SCEPTRE vs Seurat STAT3

2023-03-14

Load packages.

```
library(Seurat)
## Attaching SeuratObject
library(SeuratData)
## -- Installed datasets ------ SeuratData v0.2.2 --
## v thp1.eccite 3.1.5
## ------ Key ------ Key ------
## v Dataset loaded successfully
## > Dataset built with a newer version of Seurat than installed
## (?) Unknown version of Seurat installed
library(ggplot2)
library(patchwork)
library(scales)
library(dplyr)
## Attaching package: 'dplyr'
## The following objects are masked from 'package:stats':
##
##
      filter, lag
## The following objects are masked from 'package:base':
##
##
      intersect, setdiff, setequal, union
library(reshape2)
library(mixtools)
## mixtools package, version 2.0.0, Released 2022-12-04
## This package is based upon work supported by the National Science Foundation under Grant No. SES-051
```

```
library(stringr)
library(enrichR)
## Welcome to enrichR
## Checking connection ...
## Enrichr ... Connection is Live!
## FlyEnrichr ... Connection is available!
## WormEnrichr ... Connection is available!
## YeastEnrichr ... Connection is available!
## FishEnrichr ... Connection is available!
## OxEnrichr ... Connection is available!
library(kableExtra)
##
## Attaching package: 'kableExtra'
## The following object is masked from 'package:dplyr':
##
##
       group_rows
library(varhandle)
#using absolute paths to download results since files exist on github
code_dir = .get_config_path("LOCAL_CODE_DIR")
data.dir = paste0(code_dir,"/sceptre2-manuscript/writeups/papalexi_analysis/")
sceptre_path = paste0(data.dir,
                  'sceptre_full_mrna_results_with_effect_size.rds')
seurat_stat3_path = paste0(data.dir,
                   'seurat_STAT3_results.rds')
seurat_nfkbia_path = paste0(data.dir,
                   'seurat_NFKBIA_results.rds')
seurat_etv7_path = paste0(data.dir,
                   'seurat_ETV7_results.rds')
sceptre = readRDS(sceptre_path)
seurat_stat3 = readRDS(seurat_stat3_path)
seurat nfkbia = readRDS(seurat nfkbia path)
seurat_etv7 = readRDS(seurat_etv7_path)
#get significant genes
seurat_etv7_sig = rownames(subset(seurat_etv7,p_val_adj < 0.1))</pre>
seurat_stat3_sig = rownames(subset(seurat_stat3,p_val_adj < 0.1))</pre>
seurat_nfkbia_sig = rownames(subset(seurat_nfkbia,p_val_adj < 0.1))</pre>
```

We see that Seurat-DE finds almost no significant genes after a bonferroni adjustment.

```
#get number of response genes analyzed per perturbation
grna = unfactor(unique(sceptre$grna_group))
```

Table 1: SCEPTRE: Number of Significant Genes vs Perturbation (FDR = 0.1)

grna	num_significant
CUL3	635
CMTM6	1
ATF2	0
BRD4	748
CAV1	0
CD86	0
ETV7	1
IFNGR1	5472
IFNGR2	5584
IRF1	2579
IRF7	1
JAK2	5574
MARCH8	0
MYC	171
NFKBIA	2
PDCD1LG2	0
POU2F2	2
SMAD4	4389
SPI1	26
STAT1	4790
STAT2	346
STAT3	2
STAT5A	0
TNFRSF14	1
UBE2L6	1

```
n = c()
for(val in grna){
    sceptre_stat3 = subset(sceptre,grna_group == val)
    sceptre_stat3$p_value_adj = p.adjust(sceptre_stat3$p_value,method = "BH")
    sceptre_stat3 = subset(sceptre_stat3,p_value_adj < 0.1)
    n = c(n,nrow(sceptre_stat3))
}</pre>
```

```
summary_sceptre = data.frame(grna = grna, num_significant = n)
results = kable(summary_sceptre,booktabs = TRUE, linesep = "",
caption = "SCEPTRE: Number of Significant Genes vs Perturbation (FDR = 0.1)")
kable_styling(results,position = "center")
```