## QBS 103 Final Project

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#### Variables used to generate plots

Gene:LCN2

Continuous variable: Age

Categorical Variables: Sex and Disease Status

The LCN2 gene encodes the protein Lipocalin 2 which plays a role in innate immunity by limiting bacterial growth through the sequestering of iron-containing siderophores. This protein is thought to be be involved in multiple cellular processes: the maintenance of skin homeostasis and suppression of invasiveness and metastasis.

In relation to COVID-19, researchers found that LCN2 is overexpressed in COVID-19 patients. They also found that this gene is linked to neutrophil and virus response activities, so higher expression of this gene leads to inflammatory responses and cilium movement.

#### Sources:

https://www.genecards.org/cgi-bin/carddisp.pl?gene=LCN2

https://www.ncbi.nlm.nih.gov/pmc/articles/PMC7714049/

```
setwd("C:/Users/Student/Desktop/QBS103/")# set working directory
```

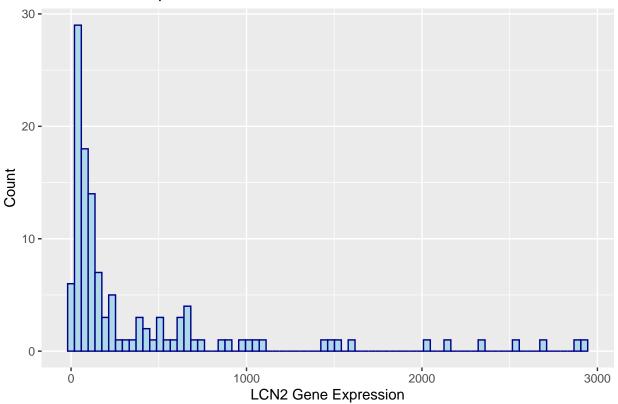
#### library(tidyverse)

head(gene\_expression)

```
## -- Attaching core tidyverse packages ----- tidyverse 2.0.0 --
## v dplyr
           1.1.2
                        v readr
                                    2.1.4
## v forcats 1.0.0
                                    1.5.0
                        v stringr
## v ggplot2 3.4.2
                        v tibble
                                    3.2.1
## v lubridate 1.9.2
                        v tidyr
                                    1.3.0
## v purrr
              1.0.1
## -- Conflicts -----
                                       ------tidyverse_conflicts() --
## x dplyr::filter() masks stats::filter()
## x dplyr::lag()
                    masks stats::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
meta_data = read.csv("QBS103_finalProject_metadata.csv")
#head(meta_data)
gene_data = read.csv("QBS103_finalProject_geneExpression.csv")
#head(gene_data)
# convert into vertical data and extract specific gene: LCN2
gene_expression = matrix(gene_data[5,2:127])
```

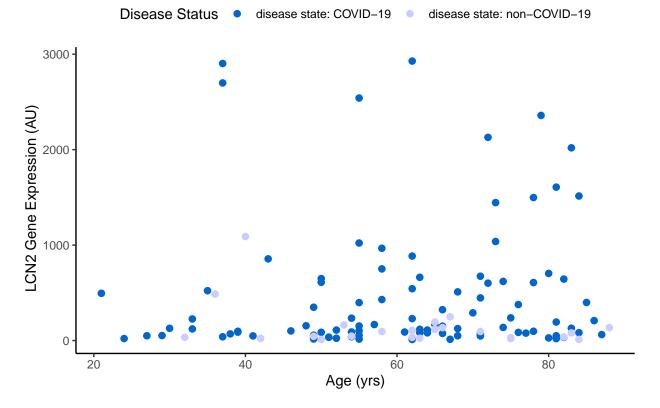
```
[,1]
##
## [1,] 87.71
## [2,] 662.59
## [3,] 121.86
## [4,] 31.73
## [5,] 16.42
## [6,] 447.98
# new data frame created to link variables from both data sets into one
myData <- data.frame('Subject ID' = seq(1:126), "Gene.Expression" = gene_expression, 'Age' = meta_data$a
head(myData)
    Subject.ID Gene.Expression Age
                                                   Disease.Status ICU.Status
##
                                      Sex
## 1
                         87.71 39 male disease state: COVID-19
             1
## 2
                         662.59 63 male disease state: COVID-19
                                                                          nο
## 3
              3
                         121.86 33 male disease state: COVID-19
                                                                          no
## 4
              4
                          31.73 49 male disease state: COVID-19
                                                                          nο
## 5
              5
                         16.42 49 male disease state: COVID-19
                                                                          no
## 6
              6
                         447.98 : male disease state: COVID-19
                                                                          nο
myData <- myData[!(row.names(myData) %in% c("6","86","104","115")),] # filter out data that do not have
# Source: https://sparkbyexamples.com/r-programming/drop-dataframe-rows-in-r/?expand_article=1
myData$Age<- as.numeric(myData$Age)# Convert age variable to numeric</pre>
myData$Gene.Expression<- as.numeric(myData$Gene.Expression) #convert gene expression variable to numeri
# Histogram for gene expression
ggplot(myData, aes(x=Gene.Expression)) +
  geom_histogram(binwidth = 39,color = "darkblue", fill = "lightblue")+
  labs(title = "LCN2 Gene Expression Distribution", x = "LCN2 Gene Expression", y = "Count")
```

### LCN2 Gene Expression Distribution



```
BlankTheme <- theme(# Remove borders and grid lines
       panel.border = element_blank(), panel.grid.major = element_blank(),
       panel.grid.minor = element_blank(),
        # Define my axis
       axis.line = element_line(colour = "black", linewidth = rel(1)),
        # Set plot background to white
       plot.background = element_rect(fill = "white"),
       panel.background = element_blank(),
        legend.key = element_rect(fill = 'white'),
        # Move legend to the top
       legend.position = 'top')
# Scatterplot for gene expression and continuous covariate
ggplot(myData, aes(x = Age, y = Gene.Expression, color = Disease.Status))+
 geom_point(size = 2)+
 labs(title = "Gene Expression of LCN2 by Age", x = "Age (yrs)", y = " LCN2 Gene Expression (AU)", col
  scale_color_manual(values = c("#0066CC", "#CCCCFF")) +
  BlankTheme
```

## Gene Expression of LCN2 by Age



```
BlankTheme <- theme(# Remove borders and grid lines
        panel.border = element_blank(), panel.grid.major = element_blank(),
       panel.grid.minor = element_blank(),
        # Define my axis
        axis.line = element_line(colour = "black", linewidth = rel(1)),
        # Set plot background to white
       plot.background = element_rect(fill = "white"),
       panel.background = element_blank(),
        legend.key = element_rect(fill = 'white'),
        # Move legend to the top
       legend.position = 'top')
# Box plot of gene expression separated by sex and disease status
ggplot(myData, aes(x = Disease.Status, y = Gene.Expression, color = Sex))+
  geom_boxplot()+
  scale_color_manual(values = c('#000066', '#669900')) +
  labs(title = "Gene Expression of LCN2 grouped by Disease status and categorized by Sex", x = "Disease
  BlankTheme
```

# Gene Expression of LCN2 grouped by Disease status and categorized by

