

eDNA

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Analysis of community sequence data to test whether the structure and composition of bacterial communities is affected by extracellular DNA

Setup Work Environment

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

## [1] "/Users/mmuscarella/GitHub/eDNA/code"
setwd("~/GitHub/eDNA/code")

# Load dependencies
require("vegan")

## Loading required package: vegan
## Loading required package: permute
## Loading required package: lattice
## This is vegan 2.4-1
require("plyr")

## Loading required package: plyr
require("car")

## Loading required package: car
require("grid")

## Loading required package: grid
require("png")

## Loading required package: png
require("ape")

## Loading required package: ape
require("picante")

## Loading required package: picante
## Loading required package: nlme
require("ade4")

## Loading required package: ade4
```

```
##
## Attaching package: 'ade4'

## The following object is masked from 'package:vegan':
##
##      cca

#require("phytools")
require("phangorn")

## Loading required package: phangorn
##
## Attaching package: 'phangorn'
## The following objects are masked from 'package:vegan':
##
##      diversity, treedist

# Source code functions
source("../bin/DiversityFunctions.R")
source("../bin/MothurTools.R")

## Loading required package: reshape
##
## Attaching package: 'reshape'
## The following objects are masked from 'package:plyr':
##
##      rename, round_any

source("../bin/phylodiversity2.R")

# Small custom functions
sem <- function(x, ...){sd(x, na.rm = TRUE)/sqrt(length(na.omit(x)))}
ttest <- function(reg, coefnum, val){
  co <- coef(summary(reg))
  tstat <- (co[coefnum,1]-val)/co[coefnum,2]
  pstat <- 2 * pt(abs(tstat), reg$df.residual, lower.tail = FALSE)
  return(list = c(t = tstat, df = reg$df.residual, p = pstat))
}

# Save Default Plot Settings
opar <- par(no.readonly = TRUE) # Saves plot defaults

# Run All: Select if all section are to be re-run
run.all <- FALSE
```

Phylogenetic Diversity Notes

A multi-fasta file was generated with representative sequences for each OTU. Representatives were picked based on the most abundant unique sequence in each OTU. FastTree was used to construct a phylogenetic tree. Note: This must be done before performing the following analyses. The following parameters were used:

Note: eDNA.bac.final.0.03.rep.fasta is larger than the github limits. Please obtain the file from SDA or

Note: The names must be renamed so that they match the OTU names:

```
> python ./bin/name_change.py "./eDNA.bac.final.0.03.rep.fasta" "./eDNA.bac.final.0.03.rep.rename.fasta"
> FastTree -gtr -nt -gamma -fastest eDNA.bac.final.0.03.rep.rename.fasta > eDNA.bac.rename.tree
```

Output:

```
ML-NNI round 14: LogLk = -5469747.038 NNIs 18759 max delta 6.95 Time 4888.39 (final)ax delta 6.954)
Optimize all lengths: LogLk = -5469663.971 Time 5006.96
Gamma(20) LogLk = -5470984.148 alpha = 1.037 rescaling lengths by 1.659
Total time: 5529.61 seconds Unique: 174292/174292 Bad splits: 903/174289 Worst delta-LogLk 10.311
```

Data Input Section

Define Inputs

```
# Design      = general design file for experiment
# shared     = OTU table from mothur with sequence similarity clustering
# tax        = Taxonomy for 97% similarity OTUs
design         <- "../data/eDNA_Design.txt"
shared        <- "../mothur/output/eDNA.bac.final.shared"
tax           <- "../mothur/output/eDNA.bac.final.0.03.taxonomy"
```

Import Design

```
design <- read.delim(design, header=T, row.names=1)
```

Import Shared, Taxonomy, and Phylogeny Files

```
OTU <- read.otu(shared = shared, cutoff = "0.03")
OTU.tax <- read.tax(taxonomy = tax, format = "rdp")
OTU.tre <- read.tree("../mothur/output/eDNA.bac.rename.tree")
```

Remove OTUs with less than two occurrences across all sites

```
OTU <- OTU[, which(colSums(OTU) >= 2)]
```

Calculate Coverage Stats

```
cov.seqs <- count.groups(OTU)
cov.mean <- mean(cov.seqs) # 222701
cov.sem <- sem(cov.seqs) # 9560
cov.min <- min(cov.seqs) # 31,475
total.seqs <- sum(cov.seqs) # 12,916,632

# Good's coverage
```

```

goods.c <- function(x = ""){
  1 - (apply(OTU, 1, function(x){sum(x == 1)})) / rowSums(x))
}

goods.c.eDNA <- goods.c(OTU)
mean.good.c <- mean(goods.c.eDNA) # 0.99
min.good.c <- min(goods.c.eDNA) # 0.98

```

Alpha Diversity

Calculate Alpha diversity using Resampling

```

# Mario's resampling code to estimate alpha diversity (used if run.all = T)

if (run.all == TRUE){
  rich <- round(richness.iter(input = OTU, size = 30000,
                             iters = 100, shared = "FALSE"), 3)
  even <- round(evenness.iter(input = OTU, size = 30000,
                              iters = 100, shared = "FALSE",
                              method = "simp_even"), 3)

  rare <- rarefy(OTU, 30000, se = FALSE, MARGIN = 1)

  # Write output to files
  write.table(rich, "../data/rich.txt", sep = "\t",
              col.names = T, row.names = T)
  write.table(even, "../data/even.txt", sep = "\t",
              col.names = T, row.names = T)
}

# Read in alpha diversity files from above
rich2 <- read.table("../data/rich.txt", sep = "\t")
even2 <- read.table("../data/even.txt", sep = "\t")

# Merge data to design and calculate mean and sem per sample
rich.data <- merge(design, rich2, by = "row.names")
row.names(rich.data) <- rich.data$Row.names
rich.data <- rich.data[sort(row.names(rich.data)), ]
rich.mean <- round(apply(rich.data[8:(7 + dim(rich2)[2])], 1, mean, na.rm = TRUE), 3)
rich.sem <- round(apply(rich.data[8:(7 + dim(rich2)[2])], 1, sem, na.rm = TRUE), 3)

even.data <- merge(design, even2, by = "row.names")
row.names(even.data) <- even.data$Row.names
even.data <- even.data[sort(row.names(even.data)), ]
even.mean <- round(apply(even.data[8:(7 + dim(even2)[2])], 1, mean, na.rm = TRUE), 3)
even.sem <- round(apply(even.data[8:(7 + dim(even2)[2])], 1, sem, na.rm = TRUE), 4)

# Make new dataframe merging design file and mean diversity
eDNA.div <- data.frame(design[sort(row.names(design)), ], rich.mean, even.mean)

# Take averages of technical reps

```

```
rich.rep.ave <- ddply(eDNA.div, .(env, treat, qPCR.match), summarize, rich = mean(rich.mean))
even.rep.ave <- ddply(eDNA.div, .(env, treat, qPCR.match), summarize, even = mean(even.mean))

# Reshape data
rich.2 <- reshape(rich.rep.ave[,1:4], timevar = "treat",
                  idvar = c("qPCR.match", "env"), direction = "wide")

even.2 <- reshape(even.rep.ave[,1:4], timevar = "treat",
                  idvar = c("qPCR.match", "env"), direction = "wide")
```

Richness: differences among sites?

```
rich.anova <- aov(rich.2$rich.C ~ rich.2$env)
summary(rich.anova)

##              Df    Sum Sq Mean Sq F value    Pr(>F)
## rich.2$env     3 34630542 11543514   14.95 5.08e-05 ***
## Residuals    17 13122506   771912
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

TukeyHSD(rich.anova)

##      Tukey multiple comparisons of means
##      95% family-wise confidence level
##
## Fit: aov(formula = rich.2$rich.C ~ rich.2$env)
##
## $`rich.2$env`
##              diff            lwr            upr            p adj
## sed-feces    2700.5429   1088.4587   4312.6271   0.0009436
## soil-feces    1379.7157   -295.6113   3055.0428   0.1276911
## water-feces   -447.6879  -2059.7721   1164.3963   0.8582474
## soil-sed     -1320.8272  -2833.0962    191.4418   0.0989233
## water-sed    -3148.2308  -4590.1228  -1706.3389   0.0000519
## water-soil   -1827.4037  -3339.6727   -315.1347   0.0151328
```

Richness: does eDNA “inflate” diversity?

```
# Take averages of technical reps
rich.rep.ave <- ddply(eDNA.div, .(env, treat, qPCR.match),
                      summarize, rich = mean(rich.mean))

# Reshape data
rich.2 <- reshape(rich.rep.ave[,1:4], timevar = "treat",
                  idvar = c("qPCR.match", "env"), direction = "wide")

# Calculate means +/- SEM of control and treated richness
rich.2.ag.mean.control <- aggregate(rich.C ~ env, rich.2, mean)
rich.2.ag.sem.control <- aggregate(rich.C ~ env, rich.2, sem)
rich.2.ag.mean.treatment <- aggregate(rich.E ~ env, rich.2, mean)
```

```

rich.2.ag.sem.treatment <- aggregate(rich.E ~ env, rich.2, sem)

# Calculate ratios
rich.2$rich.ratio <- rich.2$rich.E / rich.2$rich.C

# Richness table
rich.2.ag.mean <- aggregate(rich.ratio ~ env, rich.2, mean)
rich.2.ag.sem <- aggregate(rich.ratio ~ env, rich.2, sem)
rich.2.ag.95 <- aggregate(rich.ratio ~ env, rich.2,
                          FUN = function(x) t.test(x)$conf.int[1:2])
rich.table <- data.frame(rich.2.ag.mean$env, rich.2.ag.mean$rich.ratio,
                          rich.2.ag.sem$rich.ratio, rich.2.ag.95$rich.ratio)
colnames(rich.table) <- c("env", "mean", "sem", "LCI", "UCI")

# Regression: richness ratio vs. proportion eDNA
eDNA.prop <- read.table("../data/eDNA.prop.txt", sep = "\t", header = T)

eDNA.prop.sub <- eDNA.prop[eDNA.prop$sample.number %in% rich.2$qPCR.match, ]
eDNA.prop.sub2 <- eDNA.prop.sub[order(eDNA.prop.sub$sample.number), ]

rich.reg <- lm(rich.2[order(rich.2$qPCR.match), ]$rich.ratio ~ eDNA.prop.sub2$env + eDNA.prop.sub2$prop)

rich.reg <- lm(rich.2[order(rich.2$qPCR.match), ]$rich.ratio ~ eDNA.prop.sub2$prop)

summary(rich.reg)

##
## Call:
## lm(formula = rich.2[order(rich.2$qPCR.match), ]$rich.ratio ~
##     eDNA.prop.sub2$prop)
##
## Residuals:
##      Min       1Q   Median       3Q      Max
## -0.30939 -0.06784 -0.00313  0.07611  0.38702
##
## Coefficients:
##              Estimate Std. Error t value Pr(>|t|)
## (Intercept)      0.96384     0.07803  12.352 1.59e-10 ***
## eDNA.prop.sub2$prop -0.03855     0.17682  -0.218    0.83
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 0.1666 on 19 degrees of freedom
## Multiple R-squared:  0.002496, Adjusted R-squared:  -0.05
## F-statistic: 0.04754 on 1 and 19 DF, p-value: 0.8297

Anova(rich.reg)

## Anova Table (Type II tests)
##
## Response: rich.2[order(rich.2$qPCR.match), ]$rich.ratio
##              Sum Sq Df F value Pr(>F)
## eDNA.prop.sub2$prop 0.00132  1  0.0475 0.8297
## Residuals          0.52712 19

```

```
ttest(rich.reg, 1, 1)

##           t           df           p
## -0.4633891 19.0000000  0.6483497

ttest(rich.reg, 2, 0)

##           t           df           p
## -0.2180371 19.0000000  0.8297248

# MEM: what is the Anova() function? = prints the anova like table for the model
#                                     Anova uses type II sums of squares
# answer => no correlation between proportion eDNA and richness ratio.
# maybe a little in sediment
```

Evenness: differences among sites?

```
even.anova <- aov(even.2$even.C ~even.2$env)
summary(even.anova)

##           Df Sum Sq Mean Sq F value    Pr(>F)
## even.2$env    3  0.1247  0.04158    6.497 0.00397 **
## Residuals   17  0.1088  0.00640
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

TukeyHSD(even.anova)

##      Tukey multiple comparisons of means
##      95% family-wise confidence level
##
## Fit: aov(formula = even.2$even.C ~ even.2$env)
##
## $`even.2$env`
##           diff           lwr           upr           p adj
## sed-feces  -0.17916667 -0.32595852 -0.03237482 0.0140884
## soil-feces -0.12695000 -0.27950056  0.02560056 0.1222474
## water-feces -0.21808333 -0.36487518 -0.07129148 0.0029099
## soil-sed    0.05221667 -0.08548629  0.18991963 0.7071690
## water-sed   -0.03891667 -0.17021129  0.09237795 0.8335247
## water-soil  -0.09113333 -0.22883629  0.04656963 0.2723574
```

Evenness: does eDNA “inflate” diversity?

```
# Take averages of technical reps
even.rep.ave <- ddpby(eDNA.div, .(env, treat, qPCR.match),
                     summarize, even = mean(even.mean))

# Reshape data
even.2 <- reshape(even.rep.ave[,1:4], timevar = "treat",
                  idvar = c("qPCR.match", "env"), direction = "wide")

# Calcualte means +/- SEM of control and treated richness
```

```

even.2.ag.mean.control <- aggregate(even.C ~ env, even.2, mean)
even.2.ag.sem.control <- aggregate(even.C ~ env, even.2, sem)
even.2.ag.mean.treatment <- aggregate(even.E ~ env, even.2, mean)
even.2.ag.sem.treatment <- aggregate(even.E ~ env, even.2, sem)

# Calculate ratios
even.2$even.ratio <- even.2$even.E / even.2$even.C

# Evennes table
even.2.ag.mean <- aggregate(even.ratio ~ env, even.2, mean)
even.2.ag.sem <- aggregate(even.ratio ~ env, even.2, sem)
even.2.ag.95 <- aggregate(even.ratio ~ env, even.2,
  FUN = function(x) t.test(x)$conf.int[1:2])
even.table <- data.frame(even.2.ag.mean$env, even.2.ag.mean$even.ratio,
  even.2.ag.sem$even.ratio, even.2.ag.95$even.ratio)
colnames(even.table) <- c("env", "mean", "sem", "LCI", "UCI")

# Regression: evennes ratio vs. proportion eDNA
even.reg <- lm(even.2[order(even.2$qPCR.match), ]$even.ratio ~ eDNA.prop.sub2$prop + eDNA.prop.sub2$env)
even.reg <- lm(even.2[order(even.2$qPCR.match), ]$even.ratio ~ eDNA.prop.sub2$prop)

summary(even.reg)

##
## Call:
## lm(formula = even.2[order(even.2$qPCR.match), ]$even.ratio ~
##     eDNA.prop.sub2$prop)
##
## Residuals:
##      Min       1Q   Median       3Q      Max
## -0.15051 -0.04483 -0.02383  0.02394  0.37634
##
## Coefficients:
##              Estimate Std. Error t value Pr(>|t|)
## (Intercept)      0.98745     0.05251   18.80 9.73e-14 ***
## eDNA.prop.sub2$prop 0.11423     0.11899    0.96  0.349
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 0.1121 on 19 degrees of freedom
## Multiple R-squared:  0.04626,    Adjusted R-squared:  -0.003934
## F-statistic: 0.9216 on 1 and 19 DF,  p-value: 0.3491

ttest(even.reg, 1, 1)

##           t           df           p
## -0.2389974 19.0000000  0.8136666

ttest(even.reg, 2, 0)

##           t           df           p
##  0.9600096 19.0000000  0.3491089

```



```
# answer => no correlation between proportion eDNA and evenness ratio.
```

Calculate Phylogenetic Alpha Diversity

```
# Test if all OTUs are in tree
sum(colnames(OTU) %in% OTU.tre$tip.label) == length(colnames(OTU) %in% OTU.tre$tip.label)

## [1] TRUE

# Root Tree if Needed
is.rooted(OTU.tre)

## [1] FALSE

OTU.tre.rooted <- midpoint(OTU.tre)

if (run.all == TRUE){
  OTU.2 <- rrarefy(OTU, 30000)
  OTU.2 <- OTU.2[,which(colSums(OTU.2) > 0)]
  OTU.tre.2 <- prune.sample(OTU.2, OTU.tre.rooted)

  # Calculate Faith's D
  eDNA.pd <- pd(OTU.2, OTU.tre.2, include.root = F)

  # Write output to files
  write.table(eDNA.pd, "../data/phylo.txt", sep = "\t",
              col.names = T, row.names = T)
}

# Read in alpha diversity files from above
eDNA.pd2 <- read.table("../data/phylo.txt", sep = "\t")

# Make new dataframe merging design file and phylo diversity
eDNA.phylo <- data.frame(design[sort(row.names(design)), ],
                        pd = eDNA.pd2$PD)

# Take averages of technical reps
pd.rep.ave <- ddply(eDNA.phylo, .(env, treat, qPCR.match),
                   summarize, phylo = mean(pd))

# Reshape data
phylo.2 <- reshape(pd.rep.ave[,1:4], timevar = "treat",
                  idvar = c("qPCR.match", "env"), direction = "wide")

# Calculate means +/- SEM of control and treated richness
phylo.2.ag.mean.control <- aggregate(phylo.C ~ env, phylo.2, mean)
phylo.2.ag.sem.control <- aggregate(phylo.C ~ env, phylo.2, sem)
phylo.2.ag.mean.treatment <- aggregate(phylo.E ~ env, phylo.2, mean)
phylo.2.ag.sem.treatment <- aggregate(phylo.E ~ env, phylo.2, sem)
```

Phylogenetic: differences among sites?

```
faith.anova <- aov(phylo.2$phylo.C ~ phylo.2$env)
summary(faith.anova)

##              Df Sum Sq Mean Sq F value    Pr(>F)
## phylo.2$env   3 693939  231313    21.92 4.46e-06 ***
## Residuals    17 179413   10554
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

TukeyHSD(faith.anova)

##      Tukey multiple comparisons of means
##      95% family-wise confidence level
##
## Fit: aov(formula = phylo.2$phylo.C ~ phylo.2$env)
##
## $`phylo.2$env`
##              diff             lwr             upr             p adj
## sed-feces      492.7041      304.20628      681.20192 0.0000054
## soil-feces      262.8600       66.96729      458.75265 0.0068654
## water-feces     126.1452      -62.35265      314.64299 0.2638046
## soil-sed       -229.8441     -406.67076     -53.01750 0.0088154
## water-sed      -366.5589     -535.15651    -197.96136 0.0000546
## water-soil     -136.7148     -313.54143      40.11183 0.1637189
```

Phylogenetic Diversity: does eDNA “inflate” diversity?

```
# Calculate ratios
phylo.2$phylo.ratio <- phylo.2$phylo.E / phylo.2$phylo.C

# Richness table
phylo.2.ag.mean <- aggregate(phylo.ratio ~ env, phylo.2, mean)
phylo.2.ag.sem <- aggregate(phylo.ratio ~ env, phylo.2, sem)
phylo.2.ag.95 <- aggregate(phylo.ratio ~ env, phylo.2,
                           FUN = function(x) t.test(x)$conf.int[1:2])
phylo.table <- data.frame(phylo.2.ag.mean$env, phylo.2.ag.mean$phylo.ratio,
                          phylo.2.ag.sem$phylo.ratio, phylo.2.ag.95$phylo.ratio)
colnames(phylo.table) <- c("env", "mean", "sem", "LCI", "UCI")

# Regression: PD ratio vs. proportion eDNA
phylo.reg <- lm(phylo.2[order(phylo.2$qPCR.match), ]$phylo.ratio ~ eDNA.prop.sub2$prop * eDNA.prop.sub2$prop)

phylo.reg <- lm(phylo.2[order(phylo.2$qPCR.match), ]$phylo.ratio ~ eDNA.prop.sub2$prop)

summary(phylo.reg)

##
## Call:
## lm(formula = phylo.2[order(phylo.2$qPCR.match), ]$phylo.ratio ~
```

```
##      eDNA.prop.sub2$prop)
##
## Residuals:
##      Min        1Q      Median        3Q        Max
## -0.22163 -0.04803 -0.01018  0.05867  0.25648
##
## Coefficients:
##              Estimate Std. Error t value Pr(>|t|)
## (Intercept)      0.96565     0.05670   17.030 5.79e-13 ***
## eDNA.prop.sub2$prop -0.04337     0.12849   -0.338   0.739
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 0.121 on 19 degrees of freedom
## Multiple R-squared:  0.005961, Adjusted R-squared:  -0.04636
## F-statistic: 0.1139 on 1 and 19 DF, p-value: 0.7394

ttest(phylo.reg, 1, 1)

##           t           df           p
## -0.6057934 19.0000000  0.5518208
# answer => no correlation between proportion eDNA and evenness ratio.
```

Alpha Diversity Plots

Richness plot

```
png(filename="../figures/ratio.richness.png",
      width = 800, height = 600, res = 96*2)

par(mar=c(3, 4, 1, 1) + 0.1)

rich.plot <- plot(rich.table$mean, ylim = c(0.5, 1.5),
                  xlim = c(0.5, 4.5), pch = 22, bg = "white", lwd = 2,
                  cex = 3, yaxt = "n", xaxt = "n", cex.lab = 2, cex.axis = 1.5,
                  las = 1, ylab = "", xlab = "")
box(lwd = 2)
abline(h = 1, lty = 3, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = rich.table$mean, y1 = rich.table$LCI, angle = 90,
       length = 0.1, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = rich.table$mean, y1 = rich.table$UCI, angle = 90,
       length=0.1, lwd = 2)
points(c(1:4), rich.table$mean,pch = 22, cex = 3, bg = "gray90", lwd = 2)

mtext(expression('Richness Ratio'), side = 2,
       outer = F, cex = 1.5, line = 2.5, adj = 0.5)

# Major Axes
axis(side = 2, lwd.ticks = 2, cex.axis = 1, las = 1,
     labels = T, at = c(0.5, 1, 1.5))
axis(side = 4, lwd.ticks = 2, cex.axis = 1, las = 1,
     labels = F, at = c(0.5, 1, 1.5))
```

```

axis(side = 1, lwd.ticks = 2, cex.axis = 0.9, las = 1,
     labels = c("Gut", "Sediment", "Soil", "Water"), at = c(1, 2, 3, 4))
axis(side = 3, lwd.ticks = 2, cex.axis = 1.5, las = 1,
     at = c(1, 2, 3, 4), labels = F)

# Minor ticks
axis(side = 2, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.5, 1, 1.5))
axis(side = 4, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.5, 1, 1.5))
axis(side = 1, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))
axis(side = 3, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))

# Close Plot Device
dev.off()

## pdf
## 2

graphics.off()

# Show Plot
img <- readPNG("../figures/ratio.richness.png")
grid.raster(img)

```

Evenness plot

```

png(filename="../figures/ratio.evenness.png",
     width = 800, height = 600, res = 96*2)

par(mar=c(3, 4, 1, 1) + 0.1)

even.plot <- plot(even.table$mean, ylim = c(0.5, 1.5),
                 xlim = c(0.5, 4.5), pch = 22, bg = "white", lwd = 2,
                 cex = 3, yaxt = "n", xaxt = "n", cex.lab = 2, cex.axis = 1.5,
                 las = 1, ylab = "", xlab = "")
box(lwd = 2)
abline(h = 1, lty = 3, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = even.table$mean, y1 = even.table$LCI, angle = 90,
       length = 0.1, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = even.table$mean, y1 = even.table$UCI, angle = 90,
       length=0.1, lwd = 2)
points(c(1:4), even.table$mean,pch = 22, cex = 3, bg = "gray90", lwd = 2)

mtext(expression('Evenness Ratio'), side = 2,
       outer = F, cex = 1.5, line = 2.5, adj = 0.5)
# Major Axes
axis(side = 2, lwd.ticks = 2, cex.axis = 1, las = 1,
     labels = T, at = c(0.5, 1, 1.5))

axis(side = 4, lwd.ticks = 2, cex.axis = 1, las = 1,
     labels = F, at = c(0.5, 1, 1.5))

axis(side = 1, lwd.ticks = 2, cex.axis = 0.9, las = 1,
     labels = c("Gut", "Sediment", "Soil", "Water"), at = c(1, 2, 3, 4))

```

```

axis(side = 3, lwd.ticks = 2, cex.axis = 1.5, las = 1,
     at = c(1, 2, 3, 4), labels = F)

# Minor ticks
axis(side = 2, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.5, 1, 1.5))
axis(side = 4, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.5, 1, 1.5))
axis(side = 1, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))
axis(side = 3, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))

# Close Plot Device
dev.off()

## pdf
## 2

graphics.off()

# Show Plot
img <- readPNG("../figures/ratio.evenness.png")
grid.raster(img)

```

Faith's D plot

```

png(filename="../figures/ratio.phylo.png",
     width = 800, height = 600, res = 96*2)

par(mar=c(3, 4, 1, 1) + 0.1)

phylo.plot <- plot(phylo.table$mean, ylim = c(0.5, 1.5),
                  xlim = c(0.5, 4.5), pch = 22, bg = "white", lwd = 2,
                  cex = 3, yaxt = "n", xaxt = "n", cex.lab = 2, cex.axis = 1.5,
                  las = 1, ylab = "", xlab = "")
box(lwd = 2)
abline(h = 1, lty = 3, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = phylo.table$mean, y1 = phylo.table$LCI, angle = 90,
       length = 0.1, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = phylo.table$mean, y1 = phylo.table$UCI, angle = 90,
       length=0.1, lwd = 2)
points(c(1:4), phylo.table$mean, pch = 22, cex = 3, bg = "gray90", lwd = 2)

mtext(expression('Faiths D Ratio'), side = 2,
       outer = F, cex = 1.5, line = 2.5, adj = 0.5)
# Major Axes
axis(side = 2, lwd.ticks = 2, cex.axis = 1, las = 1,
     labels = T, at = c(0.5, 1, 1.5))

axis(side = 4, lwd.ticks = 2, cex.axis = 1, las = 1,
     labels = F, at = c(0.5, 1, 1.5))

axis(side = 1, lwd.ticks = 2, cex.axis = 0.9, las = 1,
     labels = c("Gut", "Sediment", "Soil", "Water"), at = c(1, 2, 3, 4))

```

```

axis(side = 3, lwd.ticks = 2, cex.axis = 1.5, las = 1,
     at = c(1, 2, 3, 4), labels = F)

# Minor ticks
axis(side = 2, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.5, 1, 1.5))
axis(side = 4, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.5, 1, 1.5))
axis(side = 1, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))
axis(side = 3, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))

# Close Plot Device
dev.off()

## pdf
## 2

graphics.off()

# Show Plot
img <- readPNG("../figures/ratio.phylo.png")
grid.raster(img)

```

Alpha Diversity Multi-Panel Plot

```

png(filename="../figures/alpha.ratios.png",
     width = 800, height = 1600, res = 96*2)

layout(matrix(c(1:3), byrow = T))
par(mar = c(0.5, 4, 0.5, 1), oma = c(3, 1, 1, 1))

# Richness Panel
rich.plot <- plot(rich.table$mean, ylim = c(0.5, 1.5),
                 xlim = c(0.5, 4.5), pch = 22, bg = "white", lwd = 2,
                 cex = 3, yaxt = "n", xaxt = "n", cex.lab = 2, cex.axis = 1.5,
                 las = 1, ylab = "", xlab = "")
box(lwd = 2)
abline(h = 1, lty = 3, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = rich.table$mean, y1 = rich.table$LCI, angle = 90,
       length = 0.1, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = rich.table$mean, y1 = rich.table$UCI, angle = 90,
       length=0.1, lwd = 2)
points(c(1:4), rich.table$mean, pch = 22, cex = 3, bg = "gray90", lwd = 2)

mtext(expression('Richness Ratio'), side = 2,
       outer = F, cex = 1.5, line = 3, adj = 0.5)

# Major Axes
axis(side = 2, lwd.ticks = 2, tck = -0.02, cex.axis = 1.5, las = 1,
     labels = T, at = c(0.5, 1, 1.5))
axis(side = 4, lwd.ticks = 2, cex.axis = 1, las = 1,
     labels = F, at = c(0.5, 1, 1.5))
axis(side = 1, lwd.ticks = 2, tck = -0.02, cex.axis = 0.9, las = 1,
     labels = F, at = c(1, 2, 3, 4))
axis(side = 3, lwd.ticks = 2, cex.axis = 1.5, las = 1,

```

```

    at = c(1, 2, 3, 4), labels = F)

# Minor ticks
axis(side = 2, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.5, 1, 1.5))
axis(side = 4, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.5, 1, 1.5))
axis(side = 1, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))
axis(side = 3, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))

# Evenness Panel
even.plot <- plot(even.table$mean, ylim = c(0.5, 1.5),
  xlim = c(0.5, 4.5), pch = 22, bg = "white", lwd = 2,
  cex = 3, yaxt = "n", xaxt = "n", cex.lab = 2, cex.axis = 1.5,
  las = 1, ylab = "", xlab = "")
  box(lwd = 2)
abline(h = 1, lty = 3, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = even.table$mean, y1 = even.table$LCI, angle = 90,
  length = 0.1, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = even.table$mean, y1 = even.table$UCI, angle = 90,
  length=0.1, lwd = 2)
points(c(1:4), even.table$mean,pch = 22, cex = 3, bg = "gray90", lwd = 2)

mtext(expression('Evenness Ratio'), side = 2,
  outer = F, cex = 1.5, line = 3, adj = 0.5)

# Major Axes
axis(side = 2, lwd.ticks = 2, cex.axis = 1.5, las = 1,
  labels = T, at = c(0.5, 1, 1.5))

axis(side = 4, lwd.ticks = 2, cex.axis = 1, las = 1,
  labels = F, at = c(0.5, 1, 1.5))

axis(side = 1, lwd.ticks = 2, tck = -0.02, cex.axis = 0.9, las = 1,
  labels = F, at = c(1, 2, 3, 4))

axis(side = 3, lwd.ticks = 2, tck = -0.02, cex.axis = 1.5, las = 1,
  at = c(1, 2, 3, 4), labels = F)

# Minor ticks
axis(side = 2, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.5, 1, 1.5))
axis(side = 4, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.5, 1, 1.5))
axis(side = 1, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))
axis(side = 3, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))

# Phylo Panel
phylo.plot <- plot(phylo.table$mean, ylim = c(0.5, 1.5),
  xlim = c(0.5, 4.5), pch = 22, bg = "white", lwd = 2,
  cex = 3, yaxt = "n", xaxt = "n", cex.lab = 2, cex.axis = 1.5,
  las = 1, ylab = "", xlab = "")
  box(lwd = 2)
abline(h = 1, lty = 3, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = phylo.table$mean, y1 = phylo.table$LCI, angle = 90,
  length = 0.1, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = phylo.table$mean, y1 = phylo.table$UCI, angle = 90,
  length=0.1, lwd = 2)

```

```

points(c(1:4), phylo.table$mean, pch = 22, cex = 3, bg = "gray90", lwd = 2)

mtext(expression('Faiths D Ratio'), side = 2,
        outer = F, cex = 1.5, line = 3, adj = 0.5)
# Major Axes
axis(side = 2, lwd.ticks = 2, cex.axis = 1.5, las = 1,
      labels = T, at = c(0.5, 1, 1.5))

axis(side = 4, lwd.ticks = 2, cex.axis = 1, las = 1,
      labels = F, at = c(0.5, 1, 1.5))

axis(side = 1, lwd.ticks = 2, cex.axis = 1.5, las = 1,
      labels = c("Gut", "Sediment", "Soil", "Water"), at = c(1, 2, 3, 4))

axis(side = 3, lwd.ticks = 2, tck = -0.02, cex.axis = 1.5, las = 1,
      at = c(1, 2, 3, 4), labels = F)

# Minor ticks
axis(side = 2, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.5, 1, 1.5))
axis(side = 4, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.5, 1, 1.5))
axis(side = 1, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))
axis(side = 3, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))

# Close Plot Device
dev.off()

## pdf
## 2

graphics.off()

# Show Plot
img <- readPNG("../figures/alpha.ratios.png")
grid.raster(img)

```

Beta diversity

Taxonomic Beta Diversity

```

# Make presence-absence matrix
OTU.PA <- (OTU > 0) * 1

# Make relative abundance matrix
OTU.REL <- OTU
for (i in 1:dim(OTU)[1]){
  OTU.REL[i,] <- OTU[i,]/sum(OTU[i,])
}

# Log-transform relative abundances
OTU.REL.log <- decostand(OTU, method="log")

eDNA.bc.dis <- vegdist(OTU.REL.log, method = "bray", binary = "FALSE")

```



```

eDNA.dis.mean <- mean(eDNA.bc.dis)

# Principal Coordinates Analysis (PCoA)
eDNA.PCoA <- cmdscale(eDNA.bc.dis, eig = TRUE, k = 3)
explainvar1 <- round(eDNA.PCoA$eig[1] / sum(eDNA.PCoA$eig), 3) * 100
explainvar2 <- round(eDNA.PCoA$eig[2] / sum(eDNA.PCoA$eig), 3) * 100
explainvar3 <- round(eDNA.PCoA$eig[3] / sum(eDNA.PCoA$eig), 3) * 100
sum.eig <- sum(explainvar1, explainvar2, explainvar3)

# OTU Scores
otu.scores <- t(cor(eDNA.PCoA$points, OTU.REL))
otu.scores <- as.matrix(otu.scores)[,1:2]
otu.scores <- otu.scores[abs(otu.scores[,1]) > 0.7 | abs(otu.scores[,2]) > 0.7,]

# Average BC Distance Between Treatments
eDNA.bc.dis.m <- as.matrix(eDNA.bc.dis)
all.equal(row.names(eDNA.div), rownames(eDNA.bc.dis.m))

## [1] TRUE

pair.div <- unique(eDNA.div$pair)
pair.dis <- rep(NA, length(pair.div))
for(i in 1:length(pair.div)){
  temp <- row.names(eDNA.div[eDNA.div$pair == pair.div[i], ])
  pair.dis[i] <- eDNA.bc.dis.m[temp[1], temp[2]]
}

mean(pair.dis)

## [1] 0.4230494

# Average BC Distance Between Replicates
reps <- names(which(table(eDNA.div$qPCR.match) == 4))
reps.dis <- rep(NA, length(reps))

for(i in 1:length(reps)){
  temp <- eDNA.div[eDNA.div$qPCR.match == reps[i] & eDNA.div$treat == "C", ]
  temp.names <- row.names(temp)
  reps.dis[i] <- eDNA.bc.dis.m[temp.names[1], temp.names[2]]
}

mean(reps.dis)

## [1] 0.4314466

t.test(pair.dis, reps.dis)

##
## Welch Two Sample t-test
##
## data: pair.dis and reps.dis
## t = -0.16259, df = 11.331, p-value = 0.8737
## alternative hypothesis: true difference in means is not equal to 0
## 95 percent confidence interval:
## -0.1216639 0.1048695
## sample estimates:
## mean of x mean of y

```

```
## 0.4230494 0.4314466
```

Phylogenetic Beta Diversity

```
phylo.dist <- read.delim("../mothur/output/eDNA.bac.tree.weighted.phylip.dist",  
                        skip = 1, row.names = 1, header = F, strip.white = T)
```

```
phylo.dist <- as.dist(phylo.dist)
```

```
# Principal Coordinates Analysis (PCoA)
```

```
eDNA.phylo.PCoA <- cmdscale(phylo.dist, eig = TRUE, k = 3)
```

```
explainvar1 <- round(eDNA.phylo.PCoA$eig[1] /  
                    sum(eDNA.phylo.PCoA$eig), 3) * 100
```

```
explainvar2 <- round(eDNA.phylo.PCoA$eig[2] /  
                    sum(eDNA.phylo.PCoA$eig), 3) * 100
```

```
explainvar3 <- round(eDNA.phylo.PCoA$eig[3] /  
                    sum(eDNA.phylo.PCoA$eig), 3) * 100
```

```
sum.eig <- sum(explainvar1, explainvar2, explainvar3)
```

```
# OTU Scores
```

```
otu.scores <- t(cor(eDNA.phylo.PCoA$points, OTU.REL))
```

```
otu.scores <- as.matrix(otu.scores)[,1:2]
```

```
otu.scores <- otu.scores[abs(otu.scores[,1]) > 0.7 |  
                      abs(otu.scores[,2]) > 0.7,]
```

```
# Average UniFrac Distance Between Treatments
```

```
eDNA.phylo.m <- as.matrix(phylo.dist)
```

```
all.equal(row.names(eDNA.div), rownames(eDNA.phylo.m))
```

```
## [1] TRUE
```

```
pair.div <- unique(eDNA.div$pair)  
pair.dis <- rep(NA, length(pair.div))  
for(i in 1:length(pair.div)){  
  temp <- row.names(eDNA.div[eDNA.div$pair == pair.div[i], ])  
  pair.dis[i] <- eDNA.phylo.m[temp[1], temp[2]]  
}
```

```
mean(pair.dis)
```

```
## [1] 0.07332018
```

```
# Average BC Distance Between Replicates
```

```
reps <- names(which(table(eDNA.div$qPCR.match) == 4))
```

```
reps.dis <- rep(NA, length(reps))
```

```
for(i in 1:length(reps)){  
  temp <- eDNA.div[eDNA.div$qPCR.match == reps[i] & eDNA.div$treat == "C", ]  
  temp.names <- row.names(temp)  
  reps.dis[i] <- eDNA.phylo.m[temp.names[1], temp.names[2]]  
}
```

```
mean(reps.dis)
```

```
## [1] 0.05628541
```

```
t.test(pair.dis, reps.dis)

##
## Welch Two Sample t-test
##
## data: pair.dis and reps.dis
## t = 1.7885, df = 18.432, p-value = 0.09014
## alternative hypothesis: true difference in means is not equal to 0
## 95 percent confidence interval:
## -0.002941964 0.037011504
## sample estimates:
## mean of x mean of y
## 0.07332018 0.05628541
```

PCoA Plots

Taxonomic PcoA Plot (Supplemental)

```
png(filename="../figures/ordination.png",
      width = 1200, height = 800, res = 96*2)
layout(matrix(1:2, 1, 2), widths = c(9, 3))
par(mar = c(4, 5, 1, 0) + 0.5)

plot(eDNA.PCoA$points[,1], eDNA.PCoA$points[,2],
      ylim = c(-0.4, 0.4), xlim = c(-0.5, 0.4),
      xlab = paste("PCoA 1 (", explainvar1, "%)", sep = ""),
      ylab = paste("PCoA 2 (", explainvar2, "%)", sep = ""),
      #xlab = "", ylab = "", xaxt = "n", yaxt = "n",
      pch = 22, cex = 2.0, type = "n", cex.lab = 1.5, cex.axis = 1,
      axes = FALSE)

# Add Axes
axis(side = 1, labels = T, lwd.ticks = 2, cex.axis = 1.25, las = 1)
axis(side = 2, labels = T, lwd.ticks = 2, cex.axis = 1.25, las = 1)
axis(side = 3, labels = F, lwd.ticks = 2, cex.axis = 1, las = 1, tck=-0.02)
axis(side = 4, labels = F, lwd.ticks = 2, cex.axis = 1, las = 1, tck=-0.02)
axis(side = 1, labels = F, lwd.ticks = 2, cex.axis = 1, las = 1, tck=0.01)
axis(side = 2, labels = F, lwd.ticks = 2, cex.axis = 1, las = 1, tck=0.01)
axis(side = 3, labels = F, lwd.ticks = 2, cex.axis = 1, las = 1, tck=0.01)
axis(side = 4, labels = F, lwd.ticks = 2, cex.axis = 1, las = 1, tck=0.01)
abline(h = 0, v = 0, lty = 3)
box(lwd = 2)

# Subset data
eDNA.div.sort <- eDNA.div[order(eDNA.div[,1]),]
all.equal(row.names(eDNA.PCoA$points), rownames(eDNA.div.sort))
eDNA.points <- data.frame(eDNA.PCoA$points, eDNA.div.sort)

# Gut
eDNA.gut.C <- eDNA.points[ which(eDNA.points$env == "feces" &
                                eDNA.points$treat == "C"), ]
```

```

eDNA.gut.E <- eDNA.points[ which(eDNA.points$env == "feces" &
                                eDNA.points$treat == "E"), ]

# Sed
eDNA.sed.C <- eDNA.points[ which(eDNA.points$env == "sed" &
                                eDNA.points$treat == "C"), ]
eDNA.sed.E <- eDNA.points[ which(eDNA.points$env == "sed" &
                                eDNA.points$treat == "E"), ]

# Soil
eDNA.soil.C <- eDNA.points[ which(eDNA.points$env == "soil" &
                                eDNA.points$treat == "C"), ]
eDNA.soil.E <- eDNA.points[ which(eDNA.points$env == "soil" &
                                eDNA.points$treat == "E"), ]

# Water
eDNA.water.C <- eDNA.points[ which(eDNA.points$env == "water" &
                                eDNA.points$treat == "C"), ]
eDNA.water.E <- eDNA.points[ which(eDNA.points$env == "water" &
                                eDNA.points$treat == "E"), ]

# Add points
# Gut C
points(eDNA.gut.C[,1], eDNA.gut.C[,2], pch = 21,
       cex = 2, col = "red", bg = "white", lwd = 2)
# Gut E
points(eDNA.gut.E[,1], eDNA.gut.E[,2], pch = 21,
       cex = 2, col = "red", bg = "red", lwd = 2)
# Sed C
points(eDNA.sed.C[,1], eDNA.sed.C[,2], pch = 22,
       cex = 2, col = "darkgreen", bg = "white", lwd = 2)
# Sed E
points(eDNA.sed.E[,1], eDNA.sed.E[,2], pch = 22,
       cex = 2, col = "darkgreen", bg = "darkgreen", lwd = 2)
# Soil C
points(eDNA.soil.C[,1], eDNA.soil.C[,2], pch = 24,
       cex = 2, col = "brown", bg = "white", lwd = 2)
# Soil E
points(eDNA.soil.E[,1], eDNA.soil.E[,2], pch = 24,
       cex = 2, col = "brown", bg = "brown", lwd = 2)
# Water C
points(eDNA.water.C[,1], eDNA.water.C[,2], pch = 23,
       cex = 2, col = "blue", bg = "white", lwd = 2)
# Water E
points(eDNA.water.E[,1], eDNA.water.E[,2], pch = 23,
       cex = 2, col = "blue", bg = "blue", lwd = 2)

# Add Legend Outside
par(mar = c(4, 0, 1, 1) + 0.5)
plot.new()
legend(0, 1, c("Gut", "Sediment", "Soil", "Water"),
      pch = c(21, 22, 24, 23),
      pt.bg = c("red", "darkgreen", "brown", "blue"),
      bty = "n", y.intersp = 1.5)
legend(0, 0.25, c("Control", "DNase"), pch = 22,
      pt.bg = c("white", "black"),

```

```

        bty = "n", y.intersp = 1.5)

# Close Plot Device
dev.off()
graphics.off()

# Show Plot
img <- readPNG("../figures/ordination.png")
grid.raster(img)

```

Phylogenetic PcoA Plot (Supplemental)

```

png(filename="../figures/phylo.ordination.png",
     width = 1200, height = 800, res = 96*2)
layout(matrix(1:2, 1, 2), widths = c(9, 3))
par(mar = c(4, 5, 1, 0) + 0.5)

plot(eDNA.phylo.PCoA$points[,1], eDNA.phylo.PCoA$points[,2],
     ylim = c(-0.3, 0.5), xlim = c(-0.3, 0.6),
     xlab = paste("PCoA 1 (", explainvar1, "%)", sep = ""),
     ylab = paste("PCoA 2 (", explainvar2, "%)", sep = ""),
     #xlab = "", ylab = "", xaxt = "n", yaxt = "n",
     pch = 22, cex = 2.0, type = "n", cex.lab = 1.5, cex.axis = 1,
     axes = FALSE)

# Add Axes
axis(side = 1, labels = T, lwd.ticks = 2, cex.axis = 1.25, las = 1)
axis(side = 2, labels = T, lwd.ticks = 2, cex.axis = 1.25, las = 1)
axis(side = 3, labels = F, lwd.ticks = 2, cex.axis = 1, las = 1, tck=-0.02)
axis(side = 4, labels = F, lwd.ticks = 2, cex.axis = 1, las = 1, tck=-0.02)
axis(side = 1, labels = F, lwd.ticks = 2, cex.axis = 1, las = 1, tck=0.01)
axis(side = 2, labels = F, lwd.ticks = 2, cex.axis = 1, las = 1, tck=0.01)
axis(side = 3, labels = F, lwd.ticks = 2, cex.axis = 1, las = 1, tck=0.01)
axis(side = 4, labels = F, lwd.ticks = 2, cex.axis = 1, las = 1, tck=0.01)
abline(h = 0, v = 0, lty = 3)
box(lwd = 2)

# Subset data
eDNA.div.sort <- eDNA.div[order(eDNA.div[,1]),]
all.equal(row.names(eDNA.phylo.PCoA$points), rownames(eDNA.div.sort))
eDNA.points <- data.frame(eDNA.phylo.PCoA$points, eDNA.div.sort)

# Gut
eDNA.gut.C <- eDNA.points[ which(eDNA.points$env == "feces" &
                               eDNA.points$treat == "C"), ]
eDNA.gut.E <- eDNA.points[ which(eDNA.points$env == "feces" &
                               eDNA.points$treat == "E"), ]

# Sed
eDNA.sed.C <- eDNA.points[ which(eDNA.points$env == "sed" &
                               eDNA.points$treat == "C"), ]
eDNA.sed.E <- eDNA.points[ which(eDNA.points$env == "sed" &
                               eDNA.points$treat == "E"), ]

```

```

# Soil
eDNA.soil.C <- eDNA.points[ which(eDNA.points$env == "soil" &
                                eDNA.points$treat == "C"), ]
eDNA.soil.E <- eDNA.points[ which(eDNA.points$env == "soil" &
                                eDNA.points$treat == "E"), ]

# Water
eDNA.water.C <- eDNA.points[ which(eDNA.points$env == "water" &
                                eDNA.points$treat == "C"), ]
eDNA.water.E <- eDNA.points[ which(eDNA.points$env == "water" &
                                eDNA.points$treat == "E"), ]

# Add points
# Gut C
points(eDNA.gut.C[ ,1], eDNA.gut.C[ ,2], pch = 21,
       cex = 2, col = "red", bg = "white", lwd = 2)
# Gut E
points(eDNA.gut.E[ ,1], eDNA.gut.E[ ,2], pch = 21,
       cex = 2, col = "red", bg = "red", lwd = 2)
# Sed C
points(eDNA.sed.C[ ,1], eDNA.sed.C[ ,2], pch = 22,
       cex = 2, col = "darkgreen", bg = "white", lwd = 2)
# Sed E
points(eDNA.sed.E[ ,1], eDNA.sed.E[ ,2], pch = 22,
       cex = 2, col = "darkgreen", bg = "darkgreen", lwd = 2)
# Soil C
points(eDNA.soil.C[ ,1], eDNA.soil.C[ ,2], pch = 24,
       cex = 2, col = "brown", bg = "white", lwd = 2)
# Soil E
points(eDNA.soil.E[ ,1], eDNA.soil.E[ ,2], pch = 24,
       cex = 2, col = "brown", bg = "brown", lwd = 2)
# Water C
points(eDNA.water.C[ ,1], eDNA.water.C[ ,2], pch = 23,
       cex = 2, col = "blue", bg = "white", lwd = 2)
# Soil E
points(eDNA.water.E[ ,1], eDNA.water.E[ ,2], pch = 23,
       cex = 2, col = "blue", bg = "blue", lwd = 2)

# Add Labels to Test Outgroups
# text(eDNA.gut.C[ ,1], eDNA.gut.C[ ,2], labels = eDNA.gut.C$label)
# text(eDNA.gut.E[ ,1], eDNA.gut.E[ ,2], labels = eDNA.gut.E$label)

# Add Legend Outside
par(mar = c(4, 0, 1, 1) + 0.5)
plot.new()
legend(0, 1, c("Gut", "Sediment", "Soil", "Water"),
      pch = c(21, 22, 24, 23),
      pt.bg = c("red", "darkgreen", "brown", "blue"),
      bty = "n", y.intersp = 1.5)
legend(0, 0.25, c("Control", "DNase"), pch = 22,
      pt.bg = c("white", "black"),
      bty = "n", y.intersp = 1.5)

# Close Plot Device

```

```
dev.off()
graphics.off()

# Show Plot
img <- readPNG("../figures/phylo.ordination.png")
grid.raster(img)
```

Beta Diversity Statistics

PERMANOVA: Taxonomic

```
# Check Order of Dataframes
all.equal(row.names(eDNA.div), row.names(OTU.REL.log))

## [1] TRUE

eDNA.permanova <- adonis(OTU.REL.log ~ eDNA.div$env * eDNA.div$treat,
                        method = "bray", binary = FALSE)
eDNA.permanova

##
## Call:
## adonis(formula = OTU.REL.log ~ eDNA.div$env * eDNA.div$treat,          method = "bray", binary = FALSE)
##
## Permutation: free
## Number of permutations: 999
##
## Terms added sequentially (first to last)
##
##              Df SumsOfSqs MeanSqs F.Model      R2 Pr(>F)
## eDNA.div$env      3      7.6829  2.56095   9.3326 0.35195  0.001
## eDNA.div$treat     1      0.1142  0.11422   0.4163 0.00523  1.000
## eDNA.div$env:eDNA.div$treat  3      0.3119  0.10398   0.3789 0.01429  1.000
## Residuals        50     13.7205  0.27441           0.62853
## Total            57     21.8295           1.00000
##
## eDNA.div$env      ***
## eDNA.div$treat
## eDNA.div$env:eDNA.div$treat
## Residuals
## Total
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

eDNA.bc.dis <- vegdist(OTU.REL.log, method = "bray", binary = "FALSE")
beta.disp <- betadisper(d = eDNA.bc.dis, group = eDNA.div$env)
permutest(beta.disp, 99)

##
## Permutation test for homogeneity of multivariate dispersions
## Permutation: free
## Number of permutations: 999
##
```

```
## Response: Distances
##           Df Sum Sq Mean Sq    F N.Perm Pr(>F)
## Groups      3 0.2137 0.071233 13.787    999 0.001 ***
## Residuals 54 0.2790 0.005167
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## Pairwise comparisons:
## (Observed p-value below diagonal, permuted p-value above diagonal)
##           feces      sed      soil water
## feces           1.0000e-03 1.0000e-03 0.001
## sed    5.7913e-04           3.8000e-02 0.215
## soil   2.0056e-05 4.2912e-02           0.001
## water  2.2807e-03 2.2940e-01 1.9272e-03
```

```
TukeyHSD(beta.disp)
```

```
## Tukey multiple comparisons of means
## 95% family-wise confidence level
##
## Fit: aov(formula = distances ~ group, data = df)
##
## $group
##           diff          lwr          upr      p adj
## sed-feces -0.11233229 -0.18206456 -0.04260002 0.0004509
## soil-feces -0.17047994 -0.24324545 -0.09771443 0.0000005
## water-feces -0.08725668 -0.15462445 -0.01988891 0.0061528
## soil-sed    -0.05814765 -0.13310759 0.01681228 0.1806561
## water-sed    0.02507561 -0.04465666 0.09480788 0.7762587
## water-soil   0.08322326 0.01045775 0.15598877 0.0189601
```

PERMANOVA: Phylogenetic

```
# Check Order of Dataframes
all.equal(row.names(eDNA.div), row.names(as.matrix(phylo.dist)))
```

```
## [1] TRUE
```

```
eDNA.permanova <- adonis(phylo.dist ~ eDNA.div$env * eDNA.div$treat,
                        binary = FALSE)
```

```
eDNA.permanova
```

```
##
## Call:
## adonis(formula = phylo.dist ~ eDNA.div$env * eDNA.div$treat,      binary = FALSE)
##
## Permutation: free
## Number of permutations: 999
##
## Terms added sequentially (first to last)
##
##           Df SumsOfSqs MeanSqs F.Model      R2 Pr(>F)
## eDNA.div$env      3    4.1920 1.39733   31.821 0.65359 0.001
## eDNA.div$treat     1    0.0091 0.00907    0.207 0.00141 0.984
```



```
## eDNA.div$env:eDNA.div$treat 3 0.0171 0.00571 0.130 0.00267 1.000
## Residuals 50 2.1956 0.04391 0.34233
## Total 57 6.4138 1.00000
##
## eDNA.div$env ***
## eDNA.div$treat
## eDNA.div$env:eDNA.div$treat
## Residuals
## Total
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
beta.disp <- betadisper(d = phylo.dist, group = eDNA.div$env)
permutest(beta.disp, 99)
```

```
##
## Permutation test for homogeneity of multivariate dispersions
## Permutation: free
## Number of permutations: 999
##
## Response: Distances
## Df Sum Sq Mean Sq F N.Perm Pr(>F)
## Groups 3 0.26109 0.087030 13.403 999 0.001 ***
## Residuals 54 0.35064 0.006493
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## Pairwise comparisons:
## (Observed p-value below diagonal, permuted p-value above diagonal)
## feces sed soil water
## feces 0.00100000 0.00200000 0.001
## sed 0.00025791 0.42600000 0.995
## soil 0.00165032 0.43147784 0.497
## water 0.00016369 0.99827983 0.49095914
```

```
TukeyHSD(beta.disp)
```

```
## Tukey multiple comparisons of means
## 95% family-wise confidence level
##
## Fit: aov(formula = distances ~ group, data = df)
##
## $group
## diff lwr upr p adj
## sed-feces -1.535453e-01 -0.23171919 -0.07537138 0.0000179
## soil-feces -1.398666e-01 -0.22144095 -0.05829226 0.0001797
## water-feces -1.535844e-01 -0.22910759 -0.07806125 0.0000093
## soil-sed 1.367868e-02 -0.07035574 0.09771310 0.9728051
## water-sed -3.913702e-05 -0.07821304 0.07813477 1.0000000
## water-soil -1.371782e-02 -0.09529216 0.06785653 0.9701593
```

Mantel Test: Taxonomic

```

# Check order of design
all.equal(row.names(eDNA.div), row.names(OTU.REL.log))

## [1] TRUE

# Subset OTU Matrix for Each Molecule
OTU.REL.c <- OTU.REL.log[which(eDNA.div$treat == "C"), ]
OTU.REL.e <- OTU.REL.log[which(eDNA.div$treat == "E"), ]

# Subset the Design Matrix
eDNA.div.c <- eDNA.div[which(eDNA.div$treat == "C"), ]
eDNA.div.e <- eDNA.div[which(eDNA.div$treat == "E"), ]

# Check Order of Subsets
all.equal(sort(eDNA.div.c$samp.code), sort(eDNA.div.e$samp.code))

## [1] TRUE

# Make Sure Subset OTU Matrices are Aligned by Sample Code
OTU.REL.c2 <- OTU.REL.c[order(eDNA.div.c$samp.code), ]
OTU.REL.e2 <- OTU.REL.e[order(eDNA.div.e$samp.code), ]

# Calculate Bray-Curtis Distances
dist.c <- vegdist(OTU.REL.c2, "bray")
dist.e <- vegdist(OTU.REL.e2, "bray")

# Run Mantel Test of Distance Matrix
mantel.rtest(dist.c, dist.e, nrepet = 999)

## Monte-Carlo test
## Observation: 0.9589581
## Call: mantel.rtest(m1 = dist.c, m2 = dist.e, nrepet = 999)
## Based on 999 replicates
## Simulated p-value: 0.001

```

Mantel Test: Phylogenetic

```

# Turn into square matrix
phylo.dist.m <- as.matrix(phylo.dist)

# Check order of design
all.equal(row.names(eDNA.div), row.names(phylo.dist.m))

## [1] TRUE

# Subset Phylo Distance Matrix Based on Design
dist.c <- phylo.dist.m[which(eDNA.div$treat == "C"),
                        which(eDNA.div$treat == "C")]
dist.e <- phylo.dist.m[which(eDNA.div$treat == "E"),
                        which(eDNA.div$treat == "E")]

# Define Order Based on Pairs
ord.c <- order(as.numeric(eDNA.div$pair[which(eDNA.div$treat == "C")]))
ord.e <- order(as.numeric(eDNA.div$pair[which(eDNA.div$treat == "E")]))

```

```

# Reorder Distance Matrices
dist.c <- dist.c[ord.c, ord.c]
dist.e <- dist.e[ord.e, ord.e]

# Turn Square into Lower Triangle Matrix
dist.c <- as.dist(dist.c)
dist.e <- as.dist(dist.e)

# Run Mantel Test of Distance Matrix
mantel.rtest(dist.c, dist.e, nrepet = 999)

## Monte-Carlo test
## Observation: 0.9955652
## Call: mantelnoneuclid(m1 = m1, m2 = m2, nrepet = nrepet)
## Based on 999 replicates
## Simulated p-value: 0.001

```

Centroid Distances Ratios: Taxonomic

```

# Data Check
all.equal(row.names(eDNA.div), row.names(OTU.REL.log))

## [1] TRUE

eDNA.dist <- vegdist(OTU.REL.log, method = "bray")
eDNA.env <- eDNA.div$env
eDNA.group <- paste(eDNA.div$env, eDNA.div$treat, sep = "")

disp.env <- betadisper(eDNA.dist, eDNA.env)
disp.group <- betadisper(eDNA.dist, eDNA.group)

# Soil
env soi <- which(eDNA.env == "soil")
cent.soil <- as.data.frame(matrix(NA, nrow = length(env soi), ncol = 3))
colnames(cent.soil) <- c("T", "C", "E")
rownames(cent.soil) <- rownames(eDNA.div)[env soi]

for (i in 1:length(env soi)){
  cent.soil[i, 1] <- dist(rbind(disp.env$vectors[env soi[i]],
    disp.env$centroids[3, ]), method = "euclidean")
  cent.soil[i, 2] <- dist(rbind(disp.group$vectors[env soi[i]],
    disp.group$centroids[5, ]), method = "euclidean")
  cent.soil[i, 3] <- dist(rbind(disp.group$vectors[env soi[i]],
    disp.group$centroids[6, ]), method = "euclidean")
}

pairs.soil <- eDNA.div[env soi, c(3,4,5,7)]

cent.soil.ratio <- as.data.frame(matrix(NA,
  nrow = length(unique(pairs.soil$pair)), ncol = 3))
colnames(cent.soil.ratio) <- c("T", "C", "E")
rownames(cent.soil.ratio) <- paste(pairs.soil$env, pairs.soil$pair,
  sep = "")[which(pairs.soil$treat == "E") ]

```

```

for (i in 1:dim(cent.soil.ratio)[1]){
  pair <- as.numeric(gsub("soil", "", rownames(cent.soil.ratio)[i]))
  ratio.temp.1 <- (cent.soil[, 1][which(pairs.soil$pair == pair &
                                       pairs.soil$treat == "C")]) /
    (cent.soil[, 1][which(pairs.soil$pair == pair &
                           pairs.soil$treat == "E")])
  cent.soil.ratio[i, 1] <- ratio.temp.1
  ratio.temp.2 <- (cent.soil[, 2][which(pairs.soil$pair == pair &
                                       pairs.soil$treat == "C")]) /
    (cent.soil[, 2][which(pairs.soil$pair == pair &
                           pairs.soil$treat == "E")])
  cent.soil.ratio[i, 2] <- ratio.temp.2
  ratio.temp.3 <- (cent.soil[, 3][which(pairs.soil$pair == pair &
                                       pairs.soil$treat == "C")]) /
    (cent.soil[, 3][which(pairs.soil$pair == pair &
                           pairs.soil$treat == "E")])
  cent.soil.ratio[i, 3] <- ratio.temp.3
}

# Water
env.wat <- which(eDNA.env == "water")
cent.water <- as.data.frame(matrix(NA, nrow = length(env.wat), ncol = 3))
colnames(cent.water) <- c("T", "C", "E")
rownames(cent.water) <- rownames(eDNA.div)[env.wat]

for (i in 1:length(env.wat)){
  cent.water[i, 1] <- dist(rbind(dispenv$centroids[env.wat[i]],
                                dispenv$centroids[4, ]), method = "euclidean")
  cent.water[i, 2] <- dist(rbind(dispgroup$centroids[env.wat[i]],
                                dispgroup$centroids[7, ]), method = "euclidean")
  cent.water[i, 3] <- dist(rbind(dispgroup$centroids[env.wat[i]],
                                dispgroup$centroids[8, ]), method = "euclidean")
}

pairs.water <- eDNA.div[env.wat, c(3,4,5,7)]

cent.water.ratio <- as.data.frame(matrix(NA,
                                         nrow = length(unique(pairs.water$pair)), ncol = 3))
colnames(cent.water.ratio) <- c("T", "C", "E")
rownames(cent.water.ratio) <- paste(pairs.water$env, pairs.water$pair,
                                   sep = "")[which(pairs.water$treat == "E")]

for (i in 1:dim(cent.water.ratio)[1]){
  pair <- as.numeric(gsub("water", "", rownames(cent.water.ratio)[i]))
  ratio.temp.1 <- (cent.water[, 1][which(pairs.water$pair == pair &
                                       pairs.water$treat == "C")]) /
    (cent.water[, 1][which(pairs.water$pair == pair &
                           pairs.water$treat == "E")])
  cent.water.ratio[i, 1] <- ratio.temp.1
  ratio.temp.2 <- (cent.water[, 2][which(pairs.water$pair == pair &
                                       pairs.water$treat == "C")]) /
    (cent.water[, 2][which(pairs.water$pair == pair &
                           pairs.water$treat == "E")])
  cent.water.ratio[i, 2] <- ratio.temp.2
  ratio.temp.3 <- (cent.water[, 3][which(pairs.water$pair == pair &
                                       pairs.water$treat == "C")]) /
    (cent.water[, 3][which(pairs.water$pair == pair &
                           pairs.water$treat == "E")])
  cent.water.ratio[i, 3] <- ratio.temp.3
}

```

```

cent.water.ratio[i, 2] <- ratio.temp.2
ratio.temp.3 <- (cent.water[, 3][which(pairs.water$pair == pair &
                                     pairs.water$treat == "C")]) /
               (cent.water[, 3][which(pairs.water$pair == pair &
                                     pairs.water$treat == "E")])
cent.water.ratio[i, 3] <- ratio.temp.3
}

# Sediments
env.sed <- which(eDNA.env == "sed")
cent.sed <- as.data.frame(matrix(NA, nrow = length(env.sed), ncol = 3))
colnames(cent.sed) <- c("T", "C", "E")
rownames(cent.sed) <- rownames(eDNA.div)[env.sed]

for (i in 1:length(env.sed)){
  cent.sed[i, 1] <- dist(rbind(dispenv$centroids[env.sed[i]],
                              dispenv$centroids[2, ]), method = "euclidean")
  cent.sed[i, 2] <- dist(rbind(dispgroup$centroids[env.sed[i]],
                              dispgroup$centroids[3, ]), method = "euclidean")
  cent.sed[i, 3] <- dist(rbind(dispgroup$centroids[env.sed[i]],
                              dispgroup$centroids[4, ]), method = "euclidean")
}

pairs.sed <- eDNA.div[env.sed, c(3,4,5,7)]

cent.sed.ratio <- as.data.frame(matrix(NA,
                                     nrow = length(unique(pairs.sed$pair)), ncol = 3))
colnames(cent.sed.ratio) <- c("T", "C", "E")
rownames(cent.sed.ratio) <- paste(pairs.sed$env, pairs.sed$pair,
                                sep = "")[which(pairs.sed$treat == "E")]

for (i in 1:dim(cent.sed.ratio)[1]){
  pair <- as.numeric(gsub("sed", "", rownames(cent.sed.ratio)[i]))
  ratio.temp.1 <- (cent.sed[, 1][which(pairs.sed$pair == pair &
                                     pairs.sed$treat == "C")]) /
                (cent.sed[, 1][which(pairs.sed$pair == pair &
                                     pairs.sed$treat == "E")])
  cent.sed.ratio[i, 1] <- ratio.temp.1
  ratio.temp.2 <- (cent.sed[, 2][which(pairs.sed$pair == pair &
                                     pairs.sed$treat == "C")]) /
                (cent.sed[, 2][which(pairs.sed$pair == pair &
                                     pairs.sed$treat == "E")])
  cent.sed.ratio[i, 2] <- ratio.temp.2
  ratio.temp.3 <- (cent.sed[, 3][which(pairs.sed$pair == pair &
                                     pairs.sed$treat == "C")]) /
                (cent.sed[, 3][which(pairs.sed$pair == pair &
                                     pairs.sed$treat == "E")])
  cent.sed.ratio[i, 3] <- ratio.temp.3
}

# Feces
env.fec <- which(eDNA.env == "feces")
cent.feces <- as.data.frame(matrix(NA, nrow = length(env.fec), ncol = 3))

```

```

colnames(cent.feces) <- c("T", "C", "E")
rownames(cent.feces) <- rownames(eDNA.div)[env.fec]

for (i in 1:length(env.fec)){
  cent.feces[i, 1] <- dist(rbind(dispenv$vector[env.fec[i]],
                                dispenv$centroid[1, ]), method = "euclidean")
  cent.feces[i, 2] <- dist(rbind(dispgroup$vector[env.fec[i]],
                                dispgroup$centroid[1, ]), method = "euclidean")
  cent.feces[i, 3] <- dist(rbind(dispgroup$vector[env.fec[i]],
                                dispgroup$centroid[2, ]), method = "euclidean")
}

pairs.feces <- eDNA.div[env.fec, c(3,4,5,7)]

cent.feces.ratio <- as.data.frame(matrix(NA,
                                         nrow = length(unique(pairs.feces$pair)), ncol = 3))
colnames(cent.feces.ratio) <- c("T", "C", "E")
rownames(cent.feces.ratio) <- paste(pairs.feces$env, pairs.feces$pair,
                                    sep = "")[which(pairs.feces$treat == "E") ]

for (i in 1:dim(cent.feces.ratio)[1]){
  pair <- as.numeric(gsub("feces", "", rownames(cent.feces.ratio)[i]))
  ratio.temp.1 <- (cent.feces[, 1][which(pairs.feces$pair == pair &
                                         pairs.feces$treat == "C")]) /
    (cent.feces[, 1][which(pairs.feces$pair == pair &
                           pairs.feces$treat == "E")])
  cent.feces.ratio[i, 1] <- ratio.temp.1
  ratio.temp.2 <- (cent.feces[, 2][which(pairs.feces$pair == pair &
                                         pairs.feces$treat == "C")]) /
    (cent.feces[, 2][which(pairs.feces$pair == pair &
                           pairs.feces$treat == "E")])
  cent.feces.ratio[i, 2] <- ratio.temp.2
  ratio.temp.3 <- (cent.feces[, 3][which(pairs.feces$pair == pair &
                                         pairs.feces$treat == "C")]) /
    (cent.feces[, 3][which(pairs.feces$pair == pair &
                           pairs.feces$treat == "E")])
  cent.feces.ratio[i, 3] <- ratio.temp.3
}

# Ratio Data Table
ratio.list <- list(cent.feces.ratio[,1], cent.sed.ratio[,1],
                  cent.soil.ratio[,1], cent.water.ratio[,1])
cent.dist.table <- as.data.frame(matrix(NA, nrow = 4, ncol = 5))
colnames(cent.dist.table) <- c("env", "mean", "sem", "LCI", "UCI")
cent.dist.table$env <- c("Gut", "Sediment", "Soil", "Water")
cent.dist.table$mean <- unlist(lapply(ratio.list, mean))
cent.dist.table$sem <- unlist(lapply(ratio.list, sem))
cent.dist.table$LCI <- unlist(lapply(ratio.list, FUN = function(x) t.test(x)$conf.int[1]))
cent.dist.table$UCI <- unlist(lapply(ratio.list, FUN = function(x) t.test(x)$conf.int[2]))
t.cent.dist.table <- as.data.frame(cent.dist.table)

# Regression: centroid ratio vs. proportion eDNA
# Combine Centroid Distances

```

```

centroid.dists <- c(cent.feces.ratio$T, cent.sed.ratio$T, cent.soil.ratio$T,
  cent.water.ratio$T)
names(centroid.dists) <- c(rownames(cent.feces.ratio), rownames(cent.sed.ratio),
  rownames(cent.soil.ratio), rownames(cent.water.ratio))

# Reorganize
names(centroid.dists) %in% paste(eDNA.div.c$env, eDNA.div.c$pair, sep = "")

## [1] TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE
## [15] TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE
## [29] TRUE

all.equal(names(centroid.dists), paste(eDNA.div.c$env, eDNA.div.c$pair, sep = ""))

## [1] "28 string mismatches"

cent.dist <- centroid.dists[match(names(centroid.dists),
  paste(eDNA.div.c$env, eDNA.div.c$pair, sep = ""))]

# Combine Centroid Distance and Design Information
centroid.div <- cbind(eDNA.div.c[,c(1:3, 5:7)], cent.dist)

centroid.div$qPCR.match %in% eDNA.prop$sample.number

## [1] TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE
## [15] TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE
## [29] TRUE

centroid.div$prop <- NA
for (i in 1:length(centroid.div$prop)){
  centroid.div$prop[i] <- eDNA.prop$prop[which(eDNA.prop$sample.number ==
    centroid.div$qPCR.match[i])]
}

# Take averages of technical reps
centroid.div.2 <- ddply(centroid.div, .(env, qPCR.match, prop),
  summarize, cent.dist = mean(cent.dist))

t.cent.reg <- lm(cent.dist ~ prop * env, data = centroid.div.2)
t.cent.reg <- lm(cent.dist ~ prop, data = centroid.div.2)

summary(t.cent.reg)

##
## Call:
## lm(formula = cent.dist ~ prop, data = centroid.div.2)
##
## Residuals:
##      Min       1Q   Median       3Q      Max
## -0.49248 -0.04018 -0.01482  0.07151  0.84292
##
## Coefficients:
##              Estimate Std. Error t value Pr(>|t|)
## (Intercept)   1.00409    0.12661   7.930 1.91e-07 ***
## prop          0.02031    0.28691   0.071   0.944

```

```
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 0.2703 on 19 degrees of freedom
## Multiple R-squared:  0.0002635, Adjusted R-squared:  -0.05235
## F-statistic: 0.005009 on 1 and 19 DF,  p-value: 0.9443
ttest(t.cent.reg, 1, 1)

##           t           df           p
## 0.03229089 19.00000000 0.97457688
# answer => no correlation between proportion eDNA and centroid dist ratio.
```

Centroid Distances Ratios: Phylogenetic

```
# Data Check
all.equal(row.names(eDNA.div), row.names(phylo.dist.m))

## [1] TRUE

eDNA.dist <- as.dist(phylo.dist.m)
eDNA.env <- eDNA.div$env
eDNA.group <- paste(eDNA.div$env, eDNA.div$treat, sep = "")

disp.env <- betadisper(eDNA.dist, eDNA.env)
disp.group <- betadisper(eDNA.dist, eDNA.group)

disp.group$vectors
```

	PCoA1	PCoA2	PCoA3	PCoA4	PCoA5
## eDNA_01	-0.13887108	-0.113744265	0.148915828	-0.0187348393	3.229884e-04
## eDNA_02	-0.14432245	0.045760113	-0.092823923	0.0911463532	1.329472e-02
## eDNA_03	-0.13347184	0.019785511	-0.081307708	0.0964246861	2.062611e-02
## eDNA_04	-0.17372768	0.004956970	-0.067521221	0.1492361538	2.050991e-02
## eDNA_05	-0.14740165	-0.084168080	0.182728764	-0.0313473532	-1.818740e-02
## eDNA_06	0.45819121	-0.035639120	-0.015356712	0.0005375485	1.263160e-01
## eDNA_07	-0.08974623	0.018964339	-0.104419835	0.1466444586	1.304599e-02
## eDNA_08	-0.15040514	-0.087445637	0.180724842	-0.0264791122	-1.634051e-02
## eDNA_09	-0.15576427	-0.064524158	0.099536567	0.0002918775	-2.659837e-03
## eDNA_10	0.35786453	-0.126809628	-0.012969061	-0.0619094968	1.152163e-01
## eDNA_11	0.35186294	0.384284549	0.175650888	0.0566234249	-7.220159e-02
## eDNA_12	-0.14703301	-0.078771528	0.147484251	-0.0465827280	-9.900697e-03
## eDNA_13	0.39127600	-0.129312846	-0.024796196	-0.0223439585	1.221936e-01
## eDNA_14	0.44206721	-0.082891258	-0.032157599	-0.0007412091	1.104368e-01
## eDNA_15	0.45496212	-0.082913477	-0.020162441	0.0179377124	1.292853e-01
## eDNA_16	0.46853197	-0.054256805	-0.008367318	0.0156497209	1.437989e-01
## eDNA_17	-0.17608872	0.013122425	-0.062174286	0.0997215035	1.575171e-02
## eDNA_18	-0.14807533	0.086492529	-0.121935470	-0.0054977632	9.192956e-03
## eDNA_19	-0.14272123	-0.089141638	0.154927597	-0.0565422183	-8.919142e-03
## eDNA_20	-0.15465025	0.077572001	-0.115958902	0.0204696922	1.074766e-02
## eDNA_21	-0.16578271	-0.049511758	0.098578089	-0.0115604085	-4.909437e-03
## eDNA_22	-0.17219486	0.031807886	-0.098361623	0.1141303986	1.857999e-02
## eDNA_23	-0.11041976	-0.102187826	0.055647846	0.0324638239	1.609050e-02
## eDNA_24	-0.11612508	-0.093058323	0.042926097	0.0351115733	1.568372e-02


```

## eDNA_25 0.30785981 -0.171408873 -0.103526202 0.0032092297 -2.338961e-01
## eDNA_26 0.33403702 0.347440988 0.160835874 0.0541657651 -5.099250e-02
## eDNA_27 -0.06006449 0.135072027 -0.126146457 -0.1842597747 4.697203e-03
## eDNA_28 -0.16559047 -0.078716662 0.152356549 -0.0227536927 -4.244886e-03
## eDNA_29 -0.15756180 0.051118855 -0.038334090 -0.1083715400 -1.755319e-03
## eDNA_30 -0.15838179 0.051663319 -0.035416894 -0.1117616115 -2.455067e-03
## eDNA_31 0.30650883 -0.183486778 -0.100735851 0.0062625963 -2.427458e-01
## eDNA_32 0.34489255 -0.156251158 -0.096239424 0.0074124011 -2.159014e-01
## eDNA_33 -0.05836522 0.133277549 -0.118453616 -0.1949018342 2.492396e-03
## eDNA_34 -0.17145252 0.033055144 -0.099688981 0.1136270064 1.894753e-02
## eDNA_35 -0.16682467 -0.083335244 0.151747418 -0.0143969314 -2.445223e-03
## eDNA_36 -0.15623972 0.082634741 -0.081942896 -0.1012535080 -8.565257e-05
## eDNA_37 -0.14079384 0.088262193 -0.069853679 -0.1696581338 -9.686604e-03
## eDNA_38 -0.14860794 -0.080698010 0.174314413 -0.0321870685 -1.639609e-02
## eDNA_39 -0.15272153 -0.083483936 0.171573213 -0.0273405802 -1.455097e-02
## eDNA_40 0.34946122 0.365871669 0.177840357 0.0804402896 -5.966594e-02
## eDNA_41 -0.17339488 0.016381250 -0.063427103 0.0836172199 1.686201e-02
## eDNA_42 -0.15147358 0.086708077 -0.083081911 -0.1209415415 -2.849224e-03
## eDNA_43 -0.14791033 0.087767347 -0.074780218 -0.1562359456 -6.082842e-03
## eDNA_44 -0.16432929 -0.076944554 0.102897286 -0.0255434767 5.510990e-03
## eDNA_45 0.31485364 -0.185455321 -0.100591424 0.0074105109 -2.359080e-01
## eDNA_46 -0.14801951 -0.111011226 0.139661885 -0.0097495967 1.457497e-03
## eDNA_47 -0.04003714 0.137473461 -0.117780281 -0.2150447065 -1.300527e-03
## eDNA_48 0.33927146 -0.138505246 -0.036791434 -0.0490148993 1.000670e-01
## eDNA_49 -0.14586016 0.026727690 -0.081840798 0.1123185564 1.398562e-02
## eDNA_50 -0.16386689 0.009824799 -0.080469406 0.1590369626 2.147929e-02
## eDNA_51 -0.17180480 0.005516216 -0.055797040 0.1106095390 1.783409e-02
## eDNA_52 -0.07099080 0.050005250 -0.115286169 0.1086473579 9.079120e-03
## eDNA_53 -0.06926685 0.129146016 -0.110350150 -0.1983343776 3.201907e-03
## eDNA_54 0.33412287 0.321800217 0.157048370 0.0715676929 -4.239369e-02
## eDNA_55 -0.14776790 -0.111428197 0.143654063 -0.0194012192 1.303598e-03
## eDNA_56 0.37909964 -0.121710938 -0.021892683 -0.0380869543 1.179231e-01
## eDNA_57 -0.17028834 0.009121198 -0.073849899 0.1277328518 2.068884e-02
## eDNA_58 -0.16644730 0.005198155 -0.074461296 0.1585295727 1.985107e-02
##
## PCoA6 PCoA7 PCoA8 PCoA9
## eDNA_01 0.0016338620 -0.019609578 0.0096218904 2.688449e-02
## eDNA_02 -0.0359885813 0.045306409 0.0424549814 -1.993553e-04
## eDNA_03 0.0644458905 -0.014279183 -0.0578159263 -1.163173e-02
## eDNA_04 0.0109430733 -0.029852960 -0.0074591384 -1.504662e-02
## eDNA_05 -0.0327992221 -0.001837232 0.0228021420 -6.351836e-02
## eDNA_06 -0.0439166536 -0.112445989 -0.0514265136 -2.009659e-02
## eDNA_07 -0.1438317639 0.040989519 0.0552596389 -4.419240e-03
## eDNA_08 -0.0249328895 -0.012566976 0.0266094086 -5.353080e-02
## eDNA_09 -0.0423399252 0.031796370 -0.1175743027 1.009166e-01
## eDNA_10 0.0224962703 0.078631555 0.0413615196 2.255719e-02
## eDNA_11 0.0236773398 0.008632553 0.0101957501 1.982284e-02
## eDNA_12 -0.0475121701 0.047841935 -0.0678258915 7.906557e-03
## eDNA_13 0.0299005460 0.083011266 0.0274643851 5.858388e-03
## eDNA_14 -0.0181826766 -0.058355828 -0.0400860786 -2.410674e-02
## eDNA_15 0.0021803014 -0.028387709 -0.0409299865 -3.045176e-02
## eDNA_16 -0.0092941503 -0.056403098 -0.0452681407 -2.643809e-02
## eDNA_17 0.0793423243 -0.054255517 -0.0282188398 -2.716187e-02
## eDNA_18 0.0175033047 0.069429834 -0.0270078907 3.038935e-02
## eDNA_19 -0.0299253467 0.032763640 -0.0365545944 -1.864893e-02

```

##	eDNA_20	0.0254975104	0.052554475	-0.0118883334	1.566946e-02
##	eDNA_21	-0.0518326927	0.044313159	-0.1233029290	8.701100e-02
##	eDNA_22	-0.0020715093	0.001392134	-0.0438221591	7.635016e-05
##	eDNA_23	0.0876429650	-0.057646355	0.1184035282	8.234861e-02
##	eDNA_24	0.0886070212	-0.075055880	0.1205692711	8.551374e-02
##	eDNA_25	-0.0006266059	-0.002691174	0.0005298785	1.982630e-03
##	eDNA_26	0.0121422473	0.007255251	0.0052440133	7.487237e-03
##	eDNA_27	-0.0575081976	-0.074986634	0.0010362742	2.528342e-02
##	eDNA_28	-0.0203506990	0.008541791	0.0187765106	-5.127172e-02
##	eDNA_29	0.0612139171	0.036420313	-0.0052891164	-4.745949e-02
##	eDNA_30	0.0616921536	0.036355673	-0.0053040353	-4.904813e-02
##	eDNA_31	0.0087061259	0.001320261	-0.0088155731	-4.092760e-03
##	eDNA_32	-0.0055816087	-0.017989604	-0.0076552118	-3.005899e-03
##	eDNA_33	-0.0539545420	-0.076858623	0.0129230600	1.734522e-02
##	eDNA_34	-0.0027078941	0.001119881	-0.0405281115	-1.883438e-04
##	eDNA_35	-0.0116048566	0.003088284	0.0022183821	-3.488150e-02
##	eDNA_36	0.0525239958	0.054776318	-0.0146238296	-2.535990e-04
##	eDNA_37	0.0350047431	0.038013059	0.0094035294	-3.208103e-02
##	eDNA_38	-0.0270245339	-0.009068394	0.0301552862	-6.292899e-02
##	eDNA_39	-0.0141169866	-0.017366085	0.0339065498	-5.270927e-02
##	eDNA_40	0.0157702262	0.011906395	0.0092340166	1.029985e-02
##	eDNA_41	0.0832517121	-0.020872502	-0.0499166824	-1.007866e-02
##	eDNA_42	0.0523634220	0.057787377	-0.0100838524	-1.028667e-02
##	eDNA_43	0.0367028127	0.030760164	0.0065317993	-2.565771e-02
##	eDNA_44	-0.0018409376	-0.042950918	-0.0008169888	3.966586e-02
##	eDNA_45	0.0105136869	0.002111162	-0.0102841515	-5.005653e-03
##	eDNA_46	0.0080643443	-0.029185149	-0.0017913136	4.889566e-02
##	eDNA_47	-0.0606663523	-0.076225550	0.0188985866	2.371086e-02
##	eDNA_48	0.0167548754	0.081713448	0.0418468290	1.957848e-02
##	eDNA_49	-0.0332293256	0.026244261	0.0463037317	-5.603277e-03
##	eDNA_50	-0.0324355950	-0.009390412	0.0137849007	-9.808737e-03
##	eDNA_51	0.0830666471	-0.059998801	-0.0163599305	-2.347848e-02
##	eDNA_52	-0.1492251202	0.051971011	0.0819823207	7.237150e-03
##	eDNA_53	-0.0507990107	-0.060625882	0.0154873003	1.223096e-02
##	eDNA_54	0.0058410983	0.013029014	0.0069882220	-3.747693e-03
##	eDNA_55	0.0063753091	-0.017627928	0.0016573764	3.501957e-02
##	eDNA_56	0.0208993765	0.075881969	0.0361576063	1.752844e-02
##	eDNA_57	0.0087042854	-0.017419553	-0.0142579865	-1.195463e-02
##	eDNA_58	-0.0291615407	-0.021004966	0.0170988189	-1.242761e-02
##		PCoA10	PCoA11	PCoA12	PCoA13
##	eDNA_01	0.024021703	-5.133890e-03	0.0204391920	0.0238877249
##	eDNA_02	0.047570762	-3.511342e-02	0.0122560834	0.0012001612
##	eDNA_03	-0.028605292	-5.201966e-02	0.0388949314	-0.0496491757
##	eDNA_04	-0.031173906	2.626122e-02	-0.0157379851	0.0379134624
##	eDNA_05	-0.014928752	-3.969598e-02	-0.0501515521	-0.0097030325
##	eDNA_06	0.061488825	-2.977847e-03	0.0096768923	-0.0115576810
##	eDNA_07	0.022538566	1.904297e-02	0.0104337559	-0.0400070322
##	eDNA_08	-0.014849757	-4.039063e-02	-0.0440502964	-0.0088334406
##	eDNA_09	-0.000925671	6.555052e-03	-0.0497429321	-0.0242836078
##	eDNA_10	-0.041229208	5.353806e-03	0.0044348366	-0.0041321481
##	eDNA_11	0.007808336	2.996740e-03	0.0191019643	0.0182865631
##	eDNA_12	-0.002418567	4.075164e-02	0.0657996127	-0.0326053527
##	eDNA_13	-0.054537698	-5.427675e-03	-0.0037307145	0.0109396602
##	eDNA_14	0.032199603	-1.886252e-03	-0.0066401903	-0.0030182660

##	eDNA_15	0.052377897	6.331978e-03	-0.0155192305	0.0070618883
##	eDNA_16	0.062933609	6.415434e-03	-0.0084675402	0.0067460314
##	eDNA_17	-0.040562438	-8.537188e-03	0.0072450596	-0.0326600194
##	eDNA_18	0.048578645	-6.238044e-02	-0.0010375193	0.0117862948
##	eDNA_19	0.002827383	4.230367e-02	0.0575438216	-0.0277999782
##	eDNA_20	0.040416997	-5.875014e-02	0.0009568382	0.0051811642
##	eDNA_21	-0.005528948	3.539678e-03	-0.0518393381	-0.0098969066
##	eDNA_22	-0.034632985	2.713478e-02	-0.0317086812	0.0052169946
##	eDNA_23	0.035195091	2.520837e-02	-0.0164828310	-0.0305404659
##	eDNA_24	0.034094661	1.993170e-02	-0.0161321524	-0.0260339489
##	eDNA_25	-0.002359715	-3.892424e-03	0.0037291865	-0.0030874771
##	eDNA_26	0.002855288	-1.910143e-03	-0.0012313651	0.0010058764
##	eDNA_27	-0.048880696	-6.566685e-03	0.0094698828	-0.0008173060
##	eDNA_28	-0.010930560	6.698701e-03	0.0489274553	0.0186775580
##	eDNA_29	0.025796417	7.897704e-02	-0.0189223324	-0.0275191749
##	eDNA_30	0.023503912	7.851801e-02	-0.0152869830	-0.0241148289
##	eDNA_31	-0.003575084	6.303433e-04	-0.0025123240	0.0023996388
##	eDNA_32	0.019308391	1.036123e-03	0.0065733879	-0.0004516448
##	eDNA_33	-0.044476243	-4.475989e-03	0.0012826494	-0.0061104058
##	eDNA_34	-0.034016659	2.456598e-02	-0.0284882973	0.0068106459
##	eDNA_35	-0.008968516	-4.080839e-03	0.0459403010	0.0260533076
##	eDNA_36	0.036963668	-1.509006e-02	-0.0038635281	0.0176985469
##	eDNA_37	0.035698568	1.800599e-02	-0.0110930379	0.0164298236
##	eDNA_38	-0.009687343	-3.440764e-02	-0.0468034972	-0.0085645510
##	eDNA_39	-0.008761610	-3.334874e-02	-0.0393323410	-0.0119547055
##	eDNA_40	-0.021209297	3.019325e-03	-0.0009720696	0.0046949794
##	eDNA_41	-0.016778265	-4.673119e-02	0.0411543850	-0.0135517766
##	eDNA_42	0.050583194	-2.527828e-02	-0.0025768266	0.0235775903
##	eDNA_43	0.020856172	1.464289e-02	-0.0103891609	0.0131672443
##	eDNA_44	-0.005429068	-5.949592e-03	0.0172307806	0.0352623418
##	eDNA_45	-0.005050847	2.848692e-03	-0.0068455581	0.0074661650
##	eDNA_46	0.018039550	-6.061671e-03	0.0164227574	0.0351859382
##	eDNA_47	-0.036889539	-1.457880e-02	0.0041773152	-0.0031623639
##	eDNA_48	-0.076918032	-1.052199e-02	0.0040022590	0.0001543353
##	eDNA_49	0.034055620	-2.007944e-02	0.0024526508	-0.0198073878
##	eDNA_50	-0.016197872	3.128045e-02	-0.0069956224	0.0519447672
##	eDNA_51	-0.037418987	-6.627373e-03	0.0204329234	-0.0319678492
##	eDNA_52	0.030819458	2.305175e-03	0.0183681417	-0.0254556754
##	eDNA_53	-0.024836825	-9.155484e-04	0.0081023596	0.0045598276
##	eDNA_54	-0.018624609	7.263762e-03	-0.0127472657	-0.0215079850
##	eDNA_55	0.024043836	-7.662799e-05	0.0344316338	0.0332837763
##	eDNA_56	-0.050779271	-4.591402e-03	0.0056484865	0.0033987346
##	eDNA_57	-0.023513763	1.454861e-02	-0.0054345510	0.0341442765
##	eDNA_58	-0.019880128	4.132942e-02	-0.0103938205	0.0446588687
##		PCoA14	PCoA15	PCoA16	PCoA17
##	eDNA_01	0.0303783157	1.671978e-03	-1.365818e-02	0.0079074975
##	eDNA_02	-0.0037273997	1.686180e-02	-3.566190e-03	0.0021587737
##	eDNA_03	0.0320913404	-1.583357e-02	1.275687e-02	0.0216122082
##	eDNA_04	-0.0137822141	2.073378e-02	9.745178e-03	-0.0084722679
##	eDNA_05	-0.0070190395	3.669966e-03	6.141829e-03	0.0021387138
##	eDNA_06	-0.0152223091	-7.075294e-03	4.085037e-02	-0.0387231939
##	eDNA_07	0.0182663388	2.277032e-04	4.652688e-03	0.0028862241
##	eDNA_08	-0.0028007672	4.917110e-03	4.172632e-03	0.0060222743
##	eDNA_09	0.0120969579	1.246926e-02	1.761961e-02	0.0050863652

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## eDNA_10 0.0045415537 -3.541527e-03 7.752090e-03 0.0016864592
## eDNA_11 -0.0025885802 -8.133641e-03 7.020893e-02 0.0169344185
## eDNA_12 -0.0573323749 2.711996e-02 -1.492742e-02 0.0193567099
## eDNA_13 0.0086359605 7.186226e-03 -8.545110e-03 0.0276488375
## eDNA_14 -0.0056447818 3.866801e-03 -1.916339e-02 -0.0229622695
## eDNA_15 0.0059632082 1.547086e-03 -2.388273e-02 0.0272413402
## eDNA_16 -0.0001654146 -3.249077e-03 -2.196808e-03 0.0290236866
## eDNA_17 -0.0102194529 3.382606e-02 -7.789082e-03 -0.0101093054
## eDNA_18 -0.0237280131 -2.949708e-02 -5.712462e-03 -0.0162720650
## eDNA_19 -0.0496065294 -4.468569e-03 2.471389e-03 0.0118236040
## eDNA_20 -0.0284776381 -1.269781e-02 -2.194183e-02 -0.0165757847
## eDNA_21 0.0169730027 1.318495e-02 1.299468e-02 -0.0159515412
## eDNA_22 -0.0274803080 -6.045582e-02 -9.523255e-03 0.0087545229
## eDNA_23 -0.0230329017 -1.228629e-02 8.485747e-03 0.0033510692
## eDNA_24 -0.0274048962 -1.096271e-02 -3.151843e-03 -0.0017659857
## eDNA_25 -0.0013155961 -1.531829e-04 9.503385e-03 -0.0093841995
## eDNA_26 -0.0011677689 9.168207e-05 -7.089620e-03 -0.0011159685
## eDNA_27 -0.0051513020 -2.484128e-03 -6.559343e-03 -0.0002200829
## eDNA_28 -0.0104231598 -3.012915e-02 3.379169e-03 -0.0228405261
## eDNA_29 0.0382586917 -1.442134e-03 1.087475e-04 -0.0226502736
## eDNA_30 0.0410670361 -7.858714e-03 -7.228448e-04 -0.0211135004
## eDNA_31 -0.0021352291 -1.934564e-04 1.556784e-03 -0.0016057074
## eDNA_32 0.0043792043 -8.239723e-04 -9.188319e-03 0.0133543996
## eDNA_33 -0.0054017869 -3.553079e-03 -7.640323e-03 0.0013066181
## eDNA_34 -0.0274551099 -6.121869e-02 -1.320612e-02 0.0091424739
## eDNA_35 -0.0022557695 -2.739000e-02 4.775526e-03 -0.0269493324
## eDNA_36 -0.0169733547 3.432246e-02 -5.462585e-03 -0.0205055915
## eDNA_37 0.0044485827 -2.762839e-03 1.785803e-02 0.0308950292
## eDNA_38 -0.0015991630 -4.880816e-03 4.522081e-03 0.0034348856
## eDNA_39 -0.0022347488 8.650674e-05 4.098895e-03 0.0042127638
## eDNA_40 -0.0025243646 5.529614e-03 -8.596153e-03 -0.0027437183
## eDNA_41 0.0441344288 -1.697817e-02 9.763337e-03 0.0075328157
## eDNA_42 -0.0140318596 1.536661e-02 1.929134e-03 -0.0096077053
## eDNA_43 -0.0005842972 -1.591869e-03 8.508352e-03 0.0348376893
## eDNA_44 0.0270019643 6.260450e-03 -1.972060e-02 0.0156871748
## eDNA_45 -0.0020623979 4.288653e-05 -4.380896e-03 -0.0008448824
## eDNA_46 0.0363098271 -1.369760e-03 -1.161504e-02 -0.0065786262
## eDNA_47 -0.0033095247 3.976071e-03 5.683443e-03 -0.0078727371
## eDNA_48 -0.0063407147 5.804487e-03 1.918775e-02 -0.0315169510
## eDNA_49 -0.0242241207 3.420331e-02 1.790055e-03 0.0138907333
## eDNA_50 -0.0029269475 2.170028e-02 4.704648e-03 -0.0054096505
## eDNA_51 0.0110321023 3.404355e-02 3.286056e-03 -0.0035609080
## eDNA_52 0.0523347076 -1.352883e-02 1.800880e-03 -0.0036673430
## eDNA_53 0.0051716911 5.908209e-03 -7.638026e-03 0.0129967591
## eDNA_54 0.0099813570 2.977533e-03 -6.551865e-02 -0.0152991949
## eDNA_55 0.0296972848 -3.591946e-03 -1.272628e-02 -0.0026596414
## eDNA_56 0.0025185117 -7.985169e-04 -1.248655e-06 -0.0010090626
## eDNA_57 0.0059798158 4.739107e-03 9.037542e-03 0.0086495390
## eDNA_58 -0.0109120467 2.661518e-02 4.778526e-03 -0.0015855700
## PCoA18 PCoA19 PCoA20 PCoA21
## eDNA_01 2.343733e-02 -0.0016219990 -2.075460e-03 -0.0095080179
## eDNA_02 -2.462970e-04 -0.0412676689 -1.391355e-02 -0.0064423192
## eDNA_03 8.797513e-03 -0.0082747558 9.834250e-04 0.0184752332
## eDNA_04 1.141968e-02 0.0077990153 2.408075e-02 0.0164570858

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## eDNA_05 8.378250e-03 0.0052841336 5.010824e-03 -0.0018388133
## eDNA_06 2.080348e-02 -0.0193867576 6.520018e-03 -0.0126131484
## eDNA_07 -3.148164e-03 0.0340838758 4.482734e-03 -0.0075566945
## eDNA_08 1.368196e-02 0.0038583258 5.681936e-03 -0.0010524834
## eDNA_09 -1.571690e-02 -0.0109321226 -2.012251e-04 0.0123058243
## eDNA_10 -6.782872e-03 0.0005915104 1.882494e-02 0.0057651483
## eDNA_11 7.812327e-03 0.0155099591 -1.470718e-02 -0.0198111137
## eDNA_12 3.201339e-02 0.0095841772 5.527588e-03 -0.0032502353
## eDNA_13 1.067939e-03 0.0087693774 -1.290157e-02 -0.0142860704
## eDNA_14 -1.074974e-04 -0.0124200299 -5.511094e-03 0.0210143382
## eDNA_15 -1.678517e-02 0.0166953649 -6.213998e-03 0.0082612778
## eDNA_16 -7.790701e-03 0.0199045211 -6.071998e-03 -0.0032820279
## eDNA_17 -2.456396e-02 0.0204652410 -6.499690e-03 -0.0315984797
## eDNA_18 -1.774435e-03 0.0027565179 -4.130075e-04 -0.0043049738
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## eDNA_20 6.157044e-04 0.0091012727 -9.776719e-03 -0.0130631347
## eDNA_21 -2.843812e-02 0.0018583178 -9.874528e-04 0.0009594615
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## eDNA_24 -3.103512e-06 0.0087753002 -1.028197e-03 0.0078198021
## eDNA_25 4.305669e-04 -0.0137174856 1.361474e-02 -0.0118344920
## eDNA_26 -1.608081e-03 0.0102547283 -1.014015e-02 0.0097423522
## eDNA_27 -5.043143e-03 -0.0008706450 -8.852743e-03 -0.0102472957
## eDNA_28 -3.919410e-02 0.0009981233 -4.012874e-03 0.0065082027
## eDNA_29 1.515574e-02 -0.0050400857 -6.498082e-03 -0.0149813554
## eDNA_30 1.315360e-02 -0.0075882279 -8.583462e-03 -0.0122142429
## eDNA_31 7.896579e-03 0.0117119378 -2.902257e-02 0.0153247818
## eDNA_32 -2.015502e-02 -0.0174652617 5.426229e-02 -0.0276304303
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## eDNA_34 1.031213e-02 0.0001521340 -1.119237e-03 -0.0147784043
## eDNA_35 -4.131580e-02 0.0031410685 -1.434791e-03 0.0080443180
## eDNA_36 1.550176e-02 0.0311181305 1.475837e-02 0.0004550892
## eDNA_37 -2.626548e-02 -0.0171530510 -8.023057e-04 0.0143839173
## eDNA_38 7.183772e-03 0.0012827505 8.338928e-04 -0.0013237950
## eDNA_39 2.188750e-03 -0.0006209452 7.757656e-04 -0.0011050591
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## eDNA_41 3.576791e-03 -0.0071726546 3.571776e-03 0.0172007819
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## eDNA_43 -1.693541e-02 -0.0151096950 -4.393710e-03 0.0039498525
## eDNA_44 1.299162e-02 -0.0071201413 -1.467228e-02 -0.0275488424
## eDNA_45 8.295406e-03 0.0175827884 -3.414523e-02 0.0199581784
## eDNA_46 7.733819e-03 0.0037327633 -2.569445e-03 -0.0101088848
## eDNA_47 4.980658e-03 -0.0051388617 1.595352e-03 0.0044575580
## eDNA_48 1.570027e-02 -0.0095094223 -2.238324e-02 0.0072410002
## eDNA_49 1.316711e-03 -0.0395622840 -2.369641e-02 -0.0130179155
## eDNA_50 -1.461524e-02 -0.0093265588 -6.555181e-03 0.0044621399
## eDNA_51 -2.061655e-02 0.0140813531 9.174259e-04 -0.0059894155
## eDNA_52 1.444564e-03 0.0186517721 9.242448e-03 0.0116193998
## eDNA_53 -3.913249e-03 0.0098355198 6.939224e-03 0.0166587146
## eDNA_54 -4.253365e-03 -0.0216810647 9.754456e-05 0.0187858999
## eDNA_55 1.353369e-02 0.0012075129 -2.700215e-04 -0.0054000951
## eDNA_56 -3.977470e-03 -0.0072263339 2.351892e-02 -0.0149068424
## eDNA_57 4.184143e-02 -0.0113969675 2.171421e-02 0.0322963850
## eDNA_58 -1.701426e-02 -0.0030740152 -9.150526e-03 -0.0009360887

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##		PCoA22	PCoA23	PCoA24	PCoA25
##	eDNA_01	0.0159572420	-0.0006136865	-0.0022640925	-1.108936e-02
##	eDNA_02	-0.0108884602	-0.0257334708	0.0039570570	-9.645022e-03
##	eDNA_03	0.0117892963	-0.0100458822	0.0056996241	-8.303572e-03
##	eDNA_04	-0.0224668377	0.0136126811	0.0047160613	-1.363016e-02
##	eDNA_05	0.0036403972	0.0039656147	-0.0073605405	7.062220e-03
##	eDNA_06	0.0144142199	-0.0034558324	0.0134453574	1.124698e-02
##	eDNA_07	0.0180530648	0.0144106986	0.0089424849	-9.012854e-03
##	eDNA_08	0.0015778040	-0.0022669753	-0.0100861626	3.249333e-03
##	eDNA_09	-0.0032327896	0.0039393392	0.0007745108	5.027365e-03
##	eDNA_10	-0.0131952043	-0.0214086647	-0.0002889249	2.277982e-03
##	eDNA_11	-0.0168848539	-0.0014849796	-0.0174613970	-5.597915e-03
##	eDNA_12	0.0059814008	0.0010437719	-0.0146808927	7.539283e-04
##	eDNA_13	-0.0026871714	0.0024920215	0.0053363259	5.578391e-03
##	eDNA_14	0.0041152608	0.0061377912	-0.0255473788	-1.115286e-02
##	eDNA_15	-0.0049449866	0.0043833712	-0.0013023370	-1.135673e-02
##	eDNA_16	-0.0088221574	-0.0036194663	0.0133649126	2.799609e-03
##	eDNA_17	-0.0059092399	0.0109210090	0.0125169239	-1.164282e-03
##	eDNA_18	-0.0051992488	0.0295734553	-0.0092717040	9.417521e-03
##	eDNA_19	-0.0014326827	0.0110767060	0.0055925567	1.264658e-02
##	eDNA_20	-0.0092153376	0.0259176755	0.0026709495	1.096466e-02
##	eDNA_21	-0.0065722866	-0.0093950299	0.0055080580	-6.681769e-03
##	eDNA_22	0.0086422865	-0.0174612937	-0.0019515624	-7.980177e-03
##	eDNA_23	0.0018310355	0.0048312149	-0.0038830618	1.858929e-07
##	eDNA_24	-0.0007427253	-0.0049746148	0.0033376828	2.298615e-03
##	eDNA_25	0.0131487705	0.0176525924	-0.0074473504	-3.300648e-02
##	eDNA_26	-0.0151896699	-0.0018471470	-0.0218128838	-2.301560e-03
##	eDNA_27	-0.0138687877	0.0090070789	0.0014237785	-1.213092e-03
##	eDNA_28	-0.0122628005	-0.0122026921	0.0093308871	-1.166859e-02
##	eDNA_29	-0.0111027155	0.0023288586	-0.0009316015	5.166156e-04
##	eDNA_30	-0.0077457843	0.0015524306	-0.0039636690	7.934622e-03
##	eDNA_31	-0.0038625825	-0.0082246081	0.0077451907	1.256227e-02
##	eDNA_32	-0.0130531012	-0.0035318525	-0.0116836644	9.287602e-03
##	eDNA_33	-0.0107381210	0.0027431919	-0.0008873684	2.676446e-03
##	eDNA_34	0.0107232189	-0.0174132350	-0.0031122592	-7.203126e-03
##	eDNA_35	-0.0143416278	-0.0037954968	0.0081077568	-1.991033e-03
##	eDNA_36	0.0002169072	-0.0297903534	0.0009700752	-5.954018e-03
##	eDNA_37	0.0195240004	0.0101952079	0.0065361217	-5.867785e-03
##	eDNA_38	0.0010870242	0.0015363471	-0.0008319875	1.586098e-03
##	eDNA_39	-0.0018227355	0.0002972303	0.0029423607	3.186199e-04
##	eDNA_40	0.0344463874	0.0051070385	0.0389348850	6.947238e-03
##	eDNA_41	0.0085045243	-0.0029208336	-0.0167699332	1.921915e-02
##	eDNA_42	0.0178345968	-0.0196178526	0.0045779628	-4.682263e-03
##	eDNA_43	0.0104440170	0.0198980383	0.0040399873	-1.882698e-03
##	eDNA_44	-0.0063099733	-0.0017233928	0.0133564533	-4.222591e-03
##	eDNA_45	0.0002728595	-0.0080285148	0.0131623253	1.251212e-02
##	eDNA_46	0.0049269602	0.0083396517	-0.0024132691	-4.221016e-04
##	eDNA_47	0.0040104788	-0.0118670444	0.0057808592	1.208790e-02
##	eDNA_48	0.0195967129	0.0153915244	-0.0008172273	-1.856188e-02
##	eDNA_49	-0.0167312154	-0.0133937030	0.0040347387	-4.798884e-03
##	eDNA_50	0.0222871407	-0.0017672454	-0.0154598302	1.800763e-02
##	eDNA_51	0.0042094466	-0.0007272151	-0.0070024629	-5.101804e-03
##	eDNA_52	-0.0063515272	-0.0004492586	-0.0014082963	6.396795e-03
##	eDNA_53	0.0049305013	-0.0056562597	-0.0120577733	-1.078192e-02

## eDNA_54	-0.0012059557	-0.0007872146	-0.0015997457	-1.166162e-03
## eDNA_55	0.0078778358	0.0011157286	-0.0033127791	1.627215e-03
## eDNA_56	-0.0066523357	0.0003565591	-0.0049973532	1.750944e-02
## eDNA_57	-0.0370121234	0.0176851853	0.0196256073	-1.841584e-03
## eDNA_58	0.0204016482	-0.0013081982	-0.0158239858	1.576913e-02
##	PCoA26	PCoA27	PCoA28	PCoA29
## eDNA_01	1.752012e-02	2.328511e-03	0.0010068059	0.0024429797
## eDNA_02	5.938828e-06	9.967441e-03	-0.0033277364	0.0002462371
## eDNA_03	2.058964e-02	-8.008062e-03	0.0182083004	-0.0011379620
## eDNA_04	9.569906e-03	2.006063e-02	-0.0184327442	0.0077910817
## eDNA_05	5.224104e-03	-9.606583e-04	0.0080626657	0.0039031963
## eDNA_06	3.825808e-03	6.762535e-05	-0.0027216490	-0.0021177059
## eDNA_07	-5.265030e-03	-2.212223e-02	-0.0126656927	-0.0097128658
## eDNA_08	8.177578e-04	5.924965e-03	-0.0035796290	0.0011977772
## eDNA_09	-3.908768e-03	6.483779e-04	0.0011633468	0.0038277422
## eDNA_10	1.243135e-02	-7.771283e-03	0.0026644004	0.0013460765
## eDNA_11	4.961814e-03	-6.421282e-03	-0.0040614142	0.0006807783
## eDNA_12	3.380410e-03	8.106056e-03	-0.0067340278	0.0002831559
## eDNA_13	-1.471891e-02	8.941975e-03	-0.0044530143	-0.0155672631
## eDNA_14	1.054698e-02	-1.840085e-02	-0.0130361579	-0.0015717248
## eDNA_15	-6.503153e-03	-3.514896e-03	0.0016390638	0.0031096349
## eDNA_16	-8.079173e-03	2.163185e-02	0.0133586515	0.0034133975
## eDNA_17	1.669585e-03	-1.236705e-02	0.0060139431	0.0035577721
## eDNA_18	-1.676376e-04	1.170597e-03	-0.0007482842	0.0086238815
## eDNA_19	-8.468964e-03	-3.379651e-03	0.0105808572	0.0006129789
## eDNA_20	1.369447e-02	5.917145e-03	0.0065747392	-0.0100239082
## eDNA_21	1.106596e-03	9.064093e-04	-0.0011669006	-0.0026606441
## eDNA_22	-2.767711e-03	-1.300005e-03	-0.0036706874	-0.0007937139
## eDNA_23	1.268844e-02	1.267609e-02	0.0011660451	-0.0229636995
## eDNA_24	-1.602929e-02	-1.195389e-02	0.0013667615	0.0216525485
## eDNA_25	-2.022055e-02	7.698624e-03	0.0150515986	-0.0019986020
## eDNA_26	-6.126328e-03	-2.175581e-03	0.0086970023	-0.0057731325
## eDNA_27	5.471085e-03	1.865083e-03	-0.0016236299	0.0034951576
## eDNA_28	8.728334e-03	3.499331e-03	0.0009812832	0.0020388207
## eDNA_29	2.387338e-03	3.896724e-03	0.0016341606	-0.0011884945
## eDNA_30	-3.350184e-03	2.253776e-03	0.0031959945	0.0011329395
## eDNA_31	7.796744e-03	1.960280e-03	-0.0008120906	0.0036305769
## eDNA_32	1.903631e-03	-4.111450e-03	-0.0009602887	-0.0014552767
## eDNA_33	-3.372602e-03	1.662988e-03	0.0047861035	0.0018833314
## eDNA_34	8.450672e-04	1.918003e-03	-0.0029461940	-0.0003055332
## eDNA_35	-1.272572e-02	-3.432591e-03	-0.0030200214	-0.0044003859
## eDNA_36	-4.224703e-03	-8.295559e-03	0.0065215682	-0.0003328023
## eDNA_37	6.133158e-03	-4.374005e-03	-0.0039618002	-0.0040661467
## eDNA_38	1.024041e-03	-1.690954e-03	0.0038809539	-0.0049636871
## eDNA_39	-8.526226e-03	-1.594721e-03	-0.0063832387	0.0019176390
## eDNA_40	1.862992e-03	6.738803e-03	-0.0032853288	0.0060947441
## eDNA_41	-2.730118e-02	4.523993e-03	-0.0140885596	0.0023027462
## eDNA_42	-8.723947e-03	-9.230941e-04	-0.0015477368	-0.0033891283
## eDNA_43	5.783433e-03	-4.354852e-03	-0.0108667861	0.0084576552
## eDNA_44	3.080980e-04	-1.110979e-02	0.0028375591	0.0002445803
## eDNA_45	9.528110e-03	-3.348358e-03	-0.0116151768	-0.0002216989
## eDNA_46	-8.544899e-04	-2.585118e-03	-0.0049361480	-0.0057794316
## eDNA_47	-1.030426e-02	1.939705e-03	-0.0043550816	-0.0137172586
## eDNA_48	-7.132961e-03	5.217061e-03	0.0047631431	0.0060639147

## eDNA_49	-6.042068e-03	-3.549486e-03	-0.0057197287	0.0021401804
## eDNA_50	5.459540e-03	-5.265957e-04	0.0176101554	0.0010477156
## eDNA_51	5.193353e-03	1.431933e-02	-0.0097710462	-0.0020935335
## eDNA_52	3.350427e-03	1.494398e-02	0.0061416423	0.0096586716
## eDNA_53	4.554647e-03	2.527983e-03	0.0047894829	0.0061455124
## eDNA_54	-1.111135e-03	-6.150516e-04	-0.0015287240	-0.0017587767
## eDNA_55	-1.740288e-03	2.539289e-03	-0.0022491802	0.0022805791
## eDNA_56	1.045570e-02	-7.485193e-03	-0.0042162760	0.0072105716
## eDNA_57	-7.307375e-03	-1.571564e-02	0.0087081607	-0.0081397011
## eDNA_58	-3.845950e-03	-3.764722e-03	0.0070805854	-0.0023014976
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## eDNA_02	4.030028e-04	-0.0037923989	3.231901e-03	0.0046554555
## eDNA_03	9.742965e-04	0.0102378175	4.054576e-03	0.0022791850
## eDNA_04	-6.479899e-03	0.0107111184	1.554957e-02	0.0038172240
## eDNA_05	6.269572e-05	-0.0037541029	2.824906e-03	0.0023658219
## eDNA_06	3.708334e-04	0.0021281425	-7.094086e-04	-0.0051655740
## eDNA_07	-1.253515e-03	0.0102844641	6.877158e-04	0.0038820161
## eDNA_08	-1.143687e-03	-0.0007324861	-6.443455e-03	-0.0011309750
## eDNA_09	5.945540e-03	0.0009064582	-9.496556e-04	0.0018642604
## eDNA_10	-1.043215e-02	-0.0010902520	-6.453611e-03	-0.0062226497
## eDNA_11	3.328971e-03	-0.0072727725	2.158488e-03	0.0011898659
## eDNA_12	1.855979e-04	-0.0083654732	-6.747968e-03	-0.0057217745
## eDNA_13	1.448972e-02	0.0022320102	5.986507e-03	0.0057663896
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## eDNA_16	1.673597e-03	0.0014857583	-5.800329e-03	0.0083328718
## eDNA_17	-7.839851e-03	-0.0200870659	4.289930e-03	0.0047000682
## eDNA_18	-8.565085e-03	0.0083655107	-2.685042e-03	0.0041733317
## eDNA_19	-3.117564e-03	0.0085433950	1.109840e-02	0.0050215430
## eDNA_20	1.378171e-02	-0.0056803480	3.333029e-03	-0.0054622896
## eDNA_21	-1.400003e-03	-0.0032934524	2.452721e-03	-0.0013980905
## eDNA_22	-1.187293e-03	0.0007320226	-2.595828e-03	0.0034425440
## eDNA_23	-1.532173e-02	-0.0031592461	-6.292406e-03	0.0053714968
## eDNA_24	1.574340e-02	0.0018368588	6.520966e-03	-0.0046545708
## eDNA_25	-1.579193e-03	-0.0005281669	-7.126103e-04	-0.0035906414
## eDNA_26	-3.816155e-03	0.0105710861	3.774291e-03	-0.0028705308
## eDNA_27	-8.489779e-04	0.0053425207	-8.223348e-03	-0.0049251921
## eDNA_28	-4.873900e-04	-0.0062046328	6.186081e-03	-0.0031488570
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## eDNA_30	-2.024172e-03	-0.0010074977	-8.319373e-04	0.0010094965
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## eDNA_33	-7.185677e-03	0.0037649189	1.421026e-03	0.0054096631
## eDNA_34	2.525524e-03	-0.0047316404	-1.671494e-04	-0.0048638836
## eDNA_35	8.069926e-03	0.0069947714	-1.252485e-02	0.0030800507
## eDNA_36	-2.246951e-03	0.0085350472	-5.610598e-03	-0.0035796545
## eDNA_37	7.259065e-03	-0.0048971453	6.778469e-03	-0.0028521457
## eDNA_38	-1.466710e-03	0.0031625973	4.188345e-03	-0.0021673795
## eDNA_39	2.697266e-04	0.0008316986	-3.181576e-03	0.0005054393
## eDNA_40	-1.954772e-04	-0.0009447943	-6.537545e-03	0.0042339200
## eDNA_41	-8.472749e-03	-0.0047653440	7.912422e-04	0.0004059687
## eDNA_42	-2.871946e-03	-0.0049690298	4.413905e-03	0.0060210382
## eDNA_43	-7.035779e-03	0.0009447395	-9.229848e-03	-0.0071467147


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## eDNA_21 -3.056246e-03 -1.089145e-03 1.045181e-03 -7.687768e-04
## eDNA_22 6.997829e-03 -2.246062e-03 1.921202e-03 1.022887e-04
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## eDNA_24 -1.444391e-03 -7.214795e-04 1.795078e-04 -2.323557e-05
## eDNA_25 -1.334954e-03 2.247184e-04 3.049355e-04 -1.055351e-04
## eDNA_26 9.893725e-04 -1.764260e-03 1.103277e-03 1.124284e-03
## eDNA_27 5.988875e-05 8.872954e-04 -2.934573e-04 -1.113863e-04
## eDNA_28 -1.259158e-03 -9.625835e-04 7.473141e-04 6.825772e-04

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## eDNA_29 -3.506553e-03 -5.639291e-03 5.085772e-04 8.673078e-05
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## eDNA_31 3.513960e-03 -3.560566e-04 -7.452571e-04 -4.592162e-05
## eDNA_32 7.384105e-05 9.920519e-05 2.862984e-05 -8.663514e-05
## eDNA_33 -6.555460e-03 -1.076448e-03 1.540099e-03 4.240232e-05
## eDNA_34 -5.738621e-03 2.312931e-03 -2.099916e-03 8.308822e-06
## eDNA_35 1.226197e-03 4.356128e-04 -3.432245e-04 -2.293663e-05
## eDNA_36 -3.420226e-03 2.304702e-03 -2.711155e-04 -2.126149e-03
## eDNA_37 3.058308e-03 1.641162e-04 9.013143e-04 -1.085378e-03
## eDNA_38 1.862240e-03 -5.670748e-03 -3.541469e-03 -2.277326e-03
## eDNA_39 3.023879e-03 2.509358e-03 1.927048e-03 -1.050548e-03
## eDNA_40 -3.326348e-04 -1.886481e-04 1.196406e-04 -1.889347e-04
## eDNA_41 -2.705997e-03 -9.629111e-04 -4.130043e-05 -8.460116e-07
## eDNA_42 2.228617e-03 -1.810703e-03 2.957370e-05 2.633610e-03
## eDNA_43 -5.500647e-03 -9.465485e-04 -2.551451e-04 3.853459e-04
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## eDNA_45 -2.346566e-03 -1.188410e-05 4.483248e-04 2.428352e-04
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## eDNA_49 2.428275e-03 -1.663547e-03 1.466941e-03 -5.782592e-05
## eDNA_50 -1.961537e-04 -1.848789e-04 -1.436240e-04 -4.708494e-04
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## eDNA_03 5.160123e-04 1.349497e-03 -4.376440e-03 -1.107131e-03
## eDNA_04 5.145859e-04 -4.363245e-04 -3.944985e-03 -1.684098e-03
## eDNA_05 -6.088897e-03 1.680982e-03 -2.851872e-04 -4.930715e-03
## eDNA_06 3.940332e-05 -2.477605e-04 -1.075096e-03 -1.012437e-03
## eDNA_07 1.422394e-04 -9.085622e-04 2.693273e-03 -1.587581e-04
## eDNA_08 5.226608e-03 2.775946e-03 -4.122739e-04 5.246493e-03
## eDNA_09 4.114820e-04 -2.018829e-03 -2.870575e-03 -5.620106e-03
## eDNA_10 -4.020291e-04 -6.244517e-03 2.794069e-03 -6.103337e-03
## eDNA_11 3.728894e-05 -1.592672e-03 6.605398e-04 4.147875e-03
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## eDNA_18 -1.698485e-03 -3.968698e-03 2.169202e-03 2.195191e-03
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## eDNA_20 1.704292e-03 -1.188463e-03 -5.228731e-04 4.361153e-05
## eDNA_21 1.936506e-04 1.948327e-03 3.448928e-03 6.806011e-03
## eDNA_22 9.522456e-05 -1.014047e-04 -9.947457e-04 -5.075632e-04
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## eDNA_27 -9.043683e-04 5.243624e-03 6.883609e-03 7.334056e-05
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## eDNA_51 -1.076533e-03 1.229146e-03 6.132123e-03 -4.625175e-03
## eDNA_52 2.960430e-04 2.256179e-03 -2.287604e-03 -1.117150e-03
## eDNA_53 -5.835389e-04 -1.690085e-03 -7.905091e-04 5.439243e-03
## eDNA_54 -2.946654e-04 -2.277360e-03 9.680147e-05 3.915106e-03
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## eDNA_56 3.184968e-04 5.178729e-03 -1.486811e-03 4.248754e-03
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## eDNA_02 -0.0004453259 -1.863236e-03 -0.0025815996 0.0001761155
## eDNA_03 0.0055891449 1.316716e-03 0.0022653086 0.0054786454
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## eDNA_11 0.0019284370 2.419292e-03 0.0016675928 0.0034402425
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## eDNA_13 -0.0008075542 -2.401383e-03 0.0111087302 0.0046907779
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## eDNA_15 -0.0024783053 4.500023e-03 0.0029909960 0.0098736406
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## eDNA_17 -0.0007185688 -4.648190e-03 0.0053577594 -0.0044275296
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## eDNA_19	0.0033021237	-6.119498e-03	0.0044120522	-0.0022958374
## eDNA_20	0.0049535915	2.389095e-03	-0.0095120165	-0.0029397272
## eDNA_21	0.0012945202	3.896907e-05	-0.0023697833	-0.0050201766
## eDNA_22	0.0002242544	-2.566634e-03	0.0007839037	-0.0005480711
## eDNA_23	-0.0016250305	3.649793e-03	-0.0002200010	0.0025530880
## eDNA_24	0.0020274056	8.651351e-04	-0.0001718806	-0.0021557677
## eDNA_25	0.0014399104	3.096248e-04	-0.0025399585	-0.0011703928
## eDNA_26	-0.0037312414	-5.582648e-03	-0.0030866177	-0.0066884221
## eDNA_27	0.0089526116	9.984763e-03	0.0126035554	0.0020272563
## eDNA_28	-0.0006440640	-2.697559e-03	-0.0058273598	0.0160643089
## eDNA_29	0.0033154676	-1.759392e-03	0.0007159108	-0.0005203163
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## eDNA_33	-0.0102025149	5.286017e-03	0.0017688200	0.0064620733
## eDNA_34	-0.0014705255	3.478616e-04	-0.0004147925	-0.0022074341
## eDNA_35	-0.0038332282	6.996281e-03	0.0034833008	-0.0113848821
## eDNA_36	0.0022169025	6.498411e-04	-0.0024143192	0.0017935604
## eDNA_37	0.0018571143	5.507039e-03	0.0005242782	-0.0124904482
## eDNA_38	-0.0008743895	3.190427e-03	0.0045537189	0.0015956848
## eDNA_39	0.0084321947	-5.383755e-03	0.0015301745	0.0059203927
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## eDNA_41	0.0013354563	5.943014e-03	-0.0051832142	-0.0034801306
## eDNA_42	-0.0009887662	5.421829e-03	0.0087227855	0.0039240817
## eDNA_43	-0.0021442995	-8.151291e-03	-0.0046339728	0.0017335749
## eDNA_44	0.0009385614	2.894936e-03	-0.0074009219	-0.0132307261
## eDNA_45	-0.0008434932	2.680406e-03	-0.0003807837	0.0005499788
## eDNA_46	0.0124994358	-3.012059e-03	-0.0005578730	0.0078367676
## eDNA_47	-0.0007222173	-9.489152e-03	-0.0046159437	-0.0030765733
## eDNA_48	0.0001116027	1.292733e-03	-0.0027783494	-0.0012202267
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## eDNA_50	-0.0013740923	1.398432e-03	-0.0019702925	0.0031072125
## eDNA_51	-0.0076260215	-4.468924e-03	-0.0013849182	-0.0008959914
## eDNA_52	0.0016341830	-4.738608e-04	0.0089828927	-0.0065619506
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## eDNA_54	0.0024596902	3.190311e-03	-0.0008691261	0.0010977561
## eDNA_55	-0.0053011891	-4.838000e-03	-0.0001610223	0.0072515174
## eDNA_56	-0.0007885689	6.481633e-03	-0.0123553955	0.0054264278
## eDNA_57	-0.0046915142	-1.518844e-03	-0.0006614507	0.0022433421
## eDNA_58	0.0051412611	-5.649299e-04	0.0048308382	0.0005884309
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## eDNA_04	-1.335585e-03	-2.482543e-03	-1.202101e-03	-2.594097e-04
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## eDNA_10	-6.834679e-03	-1.617866e-02	-4.408381e-03	4.724248e-03
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## eDNA_14 8.118780e-03 7.379740e-03 7.240759e-03 2.513190e-02
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## eDNA_32 -3.118456e-03 4.183345e-03 2.910412e-02 -1.620845e-02
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## eDNA_36 5.069141e-03 7.182917e-03 1.256317e-03 -4.573286e-04
## eDNA_37 8.234949e-04 -3.931699e-03 -6.441375e-03 -9.359422e-03
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## eDNA_50 -1.121844e-03 -1.131721e-03 1.107762e-03 2.037717e-03
## eDNA_51 3.308745e-03 3.837433e-03 -2.252194e-03 -2.464624e-03
## eDNA_52 4.785175e-03 1.523459e-02 -4.812032e-03 3.928125e-05
## eDNA_53 -2.091906e-02 6.330411e-03 -9.824647e-03 -9.978128e-03
## eDNA_54 1.555507e-02 -4.488680e-03 -1.076388e-02 -2.752978e-02
## eDNA_55 2.282552e-06 -1.492797e-02 9.738000e-05 -2.784300e-03
## eDNA_56 7.033640e-03 -3.064485e-03 -2.199480e-02 9.529459e-03
## eDNA_57 9.588355e-04 6.394657e-04 7.266899e-04 -1.851723e-03
## eDNA_58 2.968322e-03 2.246524e-04 1.062543e-04 7.173720e-04

```

```
disp.group$centroids
```

```

##          PCoA1      PCoA2      PCoA3      PCoA4      PCoA5
## fecesC  0.3878748 -0.05983260 -0.00827372 -0.002572693 4.520720e-02
## fecesE  0.3846140 -0.05281179 -0.01768682 -0.003620721 2.563516e-02
## sedC    -0.1621571 0.01383030 -0.07574180 0.129742326 1.774716e-02
## sedE    -0.1536365 0.02393042 -0.08484757 0.114095079 1.808900e-02
## soilC   -0.1310322 0.09076200 -0.08660862 -0.124608286 8.663795e-05
## soile   -0.1272992 0.09265998 -0.08386905 -0.139828718 -2.432636e-03

```

```

## waterC -0.1520673 -0.08814663 0.13910634 -0.019183111 -3.623969e-03
## waterE -0.1477446 -0.09197941 0.14612311 -0.021572728 -4.926486e-03
##          PCoA6          PCoA7          PCoA8          PCoA9          PCoA10
## fecesC 0.0098589442 0.016397777 -9.495006e-04 -0.002803429 -0.0005967342
## fecesE -0.0009809179 -0.002832237 -4.382681e-03 -0.002298915 -0.0032227201
## sedC 0.0019230882 -0.018076518 -7.021642e-05 -0.012981242 -0.0190403798
## sedE -0.0016667448 0.001182872 -8.991959e-03 -0.006189935 -0.0097486094
## soilC 0.0207659427 0.014086669 -1.810293e-03 -0.007799465 0.0113697682
## soile 0.0203669867 0.021771870 3.555331e-04 -0.011077239 0.0238205868
## waterC -0.0067040487 -0.013388065 3.212156e-03 0.005526618 0.0001780628
## waterE -0.0113894267 -0.004726205 3.909079e-03 0.001114462 0.0064050182
##          PCoA11          PCoA12          PCoA13          PCoA14          PCoA15
## fecesC 0.003183113 -0.003963531 0.0020243773 0.002531363 -0.0005298405
## fecesE -0.003608571 0.002141896 0.0003440503 -0.002857064 0.0013013718
## sedC 0.014102263 -0.005045631 0.0031010032 -0.010395640 0.0137423428
## sedE -0.005464704 0.006464936 0.0060758763 0.009384677 -0.0088932577
## soilC 0.005083942 -0.005300623 0.0029716511 -0.002363583 0.0044321589
## soile 0.001742570 -0.004551787 0.0082913536 0.001480640 -0.0003408614
## waterC -0.006252494 0.002339251 0.0061153258 0.003829897 -0.0030460971
## waterE -0.004545598 0.006421805 0.0064820664 0.004093985 -0.0018753856
##          PCoA16          PCoA17          PCoA18          PCoA19
## fecesC -0.008224039 0.0084617606 -0.0053131147 0.0056286235
## fecesE 0.007316486 -0.0093475111 0.0040293886 -0.0069442978
## sedC 0.001547744 -0.0005069016 -0.0050878748 0.0032158632
## sedE 0.002813347 0.0061489241 0.0091063205 -0.0099944064
## soilC -0.002956428 0.0001524823 0.0007159359 0.0027013102
## soile 0.004296706 0.0023177234 -0.0043380515 -0.0002855788
## waterC -0.001176543 0.0004675681 -0.0026140778 -0.0015345967
## waterE -0.004077742 -0.0007148156 0.0050376444 0.0023202964
##          PCoA20          PCoA21          PCoA22          PCoA23
## fecesC 1.871564e-03 0.003013167 -0.0044011219 -0.0046322545
## fecesE -5.434625e-04 -0.003286834 0.0035344766 0.0047069736
## sedC -2.851750e-04 -0.005990463 -0.0003391233 0.0006712215
## sedE 2.273680e-03 0.009031973 -0.0010661459 -0.0051716875
## soilC -1.869136e-03 -0.004531683 -0.0027033546 0.0040171382
## soile 4.693306e-03 0.007112144 0.0090301300 -0.0002756054
## waterC -1.642061e-03 -0.001620279 -0.0026177886 0.0021203775
## waterE 6.578565e-05 -0.002305885 0.0037932998 -0.0016959900
##          PCoA24          PCoA25          PCoA26          PCoA27
## fecesC 0.0046528856 0.0044767226 0.0032243942 0.0001289074
## fecesE -0.0068589271 -0.0052939926 -0.0026994272 -0.0016560634
## sedC -0.0004938347 -0.0029892349 0.0005104500 0.0012973139
## sedE -0.0001994615 0.0018385273 -0.0008720933 -0.0006985109
## soilC 0.0015694339 -0.0002737461 0.0029289752 -0.0012816005
## soile 0.0003684005 -0.0007059869 -0.0013453850 -0.0004637273
## waterC 0.0024004003 0.0011297401 -0.0035325442 -0.0016815309
## waterE -0.0015916923 -0.0027233255 0.0046684005 0.0011911673
##          PCoA28          PCoA29          PCoA30          PCoA31
## fecesC 0.0007728879 0.0032624291 -0.003361671 0.0003789649
## fecesE -0.0014147490 -0.0029653558 0.003188719 -0.0006940785
## sedC -0.0051082562 0.0007276507 -0.001683197 0.0019425900
## sedE 0.0046446094 -0.0005163410 0.001132926 -0.0026507820
## soilC -0.0002875452 0.0016014629 -0.001186218 0.0032208943
## soile -0.0009291542 -0.0014376239 0.001516174 -0.0027138314

```


##	waterC	-0.0010066063	-0.0022936813	-0.001829522	0.0030269221
##	waterE	0.0007992513	0.0021226193	0.002051291	-0.0024826671
##		PCoA32	PCoA33	PCoA34	PCoA35
##	fecesC	-0.0013059273	-0.001943739	-0.0005138131	-0.0010967522
##	fecesE	0.0013108361	0.001941544	0.0001957928	0.0010952512
##	sedC	0.0012836986	0.001619802	-0.0020419565	0.0003208542
##	sedE	-0.0006910528	-0.001585903	0.0020532735	-0.0007074934
##	soilC	-0.0040256369	-0.002804686	-0.0013101646	0.0031776381
##	soile	0.0031338563	0.001988469	0.0011337386	-0.0022734283
##	waterC	-0.0030216159	0.001318552	-0.0018196958	-0.0012894450
##	waterE	0.0031054014	-0.001198342	0.0012177101	0.0015232918
##		PCoA36	PCoA37	PCoA38	PCoA39
##	fecesC	0.0004016044	-0.0006053044	-1.470032e-04	8.496251e-04
##	fecesE	-0.0003741919	0.0003595004	2.107090e-04	-9.186449e-04
##	sedC	-0.0010018429	-0.0008153850	-4.312031e-04	1.167712e-03
##	sedE	0.0006294952	0.0007610820	6.198902e-04	-1.258176e-03
##	soilC	0.0019571170	-0.0011021402	1.194394e-03	-1.289248e-03
##	soile	-0.0019165877	0.0006419656	-1.037696e-03	1.043581e-03
##	waterC	0.0018718789	-0.0011305950	6.683773e-05	-1.935004e-05
##	waterE	-0.0016902142	0.0010388946	-1.089476e-04	-1.002142e-04
##		PCoA40	PCoA41	PCoA42	PCoA43
##	fecesC	1.080751e-05	0.0004003521	0.0008716828	-0.0002607005
##	fecesE	-3.741882e-05	-0.0003551539	-0.0008217638	0.0002395858
##	sedC	8.338249e-04	-0.0014510852	0.0014757244	-0.0003460347
##	sedE	-5.563145e-04	0.0016365865	-0.0016387249	0.0003404978
##	soilC	-6.576134e-04	-0.0014593249	-0.0036800377	-0.0006346475
##	soile	7.302543e-04	0.0013620383	0.0031691596	0.0005411815
##	waterC	-1.379796e-03	0.0017412924	0.0005010704	0.0008559772
##	waterE	1.453107e-03	-0.0014752124	-0.0002709734	-0.0007859971
##		PCoA44	PCoA45	PCoA46	PCoA47
##	fecesC	-1.571741e-04	3.525010e-05	-2.939244e-04	0.0000696631
##	fecesE	1.512704e-04	-3.646330e-05	1.531149e-04	0.0004055686
##	sedC	4.323787e-04	7.442779e-05	-7.709223e-05	-0.0001688911
##	sedE	-5.348709e-04	-1.301651e-05	1.090552e-04	0.0008936589
##	soilC	1.924022e-04	-4.424521e-04	-2.555528e-04	0.0001607612
##	soile	-5.122269e-05	3.135134e-04	1.944972e-04	-0.0009056170
##	waterC	-6.862449e-05	4.640367e-04	7.697569e-04	0.0007055191
##	waterE	1.107679e-04	-5.111751e-04	-1.345287e-04	-0.0008686371
##		PCoA48	PCoA49	PCoA50	PCoA51
##	fecesC	0.0001470747	3.473135e-04	-0.0004111759	-0.0005314460
##	fecesE	-0.0000429035	-7.783467e-04	-0.0001489254	-0.0008428236
##	sedC	-0.0013375842	-3.095195e-04	0.0003027373	-0.0014969961
##	sedE	0.0014127665	-6.589530e-06	-0.0008708655	0.0005784394
##	soilC	-0.0009371561	1.544345e-03	0.0014810810	0.0006483433
##	soile	0.0010455551	-9.707287e-04	-0.0015220056	-0.0006286484
##	waterC	-0.0004774724	1.274815e-03	0.0019980741	-0.0013172151
##	waterE	0.0004496811	-1.219755e-03	-0.0006554063	0.0019986355
##		PCoA52	PCoA53	PCoA54	PCoA55
##	fecesC	-0.0010244745	-6.765172e-05	1.985580e-04	-0.0025363545
##	fecesE	0.0019230797	-1.467412e-04	-7.821703e-04	0.0039068044
##	sedC	0.0007216906	-5.083223e-04	1.109365e-03	-0.0018822706
##	sedE	-0.0006868360	1.677050e-04	-1.305983e-03	0.0002159500
##	soilC	-0.0008457973	1.114935e-03	-6.922759e-05	0.0010052332
##	soile	0.0016351409	1.514024e-05	1.152270e-03	0.0001913701

```

## waterC 0.0008151452 8.764571e-05 1.050853e-06 0.0008271082
## waterE -0.0003077652 -4.924643e-04 -3.484497e-04 0.0009596914
##          PCoA56          PCoA57
## fecesC -0.0027385306 0.0013860052
## fecesE 0.0010564733 -0.0026147247
## sedC -0.0012657503 -0.0022066404
## sedE 0.0005512717 0.0009778711
## soilC 0.0020974559 0.0017207758
## soilE -0.0013019968 -0.0004730252
## waterC -0.0007002882 -0.0001164310
## waterE 0.0012092374 0.0010468986

# Soil
env soi <- which(eDNA.env == "soil")
cent.soil <- as.data.frame(matrix(NA, nrow = length(env soi), ncol = 3))
colnames(cent.soil) <- c("T", "C", "E")
rownames(cent.soil) <- rownames(eDNA.div)[env soi]

# Centriod Ratio Calculaed for 1: vs Env; 2: vs C; 3 vs E
for (i in 1:length(env soi)){
  cent.soil[i, 1] <- dist(rbind(dispenv$vector[env soi[i]],
                                dispenv$centroids[3, ]), method = "euclidean")
  cent.soil[i, 2] <- dist(rbind(dispgroup$vector[env soi[i]],
                                dispgroup$centroids[5, ]), method = "euclidean")
  cent.soil[i, 3] <- dist(rbind(dispgroup$vector[env soi[i]],
                                dispgroup$centroids[6, ]), method = "euclidean")
}

pairs.soil <- eDNA.div[env soi, c(3,4,5,7)]

cent.soil.ratio <- as.data.frame(matrix(NA, nrow = length(unique(pairs.soil$pair)),
                                         ncol = 3))
colnames(cent.soil.ratio) <- c("T", "C", "E")
rownames(cent.soil.ratio) <- paste(pairs.soil$env, pairs.soil$pair,
                                   sep = "")[which(pairs.soil$treat == "E") ]

for (i in 1:dim(cent.soil.ratio)[1]){
  pair <- as.numeric(gsub("soil", "", rownames(cent.soil.ratio)[i]))
  cent.soil.ratio[i, 1] <- (cent.soil[, 1][which(pairs.soil$pair == pair &
                                                  pairs.soil$treat == "C")]) /
    (cent.soil[, 1][which(pairs.soil$pair == pair &
                          pairs.soil$treat == "E")])
  cent.soil.ratio[i, 2] <- (cent.soil[, 2][which(pairs.soil$pair == pair &
                                                  pairs.soil$treat == "C")]) /
    (cent.soil[, 2][which(pairs.soil$pair == pair &
                          pairs.soil$treat == "E")])
  cent.soil.ratio[i, 3] <- (cent.soil[, 3][which(pairs.soil$pair == pair &
                                                  pairs.soil$treat == "C")]) /
    (cent.soil[, 3][which(pairs.soil$pair == pair &
                          pairs.soil$treat == "E")])
}

# Water
env wat <- which(eDNA.env == "water")

```

```

cent.water <- as.data.frame(matrix(NA, nrow = length(env.wat), ncol = 3))
colnames(cent.water) <- c("T", "C", "E")
rownames(cent.water) <- rownames(eDNA.div)[env.wat]

for (i in 1:length(env.wat)){
  cent.water[i, 1] <- dist(rbind(disp.env$vector[env.wat[i]],
                                disp.env$centroids[4, ]), method = "euclidean")
  cent.water[i, 2] <- dist(rbind(disp.group$vector[env.wat[i]],
                                disp.group$centroids[7, ]), method = "euclidean")
  cent.water[i, 3] <- dist(rbind(disp.group$vector[env.wat[i]],
                                disp.group$centroids[8, ]), method = "euclidean")
}

pairs.water <- eDNA.div[env.wat, c(3,4,5,7)]

cent.water.ratio <- as.data.frame(matrix(NA, nrow = length(unique(pairs.water$pair)),
                                         ncol = 3))
colnames(cent.water.ratio) <- c("T", "C", "E")
rownames(cent.water.ratio) <- paste(pairs.water$env, pairs.water$pair,
                                   sep = "")[which(pairs.water$treat == "E") ]

for (i in 1:dim(cent.water.ratio)[1]){
  pair <- as.numeric(gsub("water", "", rownames(cent.water.ratio)[i]))
  cent.water.ratio[i, 1] <- (cent.water[, 1][which(pairs.water$pair == pair &
                                                    pairs.water$treat == "C")]) /
    (cent.water[, 1][which(pairs.water$pair == pair &
                            pairs.water$treat == "E")])
  cent.water.ratio[i, 2] <- (cent.water[, 2][which(pairs.water$pair == pair &
                                                    pairs.water$treat == "C")]) /
    (cent.water[, 2][which(pairs.water$pair == pair &
                            pairs.water$treat == "E")])
  cent.water.ratio[i, 3] <- (cent.water[, 3][which(pairs.water$pair == pair &
                                                    pairs.water$treat == "C")]) /
    (cent.water[, 3][which(pairs.water$pair == pair &
                            pairs.water$treat == "E")])
}

# Sediments
env.sed <- which(eDNA.env == "sed")
cent.sed <- as.data.frame(matrix(NA, nrow = length(env.sed), ncol = 3))
colnames(cent.sed) <- c("T", "C", "E")
rownames(cent.sed) <- rownames(eDNA.div)[env.sed]

for (i in 1:length(env.sed)){
  cent.sed[i, 1] <- dist(rbind(disp.env$vector[env.sed[i]],
                                disp.env$centroids[2, ]), method = "euclidean")
  cent.sed[i, 2] <- dist(rbind(disp.group$vector[env.sed[i]],
                                disp.group$centroids[3, ]), method = "euclidean")
  cent.sed[i, 3] <- dist(rbind(disp.group$vector[env.sed[i]],
                                disp.group$centroids[4, ]), method = "euclidean")
}

pairs.sed <- eDNA.div[env.sed, c(3,4,5,7)]

```

```

cent.sed.ratio <- as.data.frame(matrix(NA, nrow = length(unique(pairs.sed$pair)),
                                     ncol = 3))
colnames(cent.sed.ratio) <- c("T", "C", "E")
rownames(cent.sed.ratio) <- paste(pairs.sed$env, pairs.sed$pair,
                                sep = "")[which(pairs.sed$treat == "E") ]

for (i in 1:dim(cent.sed.ratio)[1]){
  pair <- as.numeric(gsub("sed", "", rownames(cent.sed.ratio)[i]))
  cent.sed.ratio[i, 1] <- (cent.sed[, 1][which(pairs.sed$pair == pair &
                                             pairs.sed$treat == "C")]) /
    (cent.sed[, 1][which(pairs.sed$pair == pair &
                         pairs.sed$treat == "E")])
  cent.sed.ratio[i, 2] <- (cent.sed[, 2][which(pairs.sed$pair == pair &
                                             pairs.sed$treat == "C")]) /
    (cent.sed[, 2][which(pairs.sed$pair == pair &
                         pairs.sed$treat == "E")])
  cent.sed.ratio[i, 3] <- (cent.sed[, 3][which(pairs.sed$pair == pair &
                                             pairs.sed$treat == "C")]) /
    (cent.sed[, 3][which(pairs.sed$pair == pair &
                         pairs.sed$treat == "E")])
}

# Feces
env.fec <- which(eDNA.env == "feces")
cent.feces <- as.data.frame(matrix(NA, nrow = length(env.fec), ncol = 3))
colnames(cent.feces) <- c("T", "C", "E")
rownames(cent.feces) <- rownames(eDNA.div)[env.fec]

for (i in 1:length(env.fec)){
  cent.feces[i, 1] <- dist(rbind(dispenv$centroids[env.fec[i]],
                                dispenv$centroids[1, ]), method = "euclidean")
  cent.feces[i, 2] <- dist(rbind(dispgroup$centroids[env.fec[i]],
                                dispgroup$centroids[1, ]), method = "euclidean")
  cent.feces[i, 3] <- dist(rbind(dispgroup$centroids[env.fec[i]],
                                dispgroup$centroids[2, ]), method = "euclidean")
}

pairs.feces <- eDNA.div[env.fec, c(3,4,5,7)]

cent.feces.ratio <- as.data.frame(matrix(NA, nrow = length(unique(pairs.feces$pair)),
                                     ncol = 3))
colnames(cent.feces.ratio) <- c("T", "C", "E")
rownames(cent.feces.ratio) <- paste(pairs.feces$env, pairs.feces$pair,
                                sep = "")[which(pairs.feces$treat == "E") ]

for (i in 1:dim(cent.feces.ratio)[1]){
  pair <- as.numeric(gsub("feces", "", rownames(cent.feces.ratio)[i]))
  cent.feces.ratio[i, 1] <- (cent.feces[, 1][which(pairs.feces$pair == pair &
                                                  pairs.feces$treat == "C")]) /
    (cent.feces[, 1][which(pairs.feces$pair == pair &
                         pairs.feces$treat == "E")])
  cent.feces.ratio[i, 2] <- (cent.feces[, 2][which(pairs.feces$pair == pair &
                                                  pairs.feces$treat == "C")]) /
    (cent.feces[, 2][which(pairs.feces$pair == pair &
                         pairs.feces$treat == "E")])
}

```

```

        (cent.feces[, 2][which(pairs.feces$pair == pair &
                               pairs.feces$treat == "E")])
cent.feces.ratio[i, 3] <- (cent.feces[, 3][which(pairs.feces$pair == pair &
                                                  pairs.feces$treat == "C")]) /
        (cent.feces[, 3][which(pairs.feces$pair == pair &
                               pairs.feces$treat == "E")])
}

# Ratio Data Table
ratio.list <- list(cent.feces.ratio[,1], cent.sed.ratio[,1],
                  cent.soil.ratio[,1], cent.water.ratio[,1])
cent.dist.table <- as.data.frame(matrix(NA, nrow = 4, ncol = 5))
colnames(cent.dist.table) <- c("env", "mean", "sem", "LCI", "UCI")
cent.dist.table$env <- c("Gut", "Sediment", "Soil", "Water")
cent.dist.table$mean <- unlist(lapply(ratio.list, mean))
cent.dist.table$sem <- unlist(lapply(ratio.list, sem))
cent.dist.table$LCI <- unlist(lapply(ratio.list, FUN = function(x) t.test(x)$conf.int[1]))
cent.dist.table$UCI <- unlist(lapply(ratio.list, FUN = function(x) t.test(x)$conf.int[2]))
p.cent.dist.table <- as.data.frame(cent.dist.table)

# Regression: centroid ratio vs. proportion eDNA
# Combine Centroid Distances
centroid.dists <- c(cent.feces.ratio$T, cent.sed.ratio$T, cent.soil.ratio$T,
                  cent.water.ratio$T)
names(centroid.dists) <- c(rownames(cent.feces.ratio), rownames(cent.sed.ratio),
                          rownames(cent.soil.ratio), rownames(cent.water.ratio))

# Reorganize
names(centroid.dists) %in% paste(eDNA.div.c$env, eDNA.div.c$pair, sep = "")

## [1] TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE
## [15] TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE
## [29] TRUE

all.equal(names(centroid.dists), paste(eDNA.div.c$env, eDNA.div.c$pair, sep = ""))

## [1] "28 string mismatches"

cent.dist <- centroid.dists[match(names(centroid.dists),
                                  paste(eDNA.div.c$env, eDNA.div.c$pair, sep = ""))]

# Combine Centroid Distance and Design Information
centroid.div <- cbind(eDNA.div.c[,c(1:3, 5:7)], cent.dist)

centroid.div$qPCR.match %in% eDNA.prop$sample.number

## [1] TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE
## [15] TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE TRUE
## [29] TRUE

centroid.div$prop <- NA
for (i in 1:length(centroid.div$prop)){
  centroid.div$prop[i] <- eDNA.prop$prop[which(eDNA.prop$sample.number ==
                                                centroid.div$qPCR.match[i])]
}

```

```

# Take averages of technical reps
centroid.div.2 <- ddply(centroid.div, .(env, qPCR.match, prop),
                        summarize, cent.dist = mean(cent.dist))

p.cent.reg <- lm(cent.dist ~ prop * env, data = centroid.div)
p.cent.reg <- lm(cent.dist ~ prop, data = centroid.div)
summary(p.cent.reg)

##
## Call:
## lm(formula = cent.dist ~ prop, data = centroid.div)
##
## Residuals:
##      Min       1Q   Median       3Q      Max
## -0.160695 -0.040424 -0.020653  0.009324  0.290218
##
## Coefficients:
##              Estimate Std. Error t value Pr(>|t|)
## (Intercept)  1.05946    0.04257  24.885  <2e-16 ***
## prop        -0.06268    0.09419  -0.665    0.511
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 0.101 on 27 degrees of freedom
## Multiple R-squared:  0.01614,    Adjusted R-squared:  -0.0203
## F-statistic: 0.4429 on 1 and 27 DF,  p-value: 0.5114

ttest(p.cent.reg, 1, 1)

##           t           df           p
## 1.3965772 27.0000000 0.1739165

```

answer => no correlation between proportion eDNA and centroid dist ratio.

Centroid Distance Plots

Taxonomic Centroid Distance Ratio Plot

```

png(filename="../figures/ratio.centroid.png",
     width = 800, height = 600, res = 96*2)

par(mar=c(3, 4, 1, 1) + 0.1)

cent.dist.table <- t.cent.dist.table

centroid.plot <- plot(cent.dist.table$mean, ylim = c(0.5, 1.7),
                     xlim = c(0.5, 4.5), pch = 22, bg = "white", lwd = 2,
                     cex = 3, yaxt = "n", xaxt = "n", cex.lab = 2, cex.axis = 1.5,
                     las = 1, ylab = "", xlab = "")
box(lwd = 2)
abline(h = 1, lty = 3, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = cent.dist.table$mean, y1 = cent.dist.table$LCI, angle = 90,

```

```

        length = 0.1, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = cent.dist.table$mean, y1 = cent.dist.table$UCI, angle = 90,
        length=0.1, lwd = 2)
points(c(1:4), cent.dist.table$mean, pch = 22, cex = 3, bg = "gray90", lwd = 2)

mtext(expression('Centroid Distance Ratio'), side = 2,
        outer = F, cex = 1.25, line = 2.5, adj = 0.5)
# Major Axes
axis(side = 2, lwd.ticks = 2, cex.axis = 1, las = 1,
      labels = T, at = c(0.5, 1, 1.5))

axis(side = 4, lwd.ticks = 2, cex.axis = 1, las = 1,
      labels = F, at = c(0.5, 1, 1.5))

axis(side = 1, lwd.ticks = 2, cex.axis = 0.9, las = 1,
      labels = c("Gut", "Sediment", "Soil", "Water"), at = c(1, 2, 3, 4))

axis(side = 3, lwd.ticks = 2, cex.axis = 1.5, las = 1,
      at = c(1, 2, 3, 4), labels = F)

# Minor ticks
axis(side = 2, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.5, 1, 1.5))
axis(side = 4, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.5, 1, 1.5))
axis(side = 1, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))
axis(side = 3, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))

# Close Plot Device
dev.off()

## pdf
## 2

graphics.off()

# Show Plot
img <- readPNG("../figures/ratio.centroid.png")
grid.raster(img)

```

Phylogenetic Centroid Distance Ratio Plot

```

png(filename="../figures/ratio.phylo.centroid.png",
     width = 800, height = 600, res = 96*2)

par(mar=c(3, 4, 1, 1) + 0.1)

cent.dist.table <- p.cent.dist.table

centroid.plot <- plot(cent.dist.table$mean, ylim = c(0.5, 1.5),
                     xlim = c(0.5, 4.5), pch = 22, bg = "white", lwd = 2,
                     cex = 3, yaxt = "n", xaxt = "n", cex.lab = 2, cex.axis = 1.5,
                     las = 1, ylab = "", xlab = "")
                     box(lwd = 2)
abline(h = 1, lty = 3, lwd = 2)

```

```

arrows(x0 = c(1,2,3,4), y0 = cent.dist.table$mean, y1 = cent.dist.table$LCI, angle = 90,
       length = 0.1, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = cent.dist.table$mean, y1 = cent.dist.table$UCI, angle = 90,
       length=0.1, lwd = 2)
points(c(1:4), cent.dist.table$mean, pch = 22, cex = 3, bg = "gray90", lwd = 2)

mtext(expression('Centroid Distance Ratio'), side = 2,
       outer = F, cex = 1.25, line = 2.5, adj = 0.5)
# Major Axes
axis(side = 2, lwd.ticks = 2, cex.axis = 1, las = 1,
     labels = T, at = c(0.5, 1, 1.5))

axis(side = 4, lwd.ticks = 2, cex.axis = 1, las = 1,
     labels = F, at = c(0.5, 1, 1.5))

axis(side = 1, lwd.ticks = 2, cex.axis = 0.9, las = 1,
     labels = c("Gut", "Sediment", "Soil", "Water"), at = c(1, 2, 3, 4))

axis(side = 3, lwd.ticks = 2, cex.axis = 1.5, las = 1,
     at = c(1, 2, 3, 4), labels = F)

# Minor ticks
axis(side = 2, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.5, 1, 1.5))
axis(side = 4, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.5, 1, 1.5))
axis(side = 1, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))
axis(side = 3, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))

# Close Plot Device
dev.off()

## pdf
## 2

graphics.off()

# Show Plot
img <- readPNG("../figures/ratio.phylo.centroid.png")
grid.raster(img)

```

Multipanel Centroid Distance Ratio Plot

```

png(filename="../figures/panel.ratio.centroid.png",
     width = 800, height = 1200, res = 96*2)

layout(matrix(1:2, 2, 1))
par(mar = c(0.5, 5, 0.5, 0) + 0.1, oma = c(4, 1, 1, 1))

# Bray Curtis Plot
cent.dist.table <- t.cent.dist.table
centroid.plot <- plot(cent.dist.table$mean, ylim = c(0.5, 1.6),
                    xlim = c(0.5, 4.5), pch = 22, bg = "white", lwd = 2,
                    cex = 3, yaxt = "n", xaxt = "n", cex.lab = 2, cex.axis = 1.5,
                    las = 1, ylab = "", xlab = "")

```



```

    box(lwd = 2)
abline(h = 1, lty = 3, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = cent.dist.table$mean, y1 = cent.dist.table$LCI, angle = 90,
       length = 0.1, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = cent.dist.table$mean, y1 = cent.dist.table$UCI, angle = 90,
       length=0.1, lwd = 2)
points(c(1:4), cent.dist.table$mean,pch = 22, cex = 3, bg = "gray90", lwd = 2)

mtext('Centroid Distance Ratio\n(Bray Curtis Distance)', side = 2,
      outer = F, cex = 1.25, line = 2.5, adj = 0.5)
# Major Axes
axis(side = 2, lwd.ticks = 2, cex.axis = 1, las = 1,
     labels = T, at = c(0.7, 1, 1.3, 1.6))

axis(side = 4, lwd.ticks = 2, cex.axis = 1, las = 1,
     labels = F, at = c(0.7, 1, 1.3, 1.6), tck = -0.02)

axis(side = 1, lwd.ticks = 2, cex.axis = 0.9, las = 1,
     labels = F, at = c(1, 2, 3, 4), tck = -0.02)

axis(side = 3, lwd.ticks = 2, cex.axis = 1.5, las = 1,
     at = c(1, 2, 3, 4), labels = F, tck = -0.02)

# Minor ticks
axis(side = 2, lwd.ticks = 2, tck = 0.02, labels = F,
     at = c(0.7, 1, 1.3, 1.6))
axis(side = 4, lwd.ticks = 2, tck = 0.02, labels = F,
     at = c(0.7, 1, 1.3, 1.6))
axis(side = 1, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))
axis(side = 3, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))

# Phylogenetic Plot
cent.dist.table <- p.cent.dist.table
centroid.plot <- plot(cent.dist.table$mean, ylim = c(0.6, 1.4),
                    xlim = c(0.5, 4.5), pch = 22, bg = "white", lwd = 2,
                    cex = 3, yaxt = "n", xaxt = "n", cex.lab = 2, cex.axis = 1.5,
                    las = 1, ylab = "", xlab = "")
    box(lwd = 2)
abline(h = 1, lty = 3, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = cent.dist.table$mean, y1 = cent.dist.table$LCI, angle = 90,
       length = 0.1, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = cent.dist.table$mean, y1 = cent.dist.table$UCI, angle = 90,
       length=0.1, lwd = 2)
points(c(1:4), cent.dist.table$mean,pch = 22, cex = 3, bg = "gray90", lwd = 2)

mtext('Centroid Distance Ratio\n(UniFrac Distance)', side = 2,
      outer = F, cex = 1.25, line = 2.5, adj = 0.5)
# Major Axes
axis(side = 2, lwd.ticks = 2, cex.axis = 1, las = 1,
     labels = T, at = c(0.7, 1, 1.3))

axis(side = 4, lwd.ticks = 2, cex.axis = 1, las = 1,

```

```

    labels = F, at = c(0.7, 1, 1.3), tck = -0.02)

axis(side = 1, lwd.ticks = 2, cex.axis = 1, las = 1,
     labels = c("Gut", "Sediment", "Soil", "Water"), at = c(1, 2, 3, 4))

axis(side = 3, lwd.ticks = 2, cex.axis = 1.5, las = 1,
     at = c(1, 2, 3, 4), labels = F, tck = -0.02)

# Minor ticks
axis(side = 2, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.7, 1, 1.3))
axis(side = 4, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.7, 1, 1.3))
axis(side = 1, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))
axis(side = 3, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))

# Close Plot Device
dev.off()

## pdf
## 2

graphics.off()

# Show Plot
img <- readPNG("../figures/panel.ratio.centroid.png")
grid.raster(img)

```

Old Code

```

# Delta Distance as Function of Proportion eDNA
eDNA.pairs
eDNA.prop

eDNA.pairs$prop <- eDNA.prop.sub2$prop[match(eDNA.prop.sub2$sample.number, eDNA.pairs$qPCR)]

# Reshape data
eDNA.pairs.2 <- reshape(eDNA.pairs[,1:4], timevar = "treat",
                       idvar = c("qPCR.match", "env"), direction = "wide")

# Regression: PD ratio vs. proportion eDNA
phylo.reg <- lm(phylo.2[order(phylo.2$qPCR.match), ]$phylo.ratio ~ eDNA.prop.sub2$prop * eDNA.prop.sub2$
summary(phylo.reg)

# Separate by Treatments
eDNA.div.c <- eDNA.div[which(eDNA.div$treat == "C"), ]
eDNA.div.e <- eDNA.div[which(eDNA.div$treat == "E"), ]
OTU.REL.log.c <- OTU.REL.log[which(eDNA.div$treat == "C"), ]
OTU.REL.log.e <- OTU.REL.log[which(eDNA.div$treat == "E"), ]

# Order Each by Pairings
eDNA.div.c <- eDNA.div.c[order(eDNA.div.c$pair), ]
eDNA.div.e <- eDNA.div.e[order(eDNA.div.e$pair), ]
OTU.REL.log.c <- OTU.REL.log.c[order(eDNA.div.c$pair), ]

```

```

OTU.REL.log.e <- OTU.REL.log.e[order(eDNA.div.e$pair), ]
all.equal(eDNA.div.e$env, eDNA.div.c$env)

# Create Distance Matrix for Treatments
dist.c <- vegdist(OTU.REL.log.c, method = "bray", upper = T, diag = T)
dist.e <- vegdist(OTU.REL.log.e, method = "bray", upper = T, diag = T)

pcoa.c <- cmdscale(dist.c, k = 3)
pcoa.e <- cmdscale(dist.e, k = 3)

# Procrustes Analysis
eDNA.ProC <- protest(pcoa.c, pcoa.e, scores = "sites", permutations = how(nperm = 999))
eDNA.ProC
summary(eDNA.ProC)

# Delta Distances
## Taxonomic Distances Between Pairs
# Data Check
all.equal(row.names(eDNA.div), row.names(OTU.REL.log))

# Create Raw Distance Matrix
dist.raw <- as.matrix(vegdist(OTU.REL.log, method = "bray", upper = T, diag = T))

# Calculate Average Distance For Each Environment
mean.dist.env <- matrix(NA, length(unique(eDNA.div$env)), 3)
rownames(mean.dist.env) <- unique(eDNA.div$env)
colnames(mean.dist.env) <- c("Total", "C", "E")
for (i in 1:dim(mean.dist.env)[1]){
  mean.dist.env[i, 1] <- mean(dist.raw[eDNA.div$env == rownames(mean.dist.env)[i],
                                     eDNA.div$env == rownames(mean.dist.env)[i]])
  mean.dist.env[i, 2] <- mean(dist.raw[eDNA.div$env == rownames(mean.dist.env)[i] &
                                     eDNA.div$treat == "C",
                                     eDNA.div$env == rownames(mean.dist.env)[i] &
                                     eDNA.div$treat == "C"])
  mean.dist.env[i, 3] <- mean(dist.raw[eDNA.div$env == rownames(mean.dist.env)[i] &
                                     eDNA.div$treat == "E",
                                     eDNA.div$env == rownames(mean.dist.env)[i] &
                                     eDNA.div$treat == "E"])
}

# Create Pairing Matrix
pair.names <- levels(as.factor(eDNA.div$pair))
eDNA.pairs <- as.data.frame(matrix(NA, length(pair.names), 4))
colnames(eDNA.pairs) <- c("pair", "env", "dis", "qPCR")
eDNA.pairs$pair <- pair.names

# Calculate Distance for Pairs
for (i in 1:length(pair.names)){
  pair.temp <- eDNA.pairs$pair[i]
  samp.temp <- rownames(eDNA.div)[which(eDNA.div$pair == pair.temp)]
  if (length(samp.temp) != 2){
    stop("more than two samples in pair")
  }
}

```

```

dist.temp <- dist.raw[which(colnames(dist.raw) == samp.temp[1]),
                      which(colnames(dist.raw) == samp.temp[2])]
eDNA.pairs$env[i] <- eDNA.div$env[which(row.names(eDNA.div) == samp.temp[1])]
eDNA.pairs$dis[i] <- dist.temp
eDNA.pairs$qPCR[i] <- eDNA.div$qPCR.match[which(row.names(eDNA.div) == samp.temp[1])]
}

# Organize Data
pair.dis.mean <- aggregate(dis ~ env, eDNA.pairs, mean)
pair.dis.sem <- aggregate(dis ~ env, eDNA.pairs, sem)
pair.dis.95 <- aggregate(dis ~ env, eDNA.pairs,
                          FUN = function(x) t.test(x)$conf.int[1:2])
t.pair.dis.table <- data.frame(env = pair.dis.mean$env, mean = pair.dis.mean$dis,
                               sem = pair.dis.sem$dis, ci = pair.dis.95$dis)
colnames(t.pair.dis.table) <- c("env", "mean", "sem", "LCI", "UCI")

# Organize Data as Ratio
env.means <-

## Phylogenetic Distances between Pairs

# Data Check
all.equal(row.names(eDNA.div), row.names(as.matrix(phylo.dist.m)))

# Create Raw Distance Matrix
dist.raw <- phylo.dist.m
colnames(dist.raw) <- rownames(dist.raw)

# Create Pairing Matrix
pair.names <- levels(as.factor(eDNA.div$pair))
eDNA.pairs <- as.data.frame(matrix(NA, length(pair.names), 4))
colnames(eDNA.pairs) <- c("pair", "env", "dis", "qPCR")
eDNA.pairs$pair <- pair.names

# Calculate Distance for Pairs
for (i in 1:length(pair.names)){
  pair.temp <- eDNA.pairs$pair[i]
  samp.temp <- rownames(eDNA.div)[which(eDNA.div$pair == pair.temp)]
  if (length(samp.temp) != 2){
    stop("more than two samples in pair")
  }
  dist.temp <- dist.raw[which(colnames(dist.raw) == samp.temp[1]),
                        which(colnames(dist.raw) == samp.temp[2])]
  eDNA.pairs$env[i] <- eDNA.div$env[which(row.names(eDNA.div) == samp.temp[1])]
  eDNA.pairs$dis[i] <- dist.temp
  eDNA.pairs$qPCR[i] <- eDNA.div$qPCR.match[which(row.names(eDNA.div) == samp.temp[1])]
}

pair.dis.mean <- aggregate(dis ~ env, eDNA.pairs, mean)
pair.dis.sem <- aggregate(dis ~ env, eDNA.pairs, sem)
pair.dis.95 <- aggregate(dis ~ env, eDNA.pairs,
                          FUN = function(x) t.test(x)$conf.int[1:2])

```

```

p.pair.dis.table <- data.frame(env = pair.dis.mean$env, mean = pair.dis.mean$dis,
                               sem = pair.dis.sem$dis, ci = pair.dis.95$dis)
colnames(p.pair.dis.table) <- c("env", "mean", "sem", "LCI", "UCI")

# Delta Distance Plots
## Taxonomic Plot

png(filename="../figures/delta.dist.png",
     width = 800, height = 600, res = 96*2)

par(mar=c(3, 4, 1, 1) + 0.1)

distance.plot <- plot(t.pair.dis.table$mean, ylim = c(-0.1, 0.6),
                     xlim = c(0.5, 4.5), pch = 22, bg = "white", lwd = 2,
                     cex = 3, yaxt = "n", xaxt = "n", cex.lab = 2, cex.axis = 1.5,
                     las = 1, ylab = "", xlab = "")
box(lwd = 2)
abline(h = 0, lty = 3, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = delta.dis.table$mean, y1 = delta.dis.table$LCI,
       angle = 90, length = 0.1, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = delta.dis.table$mean, y1 = delta.dis.table$UCI,
       angle = 90, length=0.1, lwd = 2)
points(c(1:4), delta.dis.table$mean, pch = 22, cex = 3, bg = "gray90", lwd = 2)

mtext(expression('Distance'), side = 2,
       outer = F, cex = 1.5, line = 3, adj = 0.5)

# Major Axes
axis(side = 2, lwd.ticks = 2, cex.axis = 1, las = 1,
     labels = T, at = c(seq(-0.1, 0.3, 0.1)))
axis(side = 4, lwd.ticks = 2, cex.axis = 1, las = 1,
     labels = F, at = c(-0.1, 0.3, 0.1))
axis(side = 1, lwd.ticks = 2, cex.axis = 0.9, las = 1,
     labels = c("Gut", "Sediment", "Soil", "Water"), at = c(1, 2, 3, 4))
axis(side = 3, lwd.ticks = 2, cex.axis = 1.5, las = 1,
     at = c(1, 2, 3, 4), labels = F)

# Minor ticks
axis(side = 2, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.5, 1, 1.5))
axis(side = 4, lwd.ticks = 2, tck = 0.02, labels = F, at = c(-0.1, 0.3, 0.1))
axis(side = 1, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))
axis(side = 3, lwd.ticks = 2, tck = 0.02, labels = F, at = c(-0.1, 0.3, 0.1))

# Close Plot Device
dev.off()
graphics.off()

# Show Plot
img <- readPNG("../figures/delta.dist.png")
grid.raster(img)

```

```

## Phylogenetic Plot

png(filename="../figures/phylo.delta.dist.png",
     width = 800, height = 600, res = 96*2)

par(mar=c(3, 4, 1, 1) + 0.1)

distance.plot <- plot(delta.dis.table$mean, ylim = c(-0.1, 0.3),
  xlim = c(0.5, 4.5), pch = 22, bg = "white", lwd = 2,
  cex = 3, yaxt = "n", xaxt = "n", cex.lab = 2, cex.axis = 1.5,
  las = 1, ylab = "", xlab = "")
  box(lwd = 2)
abline(h = 0, lty = 3, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = delta.dis.table$mean, y1 = delta.dis.table$LCI,
  angle = 90, length = 0.1, lwd = 2)
arrows(x0 = c(1,2,3,4), y0 = delta.dis.table$mean, y1 = delta.dis.table$UCI,
  angle = 90, length=0.1, lwd = 2)
points(c(1:4), delta.dis.table$mean, pch = 22, cex = 3, bg = "gray90", lwd = 2)

mtext(expression('Distance'), side = 2,
  outer = F, cex = 1.5, line = 3, adj = 0.5)

# Major Axes
axis(side = 2, lwd.ticks = 2, cex.axis = 1, las = 1,
  labels = T, at = c(seq(-0.1, 0.3, 0.1)))
axis(side = 4, lwd.ticks = 2, cex.axis = 1, las = 1,
  labels = F, at = c(-0.1, 0.3, 0.1))
axis(side = 1, lwd.ticks = 2, cex.axis = 0.9, las = 1,
  labels = c("Gut", "Sediment", "Soil", "Water"), at = c(1, 2, 3, 4))
axis(side = 3, lwd.ticks = 2, cex.axis = 1.5, las = 1,
  at = c(1, 2, 3, 4), labels = F)

# Minor ticks
axis(side = 2, lwd.ticks = 2, tck = 0.02, labels = F, at = c(0.5, 1, 1.5))
axis(side = 4, lwd.ticks = 2, tck = 0.02, labels = F, at = c(-0.1, 0.3, 0.1))
axis(side = 1, lwd.ticks = 2, tck = 0.02, labels = F, at = c(1, 2, 3, 4))
axis(side = 3, lwd.ticks = 2, tck = 0.02, labels = F, at = c(-0.1, 0.3, 0.1))

# Close Plot Device
dev.off()
graphics.off()

# Show Plot
img <- readPNG("../figures/phylo.delta.dist.png")
grid.raster(img)

## Delta-Abundances

# Check order of design
all.equal(row.names(eDNA.div), row.names(OTU.REL.log))

OTU.REL.c <- OTU.REL.log[which(eDNA.div$treat == "C"), ]

```

```

OTU.REL.e <- OTU.REL.log[which(eDNA.div$treat == "E"), ]

eDNA.div.c <- eDNA.div[which(eDNA.div$treat == "C"), ]
eDNA.div.e <- eDNA.div[which(eDNA.div$treat == "E"), ]

all.equal(sort(eDNA.div.c$samp.code), sort(eDNA.div.e$samp.code))

eDNA.div.c <- eDNA.div.c[order(eDNA.div.c$samp.code), ]
eDNA.div.e <- eDNA.div.e[order(eDNA.div.e$samp.code), ]

OTU.REL.c2 <- OTU.REL.c[order(eDNA.div.c$samp.code), ]
OTU.REL.e2 <- OTU.REL.e[order(eDNA.div.e$samp.code), ]

dim(OTU.REL.c2); dim(OTU.REL.e2)
all.equal(eDNA.div.e$samp.code, eDNA.div.c$samp.code)

delta.abund <- OTU.REL.c2 - OTU.REL.e2

eDNA.permanova <- adonis(delta.abund ~ eDNA.div.c$env,
                        method = "euclidean", binary = FALSE)
eDNA.permanova

# What is this stuff for????? Check all calculations
# Variation Parititioning

# Turn Distance Matrices into Matrices
dist.raw <- as.matrix(dist.raw)
dist.e <- as.matrix(dist.e)
dist.c <- as.matrix(dist.c)

# Seperate by Habitat
eDNA.div.gut <- eDNA.div[which(eDNA.div$env == "feces"), ]
eDNA.div.sed <- eDNA.div[which(eDNA.div$env == "sed"), ]
eDNA.div.sol <- eDNA.div[which(eDNA.div$env == "soil"), ]
eDNA.div.wat <- eDNA.div[which(eDNA.div$env == "water"), ]

OTU.REL.log.gut <- OTU.REL.log[which(eDNA.div$env == "feces"), ]
OTU.REL.log.sed <- OTU.REL.log[which(eDNA.div$env == "sed"), ]
OTU.REL.log.sol <- OTU.REL.log[which(eDNA.div$env == "soil"), ]
OTU.REL.log.wat <- OTU.REL.log[which(eDNA.div$env == "water"), ]

# Order Each by Pairings
eDNA.div.gut <- eDNA.div.gut[order(eDNA.div.gut$pair), ]
eDNA.div.sed <- eDNA.div.sed[order(eDNA.div.sed$pair), ]
eDNA.div.sol <- eDNA.div.sol[order(eDNA.div.sol$pair), ]
eDNA.div.wat <- eDNA.div.wat[order(eDNA.div.wat$pair), ]
OTU.REL.log.gut <- OTU.REL.log.gut[order(eDNA.div.gut$pair), ]
OTU.REL.log.sed <- OTU.REL.log.sed[order(eDNA.div.sed$pair), ]

```

```

OTU.REL.log.sol <- OTU.REL.log.sol[order(eDNA.div.sol$pair), ]
OTU.REL.log.wat <- OTU.REL.log.wat[order(eDNA.div.wat$pair), ]

# Create Distance Matrix for Treatments
dist.gut <- vegdist(OTU.REL.log.gut, method = "bray", upper = F, diag = F)
dist.sed <- vegdist(OTU.REL.log.sed, method = "bray", upper = F, diag = F)
dist.sol <- vegdist(OTU.REL.log.sol, method = "bray", upper = F, diag = F)
dist.wat <- vegdist(OTU.REL.log.wat, method = "bray", upper = F, diag = F)

# Turn Distance Matrices into Matrices
dist.gut2 <- as.matrix(dist.gut)
dist.sed2 <- as.matrix(dist.sed)
dist.sol2 <- as.matrix(dist.sol)
dist.wat2 <- as.matrix(dist.wat)

# what are the factors I have to work with
head(eDNA.div.gut)
levels(eDNA.div.gut$treat)
levels(as.factor(eDNA.div.gut$qPCR.match))

# Variance Partitioning
var.gut <- varpart(Y = dist.gut, as.numeric(eDNA.div.gut$treat), (eDNA.div.gut$qPCR.match))
plot(var.gut)

gut.RDA <- capscale(dist.gut ~ eDNA.div.gut$treat + as.factor(eDNA.div.gut$qPCR.match))
anova(gut.RDA, by = "term")
RsquareAdj(gut.RDA)

# Adjusted RSquare by Terms
rda.aov <- (anova(gut.RDA, by = "term"))
r2 <- (rda.aov$SumOfSqs[1:2] / sum(rda.aov$SumOfSqs[1:3]))
m <- c(length(levels(eDNA.div.gut$treat)) - 1,
       length(levels(as.factor(eDNA.div.gut$qPCR.match))) - 1)
n <- nrow(gut.RDA$CCA$u)
adjr2 <- 1 - (1 - r2) * (n - 1)/(n - m - 1)
r2; adjr2; sum(r2); sum(adjr2)

var.r2 <- as.data.frame(matrix(NA, 4, 3))
colnames(var.r2) <- c("eDNA", "replicates", "unexplained")
row.names(var.r2) <- c("gut", "sed", "sol", "wat")
for (i in 1:dim(var.r2)[1]){
  env.temp <- rownames(var.r2)[i]
  dis.temp <- get(paste("dist", env.temp, sep = "."))
  dat.temp <- get(paste("eDNA.div", env.temp, sep = "."))
  rda.temp <- capscale(dis.temp ~ dat.temp$treat +
                      as.factor(dat.temp$qPCR.match))
  aov.temp <- anova(rda.temp, by = "term")
  r2 <- (aov.temp$SumOfSqs[1:2] / sum(aov.temp$SumOfSqs[1:3]))
  var.r2[i, 1] <- r2[1]
  var.r2[i, 2] <- r2[2]
  var.r2[i, 3] <- (1 - sum(r2))
}

```



```

png(filename="../figures/variation.png",
     width = 1200, height = 800, res = 96*2)
layout(matrix(1:2, 2, 1) , heights = c(8, 2))
par(mar = c(2, 5, 1, 0) + 0.5, oma = c(0.5, 0.5, 0.5, 0.5))

barplot(t(as.matrix(var.r2)),
       names.arg = c("Gut", "Sediment", "Soil", "Water"),
       las = 1, ylab = "")

mtext("Proportion of Variation", side = 2, cex = 1.25, line = 3)

# Add Legend Outside
par(mar = c(1, 5, 0, 1) + 0.5)
plot.new()
legend("top", c("DNase", "Replicates", "Unexplained"),
      pch = 22, pt.bg = c("gray40", "gray60", "gray90"),
      bty = "n", y.intersp = 1.5, horiz = T)

# Close Plot Device
dev.off()
graphics.off()

# Show Plot
img <- readPNG("../figures/variation.png")
grid.raster(img)

# Variation based on Bray Curtis Distance

bc.var <- as.data.frame(matrix(NA, 4, 3))
colnames(bc.var) <- c("reps", "treat", "env")
rownames(bc.var) <- c("gut", "sed", "sol", "wat")
for (i in 1:dim(bc.var)[1]){
  env.temp <- rownames(bc.var)[i]
  dis.temp <- get(paste("dist", env.temp, sep = "."))
  dat.temp <- get(paste("eDNA.div", env.temp, sep = "."))
  dat.temp <- dat.temp[order(dat.temp$samp.num), ]
  if (all(attributes(dis.temp)$Labels != rownames(dat.temp))){
    stop("distance matix and data are not arranged properly")
  }
  dis.temp.C <- as.matrix(dis.temp)[which(dat.temp$treat == "C"),
                                     which(dat.temp$treat == "C")]
  dis.temp.C <- as.dist(dis.temp.C)
  bc.var[i,3] <- mean(dis.temp.C)
  dis.pair <- as.data.frame(matrix(NA, length(unique(dat.temp$pair)), 3))
  colnames(dis.pair) <- c("C", "E", "dis")
  dis.pair[,1] <- rownames(dat.temp)[dat.temp$treat == "C"]
  for (j in 1:length(dis.pair$C)){
    pair <- dat.temp$pair[which(rownames(dat.temp) == dis.pair$C[j])]
    dis.pair[j, 2] <- rownames(dat.temp)[which(dat.temp$treat == "E" &
                                                dat.temp$pair == pair)]
    dis.pair[j, 3] <- as.matrix(dis.temp)[which(rownames(dat.temp) == dis.pair[j, 1]),

```

```

                                which(rownames(dat.temp) == dis.pair[j, 2]))
}
bc.var[i,2] <- mean(dis.pair[,3])
reps <- names(which(table(dat.temp$qPCR.match) > 2))
dis.rep <- as.data.frame(matrix(NA, length(reps), 3))
colnames(dis.rep) <- c("rep1", "rep2", "dis")
for (k in 1:length(reps)){
  dis.rep[k, 1:2] <- rownames(dat.temp)[which(dat.temp$treat == "C" &
                                             dat.temp$qPCR.match == reps[k])]
  dis.rep[k, 3] <- as.matrix(dis.temp)[which(rownames(dat.temp) == dis.rep[k, 1]),
                                           which(rownames(dat.temp) == dis.rep[k, 2])]
}
bc.var[i, 1] <- mean(dis.rep[, 3])
}

bc.var

# bc.var <- bc.var/rowSums(bc.var)

# Standardized BC distances
bc.var.std <- bc.var/rowSums(bc.var)

## Distance Partitioning Plot

png(filename="../figures/BC_dist.png",
     width = 1200, height = 800, res = 96*2)
layout(matrix(1:2, 2, 1) , heights = c(8, 2))
par(mar = c(2, 5, 1, 0) + 0.5, oma = c(0.5, 0.5, 0.5, 0.5))

barplot(t(as.matrix(bc.var)),
        names.arg = c("Gut", "Sediment", "Soil", "Water"),
        las = 1, ylab = "", ylim = c(0, 1))

mtext("Bray Curtis Distance", side = 2, cex = 1.25, line = 3)

# Add Legend Outside
par(mar = c(1, 5, 0, 1) + 0.5)
plot.new()
legend("top", c("Replicates", "Treatments", "Environments"),
      pch = 22, pt.bg = c("gray40", "gray60", "gray90"),
      bty = "n", y.intersp = 1.5, horiz = T)

# Close Plot Device
dev.off()
graphics.off()

# Show Plot
img <- readPNG("../figures/BC_dist.png")
grid.raster(img)

# PERMANOVA on Delta-Distances

```

```

# Check order of design
all.equal(row.names(eDNA.div), row.names(as.matrix(phylo.dist)))

OTU.REL.c <- OTU.REL.log[which(eDNA.div$treat == "C"), ]
OTU.REL.e <- OTU.REL.log[which(eDNA.div$treat == "E"), ]

eDNA.div.c <- eDNA.div[which(eDNA.div$treat == "C"), ]
eDNA.div.e <- eDNA.div[which(eDNA.div$treat == "E"), ]

all.equal(sort(eDNA.div.c$samp.code), sort(eDNA.div.e$samp.code))

eDNA.div.c <- eDNA.div.c[order(eDNA.div.c$samp.code), ]
eDNA.div.e <- eDNA.div.e[order(eDNA.div.e$samp.code), ]

OTU.REL.c2 <- OTU.REL.c[order(eDNA.div.c$samp.code), ]
OTU.REL.e2 <- OTU.REL.e[order(eDNA.div.e$samp.code), ]

dim(OTU.REL.c2); dim(OTU.REL.e2)
all.equal(eDNA.div.e$samp.code, eDNA.div.c$samp.code)

delta.abund <- OTU.REL.c2 - OTU.REL.e2

eDNA.permanova <- adonis(delta.abund ~ eDNA.div.c$env,
                        method = "euclidean", binary = FALSE)
eDNA.permanova

rowSums(delta.abund)

# Effect Size from lda
require(MASS)
eDNA.lda <- lda(delta.abund ~ eDNA.div.c$env, method = "euclidean")
anova(eDNA.dbrDA)
anova(eDNA.dbrDA, by = "terms")

# Distance Moved in PCoA Space
eDNA.points.c <- eDNA.points[which(eDNA.points$treat == "C"), ]
eDNA.points.e <- eDNA.points[which(eDNA.points$treat == "E"), ]

eDNA.points.env <- eDNA.points.c$env

eDNA.points.c <- eDNA.points.c[order(eDNA.points.c$samp.code), c(1:3)]
eDNA.points.e <- eDNA.points.e[order(eDNA.points.e$samp.code), c(1:3)]

delta.dis <- matrix(NA, dim(eDNA.points.c)[1], 2)
rownames(delta.dis) <- eDNA.points.c$samp.code
colnames(delta.dis) <- c("Abs", "Non_Abs")

for (i in 1:dim(eDNA.points.c)[1]){
  dis.temp <- matrix(NA, 3, 2)
  for (j in 1:3){
    dis.temp[j, 1] <- abs(eDNA.points.c[i,j] - eDNA.points.e[i,j])
    dis.temp[j, 2] <- (eDNA.points.c[i,j] - eDNA.points.e[i,j])
  }
}

```

```

    delta.dis[i, 1] <- sum(abs(dis.temp[,1]))
    delta.dis[i, 2] <- sum(abs(dis.temp[,2]))
  }

delta.dis <- as.data.frame(delta.dis)

delta.dis$env <- eDNA.points.env

delta.dis.mean <- aggregate(Non_Abs ~ env, delta.dis, mean)
delta.dis.sem <- aggregate(Non_Abs ~ env, delta.dis, sem)
delta.dis.95 <- aggregate(Non_Abs ~ env, delta.dis,
  FUN = function(x) t.test(x)$conf.int[1:2])
delta.dis.table <- data.frame(env = delta.dis.mean$env, mean = delta.dis.mean$Non_Abs,
  sem = delta.dis.sem$Non_Abs, ci = delta.dis.95$Non_Abs)
colnames(delta.dis.table) <- c("env", "mean", "sem", "LCI", "UCI")

```