

### 1) Problem (1): Perform PCA ordination.

Below is just a summary of the PCA components I got-

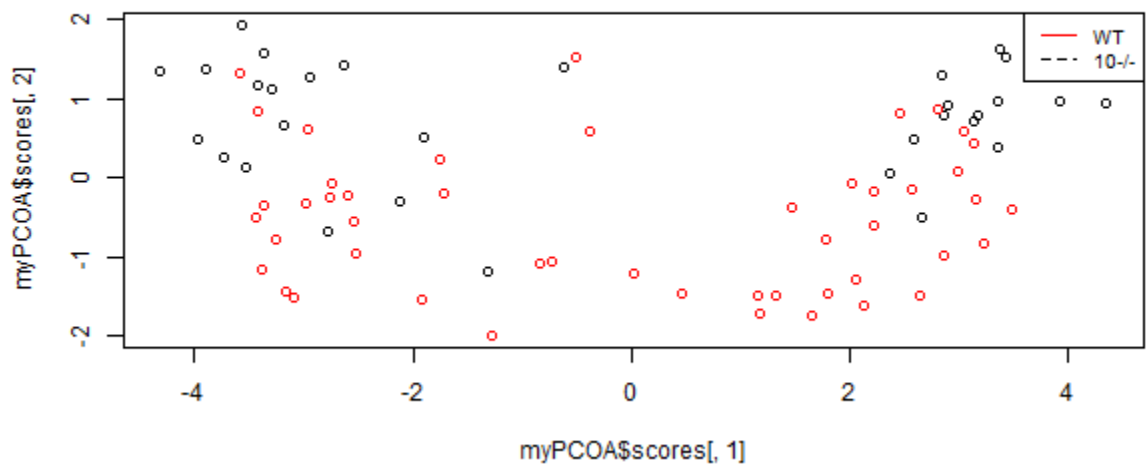
```
> print(summary(myPCOA))
```

Importance of components:

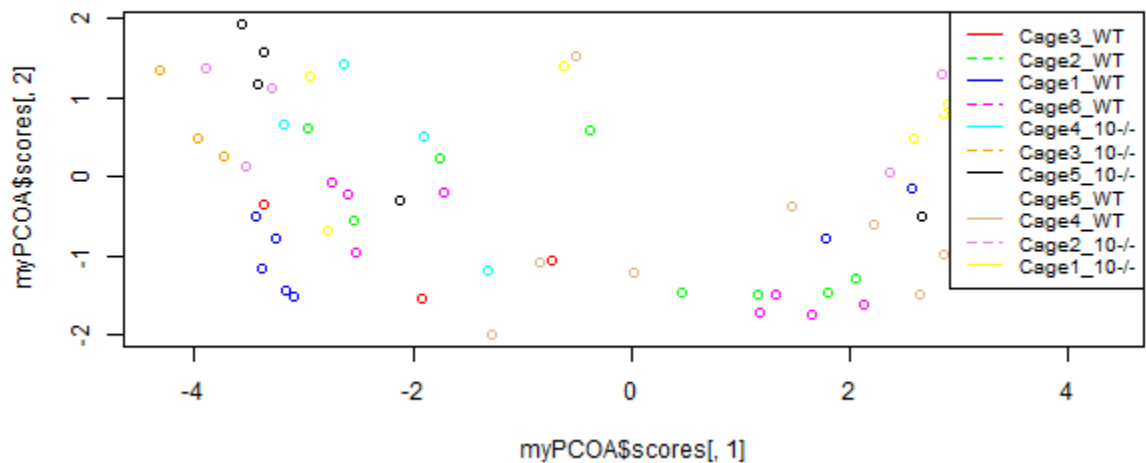
	Comp.1	Comp.2	Comp.3	Comp.4	Comp.5	Comp.6
Standard deviation	2.7823152	1.0369556	0.78000782	0.65498488	0.62347863	0.0981108931
Proportion of Variance	0.7550755	0.1048813	0.05934383	0.04184468	0.03791585	0.0009388845
Cumulative Proportion	0.7550755	0.8599567	0.91930058	0.96114526	0.99906112	1.0000000000

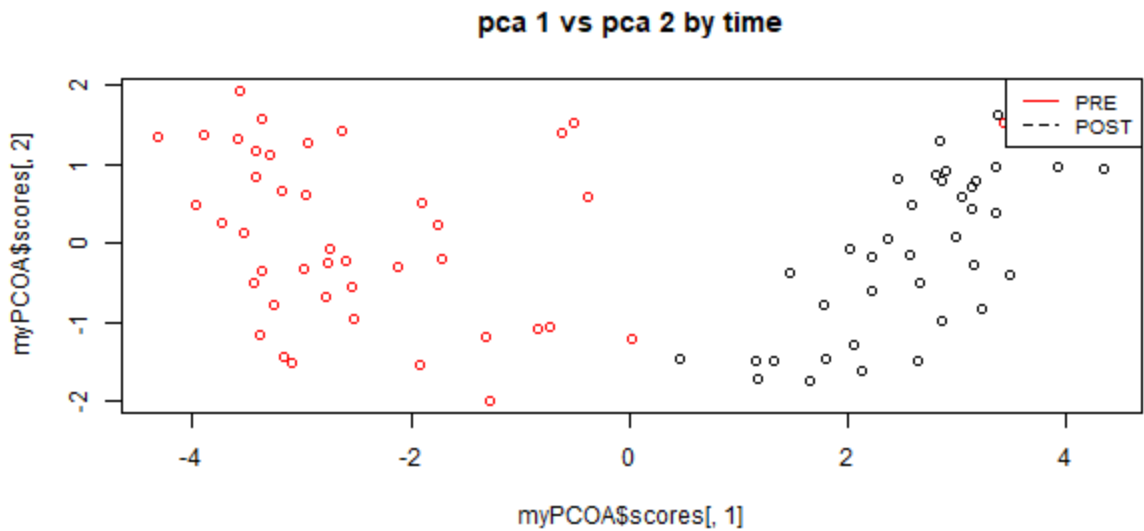
### 2) Problem (2): Graph PCA1 vs. PCA2. Make three versions of the graph. One colored by genotype, one colored by cage and one colored by timepoint (pre-vs-post).

pca 1 vs pca 2 by genotype



pca 1 vs pca 2 by cage





- 3) **Problem (3):** Fill in the following table for p-values testing the null hypothesis for PCA 1 and 2. For cage, use a way one-ANOVA. For genotype and timepoint (“pre” vs “post”) use a t-test.

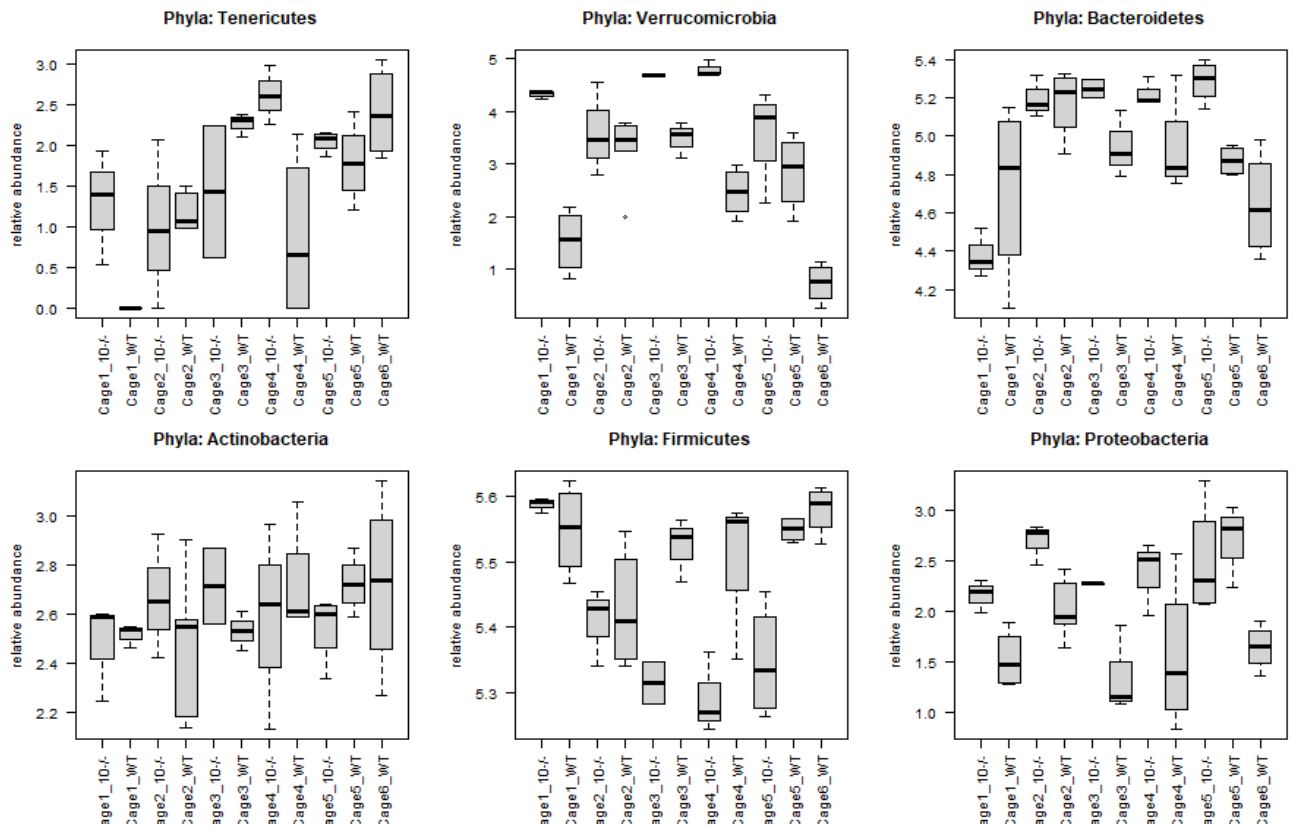
	PCA1	PCA2
Cage	<b>0.992</b>	<b>1.63e-07</b>
Genotype	<b>0.926</b>	<b>3.54e-10</b>
Time (pre vs. post)	<b>2.40e-31</b>	<b>0.428</b>

**Which variable seems to be most associated with the first PCA axis?** Time (pre vs. post) seems to be the most associated one with first PCA axis, and with cage and genotype the PCA 1 does not seem to vary or does not seem to have a slope which is significant.

**Which variable is most associated with the second PCA axis?** Genotype, even though for second PCA axis both cage and genotype are significant but it seems like genotype is a bit more associated with the second PCA axis than others.

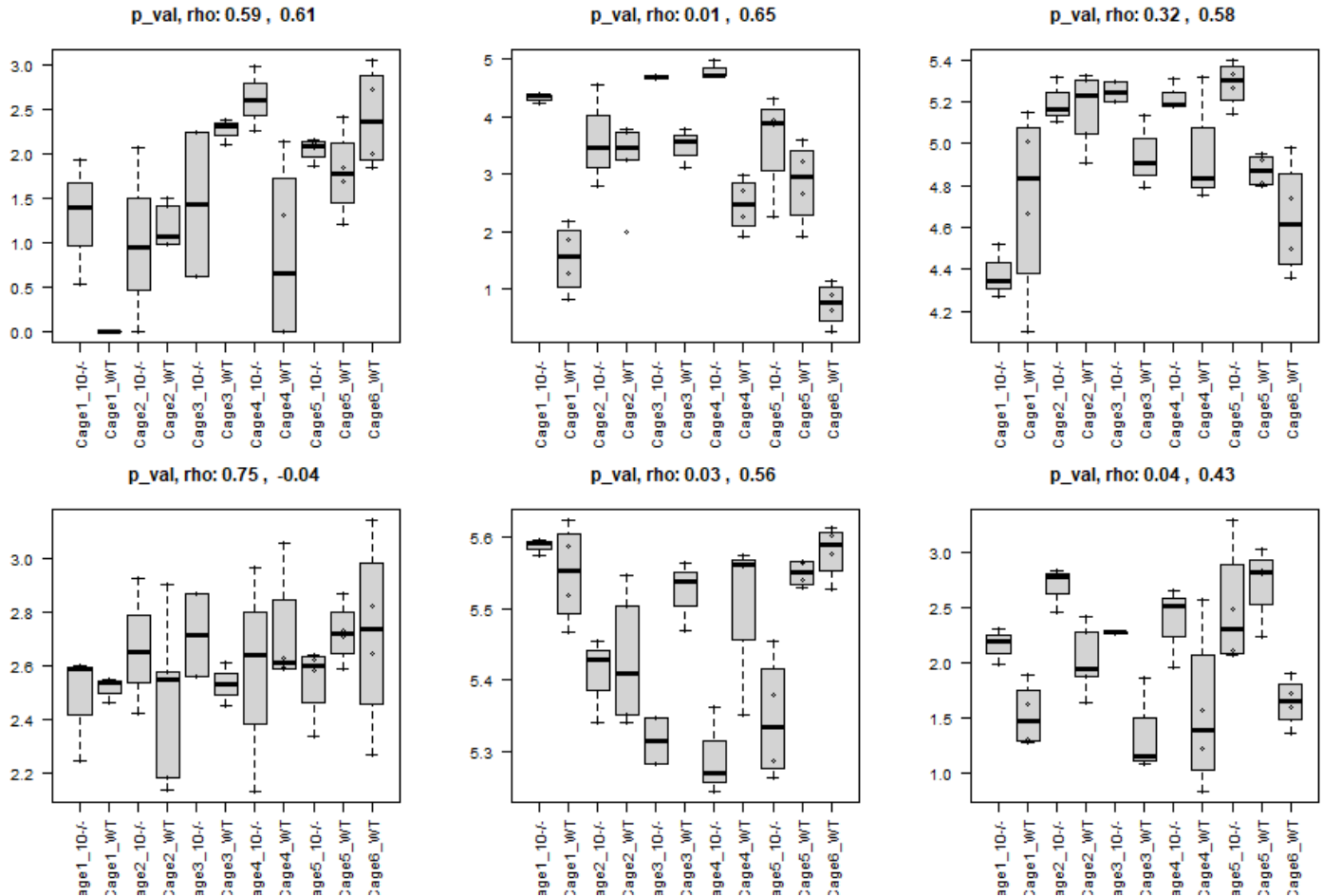
**Does cage seem to be having an effect on these data?** Not on the first PCA axis but Yes, cage seems to be having an (significant) effect on the second PCA axis.

4) Problem (4A): For each phyla, graph the relative abundance of that phyla vs. cage.



**Does there appear to be a cage effect across different phyla?** Yes, from the above boxplots it seems like there is a strong cage effect across different phyla.

- 5) **Problem (4B):** For each phyla build a mixed linear model with genotype as the fixed variable and cage as a random variable. Report the intraclass correlation coefficient for each phyla.



**Are there any phyla that are significantly different for genotype in the mixed model at a 10% false discovery rate?**

```
print(length(p_value_lme)) ## 6
print(sum(p_value_lme < 0.10)) ## unadjusted
p_value_lme_adj <- p.adjust(p_value_lme, method='BH')
print(sum(p_value_lme_adj < 0.10)) ## 3 for adjusted
```

Yes, there seems to be 3 phyla that are significantly different for genotype in the mixed model at a 10% FDR out of 6.

