

Bacterial community analysis for surfaces in a university classroom

Meadow *et al.* “Bacterial communities on classroom surfaces vary with human contact”

This document contains all statistical analyses conducted for the manuscript. Note that due to the random iterative nature of some analyses (such as beta-rarefaction, DB-RDA & PCA) some of the figure parameters will change slightly during reanalysis, though core results will remain essentially unchanged.

All data to reproduce analysis can be found here:

https://github.com/jfmeadow/Meadow_eta1_Surfaces_Analysis

Load necessary ecological analysis libraries.

```
library(vegan)
```

```
## Loading required package: permute This is vegan 2.0-8
```

```
library(labdsv)
```

```
## Loading required package: mgcv This is mgcv 1.7-22. For overview
type
## 'help("mgcv-package")'. Loading required package: MASS
##
## Attaching package: 'labdsv'
##
## The following object is masked from 'package:stats':
##
## density
```

Pull workspace from big OTU table created for Lillis Air data (Meadow et al. 2013, Indoor Air). Also removed a list of plants and 'no blast hits' as described in that manuscript.

This beginning workspace contains:

- `lillis.big.table`: OTU table created with bigger sequence dataset containing airborne bacteria. Swabs are a subset of those samples.
- `swab.map`: metadata mapping file
- `big.streptos.names`: vector of plant chloroplast OTUs - these get removed.
- `big.nbh.names`: vector of OTUs with no GreenGenes hit to bacteria level - these are also excluded from analysis.
- `lillis.big.taxa`: taxonomic assignments from GreenGenes database.

```
load("lillis.RData")
source("functions.R") #functions for formatting shortcuts.
```

Extract OTU table and remove plant sequences as well as sequences not identified as bacteria.

```
swab.table.tmp <- lillis.big.table[row.names(swab.map), -
which(colnames(lillis.big.table) %in%
      c(big.streptos.names, big.nbh.names))]
swab.taxa <- lillis.big.taxa[-which(row.names(lillis.big.taxa) %in%
c(big.streptos.names,
  big.nbh.names)), ]
```

Some important metrics:

- total sequences: 799616
- total samples: 58
- total number of OTUs in this dataset: 5718
- total number of OTUs in bigger dataset (including air): 10782

Even the most depauperate has >4000 sequences, so rarefy to that level.

```
swab.table <- rrarefy(swab.table.tmp, 4000)
swab.table <- swab.table[, which(apply(swab.table, 2, sum) > 0)]
swab.taxa <- swab.taxa[colnames(swab.table), ]
```

After rarefaction, some important metrics:

- total sequences: 2.32×10^5
- total number of OTUs: 3776

Results:

Are communities indicative of surface type? In other words, do we have 4 distinct communities on these 4 distinct surfaces?

```
swab.can <- vegdist(swab.table, "canberra")
adonis(swab.can ~ swab.map$type)
```

```
##
## Call:
## adonis(formula = swab.can ~ swab.map$type)
##
## Terms added sequentially (first to last)
##
##              Df SumsOfSqs MeanSqs F.Model    R2 Pr(>F)
## swab.map$type  3      1.91  0.637    2.31 0.114 0.001 ***
## Residuals     54     14.88  0.276
## Total        57     16.79
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Yes. Definitely.

Distance-based Redundancy Analysis (DB-RDA) for discriminant analysis as well as visualization:

```
swab.caps <- capscale(swab.can ~ swab.map$type)
anova(swab.caps)
```

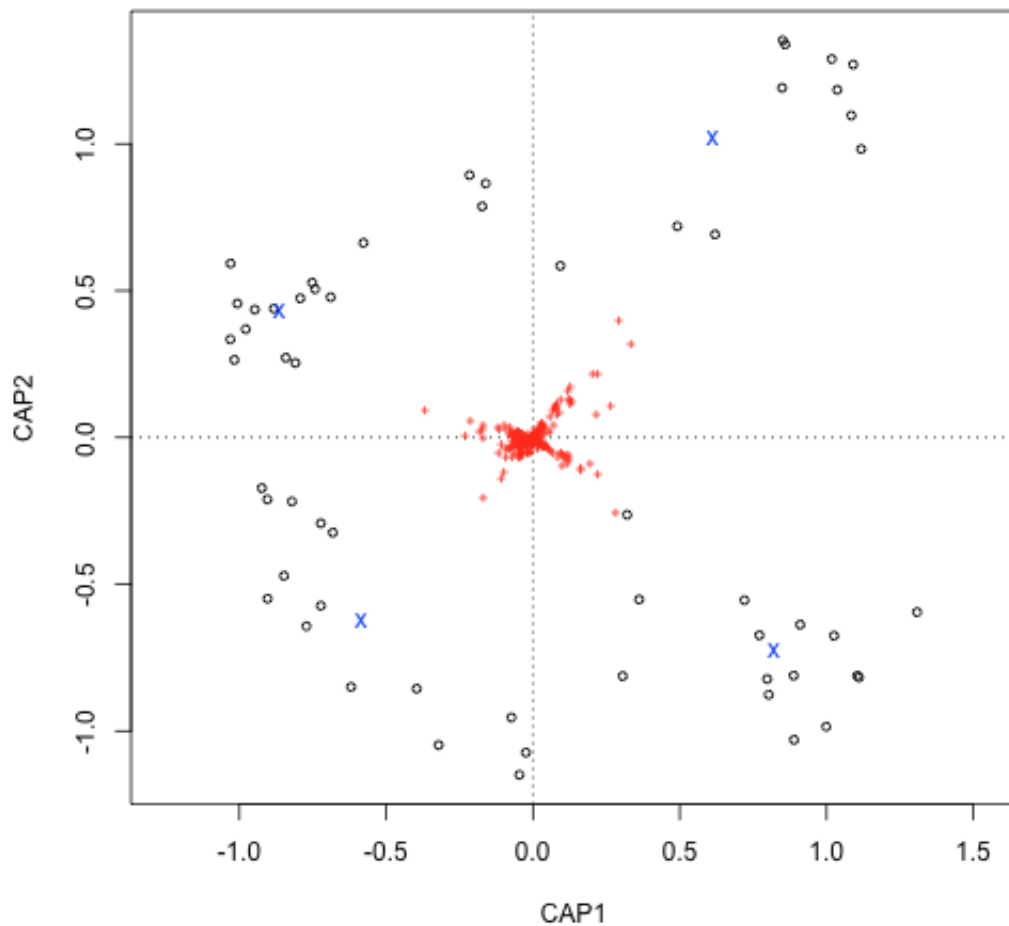
```
## Permutation test for capscale under reduced model
##
## Model: capscale(formula = swab.can ~ swab.map$type)
##              Df   Var      F N.Perm Pr(>F)
## Model         3  1.91  2.31    199 0.005 **
## Residual     54 14.88
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

This test is essentially identical, but with different iterations, so different p-value. Create a version of the DB-RDA that will have individual OTUs weighting samples:

```
swab.caps2 <- capscale(swab.table ~ swab.map$type, distance =
"canberra")
anova(swab.caps2)
```

```
## Permutation test for capscale under reduced model
##
## Model: capscale(formula = swab.table ~ swab.map$type, distance =
"canberra")
##      Df   Var    F N.Perm Pr(>F)
## Model    3  1.91 2.31   199  0.005 **
## Residual 54 14.88
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
scaps2.plot <- plot(swab.caps2)
```



This is all to get a dataframe for each set of indicator taxa sets the edit commands were used to manually change taxa names for plotting.

```
sp.caps <- data.frame(scaps2.plot$species[, c(1, 2)])
chair.caps.all <- sp.caps[which(sp.caps$CAP1 > 0 & sp.caps$CAP2 > 0),
]
desk.caps.all <- sp.caps[which(sp.caps$CAP1 > 0 & sp.caps$CAP2 < 0),
]
wall.caps.all <- sp.caps[which(sp.caps$CAP1 < 0 & sp.caps$CAP2 < 0),
]
floor.caps.all <- sp.caps[which(sp.caps$CAP1 < 0 & sp.caps$CAP2 > 0),
]
```

Function to find extant OTUs

```
hypot <- function(d.f) {  
  sqrt.d.f <- sqrt((d.f[, 1]^2) + (d.f[, 2]^2))  
  invisible(sqrt.d.f)  
}
```

Highlight those OTUs with strongest weights for each of the 4 groups. Started with top 10, then trimmed to significant indicators (from analysis below).

```
hypot.chair <- hypot(chair.caps.all)  
chair.tops <- chair.caps.all[rev(order(hypot.chair)), ][1:8, ]  
  
hypot.desk <- hypot(desk.caps.all)  
desk.tops <- desk.caps.all[rev(order(hypot.desk)), ][1:5, ]  
  
hypot.wall <- hypot(wall.caps.all)  
wall.tops <- wall.caps.all[rev(order(hypot.wall)), ][1:4, ]  
  
hypot.floor <- hypot(floor.caps.all)  
floor.tops <- floor.caps.all[rev(order(hypot.floor)), ][1:6, ]
```

Then save species scores for easier plotting.

```
sp.caps <- scores(swab.caps2)$species
```

Put taxon names on the OTUs. This requires the two functions intended to format taxonomy output. The output is a OTU classification dataframe with total abundance in the last column.

```
taxo <- cleanTaxo(makeTaxo(taxo.in = swab.taxa$taxa.names, otu.table  
= swab.table,  
  split = ";"))
```

```
## Warning: no non-missing arguments to max; returning -Inf Warning:  
no  
## non-missing arguments to max; returning -Inf Warning: no non-  
missing  
## arguments to max; returning -Inf Warning: no non-missing arguments  
to max;  
## returning -Inf Warning: no non-missing arguments to max; returning  
-Inf  
## Warning: no non-missing arguments to max; returning -Inf
```

```
head(taxo)
```

##	kingdom	phylum	class	order
## 1	Bacteria	Actinobacteria	Actinobacteria	Actinomycetales
## 7	Bacteria	SR1	-	-
## 10	Bacteria	Proteobacteria	Alphaproteobacteria	Sphingomonadales
## 11	Bacteria	Bacteroidetes	Sphingobacteria	Sphingobacteriales
## 12	Bacteria	Proteobacteria	Alphaproteobacteria	Rhizobiales
## 13	Bacteria	Cyanobacteria	Chloroplast	Chlorophyta
##	family	genus	abundance	
## 1	Corynebacteriaceae	Corynebacterium	4	
## 7	-	-	838	
## 10	Sphingomonadaceae	Sphingomonas	1640	
## 11	Flexibacteraceae	Dyadobacter	135	
## 12	-	-	10	
## 13	Trebouxiophyceae	-	184	

Put names on the right points.

```
chair.tops$taxa <- taxo[row.names(chair.tops), 6]
desk.tops$taxa <- taxo[row.names(desk.tops), 6]
wall.tops$taxa <- taxo[row.names(wall.tops), 6]
floor.tops$taxa <- taxo[row.names(floor.tops), 6]

chair.tops$pos <- 1
desk.tops$pos <- 1
wall.tops$pos <- 1
floor.tops$pos <- 1

cap.txt <- data.frame(x = c(1.3, 1.1, -0.8, -1), y = c(0.75, -0.2,
-1, 0.75))
```

Check for significance of each indicator pointed out. This uses the `indval` function in the `labdsv` package - it essentially follows the procedure outlined in Dufrene & Legendre (1998). A few steps follow to format `indval` output for downstream analysis.


```

indic <- indval(swab.table, swab.map$type)

ch.ids <- paste("X", row.names(chair.tops), sep = "")
de.ids <- paste("X", row.names(desk.tops), sep = "")
wa.ids <- paste("X", row.names(wall.tops), sep = "")
fl.ids <- paste("X", row.names(floor.tops), sep = "")

chair.tops$indic <- indic$pval[ch.ids]
desk.tops$indic <- indic$pval[de.ids]
wall.tops$indic <- indic$pval[wa.ids]
floor.tops$indic <- indic$pval[fl.ids]

all.tops <- rbind(chair.tops, desk.tops, wall.tops, floor.tops)
all.tops$surface <- factor(c(rep("chair", nrow(chair.tops)),
rep("desk", nrow(desk.tops)),
rep("wall", nrow(wall.tops)), rep("floor", nrow(floor.tops))))
for (i in 1:nrow(all.tops)) {
  all.tops$chair.ra[i] <- mean(swab.table[swab.map$type == "chair",
row.names(all.tops)[i]]/sum(swab.table[swab.map$type ==
"chair", ]))
  all.tops$desk.ra[i] <- mean(swab.table[swab.map$type == "desk",
row.names(all.tops)[i]]/sum(swab.table[swab.map$type ==
"desk", ]))
  all.tops$wall.ra[i] <- mean(swab.table[swab.map$type == "wall",
row.names(all.tops)[i]]/sum(swab.table[swab.map$type ==
"wall", ]))
  all.tops$floor.ra[i] <- mean(swab.table[swab.map$type == "floor",
row.names(all.tops)[i]]/sum(swab.table[swab.map$type ==
"floor", ]))
}

all.tops

```

##		CAP1	CAP2	taxa	pos	indic	surface
##	chair.ra						
##	17	0.29210	0.401617	Lactobacillus	1	0.001	chair
	2.482e-03						
##	141	0.33446	0.321560	Corynebacterium	1	0.001	chair
	2.737e-03						
##	639	0.22247	0.219930	Corynebacterium	1	0.001	chair
	1.807e-03						
##	10800	0.20437	0.219050	Corynebacterium	1	0.001	chair
	1.693e-03						
##	216	0.26637	0.113315	Staphylococcus	1	0.011	chair
	1.552e-03						
##	188	0.21619	0.081006	Staphylococcus	1	0.012	chair
	1.170e-03						
##	151	0.12730	0.174379	Lactobacillus	1	0.001	chair
	1.110e-03						
##	64	0.11925	0.164070	Lactobacillus	1	0.001	chair
	1.010e-03						
##	135	0.28188	-0.253388	CandidatusPhytoplasma	1	0.008	desk
	1.977e-04						
##	530	0.22282	-0.122807	Streptococcus	1	0.001	desk
	3.048e-04						
##	1961	0.19412	-0.082208	Brevundimonas	1	0.001	desk
	3.788e-04						
##	1333	0.16294	-0.107948	Streptococcus	1	0.001	desk
	1.288e-04						
##	153	0.16394	-0.101412	Streptococcus	1	0.001	desk

```

1.862e-04
## 942 -0.16849 -0.205022 Alicyclobacillus 1 0.001 wall
7.015e-04
## 314 -0.10698 -0.138775 Alicyclobacillus 1 0.001 wall
4.643e-04
## 1881 -0.09869 -0.112025 Rhodopseudomonas 1 0.001 wall
4.260e-04
## 21 -0.11687 -0.049085 Sphingomonas 1 0.054 wall
9.796e-04
## 174 -0.36685 0.094731 Salmonella 1 0.001 floor
2.283e-04
## 1312 -0.23127 0.005707 Chroococcidiopsis 1 0.104 floor
7.589e-04
## 1076 -0.21281 0.058633 Roseomonas 1 0.001 floor
1.849e-04
## 1203 -0.18086 0.024919 Chroococcidiopsis 1 0.045 floor
6.033e-04
## 69 -0.16836 0.045294 Salmonella 1 0.001 floor
9.566e-05
## 596 -0.17106 0.032129 - 1 0.001 floor
7.398e-05
## desk.ra wall.ra floor.ra
## 17 1.111e-04 1.089e-04 8.418e-05
## 141 8.211e-04 5.344e-04 4.605e-04
## 639 5.211e-04 3.133e-04 3.023e-04
## 10800 4.144e-04 2.544e-04 2.564e-04
## 216 9.644e-04 3.367e-04 3.304e-04
## 188 8.011e-04 2.156e-04 2.691e-04
## 151 6.000e-05 8.222e-05 4.209e-05
## 64 3.222e-05 4.444e-05 1.786e-05
## 135 1.880e-03 3.533e-04 2.666e-04
## 530 1.201e-03 1.267e-04 1.237e-04
## 1961 1.009e-03 1.256e-04 1.352e-04
## 1333 8.856e-04 7.444e-05 4.592e-05
## 153 9.322e-04 9.111e-05 1.186e-04
## 942 9.533e-04 2.113e-03 8.788e-04
## 314 6.589e-04 1.396e-03 5.880e-04
## 1881 4.944e-04 1.233e-03 4.592e-04
## 21 1.246e-03 1.358e-03 1.680e-03
## 174 3.600e-04 5.089e-04 2.305e-03
## 1312 4.578e-04 1.318e-03 1.365e-03
## 1076 2.033e-04 3.422e-04 1.332e-03
## 1203 3.344e-04 9.356e-04 1.122e-03
## 69 1.544e-04 2.156e-04 1.054e-03
## 596 1.744e-04 2.622e-04 1.040e-03

```

Then write a table for manual documentation of individual OTUs. This goes into Table 1.

```

write.table(all.tops, file = "surface_indicators.txt", sep = "\t",
quote = FALSE)

```

Put surface samples in context with gut, skin, soil, phyllosphere. First read in sources . csv; this is a Class-level OTU table from multiple source environments. The same approach was used in Kembel et al. (2012) to identify potential sources in airborne hospital bacterial assemblages.

```
## source environments
sources.tmp <- read.csv("sourceHabitatsBlastClass.csv")
sources <- sources.tmp[, c(2, 4, 3)]
names(sources) <- c("sample", "species", "abundance")

## prepare surface OTU table
swabClass <- aggregate(t(swab.table), by = list(taxo$class), FUN =
"sum")
row.names(swabClass) <- swabClass[, 1]
swabClass <- swabClass[, -1]
swabClass <- data.frame(t(swabClass/4000))
swabClass <- dematify(swabClass)

## combine and make longform
sourcesAll.tmp <- rbind(swabClass, sources)
sourcesAll <- matify(sourcesAll.tmp)

## make mapping factor
sourcesMap <- data.frame(env = rep("skin", nrow(sourcesAll)))
row.names(sourcesMap) <- rn <- row.names(sourcesAll)
sourcesMap$env <- as.character(sourcesMap$env)
sourcesMap$env[grepl("Swab", rn)] <- "classroom"
sourcesMap$env[grepl("Fcs", rn)] <- "gut"
sourcesMap$env[grepl("phyllosphere", rn)] <- "phyllosphere"
sourcesMap$env[grepl("soil", rn)] <- "soil"
sourcesMap$env[grepl("water", rn)] <- "water"
sourcesMap$env <- as.factor(sourcesMap$env)

sourcesMap$bg <- "transparent"
sourcesMap$bg[1:58] <- swab.map$bg
```

Then create ordination of these sources combined with surfaces samples.

```
sourcesPCA <- pca(sourcesAll)
plantskinPCA1 <- sourcesPCA$scores[, 1]
```

How much variance was explained by the axes?

```
(sourcesPCA$sdev[c(1:2)]^2)/sum(sourcesPCA$sdev^2)
```

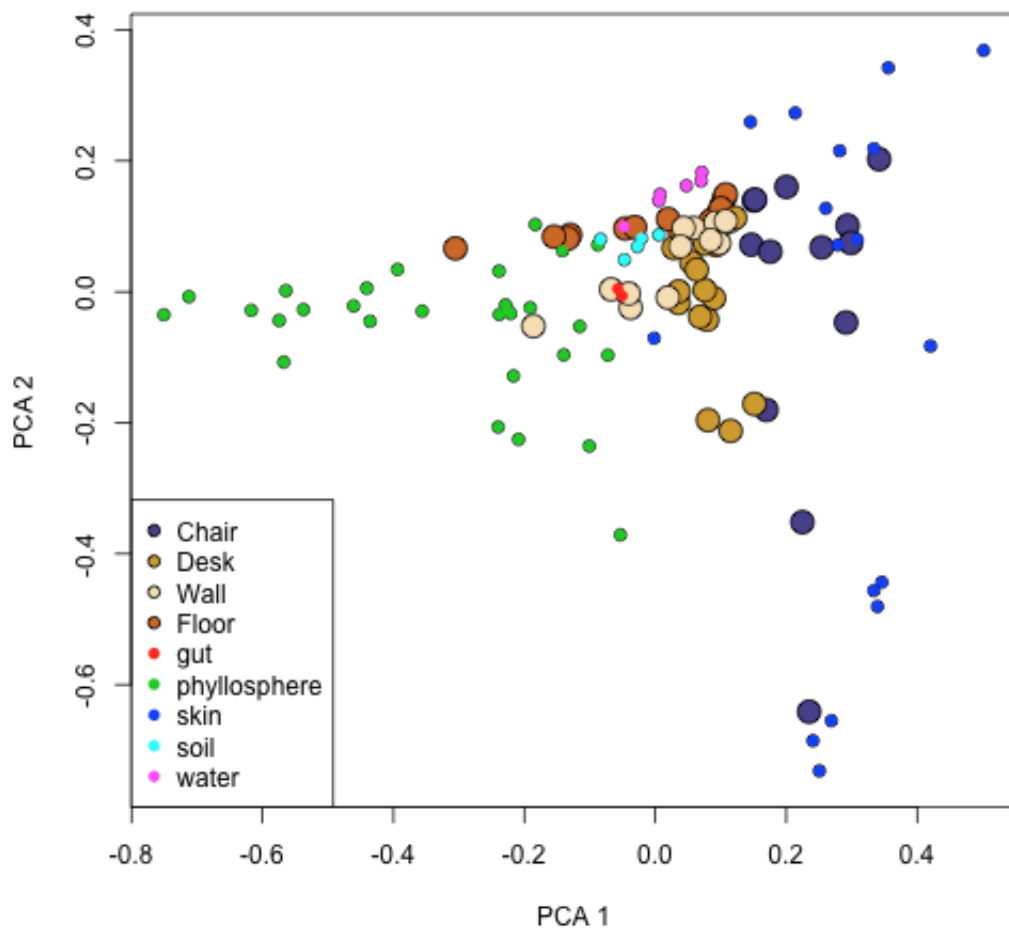
```
## [1] 0.3778 0.2320
```

Plot results

```

sourcesMap$col <- 1
for (i in 1:nlevels(sourcesMap$env)) {
  sourcesMap$col[sourcesMap$env == levels(sourcesMap$env)[i]] <- i
}
plot(sourcesPCA, type = "n")
points(sourcesPCA$scores[1:58, 1:2], pch = 21, bg =
sourcesMap$bg[1:58], cex = 2)
points(sourcesPCA$scores[-c(1:58), 1:2], col = sourcesMap$col[-
c(1:58)], pch = 16)
legend("bottomleft", legend = c("Chair", "Desk", "Wall", "Floor",
levels(sourcesMap$env)[-1]),
      pch = c(rep(21, 4), rep(16, 5)), col = c(1, 1, 1, 1, 2:6), pt.bg
= c("darkslateblue",
      "goldenrod3", "wheat", "chocolate3"))

```



```

boxType <- factor(swab.map$type, levels = c("chair", "desk", "wall",
"floor"))
library(IDPmisc)

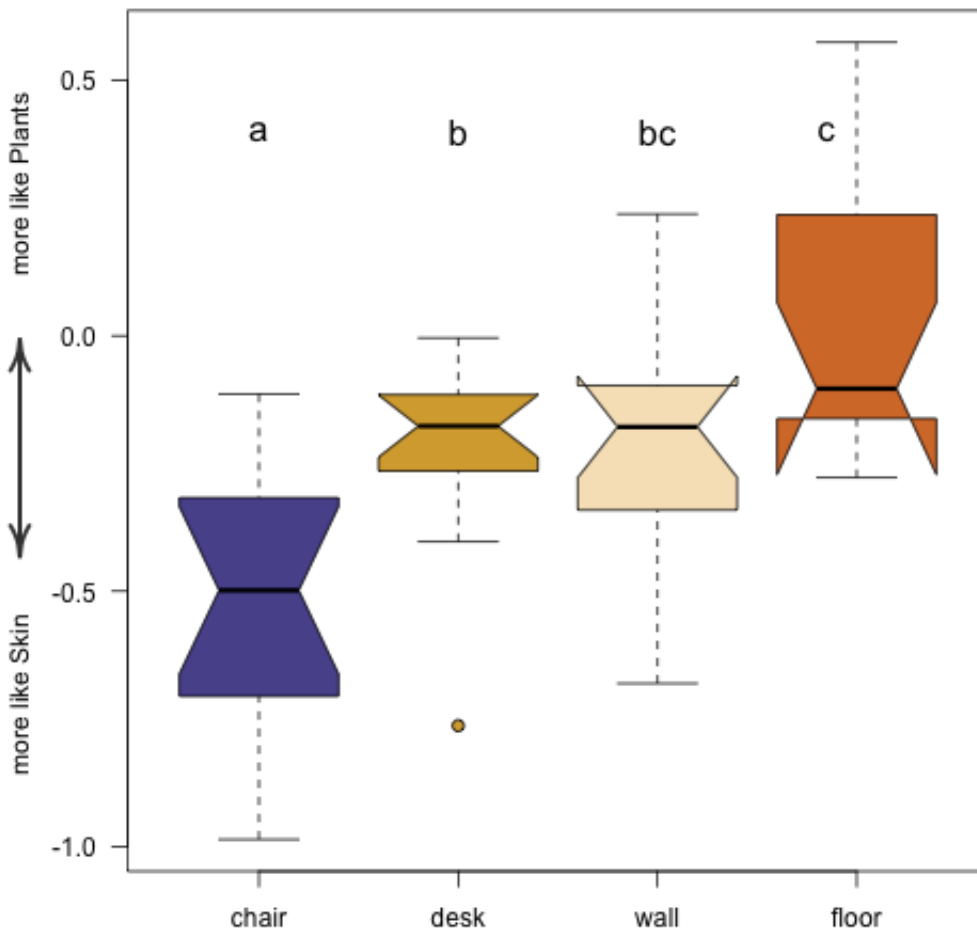
```

```
## Loading required package: grid Loading required package: lattice
```

```
par(mar = c(5, 6, 2, 1))
boxplot(swab.map$plantskin ~ boxType, notch = TRUE, col =
c("darkslateblue",
  "goldenrod3", "wheat", "chocolate3"), ylab = "", las = 1, pch =
21, bg = c("wheat",
  "goldenrod3"), pch = 1.5)
```

```
## Warning: some notches went outside hinges ('box'): maybe set
notch=FALSE
```

```
par(xpd = TRUE)
Arrows(-0.2, -0.0078, -0.2, -0.4317, sh.col = "gray20", sh.lwd = 2.5)
Arrows(-0.2, -0.4317, -0.2, -0.0078, sh.col = "gray20", sh.lwd = 2.5)
mtext(c("more like Skin", "more like Plants"), side = 2, at = c(-0.7,
0.3),
  line = 3)
text(c(1, 2, 3, 3.85), c(rep(0.4, 4)), c("a", "b", "bc", "c"), font =
1, cex = 1.4)
```



```
TukeyHSD(aov(swab.map$plantskin ~ swab.map$type))
```

```
## Tukey multiple comparisons of means
## 95% family-wise confidence level
##
## Fit: aov(formula = swab.map$plantskin ~ swab.map$type)
##
## $`swab.map$type`
##
```

	diff	lwr	upr	p adj
desk-chair	0.29182	0.06229	0.521342	0.0074
floor-chair	0.52387	0.29042	0.757322	0.0000
wall-chair	0.30429	0.07477	0.533818	0.0048
floor-desk	0.23206	0.00253	0.461581	0.0466
wall-desk	0.01248	-0.21306	0.238009	0.9989
wall-floor	-0.21958	-0.44911	0.009946	0.0656

This revised version of the figure contains information from the source PCA performed above. First some baggage from previous analysis.

```
library(IDPmisc)
boxType <- factor(swab.map$type, levels = c("chair", "desk", "wall",
"floor"))
boxTypeNum <- rep(1, nrow(swab.map))
boxTypeNum[boxType == "chair"] <- 1
boxTypeNum[boxType == "desk"] <- 2
boxTypeNum[boxType == "wall"] <- 3
boxTypeNum[boxType == "floor"] <- 4

chair.caps <- chair.tops
desk.caps <- desk.tops
wall.caps <- wall.tops
floor.caps <- floor.tops
```

```

layout(matrix(c(1, 2, 3), 1, 3), widths = c(1, 1, 0.75))
par(xpd = FALSE, mar = c(3, 3, 3, 1), las = 0)
plot(swab.caps, type = "none", xaxt = "n", yaxt = "n", xlab = "",
ylab = "")
mtext(c("CAP 1", "CAP 2"), side = c(1, 2), line = c(0.3, 0.1), adj =
c(1, 1))
points(swab.caps, "sites", pch = 21, bg = swab.map$bg, cex = 2)
text(c(cap.txt$x), c(cap.txt$y), c("Chairs", "Desks", "walls",
"Floors"), cex = 1)
mtext("a", adj = 0, font = 2, cex = 2)
plot(sp.caps, pch = 16, cex = 0.4, col = 8, xlim = c(range(sp.caps)),
ylim = c(range(sp.caps)),
xaxt = "n", yaxt = "n", xlab = "", ylab = "")
mtext(c("CAP 1", "CAP 2"), side = c(1, 2), line = c(0.3, 0.1), adj =
c(1, 1))
abline(h = 0, v = 0, lty = 3, lwd = 0.7, col = 1)
# rect(-1.5, -1.5, 2, 2)
par(xpd = TRUE)
points(chair.caps$CAP1, chair.caps$CAP2, pch = 21, bg =
"darkslateblue", cex = 1,
lwd = 0.5)
points(desk.caps$CAP1, desk.caps$CAP2, pch = 21, bg = "goldenrod3",
cex = 1,
lwd = 0.5)
points(wall.caps$CAP1, wall.caps$CAP2, pch = 21, bg = "wheat", cex =
1, lwd = 0.5)
points(floor.caps$CAP1, floor.caps$CAP2, pch = 21, bg = "chocolate3",
cex = 1,
lwd = 0.5)
text(chair.caps$CAP1, chair.caps$CAP2, as.character(chair.caps$taxa),
cex = 0.7,
pos = chair.caps$pos)
text(desk.caps$CAP1, desk.caps$CAP2, as.character(desk.caps$taxa),
cex = 0.7,
pos = desk.caps$pos)
text(wall.caps$CAP1, wall.caps$CAP2, as.character(wall.caps$taxa),
cex = 0.7,
pos = wall.caps$pos)
text(floor.caps$CAP1, floor.caps$CAP2, as.character(floor.caps$taxa),
cex = 0.7,
pos = floor.caps$pos)
mtext("b", adj = 0, font = 2, cex = 2)
par(mar = c(4, 5, 3, 1))
boxplot(swab.map$plantskin ~ boxType, notch = TRUE, col =
c("darkslateblue",
"goldenrod3", "wheat", "chocolate3"), ylab = "", las = 1, pch =
21, cex = 0,
bg = c("wheat", "goldenrod3"))

```

```

## Warning: some notches went outside hinges ('box'): maybe set
notch=FALSE

```


Create subsets of data for Mantel tests. I.e., test for geographic location within the room.

```
st.c <- swab.table[swab.map$type == "chair", ]
st.f <- swab.table[swab.map$type == "floor", ]
st.d <- swab.table[swab.map$type == "desk", ]
st.w <- swab.table[swab.map$type == "wall", ]

sm.c <- swab.map[swab.map$type == "chair", ]
sm.f <- swab.map[swab.map$type == "floor", ]
sm.d <- swab.map[swab.map$type == "desk", ]
sm.w <- swab.map[swab.map$type == "wall", ]

mantel(vegdist(st.c, "canberra"), vegdist(data.frame(sm.c$xcor,
sm.c$ycor),
"euclid"))
```

```
##
## Mantel statistic based on Pearson's product-moment correlation
##
## Call:
## mantel(xdis = vegdist(st.c, "canberra"), ydis =
vegdist(data.frame(sm.c$xcor,      sm.c$ycor), "euclid"))
##
## Mantel statistic r: -0.0308
##      Significance: 0.58
##
## Upper quantiles of permutations (null model):
##      90%      95% 97.5%      99%
## 0.190 0.240 0.284 0.352
##
## Based on 999 permutations
```

```
mantel(vegdist(st.f, "canberra"), vegdist(data.frame(sm.f$xcor,
sm.f$ycor),
"euclid"))
```

```
##
## Mantel statistic based on Pearson's product-moment correlation
##
## Call:
## mantel(xdis = vegdist(st.f, "canberra"), ydis =
vegdist(data.frame(sm.f$xcor,      sm.f$ycor), "euclid"))
##
## Mantel statistic r: 0.0393
##      Significance: 0.4
##
## Upper quantiles of permutations (null model):
##      90%      95% 97.5%      99%
## 0.205 0.248 0.288 0.320
##
## Based on 999 permutations
```

```
mantel(vegdist(st.d, "canberra"), vegdist(data.frame(sm.d$xcor,
sm.d$ycor),
"euclid"))
```

```
##
## Mantel statistic based on Pearson's product-moment correlation
##
## Call:
## mantel(xdis = vegdist(st.d, "canberra"), ydis =
vegdist(data.frame(sm.d$xcor,      sm.d$ycor), "euclid"))
##
## Mantel statistic r: -0.0671
##      Significance: 0.72
##
## Upper quantiles of permutations (null model):
##      90%      95%  97.5%    99%
## 0.156 0.211 0.256 0.295
##
## Based on 999 permutations
```

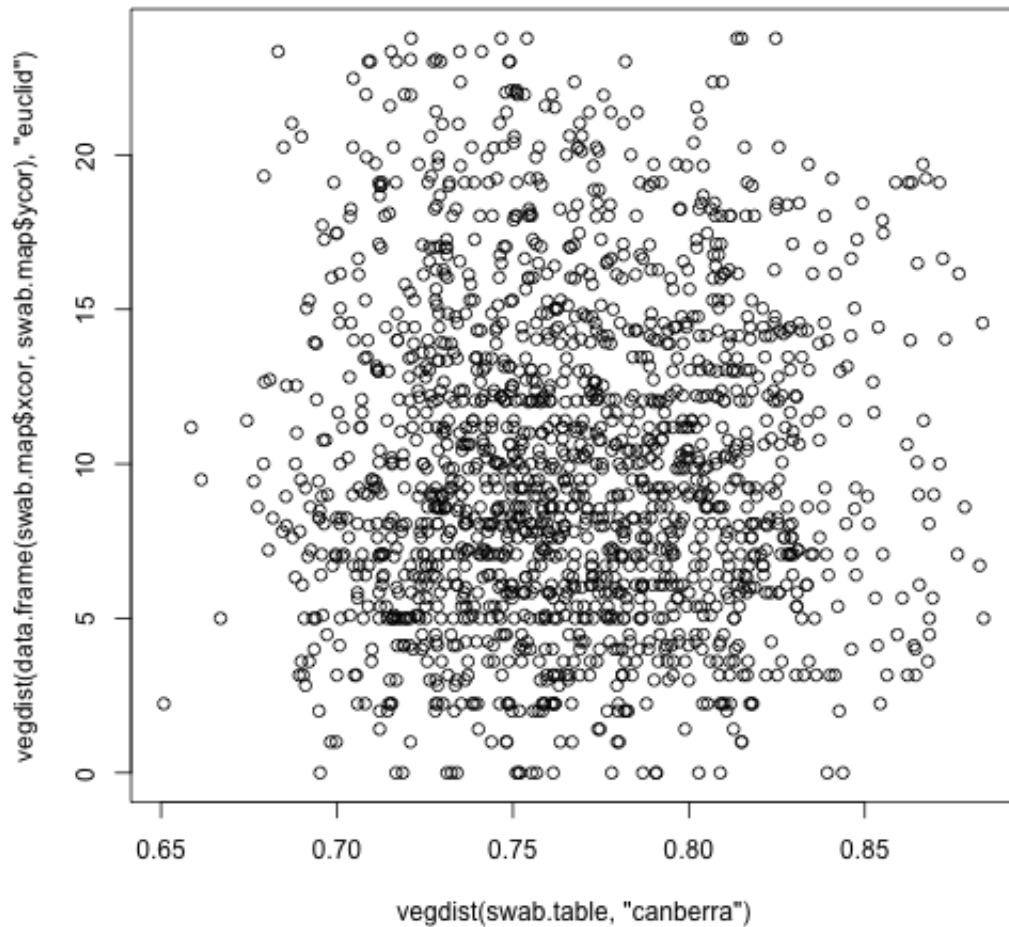
```
mantel(vegdist(st.w, "canberra"), vegdist(data.frame(sm.w$xcor,
sm.w$ycor),
"euclid"))
```

```
##
## Mantel statistic based on Pearson's product-moment correlation
##
## Call:
## mantel(xdis = vegdist(st.w, "canberra"), ydis =
vegdist(data.frame(sm.w$xcor,      sm.w$ycor), "euclid"))
##
## Mantel statistic r: -0.0896
##      Significance: 0.93
##
## Upper quantiles of permutations (null model):
##      90%      95%  97.5%    99%
## 0.121 0.177 0.223 0.288
##
## Based on 999 permutations
```

```
mantel(vegdist(swab.table, "canberra"),
vegdist(data.frame(swab.map$xcor, swab.map$ycor),
"euclid"))
```

```
##
## Mantel statistic based on Pearson's product-moment correlation
##
## Call:
## mantel(xdis = vegdist(swab.table, "canberra"), ydis =
vegdist(data.frame(swab.map$xcor,      swab.map$ycor), "euclid"))
##
## Mantel statistic r: -0.00428
##      Significance: 0.54
##
## Upper quantiles of permutations (null model):
##      90%      95%  97.5%    99%
## 0.0731 0.0922 0.1086 0.1317
##
## Based on 999 permutations
```

```
plot(vegdist(swab.table, "canberra"),
vegdist(data.frame(swab.map$xcor, swab.map$ycor),
"euclid"))
```



```
cor.test(vegdist(swab.table, "canberra"),
vegdist(data.frame(swab.map$xcor,
swab.map$ycor), "euclid"))
```

```
##
## Pearson's product-moment correlation
##
## data: vegdist(swab.table, "canberra") and
vegdist(data.frame(swab.map$xcor, swab.map$ycor), "euclid")
## t = -0.174, df = 1651, p-value = 0.8619
## alternative hypothesis: true correlation is not equal to 0
## 95 percent confidence interval:
## -0.05249 0.04394
## sample estimates:
## cor
## -0.004283
```

```
# Mantel statistic r: 0.01047 Significance: 0.399
```

This is never significant, which indicates that surface type makes more difference than location around the room (eg, proximity to door, window, ...).

And just to check, does it matter whether the walls were sampled high or low?

```
sm.w$location2 <- factor(sm.w$location2)
adonis(vegdist(st.w, "canberra") ~ sm.w$location2)
```

```
##
## Call:
## adonis(formula = vegdist(st.w, "canberra") ~ sm.w$location2)
##
## Terms added sequentially (first to last)
##
##              Df SumsOfSqs MeanSqs F.Model    R2 Pr(>F)
## sm.w$location2  1         0.31   0.308    1.1 0.078  0.18
## Residuals      13         3.65   0.281
## Total          14         3.95
```

Nope.

Make stacked barchart of OTU relative abundances. This takes the taxo table created above.

```
t.phylum <- aggregate(t(swab.table), by = list(taxo$phylum), sum)
t.class <- aggregate(t(swab.table), by = list(taxo$class), sum)
t.order <- aggregate(t(swab.table), by = list(taxo$order), sum)
```

Then order everything for visual greatness.

```
#### phylum 1% cutoff
row.names(t.phylum) <- t.phylum$Group.1
t.phylum <- t.phylum[, -1]
t.phylum <- t.phylum[order(rowSums(t.phylum)), order(swab.map$type)]
nrows <- nrow(t.phylum)
nshow <- 7
nkeep <- nrows-nshow+1
t.phylum <- rbind(colSums(t.phylum[1:nkeep-1, ]),
t.phylum[nkeep:nrows, ])
cols.8 <- c('#D73027', '#FC8D59', '#FEE090', '#FFFFBF', '#E0F3F8',
'#91BFDB', '#9475B4', '#969A97')

# class 1% cutoff
row.names(t.class) <- t.class$Group.1
t.class <- t.class[, -1]
t.class <- t.class[order(rowSums(t.class)), order(swab.map$type)]
nrows <- nrow(t.class)
nshow <- 10
nkeep <- nrows-nshow+1
t.class <- rbind(colSums(t.class[1:nkeep-1, ]), t.class[nkeep:nrows,
])
t.class <- cbind(t.class[, 1:14][, rev(order(t.phylum[8, 1:14]))],
t.class[, 15:29][, rev(order(t.phylum[8, 15:29]))],
t.class[, 30:43][, rev(order(t.phylum[8, 30:43]))],
t.class[, 44:58][, rev(order(t.phylum[8, 44:58]))])
)
cols.11 <- c('#A50026', '#D73027', '#F46D43', '#FDAE61', '#FEE090',
'#E0F3F8', '#ABD9E9', '#74ADD1', '#4575B4', '#313695',
'#969A97')

# order
row.names(t.order) <- t.order$Group.1
t.order <- t.order[, -1]
t.order <- t.order[order(rowSums(t.order)), order(swab.map$type)]
nrows <- nrow(t.order)
nshow <- 10
nkeep <- nrows-nshow+1
t.order <- rbind(colSums(t.order[1:nkeep-1, ]), t.order[nkeep:nrows,
])
t.order <- cbind(t.order[, 1:14][, rev(order(t.phylum[8, 1:14]))],
t.order[, 15:29][, rev(order(t.phylum[8, 15:29]))],
t.order[, 30:43][, rev(order(t.phylum[8, 30:43]))],
t.order[, 44:58][, rev(order(t.phylum[8, 44:58]))])
)
cols.11 <- c('#A50026', '#D73027', '#F46D43', '#FDAE61', '#FEE090',
'#E0F3F8', '#ABD9E9', '#74ADD1', '#4575B4', '#313695',
'#969A97')
# create index for bar mid-points.
bar <- barplot(as.matrix(t.order/4000), col=rev(cols.11))
```

```
#### this had to be run las to maintain phylum order in other charts
t.phylum <- cbind(t.phylum[, 1:14][, rev(order(t.phylum[8, 1:14]))],
                    t.phylum[, 15:29][, rev(order(t.phylum[8, 15:29]))],
                    t.phylum[, 30:43][, rev(order(t.phylum[8,
30:43]))]),
                    t.phylum[, 44:58][, rev(order(t.phylum[8, 44:58]))])

ph.names <- row.names(t.phylum)
ph.names[1] <- 'other (<1%)'
cl.names <- row.names(t.class)
cl.names[1] <- 'other (<1%)'
or.names <- row.names(t.order)
or.names[1] <- 'other (<4%)'
```

Figure 3. Taxonomy bar chart.

```
layout(matrix(c(1, 2, 3, 7, 1, 2, 3, 7, 4, 5, 6, 7), 4, 3), heights =
c(1, 1,
  1, 0.1))
par(mar = c(1, 4, 2, 1))
barplot(as.matrix(t.phylum/4000), col = rev(cols.8), border =
rev(cols.8), xaxt = "n",
  las = 1)
mtext("Phylum", font = 2, col = "gray20")
segments(c(mean(bar[c(14, 15)]), mean(bar[c(29, 30)]), mean(bar[c(43,
44)])),
  c(0, 0, 0), c(mean(bar[c(14, 15)]), mean(bar[c(29, 30)]),
mean(bar[c(43,
44)])), c(1, 1, 1))

mtext(c("chairs", "desks", "floors", "walls"), side = 1, at =
c(mean(bar[c(1,
14)]), mean(bar[c(15, 29)]), mean(bar[c(30, 43)]), mean(bar[c(44,
58)])),
  font = 2, col = "gray20")
barplot(as.matrix(t.class/4000), col = rev(cols.11), border =
rev(cols.11),
  xaxt = "n", las = 1)
mtext("Class", font = 2, col = "gray20")
segments(c(mean(bar[c(14, 15)]), mean(bar[c(29, 30)]), mean(bar[c(43,
44)])),
  c(0, 0, 0), c(mean(bar[c(14, 15)]), mean(bar[c(29, 30)]),
mean(bar[c(43,
44)])), c(1, 1, 1))
mtext(c("chairs", "desks", "floors", "walls"), side = 1, at =
c(mean(bar[c(1,
14)]), mean(bar[c(15, 29)]), mean(bar[c(30, 43)]), mean(bar[c(44,
58)])),
  font = 2, col = "gray20")
barplot(as.matrix(t.order/4000), col = rev(cols.11), border =
rev(cols.11),
  xaxt = "n", las = 1)
mtext("Order", font = 2, col = "gray20")
segments(c(mean(bar[c(14, 15)]), mean(bar[c(29, 30)]), mean(bar[c(43,
44)])),
  c(0, 0, 0), c(mean(bar[c(14, 15)]), mean(bar[c(29, 30)]),
mean(bar[c(43,
```

```

44)])), c(1, 1, 1))
par(xpd = TRUE)
mtext(c("chairs", "desks", "floors", "walls"), side = 1, at =
c(mean(bar[c(1,
14)]), mean(bar[c(15, 29)]), mean(bar[c(30, 43)]), mean(bar[c(44,
58)])),
font = 2, col = "gray20")
par(mar = c(0, 0, 0, 0))
plot(1, 1, type = "n", bty = "n", xlab = "", ylab = "", xaxt = "n",
yaxt = "n")
legend("left", legend = c(rev(ph.names)), pt.bg = cols.8, col =
cols.8, bty = "n",
pch = 22, cex = 1.5, pt.cex = 2.5)
plot(1, 1, type = "n", bty = "n", xlab = "", ylab = "", xaxt = "n",
yaxt = "n")
legend("left", legend = c(rev(cl.names)), pt.bg = cols.11, col =
cols.11, bty = "n",
pch = 22, cex = 1.5, pt.cex = 2.5)
plot(1, 1, type = "n", bty = "n", xlab = "", ylab = "", xaxt = "n",
yaxt = "n")
legend("left", legend = c(rev(or.names)), pt.bg = cols.11, col =
cols.11, bty = "n",
pch = 22, cex = 1.5, pt.cex = 2.5)

```

