

Week 1 Handout: Basic R

Z620: Quantitative Biodiversity, Indiana University

January 16, 2015

OVERVIEW

This handout builds from the Week 0 introduction to R and RStudio, which highlighted operators, vectors, and simple commands that will be useful to you during the course and beyond. Now, you will be introduced to some new data types, specifically matrices and dataframes. After learning how to work with these data types, you will learn about contributed packages, which will aid in the visualization of data and with performing descriptive and hypothesis-testing statistical procedures, including linear regression and ANOVA.

1) SETTING YOUR WORKING DIRECTORY

A good first step when you sit down to work in R is to clear your working directory of any variables from your workspace:

```
rm(list=ls())
```

Now we want to set the working directory. This is where your R script and output will be saved. It's also a logical place to put data files that you plan to import into R. The following command will return your current working directory:

```
getwd()
```

Use the following command to change your directory (but note that you will need to modify to reflect *your* actual directory):

```
setwd("~/GitHub/QuantitativeBiodiversity/QB-2017/Week1")
```

2) WORKING WITH MATRICES

Matrices are another data type in R. They are just two-dimensional vectors containing data of the same type (e.g., numeric, integer, character). Therefore, much of what we discussed in Week 0 about vectors translates directly into dealing with matrices.

A. Making A Matrix

There are three common ways to create a matrix in R.

Approach 1 is to combine (or **concatenate**) two or more vectors. Let's start by creating a one-dimensional vector using a new function **rnorm**.

```
j <- c(rnorm(100, mean = 50, sd = 25))
```

Create a new vector named **z** also with 100 elements, but with a different mean and standard deviation

```
z <- c(rnorm(100, mean = 500, sd = 150))
```

Now we will use the function `cbind` to create a matrix from the two one-dimensional vectors:

```
k <- cbind(z, j)
```

Use the `help` function to learn about `cbind`. Use the `dim` function to describe the matrix you just created.

```
dim(k)
```

Approach 2 to making a matrix is to use the `matrix` function along with arguments that specify the number of rows (`nrow`) and columns (`ncol`):

```
l <- matrix(c(2, 4, 3, 1, 5, 7), nrow = 3, ncol = 2, byrow = FALSE)
```

Approach 3 to making a matrix is to import or **load a dataset** from your working directory:

```
m <- as.matrix(read.table("data/matrix.txt", sep = "\t", header = FALSE))
```

In this case, we're reading in a tab-delimited file. The name of your file must be in quotes, and you need to specify that it is a tab-delimited file type using the `sep` argument. The `header` argument tells R whether or not the names of the variables are contained in the first line; in the current example, they are not.

Often, when handling datasets, we want to be able to **transpose** a matrix. This is an easy operation in R that uses the `t` function:

```
n <- t(m)
```

B. Indexing a Matrix

Frequently, you will need to **index** or retrieve a certain portion of a matrix. As with the vector example above, we will use the square brackets to return data from a matrix. Inside the square brackets, there are now two subscripts corresponding to the rows and columns, respectively, of the matrix.

The following code will create a new matrix (`n`) based on the first three rows of matrix (`m`):

```
n <- m[1:3, ]
```

Or maybe you want the first two columns of a matrix instead:

```
n <- m[, 1:2]
```

Or perhaps you want non-sequential columns of a matrix. How do we do that? It's easy when you understand how to reference data within a matrix:

```
n <- m[, c(1:2, 5)]
```

3) BASIC DATA VISUALIZATION AND STATISTICAL ANALYSIS

A. Load Aquatic Mesocosm Dataset

In the following exercise, we will use a dataset from Lennon et al. (2003) (<http://goo.gl/JplHwd>), which looked at zooplankton communities along an experimental nutrient gradient in aquatic mesocosms. Inorganic nitrogen and phosphorus were added to mesocosms for six weeks at three different levels (low, medium, and high), but we also directly measured nutrient concentrations of water samples. So we have **categorical** and **continuous** predictors that we're going to use to help explain variation in zooplankton biomass.

The first thing we're going to do is load the nutrient data:

```
meso <- read.table("data/zoop_nuts.txt", sep = "\t", header = TRUE)
```

Let's use the `str` function to look at the structure of the data.

```
str(meso)
```

```
## 'data.frame':   24 obs. of  8 variables:
## $ TANK: int   34 14 23 16 21 5 25 27 30 28 ...
## $ NUTS: Factor w/ 3 levels "H","L","M": 2 2 2 2 2 2 2 3 3 ...
## $ TP  : num   20.3 25.6 14.2 39.1 20.1 ...
## $ TN  : num   720 750 610 761 570 ...
## $ SRP : num    4.02 1.56 4.97 2.89 5.11 4.68 5 0.1 7.9 3.92 ...
## $ TIN : num   131.6 141.1 107.7 71.3 80.4 ...
## $ CHLA: num    1.52 4 0.61 0.53 1.44 1.19 0.37 0.72 6.93 0.94 ...
## $ ZP  : num    1.781 0.409 1.201 3.36 0.733 ...
```

How does this dataset differ from the `m` dataset above? Remember, **matrices** and **vectors** are only comprised of a *single* data type. In contrast, the `meso` dataset has a combination of numeric and character data; note the `Factor`, `int`, and `num` data types generated from `str` function. So, this means we're dealing with a new type of **data structure**: a **data frame**.

Here is a description of the column headers:

- TANK = unique mesocosm identifier
- NUTS = categorical nutrient treatment: "L" = low, "M" = medium, "H" = high
- TP = total phosphorus concentration ($\mu\text{g/L}$)
- TN = total nitrogen concentration ($\mu\text{g/L}$)
- SRP = soluble reactive phosphorus concentration ($\mu\text{g/L}$)
- TIN = total inorganic nutrient concentration ($\mu\text{g/L}$)
- CHLA = chlorophyll *a* concentration (proxy for algal biomass; $\mu\text{g/L}$)
- ZP = zooplankton biomass (mg/L)

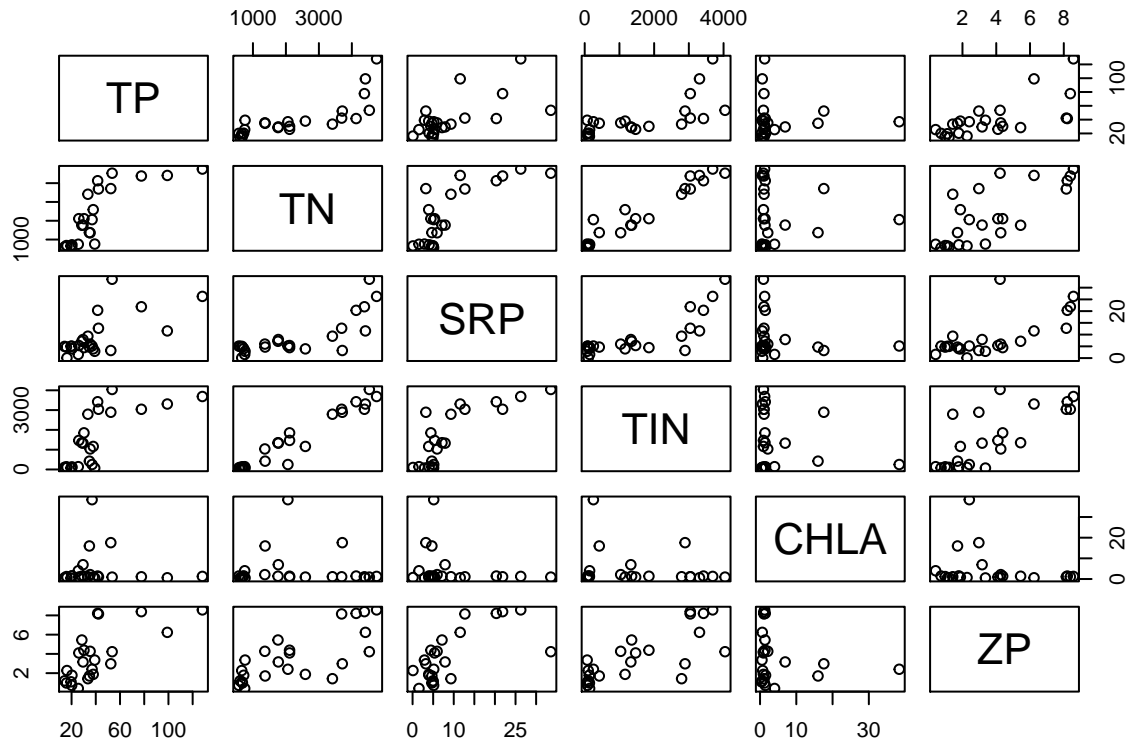
B. Correlation

A common step in data exploration is to look at correlations among variables. Before we do this, let's **index** our numerical (continuous) data in the 'meso' data frame. (Correlations typically don't work well on categorical data.)

```
meso.num <- meso[,3:8]
```

We can conveniently visualize pairwise **bi-plots** of the data using the following command:

```
pairs(meso.num)
```



Now let's conduct a simple **Pearson's correlation** analysis with the `cor()` function.

```
cor1 <- cor(meso.num)
```

C. Loading Contributed Packages

The base package in R won't always meet all of our needs. This is why there are > 6,000 **contributed packages** that have been developed for R. This may seem overwhelming, but it also means that there are tools (and web support) for just about any problem you might encounter.

When using one of the contributed packages, the first thing we need to do is **install** them along with their dependencies (i.e., other required packages). We're going to start out by using the *psych* package <http://cran.r-project.org/web/packages/psych/vignettes/overview.pdf>. The *psych* package has many features, but we're going to use it specifically for the `corr.test` function. This function generates **p-values** for each pairwise correlation. (For whatever reason, the `cor` function in the **base** package of R does not generate p-values.)

You can load an R package and its dependencies using the `require()` function. If a package is not found, i.e., not installed, you will need to install it first and then load dependencies.

```
#install.packages("psych", repos="http://cran.rstudio.com/")
require("psych")
```

Note that if a package is already installed, you can bypass the install line by placing a '#' at the beginning of it. Now, let's look at the correlations among variables and assess whether they are significant:

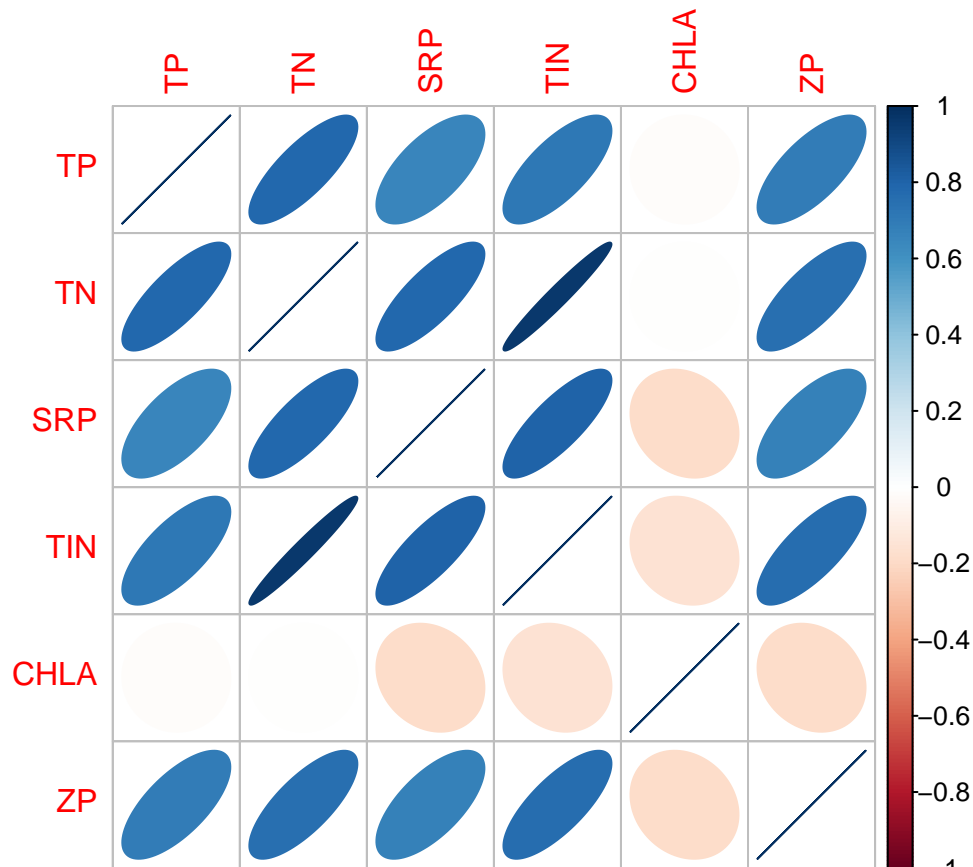
```
cor2 <- corr.test(meso.num, method = "pearson", adjust = "BH")
print(cor2, digits = 3)
```

Notes on *corr.test*:

- for rank-based correlations (i.e., non-parametric), use `method = "kendall"` or `"spearman"`. Give it a try!
- the `adjust = "BH"` statement supplies the Benjamini & Hochberg-corrected p-values in the upper right diagonal of the square matrix; the uncorrected p-values are below the diagonal. This process corrects for **false discovery rate**, which arises when making multiple comparisons.

Now, let's load another package that will let us visualize the direction and strength of the correlations:

```
install.packages("corrplot", repos="http://cran.rstudio.com/")
require("corrplot")
corrplot(cor1, method = "ellipse")
```



```
dev.off()
```

D. Linear Regression

It seems that total nitrogen (TN) is a fairly good predictor of zooplankton biomass (ZP) and this is something that we directly manipulated. This gives us license to conduct a linear regression analysis. We can do this in R using the `lm` function:

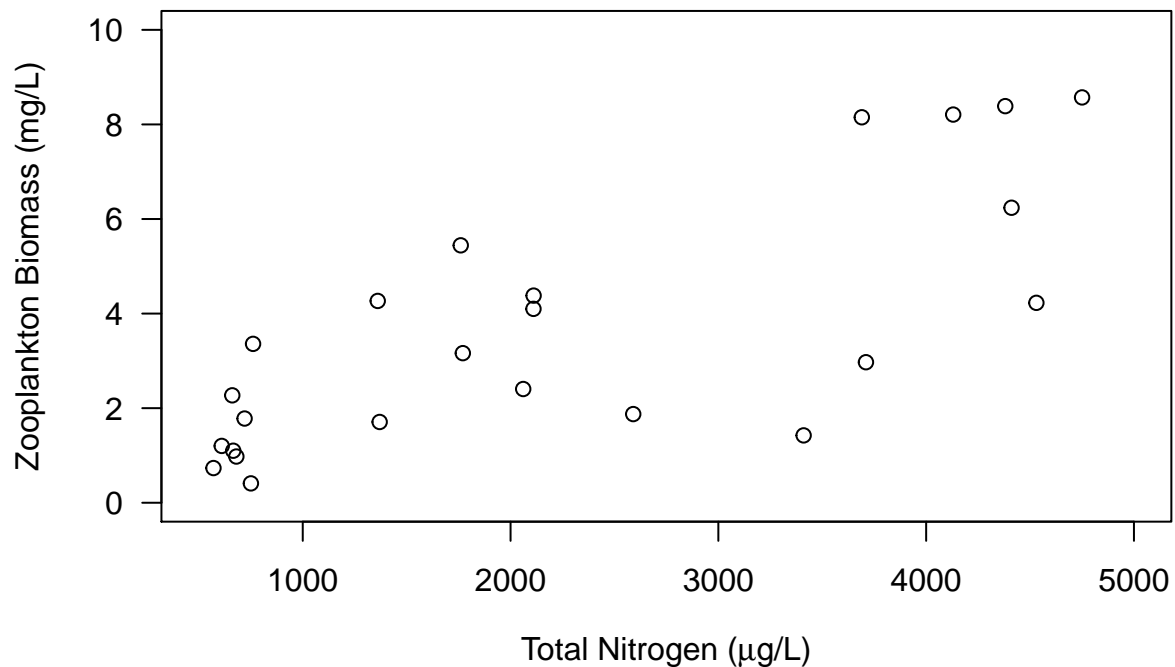
```
fitreg <- lm(ZP ~ TN, data = meso)
```

Let's examine the output of the regression model:

```
summary(fitreg)
```

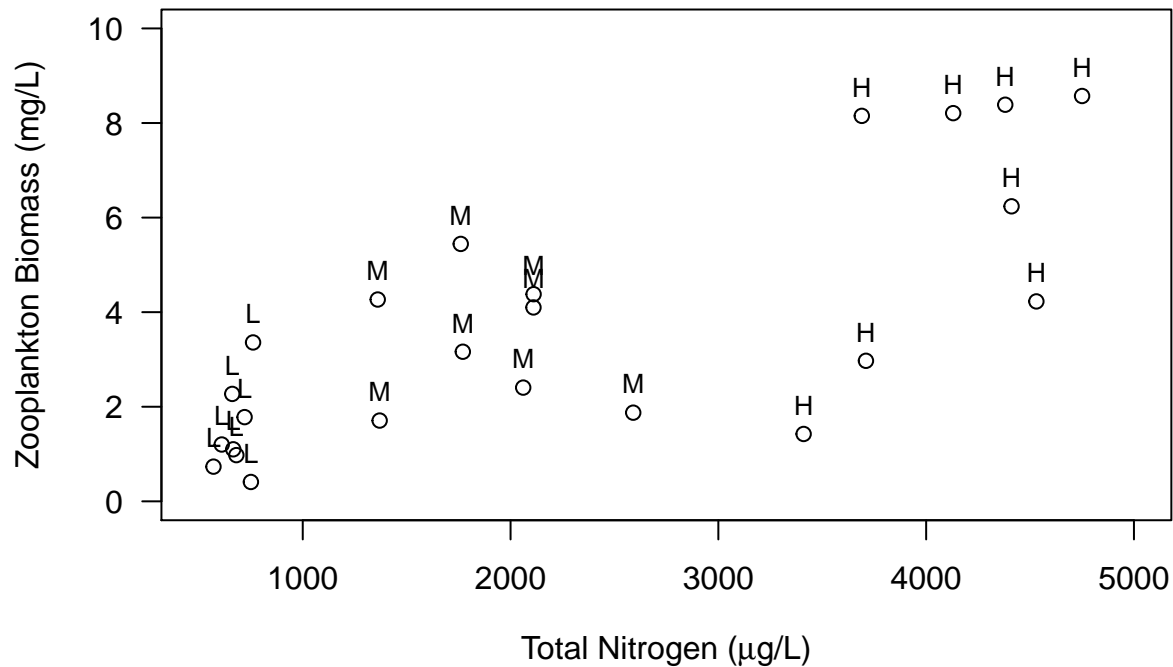
Now, let's look at a plot of the data used in the regression analysis:

```
plot(meso$TN, meso$ZP, ylim = c(0, 10), xlim = c(500, 5000),  
     xlab = expression(paste("Total Nitrogen (", mu, "g/L)")),  
     ylab = "Zooplankton Biomass (mg/L)", las = 1)
```



We can add some text to the plot to visualize the categorical nutrient treatments:

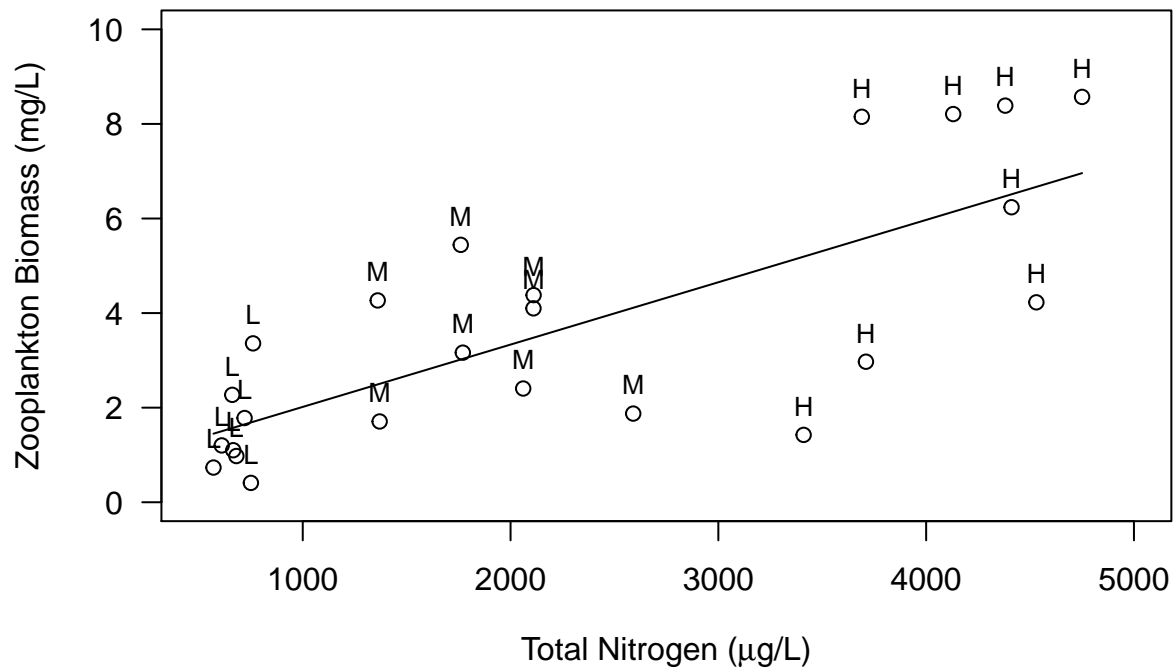
```
text(meso$TN, meso$ZP, meso$NUTS, pos = 3, cex = 0.8)
```



There is somewhat of a steep learning curve for plotting in R. However, R is also very flexible and can make beautiful figures. Perhaps one of the biggest initial obstacles is getting familiar with all of the **plotting arguments**. The following website provides a nice overview <http://www.statmethods.net/advgraphs/parameters.html> and there are plenty of other websites and books that are useful to have close by.

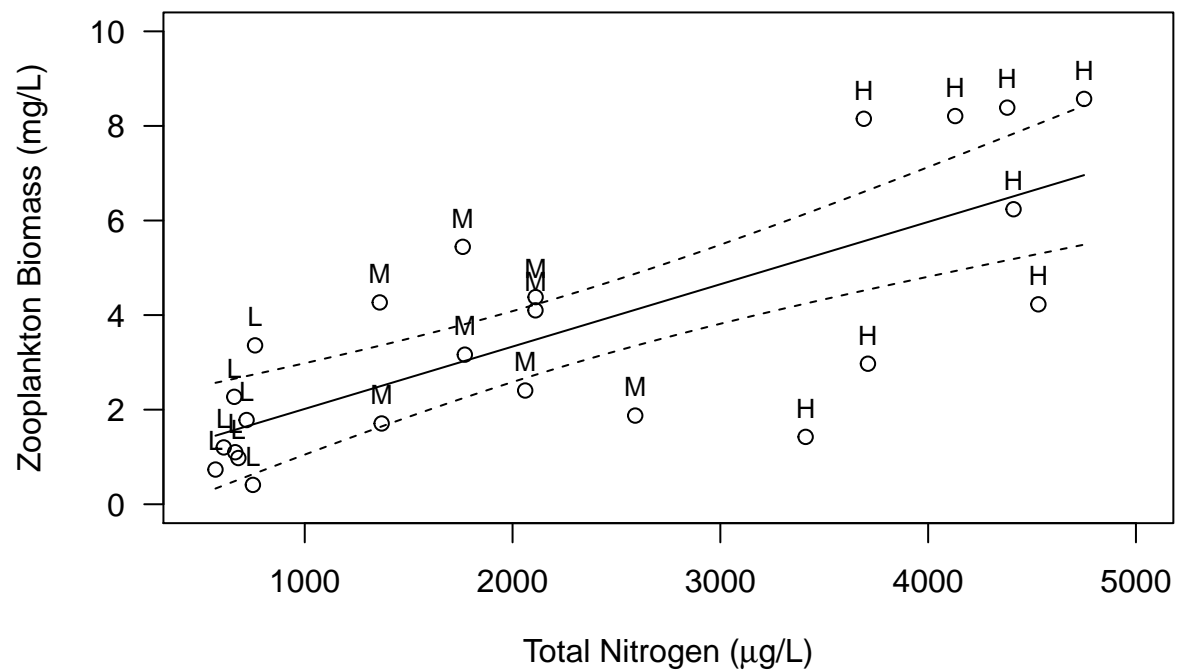
Now, let's continue with our regression analysis. To add the regression line, the first thing we need to do is identify a range of x-values and then generate the corresponding **predicted values** from our regression model:

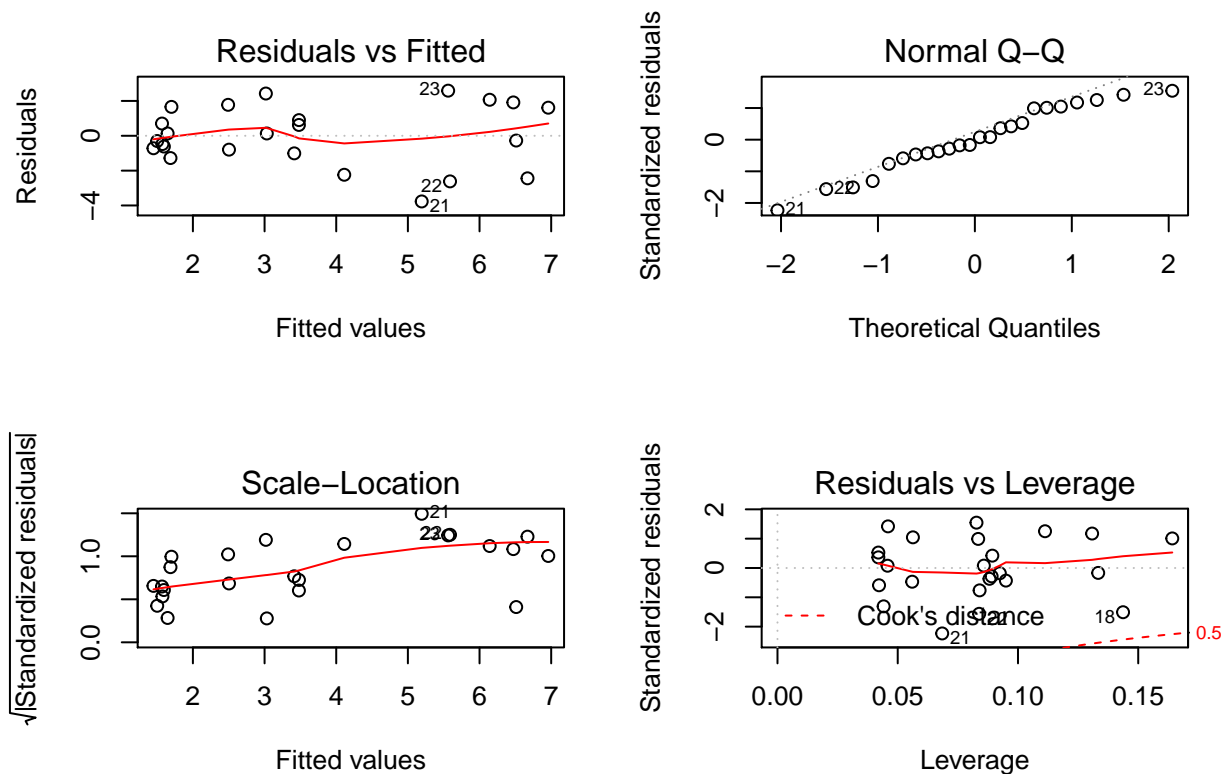
```
text(meso$TN, meso$ZP, meso$NUTS, pos = 3, cex = 0.8)
newTN <- seq(min(meso$TN), max(meso$TN), 10)
regline <- predict(fitreg, newdata = data.frame(TN = newTN))
lines(newTN, regline)
```



Now let's create and plot the **95% confidence intervals** using the same procedure as above, that is, use `newTN` to generate corresponding confidence intervals from our regression model:

```
text(meso$TN, meso$ZP, meso$NUTS, pos = 3, cex = 0.8)
newTN <- seq(min(meso$TN), max(meso$TN), 10)
regline <- predict(fitreg, newdata = data.frame(TN = newTN))
lines(newTN, regline)
# the line above calls the previous figure object
conf95 <- predict(fitreg, newdata = data.frame(TN = newTN),
                  interval = c("confidence"), level = 0.95, type = "response")
matlines(newTN, conf95[, c("lwr", "upr")], type="l", lty = 2, lwd = 1, col = "black")
```



- Upper left: is there a random distribution of the residuals around zero (horizontal line)?
- Upper right: is there a reasonably linear relationship between standardized residuals and theoretical quantiles? Try `help(qqplot)`
- Bottom left: again, looking for a random distribution of $\sqrt{\text{standardized residuals}}$
- Bottom right: leverage indicates the influence of points; contours correspond with Cook's distance, where values $> |1|$ are "suspicious"

E. Analysis of Variance (ANOVA)

We also have the option of looking at the relationship between zooplankton and nutrients where the manipulation is treated categorically (low, medium, high). First, let's order the categorical nutrient treatments from low to high (R's default is to order alphabetically):

```
NUTS <- factor(meso$NUTS, levels = c('L', 'M', 'H'))
```

Before plotting, we need to calculate the means and standard errors for zooplankton biomass in our nutrient treatments. We are going to use an important function called `tapply`. This allows us to apply a function (e.g., `mean`) to a vector (e.g., `ZP`) based on information in another column (e.g., nutrient treatment).

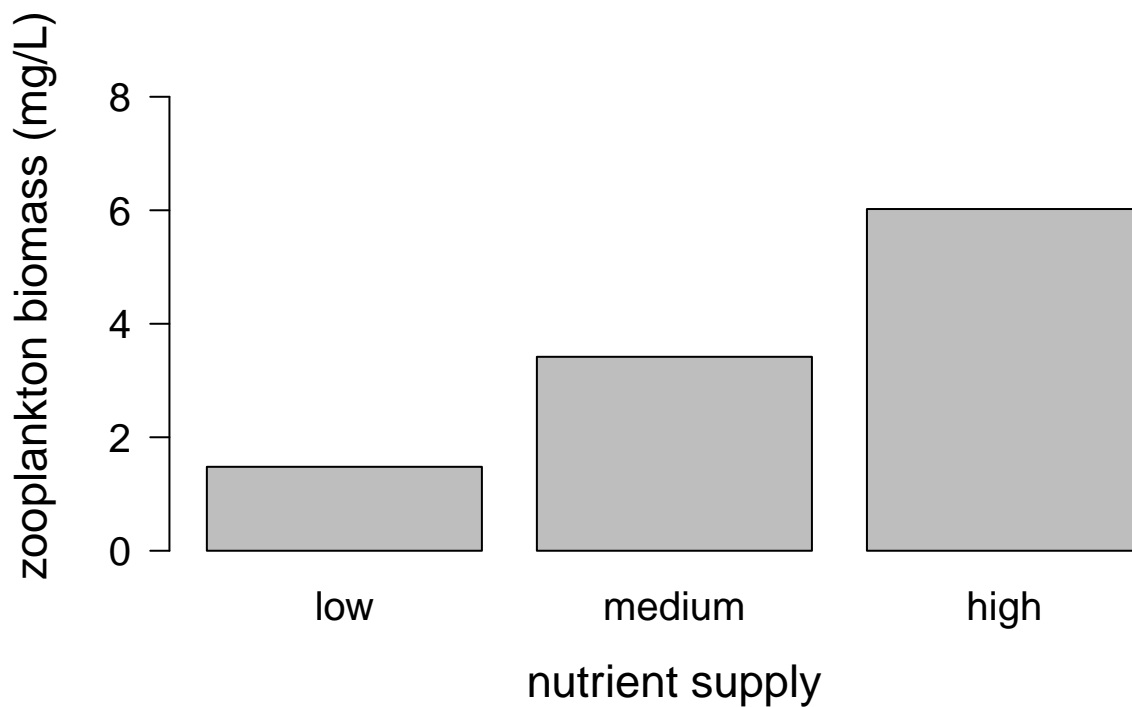
```
zp.means <- tapply(meso$ZP, NUTS, mean)

sem <- function(x){
  sd(na.omit(x))/sqrt(length(na.omit(x)))
}
```

```
zp.sem <- tapply(meso$ZP, NUTS, sem)
```

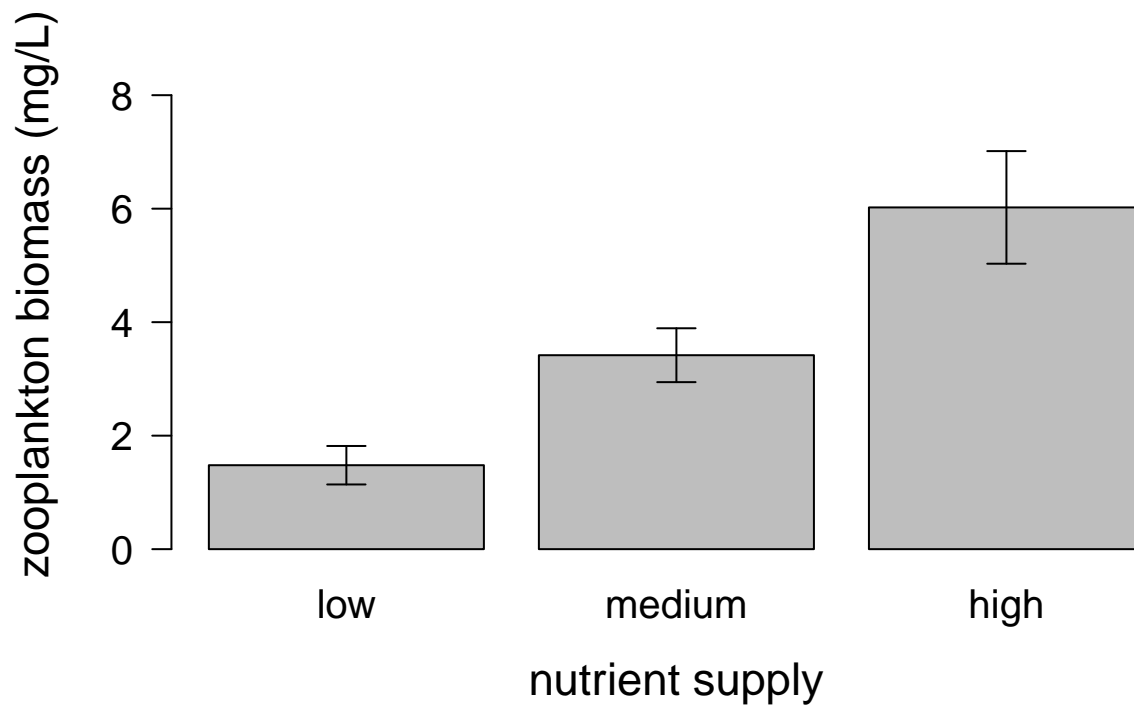
Now let's make the barplot:

```
bp <- barplot(zp.means, ylim = c(0, round(max(meso$ZP), digits = 0)),  
             pch = 15, cex = 1.25, las = 1, cex.lab = 1.4, cex.axis = 1.25,  
             xlab = "nutrient supply",  
             ylab = "zooplankton biomass (mg/L)",  
             names.arg = c("low", "medium", "high"))
```



We need to add the error bars (\pm sem) “manually” as follows:

```
arrows(x0 = bp, y0 = zp.means, y1 = zp.means - zp.sem, angle = 90,  
       length = 0.1, lwd = 1)  
arrows(x0 = bp, y0 = zp.means, y1 = zp.means + zp.sem, angle = 90,  
       length = 0.1, lwd = 1)
```



Last, we can conduct a one-way analysis of variance (ANOVA) as follows:

```
fitanova <- aov(ZP ~ NUTS, data = meso)
```

Let's look at the output in more detail (just as we did with regression):

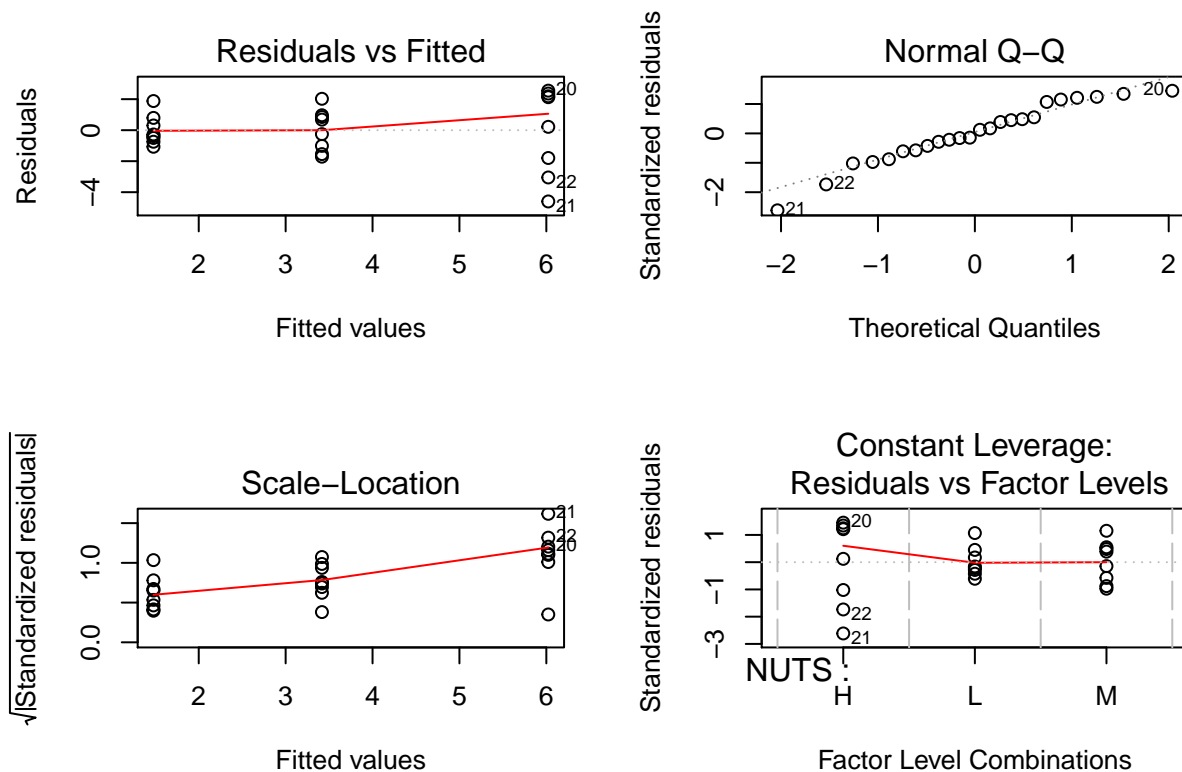
```
summary(fitanova)
```

Finally, we can conduct a post-hoc comparison of treatments using Tukey's HSD (Honest Significant Differences). This will tell us whether or not there are differences ($\alpha = 0.05$) among pairs of the three nutrient treatments

```
TukeyHSD(fitanova)
```

Just like the regression analysis above, it's good to look at the residuals:

```
par(mfrow = c(2, 2), mar = c(5.1, 4.1, 4.1, 2.1))  
plot(fitanova)
```



4) AN INTRODUCTION TO THE SITE-BY-SPECIES MATRIX

During Quantitative Biodiversity this semester, you will be introduced to a few fundamental data structures that are critical for analyzing biodiversity data. The first of these is the **site-by-species matrix**. This site-by-species matrix contains abundances, relative abundances, or the presence of species (or other taxonomic units) observed at different sampling sites. The site-by-species matrix is also very similar to other types of data that you might encounter if you are working with genomic, metagenomic, transcriptomic, or metabolomic datasets. This means that, even if you don't care about biodiversity *per se*, the tools that you are learning in Quantitative Biodiversity can be used to address questions in other systems! The table below is an example of what a typical site-by-species matrix looks like:

	Species1	Species2	Species3	Species4	...
Site1	90	76	1	0	
Site3	86	47	0	123	
Site3	23	89	0	46	
...					

Each row in the site-by-species matrix corresponds to a site (or sample), while each column corresponds to a species (or other taxonomic unit). Throughout the course, we will draw inferences about aspects of diversity from the site-by-species matrix.