# VarSelLCM

## Variable Selection for Model-Based Clustering of Mixed-Type Data Set with Missing Values 2018-08-27

#### **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, Journal of Classification, to appear.

### 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

#### Clustering

This section performs the whole analysis of the *Heart* data set. Warning the univariate margin distribution are defined by class of the features: numeric columns imply Gaussian distributions, integer columns imply Poisson distribution while factor (or ordered) columns imply multinomial distribution

```
library(VarSelLCM)
```

Attaching package: 'VarSelLCM' The following object is masked from 'package:stats':

predict

```
# Data loading:
# x contains the observed variables
# z the known status (i.e. 1: absence and 2: presence of heart disease)
data(heart)
ztrue <- heart[,"Class"]
x <- heart[,-13]
# Add a missing value artificially (just to show that it works!)
x[1,1] <- NA</pre>
```

Clustering is performed with variable selection. Model selection is done with BIC because the number of observations is large (compared to the number of features). The number of components is between 1 and 3. Do not hesitate to use parallelization (here only two cores are used).

```
# Cluster analysis without variable selection
res_without <- VarSelCluster(x, gvals = 1:3, vbleSelec = FALSE, crit.varsel = "BIC")
# Cluster analysis with variable selection (with parallelisation)</pre>
```

```
res_with <- VarSelCluster(x, gvals = 1:3, nbcores = 2, crit.varsel = "BIC")</pre>
```

Comparison of the BIC for both models: variable selection permits to improve the BIC

BIC(res\_without)

[1] -6516.216
BIC(res\_with)

[1] -6509.506

Comparison of the partition accuracy. ARI is computed between the true partition (ztrue) and its estimators. ARI is an index between 0 (partitions are independent) and 1 (partitions are equals). Variable selection permits to improve the ARI. Note that ARI cannot be used for model selection in clustering, because there is no true partition.

ARI(ztrue, fitted(res\_without))

[1] 0.2218655

ARI(ztrue, fitted(res\_with))

[1] 0.2661321

To obtained the partition and the probabilities of classification

# Estimated partition
fitted(res\_with)

```
# Estimated probabilities of classification
head(fitted(res_with, type="probability"))
```

class-1 class-2 [1,] 8.261914e-06 0.9999917 [2,] 3.665402e-01 0.6334598 [3,] 8.244720e-01 0.1755280 [4,] 4.443478e-08 1.0000000 [5,] 3.884891e-03 0.9961151 [6,] 4.521857e-02 0.9547814

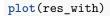
To get a summary of the selected model.

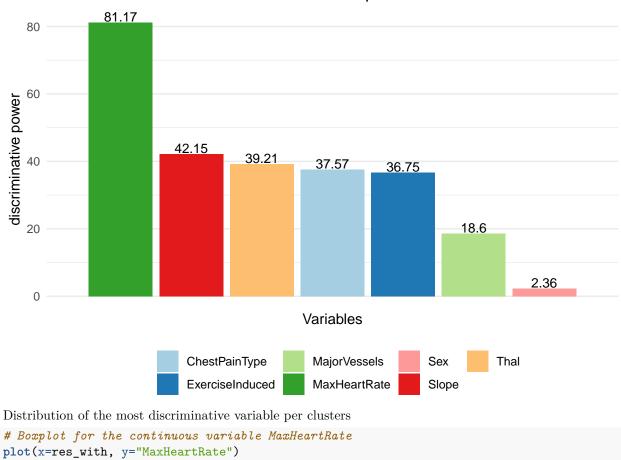
```
# Summary of the best model
summary(res_with)
```

Model:

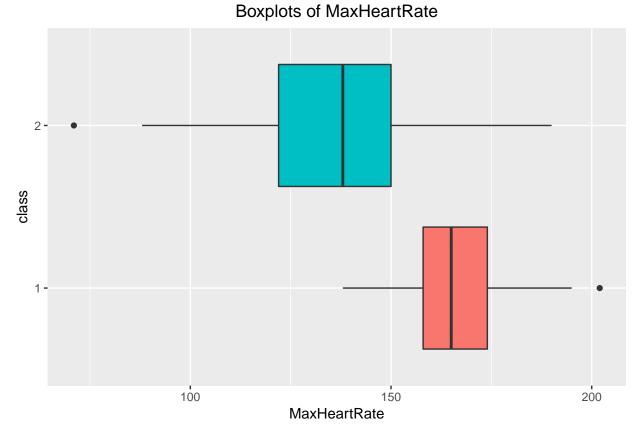
```
Number of components: 2
Model selection has been performed according to the BIC criterion
Variable selection has been performed, 8 ( 66.67 % ) of the variables are relevant for clustering
```

Discriminative power of the variables (here, the most discriminative variable is MaxHeartRate). The greater this index, the more the variable distinguishes the clusters.





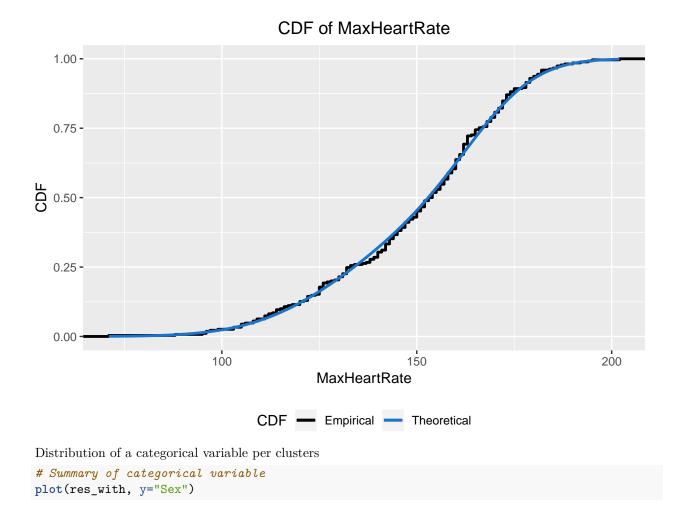
Discriminative power



Empirical and theoretical distributions of the most discriminative variable (to check that the distribution is well-fitted)

# Empirical and theoretical distributions (to check that the distribution is well-fitted)
plot(res\_with, y="MaxHeartRate", type="cdf")

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```
0.76
   0.6
                                                          0.58
probability
  0.4
                   0.42
                                     0.24
   0.2
   0.0
                             0
                                                                    1
                                              levels
                                        class
                                                         2
To have details about the selected model
# More detailed output
print(res_with)
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
To print the parameters
# Print model parameter
coef(res_with)
An object of class "VSLCMparam"
Slot "pi":
```

## Distribution per class of Sex

```
class-1
            class-2
0.4778889 0.5221111
Slot "paramContinuous":
An object of class "VSLCMparamContinuous"
Slot "pi":
numeric(0)
Slot "mu":
                   class-1 class-2
RestBloodPressure 131.3444 131.3444
SerumCholestoral 249.6593 249.6593
MaxHeartRate
                  165.2587 135.4166
Slot "sd":
                   class-1 class-2
RestBloodPressure 17.82850 17.82850
SerumCholestoral 51.59043 51.59043
MaxHeartRate
                  13.14847 20.98135
Slot "paramInteger":
An object of class "VSLCMparamInteger"
Slot "pi":
numeric(0)
Slot "lambda":
     class-1 class-2
Age 50.32062 58.11338
Slot "paramCategorical":
An object of class "VSLCMparamCategorical"
Slot "pi":
numeric(0)
Slot "alpha":
$Sex
                0
                          1
class-1 0.4166331 0.5833669
class-2 0.2358078 0.7641922
$ChestPainType
                            2
                                      3
                 1
class-1 0.05752325 0.28954333 0.4223078 0.2306256
class-2 0.08922307 0.03291638 0.1738641 0.7039965
$FastingBloodSugar
                0
                          1
class-1 0.8518519 0.1481481
class-2 0.8518519 0.1481481
$ResElectrocardiographic
                                      2
                0
                            1
```

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```
class-1 0.4851852 0.007407407 0.5074074
class-2 0.4851852 0.007407407 0.5074074
$ExerciseInduced
                0
                            1
class-1 0.9128089 0.08719107
class-2 0.4484661 0.55153394
$Slope
                1
                           2
                                      3
class-1 0.7599019 0.1933842 0.04671396
class-2 0.2266429 0.6884277 0.08492940
$MajorVessels
                0
                                      2
                           1
                                                   3
class-1 0.7915987 0.1402686 0.05987847 0.008254255
class-2 0.4104421 0.2830471 0.17928553 0.127225327
$Thal
                                        7
                3
                              6
class-1 0.8302545 5.596129e-10 0.1697455
class-2 0.3183107 9.931192e-02 0.5823774
Probabilities of classification for new observations
# Probabilities of classification for new observations
predict(res_with, newdata = x[1:3,])
          class-1
                    class-2
[1,] 8.635968e-06 0.9999914
[2,] 3.768809e-01 0.6231191
[3,] 8.307883e-01 0.1692117
The model can be used for imputation (of the clustered data or of a new observation)
# Imputation by posterior mean for the first observation
not.imputed <- x[1,]
```

```
imputed <- VarSelImputation(res_with, x[1,], method = "sampling")
rbind(not.imputed, imputed)</pre>
```

Age Sex ChestPainType RestBloodPressure SerumCholestoral 4 322 1 NA 1 130 2 69 1 4 130 322 FastingBloodSugar ResElectrocardiographic MaxHeartRate ExerciseInduced 0 2 109 0 1 2 2 0 109 0 Slope MajorVessels Thal 1 2 3 3 2 2 3 3

#### Shiny application

All the results can be analyzed by the Shiny application...

```
# Start the shiny application
VarSelShiny(res_with)
```

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