

# Ammonia Model

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## Setup

```
#load packages  
library(here)
```

```
## here() starts at /Users/summerheshong/Stats_Group_Project
```

```
library(tidyverse)
```

```
## -- Attaching core tidyverse packages ----- tidyverse 2.0.0 --  
## v dplyr      1.1.4      v readr      2.1.5  
## v forcats    1.0.0      v stringr   1.5.1  
## v ggplot2    3.5.1      v tibble    3.2.1  
## v lubridate  1.9.4      v tidyr     1.3.1  
## v purrr      1.0.2
```

```
## -- Conflicts ----- tidyverse_conflicts() --  
## x dplyr::filter() masks stats::filter()  
## x dplyr::lag()     masks stats::lag()  
## i Use the conflicted package (<http://conflicted.r-lib.org/>) to force all conflicts to become errors
```

```
library(lme4)
```

```
## Loading required package: Matrix  
##  
## Attaching package: 'Matrix'  
##  
## The following objects are masked from 'package:tidyr':  
##  
##     expand, pack, unpack
```

```
wetland <- read.csv(here('Data/Processed/Combined_Data_NArm.csv'))
```

# Research Question: What predicts ammonium levels in wetlands

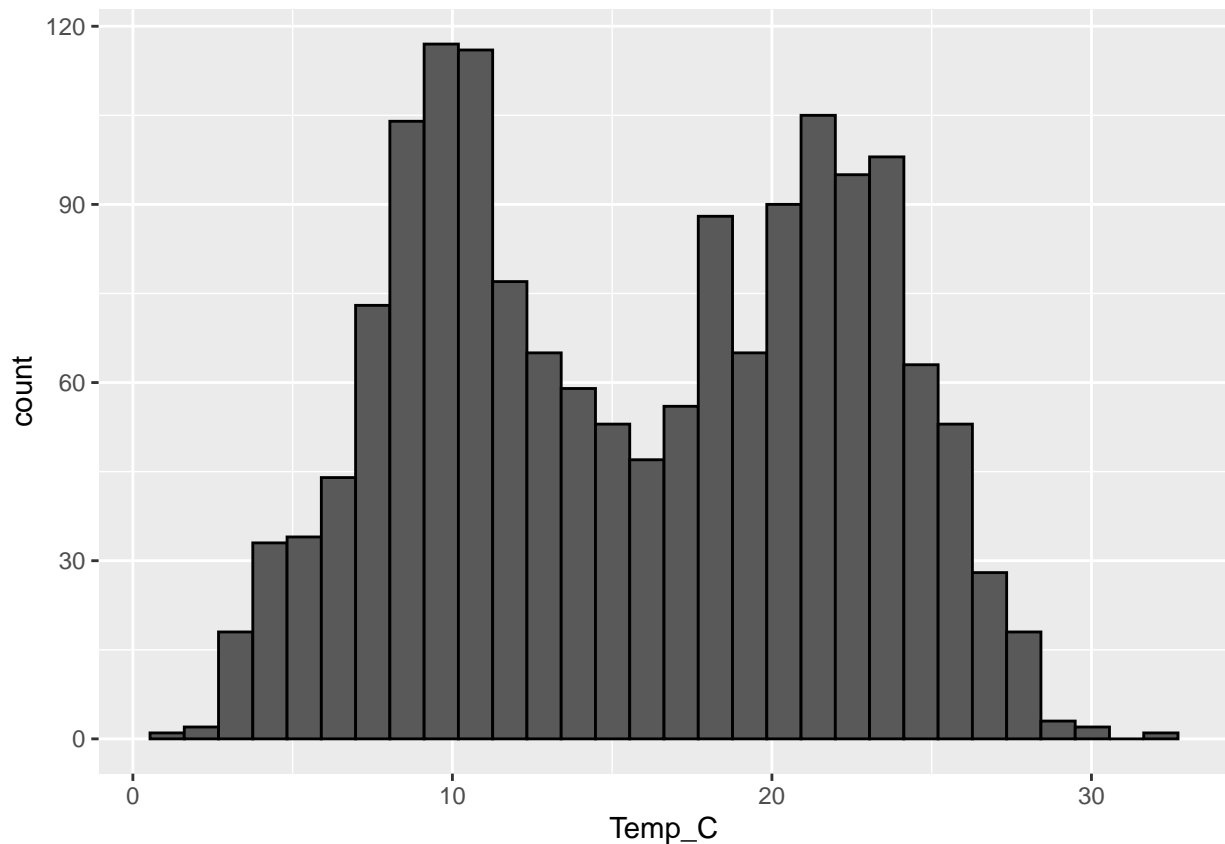
## Examine Data

display raw counts and distributions of data

```
#remove outliers: TDS 2262, row 1039; DO 224, row 840; TSS 516, row 641; fecal_coliform rows 1545 and 1549
wetland <- wetland [-c(147, 166, 641, 793, 840, 1035, 1039, 1545, 1549, 1614),]

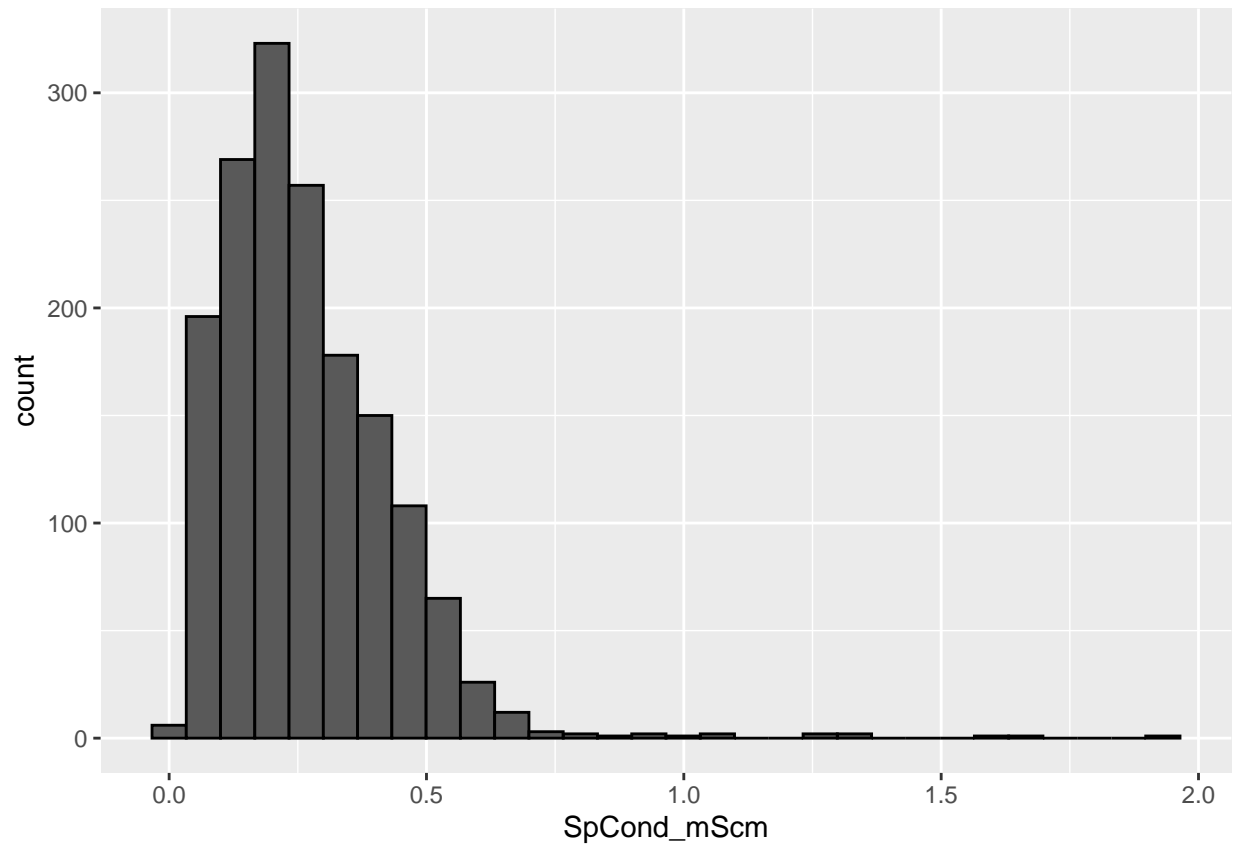
#create histograms
ggplot(wetland, aes(x = Temp_C)) +
  geom_histogram(color = 'black')

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



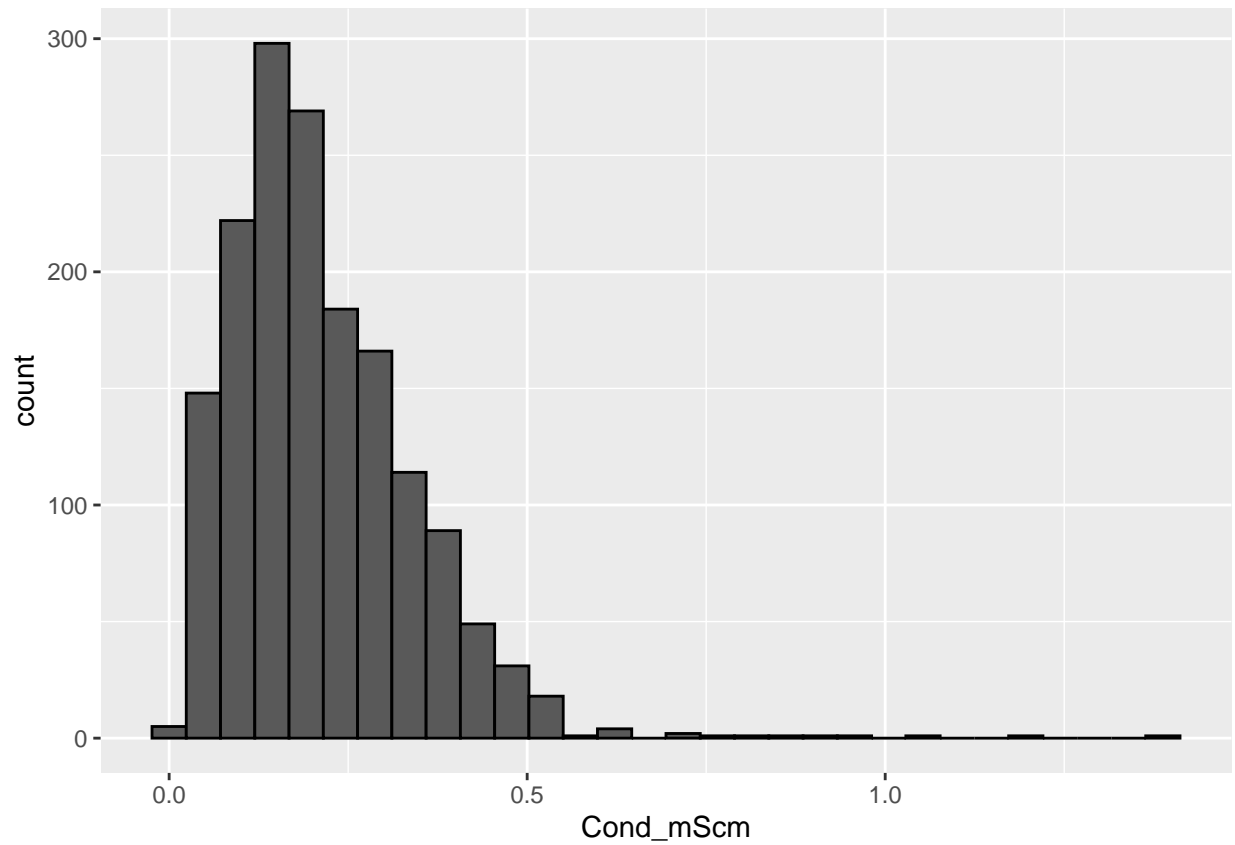
```
ggplot(wetland, aes(x = SpCond_mScm)) +
  geom_histogram(color = 'black')
```

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



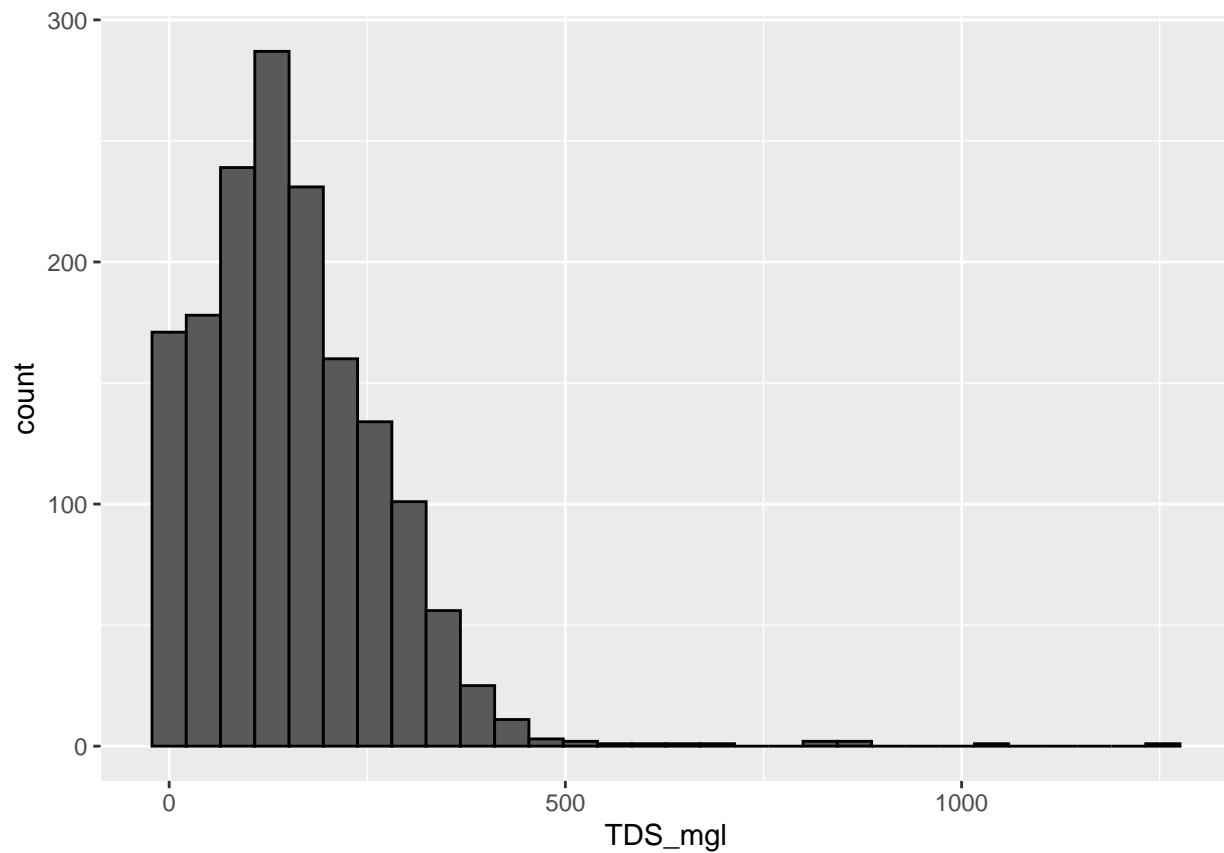
```
ggplot(wetland, aes(x = Cond_mScm)) +  
geom_histogram(color = 'black')
```

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



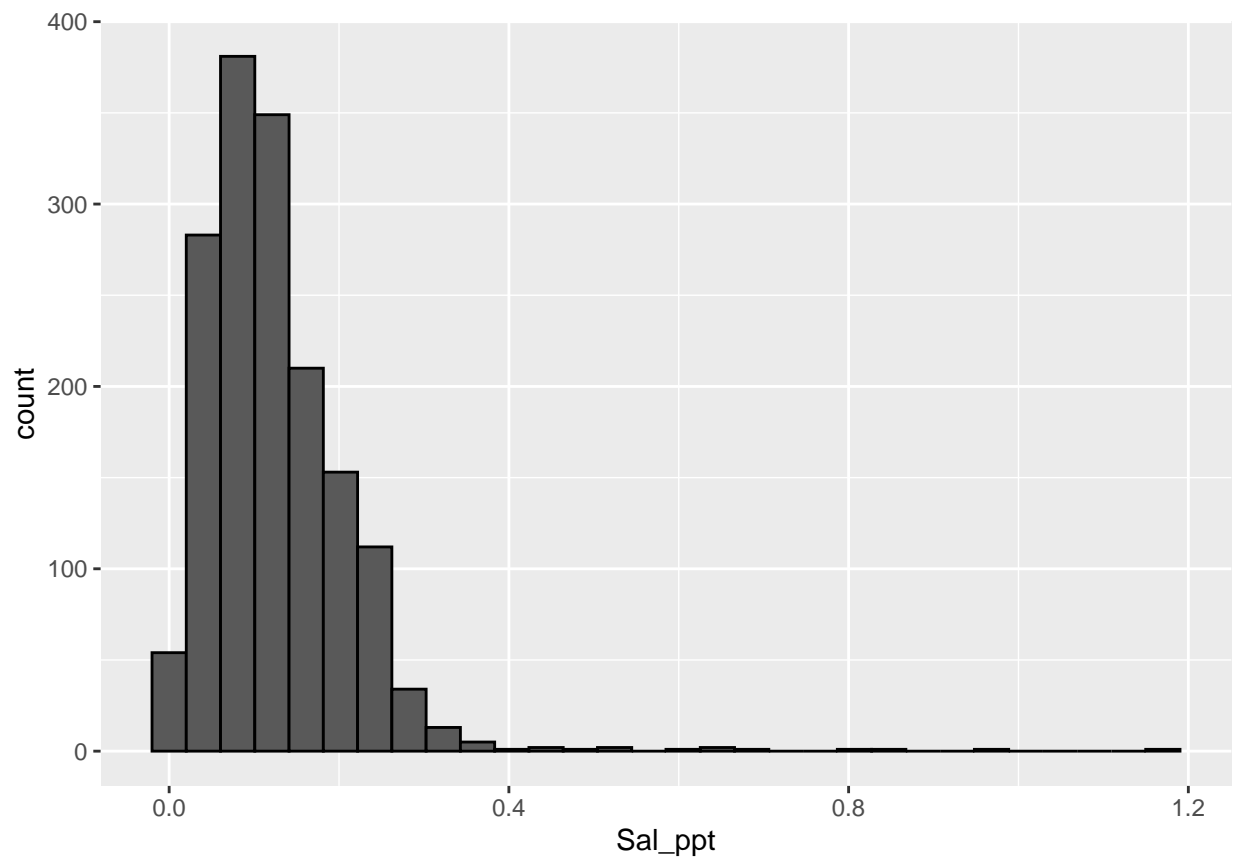
```
ggplot(wetland, aes(x = TDS_mgl)) +  
geom_histogram(color = 'black')
```

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



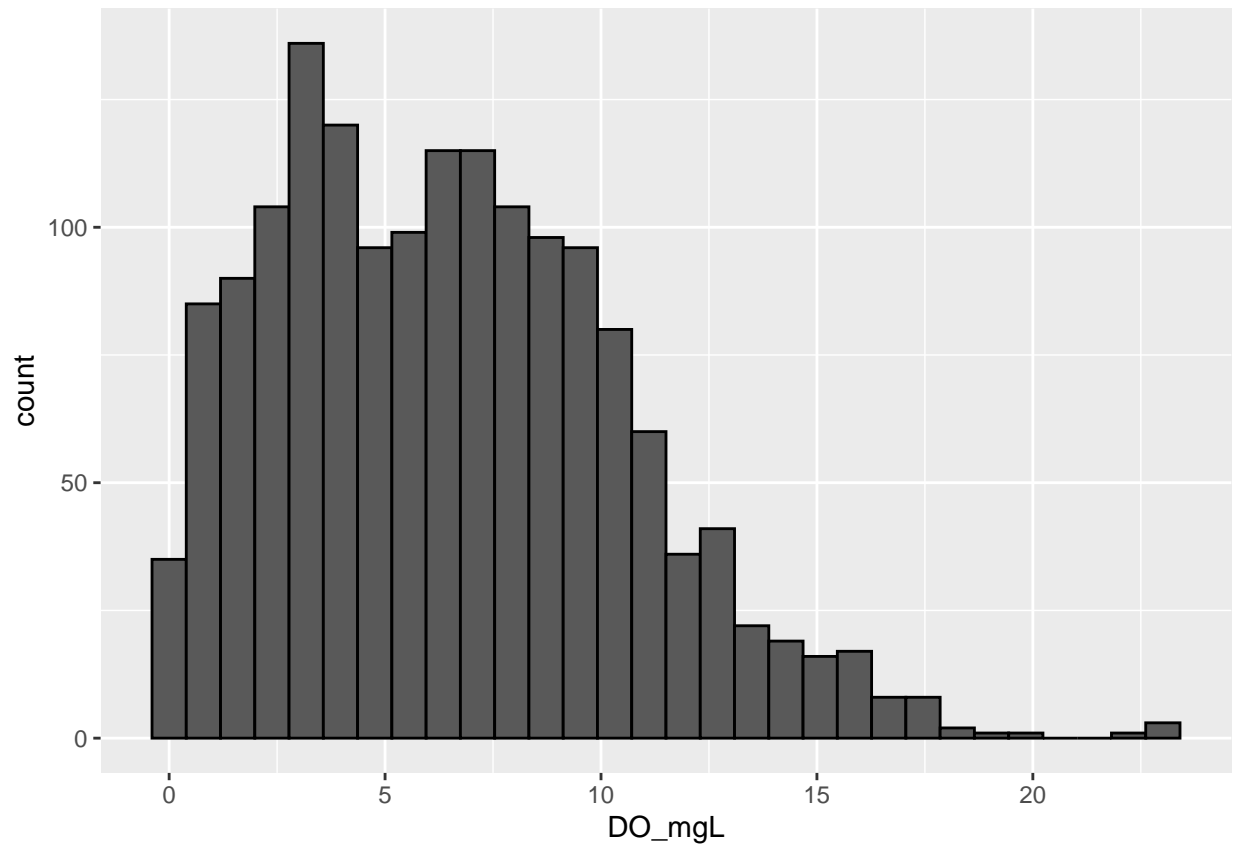
```
ggplot(wetland, aes(x = Sal_ppt)) +  
geom_histogram(color = 'black')
```

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



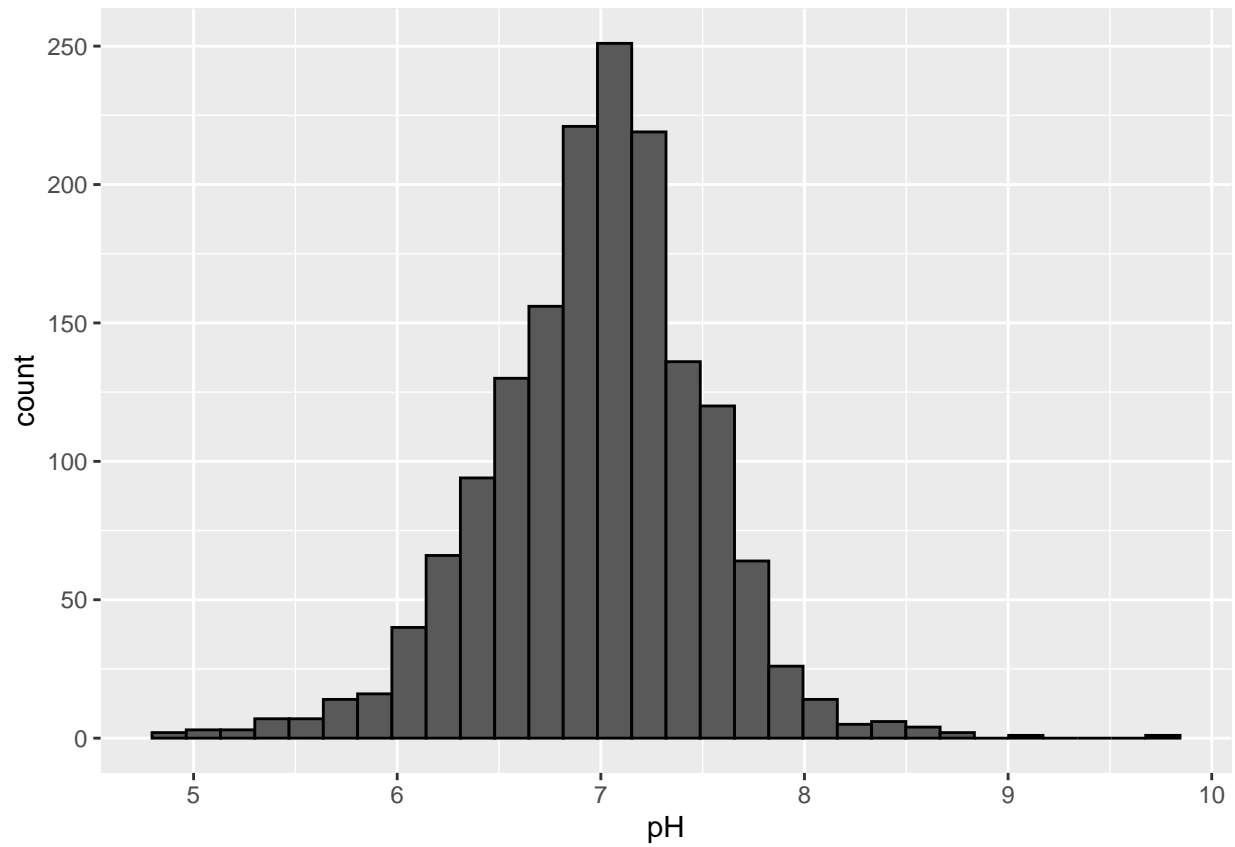
```
ggplot(wetland, aes(x = DO_mgL)) +  
geom_histogram(color = 'black')
```

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



```
ggplot(wetland, aes(x = pH)) +  
geom_histogram(color = 'black')
```

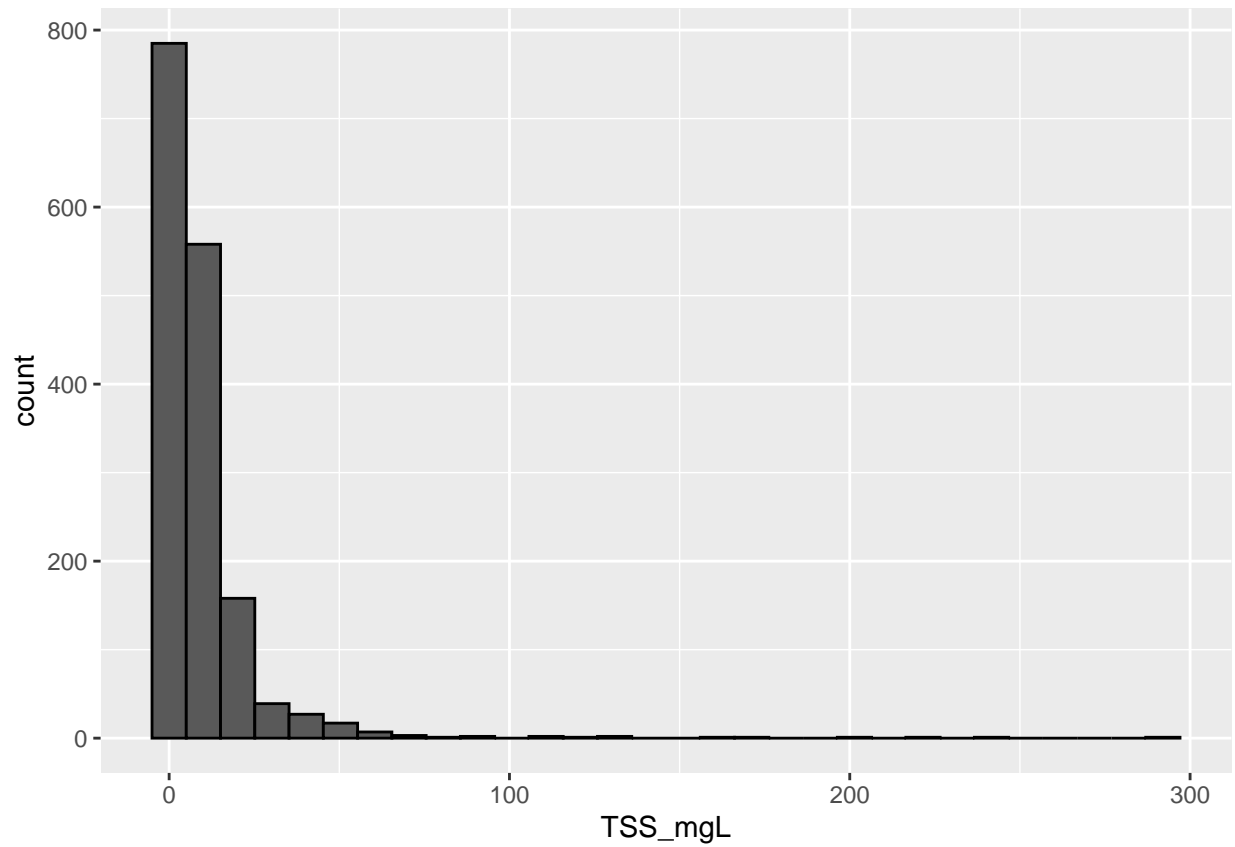
```
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
```



```
ggplot(wetland, aes(x = TSS_mgL)) +  
geom_histogram(color = 'black')
```

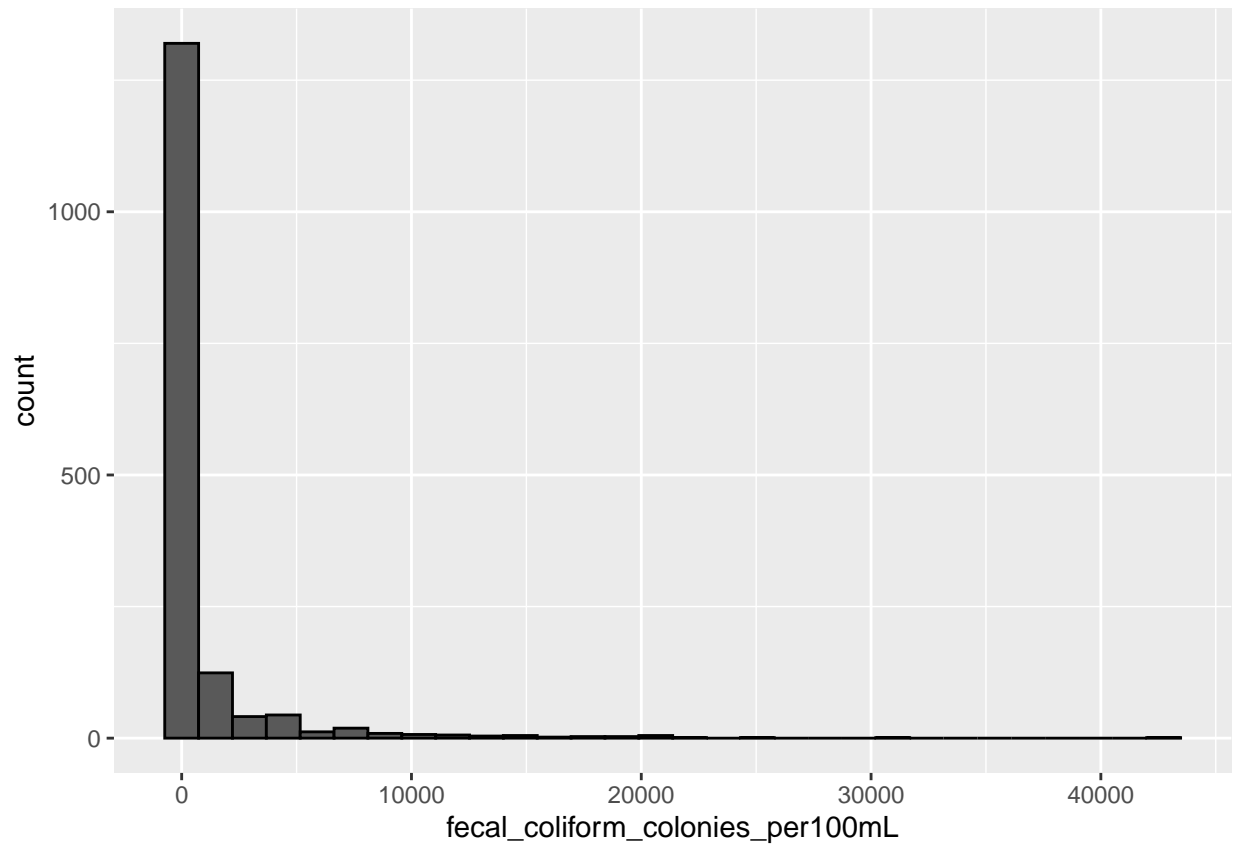
```
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
```





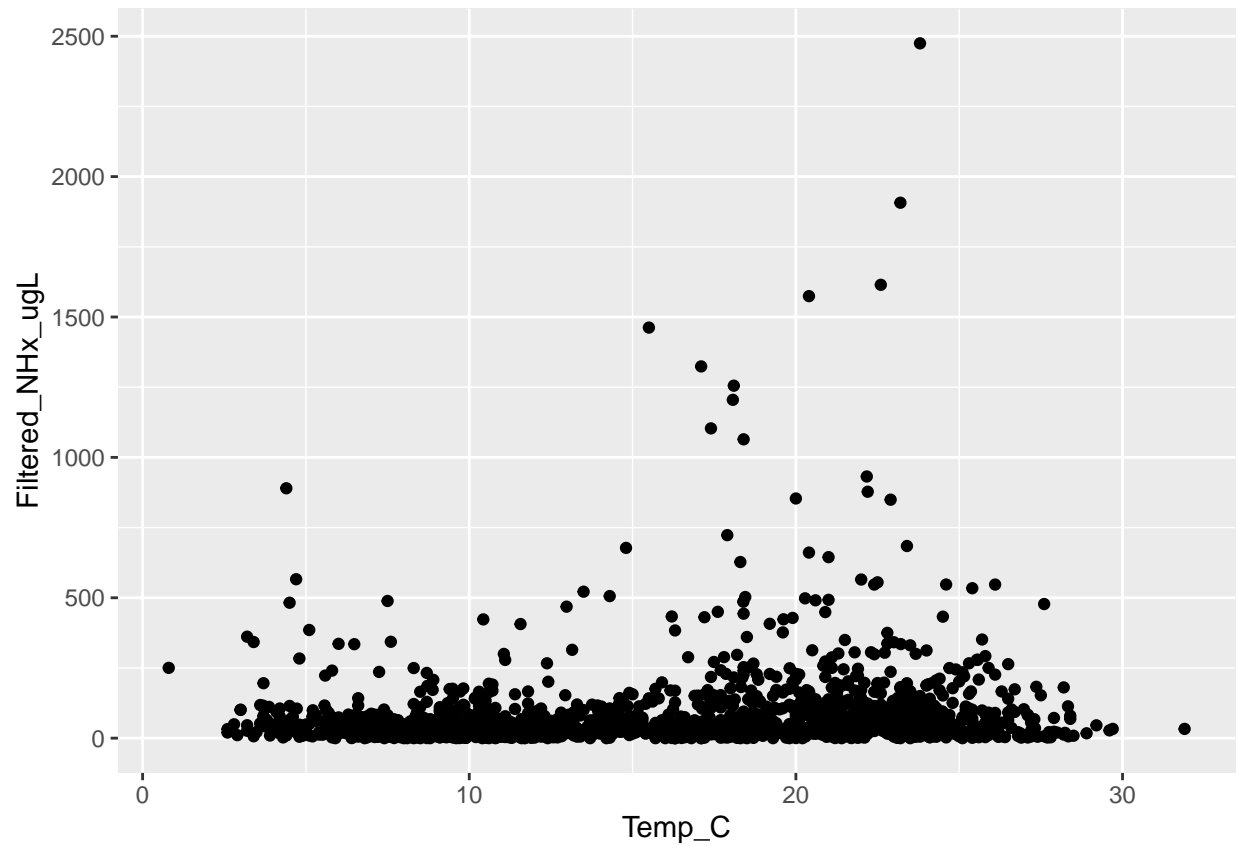
```
ggplot(wetland, aes(x = fecal_coliform_colonies_per100mL)) +  
  geom_histogram(color = 'black')
```

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

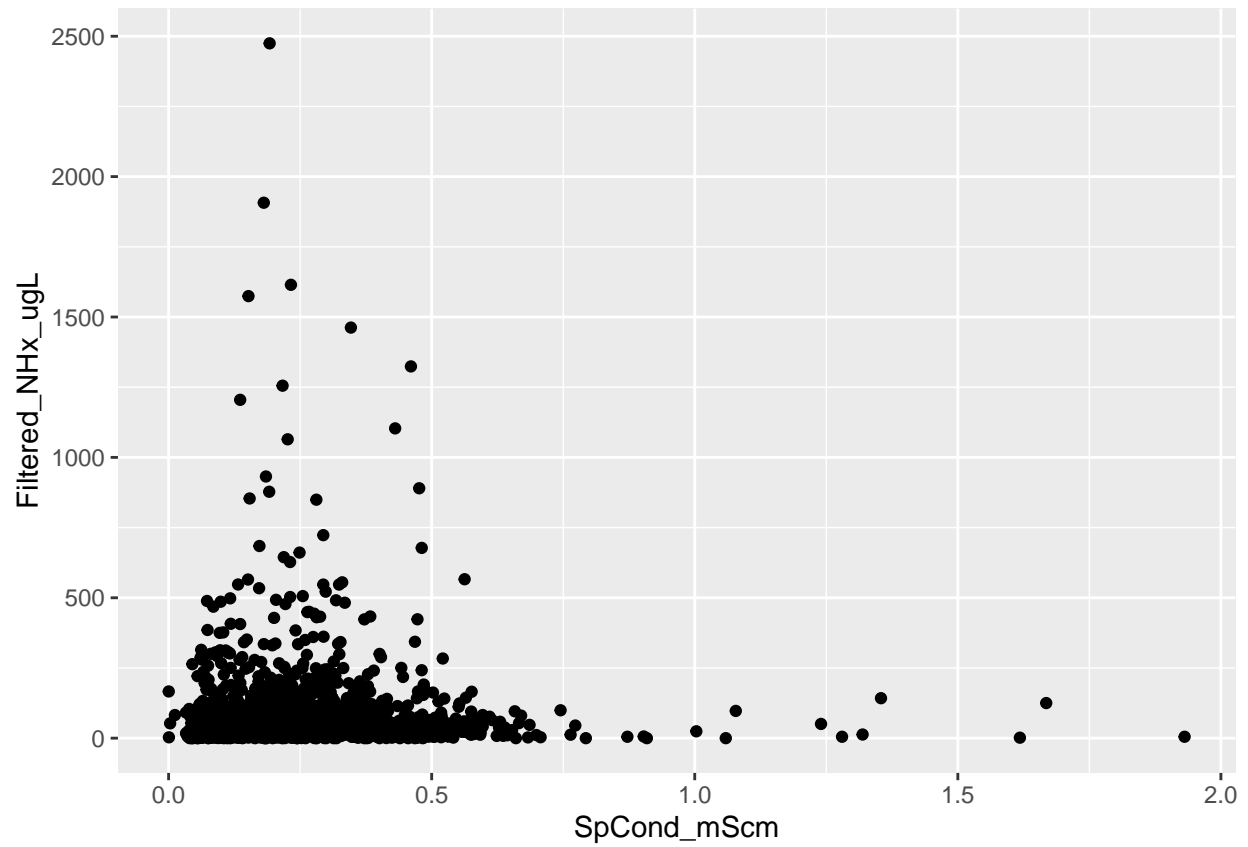


display relationships between predictor variables and outcome variable

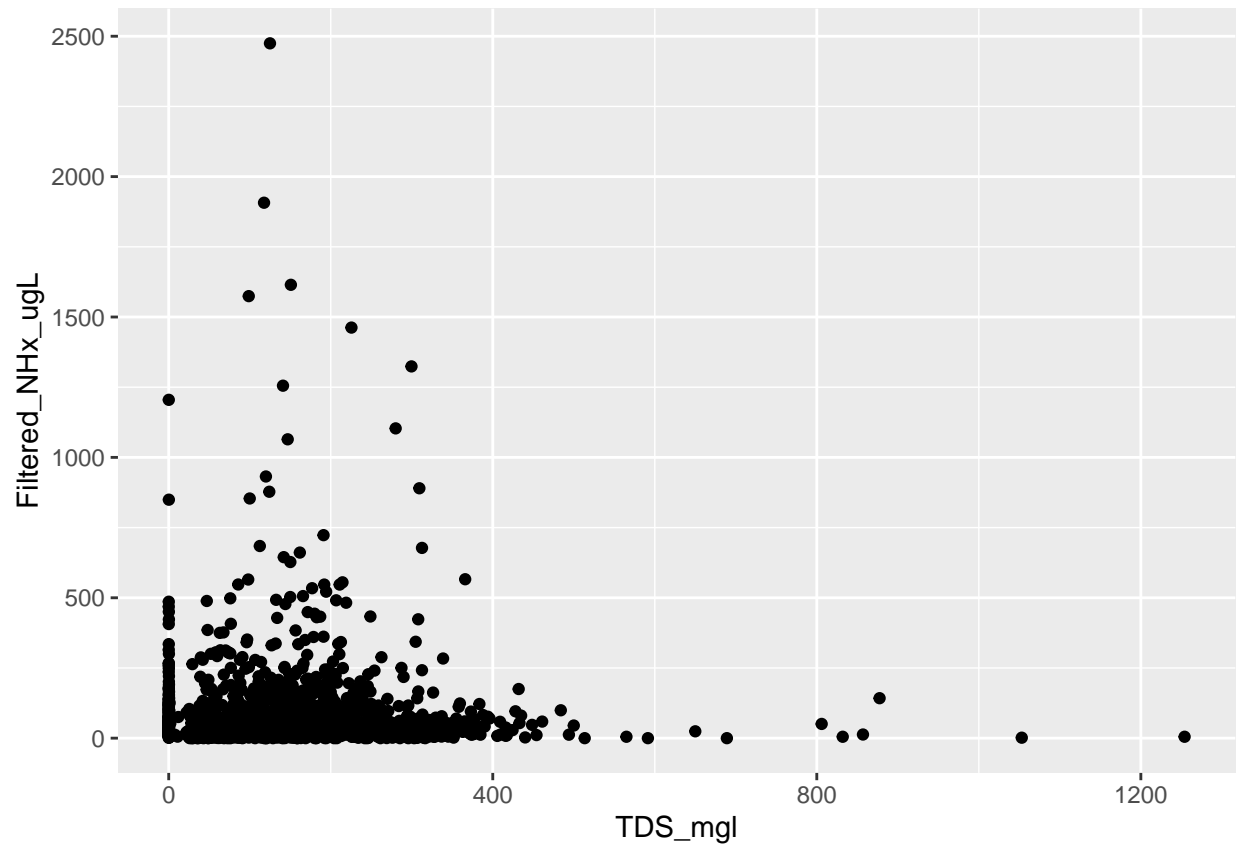
```
#create scatterplots  
ggplot(wetland, aes(x = Temp_C, y = Filtered_NHx_ugL)) +  
geom_point()
```



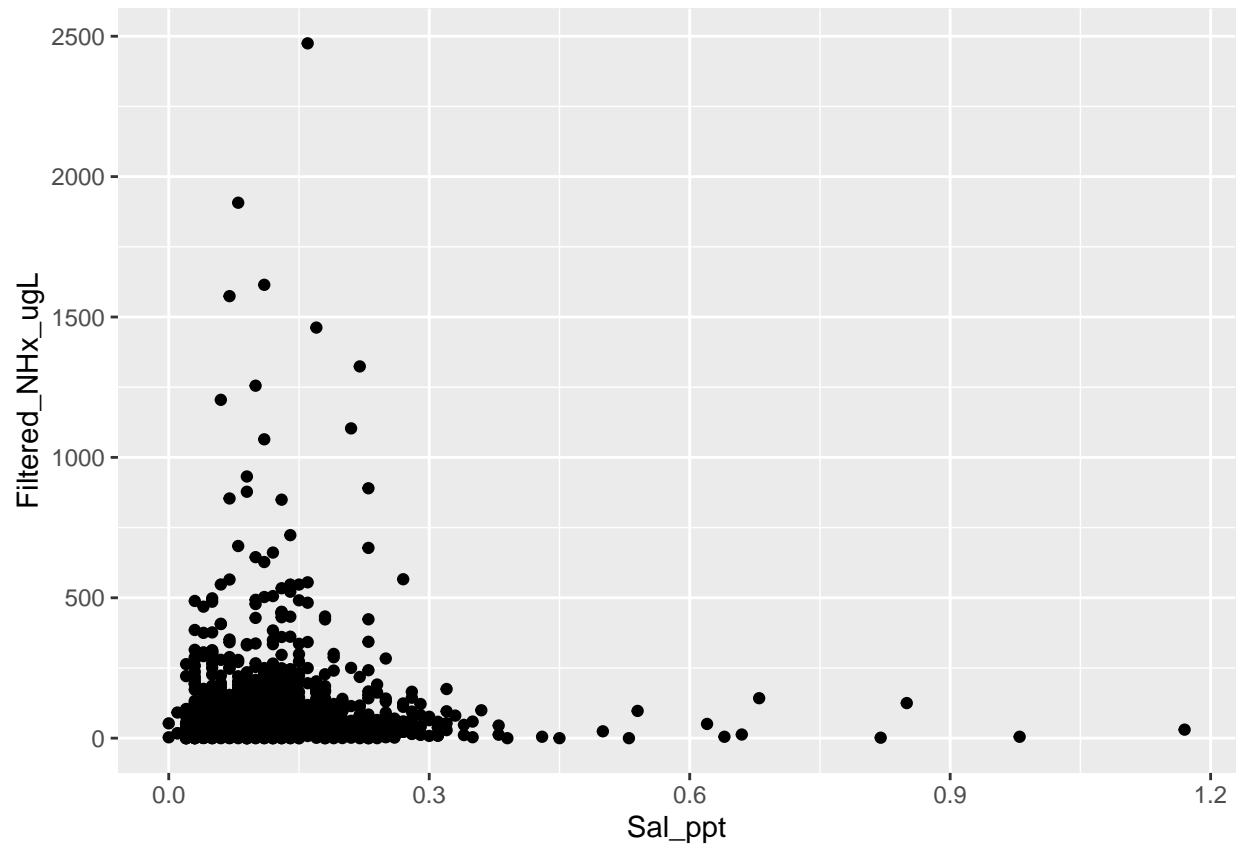
```
ggplot(wetland, aes(x = SpCond_mScm, y = Filtered_NHx_ugL)) +  
geom_point()
```



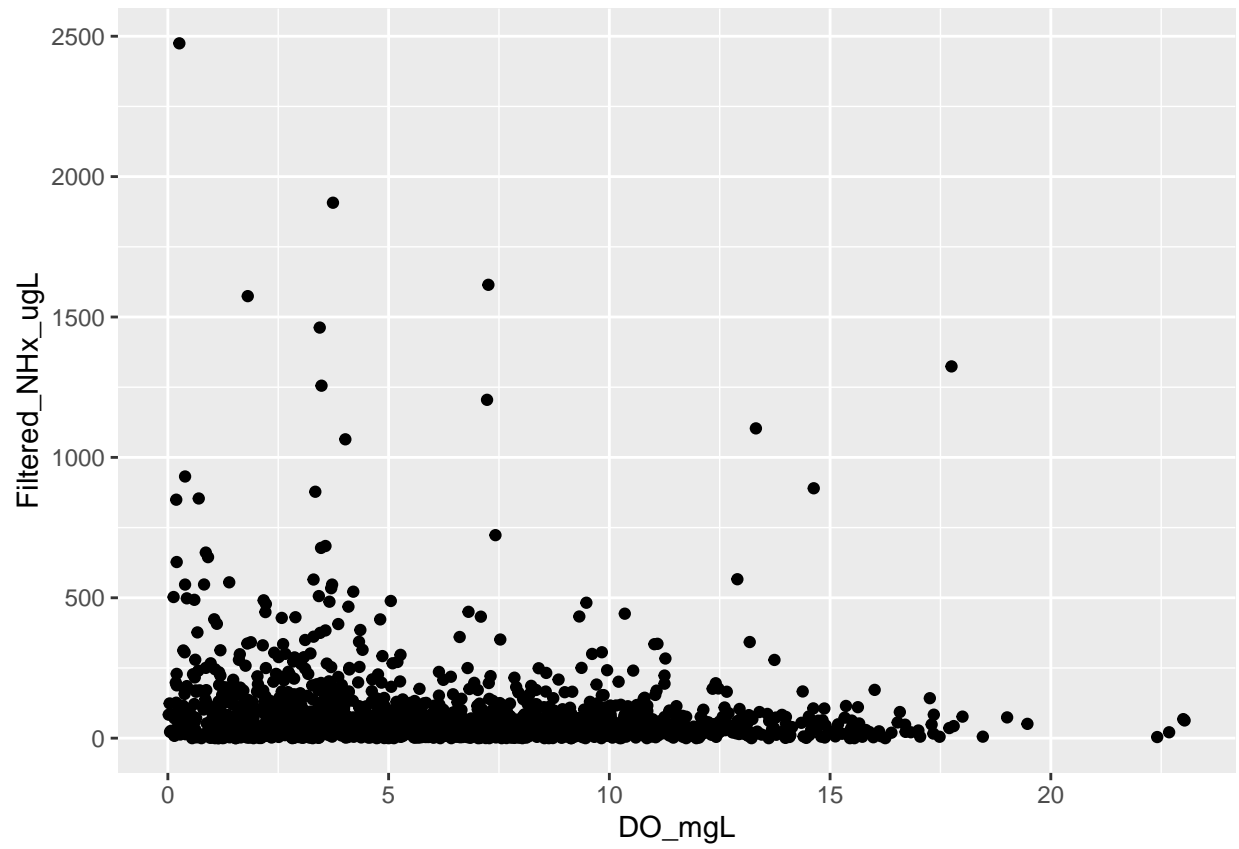
```
ggplot(wetland, aes(x = TDS_mgl, y = Filtered_NHx_ugL)) +  
geom_point()
```



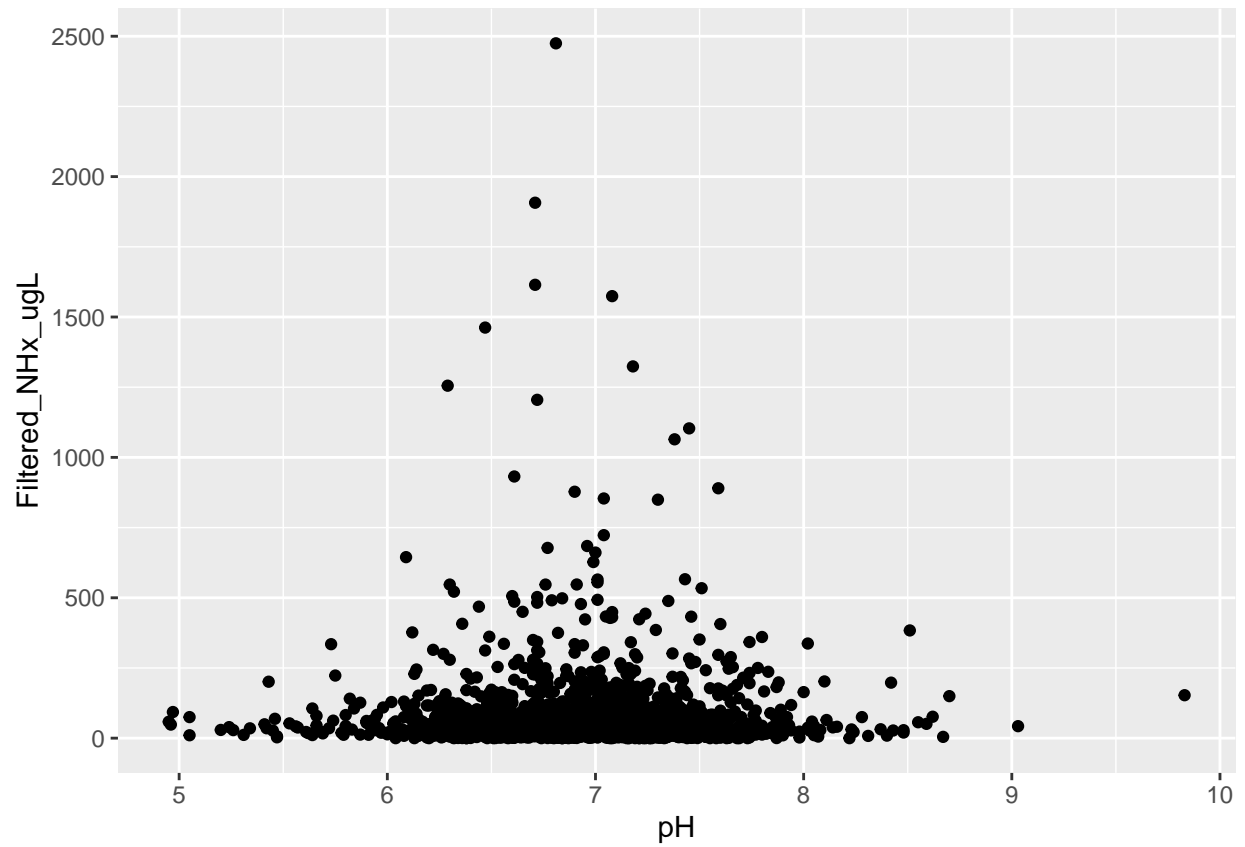
```
ggplot(wetland, aes(x = Sal_ppt, y = Filtered_NHx_ugL)) +  
geom_point()
```



```
ggplot(wetland, aes(x = DO_mgL, y = Filtered_NHx_ugL)) +  
geom_point()
```

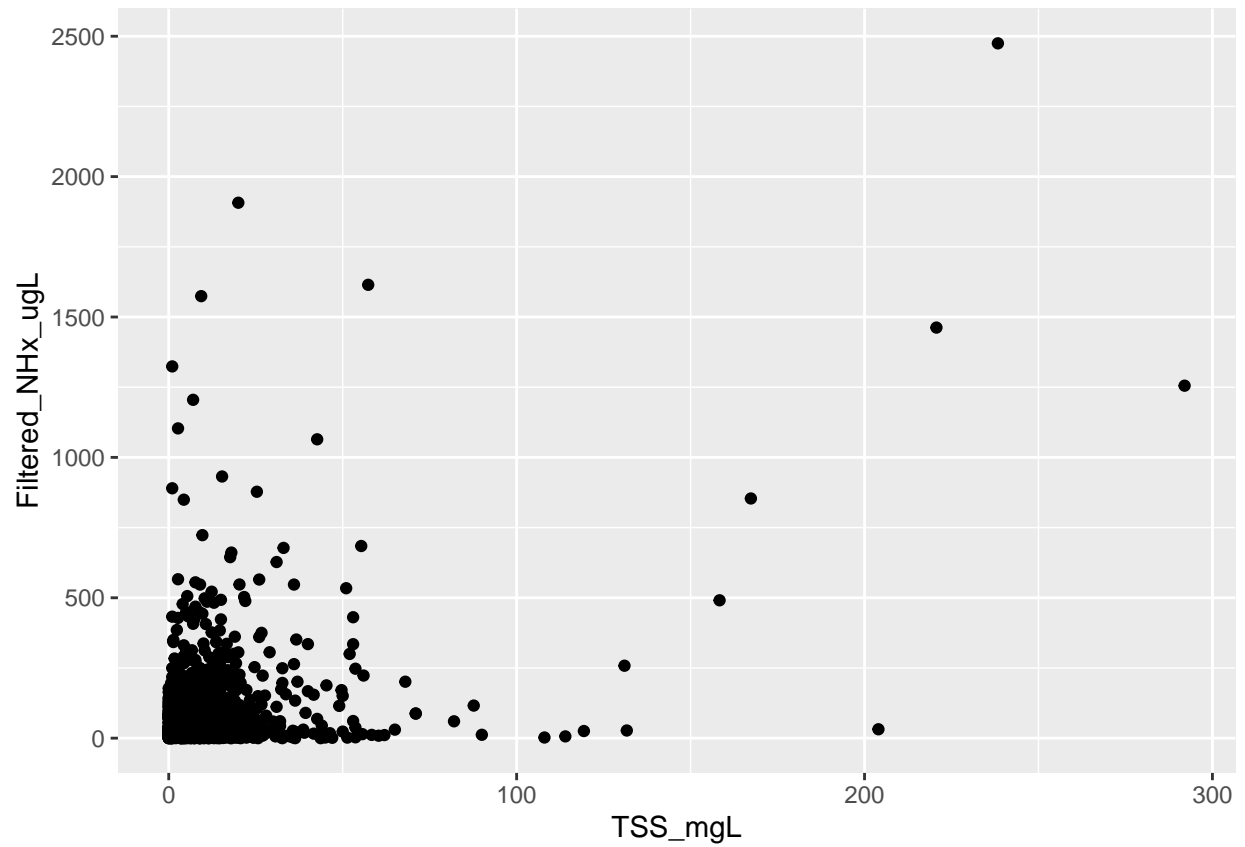


```
ggplot(wetland, aes(x = pH, y = Filtered_NHx_ugL)) +  
geom_point()
```

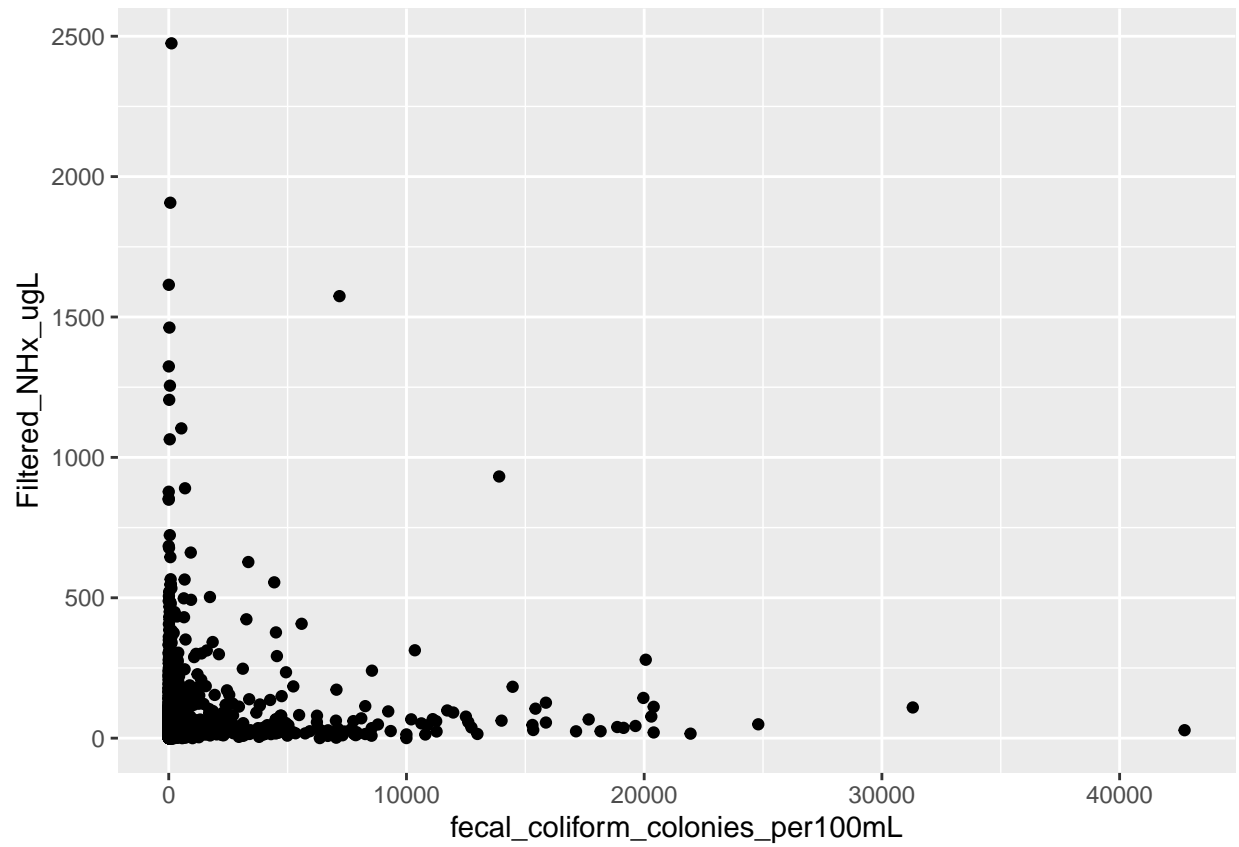


```
ggplot(wetland, aes(x = TSS_mgL, y = Filtered_NHx_ugL)) +  
geom_point()
```

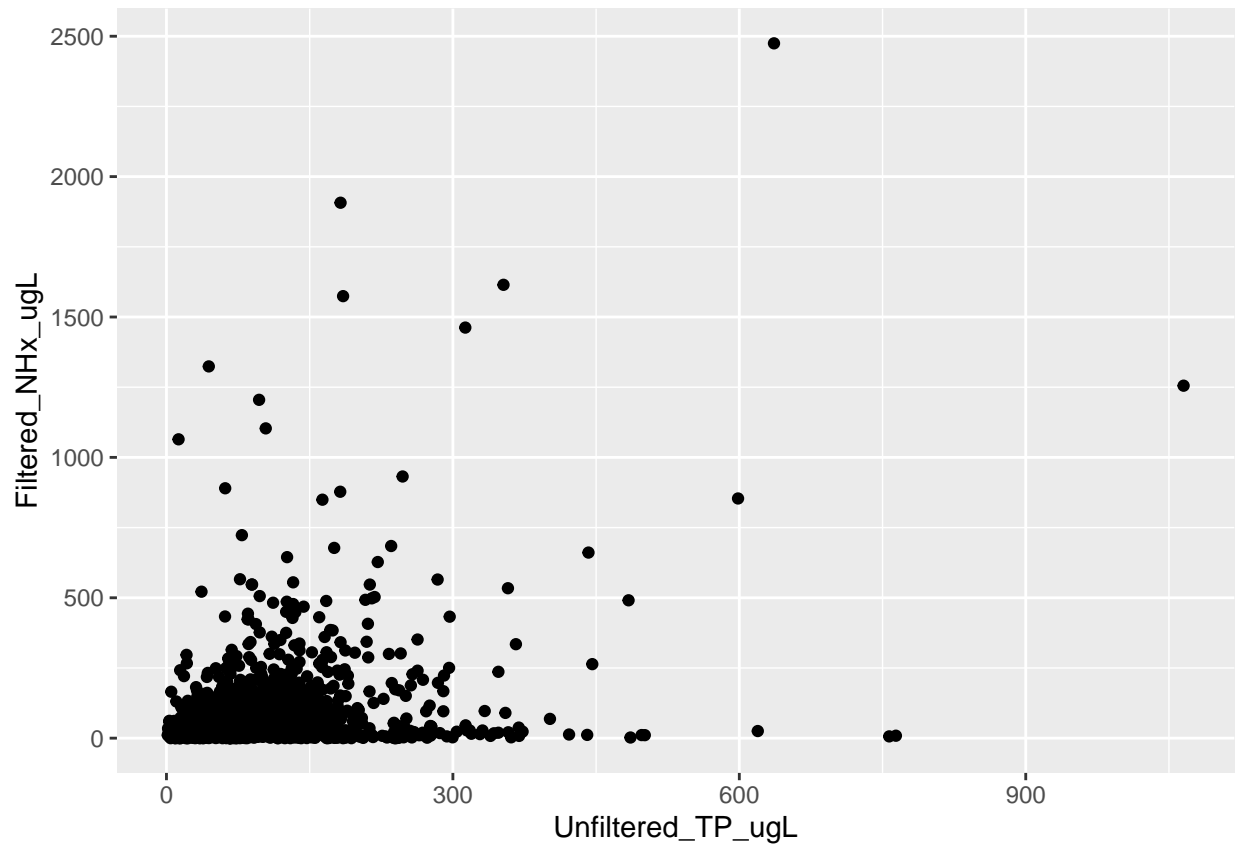




```
ggplot(wetland, aes(x = fecal_coliform_colonies_per100mL, y = Filtered_NHx_ugL)) +  
geom_point()
```



```
ggplot(wetland, aes(x = Unfiltered_TP_ugL, y = Filtered_NHx_ugL)) +  
geom_point()
```

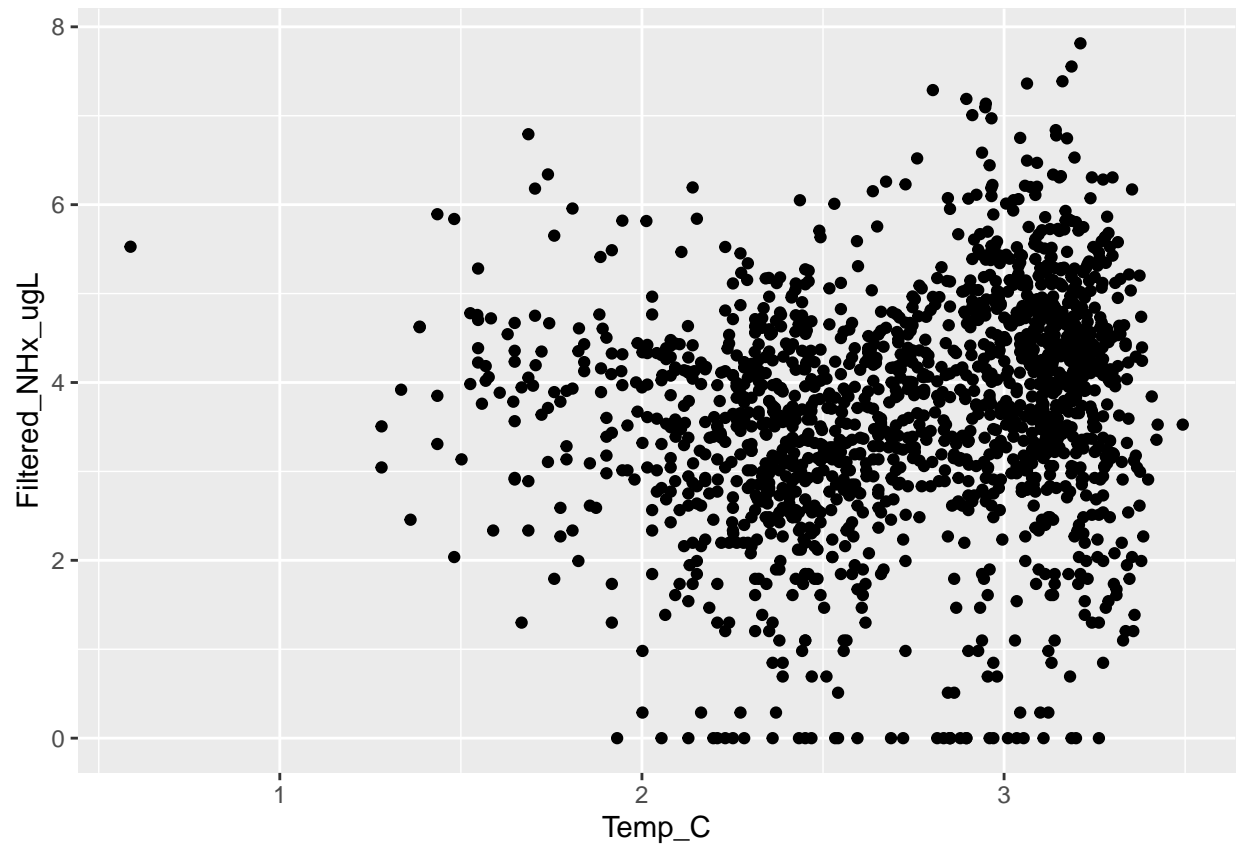


everything needs to be log transformed?

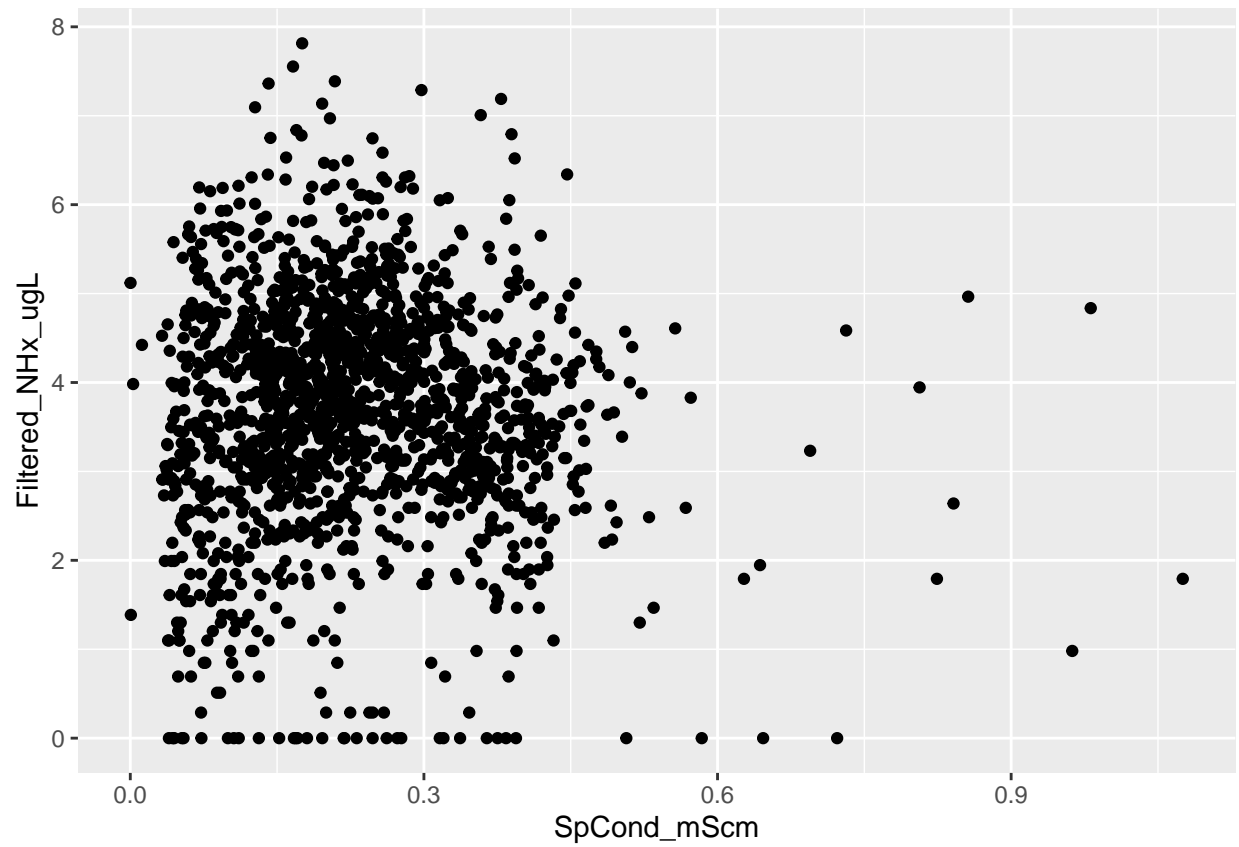
```
#log transform everything
log_wetland <- wetland %>%
  mutate(across(c(Temp_C, SpCond_mScm, Cond_mScm,
                  TDS_mgl, Sal_ppt, DO_mgL, pH,
                  Filtered_NHx_ugL, TSS_mgL,
                  fecal_coliform_colonies_per100mL, Unfiltered_TP_ugL),
    ~ log(.+1)))

#re-vistualize data

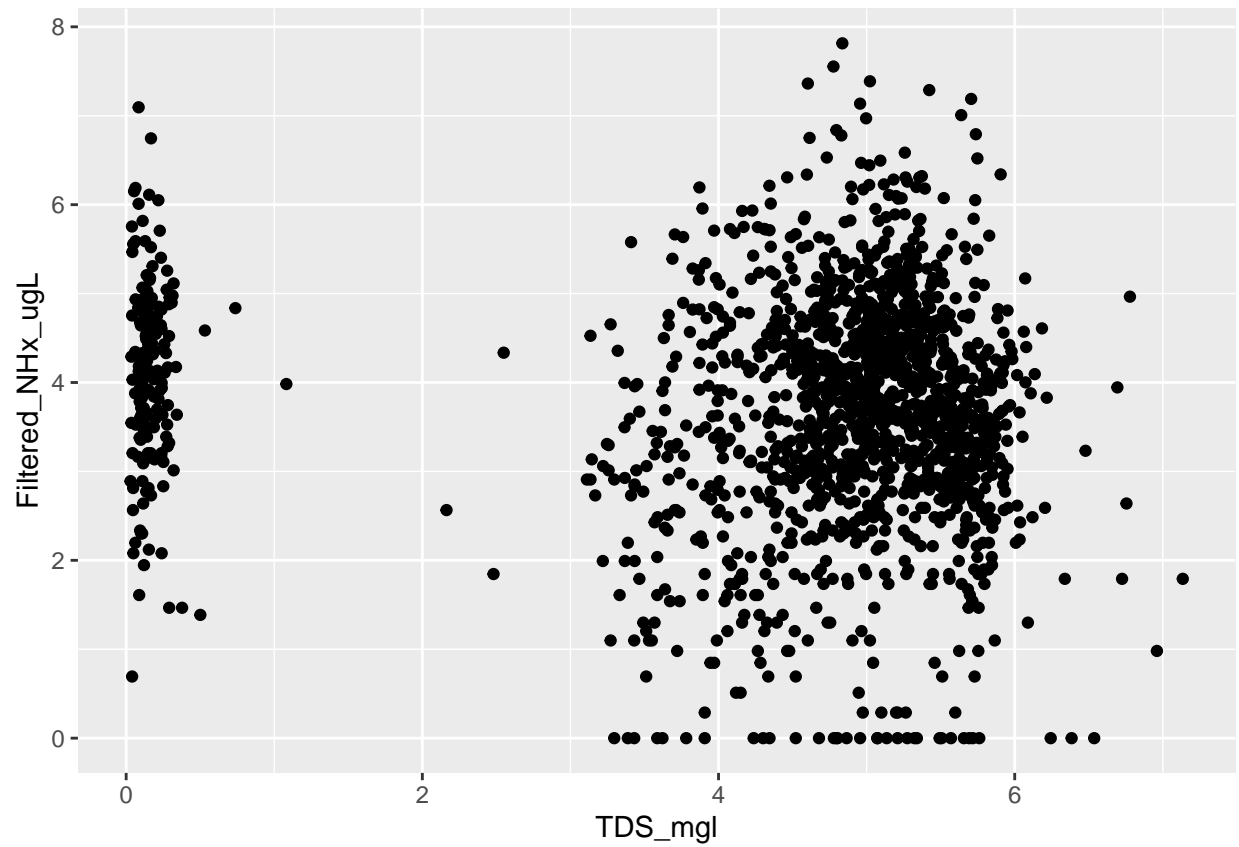
ggplot(log_wetland, aes(x = Temp_C, y = Filtered_NHx_ugL)) +
  geom_point()
```



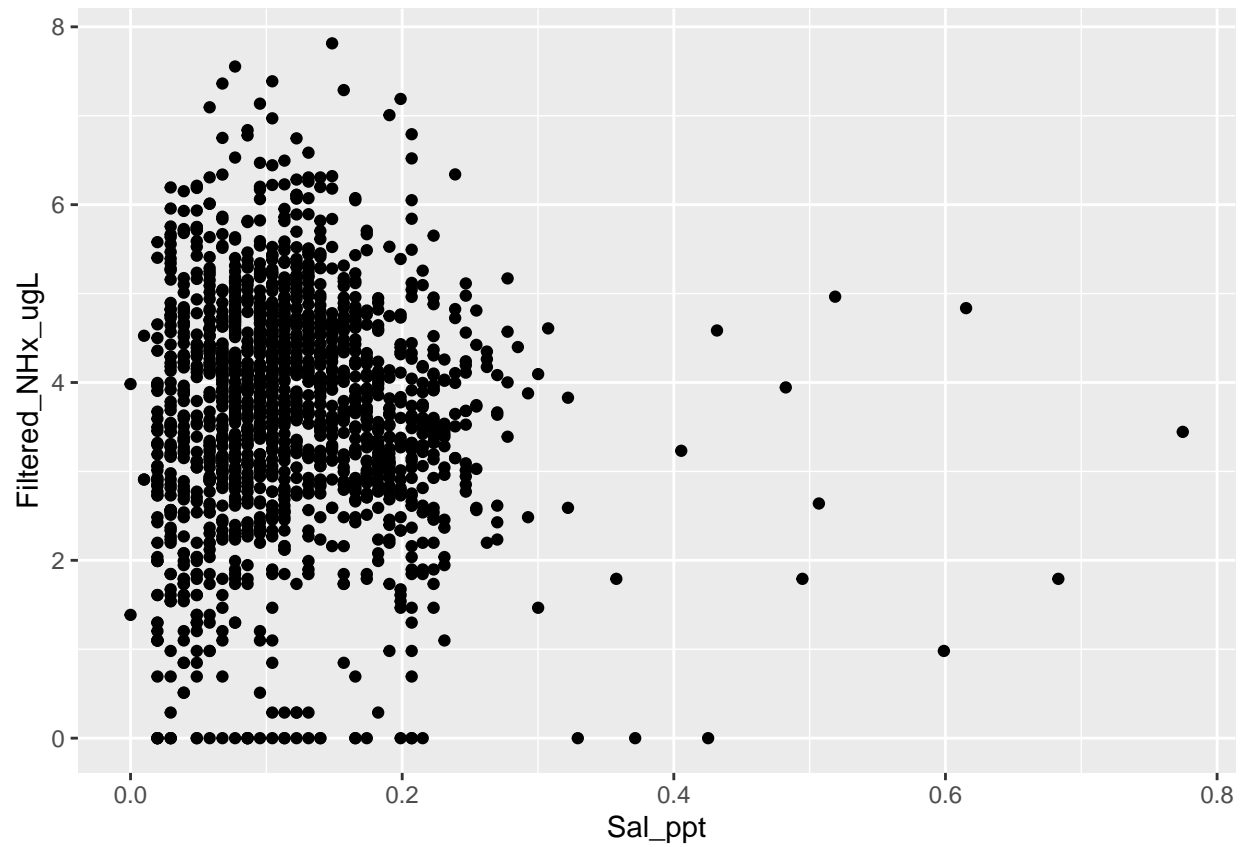
```
ggplot(log_wetland, aes(x = SpCond_mScm, y = Filtered_NHx_ugL)) +  
geom_point()
```



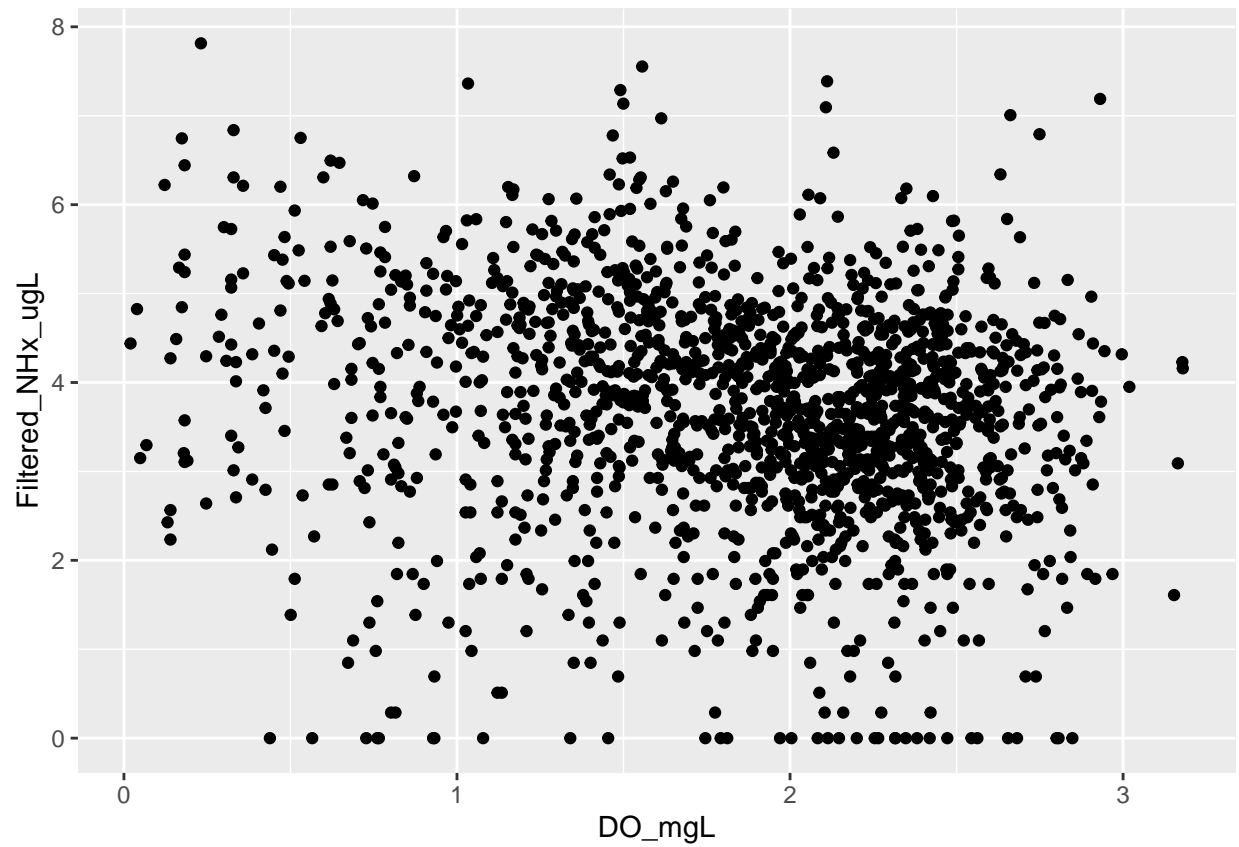
```
ggplot(log_wetland, aes(x = TDS_mgl, y = Filtered_NHx_ugL)) +  
geom_point()
```



```
ggplot(log_wetland, aes(x = Sal_ppt, y = Filtered_NHx_ugL)) +  
geom_point()
```

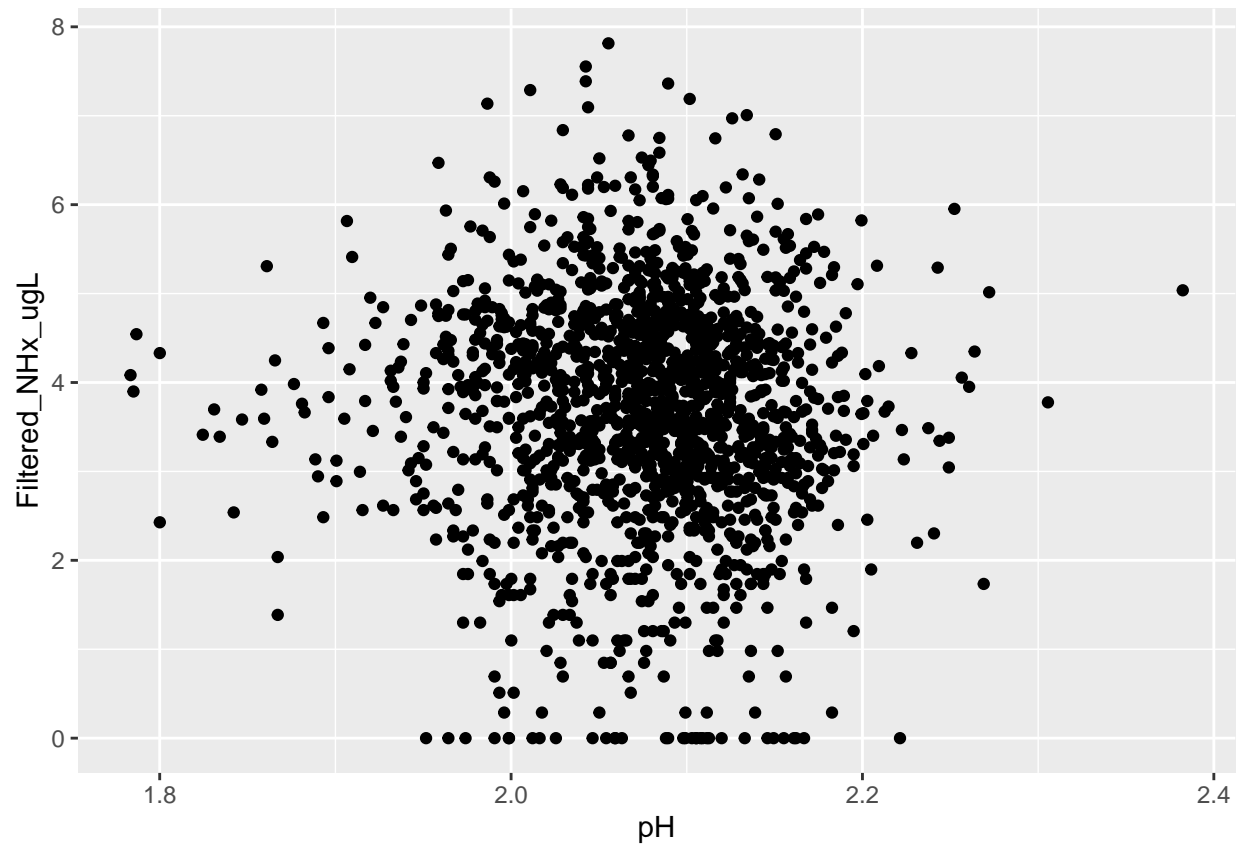


```
ggplot(log_wetland, aes(x = DO_mgL, y = Filtered_NHx_ugL)) +  
geom_point()
```

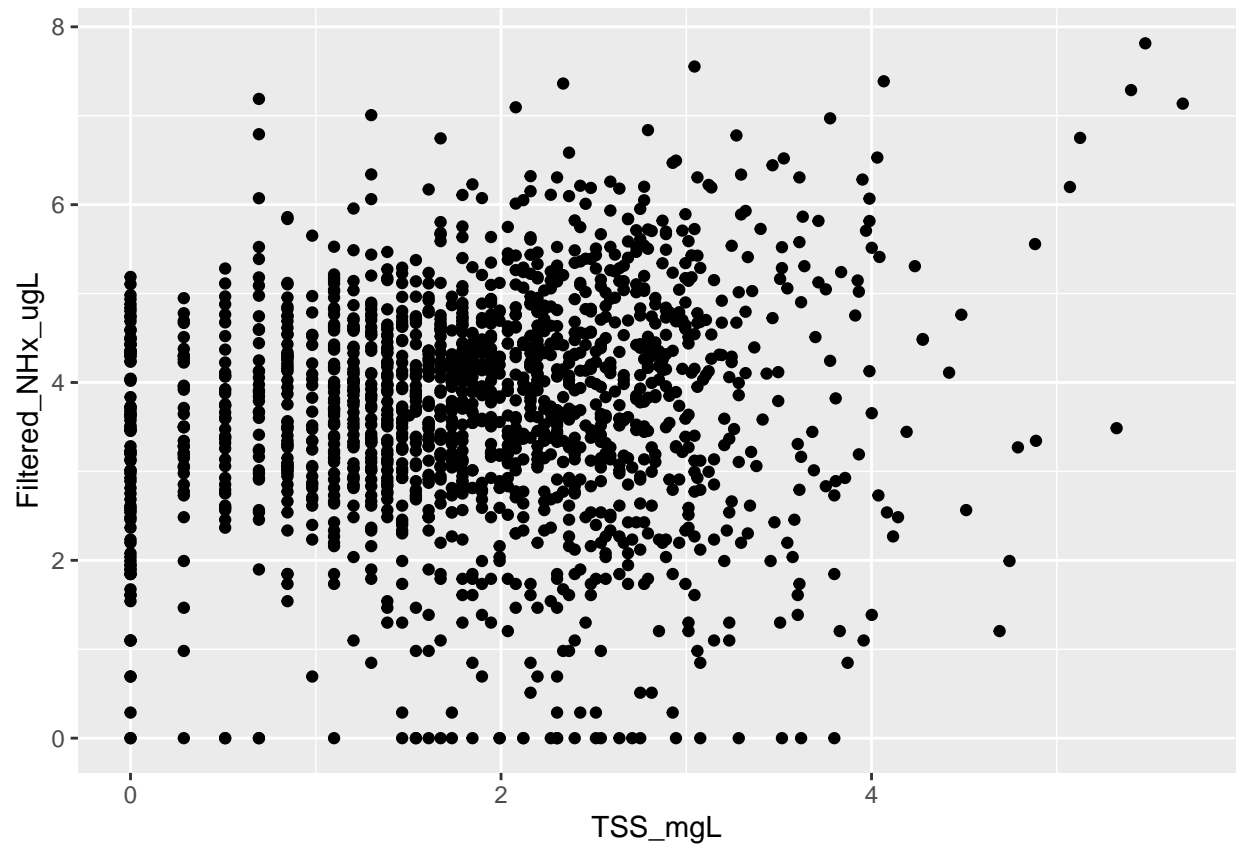


```
ggplot(log_wetland, aes(x = pH, y = Filtered_NHx_ugL)) +  
geom_point()
```

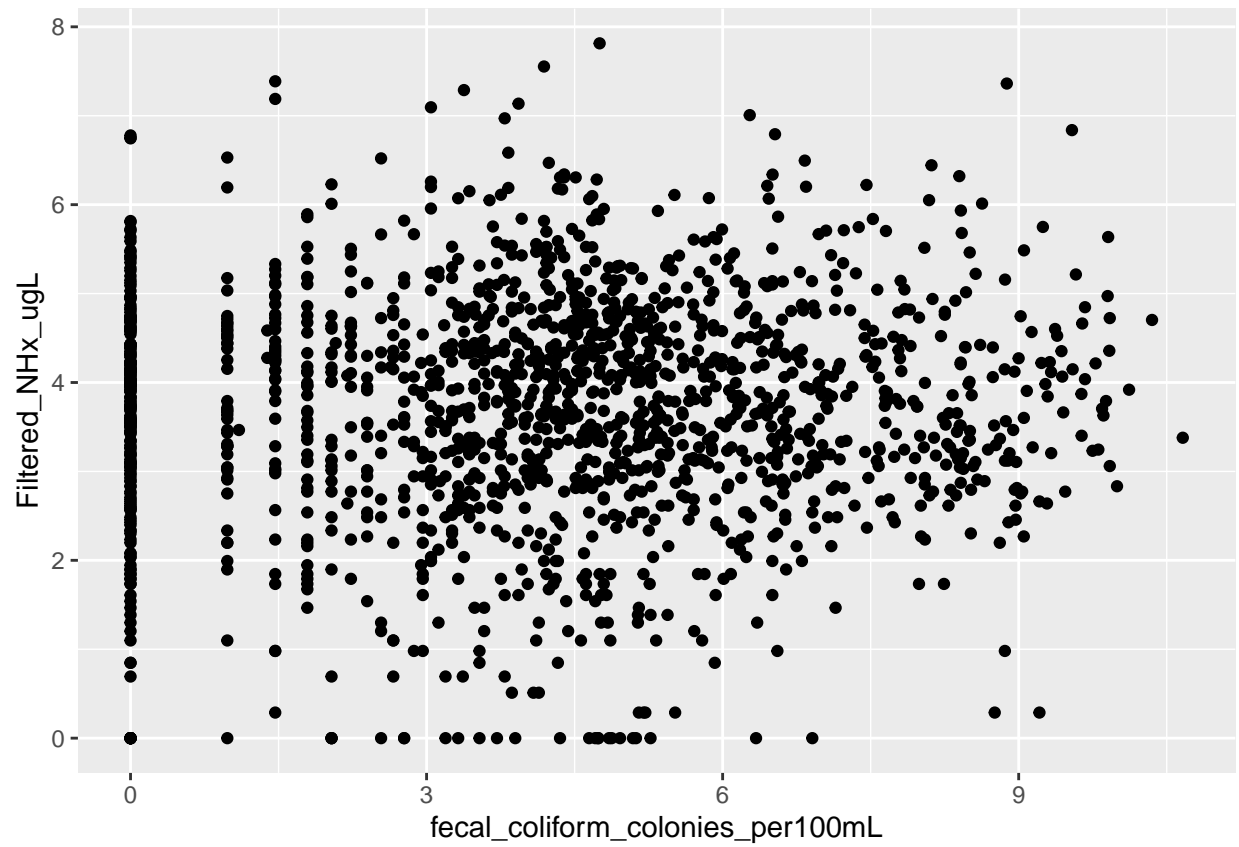




```
ggplot(log_wetland, aes(x = TSS_mgL, y = Filtered_NHx_ugL)) +  
geom_point()
```



```
ggplot(log_wetland, aes(x = fecal_coliform_colonies_per100mL, y = Filtered_NHx_ugL)) +  
geom_point()
```



```
ggplot(log_wetland, aes(x = Unfiltered_TP_ugL, y = Filtered_NHx_ugL)) +  
geom_point()
```

