Package ‘TOP’

May 1, 2024

Title  TOP Constructs Transferable Model Across Gene Expression Platforms

Version  1.4.0

Date  2022-11-09

Description  TOP constructs a transferable model across gene expression platforms for prospective experiments. Such a transferable model can be trained to make predictions on independent validation data with an accuracy that is similar to a re-substituted model. The TOP procedure also has the flexibility to be adapted to suit the most common clinical response variables, including linear response, binomial and Cox PH models.

License  GPL-3

URL  https://github.com/Harry25R/TOP

BugReports  https://github.com/Harry25R/TOP/issues

biocViews  Software, Survival, GeneExpression

Encoding  UTF-8

Roxygen  list(markdown = TRUE)

RoxygenNote  7.2.3

Imports  assertthat, caret, ClassifyR, directPA, doParallel, dplyr, ggnewscale, ggplot2, ggraph, ggrepel, ggthemes, glmnet, Hmisc, igr aph, latex2exp, limma, magrittr, methods, plotly, pROC, purrr, reshape2, stats, stringr, survival, tibble, tidygraph, tidyr, statmod

Suggests  knitr, rmarkdown, BiocStyle, Biobase, curatedOvarianData, ggbeeswarm, ggsci, survminer, tidyverse

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LazyData  false

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Repository  Bioconductor 3.19
Description

cOefNetworkPlot

Usage

cOefNetworkPlot(TRUE_model, nFeatures = 20, s = "lambda.min")

Arguments

  TOP_model  A Transferable Omincs Prediction model. The output from the TOP_model function.
  nFeatures  The number of features that will be plotted. Default: 20
  s           Lambda value for the lasso model. Default is "lambda.min"

Value

A coefNetwork plot
expit

The expit function

Description
The expit function

Usage
expit(x)

Arguments
x numeric

Value
The expit of x

Examples
curve(expit, from = -5, to = 5)
filterFeatures

Description

A function that implements feature selection, using limma, from a list of data frames with corresponding labels.

Usage

filterFeatures(
  x_list,
  y_list,
  contrast = NULL,
  nFeatures = 50,
  combinationMethod = "OSP"
)

Arguments

x_list A list of data frames, with columns corresponding to features and rows corresponding to observations.
y_list A list of factor labels.
contrast A character vector describing which order of levels to contrast in y_list ("disease - control"), Default: NULL
nFeatures Number of features to return, Default: 50
combinationMethod Which p-value combination method to use, Default: 'OSP' Options are 'Stouffer', 'OSP', 'Fisher', 'maxP'.

Details

contrast must be a character vector of length 1. If contrast is NULL, the first level of the first factor in y_list will be used as the reference level.

Value

A vector of feature names.

Examples

data(TOP_data_binary, package = "TOP")
x1 <- TOP_data_binary$x1
x2 <- TOP_data_binary$x2
x3 <- TOP_data_binary$x3
x_list <- list(x1, x2, x3)
pairwise_col_diff

Compute pairwise difference between matrix columns

Description

Compute pairwise difference between matrix columns

Usage

pairwise_col_diff(x)

Arguments

x A data matrix of size n times p. Where rows are observations and columns are features.

Value

A matrix of size n times (p choose 2), where each column is the difference between two of the original columns.

Examples

n <- 1
p <- 4
x <- matrix(rep(seq_len(p), n), nrow = n, ncol = p, byrow = TRUE)
colnames(x) <- paste0("X", seq_len(p))
pairwise_col_diff(x)
Description

A function to calculate the external performance of the Transferable Omics Prediction model.

Usage

`performance_TOP(TOP_model, newx, newy, covariates = NULL, s = "lambda.min")`

Arguments

- `TOP_model` This is the output of the function `TOP_model`.
- `newx` A matrix of the new data to be predicted. With the same number of feature columns as the original data.
- `newy` A vector of the true labels that are being predicted. With the same number of samples as `newx`.
- `covariates` A data.frame of the same covariates as the original TOP model, Default: NULL
- `s` Lambda used in the lasso model, Default: 'lambda.min'

Value

A confusion matrix that displays the performance of the classifier.

Examples

```r
data(TOP_data_binary, package = "TOP")
x1 <- TOP_data_binary$x1
x2 <- TOP_data_binary$x2
x_list <- list(x1, x2)
y_list <- list(TOP_data_binary$y1, TOP_data_binary$y2)
model <- TOP_model(x_list, y_list)
x3 <- TOP_data_binary$x3
y3 <- TOP_data_binary$y3
performance_TOP(model$models, newx = x3, newy = y3)
```
predict_TOP

Precict using the Trasferable Omics Prediction model.

Description

A prediction function for the Trasferable Omics Prediction model.

Usage

predict_TOP(TOP_model, newx, covariates = NULL, s = "lambda.min")

Arguments

- **TOP_model**: The output from the TOP_model function.
- **newx**: A matrix of the new data to be predicted. The columns should be features and the rows should be samples.
- **covariates**: A data frame of the same covariates that were used in the TOP model, Default: NULL
- **s**: Lambda value for the lasso model, Default: 'lambda.min'

Value

A vector of predictions for the new data.

Examples

data(TOP_data_binary, package = "TOP")

x1 <- TOP_data_binary$x1
x2 <- TOP_data_binary$x2
x3 <- TOP_data_binary$x3
y1 <- TOP_data_binary$y1
y2 <- TOP_data_binary$y2
y3 <- TOP_data_binary$y3

set.seed(23)
x_list <- list(x1, x2)
y_list <- list(factor(y1), factor(y2))

model <- TOP_model(x_list, y_list)
predictions <- predict_TOP(model$models, newx = x3)
Description

A function visualizes the performance of a classifier by plotting the Receiver Operating Characteristic (ROC) curve.

Usage

ROC_Plot(roc_list)

Arguments

roc_list A list of roc objects from the pROC package

Value

A ROC Plot

Examples

data(TOP_data_binary, package = "TOP")
x1 <- TOP_data_binary$x1
x2 <- TOP_data_binary$x2
x3 <- TOP_data_binary$x3
y1 <- TOP_data_binary$y1
y2 <- TOP_data_binary$y2
y3 <- TOP_data_binary$y3

set.seed(23)
x_list <- list(x1, x2)
y_list <- list(factor(y1), factor(y2))

model <- TOP_model(x_list, y_list)
pred <- predict_TOP(model$models, newx = x3)
roc <- pROC::roc(y3, pred)
ROC_Plot(list(roc))
simplenetworkPlot

Description

simplenetworkPlot

Usage

simplenetworkPlot(TOP_model, nFeatures = 50, s = "lambda.min")

Arguments

<table>
<thead>
<tr>
<th>TOP_model</th>
<th>A Transferable Omics Prediction model. The output from the TOP_model function.</th>
</tr>
</thead>
<tbody>
<tr>
<td>nFeatures</td>
<td>The number of features that will be plotted. Default: 20</td>
</tr>
<tr>
<td>s</td>
<td>Lambda value for the lasso model. Default is &quot;lambda.min&quot;</td>
</tr>
</tbody>
</table>

Value

A simple network plot

Examples

data(TOP_data_binary, package = "TOP")

x1 <- TOP_data_binary$x1
dx2 <- TOP_data_binary$x2
dx3 <- TOP_data_binary$x3
y1 <- TOP_data_binary$y1
y2 <- TOP_data_binary$y2
y3 <- TOP_data_binary$y3

set.seed(23)
x_list <- list(x1, x2)
y_list <- list(factor(y1), factor(y2))

model <- TOP_model(x_list, y_list)
simplenetworkPlot(model)
surv_top_CI Create a function to calculate the concordance index.

Description

FUNCTION DESCRIPTION

Usage

Surv_top_CI(TOP_survival, newx, newy)

Arguments

TOP_survival A TOP_survival model. See TOP_survival.
newx A new data.frame to predict the survival time.
newy A data.frame, where the first columns in each data frame is the time and the second column is the event status.

Value

An object of class concordance

Examples

data(TOP_data_binary, package = "TOP")
time <- rpois(300, c(600, 1000))
surv <- sample(c(0, 1), 300, replace = TRUE)
y <- data.frame(time, surv)
batch <- rep(paste0("y", 1:3), c(100, 100, 100))
y_list <- y |> split(batch)
x_list <- list(TOP_data_binary$x1, TOP_data_binary$x2, TOP_data_binary$x3)
surv_model <- TOP_survival(x_list[-3], y_list[-3], nFeatures = 10)
Surv_top_CI(surv_model, newx = x_list[[3]], newy = y_list[[3]])

TOPO_coefPlot

Description

TOP_coefPlot

Usage

TOP_coefPlot(TOP_model, nFeatures = 20, s = "lambda.min")
**TOP_data_binary**

**Arguments**

- **TOP_model**: A Transferable Omics Prediction model. The output from the TOP_model function.
- **nFeatures**: The number of features that will be plotted. Default: 20
- **s**: Lambda value for the lasso model, Default: 'lambda.min'

**Value**

A TOP coeff plot

**Examples**

```r
data(TOP_data_binary, package = "TOP")

x1 <- TOP_data_binary$x1
x2 <- TOP_data_binary$x2
x3 <- TOP_data_binary$x3
y1 <- TOP_data_binary$y1
y2 <- TOP_data_binary$y2
y3 <- TOP_data_binary$y3

set.seed(23)
x_list <- list(x1, x2)
y_list <- list(factor(y1), factor(y2))

model <- TOP_model(x_list, y_list)
TOP_coefPlot(model)
```

---

**Description**

A simulated binary data

**Usage**

```r
data("TOP_data_binary")
```

**Format**

A list with columns:

- **x1**: A matrix of size 100x20, each column has mean 1 and sd 1
- **x2**: A matrix of size 100x20, each column has mean 2 and sd 1
- **x3**: A matrix of size 100x20, each column has mean 3 and sd 1
y1 A factor vector of 0’s and 1’s, created by beta and x1
y2 A factor vector of 0’s and 1’s, created by beta and x2
y3 A factor vector of 0’s and 1’s, created by beta and x3
beta A vector with first 10 entries drawn from random unif(-1, 1), otherwise 0’s.

Value
The example data.

Description
TOP_lambdaPlot

Usage
TOP_lambdaPlot(
  TOP_model,
  nFeatures = 20,
  s = "lambda.min",
  interactive = FALSE,
  label = FALSE
)

Arguments

<table>
<thead>
<tr>
<th>Argument</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>TOP_model</td>
<td>A Transferable Omics Prediction model. The output from the TOP_model function.</td>
</tr>
<tr>
<td>nFeatures</td>
<td>The number of features to plot, features are ranked beta’s for lambda.min. Default: 20</td>
</tr>
<tr>
<td>s</td>
<td>Lambda value for the lasso model. Default is &quot;lambda.min&quot;</td>
</tr>
<tr>
<td>interactive</td>
<td>A boolean indicating whether the plot should be interactive. Defaults to FALSE.</td>
</tr>
<tr>
<td>label</td>
<td>A boolean indicating whether the features should be labeled on the plot. Defaults to FALSE.</td>
</tr>
</tbody>
</table>

Value
A TOP lambda plot
TOP_model

Examples

```r
data(TOP_data_binary, package = "TOP")

x1 <- TOP_data_binary$x1
x2 <- TOP_data_binary$x2
x3 <- TOP_data_binary$x3
y1 <- TOP_data_binary$y1
y2 <- TOP_data_binary$y2
y3 <- TOP_data_binary$y3

set.seed(23)
x_list <- list(x1, x2)
y_list <- list(factor(y1), factor(y2))

model <- TOP_model(x_list, y_list)
TOP_lambdaPlot(model)
```

Description
The main function of the TOP package. This function returns a glmnet model.

Usage

```r
TOP_model(
  x_list,
  y_list,
  covariates = NULL,
  dataset_weights = NULL,
  sample_weights = FALSE,
  optimiseExponent = FALSE,
  nCores = 1
)
```

Arguments

- **x_list**: a list of data frames, each containing the data for a single batch or dataset. Columns should be features and rows should be observations.
- **y_list**: a list of factors, each containing the labels for a single batch or dataset. The length of this list should be the same as the length of x_list.
- **covariates**: a list of data frames with the covariates that should be included in the model, Default: NULL
- **dataset_weights**: a list of data frames that refer to any grouping structure in the batches, Default: NULL
sample_weights Should each batch we weighted equally? This is important in unequal sample sizes, Default: FALSE

optimiseExponent Should the exponent used to modify the lasso weights be optimised using re-substitution?, Default: FALSE

nCores A numeric specifying the number of cores used if the user wants to use parallelisation, Default: 1

Value

Returns a list with the following elements: models, which is a glmnet object and features, which is a list of the features used in each model.

Examples

data(TOP_data_binary, package = "TOP")

x1 <- TOP_data_binary$x1
x2 <- TOP_data_binary$x2
x3 <- TOP_data_binary$x3
y1 <- TOP_data_binary$y1
y2 <- TOP_data_binary$y2
y3 <- TOP_data_binary$y3

set.seed(23)
x_list <- list(x1, x2)
y_list <- list(factor(y1), factor(y2))

model <- TOP_model(x_list, y_list)

Description

FUNCTION_DESCRIPTION

Usage

TOP_survival(
  x_list,
  y_list,
  nFeatures = 50,
  dataset_weights = NULL,
  sample_weights = FALSE,
  nCores = 1
)
TOP_survivalPrediction

Arguments

- **x_list**: A list of data frames, each containing the data for a single batch or dataset. Columns are features and rows are observations.
- **y_list**: A list of data frames, where the first columns in each data frame is the time and the second column is the event status. The length of this list should be the same as the length of x_list.
- **nFeatures**: Number of features to return, Default: 50
- **dataset_weights**: A list of data frames that refer to any grouping structure in the batches, Default: NULL
- **sample_weights**: Should each batch we weighted equally? This is important in unequal sample sizes, Default: FALSE
- **nCores**: A numeric specifying the number of cores used if the user wants to use parallelisation, Default: 1

Details

- DETAILS

Value

A cox net model

Examples

```r
data(TOP_data_binary, package = "TOP")
time <- rpois(300, c(600, 1000))
surv <- sample(c(0, 1), 300, replace = TRUE)
y <- data.frame(time, surv)
batch <- rep(paste0("y", 1:3), c(100, 100, 100))
y_list <- y |> split(batch)

x_list <- list(TOP_data_binary$x1, TOP_data_binary$x2, TOP_data_binary$x3)

TOP_survival(x_list[-3], y_list[-3], nFeatures = 10)
```

Description

A prediction function for TOP_survival
Usage

`TOP_survivalPrediction(TOP_survival, newx)`

Arguments

- `TOP_survival` A `TOP_survival` model. See `TOP_survival`.
- `newx` A new dataset to predict the survival time.

Value

A vector of predicted survival time.

Examples

```r
data(TOP_data_binary, package = "TOP")
time <- rpois(300, c(600, 1000))
surv <- sample(c(0, 1), 300, replace = TRUE)
y <- data.frame(time, surv)

batch <- rep(paste0("y", 1:3), c(100, 100, 100))
y_list <- y |> split(batch)

x_list <- list(TOP_data_binary$x1, TOP_data_binary$x2, TOP_data_binary$x3)
surv_model <- TOP_survival(x_list[-3], y_list[-3], nFeatures = 10)
TOP_survivalPrediction(surv_model, newx = x_list[[3]])
```
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