

Package: scAnnotate (via r-universe)

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Type Package

Title An Automated Cell Type Annotation Tool for Single-Cell RNA-Sequencing Data

Version 0.3

Description An entirely data-driven cell type annotation tools, which requires training data to learn the classifier, but not biological knowledge to make subjective decisions. It consists of three steps: preprocessing training and test data, model fitting on training data, and cell classification on test data. See Xiangling Ji, Danielle Tsao, Kailun Bai, Min Tsao, Li Xing, Xuekui Zhang.(2022)<[doi:10.1101/2022.02.19.481159](https://doi.org/10.1101/2022.02.19.481159)> for more details.

Depends R(>= 4.0.0)

License GPL-3

URL <https://doi.org/10.1101/2022.02.19.481159>

Encoding UTF-8

LazyData true

RoxygenNote 7.3.1

Suggests knitr, testthat (>= 3.0.0), rmarkdown

VignetteBuilder knitr

Imports glmnet, stats, Seurat (>= 5.0.1), harmony, SeuratObject

Config/testthat/edition 3

NeedsCompilation no

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Repository <https://ubcxzhang.r-universe.dev>

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eva_cal	<i>eva_cal</i>
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Description

calculate the F1 score of each cell population, mean of F1 score and overall accuracy

Usage

```
eva_cal(prediction, cell_label)
```

Arguments

prediction	A vector of annotate cell type labels
cell_label	A vector of original cell type labels

Value

A matrix contain the F1 score of each cell population, mean of F1 score and overall accuracy

Examples

```
data(predict_label)
data(pbmc2)
eva_cal(prediction = predict_label, cell_label = pbmc2[,1])
```

pbmc1

pbmc1

Description

A subset of human Peripheral Blood Mononuclear Cells (PBMC) scRNA-seq data that was sequenced using Drop-seq platform. The Seurat(version 4.0.5) package was used for normalized using the NormalizeData function with the "LogNormalize" method and a scale factor of 10,000. After modeling the mean-variance relationship with the FindVariableFeautre function within "vst" methods, we selected the top 2,000 highly variable genes and only used this selection going forward. The dataframe of the cell type label and a gene expression matrix of 598 cells in the row and 2,000 genes in the column.

Usage

```
data(pbmc1, package="scAnnotate")
```

Format

a data frame

References

Ding, J.et al.(2019). Systematic comparative analysis of single cellrna-sequencing methods.bioRxiv

pbmc2

pbmc2

Description

A subset of human PBMC scRNA-seq data that was sequenced using inDrops platform. The Seurat(version 4.0.5) package was used for normalized using the NormalizeData function with the "LogNormalize" method and a scale factor of 10,000. After modeling the mean-variance relationship with the FindVariableFeautre function within "vst" methods, we selected the top 2,000 highly variable genes and only used this selection going forward. The dataframe of the cell type label and a gene expression matrix of 644 cells in the row and 2,000 genes in the column.

Usage

```
data(pbmc2, package="scAnnotate")
```

Format

a data frame

References

Ding, J.et al.(2019). Systematic comparative analysis of single cellrna-sequencing methods.bioRxiv

predict_label	<i>predict_label</i>
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Description

Cell type annotation of pbmc2 data that training from pbmc1 data by 'scAnnotate'.

Usage

```
data(predict_label, package="scAnnotate")
```

Format

a data frame

scAnnotate	<i>scAnnotate</i>
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Description

Annotate cell type labels of test data using a trained mixture model from training data

Usage

```
scAnnotate(
  train,
  test,
  distribution = c("normal", "dep"),
  correction = c("auto", "harmony", "seurat"),
  screening = c("wilcox", "t.test"),
  threshold = 0,
  lognormalized = TRUE
)
```

Arguments

train	A data frame of cell type label in the first column and a gene expression matrix where each row is a cell and each column is a gene from training data
test	A data matrix where each row is a cell and each column is a gene from test data
distribution	A character string indicates the distribution assumption on positive gene expression, which should be one of "normal"(default) or "dep". "dep" refers to depth measure, which is a non-parametric distribution estimation approach.

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