Final project

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Contents

Fina	al Assignment
I	Loading data and tidying it up
	Oata Exploration
I	$\operatorname{ndices} \ldots \ldots$
A	Acknowledgement

Final Assignment

This is the second assingment for the Ecological Data Analysis in R course. In order to start, packages and functions will be loaded first. We'll use *tidiverse*, *dplyr*, *vegan*, *viridis* and a customized function that will be available in the repository.

Loading data and tidying it up.

Here, steps for tidying the dataset will be presented. Raw data was taken during completion of underwater transects, then input in Excel. Data will be imported as a .csv file.

```
\#df \leftarrow read.csv("./data/df\_daniel.csv")
df <- read.csv("D:/UABC/II Semestre/Datos R/eda finalproject/data/df daniel.csv") #this one's for knitt
# We'll only need provinces, localities, sites, transects, species and abundances
df <- df %>% select(province, locality, siteref, transect, sp, length, abundance)
head(df, n=3L)
       province
                    locality siteref transect
                                                                      sp length
## 1 Puntarenas ISLA_TORTUGA
                                                  Gnathanodon speciosus
                                                                             20
                                    Α
                                            Α1
## 2 Puntarenas ISLA_TORTUGA
                                    Α
                                            A1
                                                   Haemulon maculicauda
                                                                             15
## 3 Puntarenas ISLA_TORTUGA
                                    Α
                                             A1 Microspathodon dorsalis
                                                                             25
     abundance
## 1
             5
## 2
             4
## 3
tail(df, n=3L)
                      locality siteref transect
          province
                                                                      sp length
## 1502 Guanacaste CUAJINIQUIL
                                      G
                                              G35
                                                          Sabellidae sp
                                                                              5
                                      G
## 1503 Guanacaste CUAJINIQUIL
                                             G35 Stenorhynchus debilis
                                                                             10
## 1504 Guanacaste CUAJINIQUIL
                                      G
                                                    Toxopneustes roseus
                                             G35
                                                                              5
##
        abundance
```

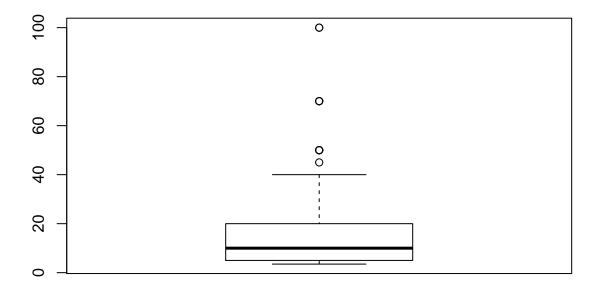
```
## 1502
## 1503
                2
## 1504
str(df)
                    1504 obs. of 7 variables:
## 'data.frame':
   \ province : Factor w/ 2 levels "Guanacaste", "Puntarenas": 2 2 2 2 2 2 2 2 2 ...
## $ locality : Factor w/ 3 levels "CUAJINIQUIL",...: 2 2 2 2 2 2 2 2 2 2 ...
## \$ siteref : Factor w/ 7 levels "A", "B", "C", "D", ...: 1 1 1 1 1 1 1 1 1 1 ...
## $ transect : Factor w/ 35 levels "A1", "A2", "A3",..: 1 1 1 1 1 1 1 1 1 1 ...
               : Factor w/ 117 levels "Abudefduf concolor",..: 46 50 67 67 2 1 107 107 23 39 ...
##
   $ length
               : num 20 15 25 30 10 5 10 15 15 15 ...
  $ abundance: int 5 4 2 3 3 5 6 8 1 1 ...
```

Before continuing, let's remember our question: will there be a difference between communities of two provinces with different fishing pressures?

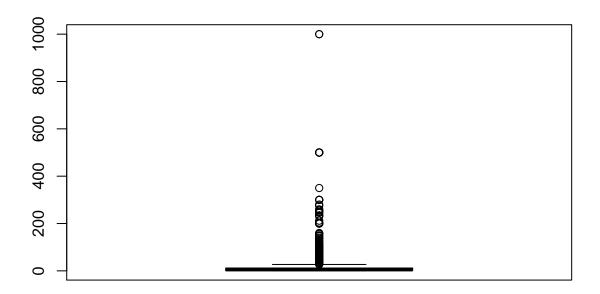
Data Exploration

1. Outliers

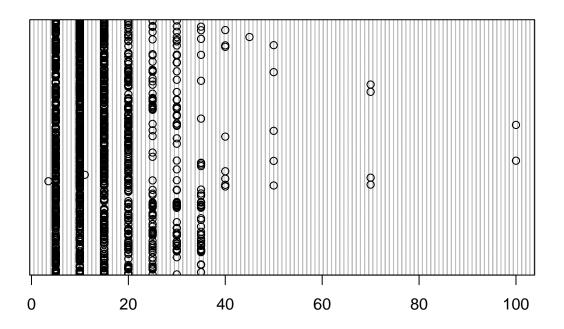
```
boxplot(df$length)
```



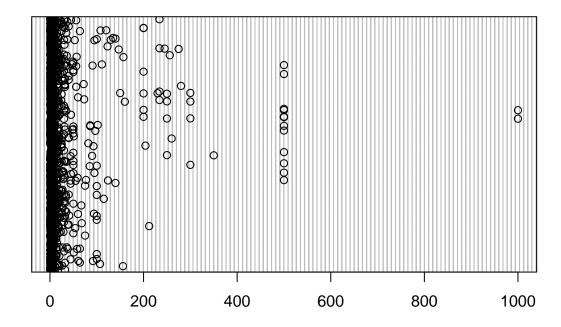
```
boxplot(df$abundance)
```



dotchart(df\$length)



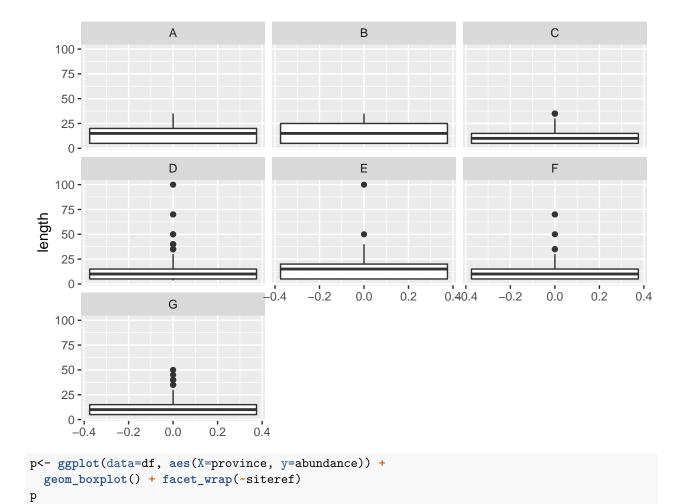
dotchart(df\$abundance)

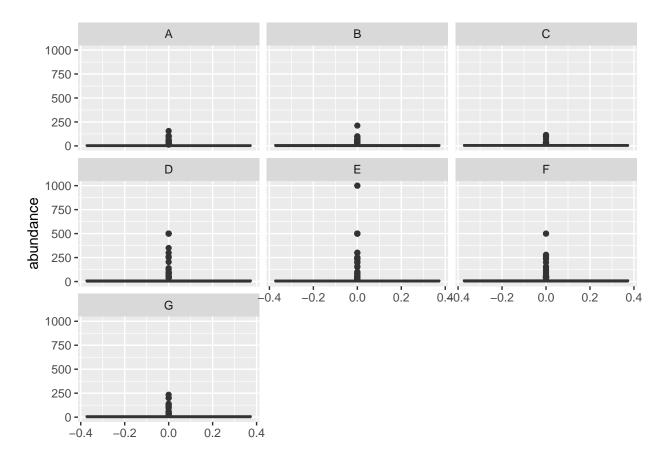


Not that uncommon to see abundances represented like that, I think. Lot's of zeroes and some outliers for abundance and no errors regarding length.

2. Homogeneity of variance

```
p <- ggplot(data=df, aes(X=province, y=length)) +
  geom_boxplot() + facet_wrap(~siteref)
p</pre>
```





3. Normality

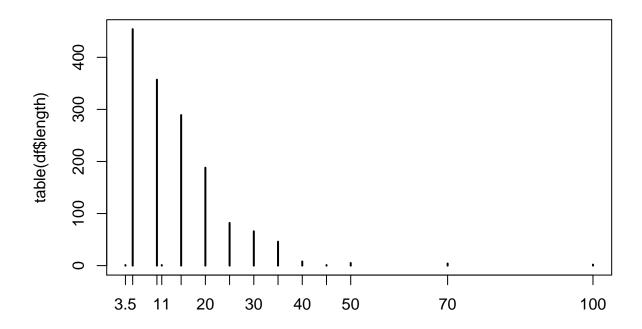
Data is not normal, as seen in histograms and confirmed by Q-Q plots.

4. Zeroes

range(df\$length)

[1] 3.5 100.0

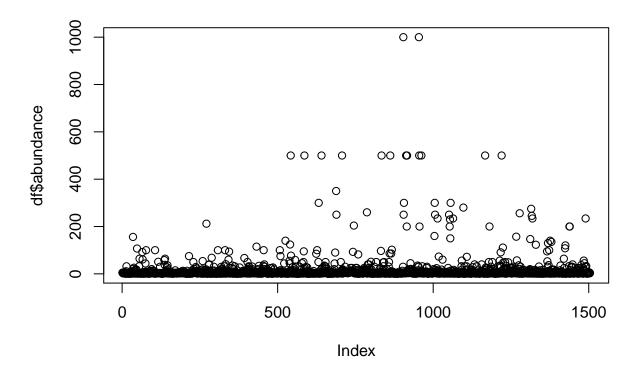
plot(table(df\$length))



range(df\$abundance)

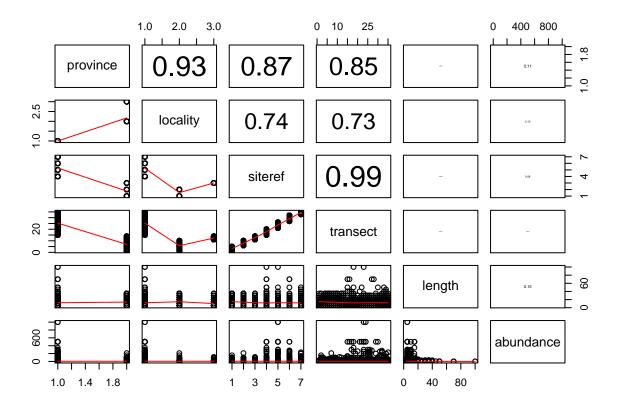
[1] 1 1000

plot(df\$abundance)

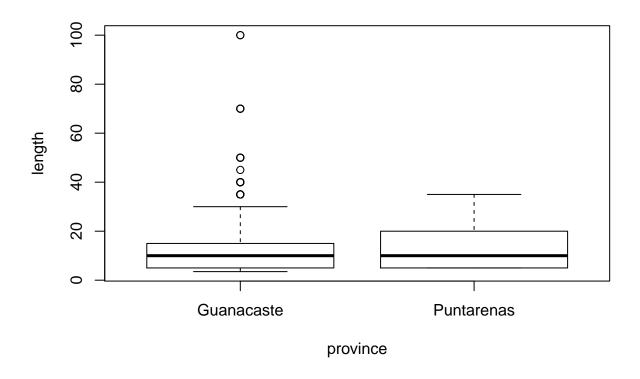


There are lots of zeroes for abundance, which we'll have to take into account when analyzing data.

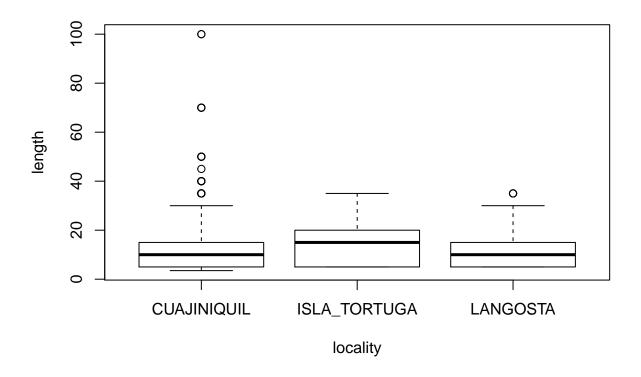
5. Collinearity X



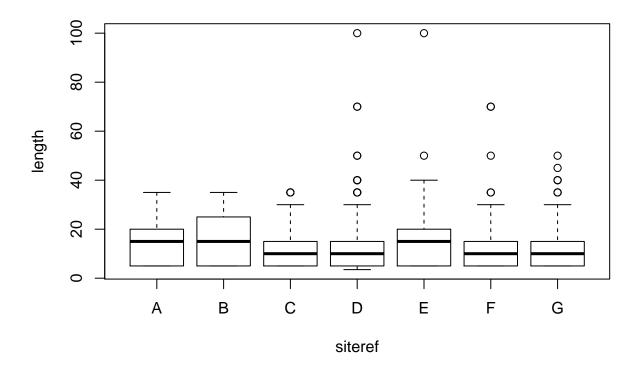
plot(length ~ province, data=df)



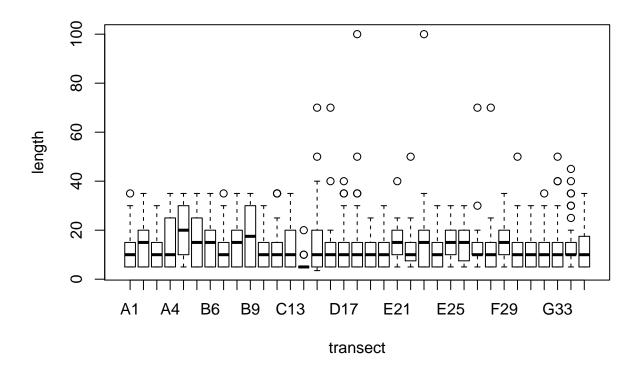
plot(length ~ locality, data=df)



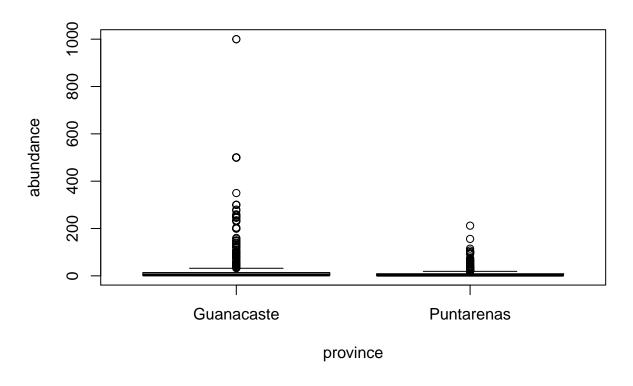
plot(length ~ siteref, data=df)



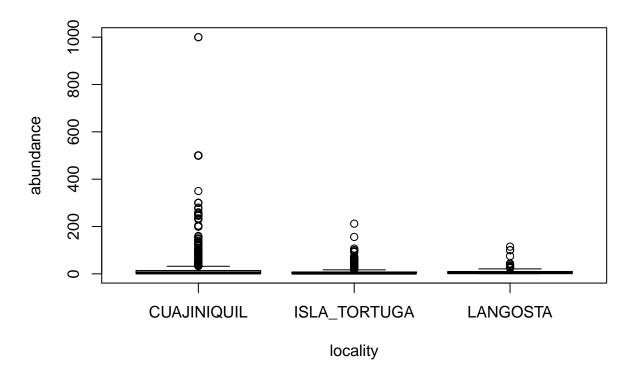
plot(length ~ transect, data=df)



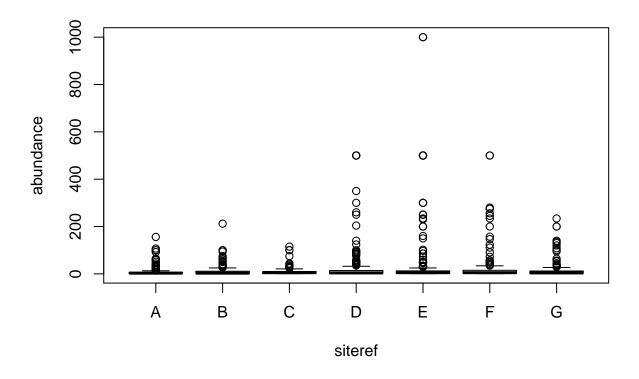
plot(abundance ~ province, data=df)



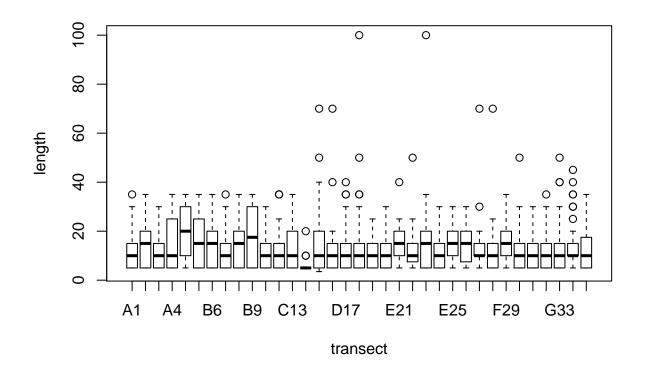
plot(abundance ~ locality, data=df)

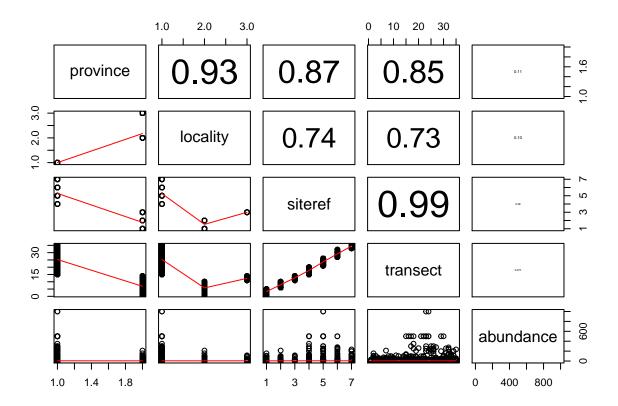


plot(abundance ~ siteref, data=df)



plot(length ~ transect, data=df)

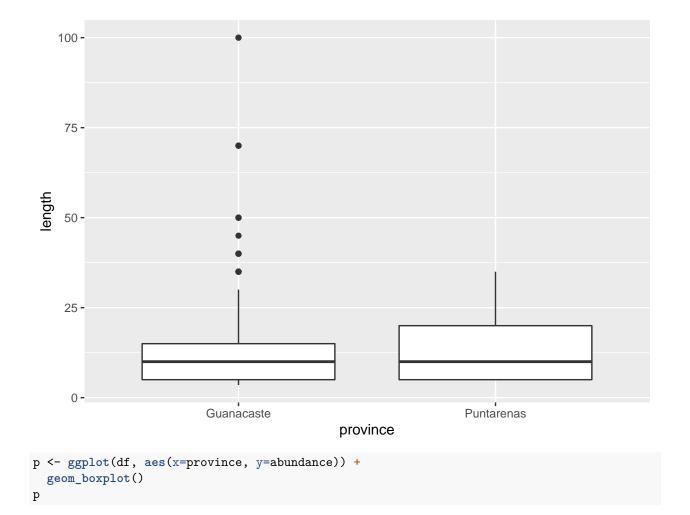


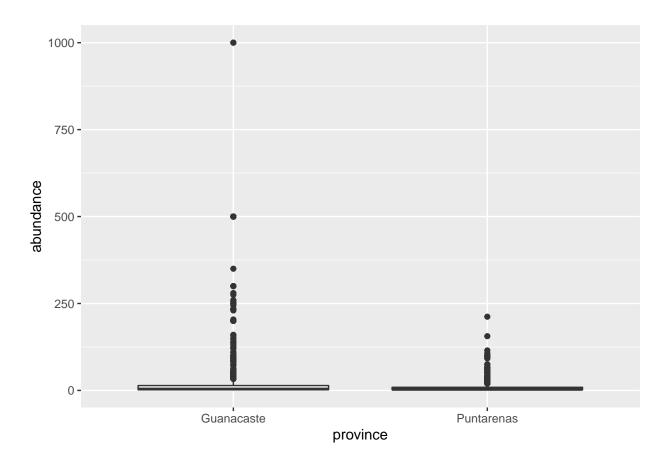


We'll take into account province, locality, site and transects as x variables and abundance as y variables; our question does not requires us to study length, so it's something we could have dropped before. We'll still explore it for the sake of it.

6. Relationships X and Y

```
p <- ggplot(df, aes(x=province, y=length)) +
   geom_boxplot()
p</pre>
```

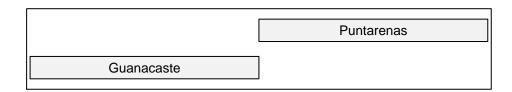


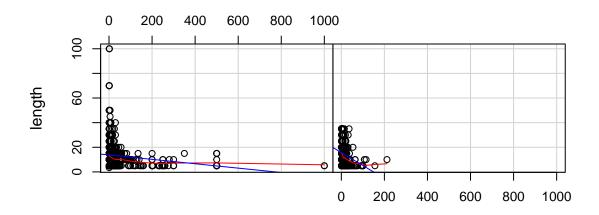


7. Interactions

```
coplot(length ~ abundance | province ,
    data=df,
    panel=function(x,y,...) {
        panel.smooth(x,y,span=0.8,iter=5,...)
        abline(lm(y ~ x), col="blue") } )
```

Given: province

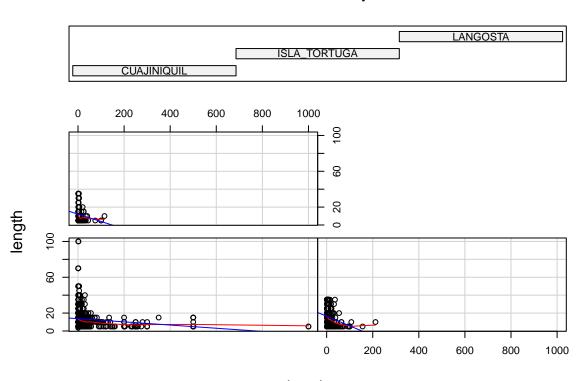




abundance

```
coplot(length ~ abundance | locality ,
    data=df,
    panel=function(x,y,...) {
        panel.smooth(x,y,span=0.8,iter=5,...)
        abline(lm(y ~ x), col="blue") } )
```

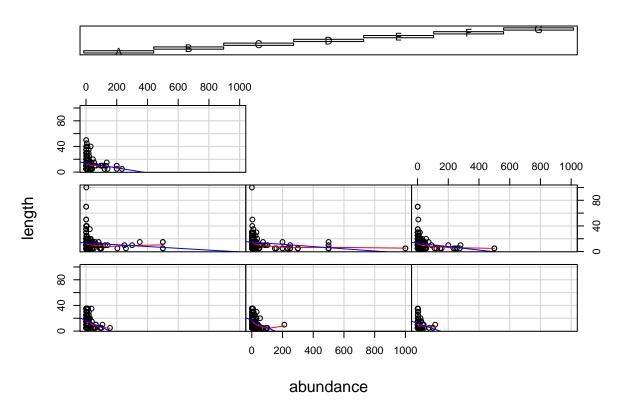
Given: locality



abundance

```
coplot(length ~ abundance | siteref ,
    data=df,
    panel=function(x,y,...) {
       panel.smooth(x,y,span=0.8,iter=5,...)
       abline(lm(y ~ x), col="blue") } )
```

Given: siteref



There seems to be no interaction.

8. Independence of Y

Both length and density could be dependent on province, locality, site because of fishing pressure effects. In transects, I'd say there could be independence.

Indices

Data setup

We'll need to create an abundance matrix in order to calculate indices. Let's create a dataframe for our indices.

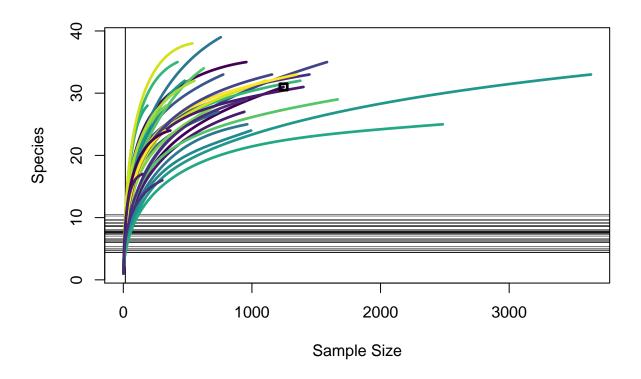
```
#Let's create a dataframe for our indices.
#We only need abundance

# first, we'll summarise species disregarding length

df_sp <- df %>% select(province, locality, siteref, transect, sp, abundance)
head(df_sp)
```

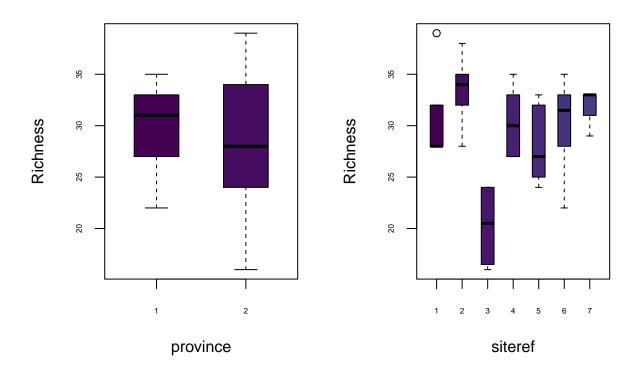
province locality siteref transect sp abundance

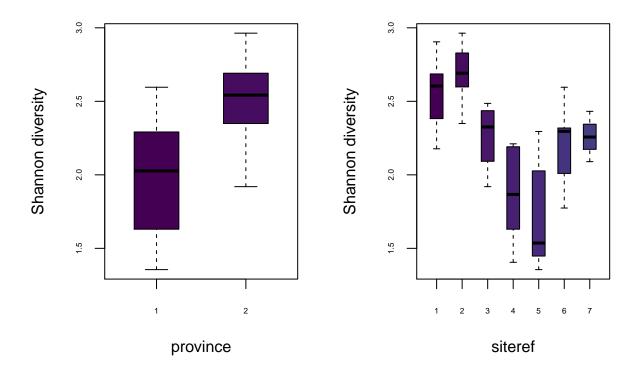
```
## 1 Puntarenas ISLA_TORTUGA
                                   Α
                                           A1
                                                 Gnathanodon speciosus
## 2 Puntarenas ISLA_TORTUGA
                                                  Haemulon maculicauda
                                                                               4
                                   Α
                                            A1
## 3 Puntarenas ISLA_TORTUGA
                                   Α
                                            A1 Microspathodon dorsalis
                                                                               2
## 4 Puntarenas ISLA_TORTUGA
                                                                               3
                                   Α
                                            A1 Microspathodon dorsalis
## 5 Puntarenas ISLA_TORTUGA
                                   Α
                                                 Abudefduf troschelii
                                                                                3
## 6 Puntarenas ISLA_TORTUGA
                                   Α
                                            A1
                                                    Abudefduf concolor
str(df_sp)
## 'data.frame': 1504 obs. of 6 variables:
## $ province : Factor w/ 2 levels "Guanacaste", "Puntarenas": 2 2 2 2 2 2 2 2 2 ...
## $ locality : Factor w/ 3 levels "CUAJINIQUIL",..: 2 2 2 2 2 2 2 2 2 ...
## $ siteref : Factor w/ 7 levels "A", "B", "C", "D", ...: 1 1 1 1 1 1 1 1 1 1 ...
## $ transect : Factor w/ 35 levels "A1", "A2", "A3",..: 1 1 1 1 1 1 1 1 1 1 ...
            : Factor w/ 117 levels "Abudefduf concolor",..: 46 50 67 67 2 1 107 107 23 39 ...
## $ abundance: int 5 4 2 3 3 5 6 8 1 1 ...
df_sp <- summarise(group_by(df_sp, province, locality, siteref, transect, sp),</pre>
                        sum abundance = sum(abundance, na.rm = TRUE))
# ab_sp contains abundance matrix for ecological indices
# we'll create it out of df; as we don't care for sizes, we'll sum all abundances for each species on f
ab_sp <- pivot_wider(df_sp, names_from = sp, values_from = sum_abundance)
#converting these factors to numeric will solve an issue that could arise later when we're creating our
ab_sp$province <- as.double(ab_sp$province)</pre>
ab_sp$locality <- as.double(ab_sp$locality)</pre>
ab_sp$siteref <-as.double(ab_sp$siteref)</pre>
ab_sp$transect <- as.double(ab_sp$transect)</pre>
# let's convert na to zeroes
ab_sp[is.na(ab_sp)] <-0
Now we can obtain ecological indices
#We'll create a dataframe to store our results
ind_sp <- ab_sp[, c("province", "locality", "siteref", "transect")]</pre>
# Richness: number of species per transect/habitat
ind_sp$richness <- rowSums(ab_sp>0)
# Shannon's diversity index: the bigger, the more diverse, basically
ind_sp$Shannon <- diversity(ab_sp)</pre>
# Rarefaction
raremax <- min(rowSums(ab_sp>0))
ind_sp$Rarefied <- c(rarefy(ab_sp[1:35,], sample=raremax))</pre>
raremax <- min(rowSums(ab_sp>0))
# Let's also visualize a rarefaction curve
#this function will create a rarefaction curve to observe species accumulation
rarecurve(ab_sp, sample = raremax, col=viridis(raremax, alpha = 1, begin = 0, end = 1, direction = 1, o
```

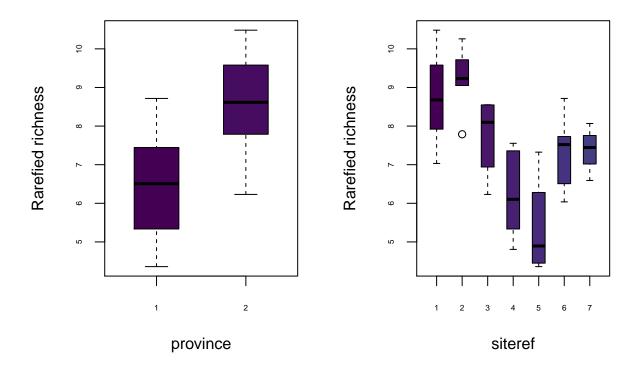


Visualization

We'll use boxplots to see differences in species diversity:







```
mfrow=c(1,1)
```

Looks like there are "some" differences, so let's explore them.

Linear modeling of indices

It appears that species diversity increases as we move from the field to the forest. We can test for differences among habitats statistically using a linear model, with Habitat as a predictor of species diversity:

```
# fit linear models (ANOVA)
mod.richness.province <- lm(richness~province, data=ind_sp)
mod.richness.siteref <- lm(richness~siteref, data=ind_sp)
mod.richness.transect <- lm(richness~transect, data=ind_sp)
mod.Shannon.province <- lm(Shannon~province, data=ind_sp)
mod.Shannon.siteref <- lm(Shannon~siteref, data=ind_sp)
mod.Shannon.transect <- lm(Shannon~transect, data=ind_sp)
mod.Rarefied.province <- lm(Rarefied~province, data=ind_sp)
mod.Rarefied.siteref <- lm(Rarefied~siteref, data=ind_sp)
mod.Rarefied.transect <- lm(Rarefied~siteref, data=ind_sp)
mod.Rarefied.transect <- lm(Rarefied~transect, data=ind_sp)
anova(mod.richness.province)</pre>
```

```
## Analysis of Variance Table
##
## Response: richness
## Df Sum Sq Mean Sq F value Pr(>F)
```

```
## province 1 8.8 8.8048 0.3208 0.5749
## Residuals 33 905.6 27.4423
anova(mod.richness.siteref)
## Analysis of Variance Table
## Response: richness
            Df Sum Sq Mean Sq F value Pr(>F)
          1 0.31 0.3054 0.011 0.917
## siteref
## Residuals 33 914.09 27.6998
anova(mod.richness.transect)
## Analysis of Variance Table
##
## Response: richness
## Df Sum Sq Mean Sq F value Pr(>F)
## transect 1 0.1 0.1011 0.0036 0.9522
## Residuals 33 914.3 27.7060
#Results show no significant differences
anova(mod.Shannon.province)
## Analysis of Variance Table
## Response: Shannon
            Df Sum Sq Mean Sq F value
## province 1 2.4871 2.48711 20.995 6.286e-05 ***
## Residuals 33 3.9092 0.11846
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
anova(mod.Shannon.siteref)
## Analysis of Variance Table
##
## Response: Shannon
           Df Sum Sq Mean Sq F value Pr(>F)
## siteref 1 1.2784 1.27842 8.2432 0.007091 **
## Residuals 33 5.1179 0.15509
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
anova(mod.Shannon.transect)
## Analysis of Variance Table
## Response: Shannon
           Df Sum Sq Mean Sq F value Pr(>F)
## transect 1 0.9268 0.92680 5.5918 0.02408 *
## Residuals 33 5.4695 0.16574
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
#There are significant differences for each one
anova(mod.Rarefied.province)
```

```
## Analysis of Variance Table
##
## Response: Rarefied
##
            Df Sum Sq Mean Sq F value
                                         Pr(>F)
## province
            1 39.387 39.387 24.783 1.967e-05 ***
## Residuals 33 52.446
                        1.589
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
anova(mod.Rarefied.siteref)
## Analysis of Variance Table
## Response: Rarefied
##
            Df Sum Sq Mean Sq F value
                                        Pr(>F)
             1 22.249 22.2491 10.552 0.002669 **
## Residuals 33 69.583 2.1086
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
anova(mod.Rarefied.transect)
## Analysis of Variance Table
## Response: Rarefied
            Df Sum Sq Mean Sq F value Pr(>F)
             1 17.048 17.0475 7.5225 0.00977 **
## Residuals 33 74.785 2.2662
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
#Same!
#Should we compare between groups? Let's try a Tukey test. Also, I'm really not sure what am I doing bu
#Tukey for Shannon
TukeyHSD(aov(lm(Shannon~as.factor(siteref), data = ind_sp)), conf.level = 0.95)
##
    Tukey multiple comparisons of means
##
      95% family-wise confidence level
##
## Fit: aov(formula = lm(Shannon ~ as.factor(siteref), data = ind_sp))
## $`as.factor(siteref)`
              diff
                           lwr
                                       upr
                                               p adj
## 2-1 0.135322169 -0.45005101 0.72069534 0.9891678
## 3-1 -0.286564766 -0.90744678 0.33431725 0.7627015
## 4-1 -0.689117231 -1.24956938 -0.12866508 0.0087555
## 5-1 -0.851372524 -1.41182467 -0.29092038 0.0008059
## 6-1 -0.335742646 -0.89619479 0.22470950 0.4966182
## 7-1 -0.291317360 -0.96724808 0.38461336 0.8140044
## 3-2 -0.421886934 -1.04276895 0.19899508 0.3501185
## 4-2 -0.824439400 -1.38489155 -0.26398725 0.0012068
## 5-2 -0.986694692 -1.54714684 -0.42624255 0.0001045
## 6-2 -0.471064815 -1.03151696  0.08938733  0.1446756
## 7-2 -0.426639528 -1.10257025 0.24929119 0.4354619
```

```
## 4-3 -0.402552465 -0.99999646 0.19489153 0.3597010
## 5-3 -0.564807758 -1.16225175 0.03263624 0.0731869
## 6-3 -0.049177881 -0.64662188
                                0.54826612 0.9999686
## 7-3 -0.004752594 -0.71165786
                                0.70215267 1.0000000
## 5-4 -0.162255292 -0.69662545
                                 0.37211486 0.9578414
                                0.88774474 0.3811865
## 6-4 0.353374585 -0.18099557
       0.397799871 -0.25666724
                                1.05226698 0.4796657
       0.515629877 -0.01874028
                                1.05000003 0.0639733
  7-5
       0.560055164 -0.09441194
                                1.21452227 0.1316368
## 7-6 0.044425287 -0.61004182 0.69889239 0.99999900
#Differences for site1-4, site1-5, site2-4, site2-5 and almost site5-6
#Tukey for Rarefied richness
TukeyHSD(aov(lm(Rarefied~as.factor(siteref), data = ind_sp)), conf.level = 0.95)
##
     Tukey multiple comparisons of means
##
       95% family-wise confidence level
##
## Fit: aov(formula = lm(Rarefied ~ as.factor(siteref), data = ind_sp))
##
## $`as.factor(siteref)`
##
              diff
                           lwr
                                      upr
                                              p adj
## 2-1 0.47024930 -1.70561859
                                2.6461172 0.9924004
## 3-1 -0.99440819 -3.30226461
                               1.3134482 0.8141752
  4-1 -2.52858373 -4.61181864 -0.4453488 0.0099268
## 5-1 -3.37119589 -5.45443079 -1.2879610 0.0003488
## 6-1 -1.40024431 -3.48347921 0.6829906 0.3624857
## 7-1 -1.37080826 -3.88328409
                               1.1416676 0.6023670
## 3-2 -1.46465750 -3.77251391
                               0.8431989 0.4290732
## 4-2 -2.99883303 -5.08206794 -0.9155981 0.0015708
## 5-2 -3.84144519 -5.92468010 -1.7582103 0.0000516
## 6-2 -1.87049361 -3.95372852 0.2127413 0.1005301
## 7-2 -1.84105756 -4.35353339
                                0.6714183 0.2679949
## 4-3 -1.53417554 -3.75491141 0.6865603 0.3314975
## 5-3 -2.37678769 -4.59752356 -0.1560518 0.0299716
## 6-3 -0.40583611 -2.62657199 1.8148998 0.9969322
## 7-3 -0.37640006 -3.00401018
                                2.2512101 0.9992154
## 5-4 -0.84261216 -2.82889870
                               1.1436744 0.8246237
       1.12833942 -0.85794712 3.1146260 0.5576441
       1.15777547 -1.27491879
                               3.5904697 0.7368895
       1.97095158 -0.01533497
                                3.9572381 0.0528165
## 7-5
       2.00038763 -0.43230664
                              4.4330819 0.1616133
## 7-6 0.02943605 -2.40325822 2.4621303 1.0000000
#Differeces for site 1-4, site1-5, site2-4, site2-5, site3-5 and almost site5-6
```

Interpretation

Results of ANOVA show no significant difference for richness, but does show it for Shannon and rarefied species richness. Seems like there's no association between province/site and species richness, but it is for Shannon's diversity index and rarefied richness.

Should we go on? Yes! Well... Seems like my data won't allow me to continue, if I'm not mistaken. I was considering doing an ordination analysis to determine which sites were similar and what species share distribution in provinces, sites and/or transects. I'd need to dampen data from abundances and divide it by

...? I don't know. So I'll leave it up to here.

Based on what I have, the answer to my question would be that there's evidence -based on my analysis- that differences in species composition could be explained by the location, site or transect. Based on Tukey test, there are some differences between provinces: sites 1-3 correspond to Puntarenas, the high fishing pressure province, and 4-7 to Guanacaste, the not-so-high fishing pressure province.

Acknowledgement

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