

PERMANOVA

1. Read in data

- Week = 13
- Two groups: 32 in PAT group and 36 in Control group

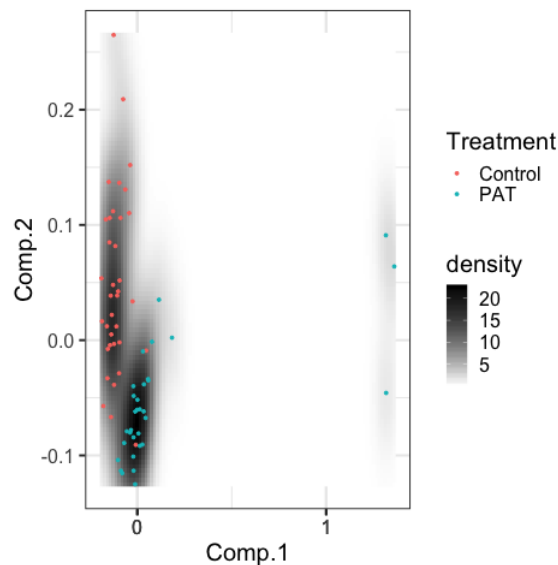
```
ind = (sample_data(phy.genus)$Week==13 &  
      (sample_data(phy.genus)$Treatment != 'STAT'))  
pseq = subset_samples(phy.genus,ind)  
pseq.rel = microbiome::transform(pseq, "compositional")  
otu = abundances(pseq.rel)  
meta = meta(pseq.rel)  
dim(otu) # 37 68; 37 kinds of microbiome, 68 subjects
```

```
## [1] 37 68
```

```
dim(meta) # 68, 53; 68 subjects and 53 features
```

```
## [1] 68 53
```

2. Visualize the population density and highlight sample groups



3. PERMANOVA significance test for group-level differences

```
permanova <- adonis(t(otu) ~ Treatment,  
                   data = meta, permutations=99, method = "bray")  
permanova
```

```
##
```

```
## Call:
```

```
## adonis(formula = t(otu) ~ Treatment, data = meta, permutations = 99,      method = "bray")
```

```
##
```

```
## Permutation: free
```

```
## Number of permutations: 99
##
## Terms added sequentially (first to last)
##
##           Df SumsOfSqs MeanSqs F.Model    R2 Pr(>F)
## Treatment  1    1.2610 1.26102  29.573 0.30943  0.01 **
## Residuals 66    2.8143 0.04264    0.69057
## Total      67    4.0753          1.00000
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

4. Checking the homogeneity condition

```
dist = vegdist(t(otu))
anova(betadisper(dist, meta$Treatment))
```

```
## Analysis of Variance Table
##
## Response: Distances
##           Df Sum Sq Mean Sq F value Pr(>F)
## Groups      1 0.00043 0.0004343  0.0319 0.8588
## Residuals 66 0.89910 0.0136228
```

5. Show coefficients for the top taxa separating the groups

