# Package 'plrs'

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<b>Title</b> Piecewise Linear Regression Splines (PLRS) for the association between DNA copy number and gene expression
Author Gwenael G.R. Leday
Maintainer Gwenael G.R. Leday to <gleday@few.vu.nl></gleday@few.vu.nl>
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<b>Description</b> The present package implements a flexible framework for modeling the relationship between DNA copy number and gene expression data using Piecewise Linear Regression Splines (PLRS).
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R topics documented:
plrs-package       2         criteria       3         modify.conf       4         neveCN17       5         neveGE17       6         plot-methods       6         plrs       9         plrs-class       9         plrs.cb       10         plrs.select       1         plrs.select-class       1         plrs.series       1
ризосию

2 plrs-package

Index																					20
	predict.plrs		•							 •					•						18
	plrs.test																				17
	plrs.sim																				16
	plrs.series-class																				15

plrs-package

Piecewise Linear Regression Splines (PLRS) for the association between DNA copy number and mRNA expression

### Description

The present package implements a framework for modeling the relationship between DNA copy number and gene expression data using Piecewise Linear Regression Splines (PLRS). It includes (point and interval) estimation, model selection and testing procedures for such models (possibly under biologically motivated constraints).

#### **Details**

The use of the present package can be divided into two approaches:

1. Analysis of a single DNA-mRNA relationship

```
Main functions are:
```

```
plrs: Fit a single plrs model.
plrs.select: Model selection based on AIC, AICC, OSAIC or BIC.
plrs.test: Likelihood ratio test for a given plrs model.
plrs.cb: Confidence bands for a plrs model.
```

2. Analysis of multiple DNA-mRNA relationships sequentially

Main function is:

plrs.series: point and interval estimation, model selection and testing of DNA-mRNA association for a series of arrays.

Note: This function extend the aforementioned univariate analysis genomewise in the same spirit as some functions of the **limma** package do.

#### Author(s)

```
Gwenael G.R. Leday
```

Maintainer: Gwenael G.R. Leday <g.g.r.leday@vu.nl>

criteria 3

#### References

Leday GGR, Van der Vaart AW, Van Wieringen WN, Van de Wiel MA. Modeling association between DNA copy number and gene expression with constrained piecewise linear regression splines. Accepted for publication. *Ann Appl Stat.* (2012).

criteria

Compute AIC, AICC, BIC and OSAIC for a given plrs model.

### **Description**

Extract AIC, AICC, BIC and OSAIC from an object of class plrs-class.

### Usage

```
criteria(obj, crit = "all")
```

### Arguments

obj	object of class plrs-class
crit	A character (vector) among "aic", "aicc", "bic", "osaic" or "all".

#### Value

A list with the following components (if specified):

aic	Akaike's information criterion
aicc	Small sample correction of AIC
bic	Bayesian Information Criterion

osaic One-Sided AIC. See Hughes and King (2003) for more details.

### Author(s)

```
Gwenael G.R. Leday < g.g.r.leday@vu.nl>
```

#### References

Hughes, A. W. and King, M. L. (2003). Model selection using AIC in the presence of one-sided information. *J Stat Plan Infer*, 115(2): 397 411.

4 modify.conf

#### **Examples**

```
# Simulate data
sim <- plrs.sim(n=80, states=4, sigma=0.5)
# Fit
model <- plrs(expr=sim$expr, cghseg=sim$seg, cghcall=sim$cal)
criteria(model)</pre>
```

modify.conf

Modify the configuration (of calls) of the plrs model

### **Description**

This function changes the discrete copy number values for a given gene in order to force a minimum number of observations per state.

### Usage

```
modify.conf(cghcall, min.obs = 3, discard = TRUE)
```

### **Arguments**

cghcall	Vector of called values
min.obs	Minimum number of observations per state
discard	Logical. Whether discrete states with few observations should be discarded from analysis.

#### **Details**

Consider that the number of observations of a given state is lower than min.obs, then:

- if discard = FALSE, observations are not discarded and a rearrangement of called values is carried out as follows. The "normal" copy number state is taken as a reference. If the minimum number of observations is not obtained, "losses" will be merged to "normals", "gains" to "normals" and "amplifications" to "gains". Note that this modifies the configuration of the model. Thus, after fitting a model using plrs, original and modified data are stored in the resulting plrs-class object, respectively under slots data and mdata.
- if discard = TRUE, states for which the number of observations is lower than min.obs are discarded (replaced by NAs).

#### Value

val Vector of new called values

neveCN17

### Note

This function is implemented within function plrs and plrs.series.

### Author(s)

```
Gwenael G.R. Leday < g.g.r.leday@vu.nl>
```

### **Examples**

```
called <- sample(c(rep(-1,5),rep(0,15),rep(1,2),rep(2,1)))
table(called)
table(modify.conf(called, min.obs=3))</pre>
```

neveCN17

Copy number for chromosome 17.

### Description

Preprocessed copy number data of Neve et al. (2006) for chromosome 17.

### Usage

neveCN17

#### **Format**

An object of class cghCall

#### **Source**

M. Neve et al. in Gray Lab at LBL. Neve2006: expression and CGH data on breast cancer cell lines. R package version 0.1.10.

#### References

Neve, R.M. et al. (2006). A collection of breast cancer cell lines for the study of functionally distinct cancer subtypes. *Cancer cell*, 10, 515-527.

### **Examples**

```
data(neveCN17)
dim(neveCN17)
head(fData(neveCN17))
```

6 plot-methods

neveGE17

mRNA expression for chromosome 17.

#### Description

Normalized gene expression data of Neve et al. (2006) for chromosome 17.

#### Usage

neveGE17

#### **Format**

An object of class ExpressionSet

#### **Source**

M. Neve et al. in Gray Lab at LBL. Neve2006: expression and CGH data on breast cancer cell lines. R package version 0.1.10.

#### References

Neve, R.M. et al. (2006). A collection of breast cancer cell lines for the study of functionally distinct cancer subtypes. *Cancer cell*, 10, 515-527.

#### **Examples**

```
data(neveGE17)
dim(neveGE17)
head(fData(neveGE17))
```

plot-methods

Plot functions in package 'plrs'

#### **Description**

Methods plot in package 'plrs'

### Usage

```
## S3 method for class 'plrs'
plot(x, col.line = "black", col.pts = c("red", "blue", "green2", "green4"),
col.cb = "yellow", xlim = c(floor(min(x@data$cghseg)), ceiling(max(x@data$cghseg))),
ylim = c(floor(min(x@data$expr)), ceiling(max(x@data$expr))),
pch = 16, lwd=4, cex = 1.2, xlab="", ylab="", main = "",
add = FALSE, lty = 1, lin = FALSE, ...)
```

plrs 7

#### **Arguments**

x	An object of class plrs-class or plrs. select-class
col.line	Color of the fitted line
col.pts	Vector of length 4, for colors associated with each state
col.cb	Color for the confidence band
xlim	The x limits of the plot
ylim	The y limits of the plot
pch	See par
lwd	See par
cex	See par
xlab	Title of the x-axis
ylab	Title of the y-axis
main	Main title for the plot
add	If the plot should be added to the current device. Default is FALSE
lty	See par
lin	Logical. Whether the simple linear model should also be plotted
	Other arguments, see par

### **Details**

plot.plrs plots the observed points, the fitted line and potentially the confidence band.

#### Methods

```
signature(x = "plrs") Plot observed points and the fitted line
signature(x = "plrs.select") Plot observed points and the fitted line of the selected model.
```

### Author(s)

```
Gwenael G.R. Leday <g.g.r.leday@vu.nl>
```

### Description

The function fits a piecewise linear regression spline to explain gene expression by the segmented DNA copy number. The called copy number values are used as a template for model building.

8 plrs

#### Usage

```
plrs(expr, cghseg, cghcall=NULL, probloss = NULL, probnorm = NULL,
probgain = NULL, probamp = NULL, knots = NULL, continuous = FALSE,
constr = TRUE, constr.slopes = 2, constr.intercepts = TRUE,
min.obs = 3, discard.obs = TRUE)
```

#### **Arguments**

expr	Vector of gene expression values
cghseg	Vector of segmented copy number values
cghcall	Vector of called copy number values. If not provided, we are reduced to a simple linear model.
probloss	Vector of call probabilities associated with state "loss". Default is NULL.
probnorm	Vector of call probabilities associated with state "normal". Default is NULL.
probgain	Vector of call probabilities associated with state "gain". Default is NULL.
probamp	Vector of call probabilities associated with state "amplification". Default is NULL.
knots	knots or change points. If NULL (default), there are estimated. See details.
continuous	Logical, whether the model is continuous (no jump) or not.
constr	Logical, whether the model is constrained or not. (this has been implemented to turn on and off easily the constraints)
constr.slopes	Type of non-negativity constraints applied on slopes. Either 1 or 2 (default). See details.
constr.interce	pts
	If TRUE (default) jumps from state to state are also constrained to be non-negative
min.obs	See modify.conf
discard.obs	See modify.conf

#### **Details**

If cghcall=NULL, discrete copy number values are omitted, which results in fitting a simple linear model.

If constr.slopes=1, all slopes are constrained to be non-negative. If constr.slopes=2, the slope associated with state "normal" is constrained to be non-negative and all others are forced to be at least equal to the latter.

Two methods are implemented for the estimation of knots. If call probabilities are provided, a knot is determined so that the sum of (the two adjacent) states membership probabilities is maximized. Otherwise, this is defined as the midpoint of the interval between the two consecutive states.

The constrained least squares problem is solved using function solve. QP of package quadprog.

plrs-class 9

#### Value

An object of class plrs-class

#### Author(s)

Gwenael G.R. Leday <g.g.r.leday@vu.nl>

### **Examples**

```
# Simulate data
sim <- plrs.sim(n=80, states=4, sigma=0.5)

# Fit a model
model <- plrs(expr=sim$expr, cghseg=sim$seg, cghcall=sim$cal)
model

# Methods
coef(model)
effects(model)
fitted(model)
knots(model)
model.matrix(model)
plot(model)
predict(model, newcghseg=seq(0,5, length.out=100))
residuals(model)
summary(model)</pre>
```

plrs-class

Class plrs

### Description

An S4 class representing the output of the plrs function.

### **Slots**

```
coefficients: Object of class numeric containing spline coefficients
fitted.values: Object of class numeric containing the fitted values
residuals: Object of class numeric containing the residuals
X: Object of class matrix containing the design matrix
data: Object of class list containing input data
mdata: Object of class list containing (possibly modified) data used to fit the model (See modify.conf).
QP: Object of class list containing input elements used for quadratic programming. If the model
is unconstrained this contains a light version of an lm object.
test: Object of class list containing results from testing.
```

10 plrs.cb

cb: Object of class list containing lower and upper bounds for predicted values.

selected: Object of class logical indicating whether the model results from a selection procedure.

type: Object of class character giving the type of model

call.arg: Object of class list containing the input arguments (for reproducibility)

#### Methods

```
coef Returns the coefficients
criteria See criteria
effects Returns matrix of effects
fitted Returns the fitted values
knots Returns the knots
model.matrix Returns the design matrix
plot See plot.plrs
predict See predict.plrs
print Print the object information
residuals Returns the residuals
show Print the object information
```

summary Print a summary of the object information

### Author(s)

Gwenael G.R. Leday <g.g.r.leday@vu.nl>

plrs.cb

Uniform confidence bands (CB) for plrs models

### Description

Determine uniform confidence intervals for predicted values of a 'plrs' model.

### Usage

```
plrs.cb(object, alpha=0.05, newcgh=NULL)
```

### Arguments

object An object of class plrs-class.

alpha Significance level

newcgh Vector of segmented values. Support for building CB.

plrs.select 11

#### **Details**

The input object of class plrs-class has to result from function plrs. test.

The problem of finding (at a given x) a confidence interval for the mean response is expressed as a semi-definite optimization problem and solved using function csdp of package **Rcsdp**.

#### Value

An object of class plrs-class that contains CB information.

#### Author(s)

```
Gwenael G.R. Leday < g.g.r.leday@vu.nl>
```

#### References

Leday GGR, Van der Vaart AW, Van Wieringen WN, Van de Wiel MA. Modeling association between DNA copy number and gene expression with constrained piecewise linear regression splines. Accepted for publication. *Ann Appl Stat.* (2012).

#### See Also

```
plrs.test
```

### **Examples**

```
# Simulate data
sim <- plrs.sim(n=80, states=4, sigma=0.5)

# Fit a model
model <- plrs(expr=sim$expr, cghseg=sim$seg, cghcall=sim$cal)

# Confidence bands
model <- plrs.test(model)
model <- plrs.cb(model, alpha=0.05)
plot(model)</pre>
```

plrs.select

Model selection

### **Description**

Selection of a model based on an information criterion (AIC, AICC, BIC or OSAIC).

#### Usage

```
plrs.select(object, crit = ifelse(object@call.arg$constr,"osaic","aic"))
```

12 plrs.select-class

#### **Arguments**

object An object of class plrs-class

crit Character corresponding to the criterion to use. See criteria.

#### Value

```
An object of class plrs. select-class
```

### Author(s)

```
Gwenael G.R. Leday < g.g.r.leday@vu.nl>
```

plrs.select-class

 ${\it Class}\, {\it plrs.select}$ 

### Description

An S4 class representing the output of the plrs. select function.

### **Slots**

table: Object of class matrix containing the criterion value for all models

model: Object of class plrs containing the selected model

crit: Object of class character containing the criterion used for model selection

#### Methods

```
plot See plot.plrs
```

print Print the object information

show Print the object information

summary Print a summary of the object information

### Author(s)

```
Gwenael G.R. Leday <g.g.r.leday@vu.nl>
```

plrs.series 13

plrs.series	Fit plrs models for a series of arrays.	
-------------	---	--

### **Description**

The function fits plrs models for a series of arrays. Model selection and testing procedures may be applied.

### Usage

```
plrs.series(expr, cghseg, cghcall=NULL,
probloss = NULL, probnorm = NULL, probgain = NULL, probamp = NULL,
control.model = list(continuous = FALSE,
                      constr = TRUE,
                      constr.slopes = 2,
                      constr.intercepts = TRUE,
                      min.obs = 3,
                      discard.obs = TRUE),
control.select = list(crit = ifelse(control.model$constr, "osaic", "aic")),
control.test = list(testing = TRUE,
                      cb = FALSE,
                      alpha = 0.05),
control.output = list(save.models = FALSE,
                      save.plots = FALSE,
                      plot.lin = FALSE,
                      type = "jpeg"))
```

#### **Arguments**

expr	Either a matrix of expression profiles or an ExpressionSet object.						
cghseg	Either a matrix of segmented copy number values or objects of class cghSeg or cghCall						
cghcall	Matrix of called copy number						
probloss	Matrix of call probabilities associated with state "loss". Default is NULL.						
probnorm	Matrix of call probabilities associated with state "normal". Default is NULL.						
probgain	Matrix of call probabilities associated with state "gain". Default is NULL.						
probamp	Matrix of call probabilities associated with state "amplification". Default is NULL.						
control.model	See details						
control.select	See details						
control.test	See details						
control.output	See details						

14 plrs.series

#### **Details**

If DNA and mRNA input data are matrices, rows should correspond to genes and columns to arrays. Alternatively, expression data may be provided as an ExpressionSet object and aCGH data as cghSeg or cghCall objects. A cghCall object contain all data from the calling step, thus arguments probloss, probnorm, probnorm and probamp can be omitted. An object of class cghSeg does not contain such data so only simple linear models will be fitted.

control.model allows the user to specify the type of model that has to be fitted. This must be a list with one or more of the following components: constr.slopes, constr.intercepts, min.obs and discard.obs. See functions plrs and modify.conf for more details.

control.select allows the user to specify whether model selection should be done and how. This must be a list with a component named crit. See function plrs.select for more details. If control.select = NULL then no model selection is done.

control.output allows the user to plot and save each plrs model. This must be a list with components:

save.models, a logical. This will create within the work directory a new directory named "plrsSeriesObjects" that will contain all objects.

save.plots, a logical. This will create within the work directory a new directory named "plrsSeries-Plots" that will contains all saved plots.

plot.lin, a logical. Whether the simple linear model should aslo be plotted.

type, a character. Format of file. To pass through function savePlot.

#### Value

An object of class plrs. series-class

#### Author(s)

Gwenael G.R. Leday < g.g.r.leday@vu.nl>

#### **Examples**

```
# Simulate data
ngenes <- 10
narray <- 48
rna <- dnaseg <- dnacal <- matrix(NA, ngenes, narray)
idx <- sample(1:4, ngenes, replace=TRUE, prob=rep(1/4,4))
for(i in 1:ngenes){
Sim <- plrs.sim(n=narray, states=idx[i], sigma=0.5)
rna[i,] <- Sim$expr
dnaseg[i,] <- Sim$seg
dnacal[i,] <- Sim$seg
}</pre>
```

# Screening procedure with linear model

plrs.series-class 15

```
series <- plrs.series(expr = rna, cghseg = dnaseg, cghcall = NULL, control.select = NULL)
# Screening procedure with full plrs model
series <- plrs.series(expr = rna, cghseg = dnaseg, cghcall = dnacal, control.select = NULL)
# Model selection
series <- plrs.series(expr = rna, cghseg = dnaseg, cghcall = dnacal)</pre>
```

plrs.series-class

Class plrs.series

### Description

An S4 class representing the output of the plrs. series function.

### **Slots**

```
coefficients: Matrix containing coefficients of models
effects: List containing effects
test: Matrix containing results from testing.
general: Matrix providing the distribution of the number genes and arrays regarding the copy number states
modelsType: List providing models' type
call.arg: List providing details on the type of models that have been fitted.
```

#### Methods

```
print Print the object informationshow Print the object informationsummary Print a summary of the object information
```

#### Author(s)

```
Gwenael G.R. Leday <g.g.r.leday@vu.nl>
```

plrs.sim

plrs.sim

Simulation of a plrs model

### **Description**

Simulation of a piecewise relationship.

The function has been only implemented for convenience of simulations and R examples.

### Usage

```
plrs.sim(n = 80, states = 4, sigma = 01, x = NULL)
```

### **Arguments**

Number of simulated data points
 states
 Number of states for the model
 sigma
 Noise
 x
 Segmented values.

#### **Details**

To be written...

### Author(s)

```
Gwenael G.R. Leday < g.g.r.leday@vu.nl>
```

### Examples

```
# Simulate 1-state model
sim <- plrs.sim(n=80, states=1, sigma=0.5)
model <- plrs(expr=sim$expr, cghseg=sim$seg, cghcall=sim$cal)
plot(model)

# Simulate 2-state model
sim <- plrs.sim(n=80, states=2, sigma=0.5)
model <- plrs(expr=sim$expr, cghseg=sim$seg, cghcall=sim$cal)
plot(model)

# Simulate 3-state model
sim <- plrs.sim(n=90, states=3, sigma=0.5)
model <- plrs(expr=sim$expr, cghseg=sim$seg, cghcall=sim$cal)
plot(model)

# Simulate 4-state model</pre>
```

plrs.test 17

```
sim <- plrs.sim(n=80, states=4, sigma=0.5)
model <- plrs(expr=sim$expr, cghseg=sim$seg, cghcall=sim$cal)
plot(model)</pre>
```

plrs.test

Likelihood ratio test for a plrs model

### Description

Test whether copy number has an effect on mRNA expression.

### Usage

```
plrs.test(object, alpha=0.05)
```

#### **Arguments**

object An object of class plrs-class

alpha Significance level

#### **Details**

Two cases present themselves:

- 1. The model is unconstrained. Thus, the model under the null hypothesis is the intercept and an F-test is performed.
- 2. The model is constrained and the following hypothesis are tested:

H0: All constraints are actives (=)

H1: At least one constraint is strict (>)

Under H0, we always have the intercept model. Indeed, if constr.slopes = 1 (or 2) and constr.intercepts = T, then the only parameter free of inequality constraint is the overall intercept. If constr.intercepts = F, the local intercepts are additionally constrained to be 0 in order to obtain the intercept model under the null. The likelihood ratio statistic (unknown variance) is asymptotically distributed as a weighted mixture of Beta distribution (cf Gromping (2010)). Calculation of p-values is based on functions ic.weights and pbetabar of package **ic.infer**. The package **mvtnorm** is also involved.

In both cases the input model is taken as the model under the alternative.

### Value

A list object with the following components:

stat Test statistic
pvalue Calculated pvalue

18 predict.plrs

wt.bar	Weights (if the mod	el is constrained)
--------	---------------------	--------------------

df.bar Degrees of freedom.

unconstr Unconstrained model of class plrs-class

qbetabar (1-alpha) quantile of the beta mixture distribution

alpha Significance level

### Author(s)

```
Gwenael G.R. Leday < g.g.r.leday@vu.nl>
```

#### References

Gromping, U. (2010). Inference with linear equality and inequality constraints using R: The package ic.infer. *J Stat Softw*, 33(i10).

### **Examples**

```
# Simulate data
sim <- plrs.sim(n=80, states=2, sigma=0.5)
# Fit a model
model <- plrs(expr=sim$expr, cghseg=sim$seg, cghcall=sim$cal)
# Testing
model <- plrs.test(model)
model</pre>
```

predict.plrs

Predict method for plrs models

#### **Description**

Determine predicted values based on a given plrs model

### Usage

```
## S3 method for class 'plrs'
predict(object, newcghseg, ...)
```

### Arguments

object An object of class plrs-class

newcghseg A vector of new segmented CGH values

... further arguments

predict.plrs 19

### Value

A vector containing the fitted values.

### Author(s)

Gwenael G.R. Leday < g.g.r.leday@vu.nl>

## **Index**

```
*Topic copy number, gene expression,
        regression splines, model
        selection, constrained
        inference.
    plrs-package, 2
cghCall, 5, 13, 14
cghSeg, 13, 14
coef, plrs-method (plrs-class), 9
criteria, 3, 10, 12
criteria,plrs-method(criteria), 3
effects, plrs-method (plrs-class), 9
ExpressionSet, 6, 13, 14
fitted, plrs-method (plrs-class), 9
knots, plrs-method (plrs-class), 9
model.matrix,plrs-method(plrs-class),9
modify.conf, 4, 8, 9, 14
neveCN17, 5
neveGE17, 6
par, 7
plot, ANY (plot-methods), 6
plot, plrs, ANY-method (plot-methods), 6
plot,plrs.select,ANY-method
        (plot-methods), 6
plot-methods, 6
plot.plrs, 10, 12
plot.plrs(plot-methods), 6
plrs, 2, 4, 5, 7, 9, 14
plrs, ANY (plrs), 7
plrs-class, 9
plrs-package, 2
plrs.cb, 2, 10
plrs.select, 2, 11, 12, 14
plrs.select, ANY (plrs.select), 11
plrs.select-class, 12
```

```
plrs.series, 2, 5, 13, 15
plrs.series-class, 15
plrs.sim, 16
plrs.test, 2, 11, 17
predict,plrs-method(plrs-class),9
predict.plrs, 10, 18
print,plrs-method(plrs-class),9
print,plrs.select-method
        (plrs.select-class), 12
print,plrs.series-method
        (plrs.series-class), 15
residuals, plrs-method (plrs-class), 9
savePlot, 14
show, plrs-method (plrs-class), 9
show,plrs.select-method
        (plrs.select-class), 12
show,plrs.series-method
        (plrs.series-class), 15
summary, plrs-method (plrs-class), 9
summary,plrs.select-method
        (plrs.select-class), 12
summary,plrs.series-method
        (plrs.series-class), 15
```