

VarSelLCM

Variable Selection for Model-Based Clustering of Mixed-Type Data Set with Missing Values.

References:

- Marbac, M. and Sedki, M. (2017), Variable selection for model-based clustering using the integrated complete-data likelihood, *Statistics and Computing*, Volume 27, Issue 4, pp 1049–1063.
- Marbac, M., Patin, E. and Sedki, M. (2018), Variable selection for mixed data clustering: Application in human population genomics, *Arxiv* 1703.02293.

Introduction

VarSelLCM permits a full model selection (detection of the relevant features for clustering and selection of the number of clusters) in model-based clustering, according to classical information criteria (BIC, MICL or AIC).

Data to analyzed can be composed of continuous, integer and/or categorical features. Moreover, missing values are managed, without any pre-processing, by the model used to cluster with the assumption that values are missing completely at random. Thus, *VarSelLCM* can also be used for data imputation via mixture models.

An R-Shiny application is implemented to easily interpret the clustering results

Mixed-type data analysis

This section performs the whole analysis of the *Heart* data set. *Warning continuous features must be stored in numeric, integer features must be stored in integer and categorical features must be stored in factor.*

```
library(VarSelLCM)
# Data loading
data("heart")
head(heart)
```

	Age	Sex	ChestPainType	RestBloodPressure	SerumCholestoral
1	70	1	4	130	322
2	67	0	3	115	564
3	57	1	2	124	261
4	64	1	4	128	263
5	74	0	2	120	269
6	65	1	4	120	177

	FastingBloodSugar	ResElectrocardiographic	MaxHeartRate	ExerciseInduced
1	0	2	109	0
2	0	2	160	0
3	0	0	141	0
4	0	0	105	1
5	0	2	121	1
6	0	0	140	0

	Slope	MajorVessels	Thal	Class
1	2	3	3	2
2	2	0	7	1
3	1	0	7	2

```

4      2      1      7      1
5      1      1      3      1
6      1      0      7      1

```

Clustering is performed with variable selection. Model selection is done with BIC because $n \gg d$. The number of components is between 1 and 4. Do not hesitate to use parallelisation (here only two cores are used).

```

# Add a missing value artificially (just to show that it works!)
heart[1,1] <- NA
# Clustering with variable selection and a number of cluster between 1 and 4
# Model selection is BIC (to use MICL, the option must be specified)
out <- VarSelCluster(heart[, -13], 1:4, nbcores = 2)

```

Now, all the results can be analyzed by the Shiny application...

```

# Start the shiny application
VarSelShiny(out)

```

... but this analysis can also be done on R.

To get a summary of the selected model.

```

# Summary of the best model
summary(out)

```

Data set:

```

Number of individuals: 270
Number of continuous variables: 3
Number of count variables: 1
Percentile of missing values for the integer variables: 0.37
Number of categorical variables: 8

```

Model:

```

Number of components: 2
Model selection has been performed according to the BIC criterion
Variable selection has been performed, 8 ( 66.67 % ) of the variables are relevant for clustering

```

Information Criteria:

```

loglike: -6403.136
AIC:     -6441.136
BIC:     -6509.506
ICL:     -6638.116

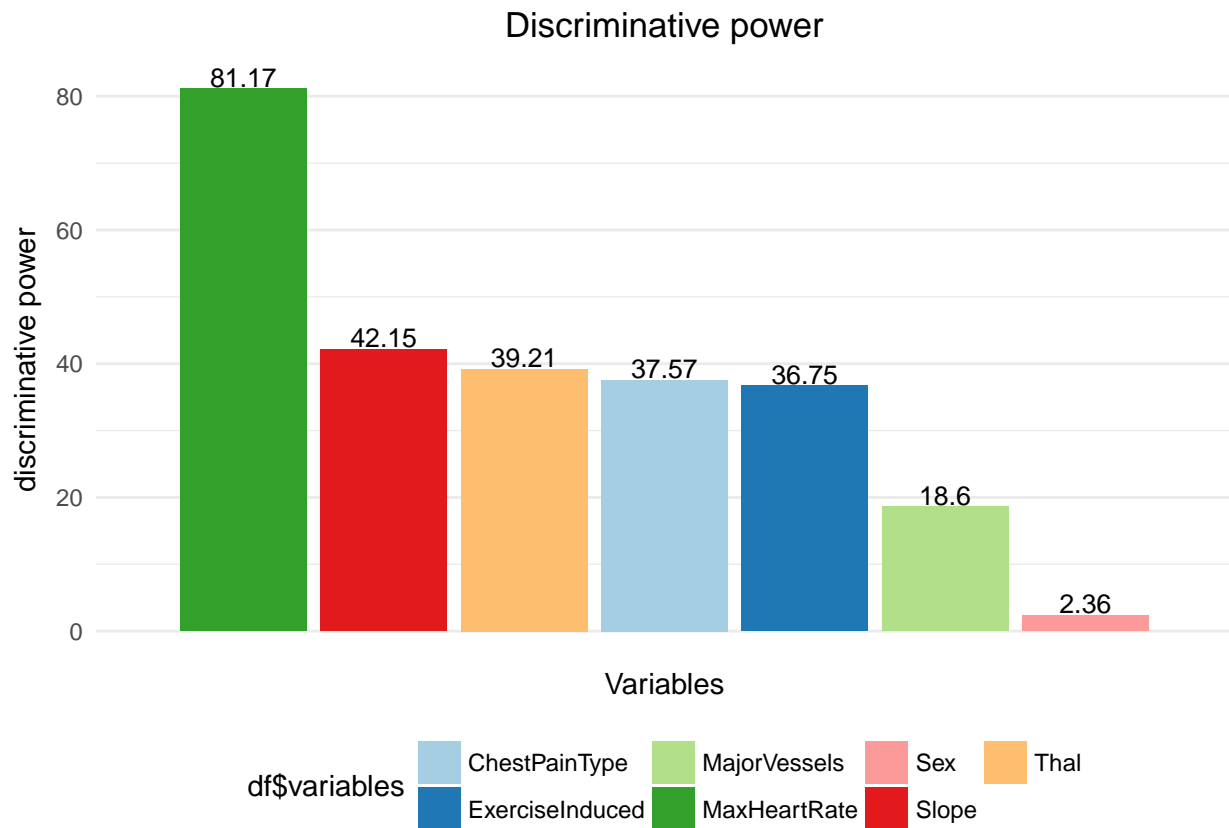
```

Model interpretation should focus on the most discriminative variables. These variables can be found with the following plot.

```

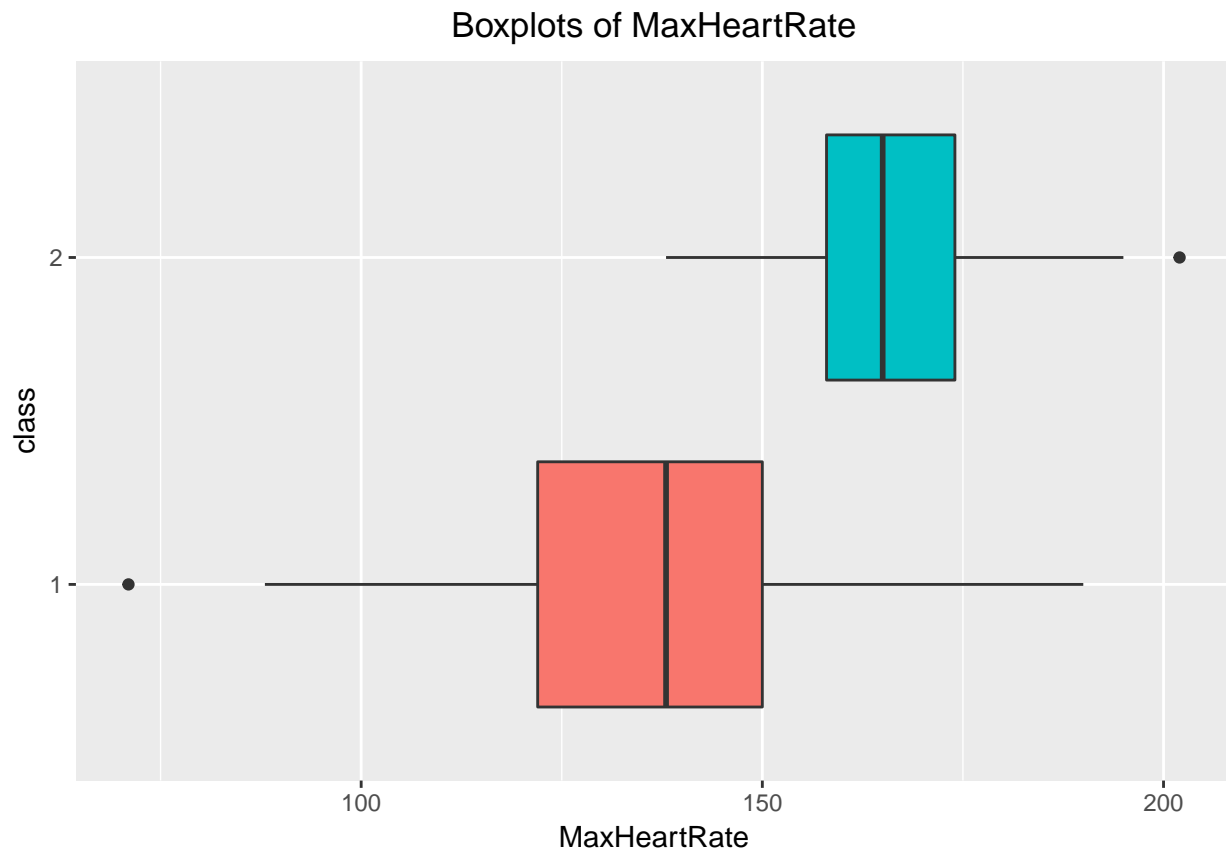
# Discriminative power of the variables (here, the most discriminative variable is MaxHeartRate)
plot(out, type="bar")

```



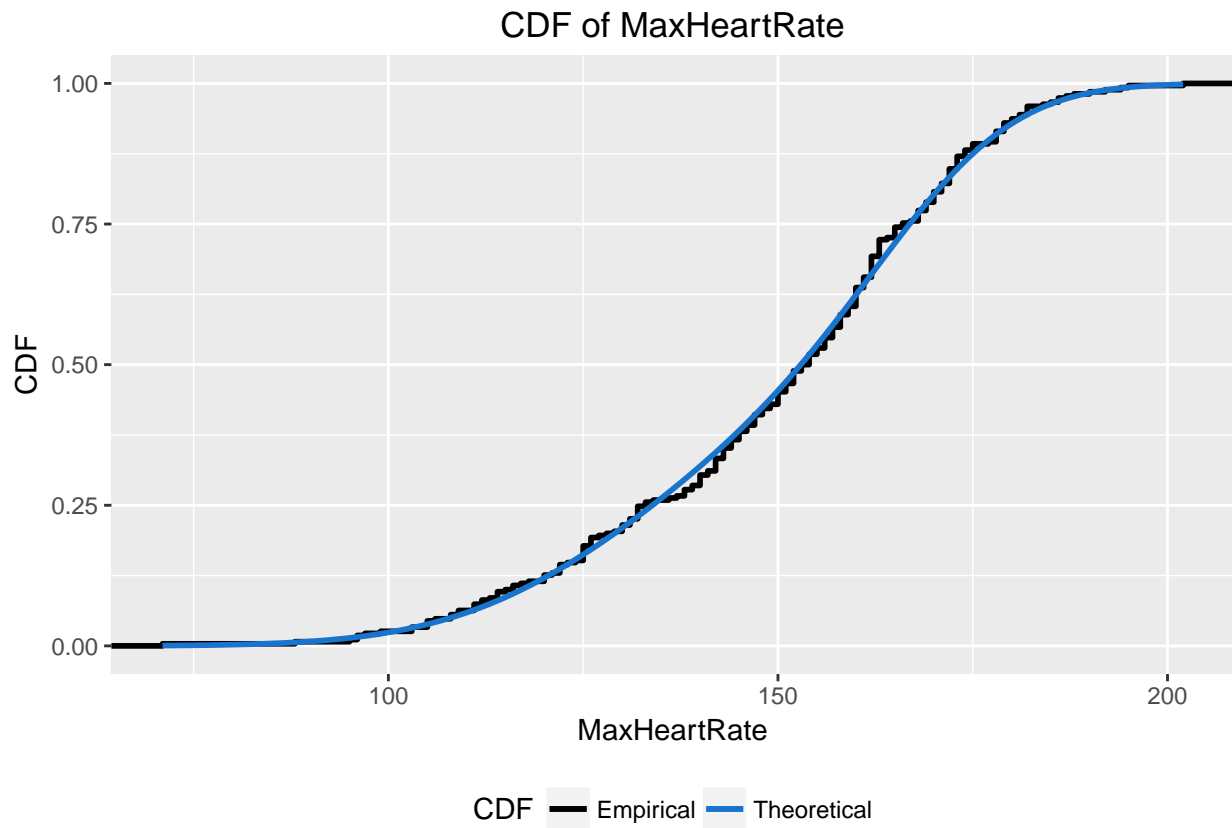
Interpretation of the most discriminative variable is based on its distribution per cluster.

```
# Boxplot for continuous (or interger) variable
plot(out, y="MaxHeartRate", type="boxplot")
```



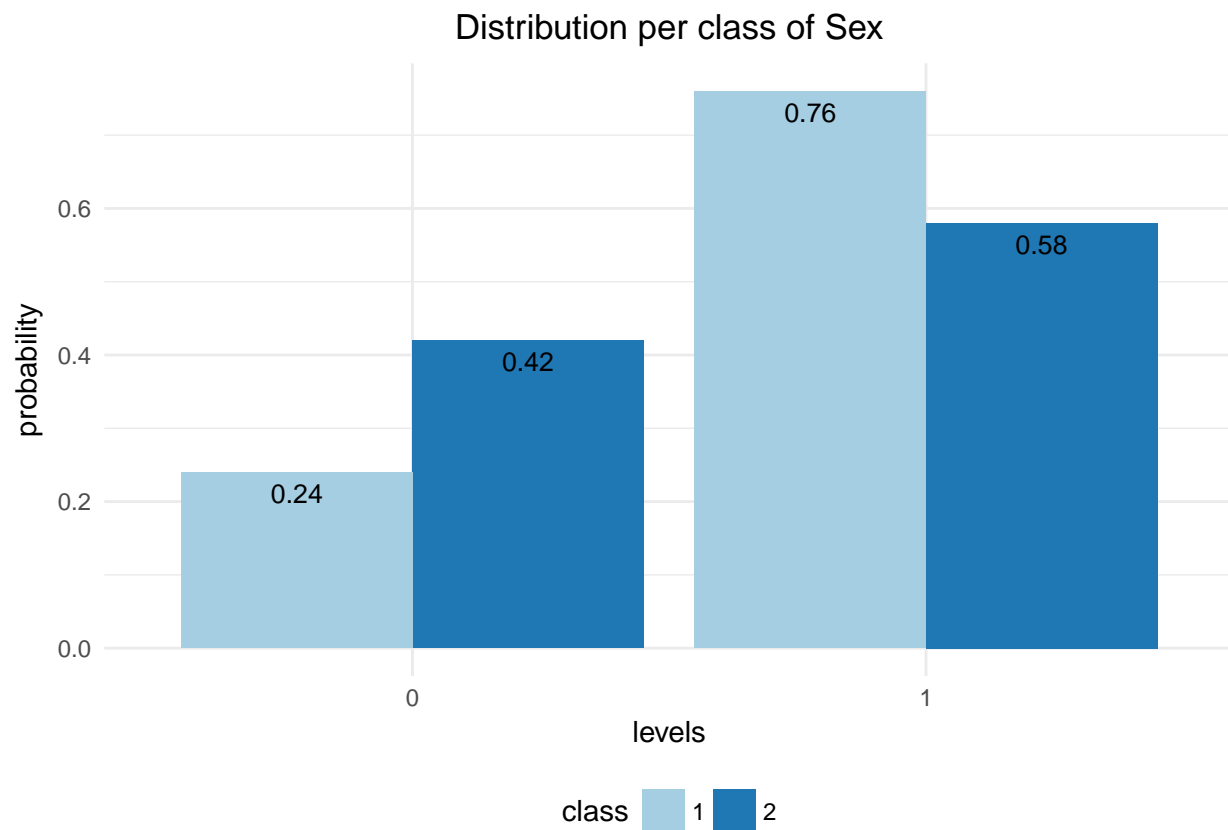
We can check that the distribution used to cluster is relevant.

```
# Empirical and theoretical distributions (to check that clustering is pertinent)  
plot(out, y="MaxHeartRate", type="cdf")
```



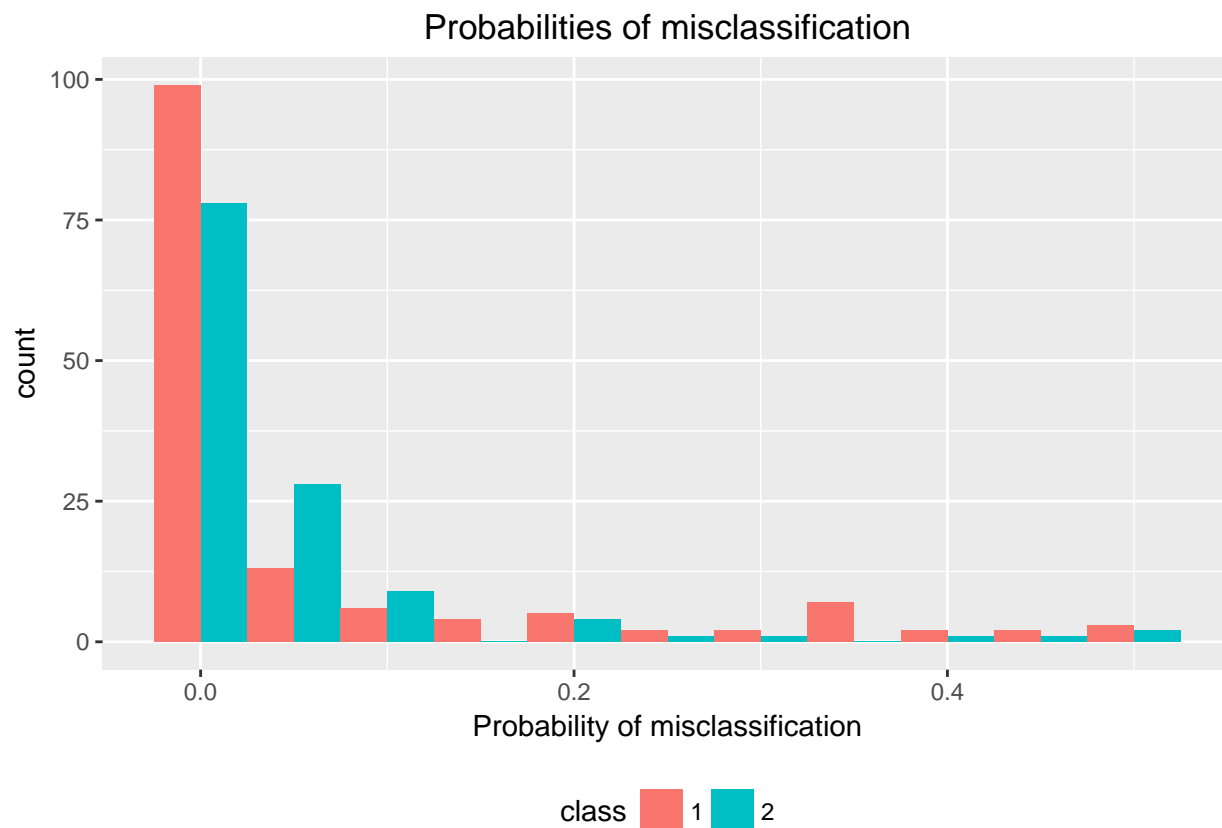
Interpretation of a categorical variable is based on its distribution per cluster.

```
# Summary of categorical variable  
plot(out, y="Sex")
```



Interpretation of the cluster overlaps by using the probabilities of missclassification.

```
# Summary of the probabilities of missclassification  
plot(out, type="probs-class")
```



Missing values can be imputed.

Imputation by posterior mean for the first observation

```
not.imputed <- heart[1,-13]
```

```
imputed <- VarSelImputation(out)[1,]
```

```
rbind(not.imputed, imputed)
```

	Age	Sex	ChestPainType	RestBloodPressure	SerumCholestoral
1	NA	1	4	130	322
2	58.11329	1	4	130	322

	FastingBloodSugar	ResElectrocardiographic	MaxHeartRate	ExerciseInduced
1	0	2	109	0
2	0	2	109	0

	Slope	MajorVessels	Thal
1	2	3	3
2	2	3	3