

# Package ‘scTGIF’

April 15, 2020

**Type** Package

**Title** Cell type annotation for unannotated single-cell RNA-Seq data

**Version** 1.0.0

**Depends** R (>= 3.6.0)

**Imports** GSEABase, Biobase, SingleCellExperiment, BiocStyle, plotly, tagcloud, rmarkdown, Rcpp, grDevices, graphics, utils, knitr, S4Vectors, SummarizedExperiment, RColorBrewer, nnTensor, methods, scales, msigdbr

**Suggests** testthat

**Description** scTGIF connects the cells and the related gene functions without cell type label.

**License** Artistic-2.0

**biocViews** DimensionReduction, QualityControl, SingleCell, Software, GeneExpression

**VignetteBuilder** knitr

**git\_url** <https://git.bioconductor.org/packages/scTGIF>

**git\_branch** RELEASE\_3\_10

**git\_last\_commit** 5e82bd9

**git\_last\_commit\_date** 2019-10-29

**Date/Publication** 2020-04-14

**Author** Koki Tsuyuzaki [aut, cre]

**Maintainer** Koki Tsuyuzaki <k.t.the-answer@hotmail.co.jp>

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scTGIF-package

*Cell type annotation for unannotated single-cell RNA-Seq data*

## Description

scTGIF connects the cells and the related gene functions without cell type label.

## Details

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Index: This package was not yet installed at build time.

`calcTGIF` function calculates what kind of cellular patterns and functional patterns are contained in single-cell RNA-seq data and `reportTGIF` function generates report of analytic result. The algorithm is based on joint NMF, which is implemented in nnTensor package.

## Author(s)

NA

Maintainer: NA

## References

Dominic Grun, Anna Lyubimova, Lennart Kester, Kay Wiebrands, Onur Basak, Nobuo Sasaki, Hans Clevers, Alexander van Oudenaarden (2015) Single-cell messenger RNA sequencing reveals rare intestinal cell types. *Nature*, **525**: 251-255

calcTGIF

*Function for connecting cellular patterns and functional patterns using jNMF*

## Description

`calcTGIF` function calculates what kind of cellular patterns and functional patterns are contained in single-cell RNA-seq data and `reportTGIF` function generates report of analytic result.

## Usage

```
calcTGIF(sce, rank)
```

## Arguments

sce	A object generated by instantiation of SingleCellExperiment-class.
rank	Rank parameter of joint NMF algorithm.

## Value

The result is saved to metadata slot of SingleCellExperiment object.

**Author(s)**

NA

**Examples**

```
showMethods("calcTGIF")
```

---

**DistalLungEpithelium**    *Gene expression matrix of DistalLungEpithelium dataset containing five cluster.*

---

**Description**

A data frame with 3397 rows (genes) with following 80 columns (cells).

The data is downloaded as supplementary information of the distal lung epithelium paper (<https://www.nature.com/article/nbt.3102.html>)

Low-expressed genes are filtered.

*All Gene ID is converted to Human Entrez Gene ID for applying the data to scTGIF.*

**Usage**

```
data("DistalLungEpithelium")
```

**Source**

<http://www.nature.com/nbt/journal/v33/n2/full/nbt.3102.html>

**References**

Treutlein, B. et al. (2014) Reconstructing lineage hierarchies of the distal lung epithelium using single-cell RNA-seq. *Nature* **509**, 371-375

**Examples**

```
data("DistalLungEpithelium")
```

---

**label.DistalLungEpithelium**    *Cellular label of DistalLungEpithelium dataset containing five cluster.*

---

**Description**

A vector containing 80 elements (cells).

**Usage**

```
data("label.DistalLungEpithelium")
```

## References

Treutlein, B. et al. (2014) Reconstructing lineage hierarchies of the distal lung epithelium using single-cell RNA-seq. *Nature* **509**, 371-375

## Examples

```
data("label.DistalLungEpithelium")
```

pca.DistalLungEpithelium

*The result of PCA of the DistalLungEpithelium dataset.*

## Description

A matrix having 80 (cells) \* 2 (PCs) elements.

## Usage

```
data("pca.DistalLungEpithelium")
```

## References

Treutlein, B. et al. (2014) Reconstructing lineage hierarchies of the distal lung epithelium using single-cell RNA-seq. *Nature* **509**, 371-375

## Examples

```
data("pca.DistalLungEpithelium")
```

reportTGIF

*Function for reporting the result of [calcTGIF](#) function*

## Description

[calcTGIF](#) function calculates what kind of cellular patterns and functional patterns are contained in single-cell RNA-seq data and [reportTGIF](#) function generates report of analytic result.

## Usage

```
reportTGIF(sce, out.dir=tempdir(), html.open=TRUE,
           title="The result of scTGIF",
           author="The person who runs this script",
           assayNames="counts")
```

**Arguments**

sce	A object generated by instantiation of SingleCellExperiment-class.
out.dir	Output directory user want to save the report (Default: tempdir()).
html.open	Whether html is opened when <code>reportTGIF</code> is finished (Default: TRUE)
title	Title of report (Default: "The result of scTGIF")
author	The name of user name (Default: "The person who runs this script")
assayNames	The unit of gene expression for using scTGIF (e.g. normcounts, cpm...etc) (Default: "counts").

**Value**

Some file is generated to output directory user specified.

**Author(s)**

NA

**Examples**

```

if(interactive()){
  # Package loading
  library("SingleCellExperiment")
  library("GSEABase")
  library("msigdbr")

  # Test data
  data("DistalLungEpithelium")
  data("pca.DistalLungEpithelium")
  data("label.DistalLungEpithelium")

  # Test data
  par(ask=FALSE)
  plot(pca.DistalLungEpithelium, col=label.DistalLungEpithelium, pch=16,
    main="Distal lung epithelium dataset", xlab="PCA1",
    ylab="PCA2", bty="n")
  text(0.1, 0.05, "AT1", col="#FF7F00", cex=2)
  text(0.07, -0.15, "AT2", col="#E41A1C", cex=2)
  text(0.13, -0.04, "BP", col="#A65628", cex=2)
  text(0.125, -0.15, "Clara", col="#377EB8", cex=2)
  text(0.09, -0.2, "Ciliated", col="#4DAF4A", cex=2)

  # Load the gmt file from MSigDB
  # Only "Entrez Gene ID" can be used in scTGIF
  # e.g. gmt <- GSEABase::getGmt(
  #       "/PATH/YOU/SAVED/THE/GMTFILES/h.all.v6.0.entrez.gmt")
  # Here we use msigdbr to retrieve mouse gene sets

  # Mouse gene set (NCBI Gene ID)
  m_df <- msigdbr(species = "Mus musculus", category = "H")[,
    c("gs_name", "entrez_gene")]

  # Convert to GeneSetCollection
  hallmark = unique(m_df$gs_name)
  gsc <- lapply(hallmark, function(h){

```

```

target = which(m_df$gs_name == h)
geneIds = unique(as.character(m_df$entrez_gene[target]))
GeneSet(setName=h, geneIds)
})
gmt <- GeneSetCollection(gsc)

# SingleCellExperiment-class
sce <- SingleCellExperiment(
  assays = list(counts = DistalLungEpithelium))
reducedDims(sce) <- SimpleList(PCA=pca.DistalLungEpithelium)

# User's Original Normalization Function
CPMED <- function(input){
  libsize <- colSums(input)
  median(libsize) * t(t(input) / libsize)
}
# Normalization
normcounts(sce) <- log10(CPMED(counts(sce)) + 1)

# Registration of required information into metadata(sce)
sce2 <- settingTGIF(sce, gmt, reducedDimNames="PCA",
  assayNames="normcounts")

# Functional Annotation based on jNMF
sce2 <- calcTGIF(sce2, rank=7)

# HTML Reprt
reportTGIF(sce2,
  html.open=TRUE,
  title="scTGIF Report for DistalLungEpithelium dataset",
  author="Koki Tsuyuzaki")
}

```

**settingTGIF***Paramter setting for scTGIF***Description**

All parameters is saved to metadata slot of SingleCellExperiment object.

**Usage**

```
settingTGIF(sce, gmt, reducedDimNames, assayNames="counts", grid.size=50)
```

**Arguments**

sce	A object generated by instantiation of SingleCellExperiment-class.
gmt	Object generated from GSEABase::getGmt function. GMT file can be downloaded from MSigDB web (site <a href="http://software.broadinstitute.org/gsea/login.jsp#msigdb">http://software.broadinstitute.org/gsea/login.jsp#msigdb</a> ). Please confirm that the gmt file contains Human Entrez Gene ID, not gene symbol. Also confirm that the DataMatrix has Human Entrez Gene ID.
reducedDimNames	The names of reducedDim(sce) that user want use in scTGIF.

assayNames	The unit of gene expression for using scTGIF (e.g. normcounts, cpm...etc) (Default: "counts").
grid.size	The grid size for segmentation of the two dimensional plot of reducedDim(sce) (Default: 50, which means 50*50 grids).

**Value**

The result is saved to metadata slot of SingleCellExperiment object.

**Author(s)**

NA

**Examples**

```
showMethods("settingTGIF")
```

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