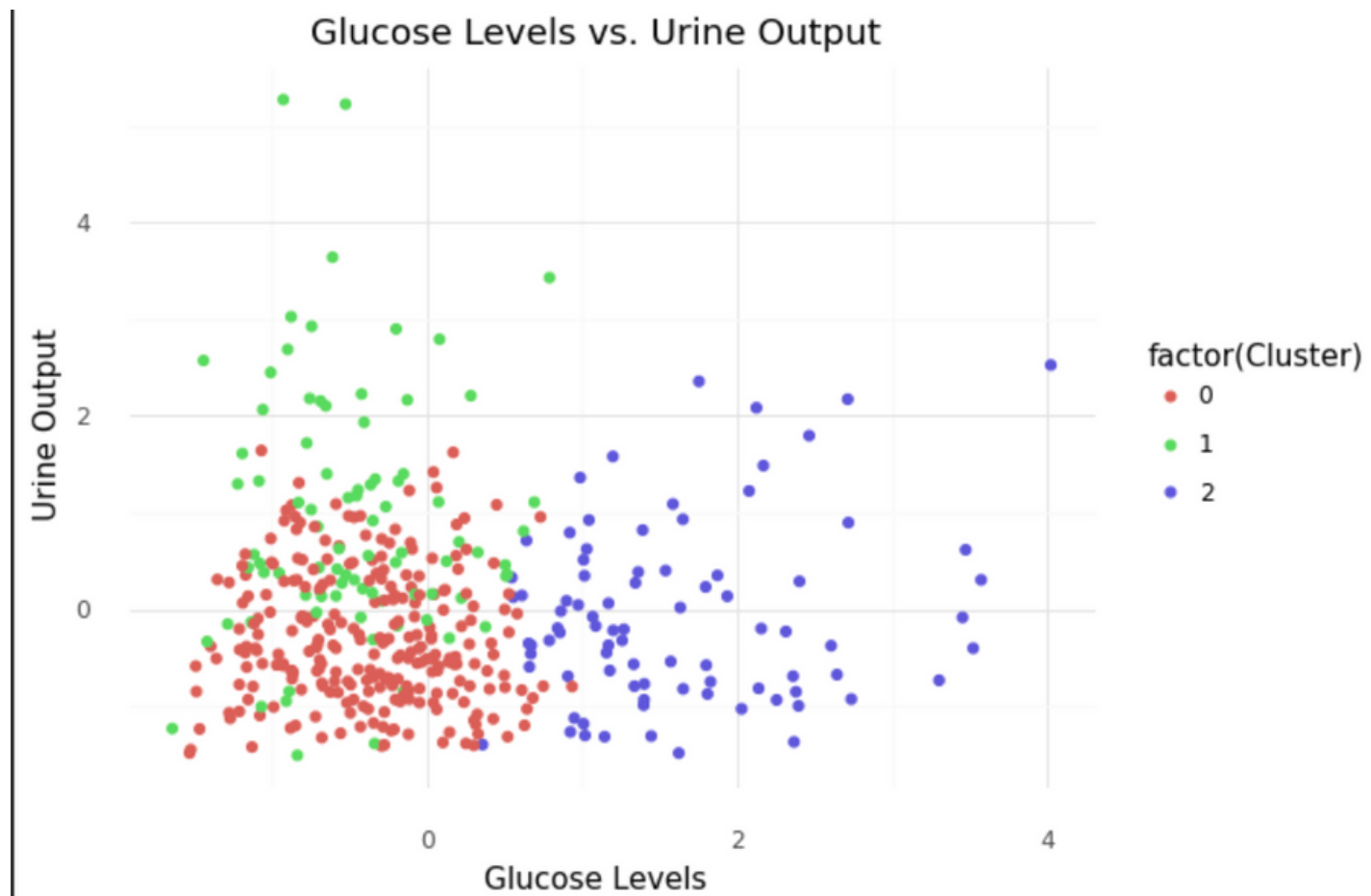
Question 2

When considering features BMI, Glucose, and Urine Output, what clusters may emerge and how can we characterize those clusters?



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Question 3

Are there any differences in the predictive performance of our model across different subgroups of patients, such as patients who are diagnosed with depression, or comorbidity status (renal failure, diabetes, hypertensive)?

Accuracy Scores:

Accuracy (Depression): 0.8255
Accuracy (Renal Failure): 0.8605
Accuracy (Diabetes): 0.8604
Accuracy (Hypertensive): 0.8372
Accuracy (All): 0.8139

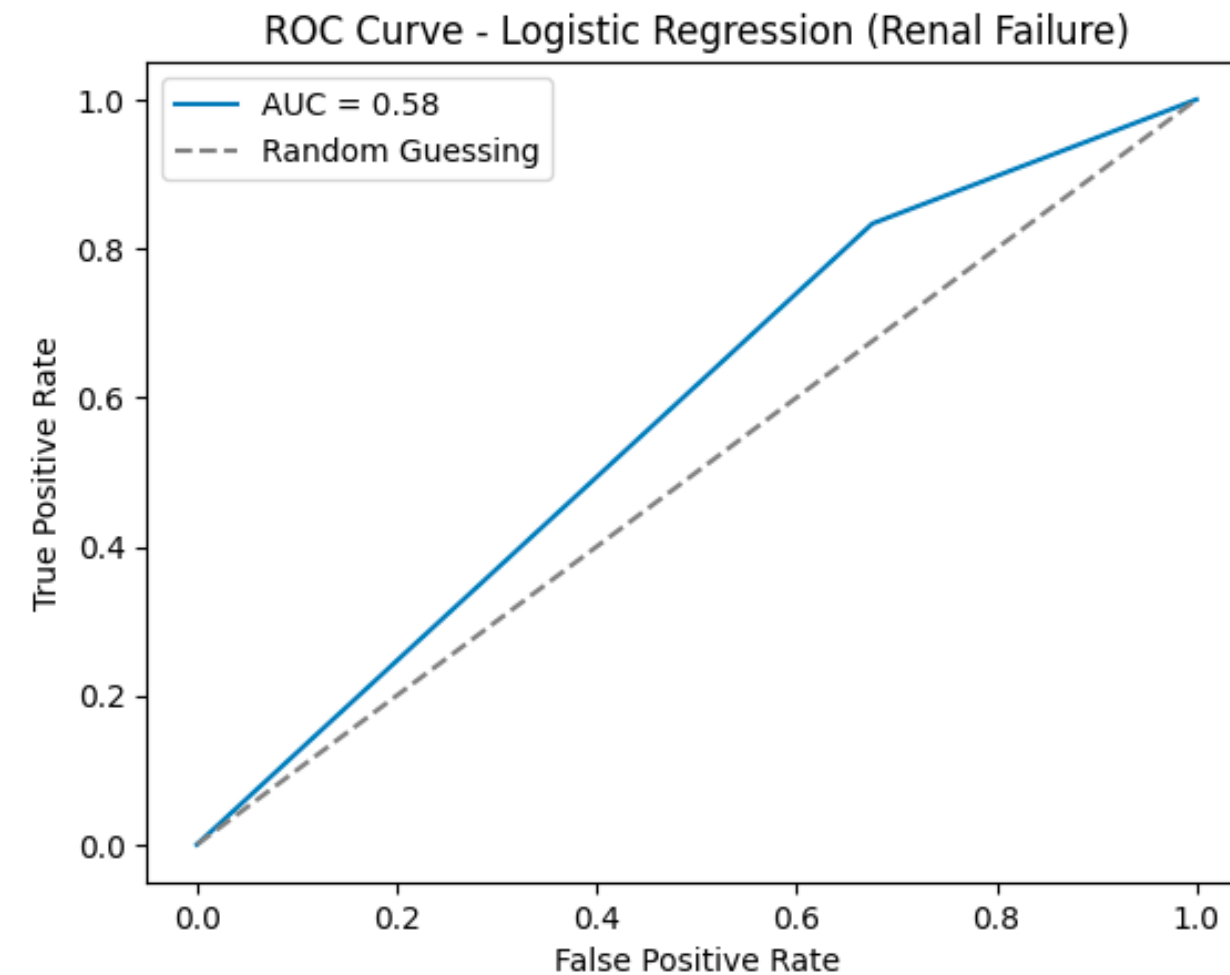
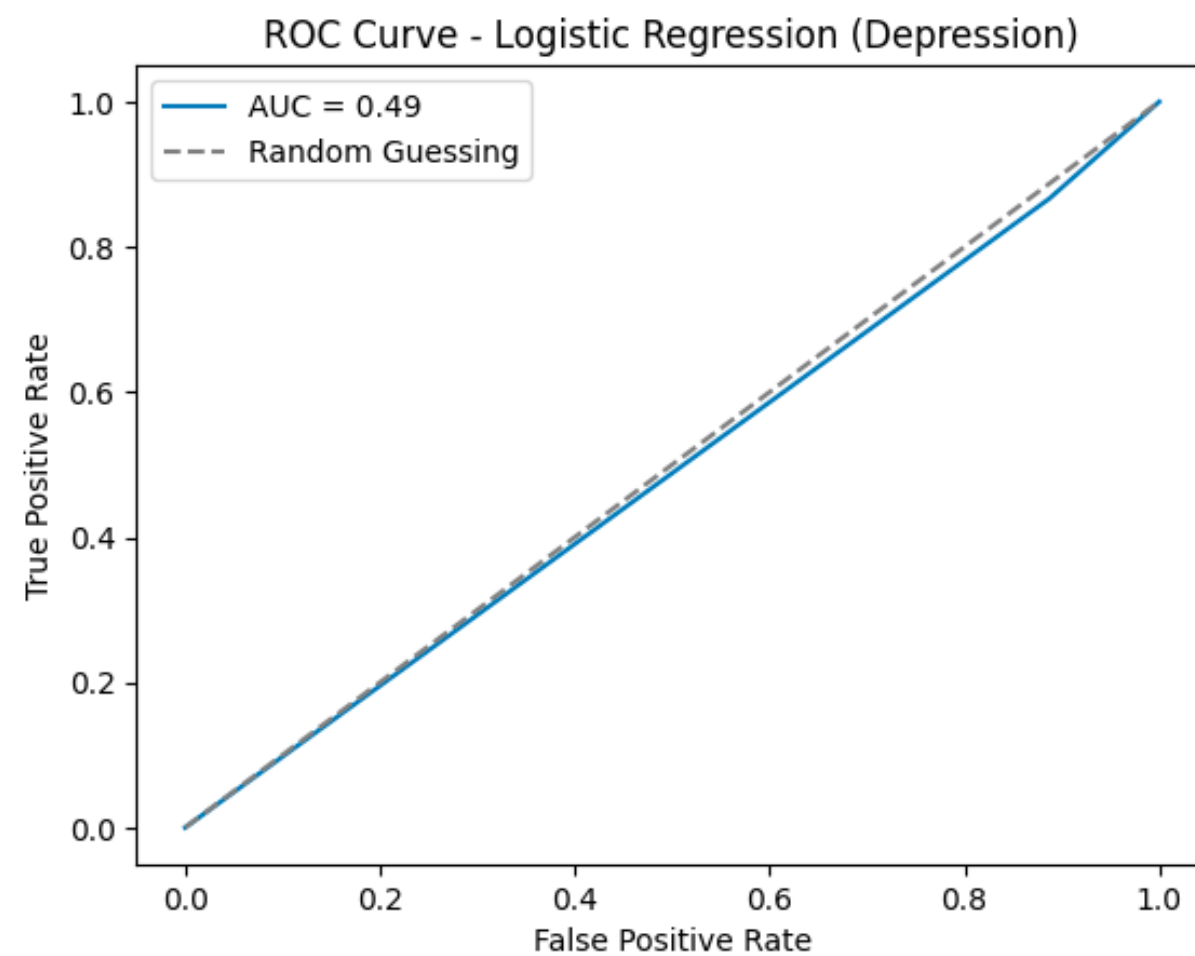
Odds Coefficients:

	Coef	Names	Odds
0	-1.232983	depression	0.291422
1	-0.741782	Renal failure	0.476264
2	-0.005079	diabetes	0.994934
3	-0.004830	hypertensive	0.995182
4	-1.444756	intercept	0.235804



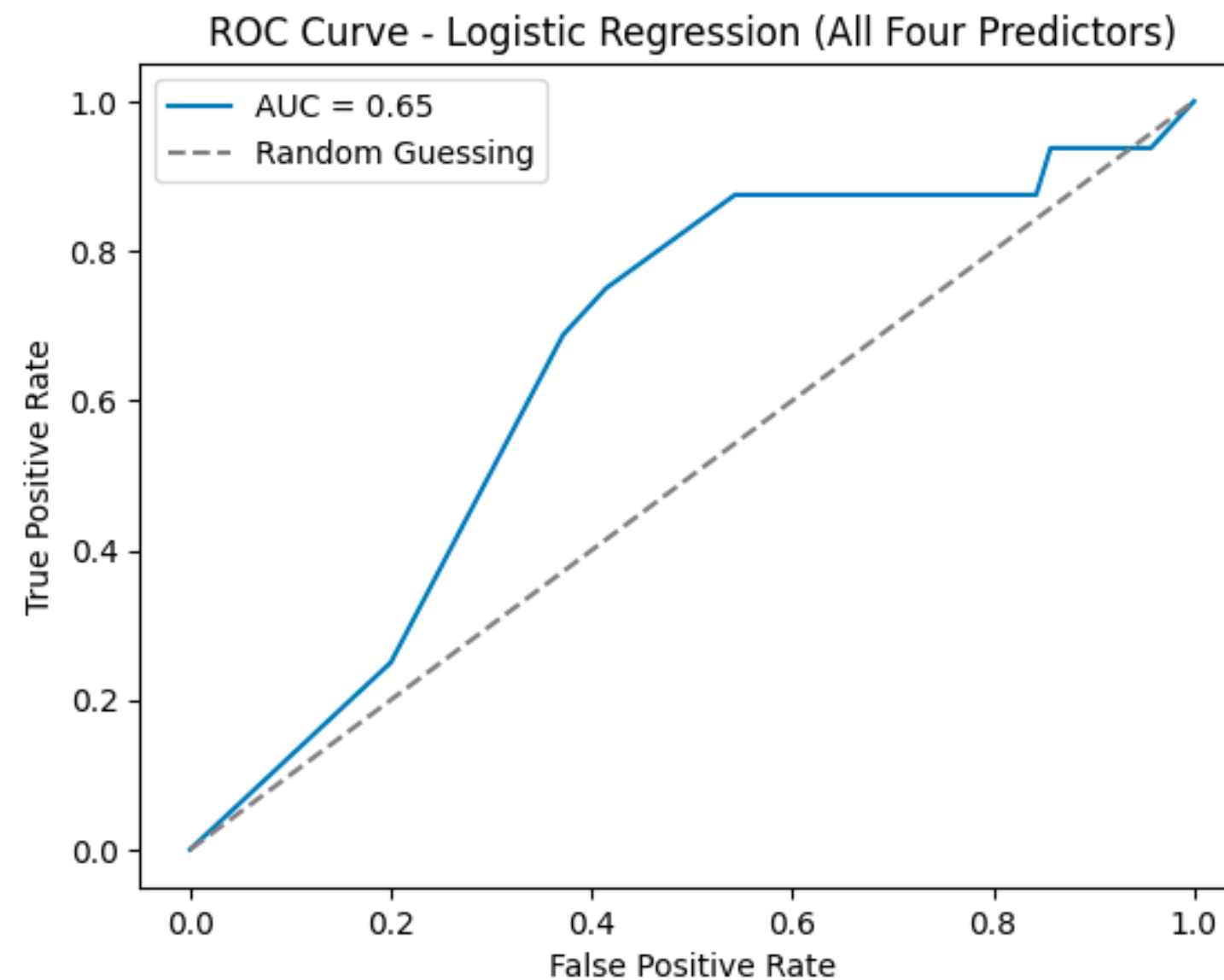
Question 3

Are there any differences in the predictive performance of our model across different subgroups of patients, such as patients who are diagnosed with depression, or comorbidity status (renal failure, diabetes, hypertensive)?



Question 3

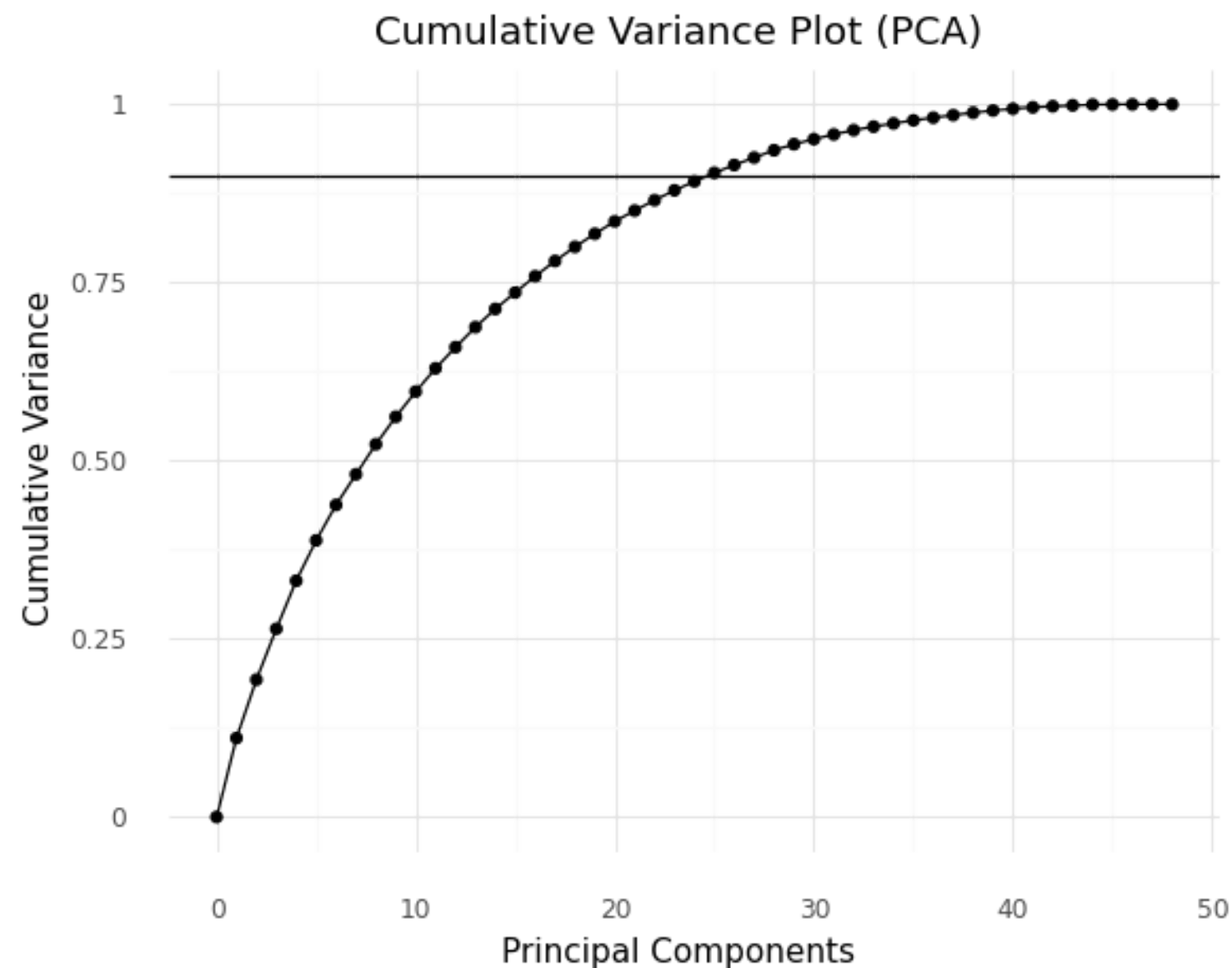
Are there any differences in the predictive performance of our model across different subgroups of patients, such as patients who are diagnosed with depression, or comorbidity status (renal failure, diabetes, hypertensive)?



Question 4

How does the mean absolute error differ between the train and test data when using Principle Component Analysis on all continuous variables, and retaining enough Principle Components to keep 90% of the variance, to predict hospital mortality with our model(s)?

	expl_var	pc	cum_var
0	0.110861	1	0.110861
1	0.081857	2	0.192718
2	0.071281	3	0.263999
3	0.067548	4	0.331547
4	0.056565	5	0.388113



**Using 25 PCs
for 90%
variance**



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How does the mean absolute error differ between the train and test data when using Principle Component Analysis on all continuous variables, and retaining enough Principle Components to keep 90% of the variance, to predict hospital mortality with our model(s)?

Logistic Regression Results:

Accuracy for Original Train Set: 0.9239

Accuracy for Original Test Set: 0.8953

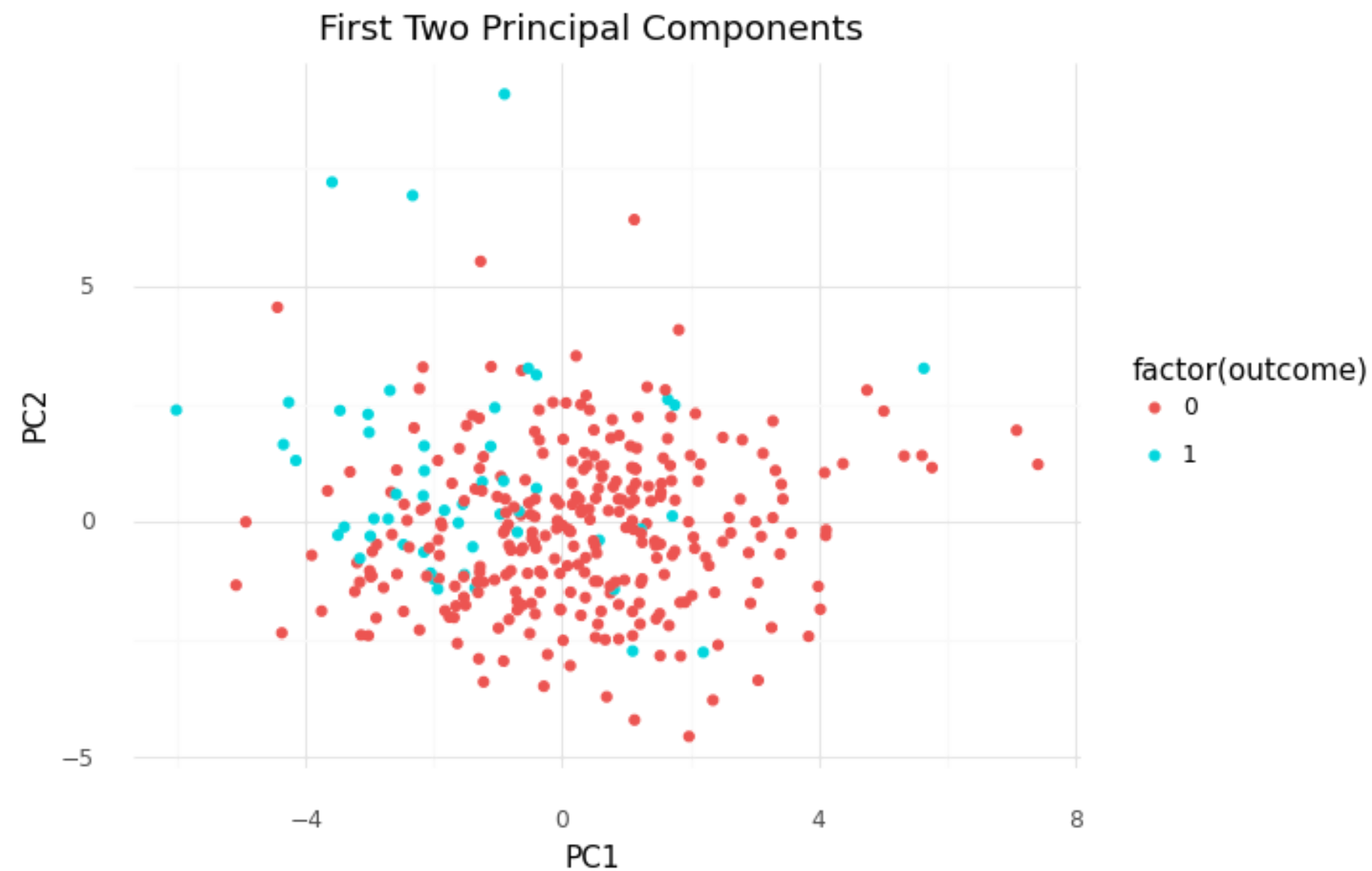
Accuracy for PCA Train Set: 0.8976

Accuracy for PCA Test Set: 0.8721



Question 4

How does the mean absolute error differ between the train and test data when using Principle Component Analysis on all continuous variables, and retaining enough Principle Components to keep 90% of the variance, to predict hospital mortality with our model(s)?



Question 5

Out of the 48 variables observed in the dataset (excluding ground and ID), how can we utilize the Regularization method LASSO to select which variables have the most impact in predicting hospital mortality?

Coefficients		Predictors
0	0.033263	age
1	0.000000	gendera
2	0.023522	BMI
3	-0.000000	hypertensive
4	-0.010378	atrialfibrillation
5	0.027015	CHD with no MI
6	-0.016475	diabetes
7	-0.000000	deficiencyanemias
8	-0.002012	depression
9	0.000000	Hyperlipemia
10	0.000000	Renal failure
11	0.000000	COPD
12	-0.000000	heart rate
13	0.009652	Systolic blood pressure
14	0.000000	Diastolic blood pressure
15	0.009747	Respiratory rate
16	-0.041035	temperature
17	-0.000000	SP O2
18	0.000000	Urine output
19	-0.016460	hematocrit
20	0.000000	RBC
21	0.022599	MCH
22	0.000355	MCHC
23	0.005712	MCV
24	-0.021458	RDW
25	0.054432	Leucocyte
26	0.011131	Platelets
27	0.000000	Neutrophils

28	0.000000	Basophils
29	-0.053990	Lymphocyte
30	0.000000	PT
31	0.064715	INR
32	0.000000	NT-proBNP
33	-0.005930	Creatine kinase
34	-0.014864	Creatinine
35	0.028233	Urea nitrogen
36	0.034051	glucose
37	0.006354	Blood potassium
38	0.000000	Blood sodium
39	0.000000	Blood calcium
40	-0.000000	Chloride
41	0.000000	Anion gap
42	0.000000	Magnesium ion
43	-0.033004	PH
44	-0.000000	Bicarbonate
45	0.000000	Lactic acid
46	-0.124233	PCO2
47	-0.000000	EF

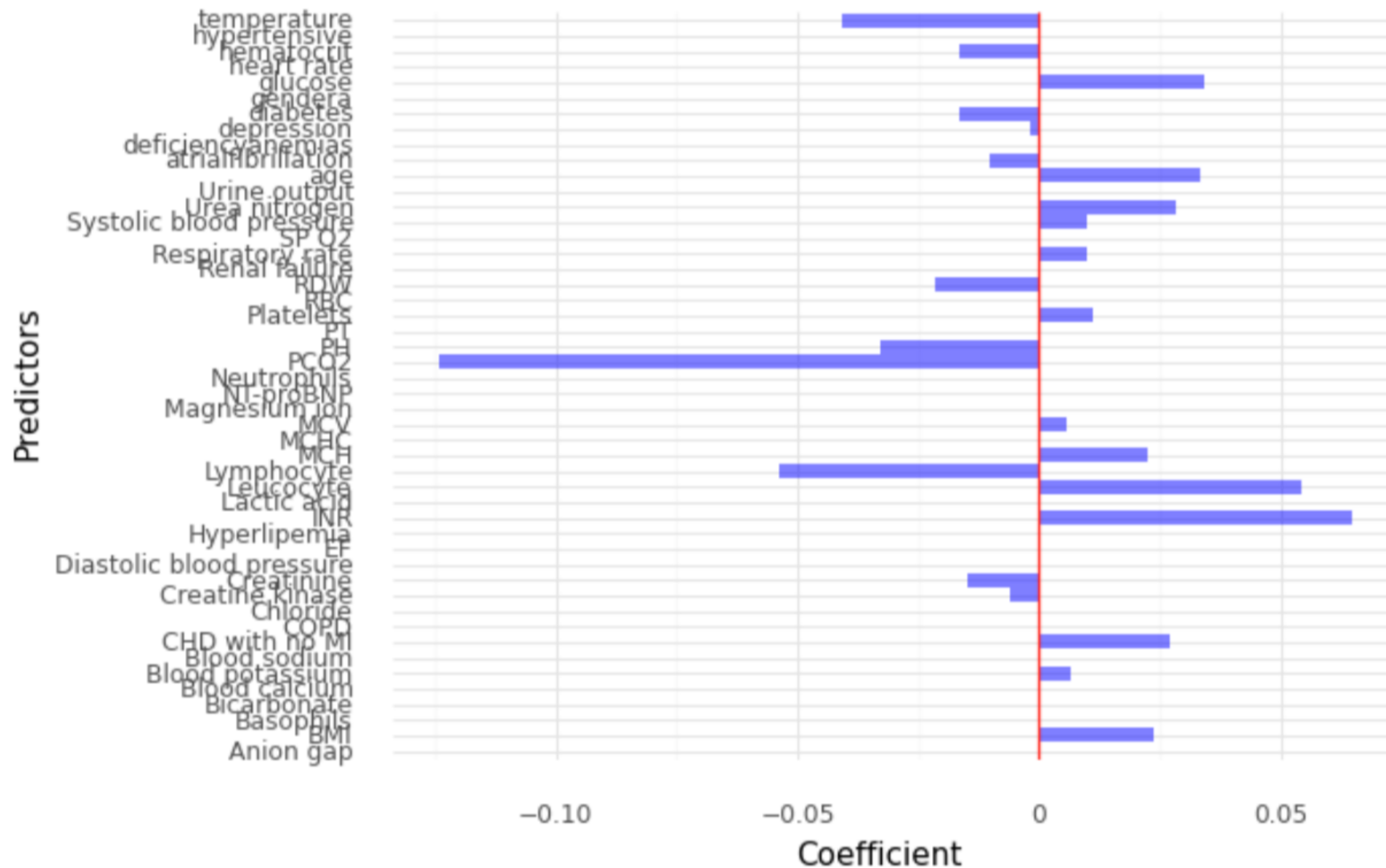
PENALTY: 0.01
TRAIN: 0.3396888666530895
TEST : 0.3918317430757958

Predictors that got shrunk to 0:

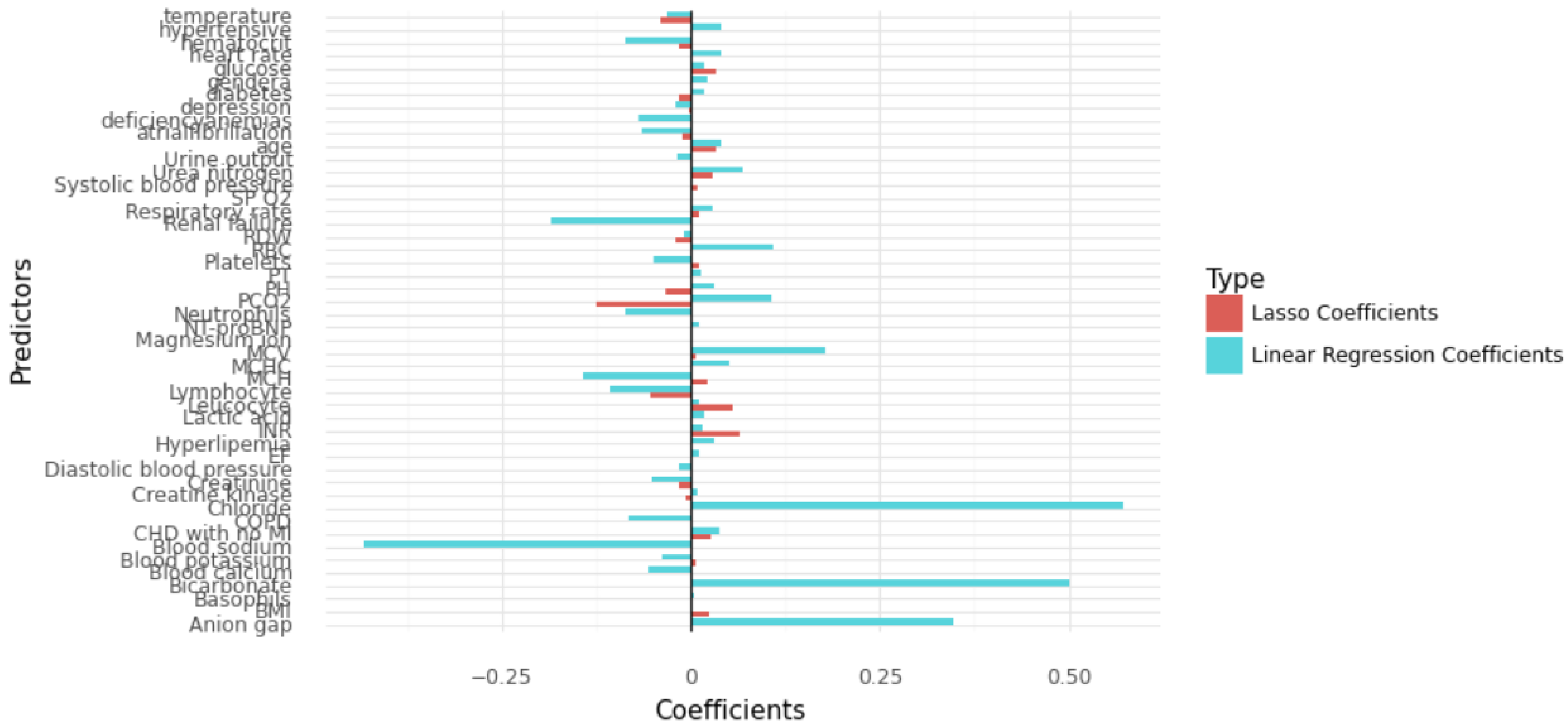
- gender
- hypertensive
- Hyperlipemia
- Renal failure
- COPD
- heart rate
- Diastolic blood pressure
- SP O2
- Urine output
- RBC
- Neutrophils
- Basophils
- PT
- NT-proBNP
- Blood sodium
- Blood calcium
- Chloride
- Anion gap
- Magnesium ion
- Bicarbonate
- Lactic acid
- EF



Lasso Regression Coefficients



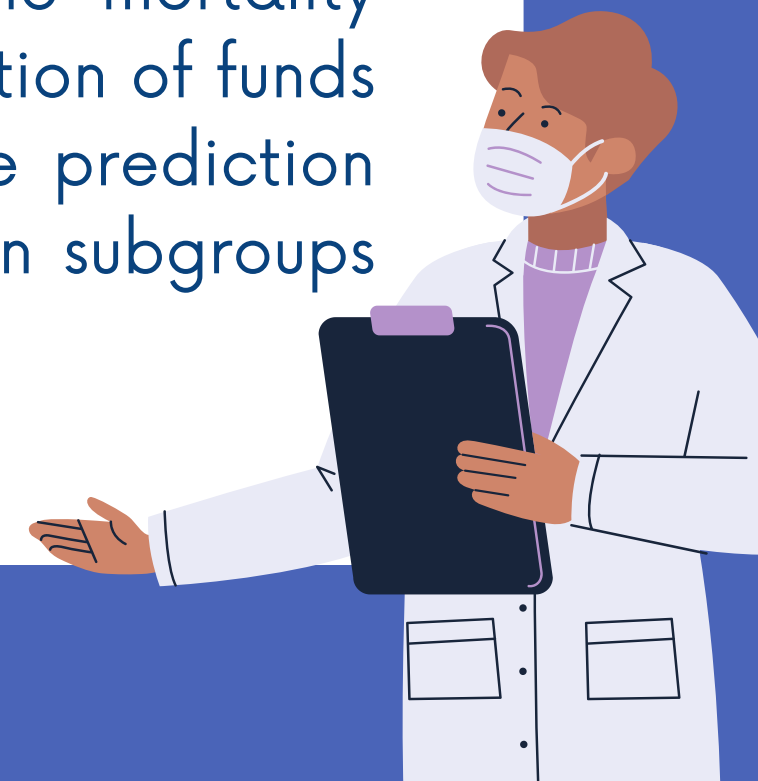
Lasso Regression Coefficients vs Linear Regression Coefficients

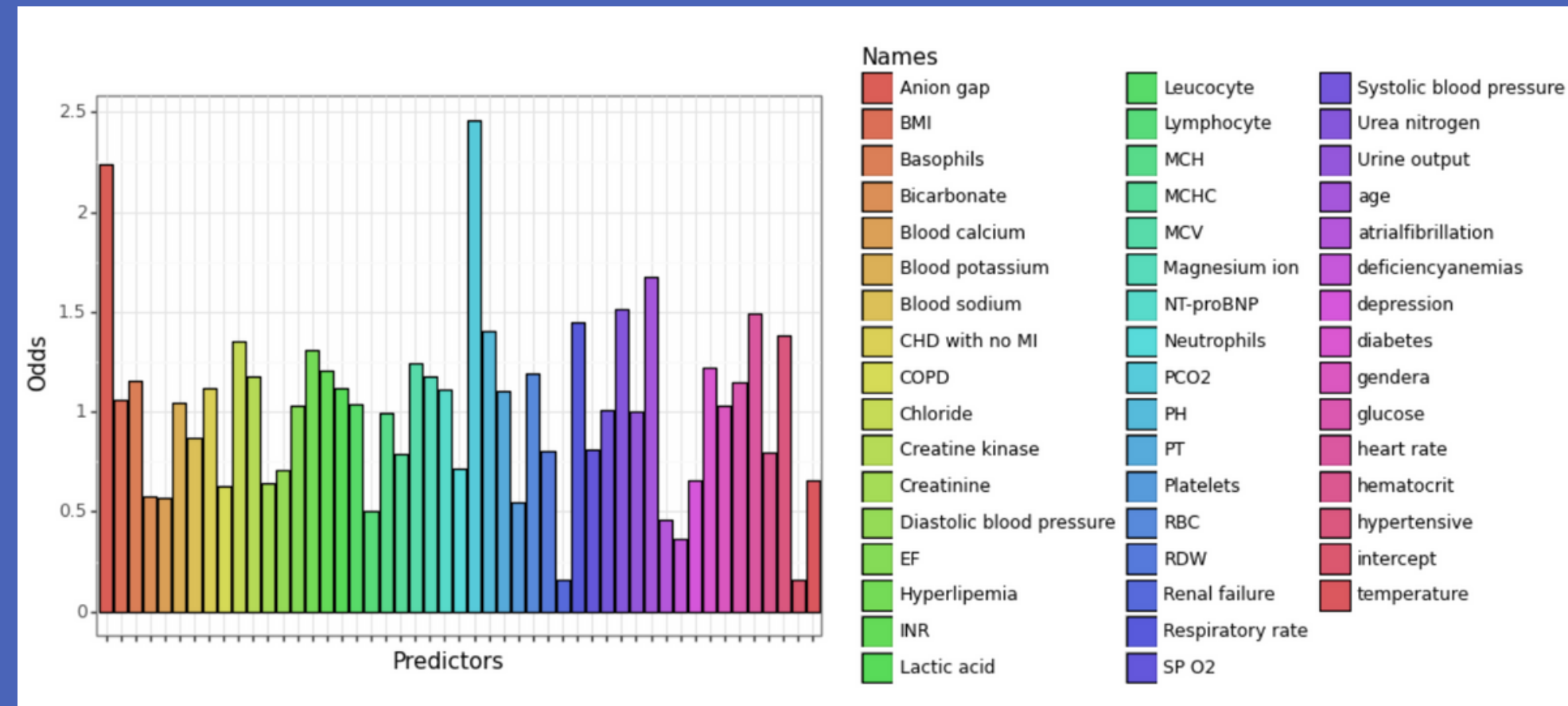


Question 6

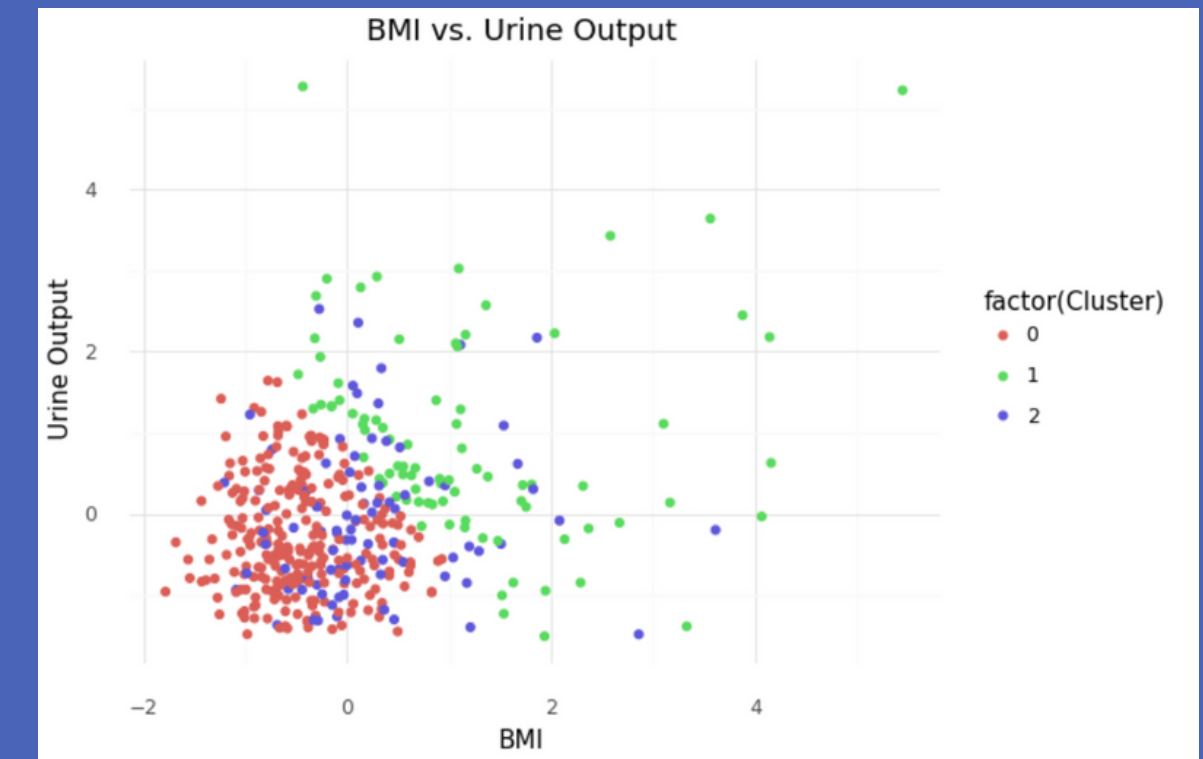
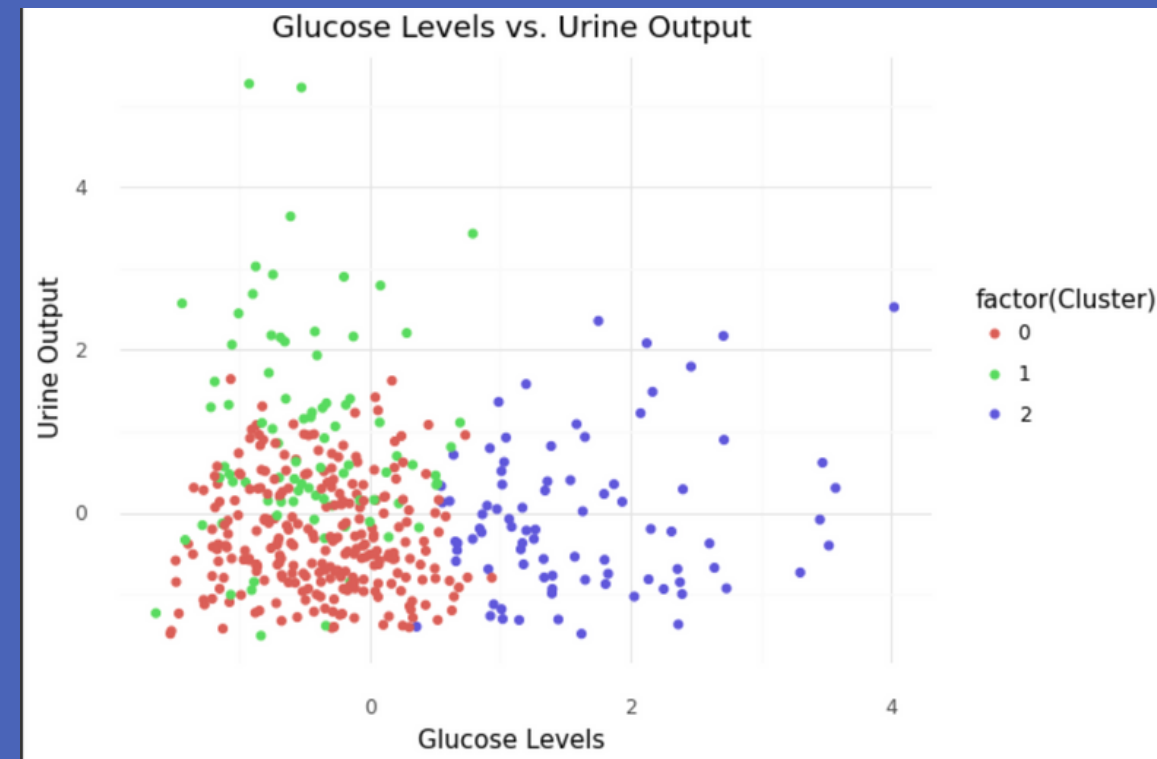
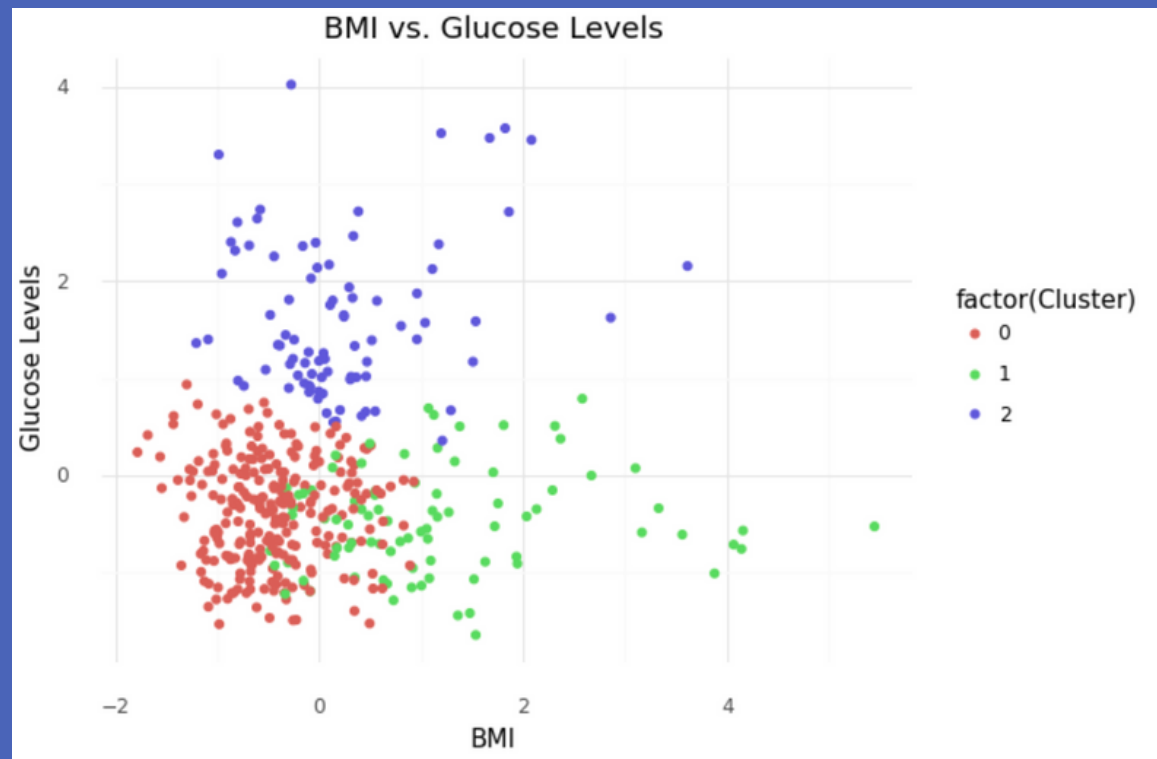
How can healthcare providers and policymakers use the results of this study to improve healthcare outcomes and reduce costs?

A model that could predict hospital mortality could be greatly beneficial for multiple parties, including healthcare providers, policymakers, and patients. First and foremost, with the prediction model, healthcare providers could easily identify patients that are at higher risk. This could allow the providers to pay extra attention to these high-risk patients and provide interventions and treatment at an early stage. Additionally, the ability to identify patients could also help healthcare providers to reduce costs and allocate resources, such as doctor/nurse staffing, medicine, and hospital equipment more efficiently. On the other hand, for policymakers, with the mortality prediction, policymakers could also utilize the results to make decisions on the distribution of funds to hospitals and healthcare providers. Moreover, policymakers could also apply the prediction results to conduct research and push for the demand to develop medicine to certain subgroups facing various medical conditions or diseases.





For example, we could use the logistic regression model from our first response to monitor the patients. According to the coefficient graph from our first response, we can see that feature "**PC O2**" is one of the health conditions that increase the odds of mortality. With that said, healthcare providers could tag those patients with these health conditions as the model indicates that they are at higher risk. In addition, with the predicted data, hospitals could delegate staffing and provide interventions/treatment more efficiently.



Another example, healthcare providers or policymakers could use the clustering model from our second response to study and develop pharmaceuticals for particular subgroups within the patients. Based on the clustering graphs from our second response, we can characterize the patients and classify them into different groups. With the classification, providers and policymakers can dive into patient groups, research their conditions for further purposes, and distribute resources and funds.