## Figure-S4C-GALNT15-Dot-Plot.R

## sokole

2024-07-28

```
# This Script Generates Figure S4C
# Script By: Eishani Kumar Sokolowski
# Empty the environment & suppress warnings
rm(list = ls())
options(warn=-1)
# Loading libraries
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(Seurat)
## Loading required package: SeuratObject
## Loading required package: sp
## 'SeuratObject' was built under R 4.4.0 but the current version is
## 4.4.1; it is recomended that you reinstall 'SeuratObject' as the ABI
## for R may have changed
##
## Attaching package: 'SeuratObject'
## The following object is masked from 'package:base':
##
##
       intersect
```

```
library(ggplot2)
library(tidyverse)
```

```
## — Conflicts — tidyverse_conflicts() —
## * dplyr::filter() masks stats::filter()
## * dplyr::lag() masks stats::lag()
## i Use the conflicted package (<http://conflicted.r-lib.org/>) to force all conflicts
to become errors
```

```
library(ggpubr)
# Load the seurat object
combined <- readRDS("./Combined_Islet_150_Islet_162_Islet_168_Islet_67_Islet_116_Islet_1</pre>
17_Cluster_All_Cell_Type_Identities_Finalized.rds")
# Assigning Idents
Idents(combined) <- "Cell Type"</pre>
# Keeping only alpha and beta cells
combined <- subset(x = combined, idents = "Acinar", invert=TRUE)</pre>
combined <- subset(x = combined, idents = "Schwann", invert=TRUE)</pre>
combined <- subset(x = combined, idents = "Immune", invert=TRUE)</pre>
combined <- subset(x = combined, idents = "Endothelial", invert=TRUE)</pre>
combined <- subset(x = combined, idents = "Activated Stellate", invert=TRUE)</pre>
combined <- subset(x = combined, idents = "Proliferating Alpha", invert=TRUE)</pre>
combined <- subset(x = combined, idents = "Ductal", invert=TRUE)</pre>
combined <- subset(x = combined, idents = "Stellate", invert=TRUE)</pre>
combined <- subset(x = combined, idents = "Delta", invert=TRUE)</pre>
combined <- subset(x = combined, idents = "Gamma", invert=TRUE)</pre>
# Making a new column
combined$Cell_Type_HTO_Label <- paste0(combined$Cell_Type,"_",combined$HTO_Label)</pre>
# Making factors
combined$Cell_Type_HTO_Label <- factor(combined$Cell_Type_HTO_Label, levels=c("Beta_24 H</pre>
r DMSO",
                                                                                   "Beta 24 H
r Thapsigargin",
                                                                                   "Beta 24 H
r Untreated",
                                                                                   "Beta_24 H
r Cytokines",
                                                                                   "Alpha 24
Hr DMSO",
                                                                                   "Alpha 24
Hr Thapsigargin",
                                                                                   "Alpha 24
Hr Untreated",
                                                                                   "Alpha 24
Hr Cytokines"))
# Assigning Idents
Idents(combined) <- "Cell Type HTO Label"</pre>
# Making the dot plot
p <- DotPlot(object = combined, features = "GALNT15", col.min = -0.5, col.max = 2) +
  theme(axis.text.x = element_text(angle=90)) +
  scale color gradient2(low="white", mid = "grey80", high="red") + coord flip()
```

```
## Scale for colour is already present.
## Adding another scale for colour, which will replace the existing scale.
```

```
# View
ggarrange(p, ncol = 1, nrow = 1)
```

