

Package ‘cellity’

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Title Quality Control for Single-Cell RNA-seq Data

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Description A support vector machine approach to identifying and filtering low quality cells from single-cell RNA-seq datasets.

License GPL (>= 2)

Depends R (>= 3.3)

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Contents

| | |
|-------------------------|----|
| cellity-package | 2 |
| assess_cell_quality_PCA | 2 |
| assess_cell_quality_SVM | 3 |
| extract_features | 4 |
| extra_human_genes | 5 |
| extra_mouse_genes | 6 |
| feature_generation | 6 |
| feature_info | 7 |
| mES1_features | 7 |
| mES1_labels | 8 |
| multiplot | 9 |
| normalise_by_factor | 9 |
| param_mES_all | 10 |
| param_mES_common | 10 |
| plot_pca | 11 |
| sample_counts | 12 |
| sample_stats | 12 |
| simple_cap | 13 |
| sum_prop | 13 |
| training_mES_features | 14 |
| training_mES_labels | 14 |
| uni.plot | 15 |

| | |
|--------------|-----------|
| Index | 16 |
|--------------|-----------|

| | |
|-----------------|---|
| cellity-package | <i>Quality Control for Single-Cell RNA-seq Data</i> |
|-----------------|---|

Description

cellity provides a support vector machine and PCA approaches to identifying and filtering low quality cells from single-cell RNA-seq datasets.

| | |
|-------------------------|--|
| assess_cell_quality_PCA | <i>ASSESS CELL QUALITY USING PCA AND OUTLIER DETECTION</i> |
|-------------------------|--|

Description

ASSESS CELL QUALITY USING PCA AND OUTLIER DETECTION

Usage

```
assess_cell_quality_PCA(features, file = "")
```

Arguments

features Input dataset containing features (cell x features)
 file Output_file where plot is saved

Details

This function applies PCA on features and uses outlier detection to determine which cells are low and which are high quality

Value

Returns a dataframe indicating which cell is low or high quality (0 or 1 respectively)

Examples

```
data(training_mES_features)
training_mES_features_all <- training_mES_features[[1]]
training_quality_PCA_allF <- assess_cell_quality_PCA(training_mES_features_all)
```

assess_cell_quality_SVM

Assess quality of a cell - SVM version

Description

Assess quality of a cell - SVM version

Usage

```
assess_cell_quality_SVM(training_set_features, training_set_labels,
    ensemble_param, test_set_features)
```

Arguments

training_set_features A training set containing features (cells x features) for prediction
 training_set_labels Annotation of each individual cell if high or low quality (1 or 0 respectively)
 ensemble_param Dataframe of parameters for SVM
 test_set_features Dataset to predict containing features (cells x features)

Details

This function takes a training set + annotation to predict a test set. It requires that hyper-parameters have been optimised.

Value

Returns a dataframe indicating which cell is low or high quality (0 or 1 respectively)
data.frame with decision on quality of cells

Examples

```
data(param_mES_all)
data(training_mES_features)
data(training_mES_labels)
data(mES1_features)
data(mES1_labels)
mES1_features_all <- mES1_features[[1]]
training_mES_features_all <- training_mES_features[[1]]
mES1_quality_SVM <- assess_cell_quality_SVM( training_mES_features_all,
training_mES_labels[,2], param_mES_all, mES1_features_all)
```

| | |
|------------------|---|
| extract_features | <i>Extracts biological and technical features for given dataset</i> |
|------------------|---|

Description

Extracts biological and technical features for given dataset

Usage

```
extract_features(counts_nm, read_metrics, prefix = "", output_dir = "",
  common_features = NULL, GO_terms = NULL, extra_genes = NULL,
  organism = "mouse")
```

Arguments

| | |
|-----------------|---|
| counts_nm | Gene expression counts dataframe (genes x cells). Either normalised by library size or TPM values |
| read_metrics | Dataframe with mapping statistics produced by python pipeline |
| prefix | Prefix of outputfiles |
| output_dir | Output directory of files |
| common_features | Subset of features that are applicable within one species, but across cell types |
| GO_terms | DataFrame with gene ontology term IDs, that will be used in feature extraction |
| extra_genes | Additional genes used for feature extraction |
| organism | The target organism to generate the features for |

Details

This function takes a combination of gene counts and mapping statistics to extract biological and technical features, which than can be used for quality data analysis

Value

a list with two elements, one providing all features, and one providing common features.

Examples

```
data(sample_counts)
data(sample_stats)
sample_counts_nm <- normalise_by_factor(sample_counts, colSums(sample_counts))
sample_features <- extract_features(sample_counts_nm, sample_stats)
```

extra_human_genes *Additional human genes that are used in feature extraction*

Description

This list contains human genes that are used for feature extraction of biological features

Usage

```
extra_human_genes
```

Format

a list containing vectors of genes. Name indicates which GO category.

Value

NULL, but makes available a list with metadata

Author(s)

Tomislav Ilicic & Davis McCarthy, 2015-03-05

Source

Wellcome Trust Sanger Institute

extra_mouse_genes *Additional mouse genes that are used in feature extraction*

Description

This list contains mouse genes that are used for feature extraction of biological features

Usage

extra_mouse_genes

Format

a list containing vectors of genes. Name indicates which GO category.

Value

NULL, but makes available a list with metadata

Author(s)

Tomislav Ilicic & Davis McCarthy, 2015-03-05

Source

Wellcome Trust Sanger Institute

feature_generation *Helper Function to create all features*

Description

Helper Function to create all features

Usage

```
feature_generation(counts_nm, read_metrics, GO_terms, extra_genes, organism)
```

Arguments

| | |
|--------------|---|
| counts_nm | Gene expression counts dataframe (genes x cells). Either normalised by library size or TPM values |
| read_metrics | Dataframe with mapping statistics produced by python pipeline |
| GO_terms | DataFrame with gene ontology term IDs, that will be used in feature extraction |
| extra_genes | Additional genes used for feature extraction |
| organism | The target organism to generate the features for |

Value

Returns the entire set of features in a data.frame

| | |
|--------------|--|
| feature_info | <i>Information which genes and GO categories should be included as features. Also defines which features are cell-type independent (common features)</i> |
|--------------|--|

Description

This list contains metadata information that is used to extract features from in the function extract_features

Usage

```
feature_info
```

Format

a list with 2 elements (GO_terms,common_features).

Value

NULL, but makes available a list with metadata

Author(s)

Tomislav Ilicic & Davis McCarthy, 2015-03-05

Source

Wellcome Trust Sanger Institute

| | |
|---------------|---|
| mES1_features | <i>Real test dataset containing all and common features from the paper (mESI)</i> |
|---------------|---|

Description

This list contains 2 dataframes where each contains features per cell (cell X features) that can be used for classification.

Usage

```
mES1_features
```

Format

a list with 2 elements (all_features, common_features).

Value

NULL, but makes available a list with 2 dataframes

Author(s)

Tomislav Ilicic & Davis McCarthy, 2015-03-05

Source

Wellcome Trust Sanger Institute

mES1_labels

Real test dataset containing annotation of cells

Description

This data frame has 2 columns: First showing cell names, the second indicating if cell is of low (0) or high (1) quality

Usage

```
mES1_labels
```

Format

a dataframe with 2 columns (cell_names, label).

Value

NULL, but makes available a dataframe with cell annotations

Author(s)

Tomislav Ilicic & Davis McCarthy, 2015-03-05

Source

Wellcome Trust Sanger Institute

| | |
|-----------|---|
| multiplot | <i>Internal multiplot function to combine plots onto a grid</i> |
|-----------|---|

Description

Internal multiplot function to combine plots onto a grid

Usage

```
multiplot(..., plotlist = NULL, file, cols = 6, layout = NULL)
```

Arguments

| | |
|----------|---|
| ... | individual plots to combine into a single plot |
| plotlist | a vector with names of plots to use in the plot |
| file | string giving filename to which pdf of plots is to be saved |
| cols | integer giving number of columns for the plot |
| layout | matrix defining the layout for the plots |

Value

a plot object

| | |
|---------------------|---|
| normalise_by_factor | <i>Internal function to normalize by library size</i> |
|---------------------|---|

Description

Internal function to normalize by library size

Usage

```
normalise_by_factor(counts, norm_factor)
```

Arguments

| | |
|-------------|---------------------------------|
| counts | matrix of counts |
| norm_factor | vector of normalisation factors |

Value

a matrix with normalized gene counts

Examples

```
data(sample_counts)
data(sample_stats)
sample_counts_nm <- normalise_by_factor(sample_counts, colSums(sample_counts))
```

param_mES_all *Parameters used for SVM classification*

Description

This data frame has 3 columns: gamma, cost, class.weights and is optimised for all features and our training data

Usage

```
param_mES_all
```

Format

a dataframe with 3 columns (gamma, cost, class.weights).

Value

NULL, but makes available a dataframe with parameters

Author(s)

Tomislav Ilicic & Davis McCarthy, 2015-03-05

Source

Wellcome Trust Sanger Institute

param_mES_common *Parameters used for SVM classification*

Description

This data frame has 3 columns: gamma, cost, class.weights and is optimised for common features and our training data

Usage

```
param_mES_common
```

Format

a dataframe with 3 columns (gamma, cost, class.weights).

Value

NULL, but makes available a dataframe with parameters

Author(s)

Tomislav Ilicic & Davis McCarthy, 2015-03-05

Source

Wellcome Trust Sanger Institute

| | |
|----------|---|
| plot_pca | <i>Plots PCA of all features. Colors high and low quality cells based on outlier detection.</i> |
|----------|---|

Description

Plots PCA of all features. Colors high and low quality cells based on outlier detection.

Usage

```
plot_pca(features, annot, pca, col, output_file)
```

Arguments

- features Input dataset containing features (cell x features)
- annot Matrix annotation of each cell
- pca PCA of features
- col color code indicating what color high and what low quality cells
- output_file where plot is stored

Details

This function plots PCA of all features + most informative features

Value

Plots of PCA

| | |
|---------------|--|
| sample_counts | <i>Sample gene expression data containing 40 cells</i> |
|---------------|--|

Description

This data frame contains genes (rows) and cells (columns) showing raw read counts

Usage

```
sample_counts
```

Format

a dataframe with genes x cells

Value

NULL, but makes available a dataframe with raw read counts

Author(s)

Tomislav Ilicic & Davis McCarthy, 2015-03-05

Source

Wellcome Trust Sanger Institute

| | |
|--------------|--|
| sample_stats | <i>Sample read statistics data containing 40 cells</i> |
|--------------|--|

Description

This data frame contains read metrics (columns) and cells (rows)

Usage

```
sample_stats
```

Format

a dataframe with cells x metrics

Value

NULL, but makes available a dataframe with read statistics

Author(s)

Tomislav Ilicic & Davis McCarthy, 2015-03-05

Source

Wellcome Trust Sanger Institute

| | |
|------------|--|
| simple_cap | <i>Converts all first letters to capital letters</i> |
|------------|--|

Description

Converts all first letters to capital letters

Usage

```
simple_cap(x)
```

Arguments

| | |
|---|--------|
| x | string |
|---|--------|

Value

a character vector in title case

| | |
|----------|--|
| sum_prop | <i>Sums up normalised values of genes to groups.</i> |
|----------|--|

Description

Supports TPM and proportion of mapped reads.

Usage

```
sum_prop(counts, genes_interest)
```

Arguments

| | |
|----------------|---|
| counts | Normalised gene expression count matrix |
| genes_interest | dataframe of genes of interest to merge |

Value

a vector of sums per group

training_mES_features *Original training dataset containing all and common features from the paper (training mES)*

Description

This list contains 2 dataframes where each contains features per cell (cell X features) that can be used for classification.

Usage

```
training_mES_features
```

Format

a list with 2 elements (all_features, common_features).

Value

NULL, but makes available a list with 2 dataframes

Author(s)

Tomislav Ilicic & Davis McCarthy, 2015-03-05

Source

Wellcome Trust Sanger Institute

training_mES_labels *Original training dataset containing annotation of cells*

Description

This data frame has 2 columns: First showing cell names, the second indicating if cell is of low (0) or high (1) quality

Usage

```
training_mES_labels
```

Format

a dataframe with 2 columns (cell_names, label).

Value

NULL, but makes available a dataframe with cell annotations

Author(s)

Tomislav Ilicic & Davis McCarthy, 2015-03-05

Source

Wellcome Trust Sanger Institute

| | |
|----------|--|
| uni.plot | <i>Internal function to detect outliers from the mvoutlier package Modified slightly so that plots are not printed</i> |
|----------|--|

Description

Internal function to detect outliers from the mvoutlier package Modified slightly so that plots are not printed

Usage

```
uni.plot(x, symb = FALSE, quan = 1/2, alpha = 0.025)
```

Arguments

| | |
|-------|----------------------------|
| x | A matrix containing counts |
| symb | Symbols |
| quan | quan |
| alpha | alpha |

Value

a list of outlier indicators

Index

`assess_cell_quality_PCA`, [2](#)
`assess_cell_quality_SVM`, [3](#)

`cellity-package`, [2](#)

`extra_human_genes`, [5](#)
`extra_mouse_genes`, [6](#)
`extract_features`, [4](#)

`feature_generation`, [6](#)
`feature_info`, [7](#)

`mES1_features`, [7](#)
`mES1_labels`, [8](#)
`multiplot`, [9](#)

`normalise_by_factor`, [9](#)

`param_mES_all`, [10](#)
`param_mES_common`, [10](#)
`plot_pca`, [11](#)

`sample_counts`, [12](#)
`sample_stats`, [12](#)
`simple_cap`, [13](#)
`sum_prop`, [13](#)

`training_mES_features`, [14](#)
`training_mES_labels`, [14](#)

`uni.plot`, [15](#)