Predictive Modeling Project Review I

Group 14

Heart Failure Prediction

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Aim

Create a model for predicting mortality caused by heart failure.

About this dataset

Cardiovascular diseases (CVDs) are the number 1 cause of death globally, taking an estimated 17.9 million lives each year, which accounts for 31% of all deaths worldwide. Heart failure is a common event caused by CVDs and this dataset contains 12 features that can be used to predict mortality by heart failure.

Most cardiovascular diseases can be prevented by addressing behavioural risk factors such as tobacco use, unhealthy diet and obesity, physical inactivity and harmful use of alcohol using population-wide strategies.

People with cardiovascular disease or who are at high cardiovascular risk (due to the presence of one or more risk factors such as hypertension, diabetes, hyperlipidaemia or already established disease) need early detection and management wherein a machine learning model can be of great help.

Objective

- Clean the dataset to remove any existing outliers and extremes.
- Train the model to predict whether the patient is likely to face mortality due to heart failure considering his age, health and other lifestyle choices.
- Train the model to determine patterns amongst the various lifestyle and health aspects that are more likely to cause mortality due to heart failure.
- Perform exploratory analysis to find patterns and insights.

Tool used

To build our model, we will use IBM SPSS Modeler.

DATA DESCRIPTION

Attributes, their descriptions, ranges and number of entries:

Age

Description: Age of patients (years)

Range: [40, 95]

Anaemia

Description: Condition where red blood cells are decreased (Boolean: 0 for false, 1 for true)

0: 170, 1: 129

• Creatinine phosphokinase

Description: Level of CPK enzyme in blood (mcg/L)

Range: [23, 7861]

Diabetes

Description: Whether the patient has diabetes (Boolean: 0 for false, 1 for true)

0: 174, 1: 125

Ejection_fraction

Description: Percentage of blood leaving from heart at each contraction (percentage)

Range: [14, 80]

High_blood_pressure

Description: Whether the patient has high blood pressure (Boolean: 0 for false, 1 for true)

0: 194, 1: 105

Platelets

Description: Platelets in blood (kiloplatelets/mL)

Range: [25100.0, 850000.0]

• Serum creatinine

Description: Level of serum creatinine in blood (mg/dL)

Range: [0.5, 9.4]

Serum_sodium

Description: Level of serum sodium in blood (mEq/L)

Range: [113, 148]

Sex

Description: Man or Woman (Boolean: 0 for man, 1 for woman)

0: 105, 1: 194

Smoking

Description: Whether the patient smokes or not (Boolean: 0 for false, 1 for true)

0: 203. 1: 96

Time

Description: Follow-up period (days)

Range: [4, 285]

Death_event

Description: If the patient dies during follow-up period (Boolean: 0 for false, 1 for true)

0: 203, 1: 96

Data Quantity

· What is the format of the data?

The data is of .csv format.

Which method is used to capture the data?

Here, the method to capture the data is unknown since we are not a part of end-to-end process.

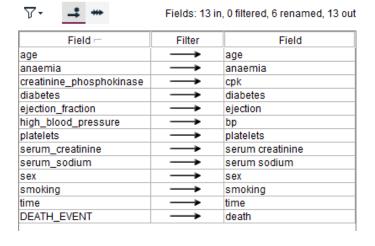
How large is the database?

The database has 299 rows and 13 columns.

Data Quality

Does the data include characteristics relevant to the business perspective?

Yes, the data does include characteristics relevant to the business perspective which can help us achieve our aim.



What data types are present?

Field ⊏	Override	Storage	Input Format
age		Integer	
anaemia		Integer	
creatinine_phosphokinase		Integer	
diabetes		Integer	
ejection_fraction		Integer	
high_blood_pressure		Integer	
platelets		Real	
serum_creatinine		Real	
serum_sodium		Integer	
sex		Integer	
smoking		Integer	
time		Integer	
DEATH_EVENT		Integer	

As we can see, We have integer and real data types and Flag and Continuous measurement levels.

• Did you compute basic statistics for the key attributes? What insight did this provide into the business question?

Yes, I did compute basic statistics for the key attributes.

Field ⊏	Measurement	Min	Max	Mean	Std. Dev	Skewness	Unique	Valid
age		40	95	60.829	11.895	0.424	-	299
anaemia	8 Flag	0	1	-			2	299
		23	7861	581.839	970.288	4.463	-	299
diabetes	8 Flag	0	1				2	299
ejection		14	80	38.084	11.835	0.555		299
☼ bp		0	1	0.351	0.478	0.627		299
platelets		25100.000	850000.000	263358.029	97804.237	1.462		299
serum creatinine		0.500	9.400	1.394	1.035	4.456		299
serum sodium		113	148	136.625	4.412	-1.048		299
sex	8 Flag	0	1				2	299
smoking \$\infty\$	🖁 Flag	0	1				2	299
time tim		4	285	130.261	77.614	0.128		299
death death	8 Flag	0	1				2	299

We can see that some of the data is skewed. We will see what this provides to our business questions further.

• Are you able to prioritize relevant attributes? If not, are business analysts available to provide further insights?

Yes, I did prioritize relevant attributes. All the fields can somehow be the cause of death. Hence, we are assigning the role of Input to all the attributes.

Field	Measurement	Values	Missing	Check	Role
age		[40,95]		None	➤ Input
anaemia	8 Flag	1/0		None	➤ Input
	Continuous	[23,7861]		None	➤ Input
	8 Flag	1/0		None	➤ Input
ejection	Continuous	[14,80]		None	➤ Input
♦ bp	Continuous	[0,1]		None	➤ Input
platelets	Continuous	[25100.0,850000.0]		None	➤ Input
serum creatinine	Continuous	[0.5,9.4]		None	➤ Input
serum sodium	Continuous	[113,148]		None	➤ Input
sex	🖁 Flag	1/0		None	➤ Input
smoking \$\infty\$	🖁 Flag	1/0		None	➤ Input
time	Continuous	[4,285]		None	➤ Input
death death	8 Flag	1/0		None	➤ Input

DATA EXPLORATION

Measurement levels

- Age, cpk, ejection level, blood pressure, platelets, serum creatinine, serum sodium, time and death are assigned Continuous measurement level since they all have values from a continuous range.
- Anaemia, diabetes, sex, smoking and death have been assigned Flag measurement level since they have 0/1 as their values.

Roles

- Currently, all fields have been assigned Input as their roles since we want to
 derive all the patterns among the attributes that will help build a connection
 between lifestyle and heart failure in order to prevent it rather than just predicting if
 the patient is likely to die or not.
- However, we will set death as a target in another Type node later to merely predict whether the patient is likely to die of heart failure of not.

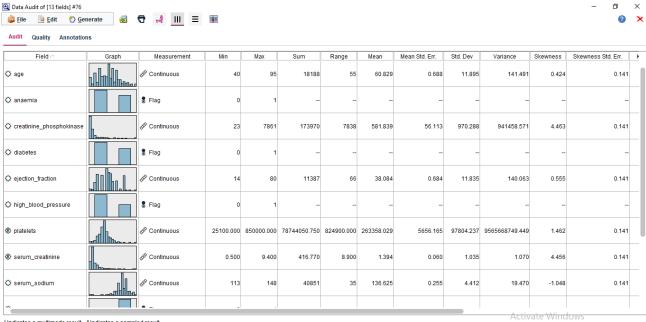
Field - Me:	asurement	Values	Missing	Check	Role
age_transfor Conti	nuous	[-1.760536		None	🔪 Input
creatinine_ph Ø Conti	nuous	[-0.757990		None	🔪 Input
ejection_fracti Conti	nuous	[-2.054378		None	🔪 Input
platelets_tran Oonti	nuous	[-2.653651		None	🔪 Input
Serum_creati Ø Conti	nuous	[-1.171258		None	🔪 Input
Serum_sodiu Conti	nuous	[-2.999999		None	🔪 Input
time_transfor Ø Conti	nuous	[-1.647126		None	🔪 Input
🜣 anaemia_tra 🚦 Flag		1/0		None	🔪 Input
🜣 diabetes_tran 🖁 Flag		1/0		None	🔪 Input
🜣 high_blood_p 🖁 Flag		1/0		None	🔪 Input
Sex_transfor SFlag		1/0		None	🔪 Input
🜣 smoking_tran 🖁 Flag		1/0		None	🔪 Input
DEATH_EVE 8 Flag		1/0		None	Target

Format

indicates that there is maximum 4 digits in that field.

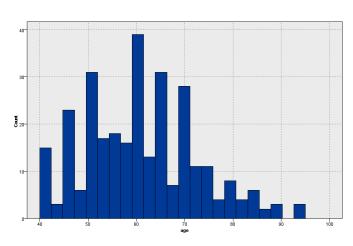
Field ⊢	Format	Justify	Column Width
age_transformed	####.###	Auto	Auto
creatinine_phosphokinase_trans	####.###	Auto	Auto
ejection_fraction_transformed	####.###	Auto	Auto
platelets_transformed	####.###	Auto	Auto
serum_creatinine_transformed	####.###	Auto	Auto
serum_sodium_transformed	####.###	Auto	Auto
time_transformed	####.###	Auto	Auto
anaemia_transformed	####	Auto	Auto
diabetes_transformed	####	Auto	Auto
high_blood_pressure_transformed	####	Auto	Auto
sex_transformed	####	Auto	Auto
smoking_transformed	####	Auto	Auto
DEATH_EVENT_transformed	####	Auto	Auto

Statistics



¹ Indicates a multimode result ² Indicates a sampled result

Age



Min value: 40 Maximum value: 95 Sum of all values: 18188

Range: 55 Mean: 60.829

Mean std. error: 0.688 Std. deviation: 11.895 Variance: 141.491 Skewness: 0.424

Skewness std. error: 0.141

Kurtosis: -0.184

Kurtosis std. error: 0.281

Median: 60 Mode: 60

Valid values: 299

Anaemia

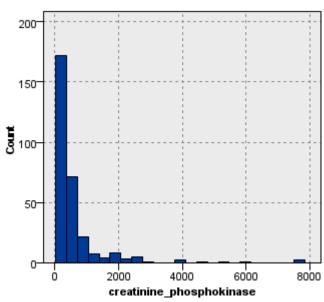
Value /	Proportion	%	Count
0		56.86	170
1		43.14	129

Min value: 0 Maximum value: 1

Mode: 0 Unique: 2

Valid values: 299

• Creatinine_phosphokinase



Min value: 23 Maximum value: 7861 Sum of all values: 173970

Range: 7838 Mean: 581.839

Mean std. error: 56.113 Std. deviation: 970.288 Variance: 941458.571 Skewness: 4.463

Skewness std. error: 0.141

Kurtosis: 25.149

Kurtosis std. error: 0.281

Median: 250 Mode: 582

Valid values: 299

Diabetes

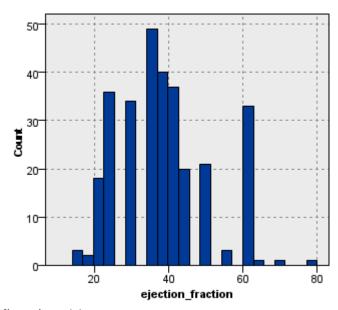
Value /	Proportion	%	Count
0		58.19	174
1		41.81	125

Min value: 0 Maximum value: 1

Mode: 0 Unique: 2

Valid values: 299

Ejection_fraction



Min value: 14 Maximum value: 80 Sum of all values: 11387

Range: 66 Mean: 38.084

Mean std. error: 0.684 Std. deviation: 11.835 Variance: 140.063 Skewness: 0.555

Skewness std. error: 0.141

Kurtosis: 0.041

Kurtosis std. error: 0.281

Median: 38 Mode: 35

Valid values: 299

• High_blood_pressure

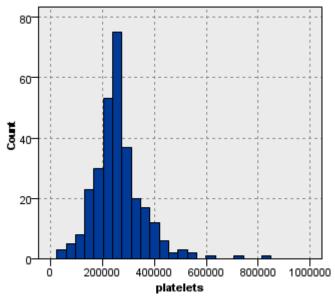
Value /	Proportion	%	Count
0		64.88	194
1		35.12	105

Min value: 0 Maximum value: 1

Mode: 0 Unique: 2

Valid values: 299

Platelets



Min value: 25100.000

Maximum value: 850000.000 Sum of all values: 78744050.750

Range: 824900.000 Mean: 263358.029

Mean std. error: 5656.165 Std. deviation: 97804.237 Variance: 9565668749.449

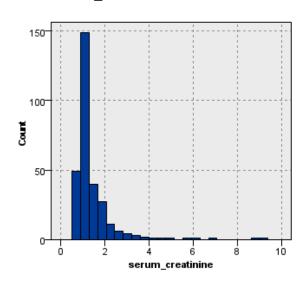
Skewness: 1.462

Skewness std. error: 0.141

Kurtosis: 6.209

Kurtosis std. error: 0.281 Median: 262000.000 Mode: 263358.030 Valid values: 299

Serum_creatinine



Min value: 0.500 Maximum value: 9.400 Sum of all values: 416.770

Range: 8.900 Mean: 1.394

Mean std. error: 0.060 Std. deviation: 1.035 Variance: 1.070 Skewness: 4.456

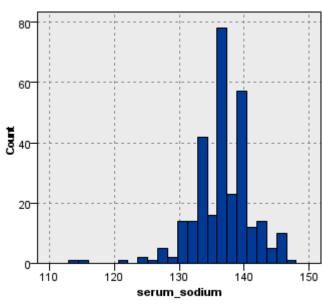
Skewness std. error: 0.141

Kurtosis: 25.828

Kurtosis std. error: 0.281

Median: 1.100 Mode: 1.000 Valid values: 299

Serum_sodium



Min value: 113 Maximum value: 148 Sum of all values: 40851

Range: 35 Mean: 136.625

Mean std. error: 0.255 Std. deviation: 4.412 Variance: 19.470

Skewness: -1.048

Skewness std. error: 0.141

Kurtosis: 4.120

Kurtosis std. error: 0.281

Median: 137 Mode: 136 Valid values: 299

Sex

Value /	Proportion	%	Count
0		35.12	105
1		64.88	194

Min value: 0
Maximum value: 1

Mode: 1 Unique: 2

Valid values: 299

• Smoking

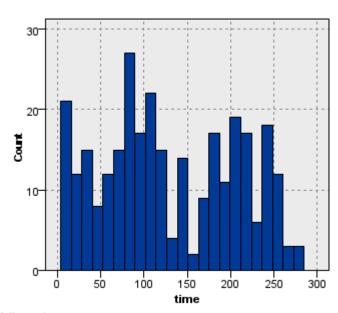
Value /	Proportion	%	Count
0		67.89	203
1		32.11	96

Min value: 0 Maximum value: 1

Mode: 0 Unique: 2

Valid values: 299

Time



Min value: 4

Maximum value: 285 Sum of all values: 38948

Range: 281 Mean: 130.261

Mean std. error: 4.489 Std. deviation: 77.614 Variance: 6023.965 Skewness: 0.128

Skewness std. error: 0.141

Kurtosis: -1.212

Kurtosis std. error: 0.281

Median: 115 Mode: 187 Valid values: 299

Death_event

Value /	Proportion	%	Count
0		67.89	203
1		32.11	96

Min value: 0 Maximum value: 1

Mode: 0 Unique: 2

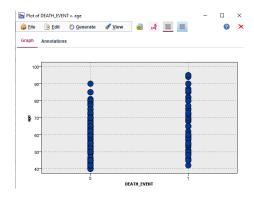
Valid values: 299

What sort of hypothesis have you formed about the data?

 Death rate caused by heart failure might decrease with the increase number of days the patient will be under observation since he would have discharged only after showing improvements.



 Deaths due to heart failure might be high in people aged 40-60 due to their unhealthy lifestyle.



- Deaths due to heart failure are related to all the health conditions in the database.
- Deaths due to heart failure might not be related to sex of the person.

Which attributes seem promising for further analysis?

Age, anaemia, creatinine_phosphokinase, diabetes, ejection_fraction, high_blood_pressure, serum_creatinine, serum_sodium and smoking are seeming to be the most promising for further analysis.

Have your explorations revealed new characteristics about your data?

Explorations helped us understand the data better but haven't revealed any new characteristics yet. It will probably happen during modeling.

How have your explorations changed your initial hypothesis?

Explorations have definitely helped us understand the data better and gave us the clarity of attributes which will be helpful during modeling. However, the initial hypothesis still stands the same.

Can you identify particular subsets of data for later use?

I am planning to segment each of my attributes with death to understand the individual patterns properly and to not miss out on anything.

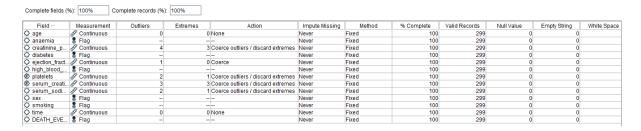
Take another look at your data mining goals. Has this exploration altered the goals?

While the goals remain the same, I definitely believe that additional data might really help to predict deaths caused by heart failures.

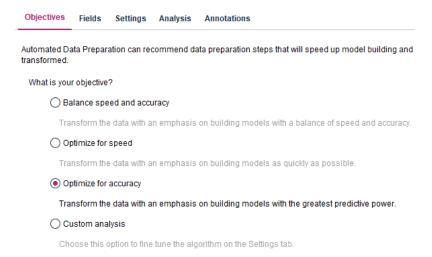
DATA QUALITY

Using data audit node, we can see there are some outliers and extremes that are skewing our data. Let us handle them.

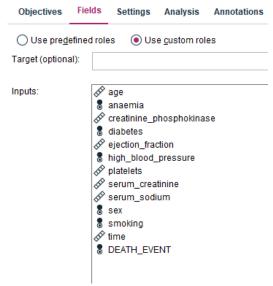
Here, I have coerced the outliers and discarded the extremes and then connected the super node (newly generated) to auto data prep node.



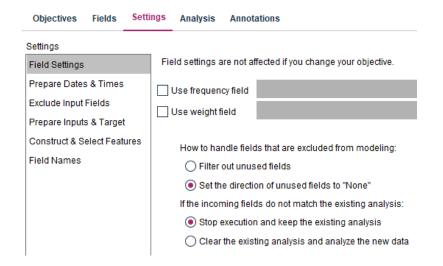
Let us optimize for accuracy rather then speed.



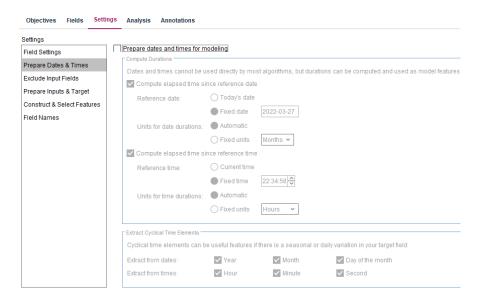
We have taken all fields as input and none as target.



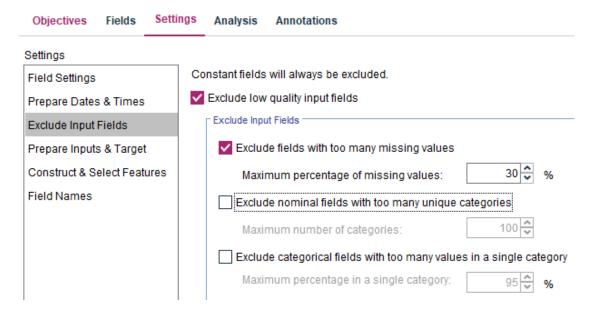
I have made a few changes in settings according to my dataset and requirements as shown.



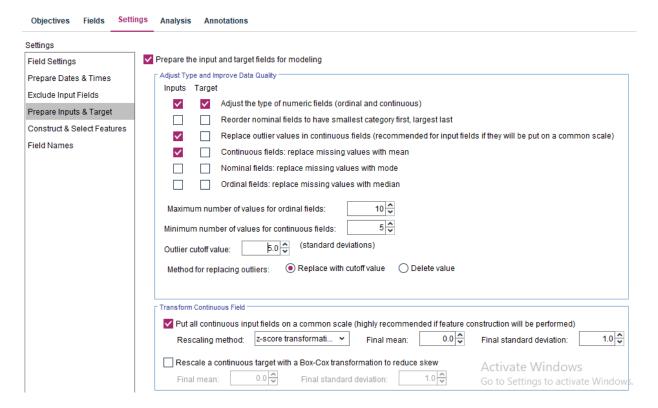
I do not need this because I have no dates and times in my data.



I unselected nominal and ordinal settings because I do not have data with those measurement levels.

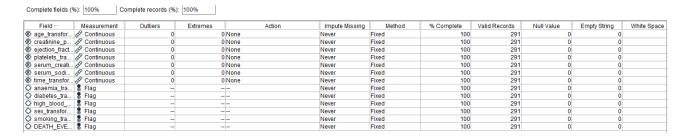


Again, I only kept the ones having my measurement levels in it. I have kept the remaining settings unchanged since they did not require to be changed.



Now connect a data audit node to this auto data prep node.

We can see how the records having extremes have been removed from the data and how the outliers have been coerced.



This shows that the data has been cleaned according to the requirements.

Have you identified missing attributes and blank fields? If so, is there meaning behind such missing values?

No, there are no missing attributes and blank fields. However, the labels in sex attribute were not mentioned but I did my research and added them.

Are there spelling inconsistencies that may cause problems in later merges or transformations?

No, there are no spelling inconsistencies that may cause problems in later merges or transformations since the data has no alphabets or characters.

Have you explored deviations to determine whether they are "noise" or phenomena worth analysing further?

There is some skewness in a few attributes which can later be reduced to clean the noise.

Have you conducted a plausibility check for values? Take notes on any apparent conflicts (such as teenagers with high income levels).

Yes, I did conduct a plausibility check for values. There is no apparent conflict as such.

Have you considered excluding data that has no impact on your hypothesis? Yes, I have. It concluded with the fact that all data is important for the analysis since each and every attribute might add up to the death caused by heart failure. So, I will not exclude any data.

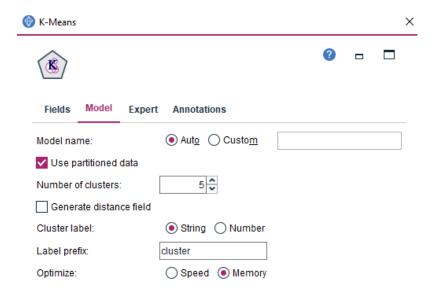
Are the data stored in flat files? If so, are the delimiters consistent among files? Does each record contain the same number of fields?

Yes. The delimiters are consistent among files. Each record contains the same number of fields.

We have managed to successfully clean the data and will now move forward towards modeling.

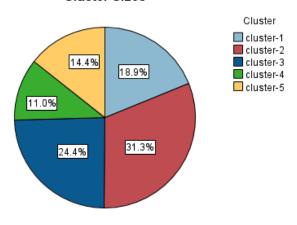
MODELING

A. Segmentation (without target)

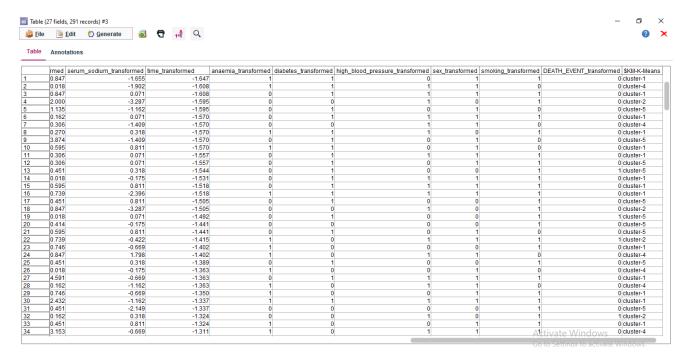


Here, we will use K-Means node and will create 5 clusters. We can see that the clusters have been created.





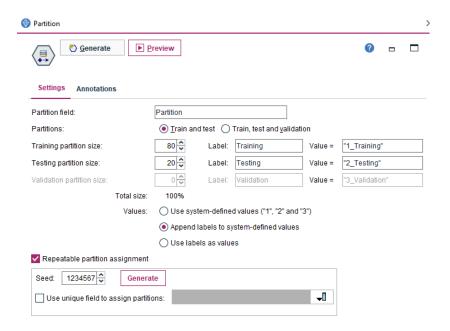
Size of Smallest Cluster	32 (11%)
Size of Largest Cluster	91 (31.3%)
Ratio of Sizes: Largest Cluster to Smallest Cluster	2.84



These clusters will group the data for us, as to which group of lifestyle patterns is more likely to experience a heart failure.

B. Supervised (with target)

Let us first partition our data in 80% training and 20% testing.



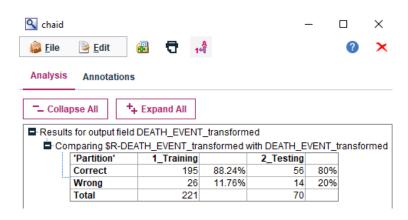
Now, let us check the accuracy of all supervised models and see some of the best:

Analysis of Auto Classifier:



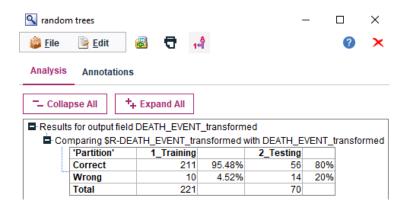
We are getting 78.57% accuracy.

Analysis of CHAID:



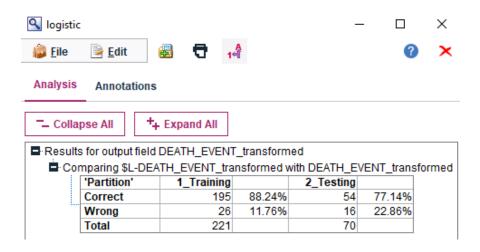
We are getting 80% accuracy.

Analysis of Random Trees:



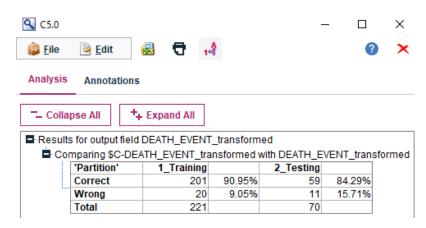
We are getting 80% accuracy.

Analysis of Logistic regression:



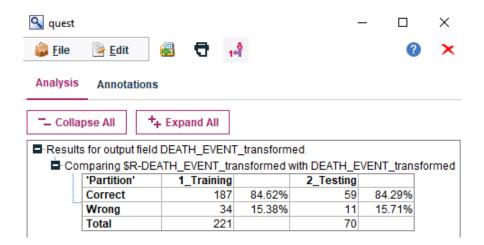
We are getting 77.14% accuracy.

Analysis of C5.0:



We are getting 84.29% accuracy.

Analysis of Quest:



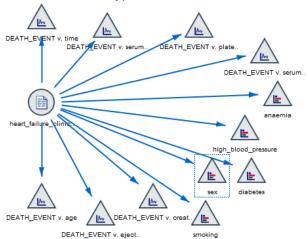
We are getting 84.29% accuracy.

Here, we are getting highest accuracy on testing data of 84.29% by C5.0 and Quest models.

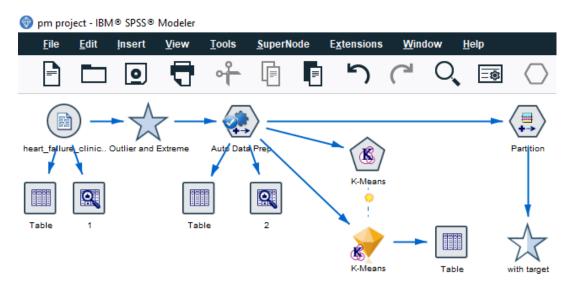
We can use this model to predict whether the person is likely to face mortality due to heart failure or not as per his lifestyle data/records.

FINAL STREAM

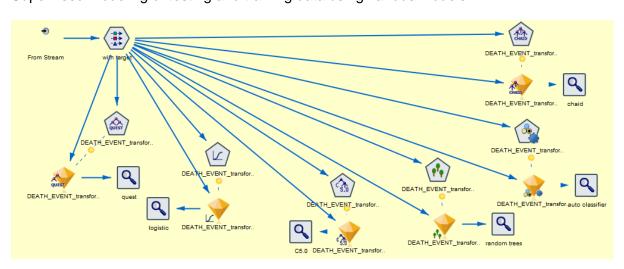
Visualization for hypothesis:



Data cleaning and segmentation using K-Means:



Supervised modeling of testing and training data using various models:



CONCLUSION

- 1. We have managed to successfully clean the data.
- 2. We segmented our data using K-Means model to divide it in 5 different clusters to predict and study the patterns of lifestyle that contributes to the heart failure.
- 3. We compared all the supervised modelling nodes where C5.0 and Quest models give the best accuracy of 84.29%.
- 4. We can further use these models for deployment and to predict the mortality caused by heart failure.