EC_figure

Ben Thomson

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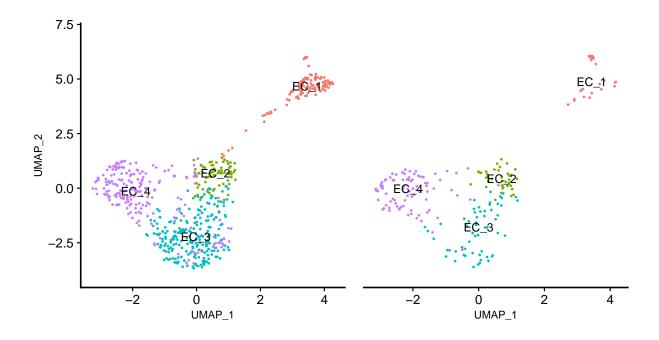
```
ec.dataset <- readRDS("../data/6wk_endos_10-1-20.rds.gz")
lec.dataset <- readRDS("../data/6wk_LECs_10-1-20.rds.gz")

panel.a <- RenameIdents(ec.dataset, SC = "EC_1", LEC = "EC_1")

panel.a.fig.wt <- DimPlot(object = subset(panel.a, subset = genotype ==
    "WT"), reduction = "umap", label = TRUE, pt.size = 0.5, label.size = 4) +
    NoLegend() + coord_fixed() + theme(aspect.ratio = 1) + theme(axis.line = element_line(color = "black"
    size = 0.7)) + theme(axis.title = element_text(size = 10)) +
    ylim(c(-4, 7))

panel.a.fig.ko <- DimPlot(object = subset(panel.a, subset = genotype ==
    "KO"), reduction = "umap", label = TRUE, pt.size = 0.5, label.size = 4) +
    NoLegend() + coord_fixed() + theme(aspect.ratio = 1) + theme(axis.line = element_line(color = "black"
    size = 0.7)) + theme(axis.line.y = element_blank(), axis.ticks.y = element_blank(),
    axis.text.y = element_blank()) + theme(axis.title.x = element_text(size = 10)) +
    theme(axis.title.y = element_blank()) + ylim(c(-4, 7))</pre>
```

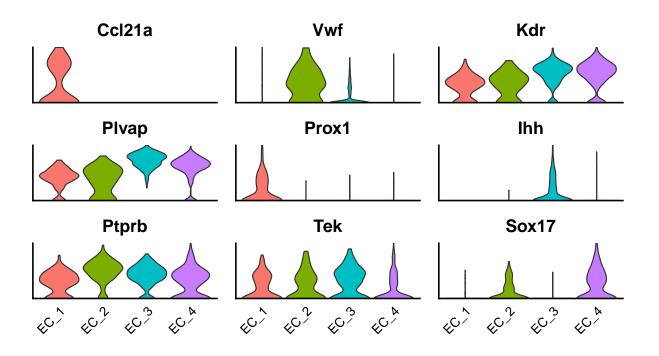
panel.a.fig.wt + panel.a.fig.ko



```
pdf(file = "../figures/EC_figure_a.pdf", width = 5, height = 2)
plot(panel.a.fig.wt + panel.a.fig.ko)
dev.off()
## pdf
##
     2
panel.b <- RenameIdents(ec.dataset, SC = "EC_1", LEC = "EC_1")</pre>
plot.list.b <- VlnPlot(panel.b, assay = "RNA", features = c("Ccl21a",</pre>
  "Vwf", "Kdr", "Plvap", "Prox1", "Ihh", "Ptprb", "Tek", "Sox17"),
  pt.size = 0, combine = F)
for (i in seq_along(plot.list.b)) {
  plot.list.b[[i]] <- plot.list.b[[i]] + theme(axis.ticks = element_blank(),</pre>
    axis.text.y = element_blank(), axis.title.x = element_blank()) +
    theme(title = element_text(family = "sans", face = "italic")) +
    theme(axis.line = element_line(size = 0.7)) + NoLegend() +
    scale_y_continuous(expand = c(0, 0)) + theme(aspect.ratio = 0.3) +
    theme(axis.title.y = element_blank())
  if (i > (length(plot.list.b) - 3)) {
    plot.list.b[[i]] <- plot.list.b[[i]] + theme(axis.text.x = element_text())</pre>
```

```
} else {
    plot.list.b[[i]] <- plot.list.b[[i]] + theme(axis.text.x = element_blank())
}

wrap_plots(plot.list.b)</pre>
```



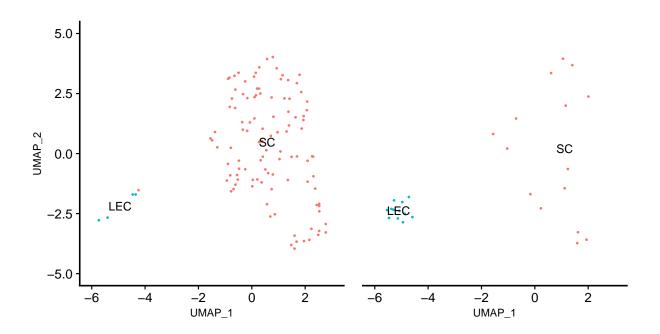
```
pdf(file = "../figures/EC_figure_b.pdf", width = 5, height = 5)
plot(wrap_plots(plot.list.b))
dev.off()

## pdf
## 2

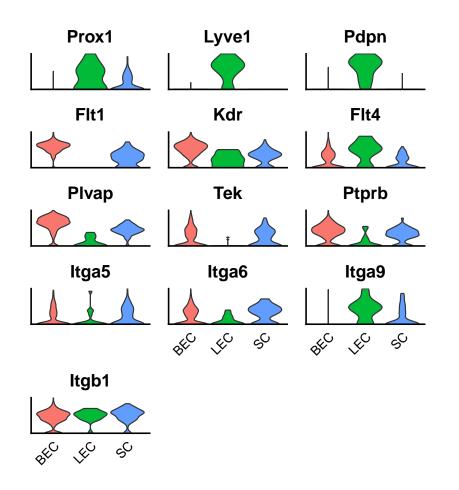
panel.c <- lec.dataset

panel.c.fig.wt <- DimPlot(object = subset(panel.c, subset = genotype ==
    "WT"), reduction = "umap", label = TRUE, pt.size = 0.5, label.size = 4) +
    NoLegend() + coord_fixed() + theme(aspect.ratio = 1) + theme(axis.line = element_line(color = "black"
    size = 0.7)) + theme(axis.title = element_text(size = 10)) +
    xlim(c(-6, 3)) + ylim(c(-5, 5))</pre>
```

```
panel.c.fig.ko <- DimPlot(object = subset(panel.c, subset = genotype ==
   "KO"), reduction = "umap", label = TRUE, pt.size = 0.5, label.size = 4) +
   NoLegend() + coord_fixed() + theme(aspect.ratio = 1) + theme(axis.line = element_line(color = "black"
   size = 0.7)) + theme(axis.line.y = element_blank(), axis.ticks.y = element_blank(),
   axis.text.y = element_blank()) + theme(axis.title.x = element_text(size = 10)) +
   theme(axis.title.y = element_blank()) + xlim(c(-6, 3)) +
   ylim(c(-5, 5))</pre>
```

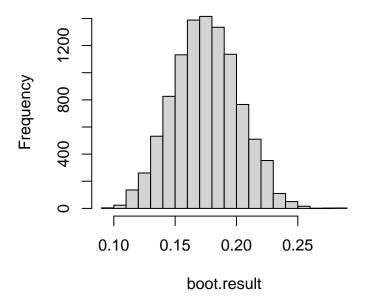


```
plot.list.d <- VlnPlot(panel.d, assay = "RNA", features = c("Prox1",</pre>
  "Lyve1", "Pdpn", "Flt1", "Kdr", "Flt4", "Plvap", "Tek", "Ptprb",
  "Itga5", "Itga6", "Itga9", "Itgb1"), pt.size = 0, combine = FALSE)
for (i in seq_along(plot.list.d)) {
  plot.list.d[[i]] <- plot.list.d[[i]] + theme(axis.ticks = element_blank(),</pre>
    axis.text.y = element_blank(), axis.title.x = element_blank()) +
    theme(title = element_text(family = "sans", face = "italic")) +
    theme(axis.line = element_line(size = 0.7)) + NoLegend() +
    scale_y_continuous(expand = c(0, 0)) + theme(aspect.ratio = 0.3) +
    theme(axis.title.y = element_blank())
  if (i > (length(plot.list.d) - 3)) {
    plot.list.d[[i]] <- plot.list.d[[i]] + theme(axis.text.x = element_text())</pre>
  } else {
    plot.list.d[[i]] <- plot.list.d[[i]] + theme(axis.text.x = element_blank())</pre>
  }
}
wrap_plots(plot.list.d, ncol = 3)
```



```
pdf(file = "../figures/EC_figure_d.pdf", width = 5, height = 6)
plot(wrap_plots(plot.list.d, ncol = 3))
dev.off()
## pdf
## 2
boot.data <- FetchData(subset(ec.dataset, subset = genotype ==</pre>
 "WT"), vars = "ident")$ident
ko.data <- FetchData(subset(ec.dataset, subset = genotype ==</pre>
  "KO"), vars = "ident")$ident
pct.gt <- function(data, value) {</pre>
  total <- length(data)</pre>
  selected <- length(data[data == value])</pre>
 return(selected/total)
}
nboots <- 10000
boot.result <- numeric(nboots)</pre>
set.seed(42)
# number of KO endothelial cells
sample.size <- length(ko.data)</pre>
for (i in 1:nboots) {
  boot.sample <- sample(boot.data, sample.size, replace = TRUE)</pre>
  boot.result[i] <- pct.gt(boot.sample, "SC")</pre>
}
hist(boot.result)
```

Histogram of boot.result



```
quantile(boot.result, c(0.025, 0.975))

## 2.5% 97.5%
## 0.1244019 0.2296651

mean(boot.result)
```

[1] 0.1751062

```
bar.data <- data.frame(row.names = c("WT", "KO"))
KO <- pct.gt(ko.data, "SC")
WT <- boot.result

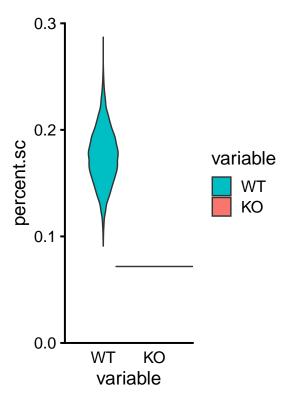
bar.data <- data.frame(WT, KO)
bar.data <- melt(bar.data, value.name = "percent.sc")

p1 <- ggplot(data = bar.data, aes(x = variable, y = percent.sc,
    fill = variable, width = 1.5)) + geom_violin() +

theme(axis.line = element_line(size = 0.7)) + theme(axis.ticks.y = element_line(size = 0.7),
    axis.ticks.x = element_blank()) + theme(aspect.ratio = 2.5)

p1 <- p1 + scale_fill_manual(values = c("#00bfc4", "#f8766d"))
p1 <- p1 + scale_y_continuous(expand = c(0, 0), limits = c(0, 0.3), breaks = c(0, 0.1, 0.2, 0.3))</pre>
```

```
plot(p1)
```

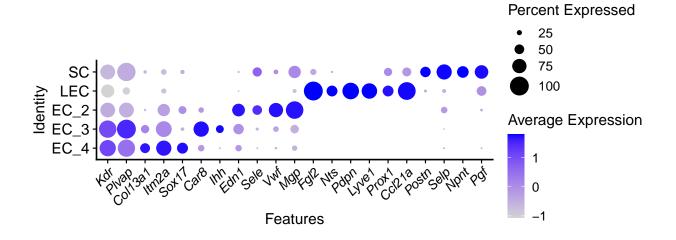


```
pdf(file = "../figures/EC_figure_e.pdf", width = 4, height = 4)
plot(p1)
dev.off()

## pdf
## 2

panel.g <- ec.dataset
panel.g@active.ident <- factor(x = panel.g@active.ident, levels = c("EC_4",
    "EC_3", "EC_2", "LEC", "SC"))
genelist <- read.csv("../figures/genes_for_ec_heatmap.csv", stringsAsFactors = F,
    fileEncoding = "UTF-8-BOM")$gene

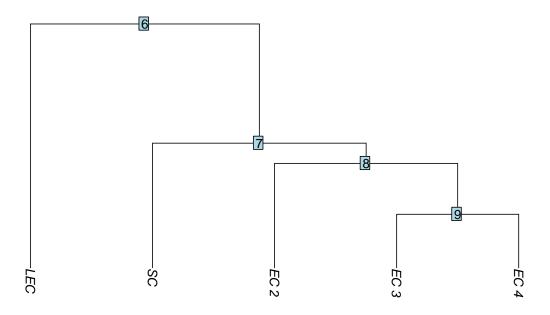
dot <- DotPlot(panel.g, features = genelist, assay = "RNA", dot.scale = 7.5,
    dot.min = 0.05) + theme(axis.line = element_line(size = 0.7)) +
    theme(aspect.ratio = 0.25) + theme(axis.text.x = element_text(angle = 45,
    face = "italic", vjust = 1, hjust = 1)) + theme(axis.text.y = element_text(size = 14))
plot(dot)</pre>
```



```
pdf(file = "../figures/EC_figure_g.pdf", width = 8, height = 4)
plot(dot)
dev.off()

## pdf
## 2

panel.f <- BuildClusterTree(ec.dataset, assay = "RNA", reorder = T)
PlotClusterTree(panel.f)</pre>
```

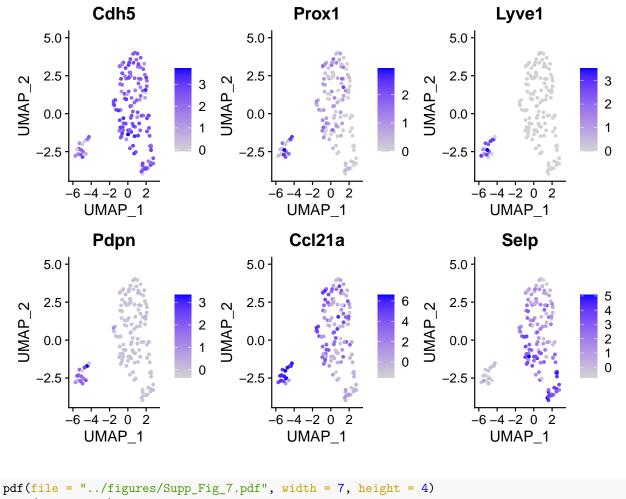


```
pdf(file = "../figures/EC_figure_f.pdf", width = 4, height = 4)
PlotClusterTree(panel.f)
dev.off()
```

pdf ## 2

The data from Panel C was also used to generate Supplemental figure 7.

```
supp.fig.7 <- FeaturePlot(panel.c, features = c("Cdh5", "Prox1",
    "Lyve1", "Pdpn", "Ccl21a", "Selp"), ncol = 3)
plot(supp.fig.7)</pre>
```



```
pdf(file = "../figures/Supp_Fig_7.pdf", width = 7, height = 4)
plot(supp.fig.7)
dev.off()
```

pdf ## 2