

# R Projects

2024-07-25

## R Markdown

This is an R Markdown document. Markdown is a simple formatting syntax for authoring HTML, PDF, and MS Word documents. For more details on using R Markdown see <http://rmarkdown.rstudio.com>.

When you click the **Knit** button a document will be generated that includes both content as well as the output of any embedded R code chunks within the document. You can embed an R code chunk like this:

```
# READ IN CSV FILES
```

```
setwd("/Users/sarahmirza/Documents/GitHub/QBS103_Repository/") # set the working directory
genes.df <- read.csv(file = "QBS103_GSE157103_genes.csv")
#head(genes.df) # see if it worked
```

```
matrix.df <- read.csv(file="QBS103_GSE157103_series_matrix.csv")
#head(matrix.df)
```

```
# GENES DATA FRAME - transpose
```

```
# remove column names and rows before transposing then add them back
```

```
genes_transpose <- t(genes.df)
```

```
genes.df <- as.data.frame(genes_transpose) # transpose, convert rows to columns and columns to rows
```

```
names(genes.df) <- genes.df[1,] #set the first row in the data frame and set it to the column names for
```

```
genes.df <- genes.df[-1,] # remove the first row in the data frame so that the names are no longer a row
```

```
# FROM TUTORIALSPPOINT - the [] maintains the data frame as a data frame because the lapply works on a list
```

```
# has been converted to a numeric because transpose makes them character
```

```
genes.df[] <- lapply(genes.df, function(x) as.numeric(as.character(x)))
```

```
genes.df <- na.omit(genes.df) # get rid of NA
```

```
genes.df$participant_id <- row.names(genes.df) # make new row called participant_id and with the gene.d
```

```
merged_matrix <- merge(genes.df,matrix.df,by = "participant_id") # merge the two data frames using participant_id
```

```
#head(merged_matrix) # see if it worked
```

```
# Load required package
```

```
library(ggplot2)
```

```
# Convert AAMP to numeric to read into histogram easier
```

```
merged_matrix$AAMP <- as.numeric(merged_matrix$AAMP)
```

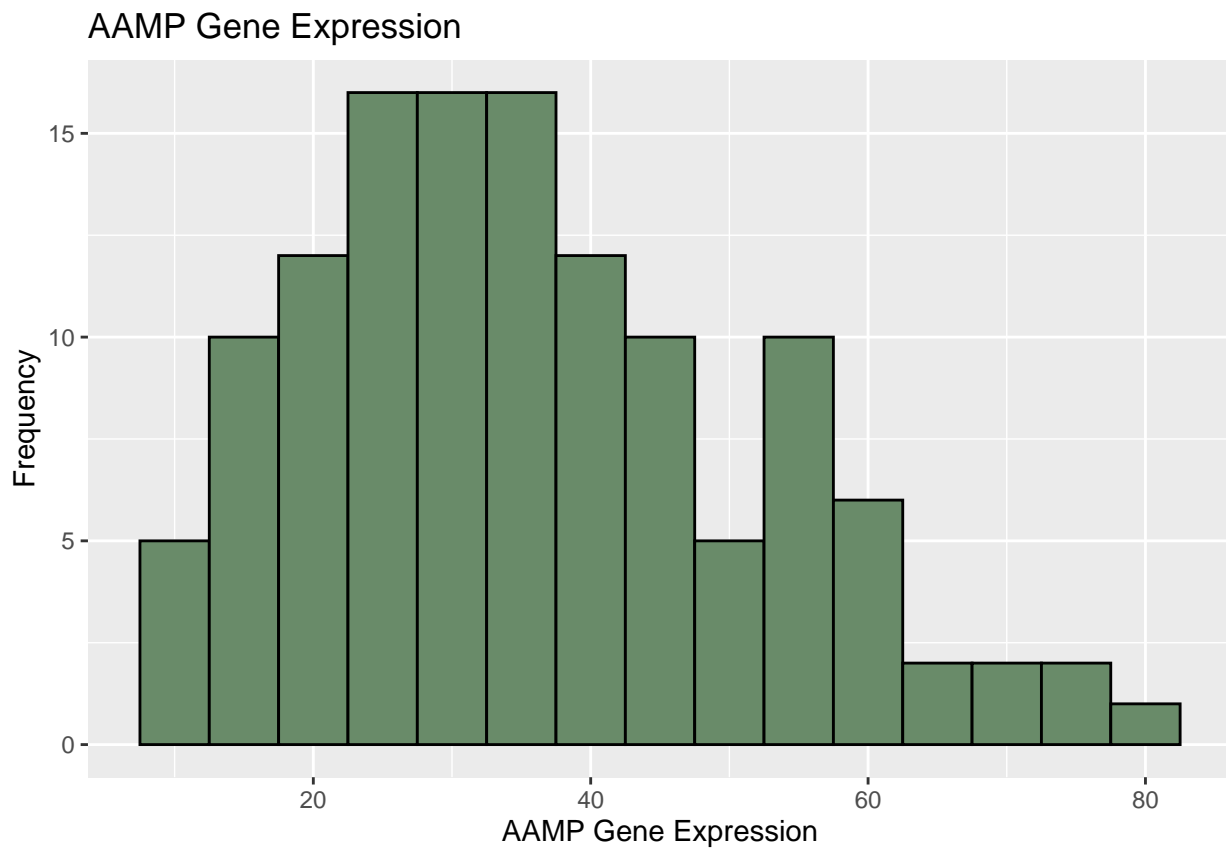
```
# Create histogram using the new data frame, the gene chosen was AAMP - changed binwidth per suggestion
```

```
histo <- ggplot(merged_matrix, aes(x = AAMP)) +
```

```
  geom_histogram(binwidth = 5, fill = "darkseagreen4", color = "black") +
```

```
  labs(title = "AAMP Gene Expression",
```

```
x = "AAMP Gene Expression",
y = "Frequency")
plot(histo)
```



```
library(ggplot2)
setwd("/Users/sarahmirza/Documents/GitHub/QBS103_Repository/")
```

```
# Convert columns to numeric for a gradient label
merged_matrix$AAMP <- as.numeric(merged_matrix$AAMP)
merged_matrix$age <- as.numeric(merged_matrix$age)
```

```
## Warning: NAs introduced by coercion
```

```
# Create scatterplot of AAMP expression vs. age
scatter <- ggplot(merged_matrix, aes(x = age, y = AAMP, color = age)) + # color = age gives the color bar
  geom_point() +
  scale_color_gradient(low = "blue", high = "red") +
  labs(title = "AAMP Expression vs. Age",
       x = "Age",
       y = "AAMP Expression")
plot(scatter)
```

```
## Warning: Removed 2 rows containing missing values or values outside the scale range
## (`geom_point()`).
```



```
# geom_smooth for best fit line
```

```
# Save the plot to a file
```

```
ggsave("AAMP_gene_expression_vs_age.pdf", plot = scatter, width = 8, height = 5)
```

```
## Warning: Removed 2 rows containing missing values or values outside the scale range
```

```
## (`geom_point()`).
```

```
library(ggplot2)
```

```
setwd("/Users/sarahmirza/Documents/GitHub/QBS103_Repository/")
```

```
# a boxplot of AAMP expression separated by sex and ICU Status
```

```
ggplot(merged_matrix, aes(x = icu_status, y = AAMP, fill = sex)) +
```

```
  geom_boxplot() +
```

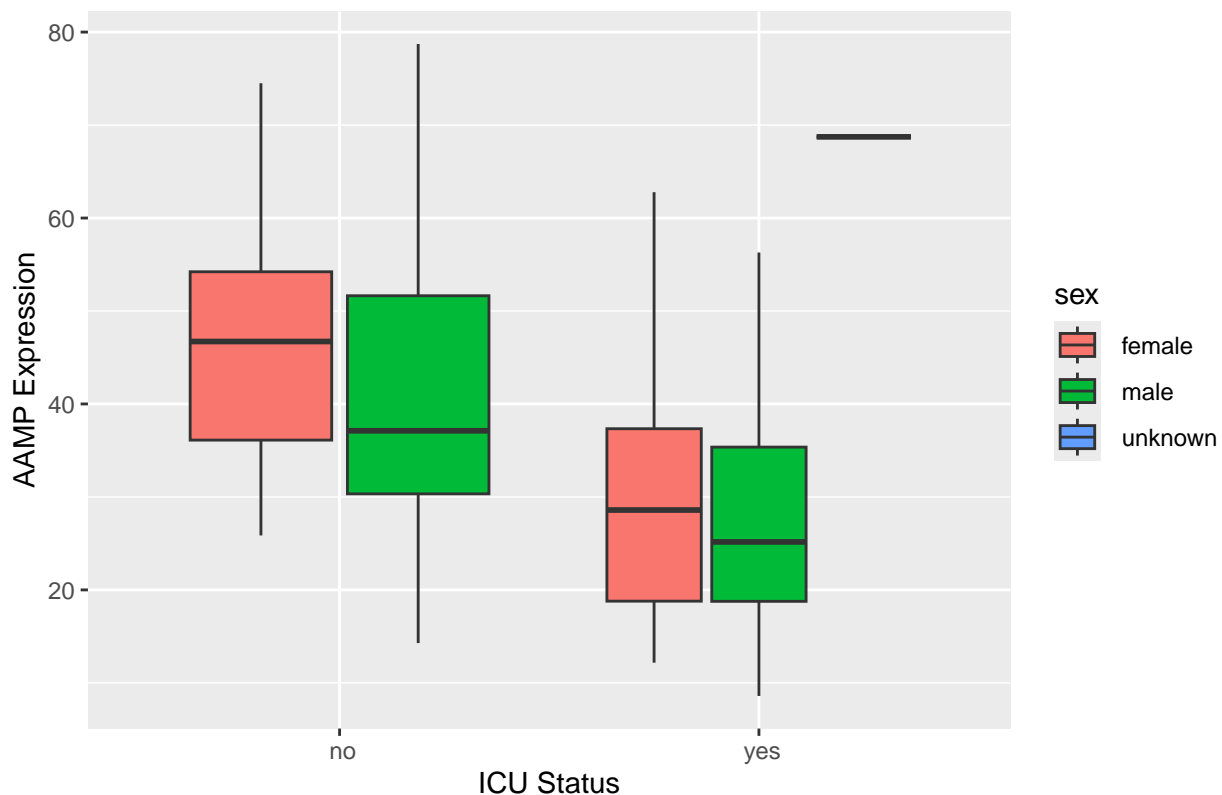
```
  labs(title = "AAMP Expression by ICU Status and Sex",
```

```
        x = "ICU Status",
```

```
        y = "AAMP Expression",
```

```
        fill = "sex")
```

## AAMP Expression by ICU Status and Sex



```
#merged_matrix$ferritin.ng.ml. <- as.factor(merged_matrix$ferritin.ng.ml.)
#merged_matrix$icu_status <- as.factor(merged_matrix$icu_status)

# Create a boxplot of AAMP expression separated by icu status and ferritin.ng.ml.
#ggplot(merged_matrix, aes(x = icu_status, y = AAMP, fill = ferritin.ng.ml.)) +
#  geom_boxplot() +
#  labs(title = "AAMP Expression by ICU Status and ferritin.ng.ml.",
#        x = "ICU Status",
#        y = "AAMP Expression",
#        fill = "ferritin.ng.ml.")

# Create a boxplot of AAMP expression separated by source_name_ch1 and icu status
#ggplot(merged_matrix, aes(icu_status, y = AAMP, fill = source_name_ch1)) +
#  geom_boxplot() +
#  labs(title = "AAMP Expression by ICU Status and source_name_ch1",
#        x = "icu_status",
#        y = "AAMP Expression",
#        fill = "source_name_ch1")
```

```
library(harrypotter)
```

```
# function to plot all three plots
fun_stats_pretty_plots <- function(matrix, gene_name, continuous_name, categorical1_name, categorical2_name) {
  # print(matrix)
  matrix$gene <- matrix[, gene_name] # dummy variable that creates a new column with the name of the gene
  # this is used for plotting, and the function is fed a string that I use for labeling which is the gene name
  #matrix$gene <- as.numeric(matrix$gene)
```

```

matrix$continuous <- matrix[,continuous_name] # do the same for all variables so they are read in a
matrix$categorical1 <- matrix[,categorical1_name]
matrix$categorical2 <- matrix[,categorical2_name]

histogram <- ggplot(matrix, aes(x=gene)) + geom_histogram(bins = 20, fill = "chartreuse2", color =
labs(title = paste0(gene_name, " Gene Expression"),
  x = paste0(gene_name, " Gene Expression"),
  y = "Frequency")

scatterplot <- ggplot(matrix, aes(x=continuous, y=gene, color = continuous)) +
geom_point() + scale_color_gradient(low = "blue", high = "red", name = (paste0(continuous_name))) +
labs(title = paste0(gene_name, " Expression vs. ", continuous_name),
  x = paste0(continuous_name),
  y = paste0(gene_name, " Expression"))

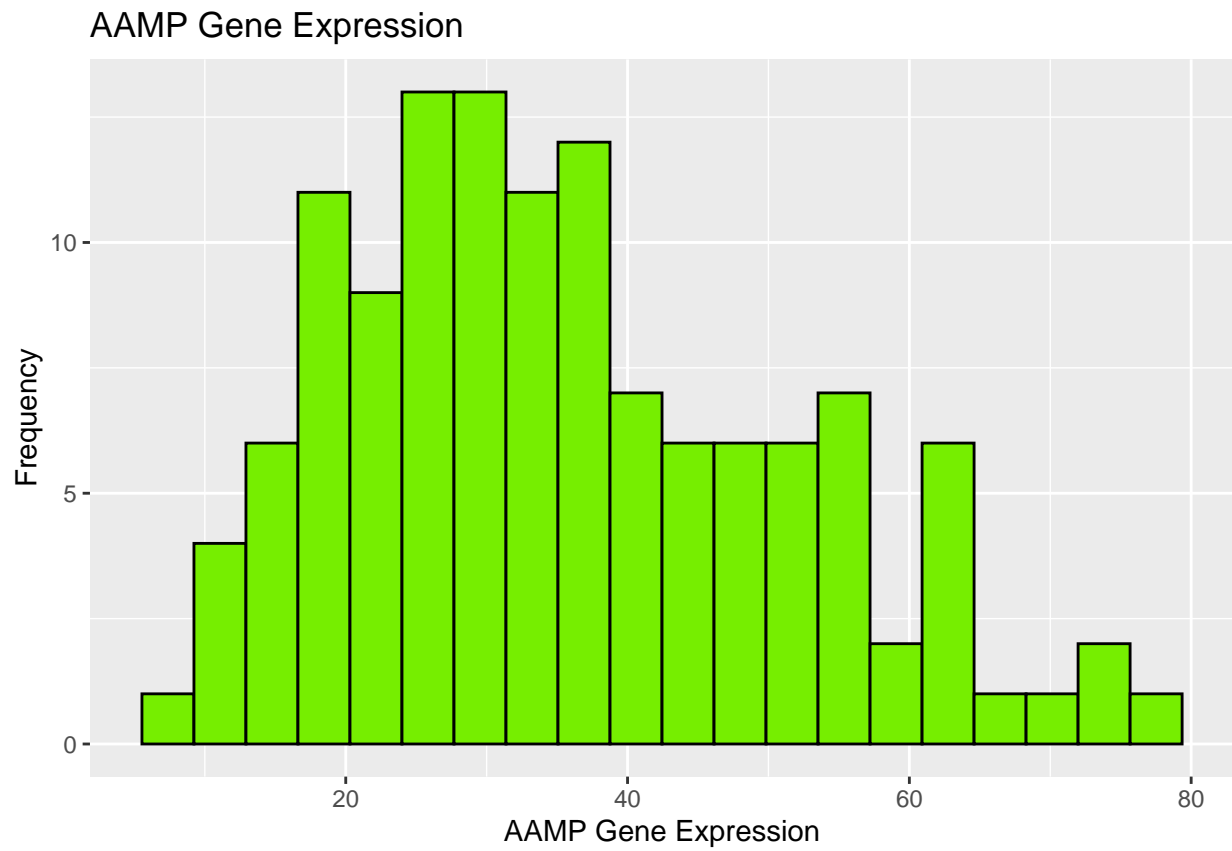
boxplot <- ggplot(matrix, aes(x=categorical1, y=gene, fill = categorical2)) +
geom_boxplot() + scale_fill_hpd(option = "lunalevegood") +
labs (title = paste0(gene_name, " Expression by ", categorical1_name, " and ", categorical2_name),
  x = paste0(categorical1_name),
  y = paste0(gene_name, " Expression"),
  fill = paste0(categorical2_name))

plot(histogram)
plot(scatterplot)
plot(boxplot)
}

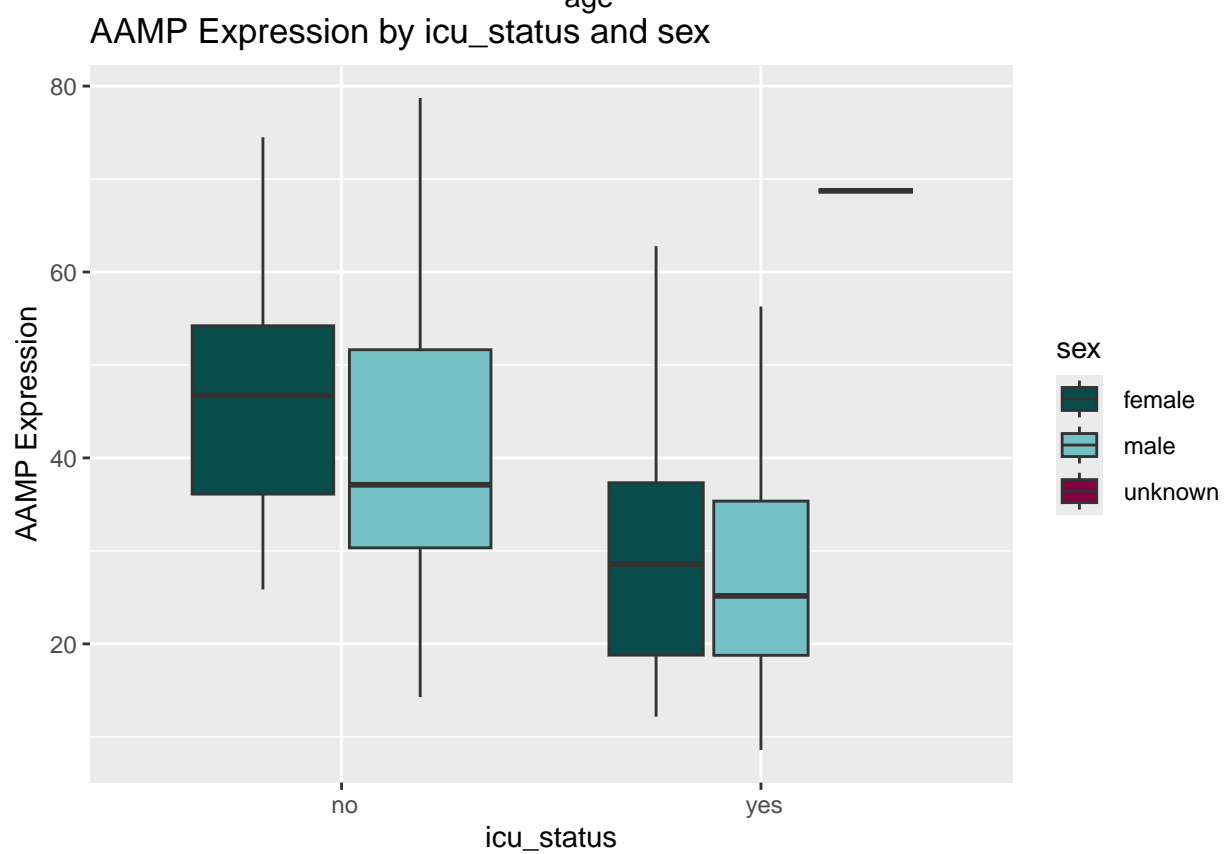
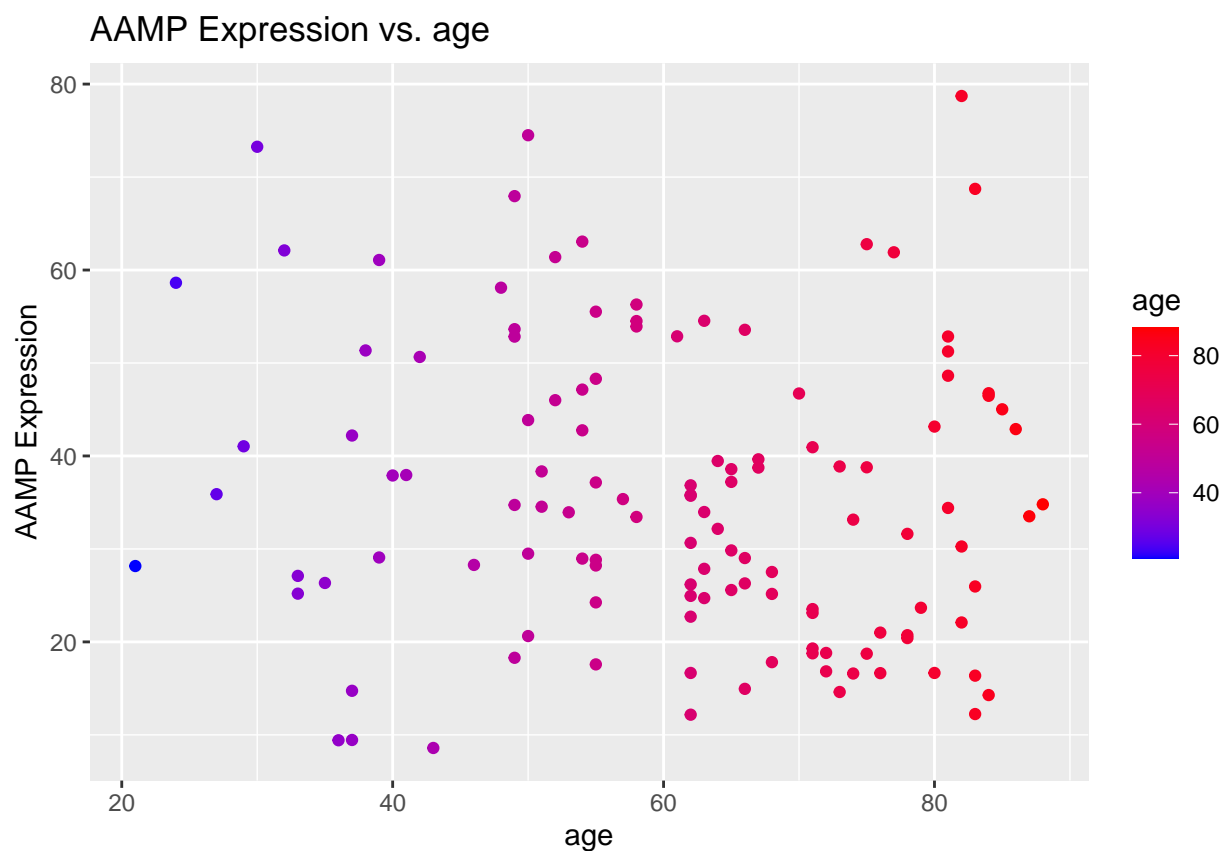
# genes to be used during plotting - stored in a list
plot_genes = c("AAMP", "AAK1", "ABCA7")

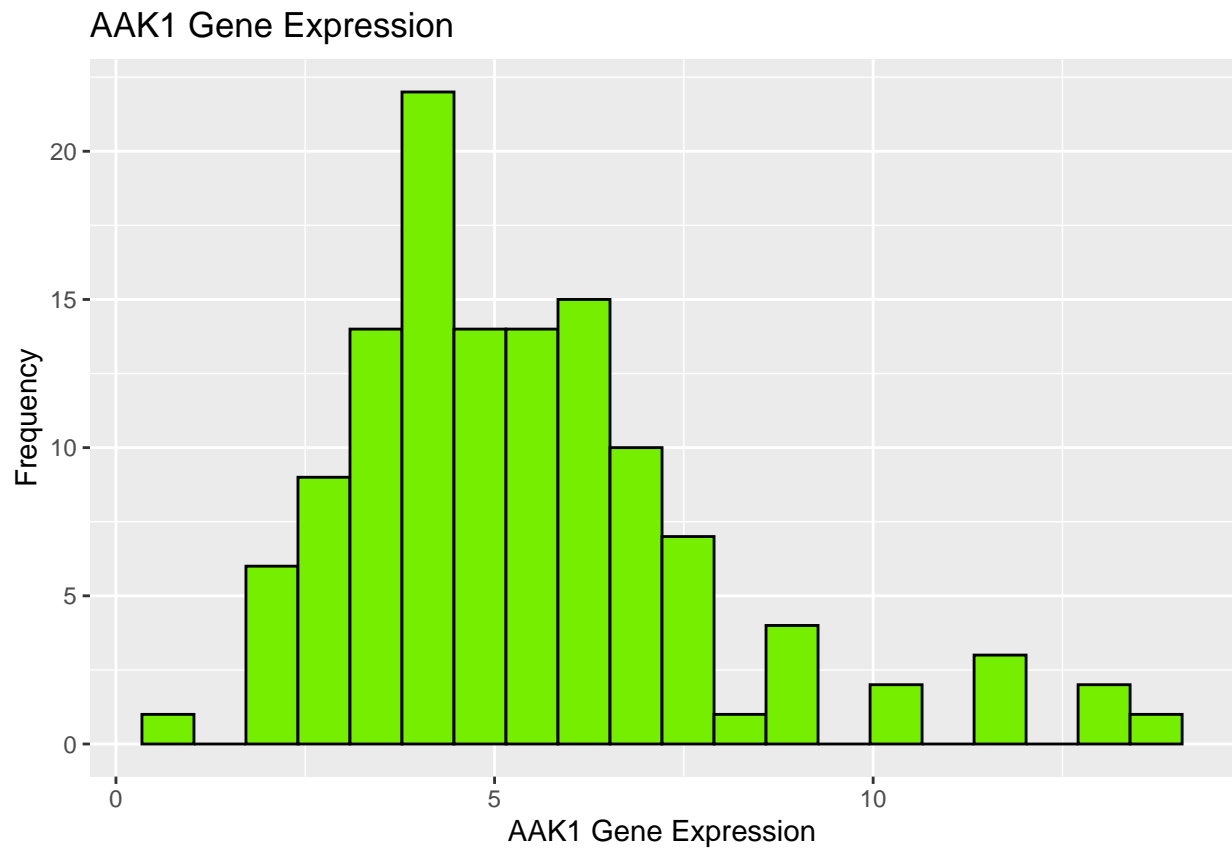
# for loop - replace gene name with gene in list
for (g in plot_genes) {
  fun_stats_pretty_plots(matrix = merged_matrix, gene_name=g, continuous_name = 'age', categorical1_name =
}

```



```
## Warning: Removed 2 rows containing missing values or values outside the scale range
## (`geom_point()`).
```

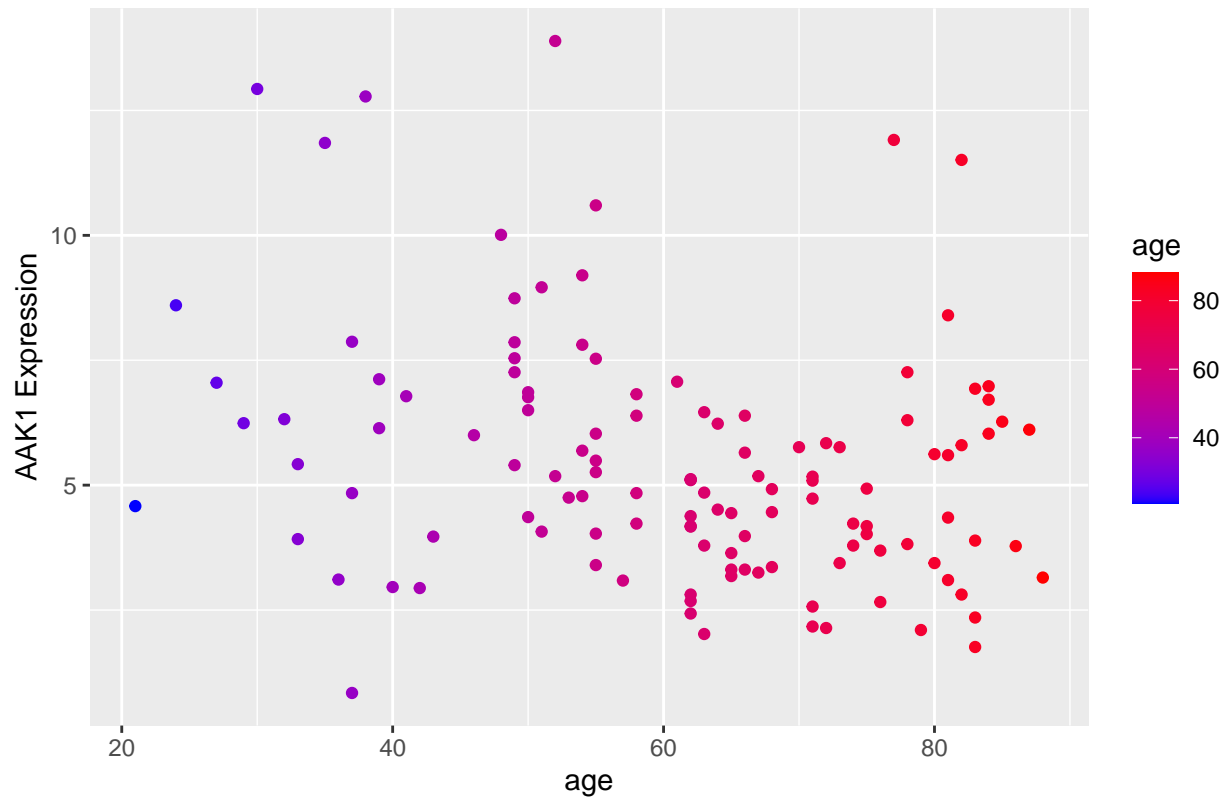




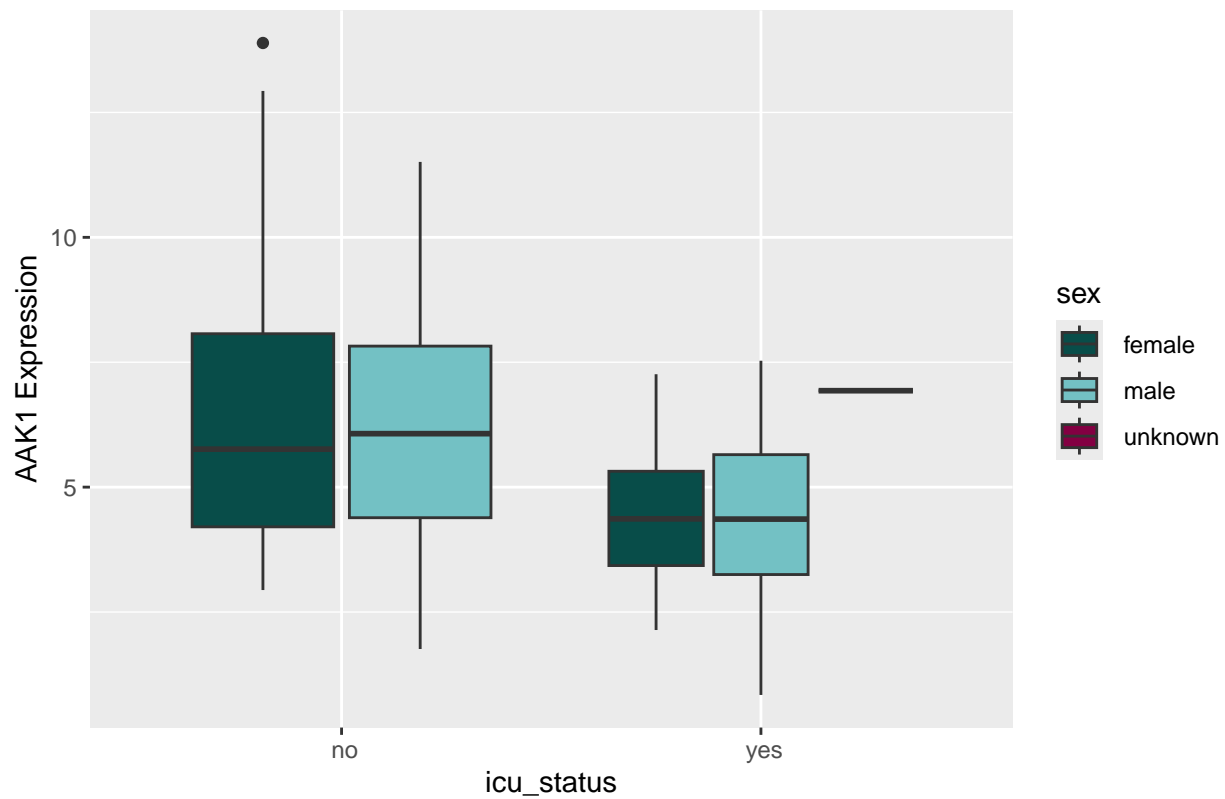
```
## Warning: Removed 2 rows containing missing values or values outside the scale range
## (`geom_point()`).
```

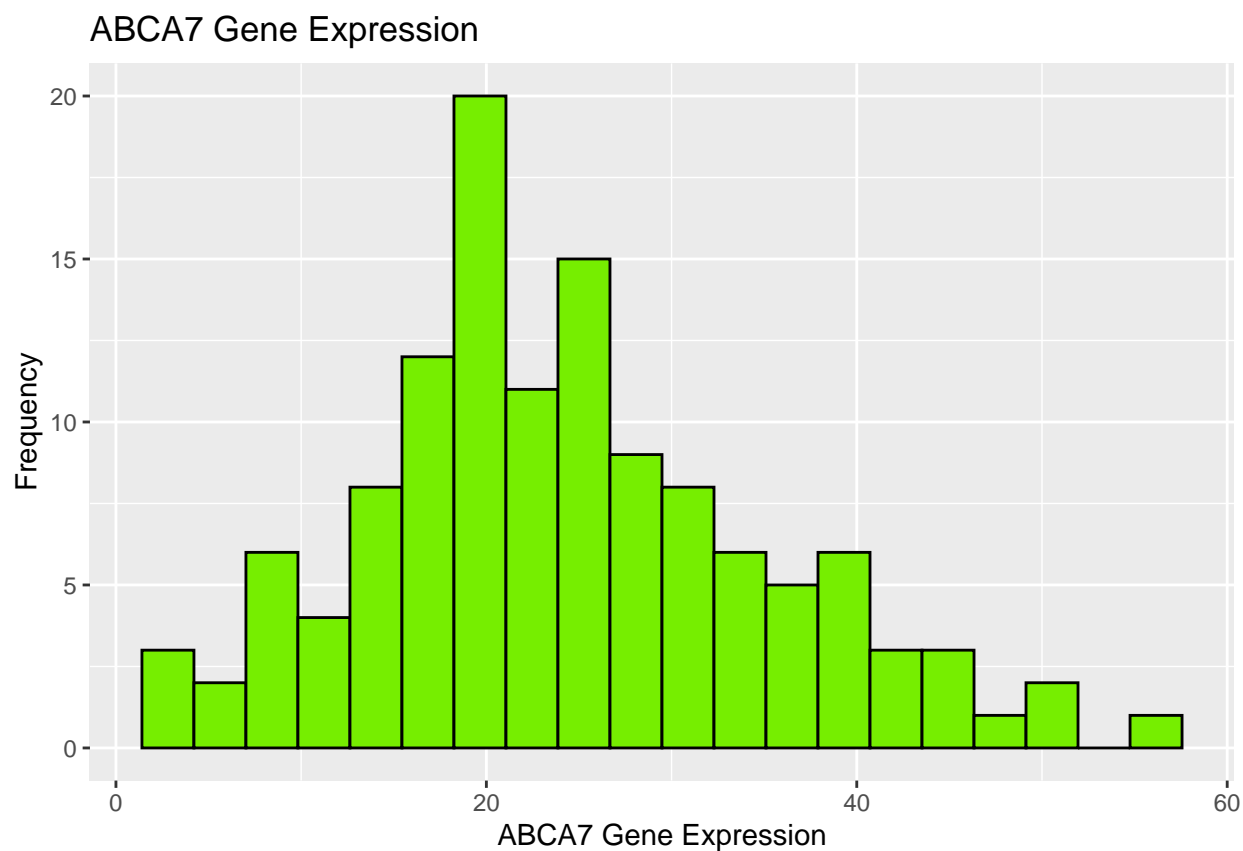


AAK1 Expression vs. age

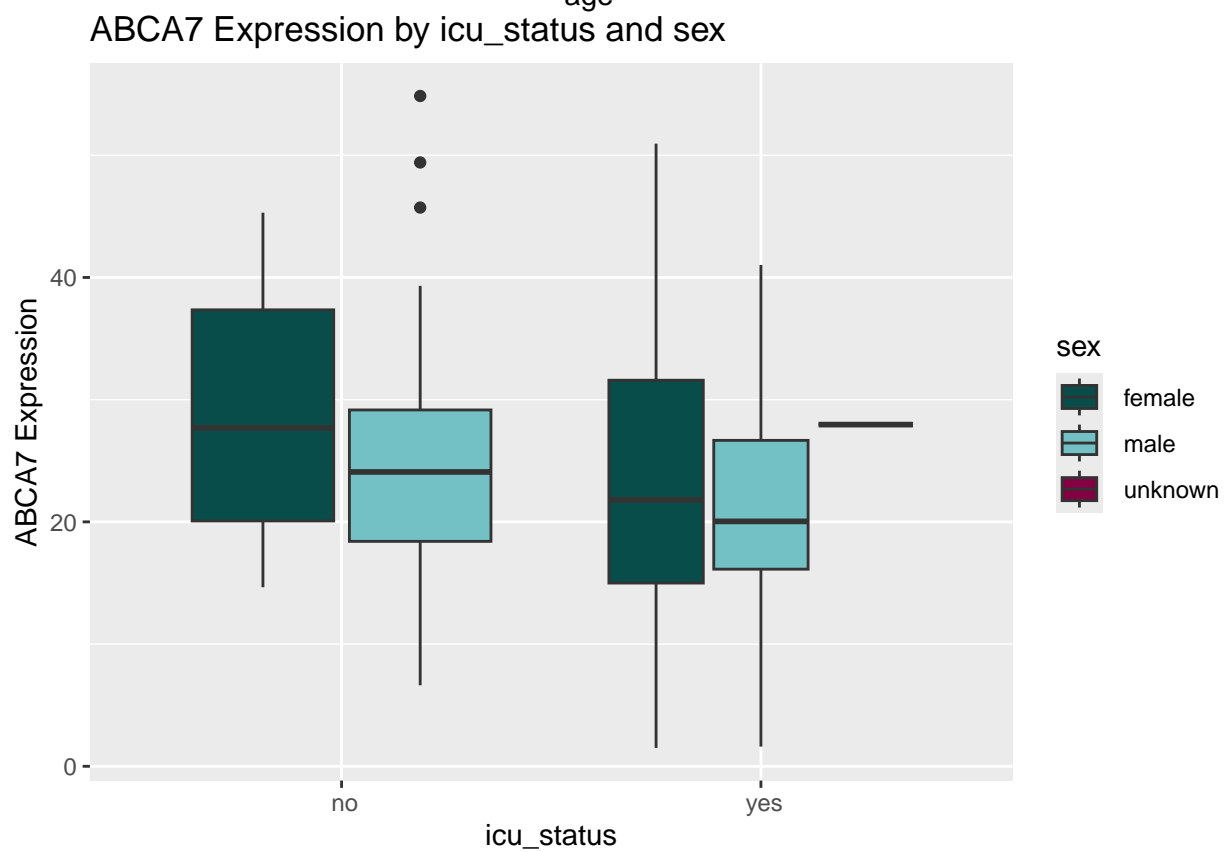


AAK1 Expression by icu\_status and sex





```
## Warning: Removed 2 rows containing missing values or values outside the scale range
## (`geom_point()`).
```



```
# gene 1
#fun_stats_pretty_plots(matrix = merged_matrix, gene_name='AAMP', continuous_name = 'age', categorical1_name = 'sex')

# gene 2
#fun_stats_pretty_plots(matrix = merged_matrix, gene_name='AAS', continuous_name = 'age', categorical1_name = 'sex')

# gene 3
#fun_stats_pretty_plots(matrix = merged_matrix, gene_name='ABHD14A-ACY1', continuous_name = 'age', categorical1_name = 'sex')
```