Final Project 3

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```
setwd("/Users/annieglenning/Documents/Dartmouth/SU24/QBS_103/Data") # set my working directory to where
original_gene_data <- read.table("QBS103_GSE157103_genes.csv", header = TRUE, sep = ",") # reading in t
series_data <- read.table("QBS103_GSE157103_series_matrix.csv", header = TRUE, sep = ",") # reading in

# transposing the gene data
original_gene_data <- as.data.frame(original_gene_data)
if (is.character(original_gene_data[1, 1])) {
   gene_data <- t(original_gene_data)
   colnames(gene_data) <- gene_data[1, ]
   gene_data <- as.data.frame(gene_data)
}</pre>
```

Identify one gene, one continuous covariate, and two categorical covariates

```
AAGAB <- as.numeric(gene_data$AAGAB) # one gene
```

Info for Table of Summary Statistics

```
# Install table1 package if not already installed
#install.packages("table1")

# Load required packages
library(table1)

##
## Attaching package: 'table1'

## The following objects are masked from 'package:base':
##
## units, units<-
library(dplyr)

##
## Attaching package: 'dplyr'</pre>
```

```
## The following objects are masked from 'package:stats':
##
       filter, lag
##
## The following objects are masked from 'package:base':
##
##
       intersect, setdiff, setequal, union
# Example dataset preparation (you should replace this with your actual data)
# Assuming `data` is your data frame and contains the relevant variables
# Convert categorical variables to factors
#table_data <- series_data %>%
# mutate(sex = factor(sex),
          disease_status = factor(disease_status),
#
          icu_status = factor(icu_status))  # Add another categorical variable here
# Example of selecting 3 continuous variables and 3 categorical variables
#continuous_vars <- c("age", "ferritin.ng.ml.", "procalcitonin.ng.ml..")</pre>
#categorical_vars <- c("sex", "disease_status", "icu_status")</pre>
# Define a custom label for your variables if desired
#labels <- list(
# age = "Age (years)",
# ferritin.ng.ml. = "Ferritin (ng/ml)",
# procalcitonin.ng.ml.. = "Procalcitonin (ng/ml)",
\# sex = "Sex",
# disease status = "Disease Status",
# icu_status = "ICU Status"
#)
# Create the table stratified by a categorical variable, e.g., `disease_status`
#table <- table1(~ age + ferritin.ng.ml. + procalcitonin.ng.ml.. + sex + icu_status | disease_status,
                 data = table_data,
#
                 render.categorical = function(x) #sprintf("%d (%0.1f%%)", sum(x), mean(x) * 100),
#
                 render.continuous = function(x) {
#
                     if (length(x) \le 5)  {
                         sprintf("\%0.1f \ [\%0.1f, \ \%0.1f]", \ median(x), \ quantile(x, \ 0.25), \ quantile(x, \ 0.75)
#
#
                     } else {
#
                         sprintf("%0.1f(%.1f)", mean(x), sd(x))
                     7
#
#
                 },
#
                 overall = FALSE, # Set to TRUE if you want overall summary stats
#
                 topclass = "Rtable1-zebra")
```

sex

```
# percent
#length(series_data$age[series_data$sex == ' male'])/length(series_data$age)*100
# percent
#length(series_data$age[series_data$sex == ' female'])/length(series_data$age)*100
#length(series_data$age[series_data$sex == ' unknown'])/length(series_data$age)*100
```

```
#mean(as.numeric(series_data$age[series_data$sex == ' male']), na.rm = TRUE) # note getting rid of >89
#sd(as.numeric(series_data$age[series_data$sex == ' male']), na.rm = TRUE)
#mean(as.numeric(series_data$age[series_data$sex == ' female']), na.rm = TRUE)
#sd(as.numeric(series_data$age[series_data$sex == ' female']), na.rm = TRUE)
#mean(as.numeric(series_data$age[series_data$sex == ' unknown']), na.rm = TRUE)
#sd(as.numeric(series_data$age[series_data$sex == ' unknown']), na.rm = TRUE)
#mean(as.numeric(series_data$ferritin.ng.ml.[series_data$sex == ' male']), na.rm = TRUE) # 11 unknown
#sd(as.numeric(series_data$ferritin.ng.ml.[series_data$sex == ' female']), na.rm = TRUE)
#mean(as.numeric(series_data$ferritin.ng.ml.[series_data$sex == ' female']), na.rm = TRUE)
#series_data$ferritin.ng.ml.[series_data$sex == ' female']), na.rm = TRUE)
#series_data$ferritin.ng.ml.[series_data$sex == ' female']), na.rm = TRUE)
#series_data$ferritin.ng.ml.[series_data$sex == ' unknown'] # 1 unknown
```

icu_status

```
#length(series_data$age[series_data$icu_status == ' yes'])/length(series_data$age) *100
#length(series_data$age[series_data$icu_status == ' no'])/length(series_data$age) *100

#mean(as.numeric(series_data$age[series_data$icu_status == ' yes']), na.rm = TRUE)
#sd(as.numeric(series_data$age[series_data$icu_status == ' yes']), na.rm = TRUE)
#mean(as.numeric(series_data$age[series_data$icu_status == ' no'] ), na.rm = TRUE) # gets rid of " :" a
#sd(as.numeric(series_data$age[series_data$icu_status == ' no']), na.rm = TRUE)

#mean(as.numeric(series_data$ferritin.ng.ml.[series_data$icu_status == ' yes']), na.rm = TRUE) # 7 unk
#sd(as.numeric(series_data$ferritin.ng.ml.[series_data$icu_status == ' yes']), na.rm = TRUE)
#mean(as.numeric(series_data$ferritin.ng.ml.[series_data$icu_status == ' no']), na.rm = TRUE) # 9 unkn
#sd(as.numeric(series_data$ferritin.ng.ml.[series_data$icu_status == ' no']), na.rm = TRUE)
```

disease_status

```
#length(series_data$age[series_data$disease_status == "disease state: COVID-19"])/length(series_data$ag
#length(series_data$age[series_data$disease_status == "disease state: non-COVID-19"])/length(series_dat
#mean(as.numeric(series_data$age[series_data$disease_status == "disease state: COVID-19"]), na.rm = TRU.
```

#sd(as.numeric(series_data\$age[series_data\$disease_status == "disease state: COVID-19"]), na.rm = TRUE)
#mean(as.numeric(series_data\$age[series_data\$disease_status == "disease state: non-COVID-19"]), na.rm =
#sd(as.numeric(series_data\$age[series_data\$disease_status == "disease state: non-COVID-19"]), na.rm = T

 $\# mean (as.numeric (series_data\$procalcitonin.ng.ml.. [series_data\$disease_status == "disease state: COVID-19" \\ \# sd (as.numeric (series_data\$procalcitonin.ng.ml.. [series_data\$disease_status == "disease state: COVID-19" \\ \# mean (as.numeric (series_data\$procalcitonin.ng.ml.. [series_data\$disease_status == "disease state: non-COVID-19" \\ \# sd (as.numeric (series_data\$procalcitonin.ng.ml.. [series_data\$disease_status == "disease state: non-COVID-19" \\ \# sd (as.numeric (series_data\$procalcitonin.ng.ml.. [series_data\$disease_status == "disease state: non-COVID-19" \\ \# sd (as.numeric (series_data\$procalcitonin.ng.ml.. [series_data\$disease_status == "disease state: non-COVID-19" \\ \# sd (as.numeric (series_data\$procalcitonin.ng.ml.. [series_data\$disease_status == "disease state: non-COVID-19" \\ \# sd (as.numeric (series_data\$procalcitonin.ng.ml.. [series_data\$disease_status == "disease state: non-COVID-19" \\ \# sd (as.numeric (series_data\$procalcitonin.ng.ml.. [series_data\$disease_status == "disease state: non-COVID-19" \\ \# sd (as.numeric (series_data\$procalcitonin.ng.ml.. [series_data\$disease_status == "disease state: non-COVID-19" \\ \# sd (as.numeric (series_data\$procalcitonin.ng.ml.. [series_data\$disease_status == "disease state: non-COVID-19" \\ \# sd (as.numeric (series_data\$procalcitonin.ng.ml.. [series_data\$disease_status == "disease state: non-COVID-19" \\ \# sd (as.numeric (series_data\$procalcitonin.ng.ml.. [series_data\$disease_status == "disease state: non-COVID-19" \\ \# sd (as.numeric (series_data\$procalcitonin.ng.ml.. [series_data\$disease_status == "disease state: non-COVID-19" \\ \# sd (as.numeric (series_data\$procalcitonin.ng.ml.. [series_data\$disease_status == "disease state: non-COVID-19" \\ \# sd (as.numeric (series_data\$procalcitonin.ng.ml.. [series_data\$disease_status == "disease state: non-COVID-19" \\ \# sd (as.numeric (series_data\$procalcitonin.ng.ml.. [series_data\$procalcitonin.ng.ml.. [series_data\$procalcitonin.ng.ml.. [series_data\$procalcitonin.ng.ml.. [series_data\$procalcitonin.ng.ml.. [series_data\pr

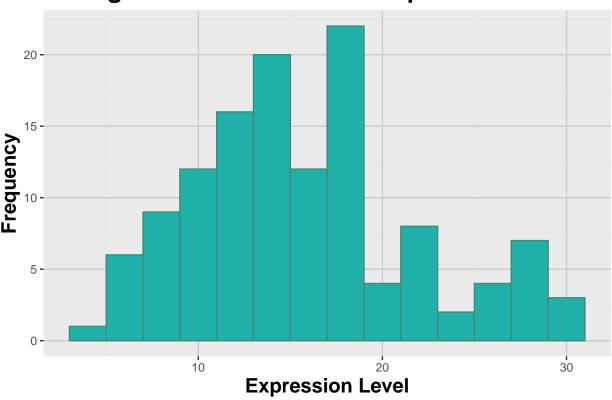
Finalizing Histogram

```
library(tidyverse)
## -- Attaching core tidyverse packages ----
                                                  ----- tidyverse 2.0.0 --
## v forcats 1.0.0
                       v readr
                                    2.1.5
## v ggplot2 3.5.1
                        v stringr
                                    1.5.1
## v lubridate 1.9.3
                        v tibble
                                    3.2.1
## v purrr
              1.0.2
                        v tidyr
                                    1.3.1
## -- Conflicts -----
                                          ## x dplyr::filter() masks stats::filter()
                    masks stats::lag()
## x dplyr::lag()
## i Use the conflicted package (<a href="http://conflicted.r-lib.org/">http://conflicted.r-lib.org/</a>) to force all conflicts to become error
library(ggplot2)
ggplot(gene_data, aes(x = as.numeric(AAGAB))) +
 geom_histogram(binwidth = 2, fill = "lightseagreen", color = "aquamarine4") +
 labs( # labeling the title and axis
   title = "Histogram of AAGAB Gene Expression Levels",
   x = "Expression Level",
   y = "Frequency"
 ) +
 theme(
   plot.title = element_text(size = 19, face = "bold"), # title
   axis.title = element_text(size = 15, face = "bold"), # axis
```

panel.grid.major = element_line(color = "grey80"), # background

panel.grid.minor = element_line(color = "grey90")

Histogram of AAGAB Gene Expression Levels



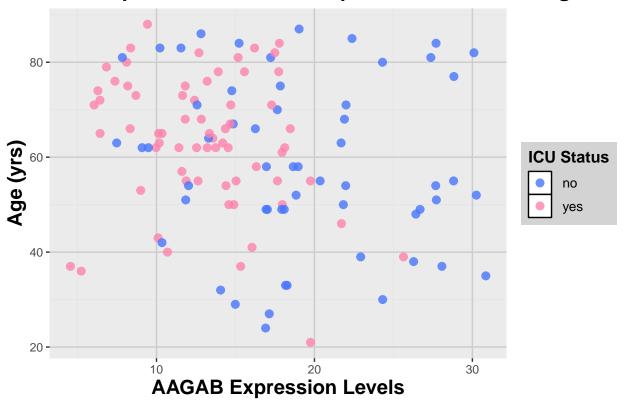
Finalizing Scatterplot

```
ggplot(series_data, aes(x = as.numeric(AAGAB), y = as.numeric(age),
                       color = factor(icu_status))) + # seperarting the points by icu status
  geom_point(size = 2.5, alpha = 0.8) + # size and transparency of the point
  labs( # labeling the title and axis
   title = "Scatterplot of the AAGAB Expression Levels vs. Age",
   x = "AAGAB Expression Levels",
   y = "Age (yrs)",
   color = "ICU Status" # legend title
   ) +
  scale_color_manual(values = c(" yes" = "palevioletred1", " no" = "royalblue1")) + # setting the colo
   plot.title = element_text(size = 16, face = "bold"),
   axis.title = element_text(size = 15, face = "bold"),
   legend.title = element_text(size = 12, face = "bold"), # editing the legend
   legend.text = element_text(size = 10),
   legend.background = element_rect(fill = "lightgray", color = NA),
   legend.key = element_rect(fill = "white", color = "black"),
   panel.grid.major = element_line(color = "grey80"), # background
   panel.grid.minor = element_line(color = "grey90")
```

Warning in FUN(X[[i]], ...): NAs introduced by coercion

Warning: Removed 3 rows containing missing values or values outside the scale range
('geom_point()').

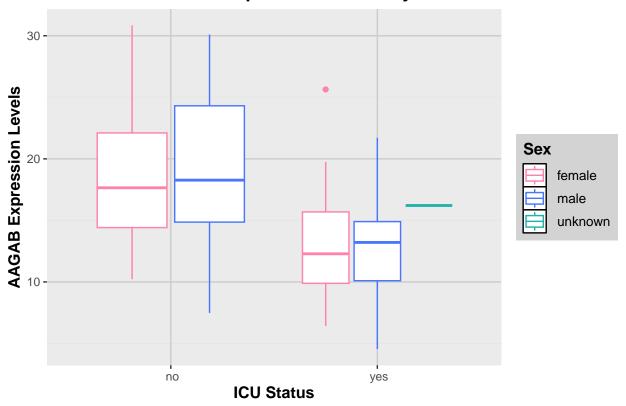
Scatterplot of the AAGAB Expression Levels vs. Age



Finalizing Box Plot

```
ggplot(series_data,aes(x = icu_status, y = as.numeric(AAGAB), color = sex)) +
  geom_boxplot() +
  labs( # labeling the title and axis
   title = "Box Plot of AAGAB Expression Levels by ICU Status and Sex",
   x = "ICU Status",
   y = "AAGAB Expression Levels",
   color = "Sex"
 ) +
  scale_color_manual(values = c(" female" = "palevioletred1", " male" = "royalblue1", " unknown" = "lig"
   plot.title = element_text(size = 14, face = "bold"),
   axis.title = element_text(size = 12, face = "bold"),
   legend.title = element_text(size = 12, face = "bold"), # editing the legend
   legend.text = element_text(size = 10),
   legend.background = element_rect(fill = "lightgray", color = NA),
   legend.key = element_rect(fill = "white", color = "black"),
   panel.grid.major = element_line(color = "grey80"), # background
   panel.grid.minor = element_line(color = "grey90")
```

Box Plot of AAGAB Expression Levels by ICU Status and Sex



```
\#ggplot(series\_data,aes(x = disease\_status, y = as.numeric(AAGAB), color = sex)) +
# geom_boxplot() +
 labs( # labeling the title and axis
    title = "Box Plot of AAGAB Expression Levels by Disease Status and Sex",
#
    x = "Disease Status",
#
   y = "AAGAB Expression Levels",
#
    color = "Sex"
# ) +
# scale_color_manual(values = c(" female" = "palevioletred1", " male" = "royalblue1", " unknown" = "li
# theme(
#
    plot.title = element_text(size = 13, face = "bold"),
     axis.title = element_text(size = 12, face = "bold"),
#
    legend.title = element_text(size = 12, face = "bold"), # editing the legend
#
#
    legend.text = element_text(size = 10),
    legend.background = element_rect(fill = "lightgray", color = NA),
#
     legend.key = element_rect(fill = "white", color = "black"),
    panel.grid.major = element_line(color = "grey80"), # background
    panel.grid.minor = element_line(color = "grey90")
```

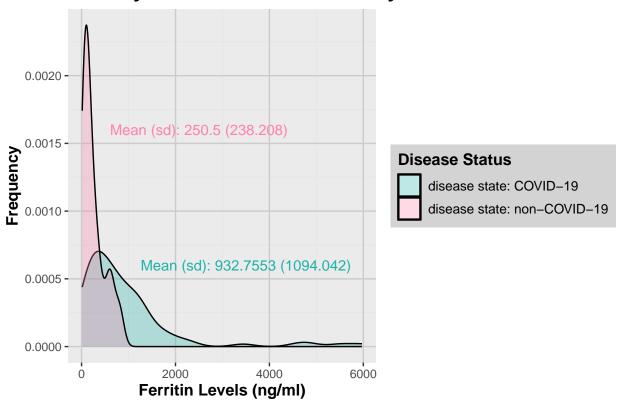
New Plot Type

```
#series_data$sex_disease_status <- paste(series_data$sex, series_data$disease_status, sep = " & ")
```

```
# Density plots with semi-transparent fill
\#ggplot(series\_data, aes(x=as.numeric(age), fill=icu\_status)) + geom\_density(alpha=.3)
\#ggplot(series\_data, aes(x=as.numeric(ferritin), fill=disease\_status)) + geom\_density(alpha=.3)
#ggplot(series_data, aes(x=as.numeric(ferritin), fill=sex)) + geom_density(alpha=.3)
#ggplot(series_data, aes(x=as.numeric(ferritin), fill=sex_disease_status)) + geom_density(alpha=.3)
mu <- series_data %>%
   group_by(disease_status) %>%
   summarize(grp.mean = mean(ferritin.ng.ml., na.rm = TRUE))
## Warning: There were 2 warnings in 'summarize()'.
## The first warning was:
## i In argument: 'grp.mean = mean(ferritin.ng.ml., na.rm = TRUE)'.
## i In group 1: 'disease_status = "disease state: COVID-19"'.
## Caused by warning in 'mean.default()':
## ! argument is not numeric or logical: returning NA
## i Run 'dplyr::last_dplyr_warnings()' to see the 1 remaining warning.
#mean(as.numeric(series_data$ferritin.ng.ml.[series_data$disease_status == "disease state: COVID-19"]),
\#sd(as.numeric(series\_data\$ferritin.ng.ml.[series\_data\$disease\_status == "disease state: COVID-19"]), note that the series is a substitution of the series of the series
#mean(as.numeric(series_data$ferritin.ng.ml.[series_data$disease_status == "disease state: non-COVID-19"
\#sd(as.numeric(series\_data\$ferritin.ng.ml.[series\_data\$disease\_status == "disease state: non-COVID-19"]
# Density plots with semi-transparent fill
ggplot(series_data, aes(x=as.numeric(series_data$ferritin.ng.ml.), fill=series_data$disease_status)) +
   labs( # labeling the title and axis
      title = "Density Plot of Ferritin Levels by Disease Status",
      x = "Ferritin Levels (ng/ml)",
      y = "Frequency",
      fill = "Disease Status"
   scale_fill_manual(values = c("disease state: COVID-19" = "lightseagreen", "disease state: non-COVID-1
   annotate(geom = 'text',x = 2500,y = 0.0016,label = 'Mean (sd): 250.5 (238.208)',color = 'palevioletre
   annotate(geom = 'text',x = 3500,y = 0.0006,label = 'Mean (sd): 932.7553 (1094.042)',color = 'lightsea
      plot.title = element_text(size = 16, face = "bold"),
       axis.title = element_text(size = 12, face = "bold"),
      legend.title = element_text(size = 12, face = "bold"), # editing the legend
      legend.text = element text(size = 10),
      legend.background = element_rect(fill = "lightgray", color = NA),
      legend.key = element_rect(fill = "white", color = "black"),
      panel.grid.major = element_line(color = "grey80"), # background
      panel.grid.minor = element_line(color = "grey90")
## Warning in FUN(X[[i]], ...): NAs introduced by coercion
## Warning: Use of 'series_data$ferritin.ng.ml.' is discouraged.
## i Use 'ferritin.ng.ml.' instead.
## Warning: Use of 'series_data$disease_status' is discouraged.
## i Use 'disease_status' instead.
```

Warning: Removed 16 rows containing non-finite outside the scale range
('stat_density()').

Density Plot of Ferritin Levels by Disease Status



Heatmap

pheatmap(

```
#install.packages('pheatmap')
library(pheatmap)

heatmap_gene_data <- gene_data %>%
    mutate(across(c(AAGAB, ABI1, ABHD5, ABHD2, AAAS, AAMDC, AAMP, AAR2, AARS1, AARSD1, AASDHPPT, AATF, AA'

# convert into a matrix
heatmap_matrix <- as.matrix(heatmap_gene_data[, c("AAGAB", "ABI1", "ABHD5", "ABHD2", "AAAS", "AAMDC", "
# adding tracking bars
tracking <- data.frame(
    ICU_Status = factor(series_data$cicu_status),
    Disease_Status = factor(series_data$disease_status)
)
rownames(tracking) <- rownames(heatmap_gene_data)

# plotting heatmap</pre>
```

```
heatmap_matrix,
cluster_rows = TRUE,
cluster_cols = TRUE,
annotation_row = tracking,
show_rownames = TRUE,
show_colnames = TRUE,
fontsize_row = 1,
#height = 10,
#cellheight = 1
```

