First TCGA analysis

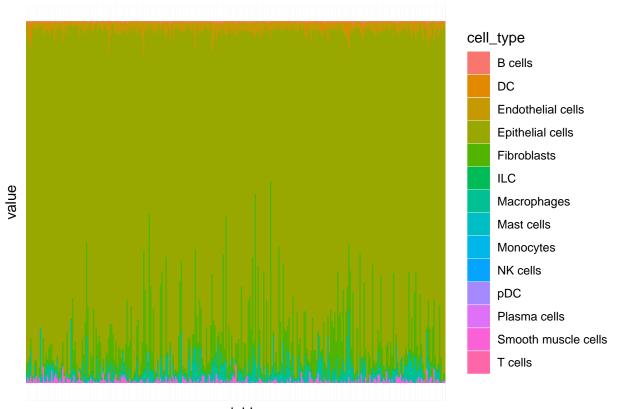
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Subtypes
Having run the TCGA RNA-seq data through Bayes Prism, this notebook compares the samples' cell type composition with their subtype annotations from the Way pipeline and the patients' survival status/time.
<pre>suppressPackageStartupMessages({ library(data.table) library(SingleCellExperiment) library(dplyr) library(yaml) library(stringr) library(ggplot2) library(survival) library(ggfortify) })</pre>
<pre>params <- read_yaml("//config.yml") data_path <- params\$data_path local_data_path <- params\$local_data_path plot_path <- params\$plot_path</pre>
<pre>tcga <- fread(paste(local_data_path, "deconvolution_output",</pre>
<pre>## Warning in melt.data.table(tcga): id.vars and measure.vars are internally ## guessed when both are 'NULL'. All non-numeric/integer/logical type columns are ## considered id.vars, which in this case are columns [cell_type,]. Consider ## providing at least one of 'id' or 'measure' vars in future.</pre>

Cell composition



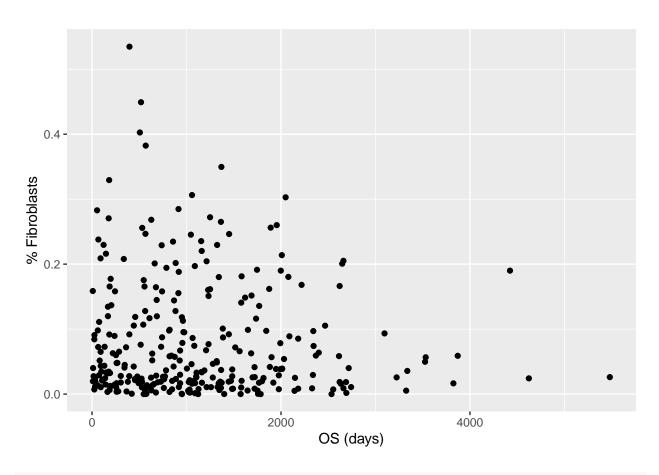
variable

```
# Switch so cell types are columns and samples are rows for easier analysis
cell_types <- tcga$cell_type

tcga$cell_type <- NULL
tcga_t <- t(as.matrix(tcga))
colnames(tcga_t) <- cell_types
tcga_t <- as.data.frame(tcga_t)</pre>
```

Cell composition by survival

```
# Load survival data
tcga_survival <- fread(paste(local_data_path, "TCGA", "TCGA_OV_survival.tsv",</pre>
                             sep = "/")
tcga_patients <- str_extract(colnames(tcga), "TCGA-\\W\\W-\\W\\W\\W")
tcga_survival <- subset(tcga_survival, tcga_survival$bcr_patient_barcode %in%
                            tcga_patients)
# Combine survival data with %
tcga_t$bcr_patient_barcode <- tcga_patients</pre>
tcga_master <- full_join(tcga_survival, tcga_t)</pre>
## Joining, by = "bcr_patient_barcode"
tcga_master$Immune <- tcga_master$Macrophages + tcga_master$Monocytes + tcga_master$`Plasma cells` +
  tcga_master$DC + tcga_master$`NK cells` + tcga_master$PDC + tcga_master$`B cells` + tcga_master$ILC +
 tcga_master$`Mast cells`
g <- ggplot(tcga_master, mapping = aes(x=tcga_master$OS.time, y=tcga_master$Fibroblasts)) +
  geom_point() + xlab("OS (days)") + ylab("% Fibroblasts")
plotfile <- paste(plot_path, "evaluation_plots", "TCGA_survival_by_fibroblasts.png", sep = "/")</pre>
png(filename = plotfile); g; dev.off()
## Warning: Removed 1 rows containing missing values (`geom_point()`).
## pdf
##
## Warning: Removed 1 rows containing missing values (`geom_point()`).
```

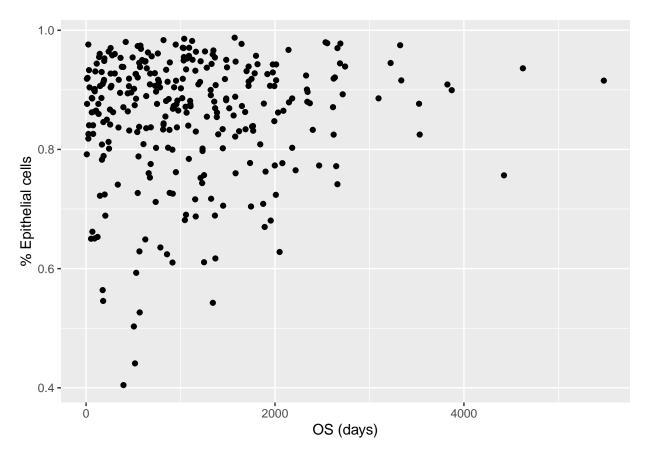


```
g <- ggplot(tcga_master, mapping = aes(x=tcga_master$0S.time, y=tcga_master$`Epithelial cells`)) +
    geom_point() + xlab("OS (days)") + ylab("% Epithelial cells")
plotfile <- paste(plot_path, "evaluation_plots", "TCGA_survival_by_epithelial.png", sep = "/")
png(filename = plotfile); g; dev.off()

## Warning: Removed 1 rows containing missing values (`geom_point()`).

## pdf
## 2</pre>
```

Warning: Removed 1 rows containing missing values (`geom_point()`).

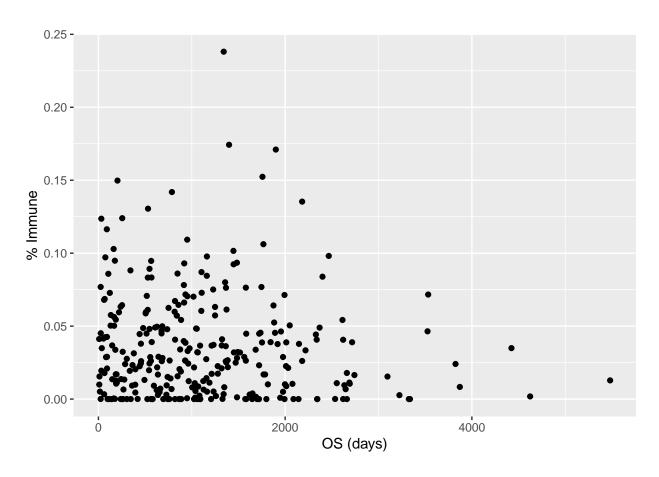


```
g <- ggplot(tcga_master, mapping = aes(x=tcga_master$OS.time, y=tcga_master$Immune)) +
    geom_point() + xlab("OS (days)") + ylab("% Immune")
plotfile <- paste(plot_path, "evaluation_plots", "TCGA_survival_by_immune.png", sep = "/")
png(filename = plotfile); g; dev.off()

## Warning: Removed 1 rows containing missing values (`geom_point()`).

## pdf
## pdf
## 2</pre>
```

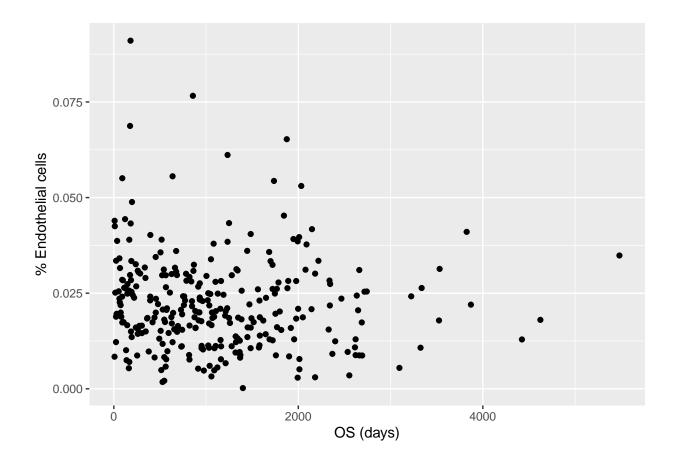
Warning: Removed 1 rows containing missing values (`geom_point()`).



```
g <- ggplot(tcga_master, mapping = aes(x=tcga_master$0S.time, y=tcga_master$Endothelial cells")) +
    geom_point() + xlab("0S (days)") + ylab("% Endothelial cells")
plotfile <- paste(plot_path, "evaluation_plots", "TCGA_survival_by_endothelial_cells.png", sep = "/")
png(filename = plotfile); g; dev.off()

## Warning: Removed 1 rows containing missing values (`geom_point()`).</pre>
## pdf
## 2
```

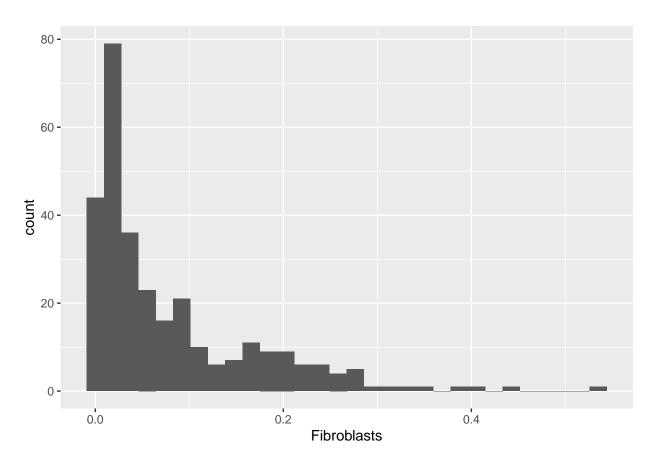
Warning: Removed 1 rows containing missing values (`geom_point()`).



Kaplan Meier curves

```
# Put the samples into quartiles based on fibroblast content
ggplot(tcga_master, mapping = aes(x=Fibroblasts)) + geom_histogram()
```

`stat_bin()` using `bins = 30`. Pick better value with `binwidth`.



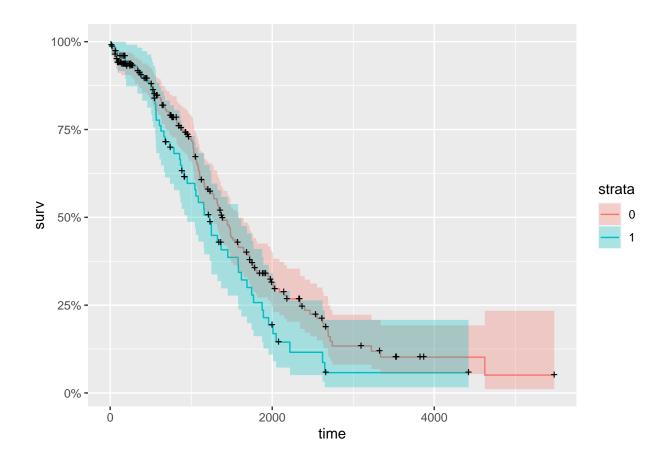
```
quantiles <- quantile(tcga_master$Fibroblasts)
q1 <- quantiles[2]
q3 <- quantiles[4]
tcga_master$high_fibro <- ifelse(tcga_master$Fibroblasts > q3, 1, 0)

# Get Kaplan-Meier curves
km <- Surv(time = tcga_master$0S.time, event = tcga_master$0S)
km_treatment<-survfit(km~high_fibro,data=tcga_master,type='kaplan-meier',conf.type='log')

plotfile <- paste(plot_path, "evaluation_plots", "TCGA_KaplanMeier_fibroblasts.png", sep = "/")
png(filename = plotfile)
autoplot(km_treatment)
dev.off()

## pdf
## 2</pre>
```

autoplot(km_treatment)

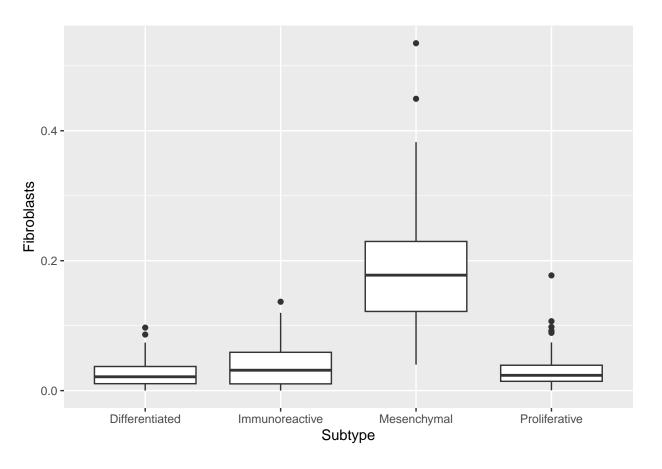


Subtypes

```
# Get rid of samples that don't have a subtype label
tcga_t <- subset(tcga_t, !is.na(tcga_t$Subtype))

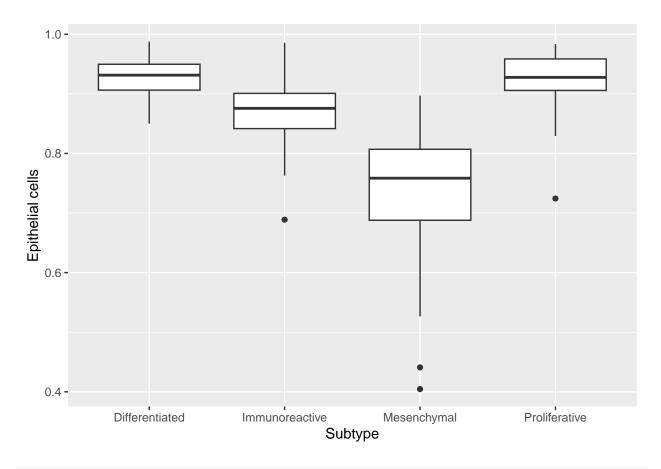
# Compare cell type proportions of subtypes
g <- ggplot(tcga_t, mapping = aes(x=Subtype, y=Fibroblasts)) + geom_boxplot()
plotfile <- paste(plot_path, "evaluation_plots", "TCGA_fibroblasts_by_subtype.png", sep = "/")
png(filename = plotfile); g; dev.off()

## pdf
## 2</pre>
g
```



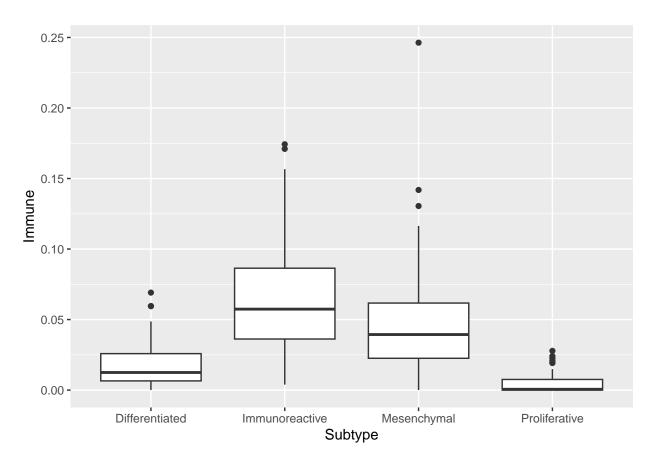
```
g <- ggplot(tcga_t, mapping = aes(x=Subtype, y=`Epithelial cells`)) + geom_boxplot()
plotfile <- paste(plot_path, "evaluation_plots", "TCGA_epithelial_by_subtype.png", sep = "/")
png(filename = plotfile); g; dev.off()

## pdf
## 2</pre>
```



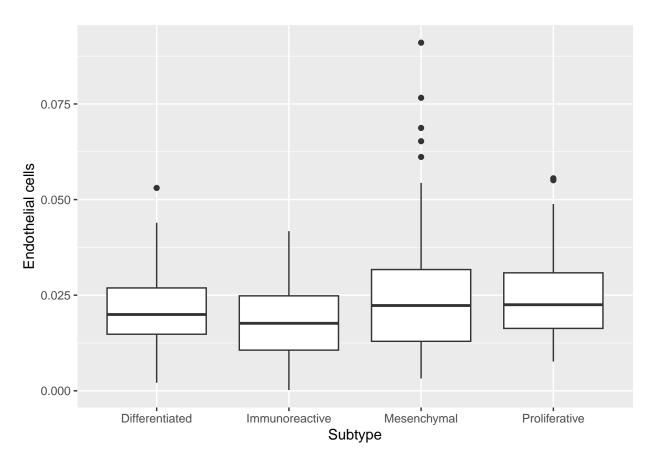
```
g <- ggplot(tcga_t, mapping = aes(x=Subtype, y=Immune)) + geom_boxplot()
plotfile <- paste(plot_path, "evaluation_plots", "TCGA_immune_by_subtype.png", sep = "/")
png(filename = plotfile); g; dev.off()</pre>
```

pdf ## 2



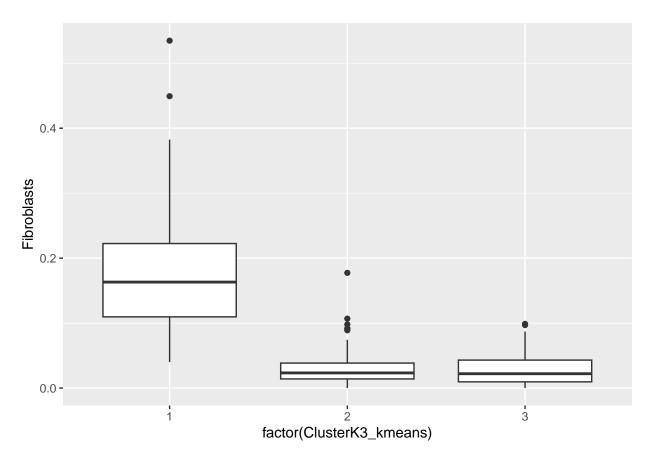
```
g <- ggplot(tcga_t, mapping = aes(x=Subtype, y=`Endothelial cells`)) + geom_boxplot()
plotfile <- paste(plot_path, "evaluation_plots", "TCGA_endothelial_by_subtype.png", sep = "/")
png(filename = plotfile); g; dev.off()

## pdf</pre>
```



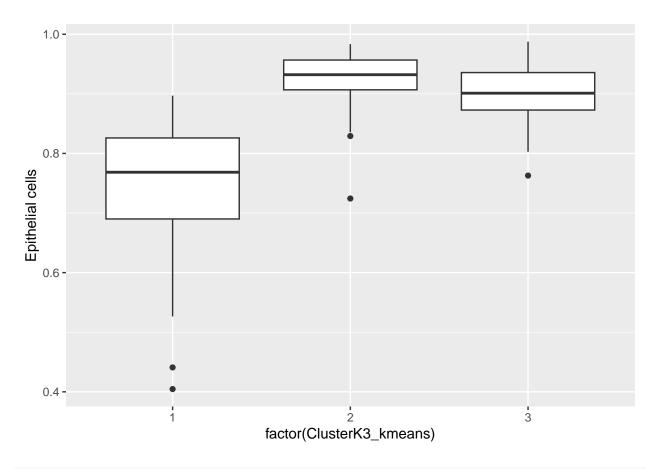
```
# Compare cell type proportions of subtypes for k=3
g <- ggplot(tcga_t, mapping = aes(x=factor(ClusterK3_kmeans), y=Fibroblasts)) + geom_boxplot()
plotfile <- paste(plot_path, "evaluation_plots", "TCGA_fibroblasts_by_subtype_k3.png", sep = "/")
png(filename = plotfile); g; dev.off()

## pdf
## pdf
## 2</pre>
```



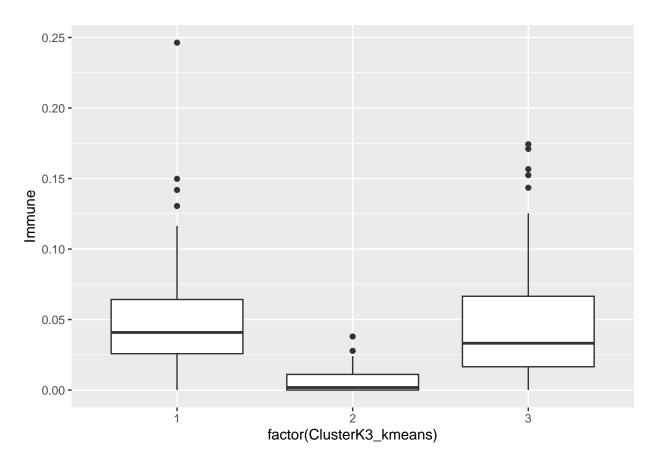
```
g <- ggplot(tcga_t, mapping = aes(x=factor(ClusterK3_kmeans), y=`Epithelial cells`)) + geom_boxplot()
plotfile <- paste(plot_path, "evaluation_plots", "TCGA_epithelial_by_subtype_k3.png", sep = "/")
png(filename = plotfile); g; dev.off()</pre>
```

pdf ## 2



```
g <- ggplot(tcga_t, mapping = aes(x=factor(ClusterK3_kmeans), y=Immune)) + geom_boxplot()
plotfile <- paste(plot_path, "evaluation_plots", "TCGA_immune_by_subtype_k3.png", sep = "/")
png(filename = plotfile); g; dev.off()</pre>
```

pdf ## 2



```
g <- ggplot(tcga_t, mapping = aes(x=factor(ClusterK3_kmeans), y=`Endothelial cells`)) + geom_boxplot()
plotfile <- paste(plot_path, "evaluation_plots", "TCGA_endothelial_by_subtype_k3.png", sep = "/")
png(filename = plotfile); g; dev.off()
## pdf</pre>
```

