Package: pRoloc (via r-universe)

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Type Package

Title A unifying bioinformatics framework for spatial proteomics

Version 1.53.0

Description The pRoloc package implements machine learning and visualisation methods for the analysis and interogation of quantitiative mass spectrometry data to reliably infer protein sub-cellular localisation.

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Imports stats4, Biobase, mclust (>= 4.3), caret, e1071, sampling, class, kernlab, lattice, nnet, randomForest, proxy, FNN, hexbin, BiocGenerics, stats, dendextend, RColorBrewer, scales, MASS, knitr, mvtnorm, LaplacesDemon, coda, mixtools, gtools, plyr, ggplot2, biomaRt, utils, grDevices, graphics, colorspace

Suggests testthat, rmarkdown, pRolocdata (>= 1.43.2), roxygen2, xtable, rgl, BiocStyle (>= 2.5.19), hpar (>= 1.41.0), dplyr, akima, fields, vegan, GO.db, AnnotationDbi, Rtsne (>= 0.13), nipals, reshape, magick, umap

LinkingTo Rcpp, RcppArmadillo

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VignetteBuilder knitr

Video https://www.youtube.com/playlist?list=PLvIXxpatSLA2loV5Srs2VBpJIYUlVJ4ow

URL https://github.com/lgatto/pRoloc

BugReports https://github.com/lgatto/pRoloc/issues

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machinelearning-functions-nnet.R
machinelearning-functions-PerTurbo.R
machinelearning-functions-plsda.R
machinelearning-functions-rf.R machinelearning-functions-svm.R
machinelearning-functions-knntl.R
machinelearning-functions-tagm-map.R
machinelearning-functions-tagm-mcmc.R
machinelearning-functions-tagm-mcmc-helper.R RcppExports.R
belief.R distances.R markers.R pRolocmarkers.R chi2.R
MLInterfaces.R clustering-framework.R MSnSet.R
clustering-kmeans.R perTurbo-algorithm.R phenodisco.R
plotting.R plotting2.R plotting3.R plottingBayes.R
plotting-ellipse.R hclust.R environment.R utils.R annotation.R
goenv.R go.R makeGoSet.R vis.R MartInterface.R dynamics.R zzz.R
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Contents

addLegend 4
addMarkers 6
AnnotationParams-class 7
checkFeatureNamesOverlap 8
checkFvarOverlap 9
chi2-methods 10
classWeights 11
clustDist 12
ClustDist-class 13
ClustDistList-class 15
empPvalues 16
fDataToUnknown 17
filterBinMSnSet 18
filterMaxMarkers 19
filterMinMarkers 20
filterZeroCols 21

GenRegRes-class . . . 21
getMarkerClasses . . . 23
getMarkers . . . 24
getNormDist . . . 25
getPredictions . . . 26
highlightOnPlot . . . 27
knnClassification . . . 29
knnOptimisation . . . 30
knntlClassification . . . 31
knntlOptimisation . . . 33
ksvmClassification . . . 35
ksvmOptimisation . . . 37
MAPParams-class . . . 38
markerMSnSet . . . 41
MartInstance-class . . . 42
mcmc_get_outliers . . . 42
MCMCChains-class . . . 43
minMarkers . . . 45
mixing_posterior_check . . . 46
MLearn-methods . . . 47
move2Ds . . . 48
mrkConsProfiles . . . 49
mrkHClust . . . 50
mrkVecToMat . . . 51
nbClassification . . . 53
nbOptimisation . . . 54
nicheMeans2D . . . 55
nndist-methods . . . 57
nnetClassification . . . 58
nnetOptimisation . . . 59
orderGoAnnotations . . . 60
orgQuants . . . 62
perTurboClassification . . . 63
perTurboOptimisation . . . 64
phenoDisco . . . 66
plot2D . . . 69
plot2Ds . . . 73
plotConsProfiles . . . 75
plotDist . . . 76
plotEllipse . . . 77
plsdaClassification . . . 78
plsdaOptimisation . . . 79
pRolocmarkers . . . 81
QSep-class . . . 83
rfClassification . . . 85
rfOptimisation . . . 86
sampleMSnSet . . . 88
setLisacol . . . 89

spatial2D 91
SpatProtVis-class 92
subsetMarkers 93
svmClassification 94
svmOptimisation 95
tagmMcmcTrain 97
testMarkers 99
testMSnSet 101
thetas 102
undocumented 103
zerosInBinMSnSet 103

Index **105**

`addLegend` *Adds a legend*

Description

Adds a legend to a `plot2D` figure.

Usage

```
addLegend(
  object,
  fcol = "markers",
 where = c("bottomleft", "bottom", "bottomright", "left", "topleft", "top", "topright",
    "right", "center", "other"),
  col,
  bg,
  palette = "light",
  t = 0.3,
  pch,
  lwd,
  bty = "n",
  unknown = "unknown",
  ...
)
```

Arguments

`object`	An instance of class `MSnSet`
`fcol`	Feature meta-data label (fData column name) defining the groups to be differentiated using different colours. Default is `markers`.
`where`	One of `"bottomleft"` (default), `"bottomright"`, `"topleft"`, `"topright"` or `"other"` defining the location of the legend. `"other"` opens a new graphics device, while the other locations are passed to `legend`.

col	A character defining point colours.
bg	background (fill) color for the open plot symbols given by pch = 21:25.
palette	A character defining which palette colour theme to use, can either defined as "light" (default) or "dark".
t	A numeric between 0 and 1. Defining the degree of lightening of the colours in the palette. Default is 0.3.
pch	A character of appropriate length defining point character.
lwd	A numeric defining the line width for drawing symbols. Default is 1.5.
bty	Box type, as in legend. Default is set to "n".
unknown	A character (default is "unknown") defining how proteins of unknown/unlabelled localisation are labelled.
...	Additional parameters passed to legend.

Details

The function has been updated in version 1.3.6 to recycle the default colours when more organelle classes are provided. See plot2D for details.

Value

Invisibly returns NULL

Author(s)

Laurent Gatto, Lisa Breckels

Examples

```
## Load an example MSnSet
library("pRolocdata")
data(dunkley2006)

## Adding a legend inside a plot
plot2D(dunkley2006)
addLegend(dunkley2006,  where = "topleft")

## Adding a legend outside a plot
par(mfrow = c(1, 2))
plot2D(dunkley2006)
addLegend(dunkley2006, where = "other")
```

addMarkers *Adds markers to the data*

Description

The function adds a 'markers' feature variable. These markers are read from a comma separated values (csv) spreadsheet file. This markers file is expected to have 2 columns (others are ignored) where the first is the name of the marker features and the second the group label. Alternatively, a markers named vector as provided by the pRolocmarkers function can also be used.

Usage

```
addMarkers(object, markers, mcol = "markers", fcol, verbose = TRUE)
```

Arguments

object	An instance of class MSnSet.
markers	A character with the name the markers' csv file or a named character of markers as provided by pRolocmarkers.
mcol	A character of length 1 defining the feature variable label for the newly added markers. Default is "markers".
fcol	An optional feature variable to be used to match against the markers. If missing, the feature names are used.
verbose	A logical indicating if number of markers and marker table should be printed to the console.

Details

It is essential to assure that featureNames(object) (or fcol, see below) and marker names (first column) match, i.e. the same feature identifiers and case fold are used.

Value

A new instance of class MSnSet with an additional markers feature variable.

Author(s)

Laurent Gatto

See Also

See pRolocmarkers for a list of spatial markers and markers for details about markers encoding.

Examples

```
library("pRolocdata")
data(dunkley2006)
atha <- pRolocmarkers("atha")
try(addMarkers(dunkley2006, atha)) ## markers already exists
fData(dunkley2006)$markers.org <- fData(dunkley2006)$markers
fData(dunkley2006)$markers <- NULL
marked <- addMarkers(dunkley2006, atha)
fvarLabels(marked)
## if 'makers' already exists
marked <- addMarkers(marked, atha, mcol = "markers2")
fvarLabels(marked)
stopifnot(all.equal(fData(marked)$markers, fData(marked)$markers2))
plot2D(marked)
addLegend(marked, where = "topleft", cex = .7)
```

AnnotationParams-class

Class "AnnotationParams"

Description

Class to store annotation parameters to automatically query a Biomart server, retrieve relevant annotation for a set of features of interest using, for example `getGOFromFeatures` and `makeGoSet`.

Objects from the Class

Objects can be created and set with the `setAnnotationParams` function. Object are created by calling without any arguments `setAnnotationParams()`, which will open an interactive interface. Depending on the value of `"many.graphics"` option, a graphical of a text-based menu will open (the text interface can be forced by setting the `graphics` argument to `FALSE`: `setAnnotationParams(graphics = FALSE)`). The menu will allow to select the species of interest first and the type of features (ENSEMBL gene identifier, Entrez id, ...) second.

The species that are available are those for which ENSEMBL data is available in Biomart and have a set of attributes of interest available. The compatible identifiers for downstream queries are then automatically filtered and displayed for user selection.

It is also possible to pass a parameter `inputs`, a character vector of length 2 containing a pattern uniquely matching the species of interest (in position 1) and a patterns uniquely matching the feature types (in position 2). If the matches are not unique, an error will be thrown.

A new instance of the `AnnotationParams` will be created to enable easy and automatic query of the `Mart` instance. The instance is invisibly returned and stored in a global variable in the **pRoloc** package's private environment for automatic retrieval. If a variable containing an `AnnotationParams` instance is already available, it can be set globally by passing it as argument to the `setAnnotationParams` function. Globally set `AnnotationParams` instances can be accessed with the `getAnnotationParams` function.

See the `pRoloc-theta` vignette for details.

Slots

`mart`: Object of class `"Mart"` from the **biomaRt** package.

`martname`: Object of class `"character"` with the name of the `mart` instance.

`dataset`: Object of class `"character"` with the data set of the `mart` instance.

`filter`: Object of class `"character"` with the filter to be used when querying the `mart` instance.

`date`: Object of class `"character"` indicating when the current instance was created.

`biomaRtVersion`: Object of class `"character"` with the **biomaRt** version used to create the `AnnotationParams` instance.

`.__classVersion__`: Object of class `"Versions"` with the version of the `AnnotationParams` class of the current instance.

Methods

show `signature(object = "AnnotationParams")`: to display objects.

Author(s)

Laurent Gatto <lg390@cam.ac.uk>

See Also

The `pRoloc-theta` vignette.

Examples

```
#data(andy2011params)
#data(dunkley2006params)
#andy2011params
#dunkley2006params

#try(setAnnotationParams(inputs = c("nomatch1", "nomatch2")))
#setAnnotationParams(inputs = c("Human genes",
#          "UniProtKB/Swiss-Prot ID"))
#getAnnotationParams()
```

---

checkFeatureNamesOverlap

Check feature names overlap

Description

Checks the marker and unknown feature overlap of two `MSnSet` instances.

Usage

```
checkFeatureNamesOverlap(x, y, fcolx = "markers", fcoly, verbose = TRUE)
```

Arguments

x	An MSnSet instance.
y	An MSnSet instance.
fcolx	The feature variable to separate unknown (fData(y)$coly == "unknown") from the marker features in the x object.
fcoly	As fcolx, for the y object. If missing, the value of fcolx is used.
verbose	If TRUE (default), the overlap is printed out on the console.

Value

Invisibly returns a named list of common markers, unique x markers, unique y markers in, common unknowns, unique x unknowns and unique y unknowns.

Author(s)

Laurent Gatto

Examples

```
library("pRolocdata")
data(andy2011)
data(andy2011goCC)
checkFeatureNamesOverlap(andy2011, andy2011goCC)
featureNames(andy2011goCC)[1] <- "ABC"
res <- checkFeatureNamesOverlap(andy2011, andy2011goCC)
res$markersX
res$markersY
```

---

checkFvarOverlap *Compare a feature variable overlap*

Description

Extracts qualitative feature variables from two MSnSet instances and compares with a contingency table.

Usage

```
checkFvarOverlap(x, y, fcolx = "markers", fcoly, verbose = TRUE)
```

Arguments

x	An MSnSet instance.
y	An MSnSet instance.
fcolx	The feature variable to separate unknown (fData(y)$coly == "unknown") from the marker features in the x object.
fcoly	As fcolx, for the y object. If missing, the value of fcolx is used.
verbose	If TRUE (default), the contingency table of the the feature variables is printed out.

Value

Invisibly returns a named list with the values of the diagonal, upper and lower triangles of the contingency table.

Author(s)

Laurent Gatto

Examples

```
library("pRolocdata")
data(dunkley2006)
res <- checkFvarOverlap(dunkley2006, dunkley2006,
                        "markers", "markers.orig")
str(res)
```

---

chi2-methods *The PCP 'chi square' method*

Description

In the original protein correlation profiling (PCP), Andersen et al. use the peptide normalised profiles along gradient fractions and compared them with the reference profiles (or set of profiles) by computing Chi^2 values, $\frac{\sum(x_i - x_p)^2}{x_p}$, where x_i is the normalised value of the peptide in fraction i and x_p is the value of the marker (from Wiese et al., 2007). The protein Chi^2 is then computed as the median of the peptide Chi^2 values. Peptides and proteins with similar profiles to the markers will have small Chi^2 values.

The `chi2` methods implement this idea and compute such Chi^2 values for sets of proteins.

Methods

`signature(x = "matrix", y = "matrix", method = "character", fun = "NULL", na.rm = "logical")` Compute `nrow(x)` times `nrow(y)` Chi^2 values, for each `x`, `y` feature pair. Method is one of `"Andersen2003"` or `"Wiese2007"`; the former (default) computed the Chi^2 as `sum(y-x)^2/length(x)`, while the latter uses `sum((y-x)^2/x)`. `na.rm` defines if missing values (`NA` and `NaN`) should be removed prior to summation. `fun` defines how to summarise the Chi^2 values; default, `NULL`, does not combine the Chi^2 values.

`signature(x = "matrix", y = "numeric", method = "character", na.rm = "logical")` Computes `nrow(x)` Chi^2 values, for all the (x_i, y) pairs. See above for the other arguments.

`signature(x = "numeric", y = "matrix", method = "character", na.rm = "logical")` Computes `nrow(y)` Chi^2 values, for all the (x, y_i) pairs. See above for the other arguments.

`signature(x = "numeric", y = "numeric", method = "character", na.rm = "logical")` Computes the Chi^2 value for the (x, y) pairs. See above for the other arguments.

Author(s)

Laurent Gatto <lg390@cam.ac.uk>

References

Andersen, J. S., Wilkinson, C. J., Mayor, T., Mortensen, P. et al., Proteomic characterization of the human centrosome by protein correlation profiling. Nature 2003, 426, 570 - 574.

Wiese, S., Gronemeyer, T., Ofman, R., Kunze, M. et al., Proteomics characterization of mouse kidney peroxisomes by tandem mass spectrometry and protein correlation profiling. Mol. Cell. Proteomics 2007, 6, 2045 - 2057.

See Also

empPvalues

Examples

```
mrk <- rnorm(6)
prot <- matrix(rnorm(60), ncol = 6)
chi2(mrk, prot, method = "Andersen2003")
chi2(mrk, prot, method = "Wiese2007")

pepmark <- matrix(rnorm(18), ncol = 6)
pepprot <- matrix(rnorm(60), ncol = 6)
chi2(pepmark, pepprot)
chi2(pepmark, pepprot, fun = sum)
```

---

classWeights *Calculate class weights*

Description

Calculates class weights to be used for parameter optimisation and classification such as svmOptimisation or svmClassification - see the *pRoloc tutorial* vignette for an example. The weights are calculated for all non-*unknown* classes the inverse of the number of observations.

Usage

```
classWeights(object, fcol = "markers")
```

Arguments

object	An instance of class MSnSet
fcol	The name of the features to be weighted

Value

A table of class weights

Author(s)

Laurent Gatto

Examples

```
library("pRolocdata")
data(hyperLOPIT2015)
classWeights(hyperLOPIT2015)
data(dunkley2006)
classWeights(dunkley2006)
```

---

clustDist *Pairwise Distance Computation for Protein Information Sets*

Description

This function computes the mean (normalised) pairwise distances for pre-defined sets of proteins.

Usage

```
clustDist(object, k = 1:5, fcol = "GOAnnotations", n = 5, verbose = TRUE, seed)
```

Arguments

object	An instance of class "MSnSet".
k	The number of clusters to try fitting to the protein set. Default is k = 1:5.
fcol	The feature meta-data containing matrix of protein sets/ marker definitions. Default is GOAnnotations.
n	The minimum number of proteins per set. If protein sets contain less than n instances they will be ignored. Defualt is 5.
verbose	A logical defining whether a progress bar is displayed.
seed	An optional seed for the random number generator.

Details

The input to the function is a MSnSet dataset containing a matrix appended to the feature data slot identifying the membership of protein instances to a pre-defined set(s) e.g. a specific Gene Ontology term etc.

For each protein set, the clustDist function (i) extracts all instances belonging to the set, (ii) using the kmeans algorithm fits and tests k = c(1:5) (default) cluster components to each set, (iii) calculates the mean pairwise distance for each k tested.

Note: currently distances are calcualted in Euclidean space, but other distance metrics will be supported in the future).

The output is a list of ClustDist objects, one per information cluster. The ClustDist class summarises the algorithm information such as the number of k's tested for the kmeans, and mean and normalised pairwise Euclidean distances per numer of component clusters tested. See ?ClustDist for more details.

Value

An instance of "ClustDistList" containing a "ClustDist" instance for every protein set, which summarises the algorithm information such as the number of k's tested for the kmeans, and mean and normalised pairwise Euclidean distances per numer of component clusters tested.

Author(s)

Lisa Breckels

See Also

For class definitions see "ClustDistList" and "ClustDist".

Examples

```
library(pRolocdata)
data(dunkley2006)
## Convert annotation data e.g. markers, to a matrix e.g. MM
xx <- mrkVecToMat(dunkley2006, vfcol = "markers", mfcol = "MM")
## get distances for protein sets
dd <- clustDist(xx, fcol = "MM", k = 1:3)
## plot clusters for first 'ClustDist' object
## in the 'ClustDistList'
plot(dd[[1]], xx)
## plot normalised distances for all protein sets
plot(dd)
## plot mean distances for all protein sets
plot(dd, method = "mean")
##' ## plot raw distances for all protein sets
plot(dd, method = "raw")
## Extract normalised distances
## Normalisation factor default is n^1/3
minDist <- getNormDist(dd)
## Get new order according to lowest distance
o <- order(minDist)
## Re-order annotations
fData(xx)$MM <- fData(xx)$MM[, o]
if (interactive()) {
pRolocVis(xx, fcol = "MM")
}
```

ClustDist-class *Class* "ClustDist"

Description

The ClustDist summaries algorithm information, from running the clustDist function, such as the number of k's tested for the kmeans, and mean and normalised pairwise (Euclidean) distances per numer of component clusters tested.

Objects from the Class

Object of this class are created with the `clustDist` function.

Slots

`k`: Object of class `"numeric"` storing the number of k clusters tested.

`dist`: Object of class `"list"` storing the list of distance matrices.

`term`: Object of class `"character"` describing GO term name.

`nrow`: Object of class `"numeric"` showing the number of instances in the set

`clustsz`: Object of class `"list"` describing the number of instances for each cluster for each k tested

`components`: Object of class `"vector"` storing the class membership of each protein for each k tested.

`fcol`: Object of class `"character"` showing the feature column name in the corresponding `MSnSet` where the protein set information is stored.

Methods

plot Plots the kmeans clustering results.

show Shows the object.

Author(s)

Lisa M Breckels <lms79@cam.ac.uk>

Examples

```
showClass("ClustDist")

library(pRolocdata)
data(dunkley2006)

## Convert annotation data e.g. markers, to a matrix e.g. MM
xx <- mrkVecToMat(dunkley2006, vfcol = "markers", mfcol = "MM")

## get distances for protein sets
dd <- clustDist(xx, fcol = "MM", k = 1:3)

## filter
xx <- filterMinMarkers(xx, n = 50, fcol = "MM")
xx <- filterMaxMarkers(xx, p = .25, fcol = "MM")

## get distances for protein sets
dd <- clustDist(xx, fcol = "MM")

## plot clusters for first 'ClustDist' object
## in the 'ClustDistList'
plot(dd[[1]], xx)
```

```
## plot distances for all protein sets
plot(dd)
```

ClustDistList-class *Storing multiple ClustDist instances*

Description

A class for storing lists of ClustDist instances.

Objects from the Class

Object of this class are created with the clustDist function.

Slots

x: Object of class list containing valid ClustDist instances.

log: Object of class list containing an object creation log, containing among other elements the call that generated the object.

.__classVersion__: The version of the instance. For development purposes only.

Methods

"[[" Extracts a single ClustDist at position.

"[" Extracts one of more ClustDists as ClustDistList.

length Returns the number of ClustDists.

names Returns the names of ClustDists, if available. The replacement method is also available.

show Display the object by printing a short summary.

lapply(x, FUN, ...) Apply function FUN to each element of the input x. If the application of FUN returns and ClustDist, then the return value is an ClustDistList, otherwise a list.

plot Plots a boxplot of the distance results per protein set.

Author(s)

Lisa M Breckels <lms79@cam.ac.uk>

Examples

```
library(pRolocdata)
data(dunkley2006)

## Convert annotation data e.g. markers, to a matrix e.g. MM
xx <- mrkVecToMat(dunkley2006, vfcol = "markers", mfcol = "MM")

## get distances for protein sets
```

```
dd <- clustDist(xx, fcol = "MM", k = 1:3)

## filter
xx <- filterMinMarkers(xx, n = 50, fcol = "MM")
xx <- filterMaxMarkers(xx, p = .25, fcol = "MM")

## get distances for protein sets
dd <- clustDist(xx, fcol = "MM")

## plot distances for all protein sets
plot(dd)

names(dd)

## Extract a sub-list of ClustDist objects
dd[1]

## Extract 1st ClustDist object
dd[[1]]
```

`empPvalues` *Estimate empirical p-values for $Chi\hat{}2$ protein correlations.*

Description

Andersen et al. (2003) used a fixed Chi^2 threshold of 0.05 to identify organelle-specific candidates. This function computes empirical p-values by permutation the markers relative intensities and computed null Chi^2 values.

Usage

```
empPvalues(marker, corMatrix, n = 100, ...)
```

Arguments

`marker`	A `numerics` with markers relative intensities.
`corMatrix`	A `matrix` of `nrow(corMatrix)` protein relative intensities to be compares against the marker.
`n`	The number of iterations.
`...`	Additional parameters to be passed to `chi2`.

Value

A `numeric` of length `nrow(corMatrix)`.

Author(s)

Laurent Gatto <lg390@cam.ac.uk>

References

Andersen, J. S., Wilkinson, C. J., Mayor, T., Mortensen, P. et al., Proteomic characterization of the human centrosome by protein correlation profiling. Nature 2003, 426, 570 - 574.

See Also

chi2 for Chi^2 calculation.

Examples

```
set.seed(1)
mrk <- rnorm(6, 5, 1)
prot <- rbind(matrix(rnorm(120, 5, 1), ncol = 6),
              mrk + rnorm(6))
mrk <- mrk/sum(mrk)
prot <- prot/rowSums(prot)
empPvalues(mrk, prot)
```

---

fDataToUnknown *Update a feature variable*

Description

This function replaces a string or regular expression in a feature variable using the sub function.

Usage

```
fDataToUnknown(object, fcol = "markers", from = "^$", to = "unknown", ...)
```

Arguments

object	An instance of class MSnSet.
fcol	Feature variable to be modified. Default is "markers". If NULL, all feature variables will updated.
from	A character defining the string or regular expression of the pattern to be replaced. Default is the empty string, i.e. the regular expression "^$". See sub for details. If NA, then NA values are replaced by to.
to	A replacement for matched pattern. Default is "unknown". See sub for details.
...	Additional arguments passed to sub.

Value

An updated MSnSet.

Author(s)

Laurent Gatto

Examples

```
library("pRolocdata")
data(dunkley2006)
getMarkers(dunkley2006, "markers")
dunkley2006 <- fDataToUnknown(dunkley2006,
                              from = "unknown", to = "unassigned")
getMarkers(dunkley2006, "markers")
```

`filterBinMSnSet` *Filter a binary MSnSet*

Description

Removes columns or rows that have a certain proportion or absolute number of 0 values.

Usage

```
filterBinMSnSet(object, MARGIN = 2, t, q, verbose = TRUE)
```

Arguments

`object`	An `MSnSet`
`MARGIN`	1 or 2. Default is 2.
`t`	Rows/columns that have `t` or less 1s, it will be filtered out. When `t` and `q` are missing, default is to use `t = 1`.
`q`	If a row has a higher quantile than defined by `q`, it will be filtered out.
`verbose`	A `logical` defining of a message is to be printed. Default is `TRUE`.

Value

A filtered `MSnSet`.

Author(s)

Laurent Gatto

See Also

`zerosInBinMSnSet`, `filterZeroCols`, `filterZeroRows`.

Examples

```
set.seed(1)
m <- matrix(sample(0:1, 25, replace=TRUE), 5)
m[1, ] <- 0
m[, 1] <- 0
rownames(m) <- colnames(m) <- letters[1:5]
fd <- data.frame(row.names = letters[1:5])
x <- MSnSet(exprs = m, fData = fd, pData = fd)
exprs(x)
## Remove columns with no 1s
exprs(filterBinMSnSet(x, MARGIN = 2, t = 0))
## Remove columns with one 1 or less
exprs(filterBinMSnSet(x, MARGIN = 2, t = 1))
## Remove columns with two 1s or less
exprs(filterBinMSnSet(x, MARGIN = 2, t = 2))
## Remove columns with three 1s
exprs(filterBinMSnSet(x, MARGIN = 2, t = 3))
## Remove columns that have half or less of 1s
exprs(filterBinMSnSet(x, MARGIN = 2, q = 0.5))
```

---

filterMaxMarkers *Removes class/annotation information from a matrix of candidate markers that appear in the* fData.

Description

Removes annotation information that contain more that a certain number/percentage of proteins

Usage

```
filterMaxMarkers(object, n, p = 0.2, fcol = "GOAnnotations", verbose = TRUE)
```

Arguments

object	An instance of class MSnSet.
n	Maximum number of proteins allowed per class/information term.
p	Maximum percentage of proteins per column. Default is 0.2 i.e. remove columns that have information for greater than 20 of the total number of proteins in the dataset (note: this is useful for example, if information is GO terms, for removing very general and uninformative terms).
fcol	The name of the matrix of marker information. Default is GOAnnotations.
verbose	Number of marker candidates retained after filtering.

Value

An updated MSnSet

See Also

filterMinMarkers and example therein.

---

filterMinMarkers *Removes class/annotation information from a matrix of candidate markers that appear in the* fData.

Description

Removes annotation information that contain less that a certain number/percentage of proteins

Usage

```
filterMinMarkers(object, n = 10, p, fcol = "GOAnnotations", verbose = TRUE)
```

Arguments

object	An instance of class MSnSet.
n	Minimum number of proteins allowed per column. Default is 10.
p	Minimum percentage of proteins per column.
fcol	The name of the matrix of marker information. Default is GOAnnotations.
verbose	Number of marker candidates retained after filtering.

Value

An updated MSnSet.

Author(s)

Lisa M Breckels

Examples

```
library(pRolocdata)
data(dunkley2006)
xx <- dunkley2006
## create a matrix of markers
xx <- mrkVecToMat(xx, vfcol = "markers", mfcol = "Markers")
## Remove marker classes with less than 15 members, from matrix of markers
xx <- filterMinMarkers(xx, n = 15, fcol = "Markers")
## Remove marker classes with more than 50 members, from matrix of markers
xx <- filterMaxMarkers(xx, p = .2, fcol = "Markers")
```

filterZeroCols *Remove 0 columns/rows*

Description

Removes all assay data columns/rows that are composed of only 0, i.e. have a `colSum`/`rowSum` of 0.

Usage

```
filterZeroCols(object, verbose = TRUE)

filterZeroRows(object, verbose = TRUE)
```

Arguments

`object`	A `MSnSet` object.
`verbose`	Print a message with the number of filtered out columns/row (if any).

Value

An `MSnSet`.

Author(s)

Laurent Gatto

Examples

```
library("pRolocdata")
data(andy2011goCC)
any(colSums(exprs(andy2011goCC)) == 0)
exprs(andy2011goCC)[, 1:5] <- 0
ncol(andy2011goCC)
ncol(filterZeroCols(andy2011goCC))
```

GenRegRes-class *Class "GenRegRes" and "ThetaRegRes"*

Description

Regularisation framework containers.

Objects from the Class

Object of this class are created with the respective regularisation function: `knnOptimisation`, `svmOptimisation`, `plsdaOptimisation`, `knntlOptimisation`, ...

Slots

`algorithm`**:** Object of class `"character"` storing the machine learning algorithm name.

`hyperparameters`**:** Object of class `"list"` with the respective algorithm hyper-parameters tested.

`design`**:** Object of class `"numeric"` describing the cross-validation design, the test data size and the number of replications.

`log`**:** Object of class `"list"` with warnings thrown during the hyper-parameters regularisation.

`seed`**:** Object of class `"integer"` with the random number generation seed.

`results`**:** Object of class `"matrix"` of dimenstions `times` (see `design`) by number of hyperparameters + 1 storing the macro F1 values for the respective best hyper-parameters for each replication.

`f1Matrices`**:** Object of class `"list"` with respective `times` cross-validation F1 matrices.

`cmMatrices`**:** Object of class `"list"` with respective `times` contingency matrices.

`testPartitions`**:** Object of class `"list"` with respective `times` test partitions.

`datasize`**:** Object of class `"list"` with details about the respective inner and outter training and testing data sizes.

Only in `ThetaRegRes`:

`predictions`**:** A `list` of predictions for the optimisation iterations.

`otherWeights`**:** Alternative best theta weigts: a vector per iterations, `NULL` if no other best weights were found.

Methods

getF1Scores Returns a matrix of F1 scores for the optimisation parameters.

f1Count `signature(object = "GenRegRes", t = "numeric")` and `signature(object = "ThetaRegRes", t = "numeric")`: Constructs a table of all possible parameter combination and count how many have an F1 scores greater or equal than `t`. When `t` is missing (default), the best F1 score is used. This method is useful in conjunctin with `plot`.

getParams Returns the *best* parameters. It is however strongly recommended to inspect the optimisation results. For a `ThetaRegRes` optimisation result, the method to chose the best parameters can be `"median"` (default) or `"mean"` (the median or mean of the best weights is chosen), `"max"` (the first weights with the highest macro-F1 score, considering that multiple max scoring combinations are possible) or `"count"` (the observed weight that get the maximum number of observations, see `f1Count`). The `favourP` argument can be used to prioritise weights that favour the primary data (i.e. heigh weights). See `favourPrimary` below.

getSeed Returns the seed used for the optimisation run.

getWarnings `signature(object = "GenRegRes")`: Returns a vector of recorded warnings.

levelPlot `signature(object = "GenRegRes")`: Plots a heatmap of of the optimisation results. Only for `"GenRegRes"` instances.

plot Plots the optisisation results.

show Shows the object.

Other functions

Only for ThetaRegRes:

`combineThetaRegRes(object)` Takes a `list` of `ThetaRegRes` instances to be combined and returnes a new `ThetaRegRes` instance.

`favourPrimary(primary, auxiliary, object, verbose = TRUE)` Takes the `primary` and `auxiliary` data sources (two `MSnSet` instances) and a `ThetaRegRes` object and returns and updated `ThetaRegRes` instance containing best parameters/weigths (see the `getParams` function) favouring the primary data when multiple best theta weights are available.

Author(s)

Laurent Gatto <lg390@cam.ac.uk>

Examples

```
showClass("GenRegRes")
showClass("ThetaRegRes")
```

---

getMarkerClasses *Returns the organelle classes in an 'MSnSet'*

Description

Convenience accessor to the organelle classes in an 'MSnSet'. This function returns the organelle classes of an `MSnSet` instance. As a side effect, it prints out the classes.

Usage

```
getMarkerClasses(object, fcol = "markers", ...)
```

Arguments

`object`	An instance of class `"MSnSet"`.
`fcol`	The name of the markers column in the `featureData` slot. Default is `markers`.
`...`	Additional parameters passed to `sort` from the base package.

Value

A `character` vector of the organelle classes in the data.

Author(s)

Lisa Breckels and Laurent Gatto

See Also

getMarkers to extract the marker proteins. See markers for details about spatial markers storage and encoding.

Examples

```
library("pRolocdata")
data(dunkley2006)
organelles <- getMarkerClasses(dunkley2006)
## same if markers encoded as a matrix
dunkley2006 <- mrkVecToMat(dunkley2006, mfcol = "Markers")
organelles2 <- getMarkerClasses(dunkley2006, fcol = "Markers")
stopifnot(all.equal(organelles, organelles2))
```

---

getMarkers *Get the organelle markers in an* MSnSet

Description

Convenience accessor to the organelle markers in an MSnSet. This function returns the organelle markers of an MSnSet instance. As a side effect, it print out a marker table.

Usage

```
getMarkers(object, fcol = "markers", names = TRUE, verbose = TRUE)
```

Arguments

object	An instance of class "MSnSet".
fcol	The name of the markers column in the featureData slot. Default is "markers".
names	A logical indicating if the markers vector should be named. Ignored if markers are encoded as a matrix.
verbose	If TRUE, a marker table is printed and the markers are returned invisibly. If FALSE, the markers are returned.

Value

A character (matrix) of length (ncol) ncol(object), depending on the vector or matrix encoding of the markers.

Author(s)

Laurent Gatto

See Also

See getMarkerClasses to get the classes only. See markers for details about spatial markers storage and encoding.

Examples

```
library("pRolocdata")
data(dunkley2006)
## marker vectors
myVmarkers <- getMarkers(dunkley2006)
head(myVmarkers)
## marker matrix
dunkley2006 <- mrkVecToMat(dunkley2006, mfcol = "Markers")
myMmarkers <- getMarkers(dunkley2006, fcol = "Markers")
head(myMmarkers)
```

---

getNormDist *Extract Distances from a* "ClustDistList" *object*

Description

This function computes and outputs normalised distances from a "ClustDistList" object.

Usage

```
getNormDist(object, p = 1/3)
```

Arguments

object	An instance of class "ClustDistList".
p	The normalisation factor. Default is 1/3.

Value

An numeric of normalised distances, one per protein set in the ClustDistList.

Author(s)

Lisa Breckels

See Also

"ClustDistList", "ClustDist", and examples in clustDist.

Examples

```
library(pRolocdata)
data(dunkley2006)
## Convert annotation data e.g. markers, to a matrix e.g. MM
xx <- mrkVecToMat(dunkley2006, vfcol = "markers", mfcol = "MM")
## get distances for protein sets
dd <- clustDist(xx, fcol = "MM", k = 1:3)
## plot clusters for first 'ClustDist' object
## in the 'ClustDistList'
```

```
plot(dd[[1]], xx)
## plot normalised distances for all protein sets
plot(dd)
## plot mean distances for all protein sets
plot(dd, method = "mean")
## plot raw distances for all protein sets
plot(dd, method = "raw")
## Extract normalised distances
## Normalisation factor default is n^1/3
minDist <- getNormDist(dd)
## Get new order according to lowest distance
o <- order(minDist)
## Re-order annotations
fData(xx)$MM <- fData(xx)$MM[, o]
if (interactive()) {
pRolocVis(xx, fcol = "MM")
}
```

`getPredictions` *Returns the predictions in an 'MSnSet'*

Description

Convenience accessor to the predicted feature localisation in an 'MSnSet'. This function returns the predictions of an `MSnSet` instance. As a side effect, it prints out a prediction table.

Usage

```
getPredictions(object, fcol, scol, mcol = "markers", t = 0, verbose = TRUE)
```

Arguments

`object`	An instance of class `"MSnSet"`.
`fcol`	The name of the prediction column in the `featureData` slot.
`scol`	The name of the prediction score column in the `featureData` slot. If missing, created by pasting '.scores' after `fcol`.
`mcol`	The feature meta data column containing the labelled training data.
`t`	The score threshold. Predictions with score < t are set to 'unknown'. Default is 0. It is also possible to define thresholds for each prediction class, in which case, `t` is a named numeric with names exactly matching the unique prediction class names.
`verbose`	If `TRUE`, a prediction table is printed and the predictions are returned invisibly. If `FALSE`, the predictions are returned.

Value

An instance of class "MSnSet" with `fcol.pred` feature variable storing the prediction results according to the chosen threshold.

Author(s)

Laurent Gatto and Lisa Breckels

See Also

orgQuants for calculating organelle-specific thresholds.

Examples

```
library("pRolocdata")
data(dunkley2006)
res <- svmClassification(dunkley2006, fcol = "pd.markers",
                         sigma = 0.1, cost = 0.5)
fData(res)$svm[500:510]
fData(res)$svm.scores[500:510]
getPredictions(res, fcol = "svm", t = 0) ## all predictions
getPredictions(res, fcol = "svm", t = .9) ## single threshold
## 50% top predictions per class
ts <- orgQuants(res, fcol = "svm", t = .5)
getPredictions(res, fcol = "svm", t = ts)
```

---

highlightOnPlot *Highlight features of interest on a spatial proteomics plot*

Description

Highlights a set of features of interest given as a FeaturesOfInterest instance on a PCA plot produced by plot2D or plot3D. If none of the features of interest are found in the MSnset's featureNames, an warning is thrown.

Usage

```
highlightOnPlot(object, foi, labels, args = list(), ...)

highlightOnPlot3D(object, foi, labels, args = list(), radius = 0.1 * 3, ...)
```

Arguments

object	The main dataset described as an MSnSet or a matrix with the coordinates of the features on the PCA plot produced (and invisibly returned) by plot2D.
foi	An instance of FeaturesOfInterest, or, alternatively, a character of feautre names.
labels	A character of length 1 with a feature variable name to be used to label the features of interest. This is only valid if object is an MSnSet. Alternatively, if TRUE, then featureNames(object) (or rownames(object), if object is a matrix) are used. Default is missing, which does not add any label.s

args	A named list of arguments to be passed to plot2D if the PCA coordinates are to be calculated. Ignored if the PCA coordinates are passed directly, i.e. object is a matrix.
...	Additional parameters passed to points or text (when labels is TRUE) when adding to plot2D, or spheres3d or text3d when adding the plot3D
radius	Radius of the spheres to be added to the visualisation produced by plot3D. Default is 0.3 (i.e plot3D's radius1 * 3), to emphasise the features with regard to uknown (radius1 = 0.1) and marker (radius1 * 2) features.

Value

NULL; used for its side effects.

Author(s)

Laurent Gatto

Examples

```
library("pRolocdata")
data("tan2009r1")
x <- FeaturesOfInterest(description = "A test set of features of interest",
                        fnames = featureNames(tan2009r1)[1:10],
                        object = tan2009r1)

## using FeaturesOfInterest or feature names
par(mfrow = c(2, 1))
plot2D(tan2009r1)
highlightOnPlot(tan2009r1, x)
plot2D(tan2009r1)
highlightOnPlot(tan2009r1, featureNames(tan2009r1)[1:10])

.pca <- plot2D(tan2009r1)
head(.pca)
highlightOnPlot(.pca, x, col = "red")
highlightOnPlot(tan2009r1, x, col = "red", cex = 1.5)
highlightOnPlot(tan2009r1, x, labels = TRUE)

.pca <- plot2D(tan2009r1, dims = c(1, 3))
highlightOnPlot(.pca, x, pch = "+", dims = c(1, 3))
highlightOnPlot(tan2009r1, x, args = list(dims = c(1, 3)))

.pca2 <- plot2D(tan2009r1, mirrorX = TRUE, dims = c(1, 3))
## previous pca matrix, need to mirror X axis
highlightOnPlot(.pca, x, pch = "+", args = list(mirrorX = TRUE))
## new pca matrix, with X mirrors (and 1st and 3rd PCs)
highlightOnPlot(.pca2, x, col = "red")

plot2D(tan2009r1)
highlightOnPlot(tan2009r1, x)
highlightOnPlot(tan2009r1, x, labels = TRUE, pos = 3)
```

```
highlightOnPlot(tan2009r1, x, labels = "Flybase.Symbol", pos = 1)

## in 3 dimensions
if (interactive()) {
  plot3D(tan2009r1, radius1 = 0.05)
  highlightOnPlot3D(tan2009r1, x, labels = TRUE)
  highlightOnPlot3D(tan2009r1, x)
}
```

`knnClassification` *knn classification*

Description

Classification using for the k-nearest neighbours algorithm.

Usage

```
knnClassification(
  object,
  assessRes,
  scores = c("prediction", "all", "none"),
  k,
  fcol = "markers",
  ...
)
```

Arguments

`object`	An instance of class `"MSnSet"`.
`assessRes`	An instance of class `"GenRegRes"`, as generated by `knnOptimisation`.
`scores`	One of `"prediction"`, `"all"` or `"none"` to report the score for the predicted class only, for all classes or none.
`k`	If `assessRes` is missing, a `k` must be provided.
`fcol`	The feature meta-data containing marker definitions. Default is `markers`.
`...`	Additional parameters passed to `knn` from package `class`.

Value

An instance of class `"MSnSet"` with `knn` and `knn.scores` feature variables storing the classification results and scores respectively.

Author(s)

Laurent Gatto

Examples

```
library(pRolocdata)
data(dunkley2006)
## reducing parameter search space and iterations
params <- knnOptimisation(dunkley2006, k = c(3, 10), times = 3)
params
plot(params)
f1Count(params)
levelPlot(params)
getParams(params)
res <- knnClassification(dunkley2006, params)
getPredictions(res, fcol = "knn")
getPredictions(res, fcol = "knn", t = 0.75)
plot2D(res, fcol = "knn")
```

`knnOptimisation` *knn parameter optimisation*

Description

Classification parameter optimisation for the k-nearest neighbours algorithm.

Usage

```
knnOptimisation(
  object,
  fcol = "markers",
  k = seq(3, 15, 2),
  times = 100,
  test.size = 0.2,
  xval = 5,
  fun = mean,
  seed,
  verbose = TRUE,
  ...
)
```

Arguments

`object`	An instance of class "MSnSet".
`fcol`	The feature meta-data containing marker definitions. Default is `markers`.
`k`	The hyper-parameter. Default values are `seq(3, 15, 2)`.
`times`	The number of times internal cross-validation is performed. Default is 100.
`test.size`	The size of test data. Default is 0.2 (20 percent).
`xval`	The n-cross validation. Default is 5.
`fun`	The function used to summarise the `xval` macro F1 matrices.

seed	The optional random number generator seed.
verbose	A logical defining whether a progress bar is displayed.
...	Additional parameters passed to knn from package class.

Details

Note that when performance scores precision, recall and (macro) F1 are calculated, any NA values are replaced by 0. This decision is motivated by the fact that any class that would have either a NA precision or recall would result in an NA F1 score and, eventually, a NA macro F1 (i.e. mean(F1)). Replacing NAs by 0s leads to F1 values of 0 and a reduced yet defined final macro F1 score.

Value

An instance of class "GenRegRes".

Author(s)

Laurent Gatto

See Also

knnClassification and example therein.

Examples

```
library(pRolocdata)
data(dunkley2006)
## reducing parameter search space and iterations
params <- knnOptimisation(dunkley2006, k = c(3, 10), times = 3)
params
plot(params)
f1Count(params)
levelPlot(params)
getParams(params)
res <- knnClassification(dunkley2006, params)
getPredictions(res, fcol = "knn")
getPredictions(res, fcol = "knn", t = 0.75)
plot2D(res, fcol = "knn")
```

---

knntlClassification *knn transfer learning classification*

Description

Classification using a variation of the KNN implementation of Wu and Dietterich's transfer learning schema

Usage

```
knntlClassification(
  primary,
  auxiliary,
  fcol = "markers",
  bestTheta,
  k,
  scores = c("prediction", "all", "none"),
  seed
)
```

Arguments

primary	An instance of class "MSnSet".
auxiliary	An instance of class "MSnSet".
fcol	The feature meta-data containing marker definitions. Default is markers.
bestTheta	Best theta vector as output from knntlOptimisation, see knntlOptimisation for details
k	Numeric vector of length 2, containing the best k parameters to use for the primary and auxiliary datasets. If k k is not specified it will be calculated internally.
scores	One of "prediction", "all" or "none" to report the score for the predicted class only, for all classes or none.
seed	The optional random number generator seed.

Value

A character vector of the classifications for the unknowns

Author(s)

Lisa Breckels

See Also

knntlOptimisation

Examples

```
## Load example primary and auxiliary data from pRolocdata
library(pRolocdata)
data(andy2011)
data(andy2011goCC)

## reducing calculation time of k by pre-running knnOptimisation
x <- c(andy2011, andy2011goCC)
k <- lapply(x, function(z)
            knnOptimisation(z, times=5,
                            fcol = "markers.orig",
```

```
                            verbose = FALSE))
k <- sapply(k, function(z) getParams(z))

## Use by = 1 in optimisation i.e. give full weight to the
## primary (indicated by 1) or full weight to auxiliary
## (indicated by 0) reducing parameter search for example
## in this documentation only. See the transfer learning
## vignette for examples and details.
opt <- knntlOptimisation(andy2011, andy2011goCC,
                         fcol = "markers.orig",
                         times = 2,
                         by = 1,
                         k = k)
th <- getParams(opt)
plot(opt)

## Now perform classification after finding the best weights
res <- knntlClassification(andy2011, andy2011goCC,
                           fcol = "markers.orig",
                           th,
                           k)
```

---

knntlOptimisation *theta parameter optimisation*

Description

Classification parameter optimisation for the KNN implementation of Wu and Dietterich's transfer learning schema

Usage

```
knntlOptimisation(
  primary,
  auxiliary,
  fcol = "markers",
  k,
  times = 50,
  test.size = 0.2,
  xval = 5,
  by = 0.5,
  length.out,
  th,
  xfolds,
  BPPARAM = BiocParallel::bpparam(),
  method = "Breckels",
  log = FALSE,
  seed
)
```

Arguments

primary	An instance of class "MSnSet".
auxiliary	An instance of class "MSnSet".
fcol	The feature meta-data containing marker definitions. Default is markers.
k	Numeric vector of length 2, containing the best k parameters to use for the primary (k[1]) and auxiliary (k[2]) datasets. See knnOptimisation for generating best k.
times	The number of times cross-validation is performed. Default is 50.
test.size	The size of test (validation) data. Default is 0.2 (20 percent).
xval	The number of rounds of cross-validation to perform.
by	The increment for theta, must be one of c(1, 0.5,0.25, 0.2, 0.15, 0.1, 0.05)
length.out	Alternative to using by parameter. Specifies the desired length of the sequence of theta to test.
th	A matrix of theta values to test for each class as generated from the function thetas, the number of columns should be equal to the number of classes contained in fcol. Note: columns will be ordered according to getMarkerClasses(primary, fcol). This argument is only valid if the default method 'Breckels' is used.
xfolds	Option to pass specific folds for the cross validation.
BPPARAM	Required for parallelisation. If not specified selects a default BiocParallelParam, from global options or, if that fails, the most recently registered() back-end.
method	The k-NN transfer learning method to use. The default is 'Breckels' as described in the Breckels et al (2016). If 'Wu' is specificed then the original method implemented Wu and Dietterich (2004) is implemented.
log	A logical defining whether logging should be enabled. Default is FALSE. Note that logging produes considerably bigger objects.
seed	The optional random number generator seed.

Details

knntlOptimisation implements a variation of Wu and Dietterich's transfer learning schema: P. Wu and T. G. Dietterich. Improving SVM accuracy by training on auxiliary data sources. In Proceedings of the Twenty-First International Conference on Machine Learning, pages 871 - 878. Morgan Kaufmann, 2004. A grid search for the best theta is performed.

Value

A list of containing the theta combinations tested, associated macro F1 score and accuracy for each combination over each round (specified by times).

Author(s)

Lisa Breckels

References

Breckels LM, Holden S, Wonjar D, Mulvey CM, Christoforou A, Groen AJ, Kohlbacher O, Lilley KS, Gatto L. Learning from heterogeneous data sources: an application in spatial proteomics. bioRxiv. doi: http://dx.doi.org/10.1101/022152

Wu P, Dietterich TG. Improving SVM Accuracy by Training on Auxiliary Data Sources. Proceedings of the 21st International Conference on Machine Learning (ICML); 2004.

See Also

knntlClassification and example therein.

Examples

```
## Load example primary and auxiliary data from pRolocdata
library(pRolocdata)
data(andy2011)
data(andy2011goCC)

## reducing calculation time of k by pre-running knnOptimisation
x <- c(andy2011, andy2011goCC)
k <- lapply(x, function(z)
            knnOptimisation(z, times=5,
                            fcol = "markers.orig",
                            verbose = FALSE))
k <- sapply(k, function(z) getParams(z))

## Use by = 1 in optimisation i.e. give full weight to the
## primary (indicated by 1) or full weight to auxiliary
## (indicated by 0) reducing parameter search for example
## in this documentation only. See the transfer learning
## vignette for examples and details.
opt <- knntlOptimisation(andy2011, andy2011goCC,
                         fcol = "markers.orig",
                         times = 2,
                         by = 1,
                         k = k)
th <- getParams(opt)
plot(opt)

## Now perform classification after finding the best weights
res <- knntlClassification(andy2011, andy2011goCC,
                           fcol = "markers.orig",
                           th,
                           k)
```

---

ksvmClassification *ksvm classification*

Description

Classification using the support vector machine algorithm.

Usage

```
ksvmClassification(
  object,
  assessRes,
  scores = c("prediction", "all", "none"),
  cost,
  fcol = "markers",
  ...
)
```

Arguments

object	An instance of class "MSnSet".
assessRes	An instance of class "GenRegRes", as generated by ksvmOptimisation.
scores	One of "prediction", "all" or "none" to report the score for the predicted class only, for all classes or none.
cost	If assessRes is missing, a cost must be provided.
fcol	The feature meta-data containing marker definitions. Default is markers.
...	Additional parameters passed to ksvm from package kernlab.

Value

An instance of class "MSnSet" with ksvm and ksvm.scores feature variables storing the classification results and scores respectively.

Author(s)

Laurent Gatto

Examples

```
library(pRolocdata)
data(dunkley2006)
## reducing parameter search space and iterations
params <- ksvmOptimisation(dunkley2006, cost = 2^seq(-1,4,5), times = 3)
params
plot(params)
f1Count(params)
levelPlot(params)
getParams(params)
res <- ksvmClassification(dunkley2006, params)
getPredictions(res, fcol = "ksvm")
getPredictions(res, fcol = "ksvm", t = 0.75)
plot2D(res, fcol = "ksvm")
```

ksvmOptimisation *ksvm parameter optimisation*

Description

Classification parameter optimisation for the support vector machine algorithm.

Usage

```
ksvmOptimisation(
  object,
  fcol = "markers",
  cost = 2^(-4:4),
  times = 100,
  test.size = 0.2,
  xval = 5,
  fun = mean,
  seed,
  verbose = TRUE,
  ...
)
```

Arguments

object	An instance of class "MSnSet".
fcol	The feature meta-data containing marker definitions. Default is markers.
cost	The hyper-parameter. Default values are 2^-4:4.
times	The number of times internal cross-validation is performed. Default is 100.
test.size	The size of test data. Default is 0.2 (20 percent).
xval	The n-cross validation. Default is 5.
fun	The function used to summarise the xval macro F1 matrices.
seed	The optional random number generator seed.
verbose	A logical defining whether a progress bar is displayed.
...	Additional parameters passed to ksvm from package kernlab.

Details

Note that when performance scores precision, recall and (macro) F1 are calculated, any NA values are replaced by 0. This decision is motivated by the fact that any class that would have either a NA precision or recall would result in an NA F1 score and, eventually, a NA macro F1 (i.e. mean(F1)). Replacing NAs by 0s leads to F1 values of 0 and a reduced yet defined final macro F1 score.

Value

An instance of class "GenRegRes".

Author(s)

Laurent Gatto

See Also

ksvmClassification and example therein.

Examples

```
library(pRolocdata)
data(dunkley2006)
## reducing parameter search space and iterations
params <- ksvmOptimisation(dunkley2006, cost = 2^seq(-1,4,5), times = 3)
params
plot(params)
f1Count(params)
levelPlot(params)
getParams(params)
res <- ksvmClassification(dunkley2006, params)
getPredictions(res, fcol = "ksvm")
getPredictions(res, fcol = "ksvm", t = 0.75)
plot2D(res, fcol = "ksvm")
```

MAPParams-class *The 'logPosteriors' function can be used to extract the log-posteriors at each iteration of the EM algorithm to check for convergence.*

Description

These functions implement the T augmented Gaussian mixture (TAGM) model for mass spectrometry-based spatial proteomics datasets using the maximum a posteriori (MAP) optimisation routine.

Usage

```
## S4 method for signature 'MAPParams'
show(object)

logPosteriors(x)

tagmMapTrain(
  object,
  fcol = "markers",
  method = "MAP",
  numIter = 100,
  mu0 = NULL,
  lambda0 = 0.01,
  nu0 = NULL,
  S0 = NULL,
```

```
  beta0 = NULL,
  u = 2,
  v = 10,
  seed = NULL
)

tagmMapPredict(
  object,
  params,
  fcol = "markers",
  probJoint = FALSE,
  probOutlier = TRUE
)
```

Arguments

object	An MSnbase::MSnSet containing the spatial proteomics data to be passed to tagmMapTrain and tagmPredict.
x	An object of class 'MAPParams'.
fcol	The feature meta-data containing marker definitions. Default is markers.
method	A charachter() describing the inference method for the TAGM algorithm. Default is "MAP".
numIter	The number of iterations of the expectation-maximisation algorithm. Default is 100.
mu0	The prior mean. Default is colMeans of the expression data.
lambda0	The prior shrinkage. Default is 0.01.
nu0	The prior degreed of freedom. Default is ncol(exprs(object)) + 2
S0	The prior inverse-wishary scale matrix. Empirical prior used by default.
beta0	The prior Dirichlet distribution concentration. Default is 1 for each class.
u	The prior shape parameter for Beta(u, v). Default is 2
v	The prior shape parameter for Beta(u, v). Default is 10.
seed	The optional random number generator seed.
params	An instance of class MAPParams, as generated by tagmMapTrain().
probJoint	A logical(1) indicating whether to return the joint probability matrix, i.e. the probability for all classes as a new tagm.map.joint feature variable.
probOutlier	A logical(1) indicating whether to return the probability of being an outlier as a new tagm.map.outlier feature variable. A high value indicates that the protein is unlikely to belong to any annotated class (and is hence considered an outlier).

Details

The tagmMapTrain function generates the MAP parameters (object or class MAPParams) based on an annotated quantitative spatial proteomics dataset (object of class MSnbase::MSnSet). Both are then

passed to the tagmPredict function to predict the sub-cellular localisation of protein of unknown localisation. See the *pRoloc-bayesian* vignette for details and examples. In this implementation, if numerical instability is detected in the covariance matrix of the data a small multiple of the identity is added. A message is printed if this conditioning step is performed.

Value

tagmMapTrain returns an instance of class MAPParams().

tagmPredict returns an instance of class MSnbase::MSnSet containing the localisation predictions as a new tagm.map.allocation feature variable.

Slots

method A character() storing the TAGM method name.

priors A list() with the priors for the parameters

seed An integer() with the random number generation seed.

posteriors A list() with the updated posterior parameters and log-posterior of the model.

datasize A list() with details about size of data

Author(s)

Laurent Gatto

Oliver M. Crook

References

A Bayesian Mixture Modelling Approach For Spatial Proteomics Oliver M Crook, Claire M Mulvey, Paul D. W. Kirk, Kathryn S Lilley, Laurent Gatto bioRxiv 282269; doi: https://doi.org/10.1101/282269

See Also

The plotEllipse() function can be used to visualise TAGM models on PCA plots with ellipses. The tagmMapTrain() function to use the TAGM MAP method.

Examples

```
## Load example data
library(pRolocdata)
data(dunkley2006)

## Generate MAP parameters (use numIter = 5 for example only)
par <- tagmMapTrain(dunkley2006, numIter = 5)

## Perform classification
dunkley2006 <- tagmMapPredict(dunkley2006,
                              params = par,
                              probJoint = TRUE,
                              probOutlier = TRUE)
```

markerMSnSet *Extract marker/unknown subsets*

Description

These function extract the marker or unknown proteins into a new MSnSet.

Usage

```
markerMSnSet(object, fcol = "markers")

unknownMSnSet(object, fcol = "markers")
```

Arguments

object	An instance of class MSnSet
fcol	The name of the feature data column, that will be used to separate the markers from the proteins of unknown localisation. When the markers are encoded as vectors, features of unknown localisation are defined as fData(object)[, fcol] == "unknown". For matrix-encoded markers, unlabelled proteins are defined as rowSums(fData(object)[, fcol]) == 0. Default is "markers".

Value

An new MSnSet with marker/unknown proteins only.

Author(s)

Laurent Gatto

See Also

sampleMSnSet testMSnSet and markers for markers encoding.

Examples

```
library("pRolocdata")
data(dunkley2006)
mrk <- markerMSnSet(dunkley2006)
unk <- unknownMSnSet(dunkley2006)
dim(dunkley2006)
dim(mrk)
dim(unk)
table(fData(dunkley2006)$markers)
table(fData(mrk)$markers)
table(fData(unk)$markers)
## matrix-encoded markers
dunkley2006 <- mrkVecToMat(dunkley2006)
dim(markerMSnSet(dunkley2006, "Markers"))
```

```
stopifnot(all.equal(featureNames(markerMSnSet(dunkley2006, "Markers")),
                    featureNames(markerMSnSet(dunkley2006, "markers"))))
dim(unknownMSnSet(dunkley2006, "Markers"))
stopifnot(all.equal(featureNames(unknownMSnSet(dunkley2006, "Markers")),
                    featureNames(unknownMSnSet(dunkley2006, "markers"))))
```

MartInstance-class *Class* "MartInstance"

Description

Internal infrastructure to query/handle several individual mart instance. See `MartInterface.R` for details.

Author(s)

Laurent Gatto <lg390@cam.ac.uk>

---

mcmc_get_outliers *Number of outlier at each iteration of MCMC*

Description

Helper function to get the number of outlier at each MCMC iteration.

Helper function to get mean component allocation at each MCMC iteration.

Helper function to get mean probability of belonging to outlier at each iteration.

Wrapper for the geweke diagnostics from coda package also return p-values.

Helper function to pool chains together after processing

Helper function to burn n iterations from the front of the chains

Helper function to subsample the chains, known informally as thinning.

Produces a violin plot with the protein posterior probabilities distributions for all organelles.

Usage

```
mcmc_get_outliers(x)

mcmc_get_meanComponent(x)

mcmc_get_meanoutliersProb(x)

geweke_test(k)

mcmc_pool_chains(param)
```

```
mcmc_burn_chains(x, n = 50)

mcmc_thin_chains(x, freq = 5)

## S4 method for signature 'MCMCParams,character'
plot(x, y, ...)
```

Arguments

x	Object of class MCMCParams
k	A list of coda::mcmc objects, as returned by mcmc_get_outliers, mcmc_get_meanComponent and mcmc_get_meanoutliersProb.
param	An object of class MCMCParams.
n	integer(1) defining number of iterations to burn. The default is 50
freq	Thinning frequency. The function retains every 'freq'th iteration and is an 'integer(1)'. The default thinning frequency is '5'.
y	A 'character(1)' with a protein name.
...	Currently ignored.

Value

A list of length length(x).

A list of length length(x).

A list of length length(x).

A matrix with the test z- and p-values for each chain.

A pooled MCMCParams object.

An updated MCMCParams object.

A thinned 'MCMCParams' object.

A ggplot2 object.

Author(s)

Laurent Gatto

MCMCChains-class *Instrastructure to store and process MCMC results*

Description

The MCMCParams infrastructure is used to store and process Marchov chain Monte Carlo results for the T-Augmented Gaussian Mixture model (TAGM) from Crook et al. (2018).

Usage

```
chains(object)

## S4 method for signature 'MCMCParams'
show(object)

## S4 method for signature 'ComponentParam'
show(object)

## S4 method for signature 'MCMCChain'
show(object)

## S4 method for signature 'MCMCChains'
length(x)

## S4 method for signature 'MCMCParams'
length(x)

## S4 method for signature 'MCMCChains,ANY,ANY'
x[[i, j = "missing", drop = "missing"]]

## S4 method for signature 'MCMCParams,ANY,ANY'
x[[i, j = "missing", drop = "missing"]]

## S4 method for signature 'MCMCChains,ANY,ANY,ANY'
x[i, j = "missing", drop = "missing"]

## S4 method for signature 'MCMCParams,ANY,ANY,ANY'
x[i, j = "missing", drop = "missing"]

## S4 method for signature 'MCMCChains'
show(object)
```

Arguments

object	An instance of appropriate class.
x	Object to be subset.
i	An integer(). Should be of length 1 for [[.
j	Missing.
drop	Missing.

Details

Objects of the MCMCParams class are created with the tagmMcmcTrain() function. These objects store the *priors* of the generative TAGM model and the results of the MCMC chains, which themselves are stored as an instance of class MCMCChains and can be accessed with the chains() function. A summary of the MCMC chains (or class MCMCSummary) can be further computed with the tagmMcmcProcess() function.

See the *pRoloc-bayesian* vignette for examples.

Slots

`chains` `list()` containing the individual full MCMC chain results in an `MCMCChains` instance. Each element must be a valid `MCMCChain` instance.

`posteriorEstimates` A `data.frame` documenting the prosterior priors in an `MCMCSummary` instance. It contains N rows and columns `tagm.allocation`, `tagm.probability`, `tagm.outlier`, `tagm.probability.lowerquantile`, `tagm.probability.upperquantile` and `tagm.mean.shannon`.

`diagnostics` A `matrix` of dimensions 1 by 2 containing the `MCMCSummary` diagnostics.

`tagm.joint` A `matrix` of dimensions N by K storing the joint probability in an `MCMCSummary` instance.

`method` `character(1)` describing the method in the `MCMCParams` object.

`chains` Object of class `MCMCChains` containing the full MCMC chain results stored in the `MCMCParams` object.

`priors` `list()`

`summary` Object of class `MCMCSummary` the summarised MCMC results available in the `MCMCParams` instance.

`n` `integer(1)` indicating the number of MCMC interactions. Stored in an `MCMCChain` instance.

`K` `integer(1)` indicating the number of components. Stored in an `MCMCChain` instance.

`N` `integer(1)` indicating the number of proteins. Stored in an `MCMCChain` instance.

`Component` `matrix(N, n)` component allocation results of an `MCMCChain` instance.

`ComponentProb` `matrix(N, n, K)` component allocation probabilities of an `MCMCChain` instance.

`Outlier` `matrix(N, n)` outlier allocation results.

`OutlierProb` `matrix(N, n, 2)` outlier allocation probabilities of an `MCMCChain` instance.

See Also

The function `tagmMcmcTrain()` to construct object of this class.

`minMarkers` *Creates a reduced marker variable*

Description

This function updates an `MSnSet` instances and sets markers class to `unknown` if there are less than `n` instances.

Usage

```
minMarkers(object, n = 10, fcol = "markers")
```

Arguments

object	An instance of class "MSnSet".
n	Minumum of marker instances per class.
fcol	The name of the markers column in the featureData slot. Default is markers.

Value

An instance of class "MSnSet" with a new feature variables, named after the original fcol variable and the n value.

Author(s)

Laurent Gatto

See Also

getPredictions to filter based on classification scores.

Examples

```
library(pRolocdata)
data(dunkley2006)
d2 <- minMarkers(dunkley2006, 20)
getMarkers(dunkley2006)
getMarkers(d2, fcol = "markers20")
```

---

mixing_posterior_check

Model calibration plots

Description

Model calibration model with posterior z-scores and posterior shrinkage

Usage

```
mixing_posterior_check(object, params, priors, fcol = "markers")
```

Arguments

object	A valid object of class MSnset
params	A valid object of class MCMCParams that has been processed and checked for convergence
priors	The prior that were used in the model
fcol	The columns of the feature data which contain the marker data.

Value

Used for side effect of producing plot. Invisibily returns an ggplot object that can be further manipulated

Author(s)

Oliver M. Crook <omc25@cam.ac.uk>

Examples

```
## Not run:
library("pRoloc")
data("tan2009r1")

tanres <- tagmMcmcTrain(object = tan2009r1)
tanres <- tagmMcmcProcess(tanres)
tan2009r1 <- tagmMcmcPredict(object = tan2009r1, params = tanres, probJoint = TRUE)
myparams <- chains(e14Tagm_converged_pooled)[[1]]
myparams2 <- chains(mcmc_pool_chains(tanres))[[1]]
priors <- tanres@priors
pRoloc:::mixing_posterior_check(object = tan2009r1, params = myparams2, priors = priors)

## End(Not run)
```

MLearn-methods *The* MLearn *interface for machine learning*

Description

This method implements MLInterfaces' MLean method for instances of the class "MSnSet".

Methods

signature(formula = "formula", data = "MSnSet", .method = "learnerSchema", trainInd = "numeric")
: The learning problem is stated with the formula and applies the .method schema on the MSnSet data input using the trainInd numeric indices as train data.

signature(formula = "formula", data = "MSnSet", .method = "learnerSchema", trainInd = "xvalSpec")
: In this case, an instance of xvalSpec is used for cross-validation.

signature(formula = "formula", data = "MSnSet", .method = "clusteringSchema", trainInd = "missing")
: Hierarchical (hclustI), k-means (kmeansI) and partitioning around medoids (pamI) clustering algorithms using MLInterface's MLearn interface.

See Also

The MLInterfaces package documentation, in particular MLearn.

move2Ds *Displays a spatial proteomics animation*

Description

Given two MSnSet instances of one MSnSetList with at least two items, this function produces an animation that shows the transition from the first data to the second.

Usage

```
move2Ds(object, pcol, fcol = "markers", n = 25, hl)
```

Arguments

object	An linkS4class{MSnSet} or a MSnSetList. In the latter case, only the two first elements of the list will be used for plotting and the others will be silently ignored.
pcol	If object is an MSnSet, a factor or the name of a phenotype variable (phenoData slot) defining how to split the single MSnSet into two or more data sets. Ignored if object is a MSnSetList.
fcol	Feature meta-data label (fData column name) defining the groups to be differentiated using different colours. Default is markers. Use NULL to suppress any colouring.
n	Number of frames, Default is 25.
hl	An optional instance of class linkS4class{FeaturesOfInterest} to track features of interest.

Value

Used for its side effect of producing a short animation.

Author(s)

Laurent Gatto

See Also

plot2Ds to a single figure with the two datasets.

Examples

```
library("pRolocdata")
data(dunkley2006)

## Create a relevant MSnSetList using the dunkley2006 data
xx <- split(dunkley2006, "replicate")
xx1 <- xx[[1]]
```

```
xx2 <- xx[[2]]
fData(xx1)$markers[374] <- "Golgi"
fData(xx2)$markers[412] <- "unknown"
xx@x[[1]] <- xx1
xx@x[[2]] <- xx2

## The features we want to track
foi <- FeaturesOfInterest(description = "test",
                          fnames = featureNames(xx[[1]])[c(374, 412)])

## (1) visualise each experiment separately
par(mfrow = c(2, 1))
plot2D(xx[[1]], main = "condition A")
highlightOnPlot(xx[[1]], foi)
plot2D(xx[[2]], mirrorY = TRUE, main = "condition B")
highlightOnPlot(xx[[2]], foi, args = list(mirrorY = TRUE))

## (2) plot both data on the same plot
par(mfrow = c(1, 1))
tmp <- plot2Ds(xx)
highlightOnPlot(data1(tmp), foi, lwd = 2)
highlightOnPlot(data2(tmp), foi, pch = 5, lwd = 2)

## (3) create an animation
move2Ds(xx, pcol = "replicate")
move2Ds(xx, pcol = "replicate", hl = foi)
```

`mrkConsProfiles` *Marker consensus profiles*

Description

A function to calculate average marker profiles.

Usage

```
mrkConsProfiles(object, fcol = "markers", method = mean)
```

Arguments

`object`	An instance of class `MSnSet`.
`fcol`	Feature meta-data label (fData column name) defining the groups to be differentiated using different colours. Default is `markers`.
`method`	A `function` to average marker profiles. Default is `mean`.

Value

A `matrix` of dimensions *number of clusters* (exluding unknowns) by *number of fractions*.

Author(s)

Laurent Gatto and Lisa M. Breckels

See Also

The mrkHClust function to produce a hierarchical cluster.

Examples

```
library("pRolocdata")
data(dunkley2006)
mrkConsProfiles(dunkley2006)
mrkConsProfiles(dunkley2006, method = median)
mm <- mrkConsProfiles(dunkley2006)
## Reorder fractions
o <- order(dunkley2006$fraction)
## Plot mean organelle profiles using the
## default pRoloc colour palette.
matplot(t(mm[, o]), type = "l",
        xlab = "Fractions", ylab = "Relative intensity",
        main = "Mean organelle profiles",
        col = getStockcol(), lwd = 2, lty = 1)
## Add a legend
addLegend(markerMSnSet(dunkley2006), where = "topleft")
```

---

mrkHClust *Draw a dendrogram of subcellular clusters*

Description

This functions calculates an average protein profile for each marker class (proteins of unknown localisation are ignored) and then generates a dendrogram representing the relation between marker classes. The colours used for the dendrogram labels are taken from the default colours (see getStockcol) so as to match the colours with other spatial proteomics visualisations such as plot2D.

Usage

```
mrkHClust(
  object,
  fcol = "markers",
  distargs,
  hclustargs,
  method = mean,
  plot = TRUE,
  ...
)
```

Arguments

object	An instance of class MSnSet.
fcol	Feature meta-data label (fData column name) defining the groups to be differentiated using different colours. Default is markers.
distargs	A list of arguments to be passed to the dist function.
hclustargs	A list of arguments to be passed to the hclust function.
method	A function to average marker profiles. Default is mean.
plot	A logical defining whether the dendrogram should be plotted. Default is TRUE.
...	Additional parameters passed when plotting the dendrogram.

Value

Invisibly returns a dendrogram object, containing the hierarchical cluster as computed by hclust.

Author(s)

Laurent Gatto

Examples

```
library("pRolocdata")
data(dunkley2006)
mrkHClust(dunkley2006)
```

---

mrkVecToMat *Create a marker vector or matrix.*

Description

Functions producing a new vector (matrix) marker vector set from an existing matrix (vector) marker set.

Usage

```
mrkVecToMat(object, vfcol = "markers", mfcol = "Markers")

mrkMatToVec(object, mfcol = "Markers", vfcol = "markers")

mrkMatAndVec(object, vfcol = "markers", mfcol = "Markers")

showMrkMat(object, mfcol = "Markers")

isMrkMat(object, fcol = "Markers")

isMrkVec(object, fcol = "markers")

mrkEncoding(object, fcol = "markers")
```

Arguments

object	An MSnSet object
vfcol	The name of the *vector* marker feature variable. Default is "markers".
mfcol	The name of the *matrix* marker feature variable. Default is "Markers".
fcol	A marker feature variable name.

Details

Sub-cellular markers can be encoded in two different ways. Sets of spatial markers can be represented as character *vectors* (character or factor, to be accurate), stored as feature metadata, and proteins of unknown or uncertain localisation (unlabelled, to be classified) are marked with the "unknown" character. While very handy, this encoding suffers from some drawbacks, in particular the difficulty to label proteins that reside in multiple (possible or actual) localisations. The markers vector feature data is typically named markers. A new *matrix* encoding is also supported. Each spatial compartment is defined in a column in a binary markers matrix and the resident proteins are encoded with 1s. The markers matrix feature data is typically named Markers. If proteins are assigned unique localisations only (i.e. no multi-localisation) or their localisation is unknown (unlabelled), then both encodings are equivalent. When the markers are encoded as vectors, features of unknown localisation are defined as fData(object)[, fcol] == "unknown". For matrix-encoded markers, unlabelled proteins are defined as rowSums(fData(object)[, fcol]) == 0.

The mrkMatToVec and mrkVecToMat functions enable the conversion from matrix (vector) to vector (matrix). The mrkMatAndVec function generates the missing encoding from the existing one. If the destination encoding already exists, or, more accurately, if the feature variable of the destination encoding exists, an error is thrown. During the conversion from matrix to vector, if multiple possible label exists, they are dropped, i.e. they are converted to "unknown". Function isMrkVec and isMrkMat can be used to test if a marker set is encoded as a vector or a matrix. mrkEncoding returns either "vector" or "matrix" depending on the nature of the markers.

Value

An updated MSnSet with a new vector (matrix) marker set.

Author(s)

Laurent Gatto and Lisa Breckels

See Also

Other functions that operate on markers are getMarkers, getMarkerClasses and markerMSnSet. To add markers to an existing MSnSet, see the addMarkers function and pRolocmarkers, for a list of suggested markers.

Examples

```
library("pRolocdata")
data(dunkley2006)
dunk <- mrkVecToMat(dunkley2006)
head(fData(dunk)$Markers)
```

```
fData(dunk)$markers <- NULL
dunk <- mrkMatToVec(dunk)
stopifnot(all.equal(fData(dunkley2006)$markers,
                    fData(dunk)$markers))
```

`nbClassification` *nb classification*

Description

Classification using the naive Bayes algorithm.

Usage

```
nbClassification(
  object,
  assessRes,
  scores = c("prediction", "all", "none"),
  laplace,
  fcol = "markers",
  ...
)
```

Arguments

`object`	An instance of class `"MSnSet"`.
`assessRes`	An instance of class `"GenRegRes"`, as generated by `nbOptimisation`.
`scores`	One of `"prediction"`, `"all"` or `"none"` to report the score for the predicted class only, for all classes or none.
`laplace`	If `assessRes` is missing, a `laplace` must be provided.
`fcol`	The feature meta-data containing marker definitions. Default is `markers`.
`...`	Additional parameters passed to `naiveBayes` from package `e1071`.

Value

An instance of class `"MSnSet"` with `nb` and `nb.scores` feature variables storing the classification results and scores respectively.

Author(s)

Laurent Gatto

Examples

```
library(pRolocdata)
data(dunkley2006)
## reducing parameter search space and iterations
params <- nbOptimisation(dunkley2006, laplace = c(0, 5),  times = 3)
params
plot(params)
f1Count(params)
levelPlot(params)
getParams(params)
res <- nbClassification(dunkley2006, params)
getPredictions(res, fcol = "naiveBayes")
getPredictions(res, fcol = "naiveBayes", t = 1)
plot2D(res, fcol = "naiveBayes")
```

`nbOptimisation` *nb paramter optimisation*

Description

Classification algorithm parameter for the naive Bayes algorithm.

Usage

```
nbOptimisation(
  object,
  fcol = "markers",
  laplace = seq(0, 5, 0.5),
  times = 100,
  test.size = 0.2,
  xval = 5,
  fun = mean,
  seed,
  verbose = TRUE,
  ...
)
```

Arguments

`object`	An instance of class "MSnSet".
`fcol`	The feature meta-data containing marker definitions. Default is `markers`.
`laplace`	The hyper-parameter. Default values are `seq(0, 5, 0.5)`.
`times`	The number of times internal cross-validation is performed. Default is 100.
`test.size`	The size of test data. Default is 0.2 (20 percent).
`xval`	The n-cross validation. Default is 5.
`fun`	The function used to summarise the `xval` macro F1 matrices.

seed	The optional random number generator seed.
verbose	A logical defining whether a progress bar is displayed.
...	Additional parameters passed to naiveBayes from package e1071.

Details

Note that when performance scores precision, recall and (macro) F1 are calculated, any NA values are replaced by 0. This decision is motivated by the fact that any class that would have either a NA precision or recall would result in an NA F1 score and, eventually, a NA macro F1 (i.e. mean(F1)). Replacing NAs by 0s leads to F1 values of 0 and a reduced yet defined final macro F1 score.

Value

An instance of class "GenRegRes".

Author(s)

Laurent Gatto

See Also

nbClassification and example therein.

Examples

```
library(pRolocdata)
data(dunkley2006)
## reducing parameter search space and iterations
params <- nbOptimisation(dunkley2006, laplace = c(0, 5),  times = 3)
params
plot(params)
f1Count(params)
levelPlot(params)
getParams(params)
res <- nbClassification(dunkley2006, params)
getPredictions(res, fcol = "naiveBayes")
getPredictions(res, fcol = "naiveBayes", t = 1)
plot2D(res, fcol = "naiveBayes")
```

---

nicheMeans2D *Uncertainty plot organelle means*

Description

Produces a pca plot with uncertainty in organelle means projected onto the PCA plot with contours.

Usage

```
nicheMeans2D(
  object,
  params,
  priors,
  dims = c(1, 2),
  fcol = "markers",
  aspect = 0.5
)
```

Arguments

object	A valid object of class MSnset
params	A valid object of class MCMCParams that has been processed and checked for convergence
priors	The prior that were used in the model
dims	The PCA dimension in which to project he data, default is c(1,2)
fcol	The columns of the feature data which contain the marker data.
aspect	A argument to change the plotting aspect of the PCA

Value

Used for side effect of producing plot. Invisibily returns an ggplot object that can be further manipulated

Author(s)

Oliver M. Crook <omc25@cam.ac.uk>

Examples

```
## Not run:
library("pRolocdata")
data("tan2009r1")

tanres <- tagmMcmcTrain(object = tan2009r1)
tanres <- tagmMcmcProcess(tanres)
tan2009r1 <- tagmMcmcPredict(object = tan2009r1, params = tanres, probJoint = TRUE)
myparams <- chains(e14Tagm_converged_pooled)[[1]]
myparams2 <- chains(mcmc_pool_chains(tanres))[[1]]
priors <- tanres@priors
pRoloc:::nicheMeans2D(object = tan2009r1, params = myparams2, priors = priors)

## End(Not run)
```

nndist-methods *Nearest neighbour distances*

Description

Methods computing the nearest neighbour indices and distances for `matrix` and `MSnSet` instances.

Methods

`signature(object = "matrix", k = "numeric", dist = "character", ...)` Calculates indices and distances to the `k` (default is 3) nearest neighbours of each feature (row) in the input matrix `object`. The distance `dist` can be either of `"euclidean"` or `"mahalanobis"`. Additional parameters can be passed to the internal function `FNN::get.knn`. Output is a matrix with `2 * k` columns and `nrow(object)` rows.

`signature(object = "MSnSet", k = "numeric", dist = "character", ...)` As above, but for an `MSnSet` input. The indices and distances to the `k` nearest neighbours are added to the object's feature metadata.

`signature(object = "matrix", query = "matrix", k = "numeric", ...)` If two `matrix` instances are provided as input, the `k` (default is 3) indices and distances of the nearest neighbours of `query` in `object` are returned as a matrix of dimensions `2 * k` by `nrow(query)`. Additional parameters are passed to `FNN::get.knnx`. Only euclidean distance is available.

Examples

```
library("pRolocdata")
data(dunkley2006)

## Using a matrix as input
m <- exprs(dunkley2006)
m[1:4, 1:3]
head(nndist(m, k = 5))
tail(nndist(m[1:100, ], k = 2, dist = "mahalanobis"))

## Same as above for MSnSet
d <- nndist(dunkley2006, k = 5)
head(fData(d))

d <- nndist(dunkley2006[1:100, ], k = 2, dist = "mahalanobis")
tail(fData(d))

## Using a query
nndist(m[1:100, ], m[101:110, ], k = 2)
```

nnetClassification *nnet classification*

Description

Classification using the artificial neural network algorithm.

Usage

```
nnetClassification(
  object,
  assessRes,
  scores = c("prediction", "all", "none"),
  decay,
  size,
  fcol = "markers",
  ...
)
```

Arguments

object	An instance of class "MSnSet".
assessRes	An instance of class "GenRegRes", as generated by nnetOptimisation.
scores	One of "prediction", "all" or "none" to report the score for the predicted class only, for all classes or none.
decay	If assessRes is missing, a decay must be provided.
size	If assessRes is missing, a size must be provided.
fcol	The feature meta-data containing marker definitions. Default is markers.
...	Additional parameters passed to nnet from package nnet.

Value

An instance of class "MSnSet" with nnet and nnet.scores feature variables storing the classification results and scores respectively.

Author(s)

Laurent Gatto

Examples

```
library(pRolocdata)
data(dunkley2006)
## reducing parameter search space and iterations
params <- nnetOptimisation(dunkley2006, decay = 10^(c(-1, -5)), size = c(5, 10), times = 3)
params
```

```
plot(params)
f1Count(params)
levelPlot(params)
getParams(params)
res <- nnetClassification(dunkley2006, params)
getPredictions(res, fcol = "nnet")
getPredictions(res, fcol = "nnet", t = 0.75)
plot2D(res, fcol = "nnet")
```

`nnetOptimisation` *nnet parameter optimisation*

Description

Classification parameter optimisation for artificial neural network algorithm.

Usage

```
nnetOptimisation(
  object,
  fcol = "markers",
  decay = c(0, 10^(-1:-5)),
  size = seq(1, 10, 2),
  times = 100,
  test.size = 0.2,
  xval = 5,
  fun = mean,
  seed,
  verbose = TRUE,
  ...
)
```

Arguments

`object`	An instance of class `"MSnSet"`.
`fcol`	The feature meta-data containing marker definitions. Default is `markers`.
`decay`	The hyper-parameter. Default values are `c(0, 10^(-1:-5))`.
`size`	The hyper-parameter. Default values are `seq(1, 10, 2)`.
`times`	The number of times internal cross-validation is performed. Default is 100.
`test.size`	The size of test data. Default is 0.2 (20 percent).
`xval`	The n-cross validation. Default is 5.
`fun`	The function used to summarise the `xval` macro F1 matrices.
`seed`	The optional random number generator seed.
`verbose`	A `logical` defining whether a progress bar is displayed.
`...`	Additional parameters passed to `nnet` from package `nnet`.

Details

Note that when performance scores precision, recall and (macro) F1 are calculated, any NA values are replaced by 0. This decision is motivated by the fact that any class that would have either a NA precision or recall would result in an NA F1 score and, eventually, a NA macro F1 (i.e. mean(F1)). Replacing NAs by 0s leads to F1 values of 0 and a reduced yet defined final macro F1 score.

Value

An instance of class "GenRegRes".

Author(s)

Laurent Gatto

See Also

nnetClassification and example therein.

Examples

```
library(pRolocdata)
data(dunkley2006)
## reducing parameter search space and iterations
params <- nnetOptimisation(dunkley2006, decay = 10^(c(-1, -5)), size = c(5, 10), times = 3)
params
plot(params)
f1Count(params)
levelPlot(params)
getParams(params)
res <- nnetClassification(dunkley2006, params)
getPredictions(res, fcol = "nnet")
getPredictions(res, fcol = "nnet", t = 0.75)
plot2D(res, fcol = "nnet")
```

---

orderGoAnnotations *Orders annotation information*

Description

For a given matrix of annotation information, this function returns the information ordered according to the best fit with the data.

Usage

```
orderGoAnnotations(
  object,
  fcol = "GOAnnotations",
  k = 1:5,
```

```
  n = 5,
  p = 1/3,
  verbose = TRUE,
  seed
)
```

Arguments

object	An instance of class MSnSet.
fcol	The name of the annotations matrix. Default is GOAnnotations.
k	The number of clusters to test. Default is k = 1:5
n	The minimum number of proteins per component cluster.
p	The normalisation factor, per k tested
verbose	A logical indicating if a progress bar should be displayed. Default is TRUE.
seed	An optional random number generation seed.

Details

As there are typically many protein/annotation sets that may fit the data we order protein sets by best fit i.e. cluster tightness, by computing the mean normalised Euclidean distance for all instances per protein set.

For each protein set i.e. proteins that have been labelled with a specified term/information criteria, we find the best k cluster components for the set (the default is to testk = 1:5) according to the minimum mean normalised pairwise Euclidean distance over all component clusters. (Note: when testing k if any components are found to have less than n proteins these components are not included and k is reduced by 1).

Each component cluster is normalised by N^p (where N is the total number of proteins per component, and p is the power). Hueristally, p = 1/3 and normalising by N^1/3 has been found the optimum normalisation factor.

Candidates in the matrix are ordered according to lowest mean normalised pairwise Euclidean distance as we expect high density, tight clusters to have the smallest mean normalised distance.

This function is a wrapper for running clustDist, and getNormDist.

Value

An updated MSnSet containing the newly ordered fcol matrix.

Author(s)

Lisa M Breckels

orgQuants *Returns organelle-specific quantile scores*

Description

This function produces organelle-specific quantiles corresponding to the given classification scores.

Usage

```
orgQuants(object, fcol, scol, mcol = "markers", t, verbose = TRUE)
```

Arguments

object	An instance of class "MSnSet".
fcol	The name of the prediction column in the featureData slot.
scol	The name of the prediction score column in the featureData slot. If missing, created by pasting '.scores' after fcol.
mcol	The name of the column containing the training data in the featureData slot. Default is markers.
t	The quantile threshold.
verbose	If TRUE, the calculated threholds are printed.

Value

A named vector of organelle thresholds.

Author(s)

Lisa Breckels

See Also

getPredictions to get organelle predictions based on calculated thresholds.

Examples

```
library("pRolocdata")
data(dunkley2006)
res <- svmClassification(dunkley2006, fcol = "pd.markers",
                         sigma = 0.1, cost = 0.5)
## 50% top predictions per class
ts <- orgQuants(res, fcol = "svm", t = .5)
getPredictions(res, fcol = "svm", t = ts)
```

perTurboClassification *perTurbo classification*

Description

Classification using the PerTurbo algorithm.

Usage

```
perTurboClassification(
  object,
  assessRes,
  scores = c("prediction", "all", "none"),
  pRegul,
  sigma,
  inv,
  reg,
  fcol = "markers"
)
```

Arguments

object	An instance of class "MSnSet".
assessRes	An instance of class "GenRegRes", as generated by svmRegularisation.
scores	One of "prediction", "all" or "none" to report the score for the predicted class only, for all classes or none.
pRegul	If assessRes is missing, a pRegul must be provided. See perTurboOptimisation for details.
sigma	If assessRes is missing, a sigma must be provided. See perTurboOptimisation for details.
inv	The type of algorithm used to invert the matrix. Values are : "Inversion Cholesky" (chol2inv), "Moore Penrose" (ginv), "solve" (solve), "svd" (svd). Default value is "Inversion Cholesky".
reg	The type of regularisation of matrix. Values are "none", "trunc" or "tikhonov". Default value is "tikhonov".
fcol	The feature meta-data containing marker definitions. Default is markers.

Value

An instance of class "MSnSet" with perTurbo and perTurbo.scores feature variables storing the classification results and scores respectively.

Author(s)

Thomas Burger and Samuel Wieczorek

References

N. Courty, T. Burger, J. Laurent. "PerTurbo: a new classification algorithm based on the spectrum perturbations of the Laplace-Beltrami operator", The European Conference on Machine Learning and Principles and Practice of Knowledge Discovery in Databases (ECML-PKDD 2011), D. Gunopulos et al. (Eds.): ECML PKDD 2011, Part I, LNAI 6911, pp. 359 - 374, Athens, Greece, September 2011.

Examples

```
library(pRolocdata)
data(dunkley2006)
## reducing parameter search space
params <- perTurboOptimisation(dunkley2006,
                               pRegul = 2^seq(-2,2,2),
                               sigma = 10^seq(-1, 1, 1),
                               inv = "Inversion Cholesky",
                               reg ="tikhonov",
                               times = 3)
params
plot(params)
f1Count(params)
levelPlot(params)
getParams(params)
res <- perTurboClassification(dunkley2006, params)
getPredictions(res, fcol = "perTurbo")
getPredictions(res, fcol = "perTurbo", t = 0.75)
plot2D(res, fcol = "perTurbo")
```

---

perTurboOptimisation *PerTurbo parameter optimisation*

Description

Classification parameter optimisation for the PerTurbo algorithm

Usage

```
perTurboOptimisation(
  object,
  fcol = "markers",
  pRegul = 10^(seq(from = -1, to = 0, by = 0.2)),
  sigma = 10^(seq(from = -1, to = 1, by = 0.5)),
  inv = c("Inversion Cholesky", "Moore Penrose", "solve", "svd"),
  reg = c("tikhonov", "none", "trunc"),
  times = 1,
  test.size = 0.2,
  xval = 5,
  fun = mean,
```

```
  seed,
  verbose = TRUE
)
```

Arguments

object	An instance of class "MSnSet".
fcol	The feature meta-data containing marker definitions. Default is markers.
pRegul	The hyper-parameter for the regularisation (values are in]0,1]). If reg =="trunc", pRegul is for the percentage of eigen values in matrix. If reg =="tikhonov", then 'pRegul' is the parameter for the tikhonov regularisation. Available configurations are : "Inversion Cholesky" - ("tikhonov" / "none"), "Moore Penrose" - ("tikhonov" / "none"), "solve" - ("tikhonov" / "none"), "svd" - ("tikhonov" / "none" / "trunc").
sigma	The hyper-parameter.
inv	The type of algorithm used to invert the matrix. Values are : "Inversion Cholesky" (chol2inv), "Moore Penrose" (ginv), "solve" (solve), "svd" (svd). Default value is "Inversion Cholesky".
reg	The type of regularisation of matrix. Values are "none", "trunc" or "tikhonov". Default value is "tikhonov".
times	The number of times internal cross-validation is performed. Default is 100.
test.size	The size of test data. Default is 0.2 (20 percent).
xval	The n-cross validation. Default is 5.
fun	The function used to summarise the times macro F1 matrices.
seed	The optional random number generator seed.
verbose	A logical defining whether a progress bar is displayed.

Details

Note that when performance scores precision, recall and (macro) F1 are calculated, any NA values are replaced by 0. This decision is motivated by the fact that any class that would have either a NA precision or recall would result in an NA F1 score and, eventually, a NA macro F1 (i.e. mean(F1)). Replacing NAs by 0s leads to F1 values of 0 and a reduced yet defined final macro F1 score.

Value

An instance of class "GenRegRes".

Author(s)

Thomas Burger and Samuel Wieczorek

See Also

perTurboClassification and example therein.

Examples

```
library(pRolocdata)
data(dunkley2006)
## reducing parameter search space
params <- perTurboOptimisation(dunkley2006,
                               pRegul = 2^seq(-2,2,2),
                               sigma = 10^seq(-1, 1, 1),
                               inv = "Inversion Cholesky",
                               reg ="tikhonov",
                               times = 3)
params
plot(params)
f1Count(params)
levelPlot(params)
getParams(params)
res <- perTurboClassification(dunkley2006, params)
getPredictions(res, fcol = "perTurbo")
getPredictions(res, fcol = "perTurbo", t = 0.75)
plot2D(res, fcol = "perTurbo")
```

---

phenoDisco *Runs the* phenoDisco *algorithm.*

Description

phenoDisco is a semi-supervised iterative approach to detect new protein clusters.

Usage

```
phenoDisco(
  object,
  fcol = "markers",
  times = 100,
  GS = 10,
  allIter = FALSE,
  p = 0.05,
  ndims = 2,
  modelNames = mclust.options("emModelNames"),
  G = 1:9,
  BPPARAM,
  tmpfile,
  seed,
  verbose = TRUE,
  dimred = c("PCA", "t-SNE"),
  ...
)
```

Arguments

object	An instance of class MSnSet.
fcol	A character indicating the organellar markers column name in feature metadata. Default is markers.
times	Number of runs of tracking. Default is 100.
GS	Group size, i.e how many proteins make a group. Default is 10 (the minimum group size is 4).
allIter	logical, defining if predictions for all iterations should be saved. Default is FALSE.
p	Significance level for outlier detection. Default is 0.05.
ndims	Number of principal components to use as input for the disocvery analysis. Default is 2. Added in version 1.3.9.
modelNames	A vector of characters indicating the models to be fitted in the EM phase of clustering using Mclust. The help file for mclust::mclustModelNames describes the available models. Default model names are c("EII", "VII", "EEI", "VEI", "EVI", "VVI", "EEE","EEV", "VEV", "VVV"), as returned by mclust.options("emModelNames"). Note that using all these possible models substantially increases the running time. Legacy models are c("EEE","EEV","VEV","VVV"), i.e. only ellipsoidal models.
G	An integer vector specifying the numbers of mixture components (clusters) for which the BIC is to be calculated. The default is G=1:9 (as in Mclust).
BPPARAM	Support for parallel processing using the BiocParallel infrastructure. When missing (default), the default registered BiocParallelParam parameters are used. Alternatively, one can pass a valid BiocParallelParam parameter instance: SnowParam, MulticoreParam, DoparParam, ... see the BiocParallel package for details. To revert to the origianl serial implementation, use NULL.
tmpfile	An optional character to save a temporary MSnSet after each iteration. Ignored if missing. This is useful for long runs to track phenotypes and possibly kill the run when convergence is observed. If the run completes, the temporary file is deleted before returning the final result.
seed	An optional numeric of length 1 specifing the random number generator seed to be used. Only relevant when executed in serialised mode with BPPARAM = NULL. See BPPARAM for details.
verbose	Logical, indicating if messages are to be printed out during execution of the algorithm.
dimred	A characater defining which of Principal Component Analysis ("PCA") or t-Distributed Stochastic Neighbour Embedding ("t-SNE") should be use to reduce dimensions prior to running phenoDisco novelty detection.
...	Additional arguments passed to the dimensionality reduction method. For both PCA and t-SNE, the data is scaled and centred by default, and these parameters (scale and centre for PCA, and pca_scale and pca_center for t-SNE can't be set). When using t-SNE however, it is important to tune the perplexity and max iterations parameters. See the *Dimensionality reduction* section in the pRoloc vignette for details.

Details

The algorithm performs a phenotype discovery analysis as described in Breckels et al. Using this approach one can identify putative subcellular groupings in organelle proteomics experiments for more comprehensive validation in an unbiased fashion. The method is based on the work of Yin et al. and used iterated rounds of Gaussian Mixture Modelling using the Expectation Maximisation algorithm combined with a non-parametric outlier detection test to identify new phenotype clusters.

One requires 2 or more classes to be labelled in the data and at a very minimum of 6 markers per class to run the algorithm. The function will check and remove features with missing values using the `filterNA` method.

A parallel implementation, relying on the `BiocParallel` package, has been added in version 1.3.9. See the `BPPARAM` arguent for details.

Important: Prior to version 1.1.2 the row order in the output was different from the row order in the input. This has now been fixed and row ordering is now the same in both input and output objects.

Value

An instance of class `MSnSet` containing the `phenoDisco` predictions.

Author(s)

Lisa M. Breckels <lms79@cam.ac.uk>

References

Yin Z, Zhou X, Bakal C, Li F, Sun Y, Perrimon N, Wong ST. Using iterative cluster merging with improved gap statistics to perform online phenotype discovery in the context of high-throughput RNAi screens. BMC Bioinformatics. 2008 Jun 5;9:264. PubMed PMID: 18534020.

Breckels LM, Gatto L, Christoforou A, Groen AJ, Lilley KS and Trotter MWB. The Effect of Organelle Discovery upon Sub-Cellular Protein Localisation. J Proteomics. 2013 Aug 2;88:129-40. doi: 10.1016/j.jprot.2013.02.019. Epub 2013 Mar 21. PubMed PMID: 23523639.

Examples

```
## Not run:
library(pRolocdata)
data(tan2009r1)
pdres <- phenoDisco(tan2009r1, fcol = "PLSDA")
getPredictions(pdres, fcol = "pd", scol = NULL)
plot2D(pdres, fcol = "pd")

## to pre-process the data with t-SNE instead of PCA
pdres <- phenoDisco(tan2009r1, fcol = "PLSDA", dimred = "t-SNE")

## End(Not run)
```

plot2D *Plot organelle assignment data and results.*

Description

Generate 2 or 3 dimensional feature distribution plots to illustrate localistation clusters. Rows/features containing NA values are removed prior to dimension reduction except for the "nipals" method. For this method, it is advised to set the method argument 'ncomp' to a low number of dimensions to avoid computing all components when analysing large datasets.

Usage

```
plot2D(
  object,
  fcol = "markers",
  fpch,
  unknown = "unknown",
  dims = 1:2,
  score = 1,
  method = "PCA",
  methargs,
  axsSwitch = FALSE,
  mirrorX = FALSE,
  mirrorY = FALSE,
  col,
  bg,
  palette = "light",
  t = 0.3,
  pch,
  cex,
  lwd,
  index = FALSE,
  idx.cex = 0.75,
  addLegend,
  identify = FALSE,
  plot = TRUE,
  grid = FALSE,
  ...
)

## S4 method for signature 'MSnSet'
plot3D(
  object,
  fcol = "markers",
  dims = c(1, 2, 3),
  radius1 = 0.1,
  radius2 = radius1 * 2,
```

```
  plot = TRUE,
  ...
)
```

Arguments

object	An instance of class MSnSet.
fcol	Feature meta-data label (fData column name) defining the groups to be differentiated using different colours. Default is markers. Use NULL to suppress any colouring.
fpch	Featre meta-data label (fData column name) desining the groups to be differentiated using different point symbols.
unknown	A character (default is "unknown") defining how proteins of unknown/unlabelled localisation are labelled.
dims	A numeric of length 2 (or 3 for plot3D) defining the dimensions to be plotted. Defaults are c(1,2) and c(1, 2, 3).
score	A numeric specifying the minimum organelle assignment score to consider features to be assigned an organelle. (not yet implemented).
method	A character describe how to transform the data or what to plot. One of "PCA" (default), "MDS", "kpca", "nipals", "t-SNE", "UMAP", or "lda", defining what dimensionality reduction is applied: principal component analysis (see prcomp), classical multidimensional scaling (see cmdscale), kernel PCA (see kpca), nipals (principal component analysis by NIPALS, non-linear iterative partial least squares which support missing values; see nipals) t-SNE (see Rtsne), UMAP (see umap) or linear discriminant analysis (see lda). The last method uses fcol to defined the sub-cellular clusters so that the ration between within ad between cluster variance is maximised. All the other methods are unsupervised and make use fcol only to annotate the plot. Prior to t-SNE, duplicated features are removed and a message informs the user if such filtering is needed. "scree" can also be used to produce a scree plot. "hexbin" applies PCA to the data and uses bivariate binning into hexagonal cells from hexbin to emphasise cluster density. If the character "none" is used, the data is plotted as is, i.e. without any transformation. In this case, object can either be an MSnSet or a matrix (as invisibly returned by plot2D). This enables to re-generate the figure without computing the dimensionality reduction over and over again, which can be time consuming for certain methods. If object is a matrix, an MSnSet containing the feature metadata must be provided in methargs (see below for details). Available methods are listed in plot2Dmethods.
methargs	A list of arguments to be passed when method is called. If missing, the data will be scaled and centred prior to PCA and t-SNE (i.e. Rtsne's arguments pca_center and pca_scale are set to TRUE). If method = "none" and object is a matrix, then the first and only argument of methargs must be an MSnSet with matching features with object.
axsSwitch	A logical indicating whether the axes should be switched.
mirrorX	A logical indicating whether the x axis should be mirrored.

mirrorY	A logical indicating whether the y axis should be mirrored.
col	A character of appropriate length defining colours.
bg	Optional background (fill) color for the open plot symbols i.e. can to be used when pch = 21:25.
palette	A character defining which palette colour theme to use, can either defined as "light" (defualt) or "dark".
t	A numeric between 0 and 1. Defining the degree of lightening of the colours in the palette. Default is 0.3.
pch	A character of appropriate length defining point character. Default is 21 (filled circles). See pch for details.
cex	Character expansion: a numerical vector. This works as a multiple of par("cex").
lwd	A numeric defining the line width for drawing symbols. Default is 1.5.
index	A logical (default is FALSE, indicating of the feature indices should be plotted on top of the symbols.
idx.cex	A numeric specifying the character expansion (default is 0.75) for the feature indices. Only relevant when index is TRUE.
addLegend	A character indicating where to add the legend. See addLegend for details. If missing (default), no legend is added.
identify	A logical (default is TRUE) defining if user interaction will be expected to identify individual data points on the plot. See also identify.
plot	A logical defining if the figure should be plotted. Useful when retrieving data only. Default is TRUE.
grid	A logical indicating whether a grid should be plotted. Default is TRUE.
...	Additional parameters passed to plot and points.
radius1	A numeric specifying the radius of feature of unknown localisation. Default is 0.1, which is specidied on the data scale. See plot3d for details.
radius2	A numeric specifying the radius of marker feature. Default is radius * 2.

Details

plot3D relies on the ##' rgl package, that will be loaded automatically.

- Note that plot2D has been update in version 1.3.6 to support more organelle classes than colours defined in getStockcol. In such cases, the default colours are recycled using the default plotting characters defined in getStockpch. See the example for an illustration. The alpha argument is also depreciated in version 1.3.6. Use setStockcol to set colours with transparency instead. See example below.
- Version 1.11.3: to plot data as is, i.e. without any transformation, method can be set to "none" (as opposed to passing pre-computed values to method as a matrix, in previous versions). If object is an MSnSet, the untransformed values in the assay data will be plotted. If object is a matrix with coordinates, then a matching MSnSet must be passed to methargs.

Value

Used for its side effects of generating a plot. Invisibly returns the 2 or 3 dimensions that are plotted.

Author(s)

Laurent Gatto, Lisa Breckels

See Also

addLegend to add a legend to plot2D figures (the legend is added by default on plot3D) and plotDist for alternative graphical representation of quantitative organelle proteomics data. plot2Ds to overlay 2 data sets on the same PCA plot. The plotEllipse function can be used to visualise TAGM models on PCA plots with ellipses.

Examples

```
library("pRolocdata")
data(dunkley2006)
plot2D(dunkley2006, fcol = NULL)
plot2D(dunkley2006, fcol = NULL, col = "black")
plot2D(dunkley2006, fcol = "markers")
addLegend(dunkley2006,
          fcol = "markers",
          where = "topright",
          cex = 0.5, bty = "n", ncol = 3)
title(main = "plot2D example")

## available methods
plot2Dmethods
plot2D(dunkley2006, fcol = NULL, method = "kpca", col = "black")
plot2D(dunkley2006, fcol = NULL, method = "kpca", col = "black",
       methargs = list(kpar = list(sigma = 1)))
plot2D(dunkley2006, method = "lda")
plot2D(dunkley2006, method = "hexbin")

## Using transparent colours
setStockcol(paste0(getStockcol(), "80"))
setStockbg(paste0(getStockbg(), "80"))
plot2D(dunkley2006, fcol = "markers")
## New behavious in 1.3.6 when not enough colours
setStockcol(c("blue", "red", "green"))
getStockcol() ## only 3 colours to be recycled
getMarkers(dunkley2006)
plot2D(dunkley2006)

## Reset colours
setStockcol(NULL)
setStockbg(NULL)
plot2D(dunkley2006, method = "none") ## plotting along 2 first fractions
plot2D(dunkley2006, dims = c(3, 5), method = "none") ## plotting along fractions 3 and 5

## Using different light and dark colour themes
plot2D(dunkley2006, palette = "dark")
plot2D(dunkley2006, palette = "dark", t = .1)
plot2D(dunkley2006, palette = "light")
plot2D(dunkley2006, palette = "light", t = .6)
```

```
## Changing the point characters
plot2D(dunkley2006, pch = 22)
setUnknownpch(22)
plot2D(dunkley2006, pch = 22)
setUnknownpch(NULL) ## reset unknowns pch back to default

## pre-calculate PC1 and PC2 coordinates
pca <- plot2D(dunkley2006, plot=FALSE)
head(pca)
plot2D(pca, method = "none", methargs  = list(dunkley2006))

## Adding a legend inside a plot
plot2D(dunkley2006)
addLegend(dunkley2006,  where = "topleft")

## Adding a legend outside a plot
par(mfrow = c(1, 2))
plot2D(dunkley2006)
addLegend(dunkley2006, where = "other")

## Plotting information from the fData slot
fvarLabels(dunkley2006)
plot2D(dunkley2006, fcol = "assigned")
addLegend(dunkley2006,  where = "topleft", ncol = 2, cex = .5)

## plotting in 3 dimenstions
plot3D(dunkley2006)
plot3D(dunkley2006, radius2 = 0.3)
plot3D(dunkley2006, dims = c(2, 4, 6))
```

`plot2Ds` *Draw 2 data sets on one PCA plot*

Description

Takes 2 `linkS4class{MSnSet}` instances as input to plot the two data sets on the same PCA plot. The second data points are projected on the PC1 and PC2 dimensions calculated for the first data set.

Usage

```
plot2Ds(
  object,
  pcol,
  fcol = "markers",
  cex.x = 1,
  cex.y = 1,
  pch.x = 21,
```

```
  pch.y = 23,
  col,
  mirrorX = FALSE,
  mirrorY = FALSE,
  plot = TRUE,
  ...
)
```

Arguments

object	An MSnSet or a MSnSetList. In the latter case, only the two first elements of the list will be used for plotting and the others will be silently ignored.
pcol	If object is an MSnSet, a factor or the name of a phenotype variable (phenoData slot) defining how to split the single MSnSet into two or more data sets. Ignored if object is a MSnSetList.
fcol	Feature meta-data label (fData column name) defining the groups to be differentiated using different colours. Default is markers. Use NULL to suppress any colouring.
cex.x	Character expansion for the first data set. Default is 1.
cex.y	Character expansion for the second data set. Default is 1.
pch.x	Plotting character for the first data set. Default is 21.
pch.y	Plotting character for the second data set. Default is 23.
col	A vector of colours to highlight the different classes defined by fcol. If missing (default), default colours are used (see getStockcol).
mirrorX	A logical indicating whether the x axis should be mirrored?
mirrorY	A logical indicating whether the y axis should be mirrored?
plot	If TRUE (default), a plot is produced.
...	Additinal parameters passed to plot and points.

Value

Used for its side effects of producing a plot. Invisibly returns an object of class plot2Ds, which is a list with the PCA analyses results (see prcomp) of the first data set and the new coordinates of the second data sets, as used to produce the plot and the respective point colours. Each of these elements can be accessed with data1, data2, col1 and code2 respectively.

Author(s)

Laurent Gatto

See Also

See plot2D to plot a single data set and move2Ds for a animation.

Examples

```
library("pRolocdata")
data(tan2009r1)
data(tan2009r2)
msnl <- MSnSetList(list(tan2009r1, tan2009r2))
plot2Ds(msnl)
## tweaking the parameters
plot2Ds(list(tan2009r1, tan2009r2),
        fcol = NULL, cex.x = 1.5)
## input is 1 MSnSet containing 2 data sets
data(dunkley2006)
plot2Ds(dunkley2006, pcol = "replicate")
## no plot, just the data
res <- plot2Ds(dunkley2006, pcol = "replicate",
               plot = FALSE)
res
head(data1(res))
head(col1(res))
```

`plotConsProfiles` *Plot marker consensus profiles.*

Description

The function plots marker consensus profiles obtained from mrkConsProfile

Usage

```
plotConsProfiles(object, order = NULL, plot = TRUE)
```

Arguments

`object`	A `matrix` containing marker consensus profiles as output from `mrkConsProfiles()`.
`order`	Order for markers (optional).
`plot`	A `logical(1)` defining whether the heatmap should be plotted. Default is `TRUE`.

Value

Invisibly returns `ggplot2` object.

Author(s)

Tom Smith

Examples

```
library("pRolocdata")
data(E14TG2aS1)
hc <- mrkHClust(E14TG2aS1, plot = FALSE)
mm <- getMarkerClasses(E14TG2aS1)
ord <- levels(factor(mm))[order.dendrogram(hc)]
fmat <- mrkConsProfiles(E14TG2aS1)
plotConsProfiles(fmat, order = ord)
```

---

plotDist *Plots the distribution of features across fractions*

Description

Produces a line plot showing the feature abundances across the fractions.

Usage

```
plotDist(
  object,
  markers,
  fcol = NULL,
  mcol = "steelblue",
  pcol = getUnknowncol(),
  alpha = 0.3,
  type = "b",
  lty = 1,
  fractions = sampleNames(object),
  ylab = "Intensity",
  xlab = "Fractions",
  ylim,
  unknown = "unknown",
  ...
)
```

Arguments

object	An instance of class MSnSet.
markers	A character, numeric or logical of appropriate length and or content used to subset object and define the organelle markers.
fcol	Feature meta-data label (fData column name) defining the groups to be differentiated using different colours. If NULL (default) ignored and mcol and pcol are used.
mcol	A character define the colour of the marker features. Default is "steelblue".
pcol	A character define the colour of the non-markers features. Default is the colour used for features of unknown localisation, as returned by getUnknowncol.

alpha	A numeric defining the alpha channel (transparency) of the points, where 0 <= alpha <= 1, 0 and 1 being completely transparent and opaque.
type	Character string defining the type of lines. For example "p" for points, "l" for lines, "b" for both. See plot for all possible types.
lty	Vector of line types for the marker profiles. Default is 1 (solid). See par for details.
fractions	A character defining the phenoData variable to be used to label the fraction along the x axis. Default is to use sampleNames(object).
ylab	y-axis label. Default is "Intensity".
xlab	x-axis label. Default is "Fractions".
ylim	A numeric vector of length 2, giving the y coordinates range.
unknown	Character defining how unlabelled points are defined default is "unknown".
...	Additional parameters passed to plot.

Value

Used for its side effect of producing a feature distribution plot. Invisibly returns the data matrix.

Author(s)

Laurent Gatto

Examples

```
library("pRolocdata")
data(tan2009r1)
j <- which(fData(tan2009r1)$markers == "mitochondrion")
i <- which(fData(tan2009r1)$PLSDA == "mitochondrion")
plotDist(tan2009r1[i, ], markers = featureNames(tan2009r1)[j])
plotDist(tan2009r1[i, ], markers = featureNames(tan2009r1)[j],
         fractions = "Fractions")
## plot and colour all marker profiles
tanmrk <- markerMSnSet(tan2009r1)
plotDist(tanmrk, fcol = "markers")
```

---

plotEllipse *A function to plot probabiltiy ellipses on marker PCA plots to visualise and assess TAGM models.*

Description

Note that when running PCA, this function does not scale the data (centring is performed), as opposed to [plot2D()]. Only marker proteins are displayed; the protein of unknown location, that are not used to estimate the MAP parameters, are filtered out.

Usage

```
plotEllipse(object, params, dims = c(1, 2), method = "MAP", ...)
```

Arguments

object	An [‘MSnbase::MSnset‘] containing quantitative spatial proteomics data.
params	An [‘MAPParams‘] with the TAGM-MAP parameters, as generated by ‘tagmMapTrain‘.
dims	A ‘numeric(2)‘ with the principal components along which to project the data. Default is ‘c(1, 2)‘.
method	The method used. Currently ‘"MAP"‘ only.
...	Additional parameters passed to [plot2D()].

Value

A PCA plot of the marker data with probability ellipises. The outer ellipse contains 99 probability whilst the middle and inner ellipses contain 95 and 90 clusters are represented by black circumpunct (circled dot).

See Also

[plot2D()] to visualise spatial proteomics data using various dimensionality reduction methods. For details about TAGM models, see [tagmPredict()] and the *pRoloc-bayesian* vignette.

---

plsdaClassification *plsda classification*

Description

Classification using the partial least square distcriminant analysis algorithm.

Usage

```
plsdaClassification(
  object,
  assessRes,
  scores = c("prediction", "all", "none"),
  ncomp,
  fcol = "markers",
  ...
)
```

Arguments

object	An instance of class "MSnSet".
assessRes	An instance of class "GenRegRes", as generated by plsdaOptimisation.
scores	One of "prediction", "all" or "none" to report the score for the predicted class only, for all classes or none.
ncomp	If assessRes is missing, a ncomp must be provided.
fcol	The feature meta-data containing marker definitions. Default is markers.
...	Additional parameters passed to plsda from package caret.

Value

An instance of class "MSnSet" with plsda and plsda.scores feature variables storing the classification results and scores respectively.

Author(s)

Laurent Gatto

Examples

```
## not running this one for time considerations
library(pRolocdata)
data(dunkley2006)
## reducing parameter search space and iterations
params <- plsdaOptimisation(dunkley2006, ncomp = c(3, 10),  times = 2)
params
plot(params)
f1Count(params)
levelPlot(params)
getParams(params)
res <- plsdaClassification(dunkley2006, params)
getPredictions(res, fcol = "plsda")
getPredictions(res, fcol = "plsda", t = 0.9)
plot2D(res, fcol = "plsda")
```

---

plsdaOptimisation *plsda parameter optimisation*

Description

Classification parameter optimisation for the partial least square distcriminant analysis algorithm.

Usage

```
plsdaOptimisation(
  object,
  fcol = "markers",
  ncomp = 2:6,
  times = 100,
  test.size = 0.2,
  xval = 5,
  fun = mean,
  seed,
  verbose = TRUE,
  ...
)
```

Arguments

object	An instance of class "MSnSet".
fcol	The feature meta-data containing marker definitions. Default is markers.
ncomp	The hyper-parameter. Default values are 2:6.
times	The number of times internal cross-validation is performed. Default is 100.
test.size	The size of test data. Default is 0.2 (20 percent).
xval	The n-cross validation. Default is 5.
fun	The function used to summarise the xval macro F1 matrices.
seed	The optional random number generator seed.
verbose	A logical defining whether a progress bar is displayed.
...	Additional parameters passed to plsda from package caret.

Details

Note that when performance scores precision, recall and (macro) F1 are calculated, any NA values are replaced by 0. This decision is motivated by the fact that any class that would have either a NA precision or recall would result in an NA F1 score and, eventually, a NA macro F1 (i.e. mean(F1)). Replacing NAs by 0s leads to F1 values of 0 and a reduced yet defined final macro F1 score.

Value

An instance of class "GenRegRes".

Author(s)

Laurent Gatto

not running this one for time considerations library(pRolocdata) data(dunkley2006) ## reducing parameter search space and iterations params <- plsdaOptimisation(dunkley2006, ncomp = c(3, 10), times = 2) params plot(params) f1Count(params) levelPlot(params) getParams(params) res <- plsdaClassification(dunkley2006, params) getPredictions(res, fcol = "plsda") getPredictions(res, fcol = "plsda", t = 0.9) plot2D(res, fcol = "plsda")

See Also

plsdaClassification and example therein.

---

pRolocmarkers *Organelle markers*

Description

This function retrieves a list of organelle markers or, if no species is provided, prints a description of available marker sets. The markers can be added to and MSnSet using the addMarkers function. Several marker version are provided (see Details for additional information).

Usage

```
pRolocmarkers(species, version = "2")
```

Arguments

species	character(1) defining the species of interest. For reference species markers, this is just the species e.g. "hsap". For published marker sets this is the species and author name e.g. "hsap_geladaki".
version	character(1) defining the marker version. Default is "2".

Details

Version 1 of the markers have been contributed by various members of the Cambridge Centre for Proteomics, in particular Dr Dan Nightingale for yeast, Dr Andy Christoforou and Dr Claire Mulvey for human, Dr Arnoud Groen for Arabodopsis and Dr Claire Mulvey for mouse. In addition, original (curated) markers from the pRolocdata datasets have been extracted (see pRolocdata for details and references). Curation involved verification of publicly available subcellular localisation annotation based on the curators knowledge of the organelles/proteins considered and tracing the original statement in the literature.

Version 2 of the markers (current default) have been updated by Charlotte Hutchings from the Cambridge Centre for Proteomics. Reference species marker sets are the same as those in version 1 with minor corrections and an updated naming system. Version 2 also contains additional marker sets from spatial proteomics publications. References for the source publications are provided below:

- Geladaki, A., Britovsek, N.K., Breckels, L.M., Smith, T.S., Vennard, O.L., Mulvey, C.M., Crook, O.M., Gatto, L. and Lilley, K.S. (2019) Combining LOPIT with differential ultracentrifugation for high-resolution spatial proteomics. Nature Communications. 10 (1). doi:10.1038/s41467-018-08191-w
- Christopher, J.A., Breckels, L.M., Crook, O.M., Vazquez–Chantada, M., Barratt, D. and Lilley, K.S. (2024) Global proteomics indicates subcellular-specific anti-ferroptotic responses to ionizing radiation.p.2024.09.12.611851. doi:10.1101/2024.09.12.611851
- Itzhak, D.N., Tyanova, S., Cox, J. and Borner, G.H. (2016) Global, quantitative and dynamic mapping of protein subcellular localization. eLife. 5. doi:10.7554/elife.16950

- Villanueva, E., Smith, T., Pizzinga, M., Elzek, M., Queiroz, R.M.L., Harvey, R.F., Breckels, L.M., Crook, O.M., Monti, M., Dezi, V., Willis, A.E. and Lilley, K.S. (2023) System-wide analysis of RNA and protein subcellular localization dynamics. Nature Methods. 1-12. doi:10.1038/s41592-023-02101-9
- Christoforou, A., Mulvey, C.M., Breckels, L.M., Geladaki, A., Hurrell, T., Hayward, P.C., Naake, T., Gatto, L., Viner, R., Arias, A.M. and Lilley, K.S. (2016) A draft map of the mouse pluripotent stem cell spatial proteome. Nature Communications. 7 (1). doi:10.1038/ncomms9992
- Barylyuk, K., Koreny, L., Ke, H., Butterworth, S., Crook, O.M., Lassadi, I., Gupta, V., Tromer, E., Mourier, T., Stevens, T.J., Breckels, L.M., Pain, A., Lilley, K.S. and Waller, R.F. (2020) A Comprehensive Subcellular Atlas of the Toxoplasma Proteome via hyperLOPIT Provides Spatial Context for Protein Functions. Cell Host and Microbe. 28 (5), 752-766.e9. doi:10.1016/j.chom.2020.09.011
- Moloney, N.M., Barylyuk, K., Tromer, E., Crook, O.M., Breckels, L.M., Lilley, K.S., Waller, R.F. and MacGregor, P. (2023) Mapping diversity in African trypanosomes using high resolution spatial proteomics. Nature Communications. 14 (1), 4401. doi:10.1038/s41467-023-40125-z

Note: These markers are provided as a starting point to generate reliable sets of organelle markers but still need to be verified against any new data in the light of the quantitative data and the study conditions.

Value

Prints a description of the available marker lists if `species` is missing or a named character with organelle markers.

Author(s)

Laurent Gatto

See Also

`addMarkers` to add markers to an `MSnSet` and `markers` for more information about marker encoding.

Examples

```
pRolocmarkers()
pRolocmarkers("hsap")
table(pRolocmarkers("hsap"))

## Old markers
pRolocmarkers("hsap", version = "2")["Q9BPW9"]
pRolocmarkers("hsap", version = "1")["Q9BPW9"]
```

QSep-class *Quantify resolution of a spatial proteomics experiment*

Description

The QSep infrastructure provide a way to quantify the resolution of a spatial proteomics experiment, i.e. to quantify how well annotated sub-cellular clusters are separated from each other.

The QSep function calculates all between and within cluster average distances. These distances are then divided column-wise by the respective within cluster average distance. For example, for a dataset with only 2 spatial clusters, we would obtain

$$\begin{array}{cc} & \begin{array}{cc} c_1 & c_2 \end{array} \\ \begin{array}{c} c_1 \\ c_2 \end{array} & \begin{array}{cc} d_1 1 & d_1 2 \\ d_2 1 & d_2 2 \end{array} \end{array}$$

Normalised distance represent the ratio of between to within average distances, i.e. how much bigger the average distance between cluster c_i and c_j is compared to the average distance within cluster c_i.

$$\begin{array}{cc} & \begin{array}{cc} c_1 & c_2 \end{array} \\ \begin{array}{c} c_1 \\ c_2 \end{array} & \begin{array}{cc} 1 & \frac{d_1 2}{d_2 2} \\ \frac{d_2 1}{d_1 1} & 1 \end{array} \end{array}$$

Note that the normalised distance matrix is not symmetric anymore and the normalised distance ratios are proportional to the tightness of the reference cluster (along the columns).

Missing values only affect the fractions containing the NA when the distance is computed (see the example below) and further used when calculating mean distances. Few missing values are expected to have negligible effect, but data with a high proportion of missing data will will produce skewed distances. In QSep, we take a conservative approach, using the data as provided by the user, and expect that the data missingness is handled before proceeding with this or any other analysis.

Objects from the Class

Objects can be created by calls using the constructor QSep (see below).

Slots

x: Object of class "matrix" containing the pairwise distance matrix, accessible with qseq(., norm = FALSE).

xnorm: Object of class "matrix" containing the normalised pairwise distance matrix, accessible with qsep(.,norm = TRUE) or qsep(.).

object: Object of class "character" with the variable name of MSnSet object that was used to generate the QSep object.

.__classVersion__: Object of class "Versions" storing the class version of the object.

Extends

Class "Versioned", directly.

Methods and functions

QSeq `signature(object = "MSnSet", fcol = "character")`: constructor for QSep objects. The `fcol` argument defines the name of the feature variable that annotates the sub-cellular clusters. Non-marker proteins, that are marked as "unknown" are automatically removed prior to distance calculation.

qsep `signature{object = "QSep", norm = "logical"}`: accessor for the normalised (when `norm` is TRUE, which is default) and raw (when `norm` is FALSE) pairwise distance matrices.

names `signature{object = "QSep"}`: method to retrieve the names of the sub-celluar clusters originally defined in QSep's `fcol` argument. A replacement method `names(.) <-` is also available.

summary `signature(object = "QSep", ..., verbose = "logical")`: Invisible return all between cluster average distances and prints (when `verbose` is TRUE, default) a summary of those.

levelPlot `signature(object = "QSep", norm = "logical",...)`: plots an annotated heatmap of all normalised pairwise distances. `norm` (default is TRUE) defines whether normalised distances should be plotted. Additional arguments `...` are passed to the `levelplot`.

plot `signature(object = "QSep", norm = "logical"...)`: produces a boxplot of all normalised pairwise distances. The red points represent the within average distance and black points between average distances. `norm` (default is TRUE) defines whether normalised distances should be plotted.

Author(s)

Laurent Gatto <lg390@cam.ac.uk>

References

Assessing sub-cellular resolution in spatial proteomics experiments Laurent Gatto, Lisa M Breckels, Kathryn S Lilley bioRxiv 377630; doi: https://doi.org/10.1101/377630

Examples

```
## Test data from Christoforou et al. 2016
library("pRolocdata")
data(hyperLOPIT2015)

## Create the object and get a summary
hlq <- QSep(hyperLOPIT2015)
hlq
summary(hlq)

## mean distance matrix
qsep(hlq, norm = FALSE)

## normalised average distance matrix
```

```
qsep(hlq)

## Update the organelle cluster names for better
## rendering on the plots
names(hlq) <- sub("/", "\n", names(hlq))
names(hlq) <- sub(" - ", "\n", names(hlq))
names(hlq)

## Heatmap of the normalised intensities
levelPlot(hlq)

## Boxplot of the normalised intensities
par(mar = c(3, 10, 2, 1))
plot(hlq)

## Boxplot of all between cluster average distances
x <- summary(hlq, verbose = FALSE)
boxplot(x)

## Missing data example, for 4 proteins and 3 fractions
x <- rbind(c(1.1, 1.2, 1.3), rep(1, 3), c(NA, 1, 1), c(1, 1, NA))
rownames(x) <- paste0("P", 1:4)
colnames(x) <- paste0("F", 1:3)

## P1 is the reference, against which we will calculate distances. P2
## has a complete profile, producing the *real* distance. P3 and P4 have
## missing values in the first and last fraction respectively.
x

## If we drop F1 in P3, which represents a small difference of 0.1, the
## distance only considers F2 and F3, and increases. If we drop F3 in
## P4, which represents a large distance of 0.3, the distance only
## considers F1 and F2, and decreases.  dist(x)
```

`rfClassification` *rf classification*

Description

Classification using the random forest algorithm.

Usage

```
rfClassification(
  object,
  assessRes,
  scores = c("prediction", "all", "none"),
  mtry,
  fcol = "markers",
```

```
  ...
)
```

Arguments

object	An instance of class "MSnSet".
assessRes	An instance of class "GenRegRes", as generated by rfOptimisation.
scores	One of "prediction", "all" or "none" to report the score for the predicted class only, for all classes or none.
mtry	If assessRes is missing, a mtry must be provided.
fcol	The feature meta-data containing marker definitions. Default is markers.
...	Additional parameters passed to randomForest from package randomForest.

Value

An instance of class "MSnSet" with rf and rf.scores feature variables storing the classification results and scores respectively.

Author(s)

Laurent Gatto

Examples

```
library(pRolocdata)
data(dunkley2006)
## reducing parameter search space and iterations
params <- rfOptimisation(dunkley2006, mtry = c(2, 5, 10),  times = 3)
params
plot(params)
f1Count(params)
levelPlot(params)
getParams(params)
res <- rfClassification(dunkley2006, params)
getPredictions(res, fcol = "rf")
getPredictions(res, fcol = "rf", t = 0.75)
plot2D(res, fcol = "rf")
```

---

rfOptimisation *svm parameter optimisation*

Description

Classification parameter optimisation for the random forest algorithm.

Usage

```
rfOptimisation(
  object,
  fcol = "markers",
  mtry = NULL,
  times = 100,
  test.size = 0.2,
  xval = 5,
  fun = mean,
  seed,
  verbose = TRUE,
  ...
)
```

Arguments

object	An instance of class "MSnSet".
fcol	The feature meta-data containing marker definitions. Default is markers.
mtry	The hyper-parameter. Default value is NULL.
times	The number of times internal cross-validation is performed. Default is 100.
test.size	The size of test data. Default is 0.2 (20 percent).
xval	The n-cross validation. Default is 5.
fun	The function used to summarise the xval macro F1 matrices.
seed	The optional random number generator seed.
verbose	A logical defining whether a progress bar is displayed.
...	Additional parameters passed to randomForest from package randomForest.

Details

Note that when performance scores precision, recall and (macro) F1 are calculated, any NA values are replaced by 0. This decision is motivated by the fact that any class that would have either a NA precision or recall would result in an NA F1 score and, eventually, a NA macro F1 (i.e. mean(F1)). Replacing NAs by 0s leads to F1 values of 0 and a reduced yet defined final macro F1 score.

Value

An instance of class "GenRegRes".

Author(s)

Laurent Gatto

See Also

rfClassification and example therein.

Examples

```
library(pRolocdata)
data(dunkley2006)
## reducing parameter search space and iterations
params <- rfOptimisation(dunkley2006, mtry = c(2, 5, 10),  times = 3)
params
plot(params)
f1Count(params)
levelPlot(params)
getParams(params)
res <- rfClassification(dunkley2006, params)
getPredictions(res, fcol = "rf")
getPredictions(res, fcol = "rf", t = 0.75)
plot2D(res, fcol = "rf")
```

---

sampleMSnSet *Extract a stratified sample of an* MSnSet

Description

This function extracts a stratified sample of an MSnSet.

Usage

```
sampleMSnSet(object, fcol = "markers", size = 0.2, seed)
```

Arguments

object	An instance of class MSnSet
fcol	The feature meta-data column name containing the marker (vector or matrix) definitions on which the MSnSet will be stratified. Default is markers.
size	The size of the stratified sample to be extracted. Default is 0.2 (20 percent).
seed	The optional random number generator seed.

Value

A stratified sample (according to the defined fcol) which is an instance of class "MSnSet".

Author(s)

Lisa Breckels

See Also

testMSnSet unknownMSnSet markerMSnSet. See markers for details about markers encoding.

Examples

```
library(pRolocdata)
data(tan2009r1)
dim(tan2009r1)
smp <- sampleMSnSet(tan2009r1, fcol = "markers")
dim(smp)
getMarkers(tan2009r1)
getMarkers(smp)
```

`setLisacol` *Manage default colours and point characters*

Description

These functions allow to get/set the colours and point character that are used when plotting organelle clusters and unknown features. These values are parametrised at the session level. Two palettes are available: the default palette (previously *Lisa's colours*) containing 30 colours and the old (original) palette, containing 13 colours.

Usage

```
setLisacol()

getLisacol()

getOldcol()

setOldcol()

getStockcol()

setStockcol(cols)

getStockpch()

setStockpch(pchs)

getUnknowncol()

setUnknowncol(col)

getUnknownpch()

setUnknownpch(pch)

getStockbg()
```

```
setStockbg(bg)

getUnknownbg()

setUnknownbg(bg)
```

Arguments

cols	A vector of colour characters or NULL, which sets the colours to the default values.
pchs	A vector of numeric or NULL, which sets the point characters to the default values.
col	A colour character or NULL, which sets the colour to #E7E7E7 (grey91), the default colour for unknown features.
pch	A numeric vector of length 1 or NULL, which sets the point character to 21, the default.
bg	A colour character or NULL, which sets the background (fill) colour for open plot symbols given by pch = 21:25 to the default colour for unknown features.

Value

The set functions set (and invisibly returns) colours. The get functions returns a character vector of colours. For the pch functions, numerics rather than characters.

Author(s)

Laurent Gatto

Examples

```
## defaults for clusters
getStockcol()
getStockbg()
getStockpch()
## unknown features
getUnknowncol()
getUnknownbg()
getUnknownpch()
## an example
library(pRolocdata)
data(dunkley2006)
par(mfrow = c(2, 1))
plot2D(dunkley2006, fcol = "markers", main = 'Default colours')
setUnknowncol("black")
setUnknownbg("grey")
plot2D(dunkley2006, fcol = "markers",
      main = 'setUnknowncol("black") and setUnknownbg("grey")')
getUnknowncol()
getUnknownbg()
setUnknowncol(NULL)
```

```
setUnknownbg(NULL)
getUnknowncol()
getStockcol()
getOldcol()
```

---

spatial2D *Uncertainty plot in localisation probabilities*

Description

Produces a pca plot with spatial variation in localisation probabilities

Usage

```
spatial2D(
  object,
  dims = c(1, 2),
  cov.function = fields::wendland.cov,
  theta = 1,
  derivative = 2,
  k = 1,
  breaks = c(0.99, 0.95, 0.9, 0.85, 0.8, 0.75, 0.7),
  aspect = 0.5
)
```

Arguments

object	A valid object of class MSnset with mcmc prediction results from tagmMCMCpredict
dims	The PCA dimension in which to project he data, default is c(1,2)
cov.function	The covariance function used default is wendland.cov. See fields package.
theta	A hyperparameter to the covariance function. See fields package. Default is 1.
derivative	The number of derivative of the wendland kernel. See fields package. Default is 2.
k	A hyperparamter to the covariance function. See fields package. Default is 1.
breaks	Probability values at which to draw the contour bands. Default is c(0.99, 0.95, 0.9, 0.85, 0.8, 0.75, 0.7)
aspect	A argument to change the plotting aspect of the PCA

Value

Used for side effect of producing plot. Invisibily returns an ggplot object that can be further manipulated

Author(s)

Oliver M. Crook <omc25@cam.ac.uk>

Examples

```
## Not run:
library("pRolocdata")
data("tan2009r1")

tanres <- tagmMcmcTrain(object = tan2009r1)
tanres <- tagmMcmcProcess(tanres)
tan2009r1 <- tagmMcmcPredict(object = tan2009r1, params = tanres, probJoint = TRUE)
spatial2D(object = tan2009r1)

## End(Not run)
```

SpatProtVis-class *Class* SpatProtVis

Description

A class for spatial proteomics visualisation, that upon instantiation, pre-computes all defined visualisations. Objects can be created with the `SpatProtVis` constructor and visualised with the `plot` method.

The class is essentially a wrapper around several calls to `plot2D` that stores the dimensionality reduction outputs, and is likely to be updated in the future.

Usage

```
SpatProtVis(x, methods, dims, methargs, ...)
```

Arguments

`x`	An instance of class `MSnSet` to visualise.
`methods`	Dimensionality reduction methods to be used to visualise the data. Must be contained in `plot2Dmethods` (except `"scree"`). See `plot2D` for details.
`dims`	A list of numerics defining dimensions used for plotting. Default are `1` and `2`. If provided, the length of this list must be identical to the length of `methods`.
`methargs`	A list of additional arguments to be passed for each visualisation method. If provided, the length of this list must be identical to the length of `methods`.
`...`	Additional arguments. Currently ignored.

Slots

`vismats`**:** A `"list"` of matrices containing the feature projections in 2 dimensions.

`data`**:** The original spatial proteomics data stored as an `"MSnSet"`.

`methargs`**:** A `"list"` of additional plotting arguments.

`objname`**:** A `"character"` defining how to name the dataset. By default, this is set using the variable name used at object creation.

Methods

`plot`: Generates the figures for the respective `methods` and additional arguments defined in the constructor. If used in an interactive session, the user is prompted to press 'Return' before new figures are displayed.

`show`: A simple textual summary of the object.

Author(s)

Laurent Gatto <lg390@cam.ac.uk>

See Also

The data for the individual visualisations is created by `plot2D`.

Examples

```
library("pRolocdata")
data(dunkley2006)
## Default parameters for a set of methods
## (in the interest of time, don't use t-SNE)
m <- c("PCA", "MDS", "kpca")
vis <- SpatProtVis(dunkley2006, methods = m)
vis
plot(vis)
plot(vis, legend = "topleft")

## Setting method arguments
margs <- c(list(kpar = list(sigma = 0.1)),
           list(kpar = list(sigma = 1.0)),
           list(kpar = list(sigma = 10)),
           list(kpar = list(sigma = 100)))
vis <- SpatProtVis(dunkley2006,
                   methods = rep("kpca", 4),
                   methargs = margs)
par(mfrow = c(2, 2))
plot(vis)

## Multiple PCA plots but different PCs
dims <- list(c(1, 2), c(3, 4))
vis <- SpatProtVis(dunkley2006, methods = c("PCA", "PCA"), dims = dims)
plot(vis)
```

---

subsetMarkers *Subsets markers*

Description

Subsets a matrix of markers by specific terms

Usage

```
subsetMarkers(object, fcol = "GOAnnotations", keep)
```

Arguments

object	An instance of class MSnSet.
fcol	The name of the markers matrix. Default is GOAnnotations.
keep	Integer or character vector specifying the columns to keep in the markers matrix, as defined by fcol.

Value

An updated MSnSet

Author(s)

Lisa M Breckels

See Also

filterMinMarkers and example therein.

---

svmClassification *svm classification*

Description

Classification using the support vector machine algorithm.

Usage

```
svmClassification(
  object,
  assessRes,
  scores = c("prediction", "all", "none"),
  cost,
  sigma,
  fcol = "markers",
  ...
)
```

Arguments

object	An instance of class "MSnSet".
assessRes	An instance of class "GenRegRes", as generated by svmOptimisation.
scores	One of "prediction", "all" or "none" to report the score for the predicted class only, for all classes or none.
cost	If assessRes is missing, a cost must be provided.
sigma	If assessRes is missing, a sigma must be provided.
fcol	The feature meta-data containing marker definitions. Default is markers.
...	Additional parameters passed to svm from package e1071.

Value

An instance of class "MSnSet" with svm and svm.scores feature variables storing the classification results and scores respectively.

Author(s)

Laurent Gatto

Examples

```
library(pRolocdata)
data(dunkley2006)
## reducing parameter search space and iterations
params <- svmOptimisation(dunkley2006, cost = 2^seq(-2,2,2), sigma = 10^seq(-1, 1, 1),  times = 3)
params
plot(params)
f1Count(params)
levelPlot(params)
getParams(params)
res <- svmClassification(dunkley2006, params)
getPredictions(res, fcol = "svm")
getPredictions(res, fcol = "svm", t = 0.75)
plot2D(res, fcol = "svm")
```

---

svmOptimisation *svm parameter optimisation*

Description

Classification parameter optimisation for the support vector machine algorithm.

Usage

```
svmOptimisation(
  object,
  fcol = "markers",
  cost = 2^(-4:4),
  sigma = 10^(-3:2),
  times = 100,
  test.size = 0.2,
  xval = 5,
  fun = mean,
  seed,
  verbose = TRUE,
  ...
)
```

Arguments

object	An instance of class "MSnSet".
fcol	The feature meta-data containing marker definitions. Default is markers.
cost	The hyper-parameter. Default values are 2^-4:4.
sigma	The hyper-parameter. Default values are 10^(-2:3).
times	The number of times internal cross-validation is performed. Default is 100.
test.size	The size of test data. Default is 0.2 (20 percent).
xval	The n-cross validation. Default is 5.
fun	The function used to summarise the xval macro F1 matrices.
seed	The optional random number generator seed.
verbose	A logical defining whether a progress bar is displayed.
...	Additional parameters passed to svm from package e1071.

Details

Note that when performance scores precision, recall and (macro) F1 are calculated, any NA values are replaced by 0. This decision is motivated by the fact that any class that would have either a NA precision or recall would result in an NA F1 score and, eventually, a NA macro F1 (i.e. mean(F1)). Replacing NAs by 0s leads to F1 values of 0 and a reduced yet defined final macro F1 score.

Value

An instance of class "GenRegRes".

Author(s)

Laurent Gatto

See Also

svmClassification and example therein.

Examples

```
library(pRolocdata)
data(dunkley2006)
## reducing parameter search space and iterations
params <- svmOptimisation(dunkley2006, cost = 2^seq(-2,2,2), sigma = 10^seq(-1, 1, 1),  times = 3)
params
plot(params)
f1Count(params)
levelPlot(params)
getParams(params)
res <- svmClassification(dunkley2006, params)
getPredictions(res, fcol = "svm")
getPredictions(res, fcol = "svm", t = 0.75)
plot2D(res, fcol = "svm")
```

---

tagmMcmcTrain *Localisation of proteins using the TAGM MCMC method*

Description

These functions implement the T augmented Gaussian mixture (TAGM) model for mass spectrometry-based spatial proteomics datasets using Markov-chain Monte-Carlo (MCMC) for inference.

Usage

```
tagmMcmcTrain(
  object,
  fcol = "markers",
  method = "MCMC",
  numIter = 1000L,
  burnin = 100L,
  thin = 5L,
  mu0 = NULL,
  lambda0 = 0.01,
  nu0 = NULL,
  S0 = NULL,
  beta0 = NULL,
  u = 2,
  v = 10,
  numChains = 4L,
  BPPARAM = BiocParallel::bpparam(),
  version = 2
)

tagmMcmcPredict(
  object,
  params,
```

```
  fcol = "markers",
  probJoint = FALSE,
  probOutlier = TRUE
)

tagmPredict(
  object,
  params,
  fcol = "markers",
  probJoint = FALSE,
  probOutlier = TRUE
)

tagmMcmcProcess(params)
```

Arguments

object	An MSnbase::MSnSet containing the spatial proteomics data to be passed to tagmMcmcTrain and tagmPredict.
fcol	The feature meta-data containing marker definitions. Default is markers.
method	A charachter() describing the inference method for the TAGM algorithm. Default is "MCMC".
numIter	The number of iterations of the MCMC algorithm. Default is 1000.
burnin	The number of samples to be discarded from the begining of the chain. Default is 100.
thin	The thinning frequency to be applied to the MCMC chain. Default is 5.
mu0	The prior mean. Default is colMeans of the expression data.
lambda0	The prior shrinkage. Default is 0.01.
nu0	The prior degreed of freedom. Default is ncol(exprs(object)) + 2
S0	The prior inverse-wishart scale matrix. Empirical prior used by default.
beta0	The prior Dirichlet distribution concentration. Default is 1 for each class.
u	The prior shape parameter for Beta(u, v). Default is 2
v	The prior shape parameter for Beta(u, v). Default is 10.
numChains	The number of parrallel chains to be run. Default it 4.
BPPARAM	Support for parallel processing using the BiocParallel infrastructure. When missing (default), the default registered BiocParallelParam parameters are used. Alternatively, one can pass a valid BiocParallelParam parameter instance: SnowParam, MulticoreParam, DoparParam, ... see the BiocParallel package for details.
version	A new version that is faster and more memory efficient is implemented as default by setting version == 2. Legacy version is indicated with a 1.
params	An instance of class MCMCParams, as generated by tagmMcmcTrain().
probJoint	A logical(1) indicating whether to return the joint probability matrix, i.e. the probability for all classes as a new tagm.mcmc.joint feature variable.

probOutlier A logical(1) indicating whether to return the probability of being an outlier as a new tagm.mcmc.outlier feature variable. A high value indicates that the protein is unlikely to belong to any annotated class (and is hence considered an outlier).

Details

The tagmMcmcTrain function generates the samples from the posterior distributions (object or class MCMCParams) based on an annotated quantitative spatial proteomics dataset (object of class MSnbase::MSnSet). Both are then passed to the tagmPredict function to predict the sub-cellular localisation of protein of unknown localisation. See the *pRoloc-bayesian* vignette for details and examples. In this implementation, if numerical instability is detected in the covariance matrix of the data a small multiple of the identity is added. A message is printed if this conditioning step is performed.

Value

tagmMcmcTrain returns an instance of class MCMCParams.

tagmMcmcPredict returns an instance of class MSnbase::MSnSet containing the localisation predictions as a new tagm.mcmc.allocation feature variable. The allocation probability is encoded as tagm.mcmc.probability (corresponding to the mean of the distribution probability). In additionm the upper and lower quantiles of the allocation probability distribution are available as tagm.mcmc.probability.lowerquantile and tagm.mcmc.probability.upperquantile feature variables. The Shannon entropy is available in the tagm.mcmc.mean.shannon feature variable, measuring the uncertainty in the allocations (a high value representing high uncertainty; the highest value is the natural logarithm of the number of classes).

tagmMcmcProcess returns an instance of class MCMCParams with its summary slot populated.

References

A Bayesian Mixture Modelling Approach For Spatial Proteomics Oliver M Crook, Claire M Mulvey, Paul D. W. Kirk, Kathryn S Lilley, Laurent Gatto bioRxiv 282269; doi: https://doi.org/10.1101/282269

See Also

The plotEllipse() function can be used to visualise TAGM models on PCA plots with ellipses.

---

testMarkers *Tests marker class sizes*

Description

Tests if the marker class sizes are large enough for the parameter optimisation scheme, i.e. the size is greater that xval + n, where the default xval is 5 and n is 2. If the test is unsuccessful, a warning is thrown.

Usage

```
testMarkers(object, xval = 5, n = 2, fcol = "markers", error = FALSE)
```

Arguments

object	An instance of class "MSnSet".
xval	The number cross-validation partitions. See the xval argument in the parameter optimisation function(s). Default is 5.
n	Number of additional examples.
fcol	The name of the prediction column in the featureData slot. Default is "markers".
error	A logical specifying if an error should be thown, instead of a warning.

Details

In case the test indicates that a class contains too few examples, it is advised to either add some or, if not possible, to remove the class altogether (see minMarkers) as the parameter optimisation is likely to fail or, at least, produce unreliable results for that class.

Value

If successfull, the test invisibly returns NULL. Else, it invisibly returns the names of the classes that have too few examples.

Author(s)

Laurent Gatto

See Also

getMarkers and minMarkers

Examples

```
library("pRolocdata")
data(dunkley2006)
getMarkers(dunkley2006)
testMarkers(dunkley2006)
toosmall <- testMarkers(dunkley2006, xval = 15)
toosmall
try(testMarkers(dunkley2006, xval = 15, error = TRUE))
```

testMSnSet *Create a stratified 'test'* MSnSet

Description

This function creates a stratified 'test' MSnSet which can be used for algorihtmic development. A "MSnSet" containing only the marker proteins, as defined in fcol, is returned with a new feature data column appended called test in which a stratified subset of these markers has been relabelled as 'unknowns'.

Usage

```
testMSnSet(object, fcol = "markers", size = 0.2, seed)
```

Arguments

object	An instance of class "MSnSet"
fcol	The feature meta-data column name containing the marker definitions on which the data will be stratified. Default is markers.
size	The size of the data set to be extracted. Default is 0.2 (20 percent).
seed	The optional random number generator seed.

Value

An instance of class "MSnSet" which contains only the proteins that have a labelled localisation i.e. the marker proteins, as defined in fcol and a new column in the feature data slot called test which has part of the labels relabelled as "unknown" class (the number of proteins renamed as "unknown" is according to the parameter size).

Author(s)

Lisa Breckels

See Also

sampleMSnSet unknownMSnSet markerMSnSet

Examples

```
library(pRolocdata)
data(tan2009r1)
sample <- testMSnSet(tan2009r1)
getMarkers(sample, "test")
all(dim(sample) == dim(markerMSnSet(tan2009r1)))
```

`thetas` *Draw matrix of thetas to test*

Description

The possible weights to be considered is a sequence from 0 (favour auxiliary data) to 1 (favour primary data). Each possible combination of weights for `nclass` classes must be tested. The `thetas` function produces a weight `matrix` for `nclass` columns (one for each class) with all possible weight combinations (number of rows).

Usage

```
thetas(nclass, by = 0.5, length.out, verbose = TRUE)
```

Arguments

`nclass`	Number of marker classes
`by`	The increment of the weights. One of `1`, `0.5`, `0.25`, `2`, `0.1` or `0.05`.
`length.out`	The desired length of the weight sequence.
`verbose`	A `logical` indicating if the weight sequences should be printed out. Default is `TRUE`.

Value

A matrix with all possible theta weight combinations.

Author(s)

Lisa Breckels

Examples

```
dim(thetas(4, by = 0.5))
dim(thetas(4, by = 0.2))
dim(thetas(5, by = 0.2))
dim(thetas(5, length.out = 5))
dim(thetas(6, by = 0.2))
```

undocumented *Undocumented/unexported entries*

Description

This is just a dummy entry for methods from unexported classes that generate warnings during package checking.

Author(s)

Laurent Gatto <lg390@cam.ac.uk>

zerosInBinMSnSet *Compute the number of non-zero values in each marker classes*

Description

The function assumes that its input is a binary `MSnSet` and computes, for each marker class, the number of non-zero expression profiles. The function is meant to be used to produce heatmaps (see the example) and visualise binary (such as GO) `MSnSet` objects and assess their utility: all zero features/classes will not be informative at all (and can be filtered out with `filterBinMSnSet`) while features/classes with many annotations (GO terms) are likely not be be informative either.

Usage

```
zerosInBinMSnSet(object, fcol = "markers", as.matrix = TRUE, percent = TRUE)
```

Arguments

`object`	An instance of class `MSnSet` with binary data.
`fcol`	A `character` defining the feature data variable to be used as markers. Default is `"markers"`.
`as.matrix`	If `TRUE` (default) the data is formatted and returned as a `matrix`. Otherwise, a `list` is returned.
`percent`	If `TRUE`, percentages are returned. Otherwise, absolute values.

Value

A `matrix` or a `list` indicating the number of non-zero value per marker class.

Author(s)

Laurent Gatto

See Also

filterBinMSnSet

Examples

```
library(pRolocdata)
data(hyperLOPIT2015goCC)
zerosInBinMSnSet(hyperLOPIT2015goCC)
zerosInBinMSnSet(hyperLOPIT2015goCC, percent = FALSE)
pal <- colorRampPalette(c("white", "blue"))
library(lattice)
levelplot(zerosInBinMSnSet(hyperLOPIT2015goCC),
          xlab = "Number of non-0s",
          ylab = "Marker class",
          col.regions = pal(140))
```

Index

∗ **classes**
AnnotationParams-class, 7
ClustDist-class, 13
ClustDistList-class, 15
GenRegRes-class, 21
QSep-class, 83
SpatProtVis-class, 92
∗ **methods**
chi2-methods, 10
MLearn-methods, 47
nndist-methods, 57
.MCMCChain *(*MCMCChains-class*)*, 43
.MCMCChains *(*MCMCChains-class*)*, 43
.MCMCParams *(*MCMCChains-class*)*, 43
.MCMCSummary *(*MCMCChains-class*)*, 43
[,ClustDistList,ANY,ANY,ANY-method *(*ClustDistList-class*)*, 15
[,ClustDistList,ANY,missing,missing-method *(*ClustDistList-class*)*, 15
[,MCMCChains,ANY,ANY,ANY-method *(*MCMCChains-class*)*, 43
[,MCMCParams,ANY,ANY,ANY-method *(*MCMCChains-class*)*, 43
[,MartInstanceList,ANY,ANY,ANY-method *(*MartInstance-class*)*, 42
[,MartInstanceList,ANY,ANY-method *(*MartInstance-class*)*, 42
[,MartInstanceList-method *(*MartInstance-class*)*, 42
[[,ClustDistList,ANY,ANY-method *(*ClustDistList-class*)*, 15
[[,ClustDistList,ANY,missing-method *(*ClustDistList-class*)*, 15
[[,MCMCChains,ANY,ANY-method *(*MCMCChains-class*)*, 43
[[,MCMCParams,ANY,ANY-method *(*MCMCChains-class*)*, 43
[[,MartInstanceList,ANY,ANY-method *(*MartInstance-class*)*, 42
[[,MartInstanceList-method *(*MartInstance-class*)*, 42

addLegend, 4, *71, 72*
addMarkers, 6, *52*, *81, 82*
AnnotationParams *(*AnnotationParams-class*)*, 7
AnnotationParams-class, 7
as.data.frame.MartInstance *(*MartInstance-class*)*, 42
as.data.frame.MartInstanceList *(*MartInstance-class*)*, 42

chains *(*MCMCChains-class*)*, 43
checkFeatureNamesOverlap, 8
checkFvarOverlap, 9
chi2, *17*
chi2 *(*chi2-methods*)*, 10
chi2,matrix,matrix-method *(*chi2-methods*)*, 10
chi2,matrix,numeric-method *(*chi2-methods*)*, 10
chi2,numeric,matrix-method *(*chi2-methods*)*, 10
chi2,numeric,numeric-method *(*chi2-methods*)*, 10
chi2-methods, 10
chol2inv, *63*, *65*
class::QSep *(*QSep-class*)*, 83
class:AnnotationParams *(*AnnotationParams-class*)*, 7
class:ClustDist *(*ClustDist-class*)*, 13
class:ClustDistList *(*ClustDistList-class*)*, 15
class:GenRegRes *(*GenRegRes-class*)*, 21
class:MAPParams *(*MAPParams-class*)*, 38
class:MCMCChain *(*MCMCChains-class*)*, 43
class:MCMCChains *(*MCMCChains-class*)*, 43
class:MCMCParams *(*MCMCChains-class*)*, 43
class:MCMCSummary *(*MCMCChains-class*)*, 43

class:SpatProtVis (SpatProtVis-class), 92
class:ThetaRegRes (GenRegRes-class), 21
classWeights, 11
ClustDist, *13*, *15*, *25*
ClustDist (ClustDist-class), 13
clustDist, 12
ClustDist-class, 13
ClustDistList, *13*, *25*
ClustDistList (ClustDistList-class), 15
ClustDistList-class, 15
cmdscale, *70*
coda::mcmc, *43*
col1 (plot2Ds), 73
col2 (plot2Ds), 73
combineThetaRegRes (GenRegRes-class), 21

data1 (plot2Ds), 73
data2 (plot2Ds), 73
dendrogram, *51*
dist, *51*

empPvalues, *11*, 16

f1Count (GenRegRes-class), 21
f1Count,GenRegRes-method (GenRegRes-class), 21
f1Count,ThetaRegRes-method (GenRegRes-class), 21
favourPrimary (GenRegRes-class), 21
fDataToUnknown, 17
FeaturesOfInterest, *27*
filterAttrs (MartInstance-class), 42
filterBinMSnSet, 18, *103, 104*
filterMaxMarkers, 19
filterMinMarkers, 20
filterNA, *68*
filterZeroCols, *18*, 21
filterZeroRows, *18*
filterZeroRows (filterZeroCols), 21

GenRegRes, *29*, *31*, *36, 37*, *53*, *55*, *58*, *60*, *63*, *65*, *79, 80*, *86, 87*, *95, 96*
GenRegRes (GenRegRes-class), 21
GenRegRes-class, 21
getAnnotationParams (AnnotationParams-class), 7
getF1Scores (GenRegRes-class), 21
getF1Scores,GenRegRes-method (GenRegRes-class), 21
getF1Scores,ThetaRegRes-method (GenRegRes-class), 21
getFilterList (MartInstance-class), 42
getGOFromFeatures, *7*
getLisacol (setLisacol), 89
getMarkerClasses, 23, *24*, *52*
getMarkers, 24, *24*, *52*, *100*
getMartInstanceList (MartInstance-class), 42
getMartTab (MartInstance-class), 42
getNormDist, 25
getOldcol (setLisacol), 89
getParams (GenRegRes-class), 21
getParams,ClustRegRes-method (undocumented), 103
getParams,GenRegRes-method (GenRegRes-class), 21
getParams,ThetaRegRes-method (GenRegRes-class), 21
getPredictions, 26, *46*, *62*
getRegularisedParams (GenRegRes-class), 21
getRegularisedParams,GenRegRes-method (GenRegRes-class), 21
getRegularizedParams (GenRegRes-class), 21
getRegularizedParams,GenRegRes-method (GenRegRes-class), 21
getSeed (GenRegRes-class), 21
getSeed,GenRegRes-method (GenRegRes-class), 21
getStockbg (setLisacol), 89
getStockcol, *50*, *71*, *74*
getStockcol (setLisacol), 89
getStockpch, *71*
getStockpch (setLisacol), 89
getUnknownbg (setLisacol), 89
getUnknowncol, *76*
getUnknowncol (setLisacol), 89
getUnknownpch (setLisacol), 89
getWarnings (GenRegRes-class), 21
getWarnings,GenRegRes-method (GenRegRes-class), 21
geweke_test (mcmc_get_outliers), 42
ginv, *63*, *65*

hclust, *51*
hexbin, *70*
highlightOnPlot, 27

highlightOnPlot3D *(highlightOnPlot)*, 27

identify, *71*
isMrkMat *(mrkVecToMat)*, 51
isMrkVec *(mrkVecToMat)*, 51

knn, *29*, *31*
knnClassification, 29, *31*
knnOptimisation, *21*, *29*, 30
knnOptimization *(knnOptimisation)*, 30
knnPrediction *(knnClassification)*, 29
knnRegularisation *(knnOptimisation)*, 30
knntlClassification, 31, *35*
knntlOptimisation, *21*, *32*, 33
kpca, *70*
ksvm, *36, 37*
ksvmClassification, 35, *38*
ksvmOptimisation, *36*, 37
ksvmOptimization *(ksvmOptimisation)*, 37
ksvmPrediction *(ksvmClassification)*, 35
ksvmRegularisation *(ksvmOptimisation)*, 37

lapply,ClustDistList-method *(ClustDistList-class)*, 15
lapply,MartInstanceList,ANY-method *(MartInstance-class)*, 42
lapply,MartInstanceList-method *(MartInstance-class)*, 42
lda, *70*
legend, *4, 5*
length,ClustDistList-method *(ClustDistList-class)*, 15
length,MCMCChains-method *(MCMCChains-class)*, 43
length,MCMCParams-method *(MCMCChains-class)*, 43
levelPlot *(GenRegRes-class)*, 21
levelplot, *84*
levelPlot,ClustRegRes-method *(undocumented)*, 103
levelPlot,GenRegRes-method *(GenRegRes-class)*, 21
levelPlot,QSep-method *(QSep-class)*, 83
logPosteriors *(MAPParams-class)*, 38

makeGoSet, *7*
MAPParams, *39*
MAPParams *(MAPParams-class)*, 38
MAPParams(), *40*
MAPParams-class, 38
markerMSnSet, 41, *52*, *88*, *101*
markers, *6*, *24*, *41*, *82*, *88*
markers *(mrkVecToMat)*, 51
MartInstance *(MartInstance-class)*, 42
MartInstance-class, 42
MartInstanceList *(MartInstance-class)*, 42
MartInstanceList-class *(MartInstance-class)*, 42
mcmc_burn_chains *(mcmc_get_outliers)*, 42
mcmc_get_meanComponent *(mcmc_get_outliers)*, 42
mcmc_get_meanoutliersProb *(mcmc_get_outliers)*, 42
mcmc_get_outliers, 42
mcmc_pool_chains *(mcmc_get_outliers)*, 42
mcmc_thin_chains *(mcmc_get_outliers)*, 42
MCMCChain *(MCMCChains-class)*, 43
MCMCChain-class *(MCMCChains-class)*, 43
MCMCChains *(MCMCChains-class)*, 43
MCMCChains-class, 43
MCMCParams-class *(MCMCChains-class)*, 43
MCMCSummary *(MCMCChains-class)*, 43
MCMCSummary-class *(MCMCChains-class)*, 43
minMarkers, 45, *100*
mixing_posterior_check, 46
MLearn, *47*
MLearn,formula,MSnSet,clusteringSchema,missing-method *(MLearn-methods)*, 47
MLearn,formula,MSnSet,learnerSchema,numeric-method *(MLearn-methods)*, 47
MLearn,formula,MSnSet,learnerSchema,xvalSpec-method *(MLearn-methods)*, 47
MLearn-methods, 47
MLearnMSnSet *(MLearn-methods)*, 47
move2Ds, 48, *74*
mrkConsProfiles, 49
mrkConsProfiles(), *75*
mrkEncoding *(mrkVecToMat)*, 51
mrkHClust, 50, *50*
mrkMatAndVec *(mrkVecToMat)*, 51
mrkMatToVec *(mrkVecToMat)*, 51
mrkVecToMat, 51
MSnbase::MSnSet, *39, 40*, *98, 99*
MSnSet, *12*, *23, 24*, *26*, *29, 30*, *32*, *34*, *36, 37*, *46, 47*, *53, 54*, *58, 59*, *62, 63*, *65*, *74*,

79, *80*, *83*, *86–88*, *92*, *95*, *96*, *100*, *101*
MSnSetList, *48*, *74*
MSnSetMLean *(*MLearn-methods*)*, 47

naiveBayes, *53*, *55*
names,ClustDistList-method *(*ClustDistList-class*)*, 15
names,QSep-method *(*QSep-class*)*, 83
names<-,ClustDistList,ANY-method *(*ClustDistList-class*)*, 15
names<-,QSep,character-method *(*QSep-class*)*, 83
nbClassification, 53, *55*
nbOptimisation, *53*, 54
nbOptimization *(*nbOptimisation*)*, 54
nbPrediction *(*nbClassification*)*, 53
nbRegularisation *(*nbOptimisation*)*, 54
nDatasets *(*MartInstance-class*)*, 42
nicheMeans2D, 55
nipals, *70*
nndist *(*nndist-methods*)*, 57
nndist,matrix,matrix-method *(*nndist-methods*)*, 57
nndist,matrix,missing-method *(*nndist-methods*)*, 57
nndist,MSnSet,missing-method *(*nndist-methods*)*, 57
nndist-methods, 57
nnet, *58*, *59*
nnetClassification, 58, *60*
nnetOptimisation, *58*, 59
nnetOptimization *(*nnetOptimisation*)*, 59
nnetPrediction *(*nnetClassification*)*, 58
nnetRegularisation *(*nnetOptimisation*)*, 59

orderGoAnnotations, 60
orgQuants, *27*, 62

par, *71*, *77*
pch, *71*
perTurboClassification, 63, *65*
perTurboOptimisation, *63*, 64
perTurboOptimization *(*perTurboOptimisation*)*, 64
phenoDisco, 66
plot, *77*
plot,ClustDist,MSnSet-method *(*ClustDist-class*)*, 13
plot,ClustDistList,missing-method *(*ClustDistList-class*)*, 15
plot,ClustRegRes,missing-method *(*undocumented*)*, 103
plot,GenRegRes,missing-method *(*GenRegRes-class*)*, 21
plot,MCMCParams,character-method *(*mcmc_get_outliers*)*, 42
plot,QSep,missing-method *(*QSep-class*)*, 83
plot,QSep-method *(*QSep-class*)*, 83
plot,SpatProtVis,missing-method *(*SpatProtVis-class*)*, 92
plot,ThetaRegRes,missing-method *(*GenRegRes-class*)*, 21
plot2D, *4*, *5*, *50*, 69, *74*, *92*, *93*
plot2Dmethods, *92*
plot2Dmethods *(*plot2D*)*, 69
plot2Ds, *48*, *72*, 73
plot3d, *71*
plot3D,MSnSet-method *(*plot2D*)*, 69
plotConsProfiles, 75
plotDist, *72*, 76
plotEllipse, *72*, 77
plotEllipse(), *40*, *99*
plsda, *79*, *80*
plsdaClassification, 78, *81*
plsdaOptimisation, *21*, 79, *79*
plsdaOptimization *(*plsdaOptimisation*)*, 79
plsdaPrediction *(*plsdaClassification*)*, 78
plsdaRegularisation *(*plsdaOptimisation*)*, 79
prcomp, *70*, *74*
pRolocmarkers, *6*, *52*, 81

QSep *(*QSep-class*)*, 83
qsep *(*QSep-class*)*, 83
QSep-class, 83

randomForest, *86*, *87*
rfClassification, 85, *87*
rfOptimisation, 86, *86*
rfOptimization *(*rfOptimisation*)*, 86
rfPrediction *(*rfClassification*)*, 85
rfRegularisation *(*rfOptimisation*)*, 86

Rtsne, *70*

sampleMSnSet, *41*, 88, *101*
sapply,ClustDistList-method *(ClustDistList-class)*, 15
sapply,MartInstanceList,ANY-method *(MartInstance-class)*, 42
sapply,MartInstanceList-method *(MartInstance-class)*, 42
setAnnotationParams *(AnnotationParams-class)*, 7
setLisacol, 89
setOldcol *(setLisacol)*, 89
setStockbg *(setLisacol)*, 89
setStockcol *(setLisacol)*, 89
setStockpch *(setLisacol)*, 89
setUnknownbg *(setLisacol)*, 89
setUnknowncol *(setLisacol)*, 89
setUnknownpch *(setLisacol)*, 89
show,AnnotationParams-method *(AnnotationParams-class)*, 7
show,ClustDist-method *(ClustDist-class)*, 13
show,ClustDistList-method *(ClustDistList-class)*, 15
show,ClustRegRes-method *(undocumented)*, 103
show,ComponentParam-method *(MCMCChains-class)*, 43
show,GenRegRes-method *(GenRegRes-class)*, 21
show,MAPParams-method *(MAPParams-class)*, 38
show,MartInstance-method *(MartInstance-class)*, 42
show,MCMCChain-method *(MCMCChains-class)*, 43
show,MCMCChains-method *(MCMCChains-class)*, 43
show,MCMCParams-method *(MCMCChains-class)*, 43
show,QSep-method *(QSep-class)*, 83
show,SpatProtVis-method *(SpatProtVis-class)*, 92
show,ThetaRegRes-method *(GenRegRes-class)*, 21
showMrkMat *(mrkVecToMat)*, 51
solve, *63*, *65*
spatial2D, 91
SpatProtVis *(SpatProtVis-class)*, 92
SpatProtVis-class, 92
sub, *17*
subsetMarkers, 93
summary,QSep-method *(QSep-class)*, 83
svd, *63*, *65*
svm, *95*, *96*
svmClassification, *11*, 94, *96*
svmOptimisation, *11*, *21*, 95, *95*
svmOptimization *(svmOptimisation)*, 95
svmPrediction *(svmClassification)*, 94
svmRegularisation, *63*
svmRegularisation *(svmOptimisation)*, 95

tagmMapPredict *(MAPParams-class)*, 38
tagmMapTrain *(MAPParams-class)*, 38
tagmMapTrain(), *39*, *40*
tagmMcmcPredict *(tagmMcmcTrain)*, 97
tagmMcmcProcess *(tagmMcmcTrain)*, 97
tagmMcmcTrain, 97
tagmMcmcTrain(), *98*
tagmPredict *(tagmMcmcTrain)*, 97
testMarkers, 99
testMSnSet, *41*, *88*, 101
ThetaRegRes *(GenRegRes-class)*, 21
ThetaRegRes-class *(GenRegRes-class)*, 21
thetas, *34*, 102

umap, *70*
undocumented, 103
unknownMSnSet, *88*, *101*
unknownMSnSet *(markerMSnSet)*, 41

Versioned, *84*

xvalSpec, *47*

zerosInBinMSnSet, *18*, 103