

Package ‘wrProteo’

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Title Proteomics Data Analysis Functions

Author Wolfgang Raffelsberger [aut, cre]

Maintainer Wolfgang Raffelsberger <w.raffelsberger@gmail.com>

Description Data analysis of proteomics experiments by mass spectrometry is supported by this collection of functions mostly dedicated to the analysis of (bottom-up) quantitative (XIC) data. Fasta-formatted proteomes (eg from UniProt Consortium <doi:10.1093/nar/gky1049>) can be read with automatic parsing and multiple annotation types (like species origin, abbreviated gene names, etc) extracted. Quantitative proteomics measurements frequently contain multiple NA values, due to physical absence of given peptides in some samples, limitations in sensitivity or other reasons. The functions provided here help to inspect graphically the data to investigate the nature of NA-values via their respective replicate measurements and to help/confirm the choice of NA-replacement by low random values. Dedicated filtering and statistical testing using the framework of package 'limma' <doi:10.18129/B9.bioc.limma> can be run, enhanced by multiple rounds of NA-replacements to provide robustness towards rare stochastic events. Multi-species samples, as frequently used in benchmark-tests (eg Navarro et al 2016 <doi:10.1038/nbt.3685>, Ramus et al 2016 <doi:10.1016/j.jprot.2015.11.011>), can be run with special options separating the data into sub-groups during normalization and testing. As example the data-set from Ramus et al 2016 <doi:10.1016/j.jprot.2015.11.011> is provided quantified by MaxQuant (Tyanova et al 2016 <doi:10.1038/nprot.2016.136>), ProteomeDiscoverer, OpenMS (<doi:10.1038/nmeth.3959>) and Proline (Bouyssie et al 2020 <doi:10.1093/bioinformatics/btaa118>). Subsequently, ROC curves (Hand and Till 2001 <doi:10.1023/A:1010920819831>) can be constructed to compare multiple analysis approaches.

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AAmass	<i>Molecular mass for amino-acids</i>
--------	---------------------------------------

Description

Calculate molecular mass based on atomic composition

Usage

```
AAmass(massTy = "mono", inPept = TRUE, inclSpecAA = FALSE)
```

Arguments

massTy	(character) 'mono' or 'average'
inPept	(logical) remove H ₂ O corresponding to water loss at peptide bond formaton
inclSpecAA	(logical) include ornithine O & selenocysteine U

Value

vector with masses for all amino-acids (argument 'massTy' to switch form mono-isotopic to average mass)

See Also

[massDeFormula](#), [convToNum](#)

Examples

```
massDeFormula(c("12H12O", "HO", " 2H 1 Se, 6C 2N", "HSeCN", " ", "e"))  
AAmass()
```

AucROC	<i>AUC from ROC-curves</i>
--------	----------------------------

Description

This function calculates the AUC (area under the curve) from ROC data in matrix of specificity and sensitivity values, as provided in the output from [summarizeForROC](#). Note that numerous other packages also provide support for working with ROC-curves : Eg [rocPkgShort](#), [ROCR](#), [pROC](#) or [ROCit](#)

Usage

```
AucROC(dat, useCol = c("spec", "sens"), silent = FALSE, callFrom = NULL)
```

Arguments

dat	(matrix or data.frame) main input containing sensitivity and specificity data (from summarizeForROC)
useCol	(character or integer) column names to be used: 1st for specificity and 2nd for sensitivity count columns
silent	(logical) suppress messages
callFrom	(character) allows easier tracking of message(s) produced

Value

matrix including imputed values or list of final and matrix with number of imputed by group (plus optional plot)

See Also

preparing ROC data [summarizeForROC](#), (re)plot the ROC figure [plotROC](#)

Examples

```
set.seed(2019); test1 <- list(annot=cbind(spec=c(rep("b",35),letters[sample.int(n=3,
  size=150,replace=TRUE)])), BH=matrix(c(runif(35,0,0.01),runif(150)),ncol=1))
roc1 <- summarizeForROC(test1,spec=c("a","b","c"))
AucROC(roc1)
```

cleanListCoNames	<i>Selective batch cleaning of sample- (ie column-) names in list</i>
------------------	---

Description

This function allows to manipulate sample-names (ie colnames) from data stored as multiple matrixes or data.frames in multiple sheets of a list in a batch-wise manner. Import functions such as `readMaxQuantFile()` organize initial flat files into lists (of matrixes) of the different types of data. Many times all column names in such lists carry long names including redundant information, like the overall experiment name or date, etc. The aim of this function is to facilitate 'cleaning' the sample- (ie column-) names to obtain short and concise names. Character terms to be removed (via argument `rem`) and/or replaced/substituted (via argument `subst`) should be given as they are, characters with special behaviour in `grep` (like `'.`) will be protected internally. Note, that the character substitution part will be done first, and the removal part (without character replacement) afterwards.

Usage

```
cleanListCoNames(
  dat,
  rem = NULL,
  subst = c("-", "_"),
  lstE = c("raw", "quant", "counts"),
```

```

    silent = FALSE,
    callFrom = NULL
  )

```

Arguments

dat	(list) main input
rem	(character) character string to be removed, may be named 'left' and 'right' for more specific exact pattern matching (this part will be performed before character substitutions by subst)
subst	(character of length=2, or matrix with 2 columns) pair(s) of character-strings for replacement (1st as search-item and 2nd as replacement); this part is performed after character-removal via rem
lstE	(character, length=1) names of list-elements where colnames should be cleaned
silent	(logical) suppress messages
callFrom	(character) allow easier tracking of message(s) produced

Value

list (equivalent to input dat)

See Also

[grep](#)

Examples

```

dat1 <- matrix(1:12, ncol=4, dimnames=list(1:3, paste0("sample_R.",1:4)))
dat1 <- list(raw=dat1, quant=dat1, notes="other..")
cleanListCoNames(dat1, rem=c(left="sample_"),c(".","-"))

```

combineMultFilterNAimput

Combine multiple filters on NA-imputed data

Description

In most omics data-analysis one needs to employ a certain number of filtering strategies to avoid getting artifacts to the step of statistical testing. `combineMultFilterNAimput` takes on one side the original data and on the other side NA-imputed data to create several different filters and to finally combine them. A filter aiming to take away the least abundant values (using the imputed data) is fine-tuned by the argument `abundThr`. This step compares the means for each group and line, at least one group-mean has to be > the threshold (based on hypothesis that if all conditions represent extremely low measures their differential may not be determined with certainty). In contrast, the filter addressing the number of missing values (NA) uses the original data, the arguments

colTotNa,minSpeNo and minTotNo are used at this step. Basically, this step allows defining a minimum content of 'real' (ie non-NA) values for further considering the measurements as reliable. This part uses internally [presenceFilt](#) for filtering elevated content of NA per line. Finally, this function combines both filters (as matrix of FALSE and TRUE) on NA-imputed and original data and retruns a vector of logical values if corresponding lines passe all filter criteria.

Usage

```
combineMultFilterNAimput(
  dat,
  imputed,
  grp,
  annDat = NULL,
  abundThr = NULL,
  colRazNa = NULL,
  colTotNa = NULL,
  minSpeNo = 1,
  minTotNo = 2,
  silent = FALSE,
  callFrom = NULL
)
```

Arguments

dat	(matrix or data.frame) main data (may contain NA)
imputed	(character) same as 'dat' but with all NA imputed
grp	(character or factor) define groups of replicates (in columns of 'dat')
annDat	(matrix or data.frame) annotation data (should match lines of 'dat')
abundThr	(numeric) optional threshold filter for minimum abundance
colRazNa	(character) if razor peptides are used: column name for razor peptide count
colTotNa	(character) column name for total peptide count
minSpeNo	(integer) minimum number of specific peptides for maintaining proteins
minTotNo	(integer) minimum total ie max razor number of peptides
silent	(logical) suppress messages
callFrom	(character) allows easier tracking of message(s) produced

Value

vector of logical values if corresponding line passes filter criteria

See Also

[presenceFilt](#)

Examples

```

set.seed(2013)
datT6 <- matrix(round(rnorm(300)+3,1),ncol=6,dimnames=list(paste("li",1:50,sep=""),letters[19:24]))
datT6 <- datT6 +matrix(rep(1:nrow(datT6),ncol(datT6)),ncol=ncol(datT6))
datT6[6:7,c(1,3,6)] <- NA
datT6[which(datT6 < 11 & datT6 > 10.5)] <- NA
datT6[which(datT6 < 6 & datT6 > 5)] <- NA
datT6[which(datT6 < 4.6 & datT6 > 4)] <- NA
datT6b <- matrixNAneighbourImpute(datT6,grp=gl(2,3))
datT6c <- combineMultFilterNAimput(datT6,datT6b,grp=gl(2,3),abundThr=2)

```

convAASeq2mass

Molecular mass for amino-acids

Description

This function calculates the molecular mass of one-letter code amino-acid sequences.

Usage

```

convAASeq2mass(
  x,
  massTy = "mono",
  seqName = TRUE,
  silent = FALSE,
  callFrom = NULL
)

```

Arguments

x	(character) aminoacid sequence (single upper case letters for describing a peptide/protein)
massTy	(character) default 'mono' for mono-isotopic masses (alternative 'average')
seqName	(logical) optional (alternative) names for the content of 'x' (ie aa seq) as name (always if 'x' has no names)
silent	(logical) suppress messages
callFrom	(character) allow easier tracking of message(s) produced

Value

vector with masses for all amino-acids (argument 'massTy' to switch from mono-isotopic to average mass)

See Also

[massDeFormula](#), [AAMass](#), [convToNum](#)

Examples

```
convAASeq2mass(c("PEPTIDE", "fPROTEINES"))
pep1 <- c(aa="AAAA", de="DEFDEF")
convAASeq2mass(pep1, seqN=FALSE)
```

corColumnOrder	<i>Order columns in list of matrixes (or matrix)</i>
----------------	--

Description

This function orders columns in list of matrixes (or matrix) according to argument `sampNames`. This function can be used to adjust/correct the order of samples after reading data using `readMaxQuantFile()`, `readPDExport()` etc. The input may also be MArrayLM-type object from package `limma` or from `moderTestXgrp` or `moderTest2grp`.

Usage

```
corColumnOrder(
  dat,
  sampNames,
  useListElem = c("quant", "raw"),
  silent = FALSE,
  callFrom = NULL
)
```

Arguments

<code>dat</code>	(matrix, list or MArrayLM-object from <code>limma</code>) main input of which columns should get re-ordered, may be output from <code>moderTestXgrp</code> or <code>moderTest2grp</code> .
<code>sampNames</code>	(character) column-names in desired order for output
<code>useListElem</code>	(character) in case <code>dat</code> is list, all list-elements who's columns should get (re-)ordered
<code>silent</code>	(logical) suppress messages
<code>callFrom</code>	(character) allow easier tracking of message(s) produced

Value

object of same class as input `dat`

See Also

`moderTestXgrp` for single comparisons, `order`

Examples

```

grp <- factor(rep(LETTERS[c(3,1,4)], c(2,3,3)))
dat1 <- matrix(1:15, ncol=5, dimnames=list(NULL,c("D","A","C","E","B")))
corColumnOrder(dat1,sampNames=LETTERS[1:5])

dat1 <- list(quant=dat1,row=dat1)
dat1
corColumnOrder(dat1,sampNames=LETTERS[1:5])

```

countNoOfCommonPeptides

Compare in-silico digested proteomes for unique and shared peptides, counts per protein or as peptides Compare in-silico digested proteomes for unique and shared peptides, counts per protein or as peptides. The in-silico digestion may be performed separately using the package [R/cleaver](https://bioconductor.org/packages/release/bioc/html/cleaver.html). Note: input must be list (or multiple names lists) of proteins with their respective peptides (eg by in-silico digestion).

Description

Compare in-silico digested proteomes for unique and shared peptides, counts per protein or as peptides

Compare in-silico digested proteomes for unique and shared peptides, counts per protein or as peptides. The in-silico digestion may be performed separately using the package [cleaver](#). Note: input must be list (or multiple names lists) of proteins with their respective peptides (eg by in-silico digestion).

Usage

```

countNoOfCommonPeptides(
  ...,
  prefix = c("Hs", "Sc", "Ec"),
  sep = "_",
  silent = FALSE,
  callFrom = NULL
)

```

Arguments

...	(list) multiple lists of (in-silico) digested proteins (typically protein ID as names) with their respective peptides (AA sequence), one entry for each species
prefix	(character) optional (species-) prefix for entries in '...', will be only considered if '...' has no names
sep	(character) concatenation symbol

silent (logical) suppress messages
 callFrom (character) allows easier tracking of message(s) produced

Value

list with \$byPep as list of logical matrixes for each peptide (as line) and unique/shared/etc for each species; \$byProt as list of matrixes with count data per proten (as line) for each species; \$tab with simple summary-type count data

See Also

[readFasta2](#) and/or cleave-methods in package [cleaver](#)

Examples

```
## The example mimics a proteomics experiment where extracts form E coli and
## Saccharomyces cerevisiae were mixed, thus not all peptdes may occur unique.
(mi2 = countNoOfCommonPeptides(Ec=list(E1=letters[1:4],E2=letters[c(3:7)],
  E3=letters[c(4,8,13)],E4=letters[9]),Sc=list(S1=letters[c(2:3,6)],
  S2=letters[10:13],S3=letters[c(5,6,11)],S4=letters[c(11)],S5="n")))
## a .. uni E, b .. inteR, c .. inteR(+intra E), d .. intra E (no4), e .. inteR,
## f .. inteR +intra E (no6), g .. uni E, h .. uni E no 8), i .. uni E,
## j .. uni S (no10), k .. intra S (no11), l .. uni S (no12), m .. inteR (no13)
lapply(mi2$byProt,head)
mi2$tab
```

extractTestingResults *Extract results from moderated t-tests*

Description

This function allows convenient access to results produced using the functions [moderTest2grp](#) or [moderTestXgrp](#). The user can define the threshold which type of multiple testing correction should be used (as long as the multiple testing correction method was actually performed as part of testing).

Usage

```
extractTestingResults(
  stat,
  compNo = 1,
  statTy = "BH",
  thrsh = 0.05,
  FCthrs = 1.5,
  annotCol = c("Accession", "EntryName", "GeneName"),
  nSign = 6,
  add = "allMeans",
  filename = NULL,
  fileTy = "csvUS",
```

```

    silent = FALSE,
    callFrom = NULL
  )

```

Arguments

stat	('MArrayLM'-object or list) Designed for the output from moderTest2grp or moderTestXgrp
compNo	(integer) the comparison number/index to be used
statTy	(character) the multiple-testing correction type to be considered when looking for significant changes with threshold thrsh (depends on which have been run initially with moderTest2grp or moderTestXgrp)
thrsh	(numeric) the threshold to be applied on statTy for the result of the statistical testing (after multiple testing correction)
FCthrs	(numeric) Fold-Change threshold given as Fold-change and NOT log2(FC), default at 1.5 (for filtering at M-value =0.585)
annotCol	(character) column-names from the annotation to be included
nSign	(integer) number of significant digits whe returning results
add	(character) additional groups to add (so far only "allMeans" available) in addition to the means used in the pairwise comparison
filename	(character) optional (path and) file-name for exporting results to csv-file
fileTy	(character) file-type to be used with argument filename, may be 'csvEur' or 'csvUS'
silent	(logical) suppress messages
callFrom	(character) allow easier tracking of message(s) produced

Value

limma-type MA-object (list)

See Also

[moderTest2grp](#) for single comparisons, [moderTestXgrp](#) for multiple comparisons, [lmFit](#) and the eBayes-family of functions in package [limma](#)

Examples

```

grp <- factor(rep(LETTERS[c(3,1,4)],c(2,3,3)))
set.seed(2017); t8 <- matrix(round(rnorm(208*8,10,0.4),2), ncol=8,
  dimnames=list(paste(letters[],rep(1:8,each=26),sep=""), paste(grp,c(1:2,1:3,1:3),sep="")))
t8[3:6,1:2] <- t8[3:6,1:2] +3 # augment lines 3:6 (c-f)
t8[5:8,c(1:2,6:8)] <- t8[5:8,c(1:2,6:8)] -1.5 # lower lines
t8[6:7,3:5] <- t8[6:7,3:5] +2.2 # augment lines
## expect to find C/A in c,d,g, (h)
## expect to find C/D in c,d,e,f
## expect to find A/D in f,g,(h)
library(wrMisc) # for testing we'll use this package

```

```
test8 <- moderTestXgrp(t8, grp)
extractTestingResults(test8)
```

extrSpeciesAnnot *Extract species annotation*

Description

extrSpeciesAnnot identifies species-related annotation (as suffix to identifiers) for data containing multiple species and returns alternative (short) names. This function also suppresses extra heading or trailing space or punctuation characters. In case multiple tags are found, the last tag is reported and a message of alert may be displayed.

Usage

```
extrSpeciesAnnot(
  annot,
  spec = c("_CONT", "_HUMAN", "_YEAST", "_ECOLI"),
  shortNa = c("cont", "H", "S", "E"),
  silent = FALSE,
  callFrom = NULL
)
```

Arguments

annot	(character) vector with initial annotation
spec	(character) the tags to be identified
shortNa	(character) the final abbreviation used, order and length must fit to argument annot
silent	(logical) suppress messages
callFrom	(character) allows easier tracking of messages produced

Value

character vector with single (last of multiple) term if found in argument annot

See Also

[grep](#)

Examples

```
spec <- c("keratin_CONT", "AB_HUMAN", "CD_YEAST", "EF_G_HUMAN", "HI_HUMAN_ECOLI", "_YEAST_012")
extrSpeciesAnnot(spec)
```

isolNAneighb	<i>Isolate NA-neighbours</i>
--------------	------------------------------

Description

This functions extracts all replicate-values where at least one of the replicates is NA. Then, the non-NA values are sorted by the number of NAs which occurred in this group of replicates. A list with all NA-neighbours organized by the number of NAs gets returned.

Usage

```
isolNAneighb(mat, gr, maxHi = 3, iniCheck = TRUE, callFrom = NULL)
```

Arguments

mat	(matrix or data.frame) main data (may contain NA)
gr	(character or factor) grouping of columns of 'mat', replicate association
maxHi	(integer) maximum count of NAs to consider separately (higher ones will be counted/pooled as maxHi)
iniCheck	(logical) check at beginning if executing this function is useful (presence any NA)
callFrom	(character) allow easier tracking of message(s) produced

Value

list with NA-neighbours sorted by number of NAs in replicate group

See Also

this function gets used by [matrixNAneighbourImpute](#) and [testRobustToNAimputation](#); estimation of mode [stableMode](#); detection of NAs [na.fail](#)

Examples

```
mat1 <- c(22.2, 22.5, 22.2, 22.2, 21.5, 22.0, 22.1, 21.7, 21.5, 22, 22.2, 22.7,
  NA, NA, NA, NA, NA, NA, NA, 21.2, NA, NA, NA, NA,
  NA, 22.6, 23.2, 23.2, 22.4, 22.8, 22.8, NA, 23.3, 23.2, NA, 23.7,
  NA, 23.0, 23.1, 23.0, 23.2, 23.2, NA, 23.3, NA, NA, 23.3, 23.8)
mat1 <- matrix(mat1, ncol=12, byrow=TRUE)
gr4 <- gl(3, 4)
isolNAneighb(mat1, gr4)
```

massDeFormula	<i>molecular mass from chemical formula</i>
---------------	---

Description

Calculate molecular mass based on atomic composition

Usage

```
massDeFormula(  
  comp,  
  massTy = "mono",  
  rmEmpty = FALSE,  
  silent = FALSE,  
  callFrom = NULL  
)
```

Arguments

comp	(character) atomic composition
massTy	(character) 'mono' or 'average'
rmEmpty	(logical) suppress empty entries
silent	(logical) suppress messages
callFrom	(character) allow easier tracking of messages produced

Value

numeric vector with mass

See Also

[convToNum](#)

Examples

```
massDeFormula(c("12H12O", "HO", " 2H 1 Se, 6C 2N", "HSeCN", " ", "e"))
```

`matrixNAinspect`*Histogram of content of NAs in matrix*

Description

`matrixNAinspect` makes histograms of the full data and shows sub-population of NA-neighbour values. The aim of this function is to investigate the nature of NA values in matrix (of experimental measures) where replicate measurements are available. If a given element was measured twice, and one of these measurements revealed a NA while the other one gave a (finite) numeric value, the non-NA-value is considered a NA-neighbour. The subpopulation of these NA-neighbour values will then be highlighted in the resulting histogram. In a number of experimental settings some actual measurements may not meet an arbitrary defined baseline (as 'zero') or may be too low to be distinguishable from noise that associated measures were initially recorded as NA. In several types of measurements in proteomics and transcriptomics this may happen. So this function allows to collect all NA-neighbour values and compare them to the global distribution of the data to investigate if NA-neighbours are typically very low values. In case of data with multiple replicates NA-neighbour values may be distinguished for the case of 2 NA per group/replicate-set. The resulting plots are typically used to decide if and how NA values may get replaced by imputed random values or whether measures containing NA-values should rather be omitted. Of course, such decisions do have a strong impact on further steps of data-analysis and should be performed with care.

Usage

```
matrixNAinspect(  
  dat,  
  gr,  
  retnNA = TRUE,  
  xLab = NULL,  
  tit = NULL,  
  xLim = NULL,  
  silent = FALSE,  
  callFrom = NULL  
)
```

Arguments

<code>dat</code>	(matrix or data.frame) main numeric data
<code>gr</code>	(character or factor) grouping of columns of <code>dat</code> indicating who is a replicate of whom (ie the length of 'gr' must be equivalent to the number of columns in 'dat')
<code>retnNA</code>	(logical) report number of NAs in graphic
<code>xLab</code>	(character) custom x-label
<code>tit</code>	(character) custom title
<code>xLim</code>	(numerical,length=2) custom x-axis limits
<code>silent</code>	(logical) suppress messages
<code>callFrom</code>	(character) allow easier tracking of messages produced

Value

graphic only

See Also

[hist](#), [na.fail](#), [naOmit](#)

Examples

```
set.seed(2013)
datT6 <- matrix(round(rnorm(300)+3,1),ncol=6,dimnames=list(paste("li",1:50,sep=""),letters[19:24]))
datT6 <- datT6 +matrix(rep(1:nrow(datT6),ncol(datT6)),ncol=ncol(datT6))
datT6[6:7,c(1,3,6)] <- NA
datT6[which(datT6 < 11 & datT6 > 10.5)] <- NA
datT6[which(datT6 < 6 & datT6 > 5)] <- NA
datT6[which(datT6 < 4.6 & datT6 > 4)] <- NA
matrixNAinspect(datT6,gr=gl(2,3))
```

matrixNAneighbourImpute

Imputation of NA-values based on non-NA replicates

Description

It is assumed that NA-values appear in data when quantitation values are very low (as this appears eg in quantitative shotgun proteomics). Here, the concept of (technical) replicates is used to investigate what kind of values appear in the other replicates next to NA-values for the same line/protein. Groups of replicate samples are defined via argument `gr` which describes the columns of `dat`. Then, they are inspected for each line to gather NA-neighbour values (ie those values where NAs and regular measures are observed the same time). Eg, let's consider a line contains a set of 4 replicates. Now, if 2 of them are NA-values, the remaining 2 non-NA-values will be considered as NA-neighbours. Ultimately, the aim is to replace all NA-values based on values from a normal distribution resembling their respective NA-neighbours.

Usage

```
matrixNAneighbourImpute(
  dat,
  gr,
  imputMethod = "mode2",
  retnNA = TRUE,
  avSdH = c(NA, 0.5),
  NAneigLst = NULL,
  plotHist = c("hist", "mode"),
  xLab = NULL,
  xLim = NULL,
  yLab = NULL,
```



```

yLim = NULL,
tit = NULL,
figImputDetail = TRUE,
seedNo = NULL,
silent = FALSE,
callFrom = NULL
)

```

Arguments

dat	(matrix or data.frame) main data (may contain NA)
gr	(character or factor) grouping of columns of 'dat', replicate association
imputMethod	(character) choose the imputation method (may be 'mode2'(default), 'mode1', 'datQuant' or 'modeAdopt')
retnNA	(logical) decide (if =TRUE) only NA-substuted data should be returned, or if list with \$data, \$nNA, \$NAneighbour and \$randParam should be returned
avSdH	(numerical,length=2) population characteristics 'high' (mean and sd) for >1 NA-neighbours (per line)
NAneigLst	(list) option for repeated rounds of imputations: list of NA-neighbour values can be furnished for slightly faster processing
plotHist	(character or logical) decide if supplemental figure with histogram should be drawn, the details 'Hist','quant' (display quantile of originak data), 'mode' (display mode of original data) can be chosen explicately
xLab	(character) label on x-axis on plot
xLim	(numeric, length=2) custom x-axis limits
yLab	(character) label on y-axis on plot
yLim	(numeric, length=2) custom y-axis limits
tit	(character) title on plot
figImputDetail	(logical) display details about data (number of NAs) and imputation in graph (min number of NA-neighbours per protein and group, quantile to model, mean and sd of imputed)
seedNo	(integer) seed-value for normal random values
silent	(logical) suppress messages
callFrom	(character) allow easier tracking of message(s) produced

Details

By default a histogram gets plotted showing the initial, imputed and final distribution to check the global hypothesis that NA-values arose from very low measurements and to appreciate the impact of the imputed values to the overall final distribution.

There are a number of experimental settings where low measurements may be reported as NA. Sometimes an arbitrary defined baseline (as 'zero') may provoke those values found below being unfortunately reported as NA or as 0 (in case of MaxQuant). In quantitative proteomics (DDA-mode) the presence of numerous high-abundance peptides will lead to the fact that a number of less intense

MS-peaks don't get identified properly and will then be reported as NA in the respective samples, while the same peptides may be correctly identified and quantified in other (replicate) samples. So, if a given protein/peptide gets properly quantified in some replicate samples but reported as NA in other replicate samples one may thus speculate that similar values like in the successful quantifications may have occurred. Thus, imputation of NA-values may be done on the basis of NA-neighbours.

There are

When extracting NA-neighbours, a slightly more focussed approach gets checked, too, the 2-NA-neighbours : In case a set of replicates for a given protein contains at least 2 non-NA-values (instead of just one) it will be considered as a (min) 2-NA-neighbour as well as regular NA-neighbour. If >300 of these (min) 2-NA-neighbours get found, they will be used instead of the regular NA-neighbours. For creating a collection of normal random values one may use directly the mode of the NA-neighbours (or 2-NA-neighbours, if >300 such values available). To do so, the first value of argument `avSdH` must be set to NA. Otherwise, the first value `avSdH` will be used as quantile of all data to define the mean for the imputed data (ie as `quantile(dat, avSdH[1], na.rm=TRUE)`). The `sd` for generating normal random values will be taken from the `sd` of all NA-neighbours (or 2-NA-neighbours) multiplied by the second value in argument `avSdH` (or `avSdH`, if >300 2-NA-neighbours), since the `sd` of the NA-neighbours is usually quite high. In extremely rare cases it may happen that no NA-neighbours are found (ie if NAs occur, all replicates are NA). Then, this function replaces NA-values based on the normal random values obtained as described above.

Value

list with `$data` .. matrix of data where NA are replaced by imputed values, `$nNA` .. number of NA by group, `$randParam` .. parameters used for making random data

See Also

this function gets used by [testRobustToNAimputation](#); stimation of mode [stableMode](#); detection of NAs [na.fail](#)

Examples

```
set.seed(2013)
datT6 <- matrix(round(rnorm(300)+3,1),ncol=6, dimnames=list(paste("li",1:50,sep=""),
  letters[19:24]))
datT6 <- datT6 +matrix(rep(1:nrow(datT6), ncol(datT6)), ncol=ncol(datT6))
datT6[6:7, c(1,3,6)] <- NA
datT6[which(datT6 < 11 & datT6 > 10.5)] <- NA
datT6[which(datT6 < 6 & datT6 > 5)] <- NA
datT6[which(datT6 < 4.6 & datT6 > 4)] <- NA
datT6b <- matrixNAneighbourImpute(datT6, gr=gl(2,3))
head(datT6b$data)
```

Description

plotROC plots ROC curves based on results from [summarizeForROC](#). This function plots only, it does not return any data. It allows printing simultaneously multiple ROC curves from different studies, it is also compatible with data from 3 species mix as in proteomics benchmark. Input can be prepared using [moderTest2grp](#) followed by [summarizeForROC](#).

Usage

```
plotROC(
  dat,
  ...,
  useColumn = 2:3,
  methNames = NULL,
  col = NULL,
  pch = 1,
  bg = NULL,
  tit = NULL,
  xlim = NULL,
  ylim = NULL,
  point05 = 0.05,
  pointSi = 0.85,
  nByMeth = NULL,
  speciesOrder = NULL,
  txtLoc = NULL,
  legCex = 0.72,
  las = 1,
  addSuplT = TRUE,
  silent = FALSE,
  callFrom = NULL
)
```

Arguments

dat	(matrix) from testing (eg summarizeForROC)
...	optional additional data-sets to include as seprate ROC-curves to same plot (must be of same type of format as 'dat')
useColumn	(integer or character, length=2) columns from dat to be used for pecificity and sensitivity
methNames	(character) names of methods (data-sets) to be displayed
col	(character) custom colors for lines and text (choose one color for each different data-set)
pch	(integer) type of symbol to be used (see also par)
bg	(character) background color in plot (see also par)
tit	(character) custom title
xlim	(numeric, length=2) custom x-axis limits
ylim	(numeric, length=2) custom y-axis limits

point05	(numeric) specific point to highlight in plot (typically at alpha=0.05)
pointSi	(numeric) size of points (as expansion factor cex)
nByMeth	(integer) value of n to display
speciesOrder	(integer) custom order of species in legend
txtLoc	(numeric, length=3) location for text (x, y and proportional factor for line-offset, default is c(0.4,0.3,0.04))
legCex	(numeric) cex expansion factor for legend (see also par)
las	(numeric) factor for text-orientation (see also par)
addSuplT	(logical) add text with information about precision, accuracy and FDR
silent	(logical) suppress messages
callFrom	(character) allows easier tracking of messages produced

Value

plot with ROC curves only

See Also

[summarizeForROC](#), [moderTest2grp](#)

Examples

```
roc0 <- cbind(alph=c(2e-6, 4e-5, 4e-4, 2.7e-3, 1.6e-2, 4.2e-2, 8.3e-2, 1.7e-1, 2.7e-1, 4.1e-1, 5.3e-1,
  6.8e-1, 8.3e-1, 9.7e-1), spec=c(1, 1, 1, 1, 0.957, 0.915, 0.915, 0.809, 0.702, 0.489, 0.362, 0.234,
  0.128, 0.0426), sens=c(0, 0, 0.145, 0.942, 2.54, 2.68, 3.33, 3.99, 4.71, 5.87, 6.67, 8.04, 8.77,
  9.93)/10, n.pos.a=c(0, 0, 0, 0, 2, 4, 4, 9, 14, 24, 36, 41) )
plotROC(roc0)
```

razorNoFilter	<i>Filter based on either number of total peptides and specific peptides or number of razor peptides</i>
---------------	--

Description

razorNoFilter filters based on either a) number of total peptides and specific peptides or b) number of razor peptides. This function was designed for filtering using a minimum number of (PSM-) count values following the common practice to consider results with 2 or more peptide counts as reliable. The function be (re-)run independently on each of various questions (comparisons). Note: Non-integer data will be truncated to integer (equivalent to floor).

Usage

```
razorNoFilter(
  annot,
  speNa = NULL,
  totNa = NULL,
  minRazNa = NULL,
  minSpeNo = 1,
  minTotNo = 2,
  silent = FALSE,
  callFrom = NULL
)
```

Arguments

annot	(matrix or data.frame) main data (may contain NAs) with (PSM-) count values for each protein
speNa	(integer or character) indicate which column of 'annot' has number of specific peptides
totNa	(integer or character) indicate which column of 'annot' has number of total peptides
minRazNa	(integer or character) name of column with number of razor peptides, alternative to 'minSpeNo' & 'minTotNo'
minSpeNo	(integer) minimum number of specific peptides
minTotNo	(integer) minimum total ie max razor number of peptides
silent	(logical) suppress messages
callFrom	(character) allows easier tracking of messages produced

Value

vector of logical values if corresponding line passes filter criteria

See Also

[presenceFilt](#)

Examples

```
set.seed(2019); datT <- matrix(sample.int(20,60,replace=TRUE),ncol=6,
  dimnames=list(letters[1:10],LETTERS[1:6])) -3
datT[,2] <- datT[,2] +2
datT[which(datT <0)] <- 0
razorNoFilter(datT,speNa="A",totNa="B")
```

readFasta2	<i>Read file of protein sequences in fasta format Read fasta formatted file (from Rhrefttps:/www.uniprot.orgUniProt) to extract (protein) sequences and name. If tableOut=TRUE output may be organized as matrix for separating meta-annotation (eg GeneName, OrganismName, ProteinName) in separate columns.</i>
------------	---

Description

Read file of protein sequences in fasta format

Read fasta formatted file (from [UniProt](#)) to extract (protein) sequences and name. If tableOut=TRUE output may be organized as matrix for separating meta-annotation (eg GeneName, OrganismName, ProteinName) in separate columns.

Usage

```
readFasta2(
  filename,
  delim = "|",
  databaseSign = c("sp", "tr", "generic", "gi"),
  tableOut = FALSE,
  UniprSep = c("OS=", "OX=", "GN=", "PE=", "SV="),
  cleanCols = TRUE,
  silent = FALSE,
  callFrom = NULL,
  debug = FALSE
)
```

Arguments

filename	(character) names fasta-file to be read
delim	(character) delimiter at header-line
databaseSign	(character) characters at beginning right after the '>' (typically specifying the data-base-origin), they will be excluded from the sequence-header
tableOut	(logical) toggle to return named character-vector or matrix with enhanced parsing of fasta-header. The resulting matrix will contain the columns 'database', 'uniqueIdentifier', 'entryName', and further columns depending on argument UniprSep
UniprSep	(character) separators for further separating entry-fields if tableOut=TRUE, see also UniProt-FASTA-headers
cleanCols	(logical) remove columns with all entries NA, if tableOut=TRUE
silent	(logical) suppress messages
callFrom	(character) allows easier tracking of message(s) produced
debug	(logical) supplemental messages for debugging

Value

return (based on 'tableOut') simple character vector (of sequence) with Uniprot ID as name or matrix with columns: 'database', 'uniqueIdentifier', 'entryName', 'proteinName', 'sequence' and further columns depending on argument UniprSep

See Also

[scan](#) or `read.fasta` from the package [seqinr](#)

Examples

```
# tiny example with common contaminants
path1 <- system.file('extdata', package='wrProteo')
fiNa <- "conta1.fasta"
fasta1 <- readFasta2(file.path(path1, fiNa))
## now let's read and further separate annotation-fields
fasta2 <- readFasta2(file.path(path1, fiNa), tableOut=TRUE)
str(fasta1)
```

readMaxQuantFile

Read proteinGroups.txt files exported from MaxQuant

Description

Quantification results from **MaxQuant** can be read using this function and relevant information extracted. Input files compressed as .gz can be read as well. Besides protein abundance values (XIC) peptide counting information like number of unique razor-peptides or PSM values can be extracted, too. The protein abundance values may be normalized using multiple methods (median normalization is default), the determination of normalization values can be restricted to specific proteins (normalization to bait protein(s), or to matrix in UPS1 spike-in experiments). Besides, a graphical display of the distribution of protein abundance values may be generated. The final output is a list containing these elements: \$raw, \$quant, \$annot, \$counts, \$quantNotes, \$notes, or (if separateAnnot=FALSE) data.frame with annotation- and main quantification-content.

Usage

```
readMaxQuantFile(
  path,
  fileName = "proteinGroups.txt",
  normalizeMeth = "median",
  quantCol = "LFQ.intensity",
  contamCol = "Potential.contaminant",
  pepCountCol = c("Razor...unique.peptides.", "MS.MS.count."),
  uniqPepPat = NULL,
  refLi = NULL,
  extrColNames = c("Majority.protein.IDs", "Fasta.headers", "Number.of.proteins"),
  specPref = c(conta = "conta|CON_|LYSC_CHICK", mainSpecies = "OS=Homo sapiens"),
```

```

remRev = TRUE,
separateAnnot = TRUE,
tit = NULL,
wex = 1.6,
plotGraph = TRUE,
silent = FALSE,
callFrom = NULL
)

```

Arguments

path	(character) path of file to be read
fileName	(character) name of file to be read (default 'proteinGroups.txt' as typically generated by MaxQuant in txt folder). Gz-compressed files can be read, too.
normalizeMeth	(character) normalization method (for details see normalizeThis)
quantCol	(character or integer) exact col-names, or if length=1 content of quantCol will be used as pattern to search among column-names for \$quant using grep
contamCol	(character or integer, length=1) which columns should be used for contaminants marked by ProteomeDiscoverer
pepCountCol	(character) pattern to search among column-names for count data of PSM and NoOfPeptides
uniqPepPat	(character, length=1) deprecated, please use pepCountCol instead
refLi	(character or integer) custom specify which line of data is main species, if character (eg 'mainSpe'), the column 'SpecType' in \$annot will be searched for exact match of the (single) term given
extrColNames	(character) column names to be read (1: prefix for LFQ quantitation, default 'LFQ.intensity'; 2: column name for protein-IDs, default 'Majority.protein.IDs'; 3: column names of fasta-headers, default 'Fasta.headers', 4: column name for number of protein IDs matching, default 'Number.of.proteins')
specPref	(character) prefix to identifiers allowing to separate i) recognize contamination database, ii) species of main identifications and iii) spike-in species
remRev	(logical) option to remove all protein-identifications based on reverse-peptides
separateAnnot	(logical) if TRUE output will be organized as list with \$annot, \$abund for initial/raw abundance values and \$quant with final normalized quantitations
tit	(character) custom title to plot
wex	(numeric) relative expansion factor of the violin in plot
plotGraph	(logical) optional plot vioplot of initial and normalized data (using normalizeMeth); alternatively the argument may contain numeric details that will be passed to layout when plotting
silent	(logical) suppress messages
callFrom	(character) allow easier tracking of message produced

Details

This function has been developed using MaxQuant versions 1.6.10.x to 1.6.17.x, the format of resulting file 'proteinGroups.txt' is typically well conserved.

Value

list with \$raw (initial/raw abundance values), \$quant with final normalized quantitations, \$annot (columns), \$counts an array with 'PSM' and 'NoOfRazorPeptides', \$quantNotes and \$notes; or a data.frame with quantitation and annotation if separateAnnot=FALSE

See Also

[read.table](#), [normalizeThis](#)), [readProlineFile](#)

Examples

```
path1 <- system.file("extdata", package="wrProteo")
# Here we'll load a short/trimmed example file (thus not MaxQuant default name)
fiNa <- "proteinGroupsMaxQuant1.txt.gz"
specPr <- c(conta="conta|CON_|LYSC_CHICK", mainSpecies="YEAST",spike="HUMAN_UPS")
dataMQ <- readMaxQuantFile(path1, file=fiNa, specPref=specPr, tit="tiny MaxQuant")
summary(dataMQ$quant)
matrixNAinspect(dataMQ$quant, gr=gl(3,3))
```

readOpenMSFile

Read csv files exported by OpenMS

Description

Protein quantification results form **OpenMS** which were exported as .csv can be imported and relevant information extracted. Peptide data get summarized by protein by top3 or sum methods. The final output is a list containing the elements: \$annot, \$raw, \$quant ie normaized final quantifications, or returns data.frame with entire content of file if separateAnnot=FALSE.

Usage

```
readOpenMSFile(
  fileName = NULL,
  path = NULL,
  normalizeMeth = "median",
  refLi = NULL,
  sampleNames = NULL,
  quantCol = "Intensity",
  sumMeth = "top3",
  minPepNo = 1,
  protNaCol = "ProteinName",
  separateAnnot = TRUE,
  plotGraph = TRUE,
  tit = "OpenMS",
  wex = 1.6,
  specPref = c(conta = "LYSC_CHICK", mainSpecies = "OS=Homo sapiens"),
  silent = FALSE,
  callFrom = NULL
)
```

Arguments

fileName	(character) name of file to be read
path	(character) path of file to be read
normalizeMeth	(character) normalization method (will be sent to normalizeThis)
refLi	(character or integer) custom specify which line of data is main species, if character (eg 'mainSpe'), the column 'SpecType' in \$annot will be searched for exact match of the (single) term given
sampleNames	(character) new column-names for quantification data (by default the names from files with spectra will be used)
quantCol	(character or integer) exact col-names, or if length=1 content of quantCol will be used as pattern to search among column-names for \$quant using grep
sumMeth	(character) method for summarizing peptide data (so far 'top3' and 'sum' available)
minPepNo	(integer) minimum number of peptides to be used for retraining quantification
protNaCol	(character) column name to be read/extracted for the annotation section (default "ProteinName")
separateAnnot	(logical) if TRUE output will be organized as list with \$annot, \$abund for initial/raw abundance values and \$quant with final normalized quantitations
plotGraph	(logical) optional plot of type vioplot of initial and normalized data (using normalizeMeth); if integer, it will be passed to layout when plotting
tit	(character) custom title to plot
wex	(integer) relative expansion factor of the violin-plot (will be passed to vioplotW)
specPref	(character or list) define characteristic text for recognizing (main) groups of species (1st for contaminants - will be marked as 'conta', 2nd for main species - marked as 'mainSpe', and optional following ones for supplemental tags/species - marked as 'species2', 'species3', ...); if list and list-element has multiple values they will be used for exact matching of accessions (ie 2nd of argument annotCol)
silent	(logical) suppress messages
callFrom	(character) allow easier tracking of message(s) produced

Details

This function has been developed based on the OpenMS peptide-identification and label-free-quantification module. Csv input files may also be compressed as .gz.

Note: With this version the information about protein-modifications (PTMs) may not yet get exploited fully.

Value

list with \$raw (initial/raw abundance values), \$quant with final normalized quantitations, \$annot, \$counts an array with number of peptides, \$quantNotes, \$expSetup and \$notes; or if separateAnnot=FALSE the function returns a data.frame with annotation and quantitation only

See Also

[read.table](#), [normalizeThis](#)), [readMaxQuantFile](#), [readProlineFile](#), [readProtDiscovFile](#)

Examples

```
path1 <- system.file("extdata", package="wrProteo")
fiNa <- "OpenMS_tiny.csv.gz"
dataOM <- readOpenMSFile(file=fiNa, path=path1, tit="tiny OpenMS example")
summary(dataOM$quant)
```

readPDExport

Read tabulated files imported from Thermo ProteomeDiscoverer

Description

This function has been deprecated and replaced by [readProtDiscovFile](#). Quantification results from **Thermo ProteomeDiscoverer** exported as tabulated text can be imported and relevant information extracted. The final output is a list containing 3 elements: \$annot, \$raw and optional \$quant, or returns data.frame with entire content of file if separateAnnot=FALSE.

Usage

```
readPDExport(
  fileName,
  path = NULL,
  normalizeMeth = "median",
  sampleNames = NULL,
  read0asNA = TRUE,
  quantCol = "^Abundances*",
  annotCol = NULL,
  contamCol = "Contaminant",
  refLi = NULL,
  separateAnnot = TRUE,
  plotGraph = TRUE,
  tit = "Proteome Discoverer",
  graphTit = NULL,
  wex = 1.6,
  specPref = c(conta = "CON_|LYSC_CHICK", mainSpecies = "OS=Homo sapiens"),
  silent = FALSE,
  callFrom = NULL
)
```

Arguments

`fileName` (character) name of file to be read (default 'proteinGroups.txt' as typically generated by MaxQuant in txt folder)

path	(character) path of file to be read
normalizeMeth	(character) normalization method (will be sent to normalizeThis)
sampleNames	(character) new column-names for quantification data (ProteomeDiscoverer does not automatically use file-names from spectra)
read0asNA	(logical) decide if initial quantifications at 0 should be transformed to NA
quantCol	(character or integer) exact col-names, or if length=1 content of quantCol will be used as pattern to search among column-names for \$quant using grep
annotCol	(character) column names to be read/extracted for the annotation section (default c("Accession", "Description", "Gene", "Contaminant", "Sum.PEP.Score", "Coverage....", "X..Peptides", "X..I", "X..AAs", "MW..kDa."))
contamCol	(character or integer, length=1) which columns should be used for contaminants marked by ProteomeDiscoverer
refLi	(character or integer) custom specify which line of data is main species, if character (eg 'mainSpe'), the column 'SpecType' in \$annot will be searched for exact match of the (single) term given
separateAnnot	(logical) if TRUE output will be organized as list with \$annot, \$abund for initial/raw abundance values and \$quant with final normalized quantitations
plotGraph	(logical) optional plot of type vioplot of initial and normalized data (using normalizeMeth); if integer, it will be passed to layout when plotting
tit	(character) custom title to plot
graphTit	(character) deprecated custom title to plot, please use 'tit'
wex	(integer) relative expansion factor of the violin-plot (will be passed to vioplotW)
specPref	(character or list) define characteristic text for recognizing (main) groups of species (1st for contaminants - will be marked as 'conta', 2nd for main species-marked as 'mainSpe', and optional following ones for supplemental tags/species - maked as 'species2','species3',...); if list and list-element has multiple values they will be used for exact matching of accessions (ie 2nd of argument annotCol)
silent	(logical) suppress messages
callFrom	(character) allow easier tracking of message(s) produced

Details

This function has been developed using Thermo ProteomeDiscoverer versions 2.2 to 2.4. The format of resulting files at export also depends which columns are chosen as visible inside ProteomeDiscoverer and subsequently get chosen for export. Please make sure that 'RawAbundance' is chosen, too.

Value

list with \$raw (initial/raw abundance values), \$quant with final normalized quantitations, \$annot, \$counts an array with number of peptides, \$quantNotes and \$notes; or if separateAnnot=FALSE the function returns a data.frame with annotation and quantitation only

See Also

[read.table](#), [normalizeThis](#)), [readProlineFile](#)

Examples

```
path1 <- system.file("extdata", package="wrProteo")
fiNa <- "tinyPD_allProteins.txt.gz"
## Function is deprecated. Please use readProtDiscovFile() instead of readPDExport() !
dataPD <- readProtDiscovFile(file=fiNa, path=path1)
```

readProlineFile *Read csv or txt files exported from Proline and MS-Angel*

Description

Quantification results form MS-Angel and Proline **Proline** exported as xlsx format can be read directly. Besides, files in tsv, csv (European and US format) or tabulated txt can be read, too. Then relevant information gets extracted, the data can optionally normalized and displayed as box-plot or vioplot. The final output is a list containing 6 elements: \$raw, \$quant, \$annot, \$counts, \$quantNotes and \$notes. Alternatively, a dataframe with annotation and quantitation data may be returned if separateAnnot=FALSE. Note: There is no normalization by default since quite frequently data produced by Proline are already sufficiently normalized. The figure produced using the argument plotGraph=TRUE may help judging if the data appear sufficiently normalized (distributions should align).

Usage

```
readProlineFile(
  fileName,
  path = NULL,
  normalizeMeth = NULL,
  logConvert = TRUE,
  sampleNames = NULL,
  quantCol = "^abundance_",
  annotCol = c("accession", "description", "is_validated", "protein_set_score",
    "X.peptides", "X.specific_peptides"),
  remStrainNo = TRUE,
  pepCountCol = c("^psm_count_", "^peptides_count_"),
  trimColnames = FALSE,
  refLi = NULL,
  separateAnnot = TRUE,
  plotGraph = TRUE,
  tit = NULL,
  graphTit = NULL,
  wex = 2,
  specPref = c(conta = "_conta\\|", mainSpecies = "OS=Homo sapiens"),
```

```

    silent = FALSE,
    callFrom = NULL
)

```

Arguments

fileName	(character) name of file to read; .xlsx-, .csv-, .txt- and .tsv can be read (csv, txt and tsv may be gz-compressed). Reading xlsx requires package 'readxl'.
path	(character) optional path (note: Windows backslash should be protected or written as '/')
normalizeMeth	(character) normalization method (for details and options see normalizeThis)
logConvert	(logical) convert numeric data as log2, will be placed in \$quant
sampleNames	(character) new column-names for quantification data (ProteomeDiscoverer does not automatically use file-names from spectra); Please use with care since order of samples might be different as you expect
quantCol	(character or integer) columns with main quantitation-data : precise colnames to extract, or if length=1 content of quantCol will be used as pattern to search among column-names for \$quant using grep
annotCol	(character) precise colnames or if length=1 pattern to search among column-names for \$annot
remStrainNo	(logical) if TRUE, the organism annotation will be trimmed to uppercaseWord+space+lowercaseWord (eg Homo sapiens)
pepCountCol	(character) pattern to search among column-names for count data of PSM and NoOfPeptides
trimColnames	(logical) optional trimming of column-names of any redundant characters from beginning and end
refLi	(integer) custom decide which line of data is main species, if single character entry it will be used to choose a group of species (eg 'mainSpe')
separateAnnot	(logical) separate annotation form numeric data (quantCol and annotCol must be defined)
plotGraph	(logical or matrix of integer) optional plot vioplot of initial data; if integer, it will be passed to layout when plotting
tit	(character) custom title to plot
graphTit	(character) (deprecated custom title to plot), please use 'tit'
wex	(integer) relative expansion factor of the violin-plot (will be passed to vioplotW)
specPref	(character or list) define characteristic text for recognizing (main) groups of species (1st for contaminants - will be marked as 'conta', 2nd for main species-marked as 'mainSpe', and optional following ones for supplemental tags/species - maked as 'species2', 'species3',...); if list and list-element has multiple values they will be used for exact matching of accessions (ie 2nd of argument annotCol)
silent	(logical) suppress messages
callFrom	(character) allow easier tracking of message(s) produced

Details

This function has been developed using Proline version 1.6.1 coupled with MS-Angel 1.6.1. The format of the exported file depends on the columns chosen for export, default settings from Proline and MS-Angel work fine.

Value

list with \$raw (initial/raw abundance values), \$quant with final normalized quantitations, \$annot (columns), \$counts an array with 'PSM' and 'NoOfPeptides', \$quantNotes and \$notes; or a data.frame with quantitation and annotation if separateAnnot=FALSE

See Also

[read.table](#)

Examples

```
path1 <- system.file("extdata", package="wrProteo")
fiNa <- "exampleProlineABC.csv.gz"
dataABC <- readProlineFile(file.path(path1, fiNa))
summary(dataABC$quant)
```

readProtDiscovFile *Read tabulated files exported by ProteomeDiscoverer*

Description

Protein quantification results form **Thermo ProteomeDiscoverer** which were exported as tabulated text can be imported and relevant information extracted. The final output is a list containing 3 elements: \$annot, \$raw and optional \$quant, or returns data.frame with entire content of file if separateAnnot=FALSE.

Usage

```
readProtDiscovFile(
  fileName,
  path = NULL,
  normalizeMeth = "median",
  sampleNames = NULL,
  read0asNA = TRUE,
  quantCol = "^Abundances*",
  annotCol = NULL,
  contamCol = "Contaminant",
  refLi = NULL,
  separateAnnot = TRUE,
  plotGraph = TRUE,
  tit = "Proteome Discoverer",
```

```

graphTit = NULL,
wex = 1.6,
specPref = c(conta = "CON_|LYSC_CHICK", mainSpecies = "OS=Homo sapiens"),
silent = FALSE,
callFrom = NULL
)

```

Arguments

fileName	(character) name of file to be read
path	(character) path of file to be read
normalizeMeth	(character) normalization method (will be sent to normalizeThis)
sampleNames	(character) new column-names for quantification data (ProteomeDiscoverer does not automatically use file-names from spectra)
read@asNA	(logical) decide if initial quantifications at 0 should be transformed to NA
quantCol	(character or integer) exact col-names, or if length=1 content of quantCol will be used as pattern to search among column-names for \$quant using grep
annotCol	(character) column names to be read/extracted for the annotation section (default c("Accession","Description","Gene","Contaminant","Sum.PEP.Score","Coverage....","X..Peptides","X..I","X..AAs","MW..kDa."))
contamCol	(character or integer, length=1) which columns should be used for contaminants marked by ProteomeDiscoverer
refLi	(character or integer) custom specify which line of data is main species, if character (eg 'mainSpe'), the column 'SpecType' in \$annot will be searched for exact match of the (single) term given
separateAnnot	(logical) if TRUE output will be organized as list with \$annot, \$abund for initial/raw abundance values and \$quant with final normalized quantitations
plotGraph	(logical) optional plot of type vioplot of initial and normalized data (using normalizeMeth); if integer, it will be passed to layout when plotting
tit	(character) custom title to plot
graphTit	(character) depreciated custom title to plot, please use 'tit'
wex	(integer) relative expansion factor of the violin-plot (will be passed to vioplotW)
specPref	(character or list) define characteristic text for recognizing (main) groups of species (1st for contaminants - will be marked as 'conta', 2nd for main species - marked as 'mainSpe', and optional following ones for supplemental tags/species - maked as 'species2','species3',...); if list and list-element has multiple values they will be used for exact matching of accessions (ie 2nd of argument annotCol)
silent	(logical) suppress messages
callFrom	(character) allow easier tracking of message(s) produced

Details

This function has been developed using Thermo ProteomeDiscoverer versions 2.2 to 2.5. The format of resulting files at export also depends which columns are chosen as visible inside ProteomeDiscoverer and subsequently get chosen for export. Please make sure that 'RawAbundance' is chosen, too. This function replaces the depreciated function readPDExport.

Value

list with \$raw (initial/raw abundance values), \$quant with final normalized quantitations, \$annot, \$counts an array with number of peptides, \$quantNotes and \$notes; or if separateAnnot=FALSE the function returns a data.frame with annotation and quantitation only

See Also

[read.table, normalizeThis](#)), [readMaxQuantFile](#), [readProlineFile](#)

Examples

```
path1 <- system.file("extdata", package="wrProteo")
fiNa <- "tinyPD_allProteins.txt.gz"
dataPD <- readProtDiscovFile(file=fiNa, path=path1)
summary(dataPD$quant)
```

readUCSCTable

Read annotation files from UCSC

Description

This function allows reading and importing genomic **UCSC-annotation** data. Files can be read as default UCSC exprot or as GTF-format. In the context of proteomics we noticed that sometimes UniProt tables from UCSC are hard to match to identifiers from UniProt Fasta-files, ie many protein-identifiers won't match. For this reason additional support is given to reading 'Genes and Gene Predictions': Since this table does not include protein-identifiers, a non-redundant list of ENSxxx transcript identifiers can be exprted as file for an additional stop of conversion, eg using a batch conversion tool at the site of **UniProt**. The initial genomic annotation can then be complemented using [readUniProtExport](#). Using this more elaborate route, we found higher coverage when trying to add genomic annotation to protein-identifiers to proteomics results with annotation based on an initial Fasta-file.

Usage

```
readUCSCTable(
  fiName,
  exportFileNa = NULL,
  gtf = NA,
  simplifyCols = c("gene_id", "chr", "start", "end", "strand", "frame"),
```

```

    silent = FALSE,
    callFrom = NULL
  )

```

Arguments

<code>fileName</code>	(character) name (and path) of file to read
<code>exportFileName</code>	(character) optional file-name to be exported, if NULL no file will be written
<code>gtf</code>	(logical) specify if file <code>fileName</code> in gtf-format (see UCSC)
<code>simplifyCols</code>	(character) optional list of column-names to be used for simplification (if 6 column-headers are given) : the 1st value will be used to identify the column used as reference to summarize all lines with this ID; for the 2nd (typically chromosome names) will be taken a representative value, for the 3rd (typically gene start site) will be taken the minimum, for the 4th (typically gene end site) will be taken the maximum, for the 5th and 6th a representative values will be reported;
<code>silent</code>	(logical) suppress messages
<code>callFrom</code>	(character) allows easier tracking of message(s) produced

Value

matrix, optionally the file 'exportFileName' may be written

See Also

[readUniProtExport](#), [readPDEExport](#), [readMaxQuantFile](#),

Examples

```

path1 <- system.file("extdata", package="wrProteo")
gtfFi <- file.path(path1, "UCSC_hg38_chr11extr.gtf.gz")
# here we'll write the file for UniProt conversion to tempdir() to keep things tidy
expFi <- file.path(tempdir(), "deUcscForUniProt2.txt")
UcscAnnot1 <- readUCSCtable(gtfFi, exportFileName=expFi)

## results can be further combined with readUniProtExport()
deUniProtFi <- file.path(path1, "deUniProt_hg38chr11extr.tab")
deUniPr1 <- readUniProtExport(deUniProtFi, deUcsc=UcscAnnot1,
  targRegion="chr11:1-135,086,622")
deUniPr1[1:5,-5]

```

readUniProtExport *Read protein annotation as exported from UniProt batch-conversion*

Description

This function allows reading and importing protein-ID conversion results from **UniProt**. To do so, first copy/paste your query IDs into **UniProt** 'Retrieve/ID mapping' field called '1. Provide your identifiers' (or upload as file), verify '2. Select options'. In a typical case of 'enst000xxx' IDs you may leave default settings, ie 'Ensemble Transcript' as input and 'UniProt KB' as output. Then, 'Submit' your search and retrieve results via 'Download', you need to specify a 'Tab-separated' format ! If you download as 'Compressed' you need to decompress the .gz file before running the function readUCSCTable In addition, a file with UCSC annotation (Ensnrnt accessions and chromosomic locations, obtained using [readUCSCTable](#)) can be integrated.

Usage

```
readUniProtExport(
  UniProtFileNa,
  deUcsc = NULL,
  targRegion = NULL,
  useUniPrCol = NULL,
  silent = FALSE,
  callFrom = NULL
)
```

Arguments

UniProtFileNa	(character) name (and path) of file exported from Uniprot (tabulated text file including headers)
deUcsc	(data.frame) object produced by readUCSCTable to be combined with data from UniProtFileNa
targRegion	(character or list) optional marking of chromosomal locations to be part of a given chromosomal target region, may be given as character like chr11:1-135,086,622 or as list with a first component characterizing the chromosome and a integer-vector with start- and end- sites
useUniPrCol	(character) optional declaration which colums from UniProt exported file should be used/imported (default 'EnsID','Entry','Entry.name','Status','Protein.names','Gene.names','Length')
silent	(logical) suppress messages
callFrom	(character) allows easier tracking of message(s) produced

Details

In a typical use case, first chromosomic location annotation is extracted from UCSC for the species of interest and imported to R using [readUCSCTable](#) . However, the tables provided by UCSC don't contain Uniprot IDs. Thus, an additional (batch-)conversion step needs to get added. For this reason [readUCSCTable](#) allows writing a file with Ensemble transcript IDs which can be converted

tu UniProt IDs at the site of [UniProt](#). Then, UniProt annotation (downloaded as tab-separated) can be imported and combined with the genomic annotation using this function.

Value

data.frame (with columns \$EnsID, \$Entry, \$Entry.name, \$Status, \$Protein.names, \$Gene.names, \$Length; if deUcsc is integrated plus: \$chr, \$type, \$start, \$end, \$score, \$strand, \$Ensnot, \$avPos)

See Also

[readUCSCTable](#)

Examples

```
path1 <- system.file("extdata",package="wrProteo")
deUniProtFi <- file.path(path1,"deUniProt_hg38chr11extr.tab")
deUniPr1a <- readUniProtExport(deUniProtFi)
str(deUniPr1a)

## Workflow starting with UCSC annotation (gtf) files :
gtfFi <- file.path(path1,"UCSC_hg38_chr11extr.gtf.gz")
UcscAnnot1 <- readUCSCTable(gtfFi)
## Results of conversion at UniProt are already available (file "deUniProt_hg38chr11extr.tab")
myTargRegion <- list("chr1", pos=c(198110001,198570000))
myTargRegion2 <- "chr11:1-135,086,622" # works equally well
deUniPr1 <- readUniProtExport(deUniProtFi,deUcsc=UcscAnnot1,
  targRegion=myTargRegion)
## Now UniProt IDs and genomic locations are both available :
str(deUniPr1)
```

removeSampleInList	<i>Remove samples/columns from list of matrixes Remove samples (ie columns) from every instance of list of matrixes. Note: This function assumes same order of columns in list-elements 'listElem' !</i>
--------------------	--

Description

Remove samples/columns from list of matrixes

Remove samples (ie columns) from every instance of list of matrixes. Note: This function assumes same order of columns in list-elements 'listElem' !

Usage

```
removeSampleInList(
  dat,
  remSamp,
  listElem = c("abund", "quant"),
  silent = FALSE,
  callFrom = NULL
)
```

Arguments

dat	(list) main input to be filtered
remSamp	(integer) column number to exclude
listElem	(character) names of list-elements where columns indicated with 'remSamp' should be removed
silent	(logical) suppress messages
callFrom	(character) allows easier tracking of messages produced

Value

matrix including imputed values or list of final and matrix with number of imputed by group (plus optional plot)

See Also

[testRobustToNAimputation](#)

Examples

```
set.seed(2019)
datT6 <- matrix(round(rnorm(300)+3,1),ncol=6,dimnames=list(paste("li",1:50,sep=""),
  letters[19:24]))
datL <- list(abund=datT6,quant=datT6,annot=matrix(nrow=nrow(datT6),ncol=2))
datDelta2 <- removeSampleInList(datL,remSam=2)
```

replMissingProtNames *Complement missing EntryNames in annotation*

Description

This function helps replacing missing EntryNames (in \$annot) after reading quantification results. To do so the column-names of annCol will be used : The content of 2nd element (and optional 3rd element) will be used to replace missing content in column defined by 1st element.

Usage

```
replMissingProtNames(
  x,
  annCol = c("EntryName", "Accession", "SpecType"),
  silent = FALSE,
  callFrom = NULL
)
```

Arguments

x	(list) output of readMaxQuantFile, readProtDiscovFile or readProlineFile. This list must be a matrix and contain \$annot with the columns designated in annCol.
annCol	(character) the column-names form x\$annot) which will be used : The first column designs the column where empty fields are searched and the 2nd and (optional) 3rd will be used to fill the empty spots in the st column
silent	(logical) suppress messages
callFrom	(character) allow easier tracking of message(s) produced

Value

list as input, but with missing elements of \$annot completed (if available in other columns)

See Also

[readMaxQuantFile](#), [readProtDiscovFile](#), [readProlineFile](#)

Examples

```
dat <- list(quant=matrix(sample(11:99,9,replace=TRUE), ncol=3), annot=cbind(EntryName=c(
  "YP010_YEAST", "", ""), Accession=c("A5Z2X5", "P01966", "P35900"), SpecType=c("Yeast", NA, NA)))
replMissingProtNames(dat)
```

summarizeForROC

Summarize statistical test result for plotting ROC-curves

Description

summarizeForROC takes statistical testing results (obtained using [testRobustToNAimputation](#) or [moderTest2grp](#), based on [limma](#)) and calculates specificity and sensitivity values for plotting ROC-curves along a panel of thresholds. Based on column from test\$annot and argument 'spec' TP,FP,FN and TN are determined. Special consideration is made to 3 species mix samples as found in proteomics benchmark-tests. See also [ROC on Wikipedia](#) for explanations of TP,FP,FN and TN as well as examples. An optional plot may be produced, too. Return matrix with TP,FP,FN,TN,spec,sens,prec,accur and FDR count values along the various thresholds specified in column 'alph'. Note that numerous other packages also provide support for building and plotting ROC-curves : Eg [rocPkgShort](#), [ROCR](#), [pROC](#) or [ROCit](#)

Usage

```
summarizeForROC(
  test,
  thr = NULL,
  tyThr = "BH",
  useComp = 1,
```

```

columnTest = NULL,
FCthrs = NULL,
spec = c("H", "E", "S"),
annotCol = "spec",
tit = NULL,
color = 1,
plotROC = TRUE,
pch = 1,
bg = NULL,
overlPlot = FALSE,
silent = FALSE,
callFrom = NULL
)

```

Arguments

test	(class MArrayLM, S3-object from limma) from testing (eg testRobustToNAimputation or test2grp)
thr	(numeric) p-value threshold, if NULL a panel of 108 p-value threshold-levels values will be used for calculating specificity and sensitivity
tyThr	(character,length=1) type of test-result to be used for sensitivity and specificity calculations (eg 'BH', 'fdr' or 'p.value'), must be list-element of 'test'
useComp	(character or integer) in case multiple comparisons (ie multiple columns 'test\$tyThr'); which pairwise comparison to used
columnTest	deprecated, please use 'useComp' instead
FCthrs	(numeric) Fold-Change threshold (display as line) give as Fold-change and NOT log2(FC), default at 1.5, set to NA for omitting
spec	(character) labels for species will be matched to column 'spec' of test\$annot and used for sensitivity and specificity calculations. Important : 1st label for matrix (expected as constant) and subsequent labels for spike-ins (variable)
annotCol	(character) column name of test\$annot to use to separate species
tit	(character) optional custom title in graph
color	(character or integer) color in graph
plotROC	(logical) toggle plot on or off
pch	(integer) type of symbol to be used (see par)
bg	(character) background in plot (see par)
overlPlot	(logical) overlay to existing plot if TRUE
silent	(logical) suppress messages
callFrom	(character) allows easier tracking of message(s) produced

Value

matrix including imputed values or list of final and matrix with number of imputed by group (plus optional plot)

See Also

replot the figure [plotROC](#), calculate AUC using [AucROC](#), robust test for preparing tables [testRobustToNAimputation](#), [moderTest2grp](#), [test2grp](#), eBayes in package [limma](#), [t.test](#)

Examples

```
set.seed(2019); test1 <- list(annot=cbind(spec=c(rep("b",35),letters[sample.int(n=3,
  size=150,replace=TRUE)])),BH=matrix(c(runif(35,0,0.01),runif(150)),ncol=1))
tail(roc1 <- summarizeForROC(test1,spec=c("a","b","c")))
```

test2grp	<i>t-test each line of 2 groups of data</i>
----------	---

Description

test2grp performs t-test on two groups of data using [limma](#), this is a custom implementation of [moderTest2grp](#) for proteomics. The final object also includes the results without moderation by [limma](#) (eg BH-FDR in `$nonMod.BH`). Furthermore, there is an option to make use of package [ROTS](#) (note, this will increase the time of computations considerably).

Usage

```
test2grp(
  dat,
  questNo,
  useCol = NULL,
  grp = NULL,
  annot = NULL,
  ROTSn = 0,
  silent = FALSE,
  callFrom = NULL
)
```

Arguments

dat	(matrix or data.frame) main data (may contain NAs)
questNo	(integer) specify here which question, ie comparison should be addressed
useCol	(integer or character)
grp	(character or factor)
annot	(matrix or data.frame)
ROTSn	(integer) number of iterations ROTS runs (stabilization of results may be seen with >300)
silent	(logical) suppress messages
callFrom	(character) allow easier tracking of message produced

Value

limma-type S3 object of class 'MArrayLM' which can be accessed; multiple testing correction types or modified testing by ROTS may get included ('p.value', 'FDR', 'BY', 'lfdr' or 'ROTS.BH')

See Also

[moderTest2grp](#), [pVal2lfdr](#), [t.test](#), ROTS from the Bioconductor package [ROTS](#)

Examples

```
set.seed(2018); datT8 <- matrix(round(rnorm(800)+3,1),nc=8,dimnames=list(paste(
  "li",1:100,sep=""),paste(rep(LETTERS[1:3],c(3,3,2)),letters[18:25],sep="")))
datT8[3:6,1:2] <- datT8[3:6,1:2]+3 # augment lines 3:6 (c-f)
datT8[5:8,5:6] <- datT8[5:8,5:6]+3 # augment lines 5:8 (e-h)
grp8 <- gl(3,3,labels=LETTERS[1:3],length=8)
datL <- list(data=datT8,filt= wrMisc::presenceFilt(datT8,grp=grp8,maxGrpM=1,ratMa=0.8))
testAvB0 <- wrMisc::moderTest2grp(datT8[,1:6],gl(2,3))
testAvB <- test2grp(datL,questNo=1)
```

testRobustToNAimputation

Pair-wise testing robust to NA-imputation

Description

testRobustToNAimputation replaces NA values based on group neighbours (based on grouping of columns in argument gr), following overall assumption of close to Gaussian distribution. Furthermore, it is assumed that NA-values originate from experimental settings where measurements at or below detection limit are recorded as NA. In such cases (eg in proteomics) it is current practice to replace NA-values by very low (random) values in order to be able to perform t-tests. However, random normal values used for replacing may in rare cases deviate from the average (the 'assumed' value) and in particular, if multiple NA replacements are above the average, may look like induced biological data and be misinterpreted as so. The statistical testing uses eBayes from Bioconductor package [limma](#) for robust testing in the context of small numbers of replicates. By repeating multiple times the process of replacing NA-values and subsequent testing the results can be summarized afterwards by median over all repeated runs to remove the stochastic effect of individual NA-imputation. Thus, one may gain stability towards random-character of NA imputations by repeating imputation & test 'nLoop' times and summarize p-values by median (results stabilized at 50-100 rounds). It is necessary to define all groups of replicates in gr to obtain all possible pair-wise testing (multiple columns in \$BH, \$lfdr etc). The modified testing-procedure of Bioconductor package [ROTS](#) may optionally be included, if desired. This function returns a [limma](#)-like S3 list-object further enriched by additional fields/elements.

Usage

```
testRobustToNAimputation(
  dat,
```

```

    gr,
    annot = NULL,
    retnNA = TRUE,
    avSdH = c(0.15, 0.5),
    avSdL = NULL,
    plotHist = FALSE,
    xLab = NULL,
    tit = NULL,
    seedNo = NULL,
    multCorMeth = NULL,
    nLoop = 100,
    lfdrInclude = NULL,
    ROTSn = NULL,
    silent = FALSE,
    callFrom = NULL
  )

```

Arguments

dat	(matrix or data.frame) main data (may contain NA); if dat is list containing \$quant and \$annot as matrix, the element \$quant will be used
gr	(character or factor) replicate association
annot	(matrix or data.frame) annotation (lines must match lines of data !), if annot is NULL and argument dat is a list containing both \$quant and \$annot, the element \$annot will be used
retnNA	(logical) retain and report number of NA
avSdH	(numeric) population characteristics (mean and sd) for >1 NA neighbours 'high' (per line)
avSdL	depreciated argument, no longer used
plotHist	(logical) additional histogram of original, imputed and resultant distribution (made using matrixNAneighbourImpute)
xLab	(character) custom x-axis label
tit	(character) custom title
seedNo	(integer) seed-value for normal random values
multCorMeth	(character)
nLoop	(integer) number of runs of independent NA-imputation
lfdrInclude	(logical) depreciated, please used multCorMeth instead (include lfdr estimations, may cause warning message(s) concerning convergence if few too lines/proteins in dataset tested).
ROTSn	(integer) depreciated, please used multCorMeth instead (number of repeats by ROTs, if NULL ROTs will not be called)
silent	(logical) suppress messages
callFrom	(character) allows easier tracking of messages produced

Details

The argument `multCorMeth` allows to choose which multiple correction algorithms will be used and included to the final results. Possible options are `'lfdr'`, `'BH'`, `'BY'`, `'tValTab'`, `ROTSn='100'` (name to element necessary) or `'noLimma'` (to add initial p.values and BH to limma-results). By default `'lfdr'` (local false discovery rate from package `'fdrtools'`) and `'BH'` (Benjamini-Hochberg FDR) are chosen. The option `'BY'` refers to Benjamini-Yakuteli FDR, `'tValTab'` allows exporting all individual t-values from the repeated NA-substitution and subsequent testing.

Value

limma-type S3 object of class `'MArrayLM'` which can be accessed; multiple results of testing or multiple testing correction types may get included (`'p.value'`, `'FDR'`, `'BY'`, `'lfdr'` or `'ROTS.BH'`)

See Also

`moderTest2grp`, `pVal2lfdr`, eBayes in Bioconductor package `limma`, `t.test`, ROTS of Bioconductor package `ROTS`

Examples

```
set.seed(2015); rand1 <- round(runif(600) + rnorm(600,1,2),3)
dat1 <- matrix(rand1,ncol=6) + matrix(rep((1:100)/20,6),ncol=6)
dat1[13:16,1:3] <- dat1[13:16,1:3] +2 # augment lines 13:16
dat1[19:20,1:3] <- dat1[19:20,1:3] +3 # augment lines 19:20
dat1[15:18,4:6] <- dat1[15:18,4:6] +1.4 # augment lines 15:18
dat1[dat1 <1] <- NA # mimick some NAs for low abundance
## normalize data
boxplot(dat1, main="data before normalization")
dat1 <- wrMisc::normalizeThis(as.matrix(dat1), meth="median")
## designate replicate relationships in samples ...
grp1 <- gl(2, 3, labels=LETTERS[1:2])
## moderated t-test with repeated inputations (may take >10 sec, >60 sec if ROTSn >0 !)
PLtestR1 <- testRobustToNAimputation(dat=dat1, gr=grp1, retnNA=TRUE, nLoop=70)
names(PLtestR1)
```

Description

This type of plot is very common in high-throughput biology, see [Volcano-plot](#). Basically, this plot allows comparing the outcome of a statistical test to the differential of the group means (ie log fold-change),

Usage

```

VolcanoPlotW2(
  Mvalue,
  pValue = NULL,
  useComp = 1,
  filtFin = NULL,
  ProjNa = NULL,
  FCthrs = NULL,
  FdrList = NULL,
  FdrThrs = NULL,
  FdrType = NULL,
  subTxt = NULL,
  grayIncr = TRUE,
  col = NULL,
  pch = 16,
  compNa = NULL,
  batchFig = FALSE,
  cexMa = 1.8,
  cexLa = 1.1,
  limM = NULL,
  limp = NULL,
  annotColumn = c("SpecType", "GeneName", "EntryName", "Accession", "Species",
    "Contam"),
  annColor = NULL,
  cexPt = NULL,
  cexSub = NULL,
  cexTxLab = 0.7,
  namesNBest = NULL,
  NbestCol = 1,
  sortLeg = "descend",
  NaSpecTypeAsContam = TRUE,
  useMar = c(6.2, 4, 4, 2),
  returnData = FALSE,
  callFrom = NULL,
  silent = FALSE,
  debug = FALSE
)

```

Arguments

Mvalue	(numeric or matrix) data to plot; M-values are typically calculated as difference of log ₂ -abundance values and 'pValue' the mean of log ₂ -abundance values; M-values and p-values may be given as 2 columns of a matrix, in this case the argument pValue should remain NULL
pValue	(numeric, list or data.frame) if NULL it is assumed that 2nd column of 'Mvalue' contains the p-values to be used
useComp	(integer, length=1) choice of which of multiple comparisons to present in Mvalue (if generated using moderTestXgrp())

<code>filtFin</code>	(matrix or logical) The data may get filtered before plotting: If FALSE no filtering will get applied; if matrix of TRUE/FALSE it will be used as optional custom filter, otherwise (if Mvalue if an MArrayLM-object eg from limma) a default filtering based on the <code>filtFin</code> element will be applied
<code>ProjNa</code>	(character) custom title
<code>FCthrs</code>	(numeric) Fold-Change threshold (display as line) give as Fold-change and NOT $\log_2(\text{FC})$, default at 1.5, set to NA for omitting
<code>FdrList</code>	(numeric) FDR data or name of list-element
<code>FdrThrs</code>	(numeric) FDR threshold (display as line), default at 0.05, set to NA for omitting
<code>FdrType</code>	(character) FDR-type to extract if Mvalue is 'MArrayLM'-object (eg produced by from <code>moderTest2grp</code> etc); if NULL it will search for suitable fields/values in this order : 'FDR','BH','lfdr' and 'BY'
<code>subTxt</code>	(character) custom sub-title
<code>grayIncrem</code>	(logical) if TRUE, display overlay of points as increased shades of gray
<code>col</code>	(character) custom color(s) for points of plot (see also par)
<code>pch</code>	(integer) type of symbol(s) to plot (default=16) (see also par)
<code>compNa</code>	(character) names of groups compared
<code>batchFig</code>	(logical) if TRUE figure title and axes legends will be kept shorter for display on fewer space
<code>cexMa</code>	(numeric) font-size of title, as expansion factor (see also <code>cex</code> in par)
<code>cexLa</code>	(numeric) size of axis-labels, as expansion factor (see also <code>cex</code> in par)
<code>limM</code>	(numeric, length=2) range of axis M-values
<code>limp</code>	(numeric, length=2) range of axis FDR / p-values
<code>annotColumn</code>	(character) column names of annotation to be extracted (only if Mvalue is MArrayLM-object containing matrix \$annot). The first entry (typically 'SpecType') is used for different symbols in figure, the second (typically 'GeneName') is used as preferred text for annotating the best points (if <code>namesNBest</code> allows to do so.)
<code>annColor</code>	(character or integer) colors for specific groups of annotation (only if Mvalue is MArrayLM-object containing matrix \$annot)
<code>cexPt</code>	(numeric) size of points, as expansion factor (see also <code>cex</code> in par)
<code>cexSub</code>	(numeric) size of subtitle, as expansion factor (see also <code>cex</code> in par)
<code>cexTxLab</code>	(numeric) size of text-labels for points, as expansion factor (see also <code>cex</code> in par)
<code>namesNBest</code>	(integer or character) number of best points to add names in figure; if 'passThr' all points passing FDR and FC-filtes will be selected; if the initial object Mvalue contains a list-element called 'annot' the second of the column specified in argument <code>annotColumn</code> will be used as text
<code>NbestCol</code>	(character or integer) colors for text-labels of best points
<code>sortLeg</code>	(character) sorting of 'SpecType' annotation either ascending ('ascend') or descending ('descend'), no sorting if NULL
<code>NaSpecTypeAsContam</code>	(logical) consider lines/proteins with NA in Mvalue\$annot["SpecType"] as contaminants (if a 'SpecType' for contaminants already exists)

useMar	(numeric,length=4) custom margins (see also par)
returnData	(logical) optional returning data.frame with (ID, Mvalue, pValue, FDRvalue, passFilt)
callFrom	(character) allow easier tracking of message(s) produced
silent	(logical) suppress messages
debug	(logical) additional messages for debugging

Details

In high-throughput biology data are typically already transformed to log₂ and thus, the 'M'-value represents a relative change. Besides, output from statistical testing by [moderTest2grp](#) or [moderTestXgrp](#) can be directly read to produce Volcano plots for diagnostic reasons. Please note, that plotting a very number of points in transparency (eg >10000) may take several seconds.

Value

MA-plot only

See Also

(for PCA) [plotPCAw](#)

Examples

```
library(wrMisc)
set.seed(2005); mat <- matrix(round(runif(900),2), ncol=9)
rownames(mat) <- paste0(rep(letters[1:25],each=4), rep(letters[2:26],4))
mat[1:50,4:6] <- mat[1:50,4:6] + rep(c(-1,1)*0.1,25)
mat[3:7,4:9] <- mat[3:7,4:9] + 0.7
mat[11:15,1:6] <- mat[11:15,1:6] - 0.7
## assume 2 groups with 3 samples each
gr3 <- gl(3,3,labels=c("C","A","B"))
tRes2 <- moderTest2grp(mat[,1:6], gl(2,3), addResults = c("FDR","means"))
# Note: due to the small number of lines only FDR chosen to calculate
VolcanoPlotW2(tRes2)
## Add names of points passing custom filters
VolcanoPlotW2(tRes2, FCth=1.3, FdrThrs=0.2, namesNBest="passThr")

## assume 3 groups with 3 samples each
tRes <- moderTestXgrp(mat, gr3, addResults = c("FDR","means"))
# Note: due to the small number of lines only FDR chosen to calculate
VolcanoPlotW2(tRes)
VolcanoPlotW2(tRes, FCth=1.3, FdrThrs=0.2)
VolcanoPlotW2(tRes, FCth=1.3, FdrThrs=0.2, useComp=2)
```

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