Package ‘pRolocGUI’

May 16, 2024

Title  Interactive visualisation of spatial proteomics data

Version  2.14.0

Description  The package pRolocGUI comprises functions to interactively visualise spatial proteomics data on the basis of pRoloc, pRolocdata and shiny.

Depends  methods, R (>= 3.1.0), pRoloc (>= 1.27.6), Biobase, MSnbase (>= 2.1.11)

Imports  shiny (>= 0.9.1), scales, dplyr, DT (>= 0.1.40), graphics, utils, ggplot2, shinydashboardPlus (>= 2.0.0), colourpicker, shinyhelper, shinyWidgets, shinyjs, colorspace, stats, grDevices, grid, BiocGenerics, shinydashboard

Suggests  pRolocdata, knitr, BiocStyle (>= 2.5.19), rmarkdown, testthat (>= 3.0.0)

License  GPL-2

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

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

VignetteBuilder  knitr

Video  https://www.youtube.com/playlist?list=PLvIXxpatsLA2loV5Srs2VBpJlYUIVJ4ow

biocViews  Proteomics, Visualization, GUI

Encoding  UTF-8

RoxygenNote  7.1.1

Config/testthat/edition  3

git_url  https://git.bioconductor.org/packages/pRolocGUI

git_branch  RELEASE_3_19

git_last_commit  747fc75

git_last_commit_date  2024-04-30

Repository  Bioconductor 3.19

Date/Publication  2024-05-15
pRolocVis

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pRolocVis  
Interactive visualisation of spatial proteomics data

Description

These functions allow one to explore spatial proteomics data interactively.

Usage

pRolocVis(object, app = "explore", fcol = "markers", ...)
pRolocVis_aggregate(
  object,
  fcol = "markers",
  groupBy,
  fig.height = "700px",
  nchar = 25,
  ...
)

pRolocVis_compare(
  object,
  fcol = "markers",
  classProfiles = FALSE,
  fig.height = "400px",
  nchar = 25,
  ...
)

pRolocVis_explore(
  object,
  fcol = "markers",
  classProfiles = FALSE,
  fig.height = "700px",
  nchar = 25,
  ...
)
Arguments

object  An instance of class MSnSet, or an MSnSetList of length 2 if using "compare" application.

app     The type of application requested: "explore" (default), "compare" or "aggregate". See description below.

fcol   The feature meta-data label (fData column name) to be used for colouring. Default is "markers". If set to NULL, no annotation is expected. For the "compare" app, a character vector of length 2 is allowed to set different labels for each dataset. If only one label is specified, then this single label will be used to identify the annotation column in both datasets. Please see example herein.

...    Additional parameters passed to plot2D for the "explore" (such as the dimensionality reduction technique, and methods), "compare" apps. For the "aggregate" app this is for additional parameters to be passed to combineFeatures.

groupBy The feature meta-data label (fData column name) to be used for summarising the features to be combined.

fig.height Height of the figure.

nchar    Maximum number of characters of the subcellular class names, before their names are truncated. Default is 25.

classProfiles A logical indicating if a tab displaying individual class profile plots should be displayed. Default is FALSE.

Details

The function pRolocVis is a wrapper for pRolocVis_pca, pRolocVis_compare, and pRolocVis_aggregate. These Shiny apps allow to explore and analyse interactively spatial proteomics data.

The explore Shiny app allows exploration of quantitive data (1) visually through a projection of the dataset, (2) protein profiles, and (3) a searchable feature data table, allowing visualisation of sets of proteins of interest.

The compare Shiny app is meant for comparing protein localisation between two conditions, or two different experiments, replicates etc.

The aggregation Shiny app displays a scatter plot of the maximum or mean distances within each feature (e.g. protein group) according to its components (e.g. peptides) defined by the groupBy argument. A PCA plot of the components is also displayed. It can be used for visualising peptides, PSMs or any other features defined in the feature data of the MSnSet and their distributions.

Author(s)

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See Also

The package vignette: vignette("pRolocGUI").
Examples

library("pRoloc")
library("pRolocdata")

## Load the Explore app
data(hyperLOPIT2015)
if (interactive()) {
  pRolocVis(hyperLOPIT2015)
pRolocVis(hyperLOPIT2015, method = "t-SNE")
## store the t-SNE coords and pass a matrix to pRolocVis
xx <- plot2D(hyperLOPIT2015, method = "t-SNE")
pRolocVis(xx, method = "none", methargs = list(hyperLOPIT2015))
}

## Load the Compare app
data("hyperLOPITU2OS2018")
data("lopitdcU2OS2018")
xx <- MSnSetList(list(hyperLOPITU2OS2018, lopitdcU2OS2018))
if (interactive()) {
  pRolocVis(xx, app = "compare", fcol = c("markers", "final.assignment"))
}

## Visualise the location and distribution of peptides per protein group
data("hyperLOPIT2015ms2psm")
if (interactive()) {
  ## Combine PSM data to peptides
  hl <- combineFeatures(hyperLOPIT2015ms2psm,
                        groupBy = fData(hyperLOPIT2015ms2psm)$Sequence,
                        method = median)
  ## Visualise peptides according to protein group
  pRolocVis(hl, app = "aggregate", fcol = "markers",
            groupBy = "Protein.Group.Accessions")
}
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