Package 'scMAGeCK'

March 30, 2021

Type Package

single-cell CRISPR screening data	
Version 1.2.0	
Pate 2019-12-13	
author Wei Li, Xiaolong Cheng	
Maintainer Xiaolong Cheng <xiaolongcheng1120@gmail.com></xiaolongcheng1120@gmail.com>	
Pescription scMAGeCK is a computational model to identify genes associated with multiple expression protypes from CRISPR screening coupled with single-cell RNA sequencing data (CROP-seq)	he-
cicense BSD_2_clause	
iocViews CRISPR, SingleCell, RNASeq, PooledScreens, Transcriptomics, GeneExpression, Regression	
ReedsCompilation yes	
mports Seurat, stats, utils	
uggests knitr, rmarkdown	
'ignetteBuilder knitr	
it_url https://git.bioconductor.org/packages/scMAGeCK	
it_branch RELEASE_3_12	
it_last_commit d710baa	
it_last_commit_date 2020-10-27	
Date/Publication 2021-03-29	
R topics documented:	
scMAGeCK-package scmageck_lr scmageck_rra	2 3 4
ndex	5

2 scMAGeCK-package

scMAGeCK-package	Identify genes associated with multiple expression phenotypes in single-cell CRISPR screening data
	o o

Description

scMAGeCK is a computational model to identify genes associated with multiple expression phenotypes from CRISPR screening coupled with single-cell RNA sequencing data (CROP-seq)

Details

The DESCRIPTION file: This package was not yet installed at build time.

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

scMAGeCK is a computational model to identify genes associated with multiple expression phenotypes from CRISPR screening coupled with single-cell RNA sequencing data (CROP-seq).scMAGeCK is based on our previous MAGeCK and MAGeCK-VISPR models for pooled CRISPR screens.

The scMAGeCK manuscript can be found at bioRxiv(https://www.biorxiv.org/content/10.1101/658146v1/).

Author(s)

Wei Li, Xiaolong Cheng

Maintainer: Xiaolong Cheng < xiaolongcheng 1120@gmail.com>

Examples

```
### BARCODE file contains cell identity information, generated from
### the cell identity collection step
BARCODE <- system.file("extdata", "barcode_rec.txt", package = "scMAGeCK")
### RDS can be a Seurat object or local RDS file path that contains
### the scRNA-seq dataset
RDS <- system.file("extdata", "singles_dox_mki67_v3.RDS", package = "scMAGeCK")</pre>
### Set RRA executable file path.
    You can generate RRA executable file by following commands:
###
     wget https://bitbucket.org/weililab/scmageck/downloads/RRA_0.5.9.zip
###
     unzip RRA_0.5.9.zip
###
     cd RRA_0.5.9
     make
RRAPATH <- "/Library/RRA_0.5.9/bin/RRA"
target_gene <- "MKI67"</pre>
rra_result <- scmageck_rra(BARCODE=BARCODE, RDS=RDS, GENE=target_gene,</pre>
                           RRAPATH=RRAPATH, LABEL='dox_mki67',
                           NEGCTRL=NULL, KEEPTMP=FALSE,
                           PATHWAY=FALSE, SAVEPATH=NULL)
head(rra_result)
lr_result <- scmageck_lr(BARCODE=BARCODE, RDS=RDS, LABEL='dox_scmageck_lr',</pre>
        NEGCTRL = 'NonTargetingControlGuideForHuman', PERMUTATION = 1000,
        SAVEPATH=NULL, LAMBDA=0.01)
```

scmageck_lr 3

```
lr_score <- lr_result[1]
lr_score_pval <- lr_result[2]
head(lr_score_pval)</pre>
```

scmageck_lr

Use linear regression to test the association of gene knockout with all possible genes

Description

echo "Use linear regression to test the association of gene knockout with all possible genes"

Usage

```
scmageck_lr(BARCODE, RDS, NEGCTRL, SELECT_GENE=NULL, LABEL = NULL,
PERMUTATION = NULL, SAVEPATH = "./",LAMBDA=0.01,GENE_FRAC=0.01)
```

Arguments

BARCODE A txt file to include cell identity information, generated from the cell identity

collection step.

RDS A Seurat object or local RDS file path that contains the scRNA-seq dataset. Note

that the dataset has to be normalized and scaled.

NEGCTRL The name of the genes (separated by ",") served as negative controls.

SELECT_GENE The list of genes for regression. By default, all genes in the table are subject to

regression.

LABEL The label of the output file.

PERMUTATION The number of permutations for p value calculation.

SAVEPATH The save path of result. Default save path is the current working directory. If

you don't need save the result, set SAVEPATH as NULL.

LAMBDA A paramter for the LR model for ridge regression. Default: 0.01.

GENE_FRAC A paramter for filtering low expressed genes. By default, only genes that have

expressions in at least that fractions of cells are kept. Default: 0.01.

Value

The result for object RDS

Examples

4 scmageck_rra

scmageck_rra	Use RRA to test the association of gene knockout with certain marker expression

Description

echo "Use RRA to test the association of gene knockout with certain marker expression"

Usage

```
scmageck_rra(BARCODE, RDS, GENE, RRAPATH = NULL, LABEL = NULL, NEGCTRL = NULL,
KEEPTMP = FALSE, PATHWAY = FALSE, SAVEPATH = "./")
```

Arguments

BARCODE	A txt file to include cell identity information, generated from the cell identity collection step.
RDS	A Seurat object or local RDS file path that contains the scRNA-seq dataset. Note that the dataset has to be normalized and scaled.
GENE	Genes whose expressions are to be tested. Multiple genes can be provided, separated by ",". For example, "MKI67,TP53"
RRAPATH	The path to the RRA program, if RRA cannot be found in the PATH environment variable.
LABEL	The label of the output file.
NEGCTRL	The name of the negative control gene. For example, "NonTargetingControl-GuideForHuman". Default is NULL (do not use any negative controls).
KEEPTMP	Keep temporary files.
PATHWAY	Treat genes in –GENE option as a pathway. In other words, the averaged expression of these genes will be used for testing.
SAVEPATH	The save path of result. Default save path is the current working directory. If you don't need save the result, set SAVEPATH as NULL.

Value

The result for object RDS

Examples

Index

```
* package
        scMAGeCK-package, 2

scmageck (scMAGeCK-package), 2
scMAGeCK-package, 2
scmageck_lr, 3
scmageck_rra, 4
```