

Appendix 2.

Custom R scripts:

```
#### Loading packages and calling the data ####
```

```
library(vegan)
```

```
library(ggplot2)
```

```
setwd(choose.dir())
```

```
data<-read.table("sucessao_ABUND.txt", header=T) ### data containing abundances of  
the 10 most abundant ethnospecies in each plot
```

```
species<-data[, 3:49] ### removing the columns "Area" and "Environment"
```

```
data2<-read.table("sucessão_final_tudo3.txt", header=T) ### data containing  
abundances of all 230 ethnospecies
```

```
species2<-data2[, 3:232] ### removing the columns "Area" and "Environment"
```

```
### Gradient of relative abundance of ethnospecies ordered by the one-  
dimensional NMDS (Figure 3) ###
```

```
### 1 – Generic function that sorts data###
```

```
generic<-function(tabel,gradient,at,grad,axisY,axisX){
```

```
tabel<-as.matrix(tabel)
```

```
gradient<-as.matrix(gradient)
```

```
weighted.average <-colSums(tabel*gradient[,1])/colSums(tabel)
```

```
sub.orden<-tabel[order(gradient[,1],decreasing=F),] # sort plots according to the  
gradient
```

```
sub.orde<-sub.orden[,order(weighted.average,decreasing=T)] # put species sorted by  
the weighted average
```

```
data.pa<-matrix(0,nrow(tabel),ncol(tabel))
```

```
data.pa[tabel>0]<-1
```

```
ordered<-sub.orde[,which(colSums(data.pa)>0)] ## to delete possible empty columns  
(species that did not occur)
```

```
par(mfrow=c(ncol(ordenado)+1,1),mar=c(0,4,0.2,10),oma=c(3,1,1,6))
```

```
layout(matrix(1:(ncol(ordered)+1)),heights=c(3,rep(1,ncol(ordered))))
```

```
plot(sort(gradient[,1]),axes=F,ylab="",mfg=c(21,1),lwd=10,las=2,lend="butt",frame.plo  
t=F,xaxt="n",type="h",col="black",ylim=c(min(gradient),max(gradient)))
```

```

axis(side=2,at=c(0,max(gradient)),las=2)
mtext(grad,4,outer=T,font=2,line=-10,padj=-18.5,las=2)
for(i in 1:ncol(ordered)){
barplot(ordered[,i],bty="l",axisnames=F,axes=FALSE,col="black")
#axis(side=2,at=max(ordered[,i]),las=2)
mtext(colnames(ordered)[i],3,line=-1.0,adj=0,at=at,cex=.8,font=3)
}
mtext(axisX,1,outer=T,font=2,line=1.2)
mtext(axisY,2,font=2,outer=T,line=-2)
}

```

2 – NMDS

```

parc1<-decostand(species,"total",MARGIN=1) ### standardization method for ordering
rows (ethnospecies abundances)

```

```

resu<-metaMDS(parc1,distance = "bray", k=1,trymax=100,autotransform=F)

```

```

scor.nmds<-resu$points

```

```

summary(lm(vegdist(parc1)~vegdist(scor.nmds,"euclid")))

```

```

plot(scor.nmds, type="n", main="NMDS with Vegan") ### NMDS plot

```

```

text(scor.nmds)

```

```

generic(parc1,scor.nmds, 9.75, "") ### ordering plots by ethnospecies abundances

```

Dendrogram of the hierarchical clustering analysis to compare floristic composition (Figure 4)

```

cluster<-vegdist(species2, method="jaccard", binary=T)

```

```

phylogeny<-hclust(cluster, method="average")

```

```

plot(phylogeny, hang=-1)

```

```

matrix_cof<-cophenetic(phylogeny)

```

```

cor(matrix_cof, dist.jac) ### Cophenetic correlation coefficient

```

Comparison among plots in terms of ethnospecies richness (Figure 5)

1 – Rarefaction curves

```

rarecurve(species2, lty=1, xlab = "Number of stems", ylab="Expected richness",
          xlim=c(0,280), ylim=c(0,85), col=c("red", "black")[Environment], label=F)

### Expected richness with 95% confidence intervals ###

min(rowSums(species2)) ### minimum value of abundance showed in the rarefaction
curves (value = 135)

Srar<-rarefy(species2, 135, se=T) ### average values of expected richness for each plot
(standardized by the minimum value) and their standard errors

rarefaction <-read.table("Rarefação_Confiança.txt", header=T) ### data containing the
average values of expected richness for each plot and their 95% confidence intervals
(1.96 * standard error)

ggplot(rarefaction, aes(x=Group, y=Average, group=1)) +
  geom_errorbar(aes(ymin= Average-IC, ymax= Average+IC), width=.1) +
  geom_point() +
  theme_bw() +
  theme(panel.border = element_blank(), panel.grid.major = element_blank(),
        panel.grid.minor = element_blank(), axis.line = element_line(colour =
"black")) +
  scale_x_discrete(limits=c("1","2","3","4","5","6","7","8")) +
  scale_y_continuous(breaks=seq(0, 70, 5)) +
  labs(x="Forest age", y="Expected richness")

```

Diversity profiles using Hill's series (Figure 6)

Naret, Kuruwaty, Missão, Bikut, Dipisow, MataKoporuhu, MataKuruwaty and MataDipisowNaret are the names of the sampled forest areas

```

Naret<-data2[1, 3:232]
Kuruwaty<- data2 [2, 3:232]
Missão<- data2 [3, 3:232]
Bikut<- data2 [4, 3:232]
Dipisow<- data2[5, 3:232]
MataKoporuhu<- data2[6, 3:232]
MataKuruwaty<- data2[7, 3:232]

```

```
MataDipisowNaret<- data2[8, 3:232]
```

```
nine<-renyi(Naret, hill=T)
```

```
seventeen<-renyi(Kuruwaty, hill=T)
```

```
twentyeight<-renyi(Missão, hill=T)
```

```
seventy <-renyi(Bikut, hill=T)
```

```
onehundredforty <-renyi(Dipisow, hill=T)
```

```
oldforest1<-renyi(MataKoporuhu, hill=T)
```

```
oldforest2<-renyi(MataKuruwaty, hill=T)
```

```
oldforest3<-renyi(MataDipisowNaret, hill=T)
```

```
plot.default(nine, type="l", lty=2, xlab="Alpha", ylab="Hill number", ylim=c(0,90))
```

```
points(nine, pch=16)
```

```
points(seventeen, type="l", lty=2)
```

```
points(seventeen, pch=16)
```

```
points(twentyeight, type="l", lty=2)
```

```
points(twentyeight, pch=16)
```

```
points(seventy, type="l", lty=2)
```

```
points(seventy, pch=16)
```

```
points(onehundredforty, type="l", lty=2)
```

```
points(onehundredforty, pch=16)
```

```
points(oldforest1, type="l", lty=2, col="red")
```

```
points(oldforest1, col="red", pch=16)
```

```
points(oldforest2, type="l", lty=2, col="red")
```

```
points(oldforest2, col="red", pch=16)
```

```
points(oldforest3, type="l", lty=2, col="red")
```

```
points(oldforest3, col="red", pch=16)
```