# Project Title: Effects of hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>) on the growth of different bloom-forming cyanobacteria

# 1. Creating the "FACETED LINE PLOT" **#Load Necessary Packages** library(readxl) library(dplyr) library(writexl) library(ggplot2) library(scales) # Set the file path file\_path <- "C:/Users/ASUS/Desktop/Exp\_1.9.xlsx" # Read the data from the first sheet data <- read\_excel(file\_path, sheet = 1) # Calculate the average and standard deviation data\_summary <- data %>% mutate( Average = rowMeans(select(., R1, R2), na.rm = TRUE), SD = apply(select(., R1, R2), 1, sd, na.rm = TRUE)) # View the result print(data\_summary) # CREATING PLOT # Make sure variables are formatted correctly exp <- data\_summary %>% mutate( Treatment = as.factor(Treatment), Pigment = as.factor(Pigment), Phytoplankton = as.factor(Phytoplankton) # Create the plot ggplot(exp, aes(x = Day, y = Average, color = Treatment, fill = Treatment)) + geom\_point(size = 2) + geom\_errorbar(aes(ymin = Average - SD, ymax = Average + SD), width = 0.3) + geom\_smooth(method = "loess", se = FALSE) + facet\_grid(Pigment ~ Phytoplankton, scales = "free") + theme(text = element\_text(size = 14)) + labs( title = "Effects of Hydrogen Peroxide on Cyanobacteria",

subtitle = "Exp\_1.9",

x = "Days",

```
y = "Pigment (RFU)",
 color = "Treatment",
 shape = "Treatment"
) +
scale_y_continuous(labels = comma) +
theme_minimal()
2. ANOVA
# Load required packages
library(readxl)
library(dplyr)
library(tidyr)
library(writexl)
library(ggplot2)
library(openxlsx)
# Load data
raw_data <- read_excel("C:/Users/ASUS/Desktop/Exp_1.9.xlsx", sheet = 1)
# Convert wide to long format
data <- raw_data %>%
pivot_longer(cols = starts_with("R"), names_to = "REP", values_to = "READ") %>%
mutate(
 REP = ifelse(REP == "R1", 1, 2),
 log10READ = log10(READ)
# Step 3: Load the existing workbook
wb <- loadWorkbook("C:/Users/ASUS/Desktop/Exp_1.9.xlsx")
addWorksheet(wb, "Sheet2")
writeData(wb, sheet = "Sheet2", data)
# Step 5: Save the updated workbook
saveWorkbook(wb, "C:/Users/ASUS/Desktop/Exp_1.9.xlsx", overwrite = TRUE)
#factor Treatment
data$Treatment <- as.factor(data$Treatment)
# Filter only Chl-a pigment
chl_data <- data %>%
filter(Pigment == "Chl-a")
# Calculate growth rate: slope of ln(Chl-a) vs Day for each Phytoplankton and Treatment
growth_rates <- chl_data %>%
group_by(Phytoplankton, Treatment, REP) %>%
```

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arrange(Day) %>%
summarise(Growth_Rate = coef(lm(log10READ ~ Day))[2], .groups = "drop")
```

# # View results

print(growth\_rates)

# # Save the growth rate data to an Excel file

write\_xlsx(growth\_rates, "Cyanobacteria\_Growth\_Rates.xlsx")

#### 2.1 Filter by phytoplankton - MC first

MC\_growth<- growth\_rates %>% filter(Phytoplankton == "Microcystis LE21")

## # ANOVA (indicates that a single factor has a significant effect on your population)

MC\_aov <- aov(Growth\_Rate~Treatment, data=MC\_growth) summary(MC\_aov)

#### # Create a formatted table

anova\_table <- as.data.frame(anova(MC\_aov))</pre>

#### # Save the table as CSV

write.csv(format(anova\_table, scientific = FALSE), "MC\_anova\_results\_clean.csv")

## 2.2 Filter by phytoplankton - PLK

PLK\_growth<- growth\_rates %>% filter(Phytoplankton == "Planktothrix 1808")

#### # ANOVA

PLK\_aov <- aov(Growth\_Rate~Treatment, data=PLK\_growth) summary(PLK\_aov)

# # Create a formatted table

anova\_table <- as.data.frame(anova(PLK\_aov))</pre>

# # Save the table as CSV

write.csv(format(anova\_table, scientific = FALSE), "PLK\_anova\_results\_clean.csv")

## 2.3 Filter by phytoplankton - FIS

FIS\_growth<- growth\_rates %>%
filter(Phytoplankton == "Fischerella 1.5")

#### # ANOVA

FIS\_aov <- aov(Growth\_Rate~Treatment, data=FIS\_growth) summary(FIS\_aov)

#### # Create a formatted table

anova\_table <- as.data.frame(anova(FIS\_aov))</pre>

#### # Save the table as CSV

write.csv(format(anova\_table, scientific = FALSE), "FIS\_anova\_results\_clean.csv")