

01- Data Exploration

Directory and doc rules

```
knitr::opts_chunk$set(  
  echo = TRUE,      # Display code chunks  
  eval = TRUE,      # Evaluate code chunks  
  warning = FALSE,  # Hide warnings  
  message = FALSE,  # Hide messages  
  fig.width = 20,    # Set plot width in inches  
  fig.height = 9,    # Set plot height in inches  
  fig.align = "center" # Align plots to the center  
)
```

Load packages

```
library(tinytex)  
library(tidyr)  
library(tidyverse)  
library(vegan)
```

Load data

Weight = mg
Length, width, height = mm
p450, SOD = activity/ (mg/protein)
Condition factor, economic factor = unitless

```
getwd()
```

```
## [1] "/Users/cmantegna/Documents/WDFWmussels/code"
```

```
#data has all sites, coordinates, p450, sod, condition factor, economic factor data  
data<- read.csv("/Users/cmantegna/Documents/WDFWmussels/data/biomarkerfull.csv")
```

```
#alldata has the site names, biomarkers, condition factor, average thickness and analyte data - each row is a site  
alldata<- read.csv("/Users/cmantegna/Documents/WDFWmussels/data/alldata.csv")
```

fix zero's in the data frame and alldata frame

```
# Data contains 0's and must be adjusted in this order to preserve all usable data.
```

```
#sod
```

```
#replace any SOD values at or below 0 with half of the lower detection limit of .005 (.005*.5). Lower d
```

```
data$SOD[data$SOD <= 0] <- 0.0025
```

```
alldata$SOD[alldata$SOD <= 0] <- 0.0025
```

```
#p450
#remove any p450 values that are 0 - those are true 0's not non-detectable. I am replacing with na so I
data$p450[data$p450 <= 0] <- NA
alldata$p450[alldata$p450 <= 0] <- NA

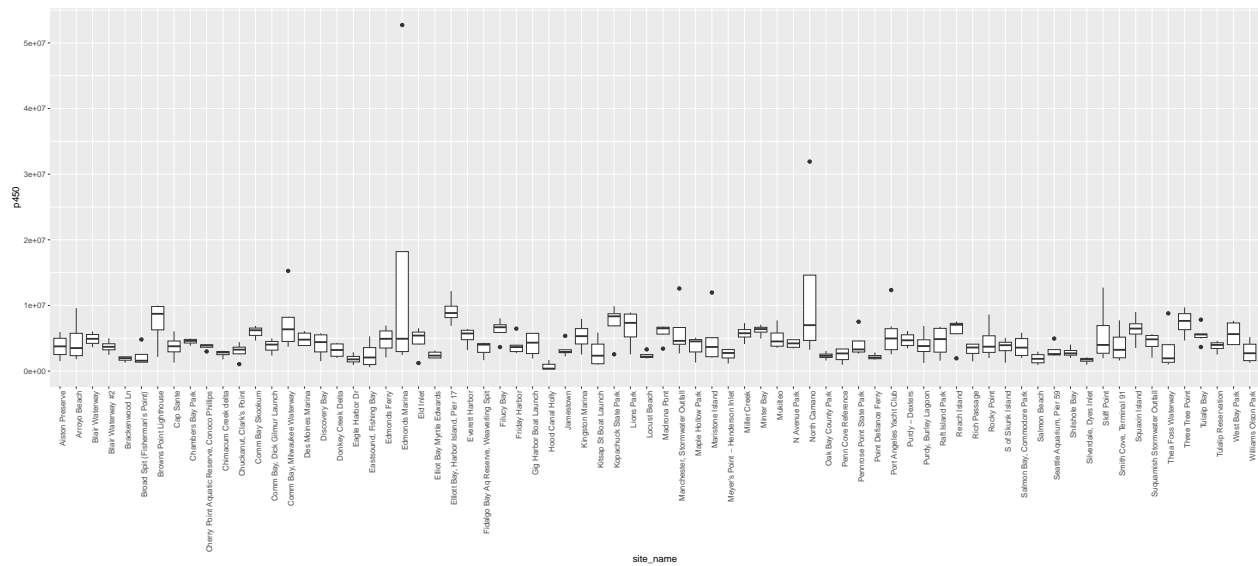
#write.csv(alldata, "/Users/cmantegna/Documents/WDFWmussels/data/alldata.csv")

# check the data frame
#summary(alldata)
#str(alldata)
```

Boxplot of biomarker data, p450 and SOD

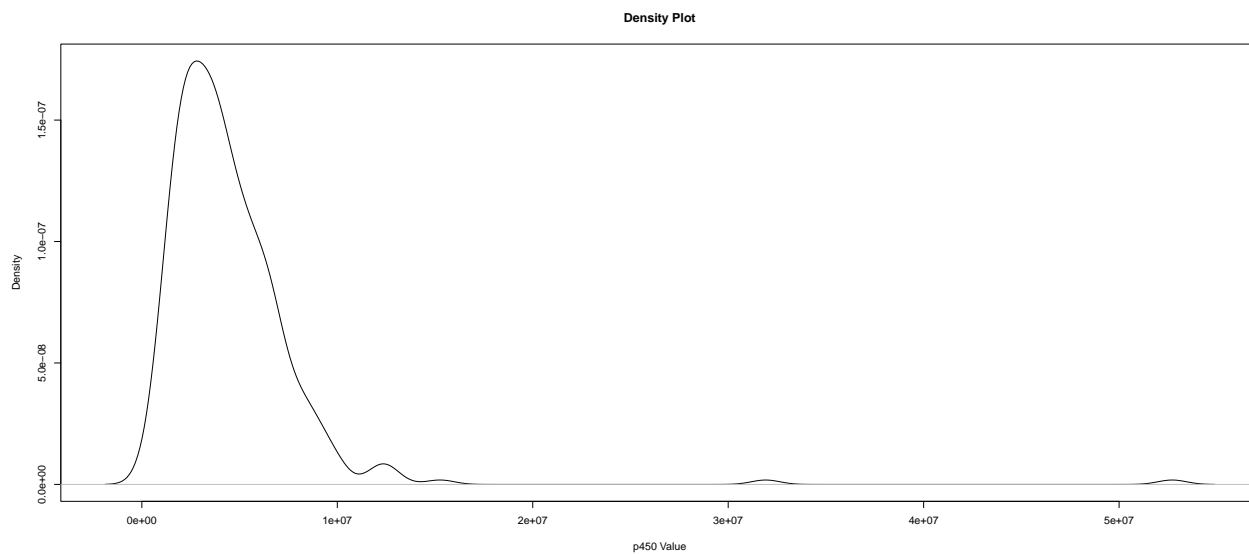
```
#p450
pplot<- ggplot(data, aes(x = site_name, y = p450)) +
  geom_boxplot() +
  theme(axis.text.x = element_text(angle = 90, hjust = 1))

print(pplot)
```

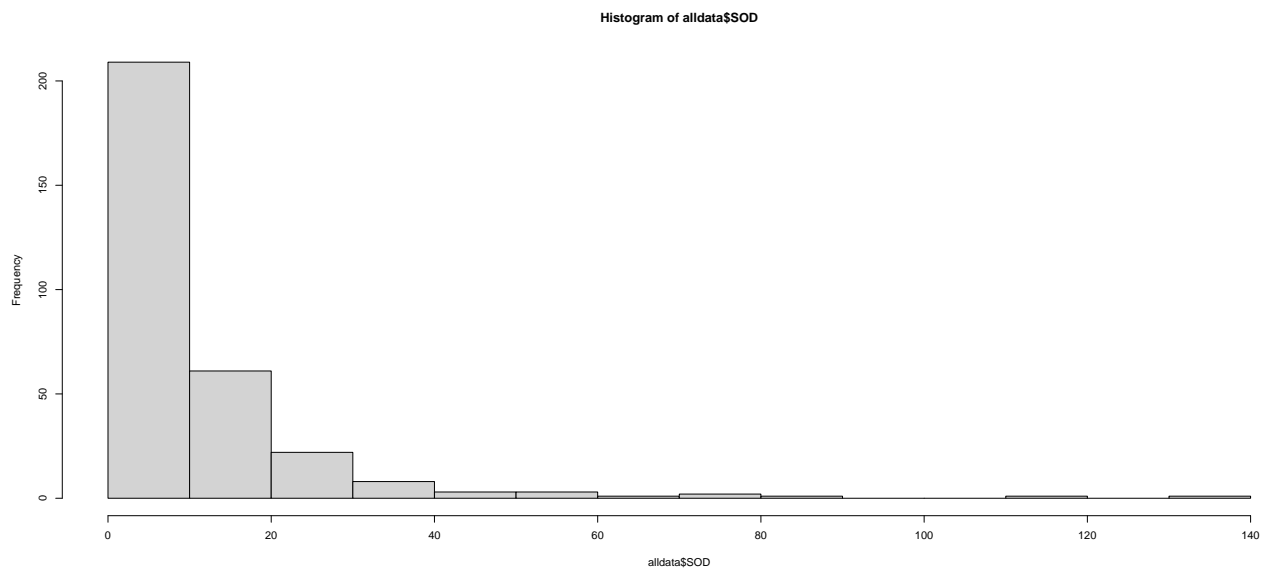


```
#SOD
splot<- ggplot(data, aes(x = site_name, y = SOD)) +
  geom_boxplot() +
  theme(axis.text.x = element_text(angle = 90, hjust = 1))

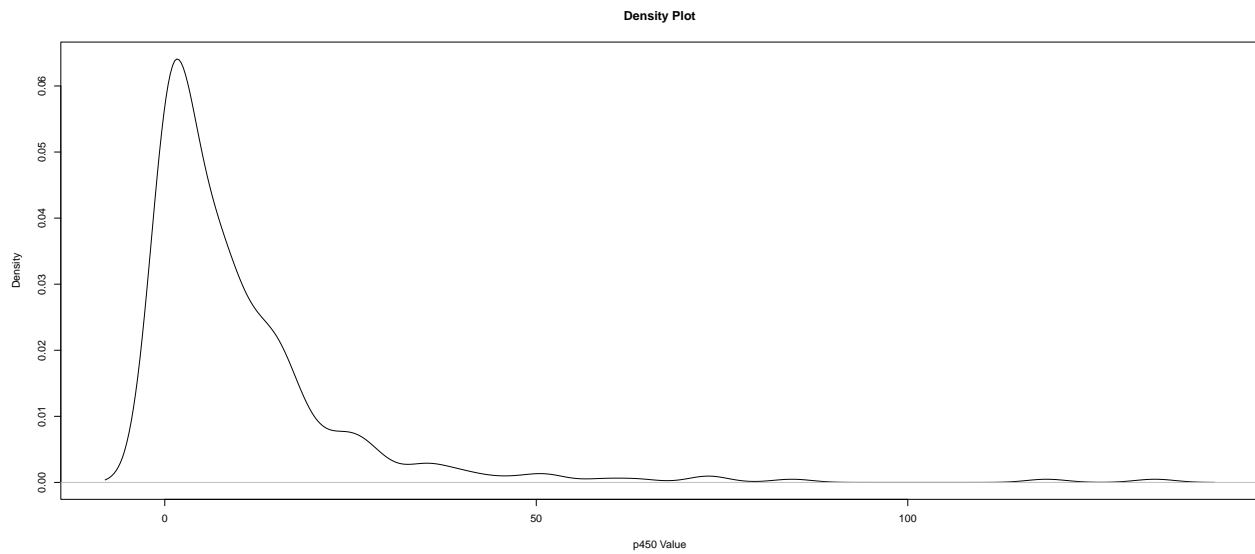
print(splot)
```

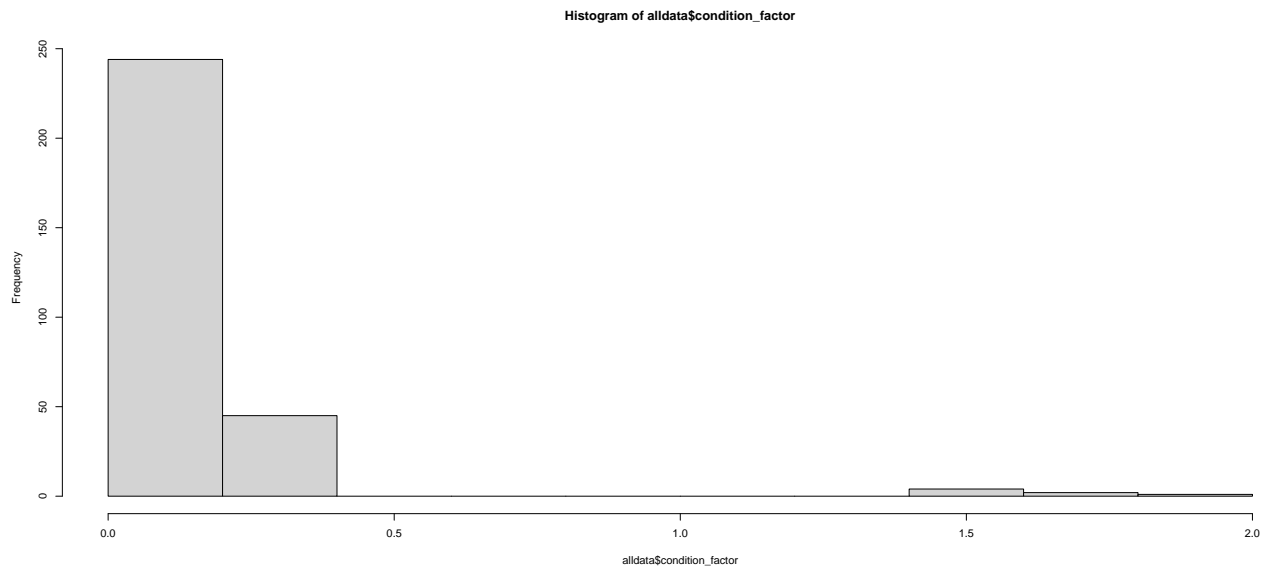
```
# SOD basic histogram + basic density plot
hist(alldata$SOD)
```



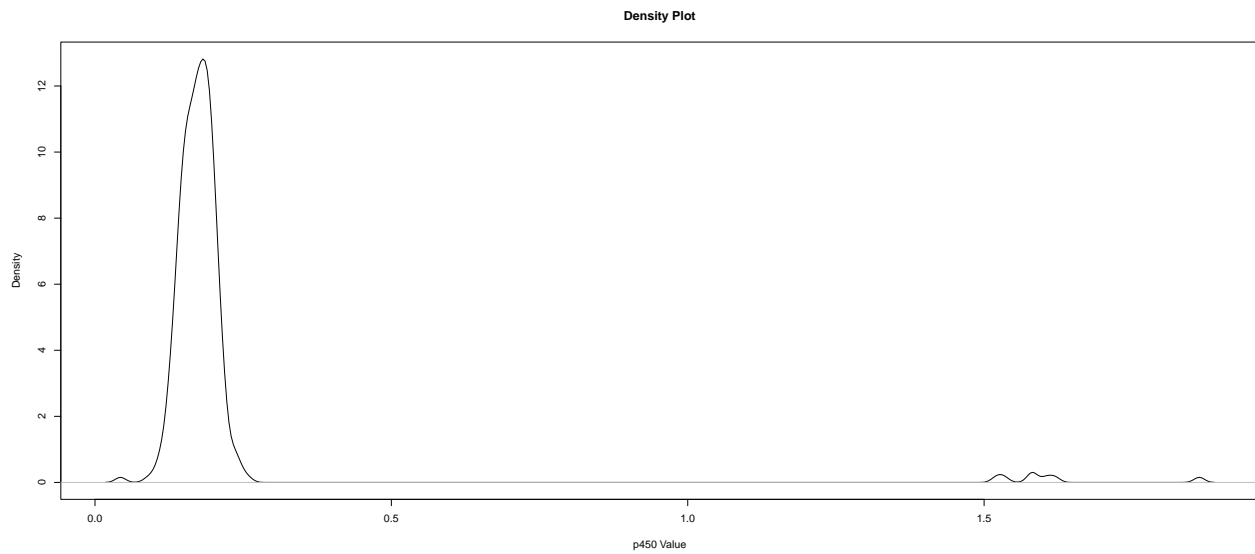
```
plot(density(alldata$SOD, na.rm= TRUE), main="Density Plot", xlab="p450 Value")
```



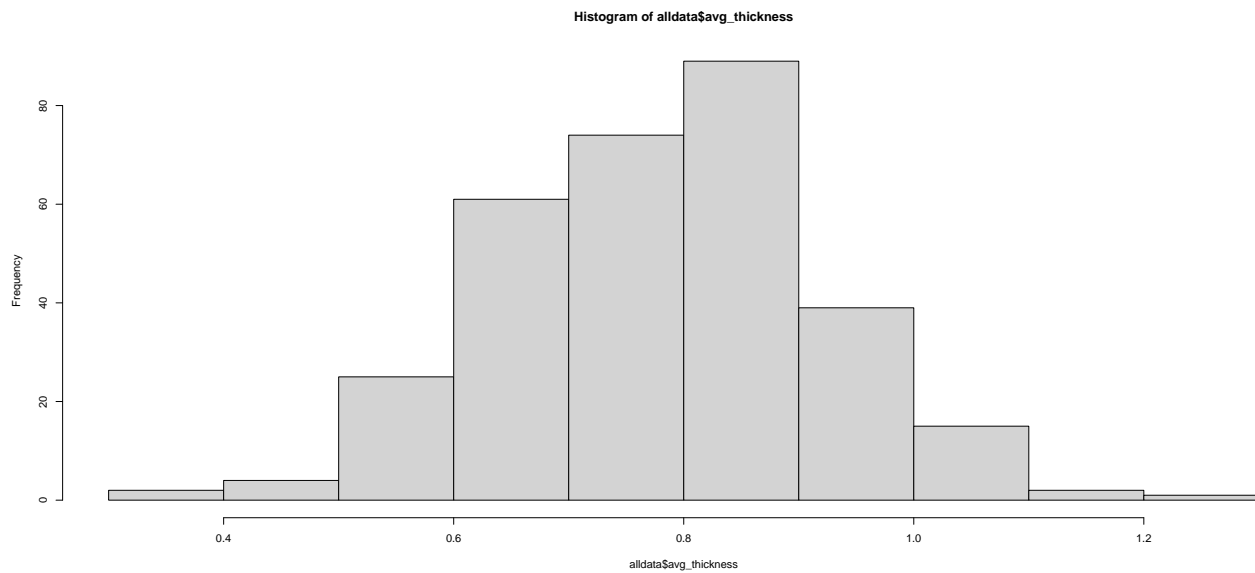
```
# Condition_factor basic histogram + basic density plot  
hist(alldata$condition_factor)
```



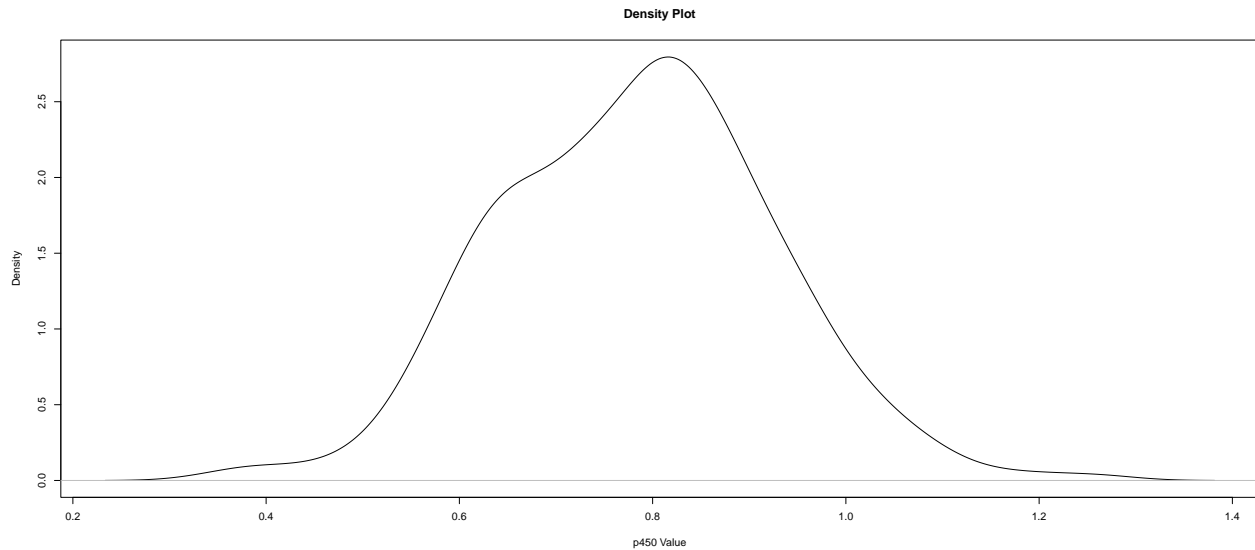
```
plot(density(alldata$condition_factor, na.rm= TRUE), main="Density Plot", xlab="p450 Value")
```



```
# Avg_thickness basic histogram + basic density plot  
hist(alldata$avg_thickness)
```



```
plot(density(alldata$avg_thickness, na.rm= TRUE), main="Density Plot", xlab="p450 Value")
```



Shapiro-Wilkes test for normality

Only avg_thickness is normally distributed.

```
shapiro.test(alldata$p450)
```

```
##  
##  Shapiro-Wilk normality test  
##  
## data:  alldata$p450  
## W = 0.55896, p-value < 2.2e-16
```

```
shapiro.test(alldata$SOD)
```

```
##  
##  Shapiro-Wilk normality test  
##  
## data:  alldata$SOD  
## W = 0.61784, p-value < 2.2e-16
```

```
shapiro.test(alldata$condition_factor)
```

```
##  
##  Shapiro-Wilk normality test  
##  
## data:  alldata$condition_factor  
## W = 0.23346, p-value < 2.2e-16
```

```
shapiro.test(alldata$avg_thickness)
```

```
##  
##  Shapiro-Wilk normality test  
##  
## data:  alldata$avg_thickness  
## W = 0.99674, p-value = 0.7814
```

Kruskal-Wallis, p450

p450 and site have a statistically significant relationship, $p < 0.0000008077$

```
kruskal.test(p450 ~ site_name, data = alldata)

##
## Kruskal-Wallis rank sum test
##
## data: p450 by site_name
## Kruskal-Wallis chi-squared = 137.23, df = 73, p-value = 8.077e-06

kruskal.test(p450 ~ SOD, data = alldata)

##
## Kruskal-Wallis rank sum test
##
## data: p450 by SOD
## Kruskal-Wallis chi-squared = 252.32, df = 254, p-value = 0.5179

kruskal.test(p450 ~ condition_factor, data = alldata)

##
## Kruskal-Wallis rank sum test
##
## data: p450 by condition_factor
## Kruskal-Wallis chi-squared = 259.49, df = 255, p-value = 0.4102

kruskal.test(p450 ~ avg_thickness, data = alldata)

##
## Kruskal-Wallis rank sum test
##
## data: p450 by avg_thickness
## Kruskal-Wallis chi-squared = 106.49, df = 110, p-value = 0.577
```

Kruskal-Wallis, SOD

SOD and site have a statistically significant relationship, $p < 0.0000005669$

```
kruskal.test(SOD ~ site_name, data = alldata)

##
## Kruskal-Wallis rank sum test
##
## data: SOD by site_name
## Kruskal-Wallis chi-squared = 138.64, df = 73, p-value = 5.669e-06

kruskal.test(SOD ~ p450, data = alldata)

##
## Kruskal-Wallis rank sum test
##
## data: SOD by p450
## Kruskal-Wallis chi-squared = 303, df = 303, p-value = 0.4892
```



```
kruskal.test(SOD ~ condition_factor, data = alldata)

##
##  Kruskal-Wallis rank sum test
##
## data:  SOD by condition_factor
## Kruskal-Wallis chi-squared = 250.73, df = 259, p-value = 0.6323
kruskal.test(SOD ~ avg_thickness, data = alldata)

##
##  Kruskal-Wallis rank sum test
##
## data:  SOD by avg_thickness
## Kruskal-Wallis chi-squared = 107.24, df = 110, p-value = 0.5567
```

Post hoc test. Kruskal-Wallac Multiple Comparisons

p450 and site_name show no true differences despite K-W test result.

SOD and site_name show a true site difference between Elliott Bay, Mrytle Edwards and Hood Canal only.

```
library(pgirmess)

mc_p450<- as.data.frame(kruskalmc(p450 ~ site_name, data = alldata, method = "bonferroni"))
mc_SOD<- as.data.frame(kruskalmc(SOD ~ site_name, data = alldata, method = "bonferroni"))
```