

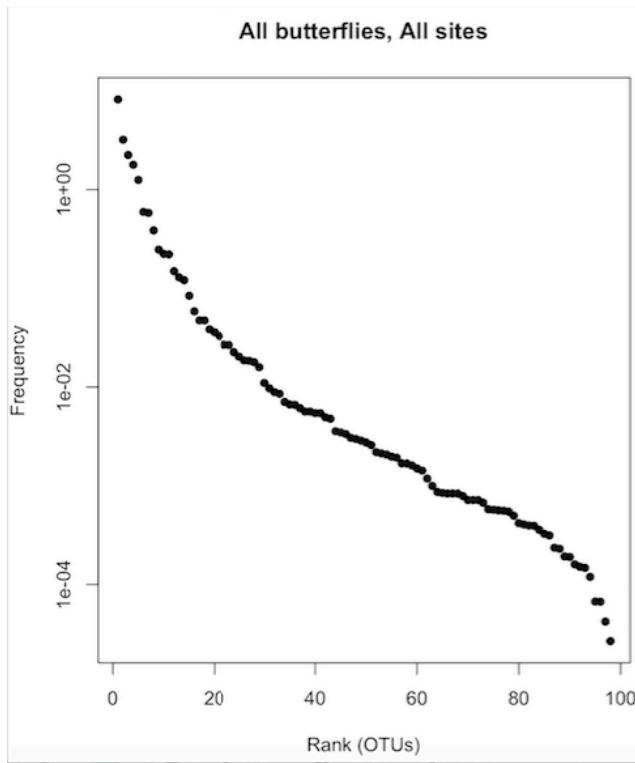
```
library(RADanalysis)
library(BiodiversityR)
library(codyn) # Explore this one...
```

Function to normalize data

```
normalize <- function(x) {
  return ((x - min(x)) / (max(x) - min(x)))
}
```

```
data=read_xlsx("/Users/Daniel/Desktop/BES project 2014/BES analysis 2020/
data1.xlsx") # prueba con Brown argus
data=as.data.frame(data)
data_original=data
data=data[,-c(1,2,3,4)]
datasites=data[1]
data[1]=datasites
#data = setNames(data.frame(t(data[,-1])), data[,1]) # Transpose a dataframe and
keep first column as column names
data_old = data %>% filter(str_detect(site, "old")) # filters old sites
data_new = data %>% filter(str_detect(site, "new")) # filters new sites
olddata = setNames(data.frame(t(data_old[,-1])), data_old[,1]) # Transpose a
dataframe and keep first column as column names
newdata = setNames(data.frame(t(data_new[,-1])), data_new[,1])
nrow(olddata)-length(which(rowSums(olddata)==0)) # Number of OTUs in old range:
55 - 90
(nrow(olddata)-length(which(rowSums(olddata)==0)))/nrow(olddata) # Same in
percentage
nrow(newdata)-length(which(rowSums(newdata)==0)) # Number of OTUs in new
range: 53 - 82
(nrow(newdata)-length(which(rowSums(newdata)==0)))/nrow(newdata) # Same in
percentage

q=colSums(data[,-1])
plot(rev(sort(q)),log="y",pch=16,xlab="Rank (OTUs)",ylab="Frequency",main="All
butterflies, All sites")
```



Per butterfly species

```

dataBA=filter(data_original,butterfly=="BrownArgus")
dataBA=dataBA[,-c(1,2,3,4)]
datasites=dataBA[1]
dataBA[1]=datasites
dataBA=dataBA[,-(which(colSums(dataBA[, -1])==0)+1)] # We consider only OTUs
reported in the butterfly's range (old+new), not OTUs from other butterflies
dataBA_old = dataBA %>% filter(str_detect(site, "old")) # filters old sites
dataBA_new = dataBA %>% filter(str_detect(site, "new")) # filters new sites
olddataBA = setNames(data.frame(t(dataBA_old[, -1])), dataBA_old[, 1]) # Transpose
a dataframe and keep first column as column names
newdataBA = setNames(data.frame(t(dataBA_new[, -1])), dataBA_new[, 1])
#nrow(olddataBA)-length(which(rowSums(olddataBA)==0)) # Number of OTUs in old
range: 47 - 73
#(nrow(olddataBA)-length(which(rowSums(olddataBA)==0)))/nrow(olddataBA) #
Same in percentage
#nrow(newdataBA)-length(which(rowSums(newdataBA)==0)) # Number of OTUs in
new range: 44 - 72
#(nrow(newdataBA)-length(which(rowSums(newdataBA)==0)))/nrow(newdataBA) #
Same in percentage
dataSW=filter(data_original,butterfly=="SpeckledWood")
dataSW=dataSW[,-c(1,2,3,4)]
datasites=dataSW[1]
dataSW[1]=datasites
dataSW=dataSW[,-(which(colSums(dataSW[, -1])==0)+1)] # We consider only OTUs
reported in the butterfly's range (old+new), not OTUs from other butterflies
dataSW_old = dataSW %>% filter(str_detect(site, "old")) # filters old sites
dataSW_new = dataSW %>% filter(str_detect(site, "new")) # filters new sites
olddataSW = setNames(data.frame(t(dataSW_old[, -1])), dataSW_old[, 1]) #

```

```

Transpose a dataframe and keep first column as column names
newdataSW = setNames(data.frame(t(dataSW_new[,-1])), dataSW_new[,1])
#nrow(olddataSW)-length(which(rowSums(olddataSW)==0)) # Number of OTUs in
old range: 54
#(nrow(olddataSW)-length(which(rowSums(olddataSW)==0)))/nrow(olddataSW) #
Same in percentage
#nrow(newdataSW)-length(which(rowSums(newdataSW)==0)) # Number of OTUs in
new range: 53
#(nrow(newdataSW)-length(which(rowSums(newdataSW)==0)))/nrow(newdataSW)
# Same in percentage
dataSSS=filter(data_original,butterfly=="SSSkipper")
dataSSS=dataSSS[,-c(1,2,3,4)]
datasites=dataSSS[1]
dataSSS[1]=datasites
dataSSS=dataSSS[,-(which(colSums(dataSSS[,-1])==0)+1)] # We consider only
OTUs reported in the butterfly's range (old+new), not OTUs from other butterflies
dataSSS_old = dataSSS %>% filter(str_detect(site, "old")) # filters old sites
dataSSS_new = dataSSS %>% filter(str_detect(site, "new")) # filters new sites
olddataSSS = setNames(data.frame(t(dataSSS_old[,-1])), dataSSS_old[,1]) #
Transpose a dataframe and keep first column as column names
newdataSSS = setNames(data.frame(t(dataSSS_new[,-1])), dataSSS_new[,1])
#nrow(olddataSSS)-length(which(rowSums(olddataSSS)==0)) # Number of OTUs in
old range: 52
#(nrow(olddataSSS)-length(which(rowSums(olddataSSS)==0)))/nrow(olddataSSS) #
Same in percentage
#nrow(newdataSSS)-length(which(rowSums(newdataSSS)==0)) # Number of OTUs
in new range: 29
#(nrow(newdataSSS)-length(which(rowSums(newdataSSS)==0)))/
nrow(newdataSSS) # Same in percentage

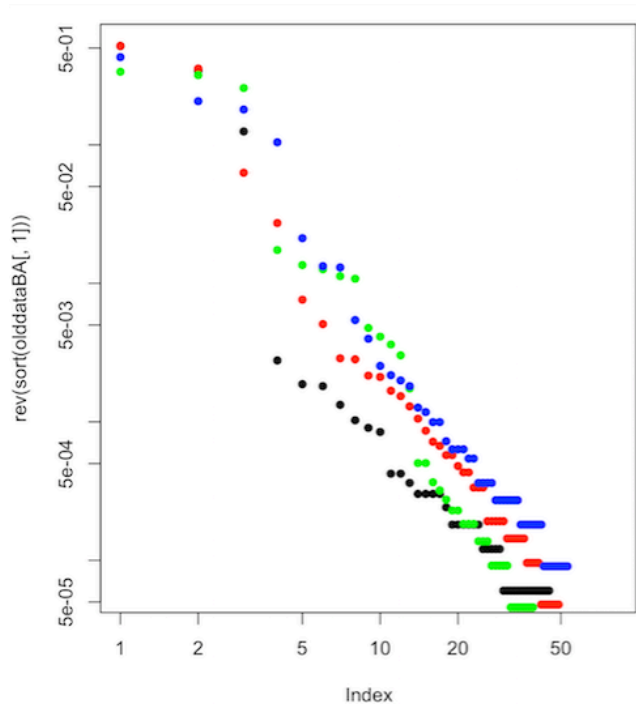
```

#####

```

#par(mfrow=c(2,2)) # Plotting 4 old x new sites
plot(rev(sort(olddataBA[,1])),log="y",pch=16,xlab="Rank",ylab="Abundance")
points(rev(sort(olddataBA[,2])),log="y",pch=16,col="red")
points(rev(sort(olddataBA[,3])),log="y",pch=16,col="green")
points(rev(sort(olddataBA[,4])),log="y",pch=16,col="blue")
points(rev(sort(newdataBA[,1])),log="y",pch=4)
points(rev(sort(newdataBA[,2])),log="y",pch=4,col="red")
points(rev(sort(newdataBA[,4])),log="y",pch=4,col="green")
points(rev(sort(newdataBA[,5])),log="y",pch=4,col="blue")

```



https://cran.r-project.org/web/packages/RADanalysis/vignettes/RADanalysis_intro.html

```
rads <- olddataBA
```

```
#plot original rads
```

```
line_cols <- c("black","green","red","blue")
```

```
#to specify different stages of subjects
```

```
sample_classes <- c(1,2,3,4)
```

```
plot(1,col = "white", xlim=c(1,60), ylim=c(0.00001,1),log = "y",axes = F,xlab =  
"Rank",ylab = "Abundance",main = "")
```

```
sfsmisc::eaxis(side = 1,at = c(1,10,20,30,50,100))
```

```
sfsmisc::eaxis(side = 2,at = c(0.00001,0.001,0.01,0.1,1))
```

```
for(i in 1:ncol(rads)){
```

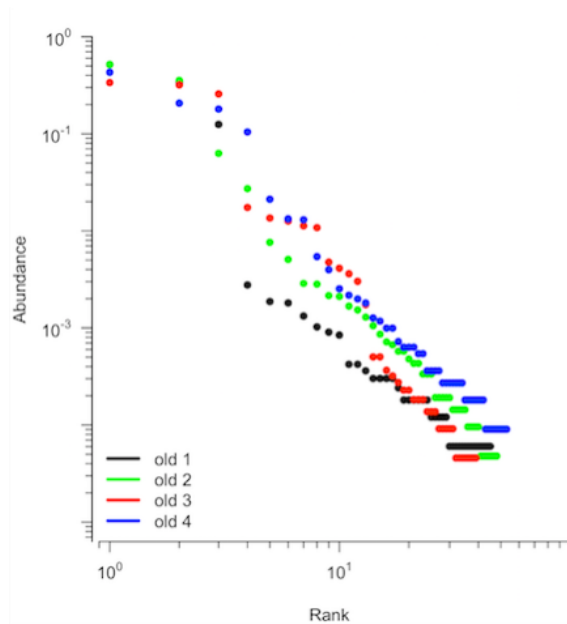
```
  temp <- sort(rads[-1,i],decreasing = TRUE)
```

```
  temp <- temp[temp>0]
```

```
  points(x=temp,lwd = 2,col = line_cols[sample_classes[i]], pch=16)
```

```
}
```

```
legend("topright",bty = "n",legend = c("old 1","old 2","old 3","old 4"),col =  
line_cols,lwd = 3)
```



Different fits

```
mod <- radfit(olddataBA[,1])
mod
plot(mod)
```

#Zipf fits: the smaller the absolute value of the parameter, the more even the distribution of abundances in the community. **In gral, parameters are higher in expanded margins → more uneven distributions in expanded ranges. This info must be taken together with the next section on RADs**

Brown argus

old 1: 0.67
 old 2: 0.66
 old 3: 0.54
 old 4: 0.55
mean old = 0.605
 new 1: 0.75
 new 2: 0.58
 new 3: 0.54
 new 4: 0.56
 new 5: 0.51
mean new: 0.6

Speckled wood

old 5: 0.62
 old 6: 0.62
 old 7: 0.58
mean old: 0.61
 new 6: 0.50
 new 7: 0.81
 new 8: 0.70
mean new: 0.67

Silver-spotted skipper

old 8: 0.56

old 9: 0.47

old 10: 0.48

mean old: 0.50

new 9: 0.59

new 10: 0.62

mean new: 0.605

#####

BROWN ARGUS

We sort the datasets in decreasing rank-abundance order:

olddataBA2=matrix(nrow=nrow(olddataBA),ncol=ncol(olddataBA))

colnames(olddataBA2)=c("old1","old2","old3","old4")

newdataBA2=matrix(nrow=nrow(newdataBA),ncol=ncol(newdataBA))

colnames(newdataBA2)=c("new1","new2","new3","new4","new5")

for(i in 1:ncol(olddataBA)) {

 olddataBA2[,i] <- sort(olddataBA[,i],decreasing = TRUE)

}

for(i in 1:ncol(newdataBA)) {

 newdataBA2[,i] <- sort(newdataBA[,i],decreasing = TRUE)

}

newdataBA2=newdataBA2[,-c(3)]# Remove BA new site #3

We calculate min and max values to have the range for the ribbon plots:

olddataBA and newdataBA have the same number of rows

oldBA=matrix(nrow=nrow(olddataBA2),ncol=4)

colnames(oldBA)=c("OTU","min","max","range")

newBA=matrix (nrow=nrow(newdataBA2),ncol=4)

colnames(newBA)=c("OTU","min","max","range")

for(i in 1:nrow(olddataBA2))

{

 oldmax=max(olddataBA2[i,])

 x=olddataBA2[i,]

 oldmin=min(x[which(x>0)])

 newmax=max(newdataBA2[i,])

 y=newdataBA2[i,]

 newmin=min(x[which(x>0)])

 oldBA[i,1]=i

 oldBA[i,2]=oldmin

 oldBA[i,3]=oldmax

 oldBA[i,4]=1 # OLD RANGE

 newBA[i,1]=i

 newBA[i,2]=newmin

 newBA[i,3]=newmax

 newBA[i,4]=2 # NEW RANGE

```

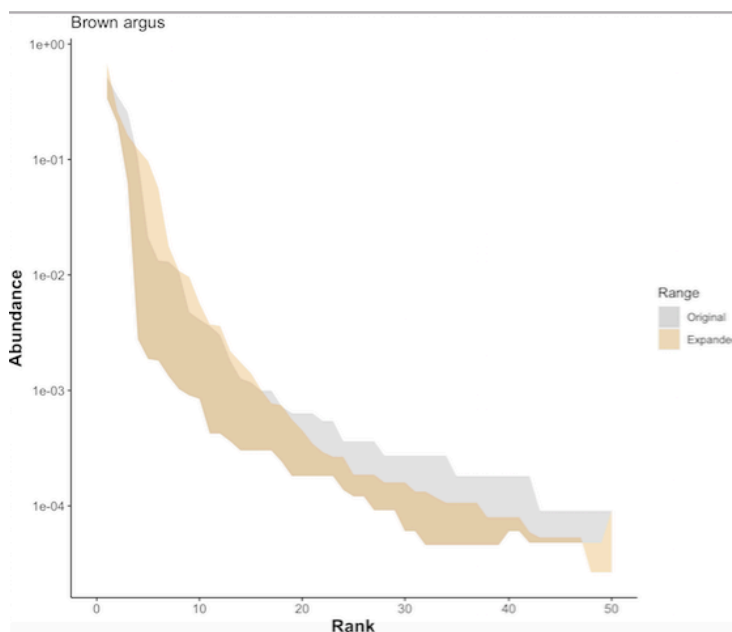
    }
    BA=rbind(oldBA,newBA)
    BA=as.data.frame(BA)
    BA[sapply(BA, is.infinite)] <- NA

```

```

ggplot(BA) + geom_ribbon(aes(x=OTU, ymin=min, ymax=max, fill=factor(range)),
alpha=0.3) +
theme(axis.text=element_text(size=10),axis.title=element_text(size=14,face="bold"))
+ theme(panel.grid.major = element_blank(), panel.grid.minor =
element_blank(),panel.background = element_blank(), axis.line =
element_line(colour = "black")) + labs(x="Rank",y="Abundance",title="Brown argus")
+ labs(color = "OTU") + scale_fill_manual(values=c("#999999", "#E69F00"),
name="Range", breaks=c("Original","Expanded")) + scale_y_log10() +
xlim(0,50)

```



SPECKLED WOOD

We sort the datasets in decreasing rank-abundance order:

```

olddataSW2=matrix(nrow=nrow(olddataSW),ncol=ncol(olddataSW))
colnames(olddataSW2)=c("old5","old6","old7")
newdataSW2=matrix(nrow=nrow(newdataSW),ncol=ncol(newdataSW))
colnames(newdataSW2)=c("new6","new7","new8")
for(i in 1:ncol(olddataSW)) {
  olddataSW2[,i] <- sort(olddataSW[,i],decreasing = TRUE)
}
for(i in 1:ncol(newdataSW)) {
  newdataSW2[,i] <- sort(newdataSW[,i],decreasing = TRUE)
}

```

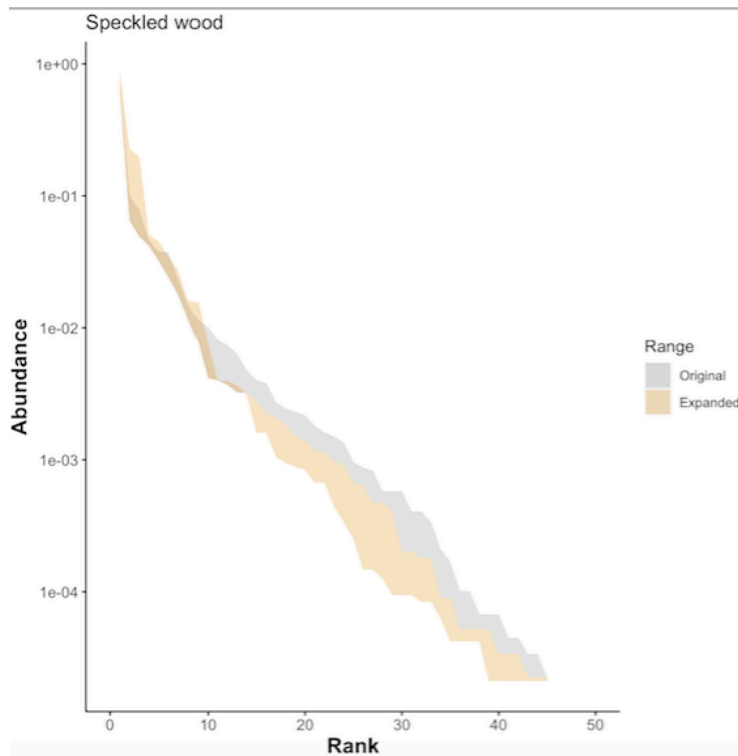
We calculate min and max values to have the range for the ribbon plots:

olddataSW and newdataSW have the same number of rows

```
oldSW=matrix(nrow=nrow(olddataSW2),ncol=4)
colnames(oldSW)=c("OTU","min","max","range")
newSW=matrix (nrow=nrow(newdataSW2),ncol=4)
colnames(newSW)=c("OTU","min","max","range")
```

```
for(i in 1:nrow(olddataSW2))
{
  oldmax=max(olddataSW2[i,])
  x=olddataSW2[i,]
  oldmin=min(x[which(x>0)])
  newmax=max(newdataSW2[i,])
  y=newdataSW2[i,]
  newmin=min(x[which(x>0)])
  oldSW[i,1]=i
  oldSW[i,2]=oldmin
  oldSW[i,3]=oldmax
  oldSW[i,4]=1 # OLD RANGE
  newSW[i,1]=i
  newSW[i,2]=newmin
  newSW[i,3]=newmax
  newSW[i,4]=2 # NEW RANGE
}
SW=rbind(oldSW,newSW)
SW=as.data.frame(SW)
SW[sapply(SW, is.infinite)] <- NA
```

```
ggplot(SW) + geom_ribbon(aes(x=OTU, ymin=min, ymax=max, fill=factor(range)),
alpha=0.3) +
theme(axis.text=element_text(size=10),axis.title=element_text(size=14,face="bold"))
+ theme(panel.grid.major = element_blank(), panel.grid.minor =
element_blank(),panel.background = element_blank(), axis.line =
element_line(colour = "black")) + labs(x="Rank",y="Abundance",title="Speckled
wood") + labs(color = "OTU") + scale_fill_manual(values=c("#999999", "#E69F00"),
name="Range", breaks=c(1,2), labels=c("Original","Expanded")) + scale_y_log10() +
xlim(0,50)
```

SILVER-SPOTTED SKIPPER

We sort the datasets in decreasing rank-abundance order:

```
olddataSSS2=matrix(nrow=nrow(olddataSSS),ncol=ncol(olddataSSS))
colnames(olddataSSS2)=c("old8","old9","old10")
newdataSSS2=matrix(nrow=nrow(newdataSSS),ncol=ncol(newdataSSS))
colnames(newdataSSS2)=c("new9","new10")
for(i in 1:ncol(olddataSSS)) {
  olddataSSS2[,i] <- sort(olddataSSS[,i],decreasing = TRUE)
}
for(i in 1:ncol(newdataSSS)) {
  newdataSSS2[,i] <- sort(newdataSSS[,i],decreasing = TRUE)
}
```

We calculate min and max values to have the range for the ribbon plots:

olddataSSS and newdataSSS have the same number of rows

```
oldSSS=matrix(nrow=nrow(olddataSSS2),ncol=4)
colnames(oldSSS)=c("OTU","min","max","range")
newSSS=matrix(nrow=nrow(newdataSSS2),ncol=4)
colnames(newSSS)=c("OTU","min","max","range")
```

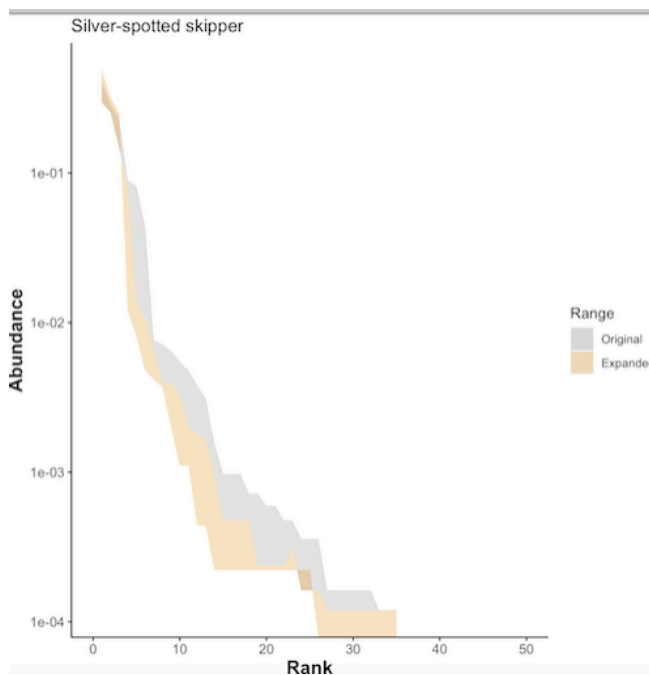
```
for(i in 1:nrow(olddataSSS2))
{
  oldmax=max(olddataSSS2[i,])
  x=olddataSSS2[i,]
  oldmin=min(x[which(x>0)])
  newmax=max(newdataSSS2[i,])
  y=newdataSSS2[i,]
  newmin=min(y[which(y>0)])
}
```

```

oldSSS[i,1]=i
oldSSS[i,2]=oldmin
oldSSS[i,3]=oldmax
oldSSS[i,4]=1 # OLD RANGE
newSSS[i,1]=i
newSSS[i,2]=newmin
newSSS[i,3]=newmax
newSSS[i,4]=2 # NEW RANGE
}
SSS=rbind(oldSSS,newSSS)
SSS=as.data.frame(SSS)
SSS[is.na(SSS)] <- NA

ggplot(SSS) + geom_ribbon(aes(x=OTU, ymin=min, ymax=max, fill=factor(range)),
alpha=0.3) +
theme(axis.text=element_text(size=10),axis.title=element_text(size=14,face="bold"))
+ theme(panel.grid.major = element_blank(), panel.grid.minor =
element_blank(),panel.background = element_blank(), axis.line =
element_line(colour = "black")) + labs(x="Rank",y="Abundance",title="Silver-spotted
skipper") + labs(color = "OTU") + scale_fill_manual(values=c("#999999",
"#E69F00"), name="Range", breaks=c(1,2), labels=c("Original","Expanded")) +
scale_y_log10() + xlim(0,50)

```



CHECK WHAT OTUs ARE MOST ABUNDANT: most species in new range are already in the old range → Abundant species in old range tend to also be abundant in the new range

Using colours for high, intermediate, low OTU abundances

Brown argus old #1

```
q1=as.data.frame(cbind(gsub("OTU", "",
rownames(olddataBA)),olddataBA[,1],rep(0,nrow(olddataBA))))
colnames(q1)=c("OTU","old1","cat")
q1$OTU=as.numeric(as.character(q1$OTU))
q1$old1=as.numeric(as.character(q1$old1))
q1$cat=as.numeric(as.character(q1$cat))
q1=q1 %>% arrange(desc(old1))
for (i in c(1:nrow(q1))) {
  if (q1[i,2]>0.01) {
    q1[i,3]="black"
  }
  if (q1[i,2]<=0.01 && q1[i,2]>0.001) {
    q1[i,3]="blue"
  }
  if (q1[i,2]<0.001) {
    q1[i,3]="red"
  }
}
```

Brown argus old #2

```
q2=as.data.frame(cbind(gsub("OTU", "",
rownames(olddataBA)),olddataBA[,2],rep(0,nrow(olddataBA))))
colnames(q2)=c("OTU","old2","cat")
q2$OTU=as.numeric(as.character(q2$OTU))
q2$old2=as.numeric(as.character(q2$old2))
q2$cat=as.numeric(as.character(q2$cat))
q2$cat=(q1 %>% arrange(OTU))[3]$cat
q2=q2 %>% arrange(desc(old2))
```

Brown argus old #3

```
q3=as.data.frame(cbind(gsub("OTU", "",
rownames(olddataBA)),olddataBA[,3],rep(0,nrow(olddataBA))))
colnames(q3)=c("OTU","old3","cat")
q3$OTU=as.numeric(as.character(q3$OTU))
q3$old3=as.numeric(as.character(q3$old3))
q3$cat=as.numeric(as.character(q3$cat))
q3$cat=(q1 %>% arrange(OTU))[3]$cat
q3=q3 %>% arrange(desc(old3))
```

Brown argus old #4

```
q4=as.data.frame(cbind(gsub("OTU", "",
rownames(olddataBA)),olddataBA[,4],rep(0,nrow(olddataBA))))
colnames(q4)=c("OTU","old4","cat")
q4$OTU=as.numeric(as.character(q4$OTU))
q4$old4=as.numeric(as.character(q4$old4))
q4$cat=as.numeric(as.character(q4$cat))
q4$cat=(q1 %>% arrange(OTU))[3]$cat
q4=q4 %>% arrange(desc(old4))
```

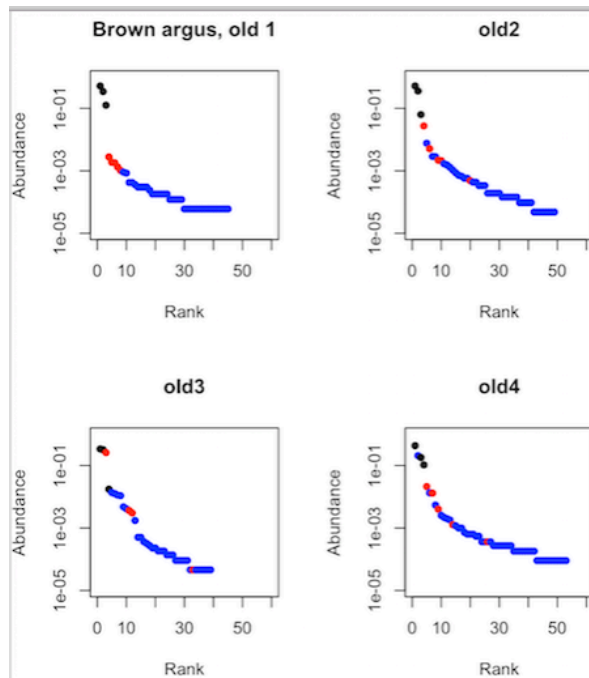
Plotting

```
par(mfrow=c(2,2))
plot(rev(sort(q1[,2])),log="y",xlim=c(0,60),ylim=c(0.00001,1),col=q1$cat,pch=16,xlab
="Rank",ylab ="Abundance",main="Brown argus, old 1")
```

```

plot(rev(sort(q2[,2])),log="y",xlim=c(0,60),ylim=c(0.00001,1),col=q2$cat,pch=16,xlab
="Rank",ylab ="Abundance",main="old2")
plot(rev(sort(q3[,2])),log="y",xlim=c(0,60),ylim=c(0.00001,1),col=q3$cat,pch=16,xlab
="Rank",ylab ="Abundance",main="old3")
plot(rev(sort(q4[,2])),log="y",xlim=c(0,60),ylim=c(0.00001,1),col=q4$cat,pch=16,xlab
="Rank",ylab ="Abundance",main="old4")

```



RUN THE FOLLOWING CODE TO CHANGE ABUNDANCE CRITERIA IN THE ANALYSIS OF CORE COMMUNITITES!!!!!! (ALSO THE PREVIOSI LINES MUST BE RUN)

Same as above, but pooling ranges (old sites together, new sites together) and comparing old Vs new

Brown argus OLD SITES

```

olddataBAall=as.data.frame(rowSums(olddataBA))
newdataBAall=as.data.frame(rowSums(newdataBA))
olddataBAall=as.data.frame(cbind(gsub("OTU", "",
rownames(olddataBAall)),olddataBAall[,1],rep(0,nrow(olddataBAall))))
colnames(olddataBAall)=c("OTU","old","cat")
olddataBAall$OTU=as.numeric(as.character(olddataBAall$OTU))
olddataBAall$old=as.numeric(as.character(olddataBAall$old))
olddataBAall$old=normalize(olddataBAall$old) # We normalize data to constraint
values to {0,1}
olddataBAall$cat=as.numeric(as.character(olddataBAall$cat))
olddataBAall=olddataBAall %>% arrange(desc(old))
for (i in c(1:nrow(olddataBAall))) {
  if (olddataBAall[i,2]>0.01) {
    olddataBAall[i,3]="black"
  }
}

```

```

if (olddataBAall[i,2]<=0.01 && olddataBAall[i,2]>0.0001) {
  olddataBAall[i,3]="blue"
}
if (olddataBAall[i,2]<=0.0001) {
  olddataBAall[i,3]="red"
}
}

```

Brown argus NEW SITES

```

newdataBAall=as.data.frame(rowSums(newdataBA))
newdataBAall=as.data.frame(rowSums(newdataBA))
newdataBAall=as.data.frame(cbind(gsub("OTU", "",
rownames(newdataBAall)),newdataBAall[,1],rep(0,nrow(newdataBAall))))
colnames(newdataBAall)=c("OTU","new","cat")
newdataBAall$OTU=as.numeric(as.character(newdataBAall$OTU))
newdataBAall$new=as.numeric(as.character(newdataBAall$new))
newdataBAall$new=normalize(newdataBAall$new) # We normalize data to
constraint values to {0,1}
newdataBAall$cat=as.numeric(as.character(newdataBAall$cat))
newdataBAall$cat=(olddataBAall %>% arrange(OTU))[3]$cat
newdataBAall=newdataBAall %>% arrange(desc(new))

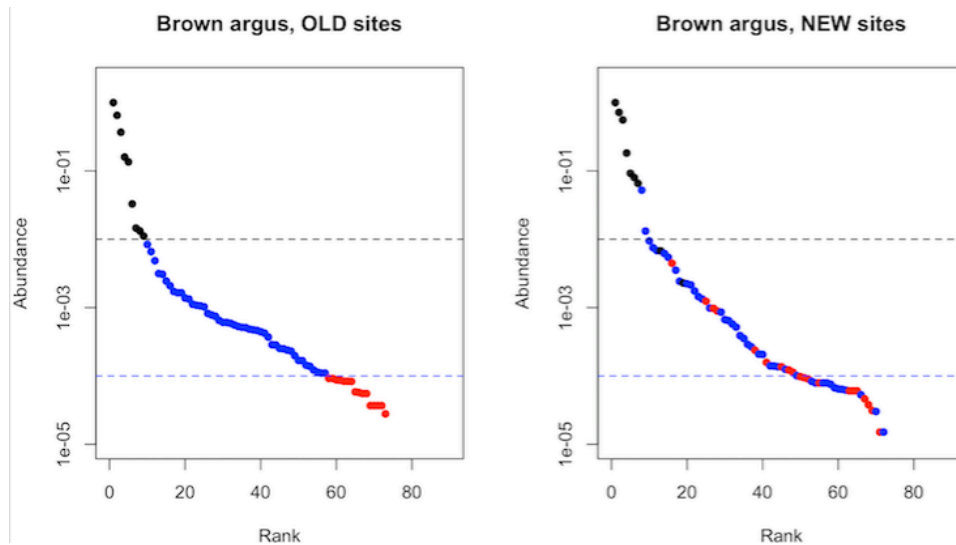
```

Plotting

```

par(mfrow=c(1,2))
plot(rev(sort(olddataBAall[,2])),log="y",xlim=c(0,90),ylim=c(0.00001,2),col=olddataBA
all$cat,pch=16,xlab="Rank",ylab ="Abundance",main="Brown argus \nOLD sites")
abline(h=0.01,col="black",lty=2)
abline(h=0.0001,col="blue",lty=2)
text(55,0.3,"High",cex=0.8)
text(65,0.001,"Intermediate",cex=0.8)
text(15,0.000025,"Low",cex=0.8)
plot(rev(sort(newdataBAall[,2])),log="y",xlim=c(0,90),ylim=c(0.00001,2),col=newdata
BAall$cat,pch=16,xlab="Rank",ylab ="Abundance",main="Brown argus \nNEW
sites")
abline(h=0.01,col="black",lty=2)
abline(h=0.0001,col="blue",lty=2)
text(55,0.3,"High",cex=0.8)
text(65,0.001,"Intermediate",cex=0.8)
text(15,0.000025,"Low",cex=0.8)

```



Speckled wood OLD SITES

```

olddataSWall=as.data.frame(rowSums(olddataSW))
newdataSWall=as.data.frame(rowSums(newdataSW))
olddataSWall=as.data.frame(cbind(gsub("OTU", "",
rownames(olddataSWall)),olddataSWall[,1],rep(0,nrow(olddataSWall))))
colnames(olddataSWall)=c("OTU","old","cat")
olddataSWall$OTU=as.numeric(as.character(olddataSWall$OTU))
olddataSWall$old=as.numeric(as.character(olddataSWall$old))
olddataSWall$old=normalize(olddataSWall$old) # We normalize data to constraint
values to {0,1}
olddataSWall$cat=as.numeric(as.character(olddataSWall$cat))
olddataSWall=olddataSWall %>% arrange(desc(old))
for (i in c(1:nrow(olddataSWall))) {
  if (olddataSWall[i,2]>0.01) {
    olddataSWall[i,3]="black"
  }
  if (olddataSWall[i,2]<=0.01 && olddataSWall[i,2]>0.0001) {
    olddataSWall[i,3]="blue"
  }
  if (olddataSWall[i,2]<=0.0001) {
    olddataSWall[i,3]="red"
  }
}

```

Speckled wood NEW SITES

```

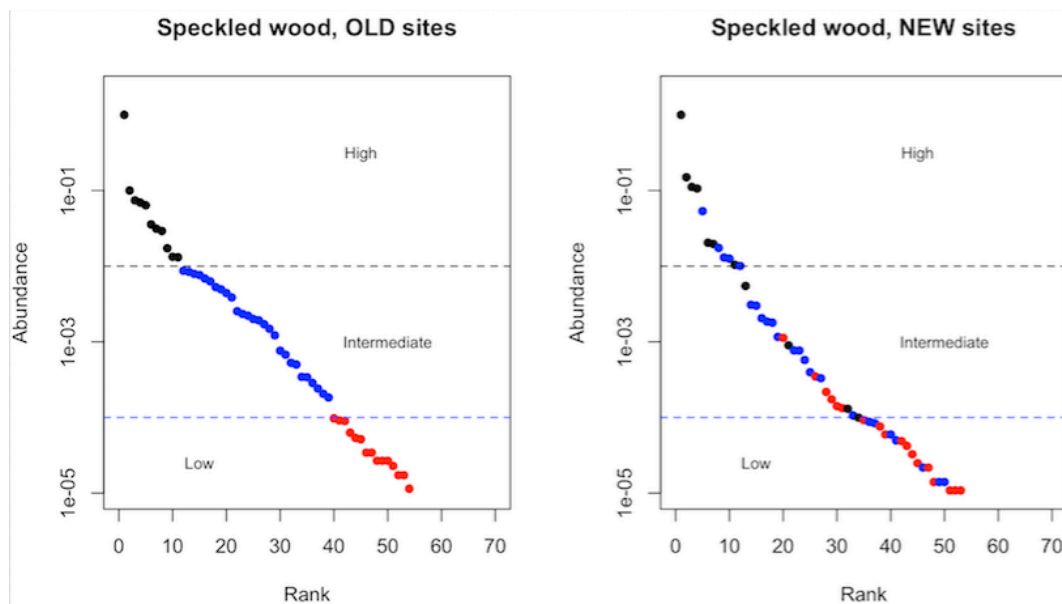
newdataSWall=as.data.frame(rowSums(newdataSW))
newdataSWall=as.data.frame(rowSums(newdataSW))
newdataSWall=as.data.frame(cbind(gsub("OTU", "",
rownames(newdataSWall)),newdataSWall[,1],rep(0,nrow(newdataSWall))))
colnames(newdataSWall)=c("OTU","new","cat")
newdataSWall$OTU=as.numeric(as.character(newdataSWall$OTU))
newdataSWall$new=as.numeric(as.character(newdataSWall$new))
newdataSWall$new=normalize(newdataSWall$new) # We normalize data to
constraint values to {0,1}
newdataSWall$cat=as.numeric(as.character(newdataSWall$cat))
newdataSWall$cat=(olddataSWall %>% arrange(OTU))[3]$cat

```

```

newdataSWall=newdataSWall %>% arrange(desc(new))
# Plotting
par(mfrow=c(1,2))
plot(rev(sort(olddataSWall[,2])),log="y",xlim=c(0,70),ylim=c(0.00001,2),col=olddataSWall$cat,pch=16,xlab="Rank",ylab="Abundance",main="Speckled wood \nOLD sites")
abline(h=0.01,col="black",lty=2)
abline(h=0.0001,col="blue",lty=2)
text(45,0.3,"High",cex=0.8)
text(50,0.001,"Intermediate",cex=0.8)
text(15,0.000025,"Low",cex=0.8)
plot(rev(sort(newdataSWall[,2])),log="y",xlim=c(0,70),ylim=c(0.00001,2),col=newdataSWall$cat,pch=16,xlab="Rank",ylab="Abundance",main="Speckled wood \nNEW sites")
abline(h=0.01,col="black",lty=2)
abline(h=0.0001,col="blue",lty=2)
text(45,0.3,"High",cex=0.8)
text(50,0.001,"Intermediate",cex=0.8)
text(15,0.000025,"Low",cex=0.8)

```



Silver-spotted skipper OLD SITES

```

olddataSSSal=as.data.frame(rowSums(olddataSSS))
newdataSSSal=as.data.frame(rowSums(newdataSSS))
olddataSSSal=as.data.frame(cbind(gsub("OTU", "",
rownames(olddataSSSal)),olddataSSSal[,1],rep(0,nrow(olddataSSSal))))
colnames(olddataSSSal)=c("OTU","old","cat")
olddataSSSal$OTU=as.numeric(as.character(olddataSSSal$OTU))
olddataSSSal$old=as.numeric(as.character(olddataSSSal$old))
olddataSSSal$old=normalize(olddataSSSal$old) # We normalize data to constraint
values to {0,1}
olddataSSSal$cat=as.numeric(as.character(olddataSSSal$cat))
olddataSSSal=olddataSSSal %>% arrange(desc(old))
# WE CHANGE CRITERIA TO ESTABLISH CATEGORIES AS THE LOW

```

ABUNDANCE OTUs WITH THE CRITERIA ABOVE DOES NOT FIND ANY OTU

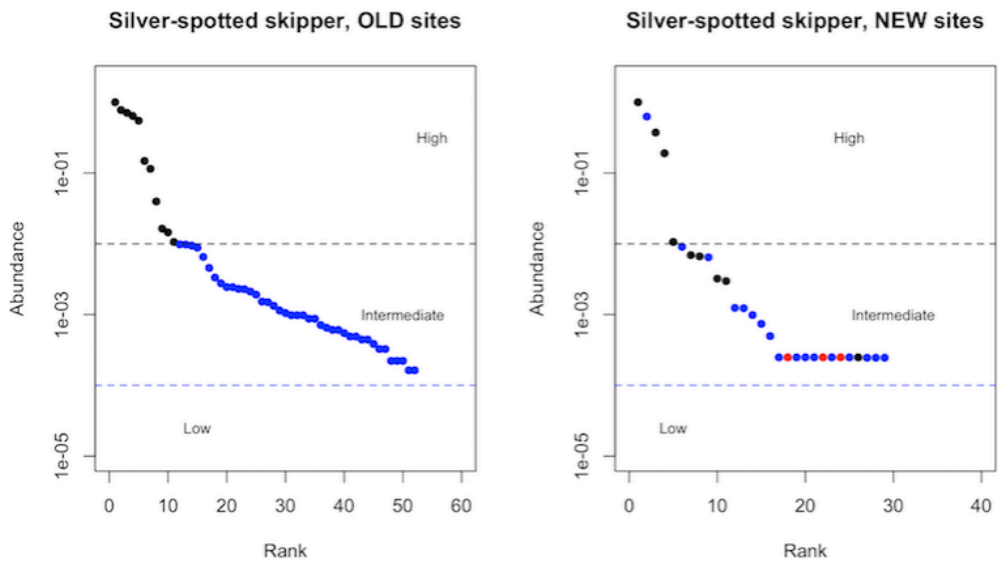
```
for (i in c(1:nrow(olddataSSSal))) {  
  if (olddataSSSal[i,2]>0.01) {  
    olddataSSSal[i,3]="black"  
  }  
  if (olddataSSSal[i,2]<=0.01 && olddataSSSal[i,2]>0.001) {  
    olddataSSSal[i,3]="blue"  
  }  
  if (olddataSSSal[i,2]<=0.001) {  
    olddataSSSal[i,3]="red"  
  }  
}
```

Silver-spotted skipper NEW SITES

```
newdataSSSal=as.data.frame(rowSums(newdataSSS))  
newdataSSSal=as.data.frame(rowSums(newdataSSS))  
newdataSSSal=as.data.frame(cbind(gsub("OTU", "",  
rownames(newdataSSSal)),newdataSSSal[,1],rep(0,nrow(newdataSSSal))))  
colnames(newdataSSSal)=c("OTU","new","cat")  
newdataSSSal$OTU=as.numeric(as.character(newdataSSSal$OTU))  
newdataSSSal$new=as.numeric(as.character(newdataSSSal$new))  
newdataSSSal$new=normalize(newdataSSSal$new) # We normalize data to  
constraint values to {0,1}  
newdataSSSal$cat=as.numeric(as.character(newdataSSSal$cat))  
newdataSSSal$cat=(olddataSSSal %>% arrange(OTU))[3]$cat  
newdataSSSal=newdataSSSal %>% arrange(desc(new))
```

Plotting

```
par(mfrow=c(1,2))  
plot(rev(sort(olddataSSSal[,2])),log="y",xlim=c(0,60),ylim=c(0.0001,2),col=olddataSSSal$cat,pch=16,xlab="Rank",ylab="Abundance",main="Silver-spotted skipper  
\\nOLD sites")  
abline(h=0.01,col="black",lty=2)  
abline(h=0.001,col="blue",lty=2)  
text(40,0.3,"High",cex=0.8)  
text(40,0.005,"Intermediate",cex=0.8)  
text(15,0.00025,"Low",cex=0.8)  
plot(rev(sort(newdataSSSal[,2])),log="y",xlim=c(0,40),ylim=c(0.0001,2),col=newdataSSSal$cat,pch=16,xlab="Rank",ylab="Abundance",main="Silver-spotted skipper  
\\nNEW sites")  
abline(h=0.01,col="black",lty=2)  
abline(h=0.001,col="blue",lty=2)  
text(25,0.3,"High",cex=0.8)  
text(30,0.005,"Intermediate",cex=0.8)  
text(5,0.00025,"Low",cex=0.8)
```

```
#####
#####
oldBA=cbind(rownames(olddataBA),olddataBA)
oldBA %>% arrange(desc(old1)) # 60,62,67
oldBA %>% arrange(desc(old2)) # 60,67,62
oldBA %>% arrange(desc(old3)) # 67,60,84,62
oldBA %>% arrange(desc(old4)) # 67,39,62,60
newBA=cbind(rownames(newdataBA),newdataBA)
newBA %>% arrange(desc(new1)) # 67,60,62
newBA %>% arrange(desc(new2)) # 67,60,62,84
newBA %>% arrange(desc(new3)) # 67,62,60
newBA %>% arrange(desc(new4)) # 60,39,18(this is exclusive to new
range!),67,84,62
newBA %>% arrange(desc(new5)) # 62,67,84,79,60
oldSW=cbind(rownames(olddataSW),olddataSW)
oldSW %>% arrange(desc(old5)) # 67,90,60
oldSW %>% arrange(desc(old6)) # 67,18,79,60
oldSW %>% arrange(desc(old7)) # 67,60,79,84
newSW=cbind(rownames(newdataSW),newdataSW)
newSW %>% arrange(desc(new6)) # 67,79,88,84
newSW %>% arrange(desc(new7)) # 67,79,84,88
newSW %>% arrange(desc(new8)) # 67,84,47,79
oldSSS=cbind(rownames(olddataSSS),olddataSSS)
oldSSS %>% arrange(desc(old8)) # 79,78,67,60
oldSSS %>% arrange(desc(old9)) # 84,79,60,106(exclusive of old range),67,62
oldSSS %>% arrange(desc(old10)) # 67,78,60,84,62,79
newSSS=cbind(rownames(newdataSSS),newdataSSS)
newSSS %>% arrange(desc(new9)) # 79,67,81
newSSS %>% arrange(desc(new10)) # 79,81,60,67
#####
#####
```

Beta-diversity analysis: comparing old Vs new sites for each

butterfly

We use betadiver(), which calculates beta diversity indices of Koleff et al (JAE 2003, Table 1).

Below, we select Whittaker's index

BROWN ARGUS

```
olddataBAall_ab1=filter(olddataBAall,cat=="black")
```

```
olddataBAall_ab2=filter(olddataBAall,cat=="blue")
```

```
olddataBAall_ab3=filter(olddataBAall,cat=="red")
```

For new sites, we need to locate most, intermediate and least abundant species. Before we use eye comparisons to see if there's changes in the RADs between old and new ranges. Now we want to do a beta-diversity analysis

```
for (i in c(1:nrow(newdataBAall))) {  
  if (newdataBAall[i,2]>0.01) {  
    newdataBAall[i,3]="black"  
  }  
  if (newdataBAall[i,2]<=0.01 && newdataBAall[i,2]>0.0001) {  
    newdataBAall[i,3]="blue"  
  }  
  if (newdataBAall[i,2]<=0.0001) {  
    newdataBAall[i,3]="red"  
  }  
}
```

```
newdataBAall_ab1=filter(newdataBAall,cat=="black")
```

```
newdataBAall_ab2=filter(newdataBAall,cat=="blue")
```

```
newdataBAall_ab3=filter(newdataBAall,cat=="red")
```

Most abundant species

```
BA_ab1=cbind(olddataBAall_ab1[,1:2],newdataBAall_ab1[,1:2])
```

```
BA_ab1 <- full_join(olddataBAall_ab1[,1:2],newdataBAall_ab1[,1:2], by = c('OTU'))
```

```
BA_ab1[is.na(BA_ab1)] <- 0
```

```
BA_ab1=as.data.frame(t(BA_ab1))
```

```
colnames(BA_ab1)=BA_ab1[,1]
```

```
BA_ab1[,1]=rownames(BA_ab1)
```

```
BA_ab1=BA_ab1[-1,]
```

```
BA_ab1.beta=betadiver(BA_ab1,method="w") #0.22
```

Intermediate abundant species

```
BA_ab2=cbind(olddataBAall_ab2[,1:2],newdataBAall_ab2[,1:2])
```

```
BA_ab2 <- full_join(olddataBAall_ab2[,1:2],newdataBAall_ab2[,1:2], by = c('OTU'))
```

```
BA_ab2[is.na(BA_ab2)] <- 0
```

```
BA_ab2=as.data.frame(t(BA_ab2))
```

```
colnames(BA_ab2)=BA_ab2[,1]
```

```
BA_ab2[,1]=rownames(BA_ab2)
```

```
BA_ab2=BA_ab2[-1,]
```

```
BA_ab2.beta=betadiver(BA_ab2,method="w") #0.34
```

High abundant species

```
BA_ab3=cbind(olddataBAall_ab3[,1:2],newdataBAall_ab3[,1:2])
```

```
BA_ab3 <- full_join(olddataBAall_ab3[,1:2],newdataBAall_ab3[,1:2], by = c('OTU'))
```

```
BA_ab3[is.na(BA_ab3)] <- 0
```

```
BA_ab3=as.data.frame(t(BA_ab3))
```

```
colnames(BA_ab3)=BA_ab3[,1]
```

```
BA_ab3[,1]=rownames(BA_ab3)
BA_ab3=BA_ab3[-1,]
BA_ab3.beta=betadiver(BA_ab3,method="w") #0.75
```

SPECKLED WOOD

```
olddataSWall_ab1=filter(olddataSWall,cat=="black")
olddataSWall_ab2=filter(olddataSWall,cat=="blue")
olddataSWall_ab3=filter(olddataSWall,cat=="red")
# For new sites, we need to locate most, intermediate and least abundant species.
BEfore we use eye comparisons to see if there's changes in the RADs between old
and new ranges. Now we want to do a beta-diversity analysis
```

```
for (i in c(1:nrow(newdataSWall))) {
  if (newdataSWall[i,2]>0.01) {
    newdataSWall[i,3]="black"
  }
  if (newdataSWall[i,2]<=0.01 && newdataSWall[i,2]>0.0001) {
    newdataSWall[i,3]="blue"
  }
  if (newdataSWall[i,2]<=0.0001) {
    newdataSWall[i,3]="red"
  }
}
newdataSWall_ab1=filter(newdataSWall,cat=="black")
newdataSWall_ab2=filter(newdataSWall,cat=="blue")
newdataSWall_ab3=filter(newdataSWall,cat=="red")
# Most abundant species
SW_ab1=cbind(olddataSWall_ab1[,1:2],newdataSWall_ab1[,1:2])
SW_ab1 <- full_join(olddataSWall_ab1[,1:2],newdataSWall_ab1[,1:2], by = c('OTU'))
SW_ab1[is.na(SW_ab1)] <- 0
SW_ab1=as.data.frame(t(SW_ab1))
colnames(SW_ab1)=SW_ab1[1,]
SW_ab1[,1]=rownames(SW_ab1)
SW_ab1=SW_ab1[-1,]
SW_ab1.beta=betadiver(SW_ab1,method="w") #0.39
# Intermediate abundant species
SW_ab2=cbind(olddataSWall_ab2[,1:2],newdataSWall_ab2[,1:2])
SW_ab2 <- full_join(olddataSWall_ab2[,1:2],newdataSWall_ab2[,1:2], by = c('OTU'))
SW_ab2[is.na(SW_ab2)] <- 0
SW_ab2=as.data.frame(t(SW_ab2))
colnames(SW_ab2)=SW_ab2[1,]
SW_ab2[,1]=rownames(SW_ab2)
SW_ab2=SW_ab2[-1,]
SW_ab2.beta=betadiver(SW_ab2,method="w") #0.48
# High abundant species
SW_ab3=cbind(olddataSWall_ab3[,1:2],newdataSWall_ab3[,1:2])
SW_ab3 <- full_join(olddataSWall_ab3[,1:2],newdataSWall_ab3[,1:2], by = c('OTU'))
SW_ab3[is.na(SW_ab3)] <- 0
SW_ab3=as.data.frame(t(SW_ab3))
colnames(SW_ab3)=SW_ab3[1,]
SW_ab3[,1]=rownames(SW_ab3)
```

```
SW_ab3=SW_ab3[-1,]  
SW_ab3.beta=betadiver(SW_ab3,method="w") #0.71
```

SILVER-SPOTTED SKIPPER

```
olddataSSSal_ab1=filter(olddataSSSal,cat=="black")  
olddataSSSal_ab2=filter(olddataSSSal,cat=="blue")  
olddataSSSal_ab3=filter(olddataSSSal,cat=="red")  
# For new sites, we need to locate most, intermediate and least abundant species.  
BEfore we use eye comparisons to see if there's changes in the RADs between old  
and new ranges. Now we want to do a beta-diversity analysis
```

```
for (i in c(1:nrow(newdataSSSal))) {  
  if (newdataSSSal[i,2]>0.01) {  
    newdataSSSal[i,3]="black"  
  }  
  if (newdataSSSal[i,2]<=0.01 && newdataSSSal[i,2]>0.001) {  
    newdataSSSal[i,3]="blue"  
  }  
  if (newdataSSSal[i,2]<=0.001) {  
    newdataSSSal[i,3]="red"  
  }  
}  
  
newdataSSSal_ab1=filter(newdataSSSal,cat=="black")  
newdataSSSal_ab2=filter(newdataSSSal,cat=="blue")  
newdataSSSal_ab3=filter(newdataSSSal,cat=="red")  
# Most abundant species  
SSS_ab1=cbind(olddataSSSal_ab1[,1:2],newdataSSSal_ab1[,1:2])  
SSS_ab1 <- full_join(olddataSSSal_ab1[,1:2],newdataSSSal_ab1[,1:2], by =  
c('OTU'))  
SSS_ab1[is.na(SSS_ab1)] <- 0  
SSS_ab1=as.data.frame(t(SSS_ab1))  
colnames(SSS_ab1)=SSS_ab1[,1]  
SSS_ab1[,1]=rownames(SSS_ab1)  
SSS_ab1=SSS_ab1[-1,]  
SSS_ab1.beta=betadiver(SSS_ab1,method="w") #0.5  
# Intermediate abundant species  
SSS_ab2=cbind(olddataSSSal_ab2[,1:2],newdataSSSal_ab2[,1:2])  
SSS_ab2 <- full_join(olddataSSSal_ab2[,1:2],newdataSSSal_ab2[,1:2], by =  
c('OTU'))  
SSS_ab2[is.na(SSS_ab2)] <- 0  
SSS_ab2=as.data.frame(t(SSS_ab2))  
colnames(SSS_ab2)=SSS_ab2[,1]  
SSS_ab2[,1]=rownames(SSS_ab2)  
SSS_ab2=SSS_ab2[-1,]  
SSS_ab2.beta=betadiver(SSS_ab2,method="w") #0.79  
# High abundant species  
SSS_ab3=cbind(olddataSSSal_ab3[,1:2],newdataSSSal_ab3[,1:2])  
SSS_ab3 <- full_join(olddataSSSal_ab3[,1:2],newdataSSSal_ab3[,1:2], by =  
c('OTU'))  
SSS_ab3[is.na(SSS_ab3)] <- 0  
SSS_ab3=as.data.frame(t(SSS_ab3))
```

```
colnames(SSS_ab3)=SSS_ab3[1,]  
SSS_ab3[,1]=rownames(SSS_ab3)  
SSS_ab3=SSS_ab3[-1,]  
SSS_ab3.beta=betadiver(SSS_ab3,method="w") #0.69
```