Package ‘ridge’

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Title  Ridge Regression with Automatic Selection of the Penalty Parameter


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Type  Package

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Repository  CRAN

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R topics documented:

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R package for fitting linear and logistic ridge regression models.

This package contains functions for fitting linear and logistic ridge regression models, including functions for fitting linear and logistic ridge regression models for genome-wide SNP data supplied as file names when the data are too big to read into R.

For a complete list of functions, use `help(package="ridge")`.

`Steffen Moritz, Erika Cule`

Simulated genetic data with a binary phenotypes

Simulated genetic data at 15 SNPs, together with simulated binary phenotypes

`data(GenBin)`
GenBin is a saved R matrix with 500 rows and 15 columns. The first column is the phenotypes and columns 2-15 contain the genotypes. Each row represents an individual. The same data are stored in flat text files in GenBin_genotypes and GenBin_phenotypes (in the directory extdata (in the installed package) or inst/extdata (in the source)).

Source

Simulated using FREGENE

References


Examples

data(GenBin)

---

GenCont

Simulated genetic data with continuous outcomes

Description

Simulated genetic data with continuous outcomes.

Usage

data(GenCont)

Format

GenCont is a saved R matrix with 500 rows and 13 columns. The first column is the phenotypes and columns 2-13 contain the genotypes. Each row represents an individual. The same data are stored in flat text files in GenCont_genotypes and GenCont_phenotypes (in the directory extdata (in the installed package) or inst/extdata (in the source)).

Details

Genotypes were simulated using FREGENE.

References


Examples

data(GenCont)
The Ten-Factor data first described by Gorman and Toman (1966).

Description

A Ten-Factor data set first described by Gorman and Toman (1966) and used by Hoerl and Kennard (1970) (and others) to investigate regression problems.

Usage

data(Gorman)

Format

Numeric matrix.

Details

The first column is the response on the log scale, the remaining columns are the predictors.

Source


References


Examples

data(Gorman)

Hald data

Description

The Hald data as used by Hoerl, Kennard and Baldwin (1975). These data are also in package wle.

Usage

data(Hald)
linearRidge

Format

Numeric matrix.

Details

The first column is the response and the remaining four columns are the predictors.

References


Examples

data(Hald)

linearRidge

Linear ridge regression.

Description

Fits a linear ridge regression model. Optionally, the ridge regression parameter is chosen automatically using the method proposed by Cule et al (2012).

Usage

linearRidge(formula, data, lambda = "automatic", nPCs = NULL, scaling = c("corrForm", "scale", "none"), ...)

## S3 method for class 'ridgeLinear'
coef(object, all.coef = FALSE, ...)

## S3 method for class 'ridgeLinear'
plot(x, y = NULL, ...)

## S3 method for class 'ridgeLinear'
predict(object, newdata, na.action = na.pass, all.coef = FALSE, ...)

## S3 method for class 'ridgeLinear'
print(x, all.coef = FALSE, ...)

## S3 method for class 'summary.ridgeLinear'
print(x, digits = max(3, getOption("digits") - 3), signif.stars = getOption("show.signif.stars"), ...)
Arguments

- **formula**: a formula expression as for regression models, of the form `response ~ predictors`. See the documentation of `formula` for other details.
- **data**: an optional data frame in which to interpret the variables occurring in `formula`.
- **lambda**: A ridge regression parameter. May be a vector. If `lambda` is "automatic" (the default), then the ridge parameter is chosen automatically using the method of Cule et al (2012).
- **nPCs**: The number of principal components to use to choose the ridge regression parameter, following the method of Cule et al (2012). It is not possible to specify both `lambda` and `nPCs`.
- **scaling**: The method to be used to scale the predictors. One of "corrform" (the default) scales the predictors to correlation form, such that the correlation matrix has unit diagonal. "scale" Standardizes the predictors to have mean zero and unit variance. "none" No scaling.
- **object**: A `ridgeLinear` object, typically generated by a call to `linearRidge`.
- **newdata**: An optional data frame in which to look for variables with which to predict. If omitted, the fitted values are used.
- **na.action**: function determining what should be done with missing values in `newdata`. The default is to predict `NA`.
- **all.coef**: Logical. Should results be returned for all ridge regression penalty parameters (`all.coef = TRUE`), or only for the ridge parameter chosen automatically using the method of Cule et al?
- **x**: An object of class `ridgeLinear` (for the `print.ridgeLinear` and `plot.ridgeLinear` functions) or an object of class `summary.ridgeLinear` (for the `print.summary.ridgeLinear` function).
- **y**: Dummy argument for compatibility with the default `plot` method. Ignored.
- **digits**: minimum number of significant digits to be used for most numbers
- **signif.stars**: logical; if `TRUE`, P-values are additionally encoded visually as significance stars in order to help scanning of long coefficient tables. It defaults to the `show.signif.stars` slot of options.
- **...**: Additional arguments to be passed to or from other methods.

Details

If an intercept is present in the model, its coefficient is not penalised. If you want to penalise an intercept, put in your own constant term and remove the intercept.

Value

An object of class "ridgeLinear", with components:

- **automatic**: Logical. Was `lambda` chosen automatically?
- **call**: The matched call.
- **coef**: A named vector of fitted coefficients.
df                  A vector of degrees of freedom of the model fit, degrees of freedom for variance, and residual degrees of freedom of the fitted model.

Inter                Was an intercept included?

isScaled              Were the predictors scaled before the model was fitted?

lambda                The ridge regression parameter(s).

scales                The scales used to standardize the predictors.

terms                 The terms object used.

x                     The scaled predictor matrix.

xm                    A vector of means of the predictors.

y                     The response.

ym                    The mean of the response.

And optionally the components

max.nPCs              The maximum number of principal components for which a ridge regression parameter was computed.

chosen.nPCs           The number of principal components used to compute the ridge parameter.

Author(s)

Erika Cule

References


See Also

logisticRidge

Examples

data(GenCont)
mod <- linearRidge(Phenotypes ~ ., data = as.data.frame(GenCont))
summary(mod)
linearRidgeGenotypes  
*Fits linear ridge regression models for genome-wide SNP data.*

**Description**

Fits linear ridge regression models for genome-wide SNP data. The SNP genotypes are not read into R but file names are passed the code directly, enabling the analysis of genome-wide scale SNP data sets.

**Usage**

```r
linearRidgeGenotypes(genotypesfilename, phenotypesfilename, lambda = -1, thinfilename = NULL, betafilename = NULL, approxfilename = NULL, permfilename = NULL, intercept = TRUE, verbose = FALSE)
```

**Arguments**

- `genotypesfilename`: character string: path to file containing SNP genotypes coded 0, 1, 2. See Input file formats.
- `phenotypesfilename`: character string: path to file containing phenotypes. See Input file formats.
- `lambda`: (optional) shrinkage parameter. If not provided, the default denotes automatic choice of the shrinkage parameter using the method of Cule & De Iorio (2012).
- `thinfilename`: (optional) character string: path to file containing three columns: SNP name, chromosome and SNP position. See Input file formats. (See details.)
- `betafilename`: (optional) character string: path to file where the output will be written. See Output file formats.
- `approxfilename`: (optional) character string: path to file where the approximate test p-values will be written. Approximate p-values are not computed unless this argument is given. Approximate p-values are computed using the method of Cule et al (2011). See Output file formats.
- `permfilename`: (optional) character string: path to file where the permutation test p-values will be written. Permutation test p-values are not computed unless this argument is given. (See warning.) See Output file formats.
- `intercept`: Logical: Should the ridge regression model be fitted with an intercept? (Defaults to TRUE)
- `verbose`: Logical: If TRUE, additional information is printed to the R output as the code runs. Defaults to FALSE.

**Details**

If a file thin is supplied, and the shrinkage parameter lambda is being computed automatically based on the data, then this file is used to thin the SNP data by SNP position. If this file is not supplied, SNPs are thinned automatically based on number of SNPs.
Value

The vector of fitted ridge regression coefficients. If betafile is given, the fitted coefficients are written to this file as well as being returned. If approxfile and/or permfile are given, results of approximate test p-values and/or permutation test p-values are written to the files given in their arguments.

Input file formats

- **genotypesfile**: A header row, plus one row for each individual, one SNP per column. The header row contains SNP names. SNPs are coded as 0, 1, 2 for minor allele count. Missing values are not accommodated. Invariant SNPs in the data cause an error, please remove these from the file before calling the function.

- **phenofile**: A single column of phenotypes with the individuals in the same order as those in the file genotypesfile.

- **thin**: (optional) Three columns and the same number of rows as there are SNPs in the file genotypesfile, one row per SNP. First column: SNP names (must match names in genotypesfile); second column: chromosome; third column: SNP position in BP.

Output file formats

All output files are optional. Whether or not betafile is provided, fitted coefficients are returned to the R workspace. If betafile is provided, fitted coefficients are written to the file specified (in addition).

- **betafile**: Two columns: First column is SNP names in same order as in genotypesfile, second column is fitted coefficients. If intercept = TRUE (the default) then the first row is the fitted intercept (with the name Intercept in the first column).

- **approxfile**: Two columns: First column is SNP names in same order as in genotypesfile, second column is approximate p-values.

- **permfile**: Two columns: First column is SNP names in same order as in genotypesfile, second column is permutation p-values.

Warning

When data are large, the permutation test p-values may take a very long time to compute. It is recommended not to request permutation test p-values (using the argument permfile) when data are large.

Author(s)

Erika Cule

References

See Also

`linearRidge` for fitting linear ridge regression models when the data are small enough to be read into R. `logisticRidge` and `logisticRidgeGenotypes` for fitting logistic ridge regression models.

Examples

```r
## Not run:
genotypesfile <- system.file("extdata","GenCont_genotypes.txt",package = "ridge")
phenotypesfile <- system.file("extdata","GenCont_phenotypes.txt",package = "ridge")
beta_linearRidgeGenotypes <- linearRidgeGenotypes(genotypesfilename = genotypesfile,
phenotypesfilename = phenotypesfile)
## compare to output of linearRidge
data(GenCont) ## Same data as in GenCont_genotypes.txt and GenCont_phenotypes.txt
beta_linearRidge <- linearRidge(Phenotypes ~ ., data = as.data.frame(GenCont))
cbind(round(coef(beta_linearRidge), 6), beta_linearRidgeGenotypes)
## End(Not run)
```

linearRidgeGenotypesPredict

`Predict phenotypes from genome-wide SNP data based on a file of coefficients`

Description

Predict phenotypes from genome-wide SNP data based on a file of coefficients. Genotypes and fitted coefficients are provided as filenames, allowing the computation of fitted probabilities when SNP data are too large to be read into R.

Usage

`linearRidgeGenotypesPredict(genotypesfilename, betafilename, phenotypesfilename = NULL, verbose = FALSE)`

Arguments

- `genotypesfilename`: character string: path to file containing SNP genotypes coded 0, 1, 2. See Input file formats.
- `betafilename`: character string: path to file containing fitted coefficients. See Input file formats.
- `phenotypesfilename`: (optional) character string: path to file in which to write out the predicted phenotypes. See Output file formats. Whether or not this argument is supplied, the fitted coefficients are also returned by the function.
- `verbose`: Logical: If TRUE, additional information is printed to the R output as the code runs. Defaults to FALSE.
Value

A vector of fitted values, the same length as the number of individuals whose data are in genotypesfilename. If phenotypesfilename is supplied, the fitted values are also written there.

Input file formats

- **genotypesfilename**: A header row, plus one row for each individual, one SNP per column. The header row contains SNP names. SNPs are coded as 0, 1, 2 for minor allele count. Missing values are not accommodated.

- **betafilename**: Two columns: First column is SNP names in same order as in genotypesfilename, second column is fitted coefficients. If the coefficients include an intercept then the first row of betafilename should contain it with the name Intercept in the first column. An Intercept thus labelled will be used appropriately in predicting the phenotypes. SNP names must match those in genotypesfilename. The format of betafilename is that of the output of linearRidgeGenotypes, meaning linearRidgeGenotypesPredict can be used to predict using coefficients fitted using linearRidgeGenotypes (see the example).

Output file format

Whether or not phenotypesfilename is provided, predicted phenotypes are returned to the R workspace. If phenotypesfilename is provided, predicted phenotypes are written to the file specified (in addition).

- **phenotypesfilename**: One column, containing predicted phenotypes, one individual per row.

Author(s)

Erika Cule

References


See Also

linearRidgeGenotypes for model fitting. logisticRidgeGenotypes and logisticRidgeGenotypesPredict for corresponding functions to fit and predict on SNP data with binary outcomes.

Examples

```r
## Not run:
genotypesfile <- system.file("extdata","GenCont_genotypes.txt",package = "ridge")
phenotypesfile <- system.file("extdata","GenCont_phenotypes.txt",package = "ridge")
betafile <- tempfile(pattern = "beta", fileext = ".dat")
beta_linearRidgeGenotypes <- linearRidgeGenotypes(genotypesfilename = genotypesfile,
                                            phenotypesfilename = phenotypesfile,
                                            betafilename = betafile)
pred_phen_geno <- linearRidgeGenotypesPredict(genotypesfilename = genotypesfile,
                                            phenotypesfilename = phenotypesfile,
                                            betafilename = betafile)
```
```r
## compare to output of linearRidge
data(GenCont) ## Same data as in GenCont_genotypes.txt and GenCont_phenotypes.txt
beta_linearRidge <- linearRidge(Phenotypes ~ ., data = as.data.frame(GenCont))
pred_phen <- predict(beta_linearRidge)
print(cbind(pred_phen_geno, pred_phen))
## Delete the temporary betafile
unlink(betafile)
## End(Not run)

logisticRidge

Logistic ridge regression.

Description

Fits a logistic ridge regression model. Optionally, the ridge regression parameter is chosen automatically using the method proposed by Cule et al (2012).

Usage

logisticRidge(formula, data, lambda = "automatic", nPCs = NULL, scaling = c("corrForm", "scale", "none"), ...)

coef(object, all.coef = FALSE, ...)

plot(x, y = NULL, ...)

predict(object, newdata = NULL, type = c("link", "response"), na.action = na.pass, all.coef = FALSE, ...)

print(x, all.coef = FALSE, ...)

summary(object, all.coef = FALSE, ...)

Arguments

formula a formula expression as for regression models, of the form response ~ predictors. See the documentation of formula for other details.
data An optional data frame in which to interpret the variables occurring in formula.

lambda A ridge regression parameter. If lambda is "automatic" (the default), then the ridge parameter is chosen automatically using the method of Cule et al (2012).

nPCs The number of principal components to use to choose the ridge regression parameter, following the method of Cule et al (2012). It is not possible to specify both lambda and nPCs.

scaling The method to be used to scale the predictors. One of "corrform" (the default) scales the predictors to correlation form, such that the correlation matrix has unit diagonal. "scale" Standardizes the predictors to have mean zero and unit variance. "none" No scaling.

object A ridgeLogistic object, typically generated by a call to linearRidge.

newdata An optional data frame in which to look for variables with which to predict. If omitted, the fitted values are used.

type the type of prediction required. The default predictions are of log-odds (probabilities on logit scale) and type = "response" gives the predicted probabilities.

na.action function determining what should be done with missing values in newdata. The default is to predict NA.

all.coef Logical. Should results be returned for all ridge regression penalty parameters (all.coef = TRUE), or only for the ridge parameter chosen automatically using the method of Cule et al?

x An object of class ridgeLogistic (for the print.ridgeLogistic and plot.ridgeLogistic functions) or an object of class summary.ridgeLogistic (for the print.summary.ridgeLogistic function)

y Dummy argument for compatibility with the default plot method. Ignored.

digits minimum number of significant digits to be used for most numbers

signif.stars logical; if TRUE, P-values are additionally encoded visually as significance stars in order to help scanning of long coefficient tables. It defaults to the show.signif.stars slot of options.

... Additional arguments to be passed to or from other methods.

Details

If an intercept is present in the model, its coefficient is not penalised. If you want to penalise an intercept, put in your own constant term and remove the intercept.

Value

An object of class "ridgeLogistic", with components:

automatic Was lambda chosen automatically?

call The matched call.

coef A named vector of fitted coefficients.

df A vector of degrees of freedom of the model fit and degrees of freedom for variance.
Inter  Was in intercept included?
isScaled  Were the predictors scaled before the model was fitted?
lambda  The ridge regression parameter.
scales  The scales used to standardize the predictors.
terms  The terms object used.
x  The scaled predictor matrix.
xm  A vector of means of the predictors.
y  The response.

And optionally the component

nPCs  The number of principal components used to compute the ridge regression parameter.

Author(s)

Erika Cule

References


See Also

linearRidge

Examples

data(GenBin)
mod <- logisticRidge(Phenotypes ~ ., data = as.data.frame(GenBin))
summary(mod)

logisticRidgeGenotypes

Fits logistic ridge regression models for genome-wide SNP data.

Description

Fits logistic ridge regression models for genome-wide SNP data. The SNP genotypes are not read into R but file names are passed to the code directly, enabling the analysis of genome-wide SNP data sets which are too big to be read into R.
Usage

logisticRidgeGenotypes(genotypesfilename, phenotypesfilename, lambda = -1,
thinfilename = NULL, betafilename = NULL, approxfilename = NULL,
permfilename = NULL, intercept =
TRUE, verbose = FALSE)

Arguments

genotypesfilename
character string: path to file containing SNP genotypes coded 0, 1, 2. See Input file formats.

phenotypesfilename
character string: path to file containing phenotypes. See Input file formats.

lambda
(optional) shrinkage parameter. If not provided, the default denotes automatic choice of the shrinkage parameter using the method of Cule & De Iorio (2012).

thinfilename
(optional) character string: path to file containing three columns: SNP name, chromosome and SNP position. See Input file formats. (See details.)

betafilename
(optional) character string: path to file where the output will be written. See Output file formats.

approxfilename
(optional) character string: path to file where the approximate test p-values will be written. Approximate p-values are not computed unless this argument is given. Approximate p-values are computed using the method of Cule et al (2011). See Output file formats.

permfilename
(optional) character string: path to file where the permutation test p-values will be written. Permutation test p-values are not computed unless this argument is given. (See warning). See Output file formats.

intercept
Logical: Should the ridge regression model be fitted with an intercept? Defaults to TRUE.

verbose
Logical: If TRUE, additional information is printed to the R output as the code runs. Defaults to FALSE.

Details

If a file thin is supplied, and the shrinkage parameter lambda is being computed automatically based on the data, then this file is used to thin the SNP data by SNP position. If this file is not supplied, SNPs are thinned automatically based on number of SNPs.

Value

The vector of fitted ridge regression coefficients. If betafilename is given, the fitted coefficients are written to this file as well as being returned. If approxfilename and/or permfilename are given, results of approximate test p-values and/or permutation test p-values are written to the files given in their arguments.
logisticRidgeGenotypes

Input file formats

**genotypesfilename:** A header row, plus one row for each individual, one SNP per column. The header row contains SNP names. SNPs are coded as 0, 1, 2 for minor allele count. Missing values are not accommodated. Invariant SNPs in the data cause an error, please remove these from the file before calling the function.

**phenofilename:** A single column of phenotypes with the individuals in the same order as those in the file genotypesfilename. Phenotypes must be coded as 0 or 1.

**thin:** (optional) Three columns and the same number of rows as there are SNPs in the file genotypesfilename, one row per SNP. First column: SNP names (must match names in genotypesfilename); second column: chromosome; third column: SNP position in BP.

Output file formats

All output files are optional. Whether or not betafilename is provided, fitted coefficients are returned to the R workspace. If betafilename is provided, fitted coefficients are written to the file specified (in addition).

**betafilename:** Two columns: First column is SNP names in same order as in genotypesfilename, second column is fitted coefficients. If intercept = TRUE (the default) then the first row is the fitted intercept (with the name Intercept in the first column).

**approxfilename:** Two columns: First column is SNP names in same order as in genotypesfilename, second column is approximate p-values.

**permfilename:** Two columns: First column is SNP names in same order as in genotypesfilename, second column is permutation p-values.

Warning

When data are large, the permutation test p-values may take a very long time to compute. It is recommended not to request permutation test p-values (using the argument permfilename) when data are large.

Author(s)

Erika Cule

References


See Also

logisticRidge for fitting logistic ridge regression models when the data are small enough to be read into R. linearRidge and linearRidgeGenotypes for fitting linear ridge regression models.
logisticRidgeGenotypesPredict

Predict fitted probabilities from genome-wide SNP data based on a file of coefficients

Examples

```r
## Not run:
genotypesfile <- system.file("extdata","GenBin_genotypes.txt",package = "ridge")
phenotypesfile <- system.file("extdata","GenBin_phenotypes.txt",package = "ridge")
beta_logisticRidgeGenotypes <-
logisticRidgeGenotypes(genotypesfilename = genotypesfile, phenotypesfilename = phenotypesfile)
## compare to output of logisticRidge
data(GenBin) ## Same data as in GenBin_genotypes.txt and GenBin_phenotypes.txt
beta_logisticRidge <- logisticRidge(Phenotypes ~ ., data = as.data.frame(GenBin))
cbind(round(coef(beta_logisticRidge), 6), beta_logisticRidgeGenotypes)
## End(Not run)
```

Description

Predict fitted probabilities from genome-wide SNP data based on a file of coefficients. Genotypes and fitted coefficients are provided as filenames, allowing the computation of fitted probabilities when SNP data are too large to be read into R.

Usage

```r
logisticRidgeGenotypesPredict(genotypesfilename, betafilename,
phenotypesfilename = NULL, verbose = FALSE)
```

Arguments

- `genotypesfilename` character string: path to file containing SNP genotypes coded 0, 1, 2. See Input file formats.
- `betafilename` character string: path to file containing fitted coefficients. See Input file formats.
- `phenotypesfilename` (optional) character string: path to file in which to write out the fitted probabilities. See Output file formats. Whether or not this argument is supplied, the fitted coefficients are also returned by the function.
- `verbose` Logical: If TRUE, additional information is printed to the R output as the code runs. Defaults to FALSE.

Value

A vector of fitted probabilities, the same length as the number of individuals whose data are in `genotypesfilename`. If `phenotypesfilename` is supplied, the fitted probabilities are also written there.
Input file formats

**genotypesfilename**: A header row, plus one row for each individual, one SNP per column. The header row contains SNP names. SNPs are coded as 0, 1, 2 for minor allele count. Missing values are not accommodated.

**betafilename**: Two columns: First column is SNP names in same order as in genotypesfilename, second column is fitted coefficients. If the coefficients include an intercept then the first row of betafilename should contain it with the name Intercept in the first column. An Intercept thus labelled will be used appropriately in predicting the phenotypes. SNP names must match those in genotypesfilename. The format of betafilename is that of the output of linearRidgeGenotypes, meaning linearRidgeGenotypesPredict can be used to predict using coefficients fitted using linearRidgeGenotypes (see the example).

Output file format

Whether or not phenotypesfilename is provided, fitted probabilities are returned to the R workspace. If phenotypesfilename is provided, fitted probabilities are written to the file specified (in addition).

**phenotypesfilename**: One column, containing fitted probabilities, one individual per row.

Author(s)

Erika Cule

References


See Also

logisticRidgeGenotypes for model fitting. linearRidgeGenotypes and linearRidgeGenotypesPredict for corresponding functions to fit and predict on SNP data with continuous outcomes.

Examples

```r
## Not run:
genotypesfile <- system.file("extdata","GenBin_genotypes.txt",package = "ridge")
phenotypesfile <- system.file("extdata","GenBin_phenotypes.txt",package = "ridge")
betafile <- tempfile(pattern = "beta", fileext = ".dat")
beta_logisticRidgeGenotypes <- logisticRidgeGenotypes(genotypesfilename = genotypesfile,
phenotypesfilename = phenotypesfile,
betafilename = betafile)
pred_phen_geno <- logisticRidgeGenotypesPredict(genotypesfilename = genotypesfile,
betafilename = betafile)

## compare to output of logisticRidge
data(GenBin) ## Same data as in GenBin_genotypes.txt and GenBin_phenotypes.txt
beta_logisticRidge <- logisticRidge(Phenotypes ~ ., data = as.data.frame(GenBin))
pred_phen <- predict(beta_logisticRidge, type="response")
print(cbind(pred_phen_geno, pred_phen))
```
pvals

Compute p-values for ridgeLinear and ridgeLogistic models

Description

Functions for computing, printing and plotting p-values for ridgeLinear and ridgeLogistic models. The p-values are computed using the significance test of Cule et al (2011).

Usage

pvals(x, ...)

## S3 method for class 'ridgeLinear'
pvals(x, ...)

## S3 method for class 'ridgeLogistic'
pvals(x, ...)

## S3 method for class 'pvalsRidgeLinear'
print(x, digits = max(3, getOption("digits") - 3),
    signif.stars = getOption("show.signif.stars"), all.coef = FALSE, ...)

## S3 method for class 'pvalsRidgeLogistic'
print(x, digits = max(3, getOption("digits") - 3),
    signif.stars = getOption("show.signif.stars"), all.coef = FALSE, ...)

## S3 method for class 'pvalsRidgeLinear'
plot(x, y = NULL, ...)

## S3 method for class 'pvalsRidgeLogistic'
plot(x, y = NULL, ...)

Arguments

x For the pvals methods, an object of class "ridgeLinear" or "ridgeLogistic", typically from a call to "linearRidge" or "logisticRidge". For the print and plot methods, an object of class "pvalsRidgeLinear" or "pvalsRidgeLogistic", typically from a call to "pvals".
digits minimum number of significant digits to be used for most numbers
signif.stars logical; if TRUE, P-values are additionally encoded visually as significance stars in order to help scanning of long coefficient tables. It defaults to the show.signif.stars slot of options.
all.coef Logical. Should p-values for all the ridge regression parameters be printed, or only the one from the ridge parameter chosen using the method of Cule et al (2012)

y Dummy argument for compatibility with the default plot method. Ignored.

... further arguments to be passed to or from other methods

Details

Standard errors, test statistics and p-values are computed using coefficients and data on the scale that was used to fit them. If the coefficients were standardized before the model was fitted, then the p-values relate to the scaled data.

Value

For the pvals methods, an object of class "pvalsRidgeLinear" or "pvalsRidgeLogistic" which is a list with elements

coeff The (scaled) regression coefficients
se The standard errors of the regression coefficients
tstat The test statistic of the regression coefficients
pval The p-values of the regression coefficients
isScaled Were the data scaled before the regression coefficients were fitted?

For the print methods, the argument x is returned invisibly.

Author(s)

Erika Cule

References


See Also

linearRidge, logisticRidge

Examples

data(GenBin)
mod <- logisticRidge(Phenotypes ~ ., data = as.data.frame(GenBin))
pvalsMod <- pvals(mod)
print(pvalsMod)
print(pvalsMod, all.coef = TRUE)
plot(pvalsMod)
**Description**

Additionally includes special functions for genome-wide single-nucleotide polymorphism (SNP) data.
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