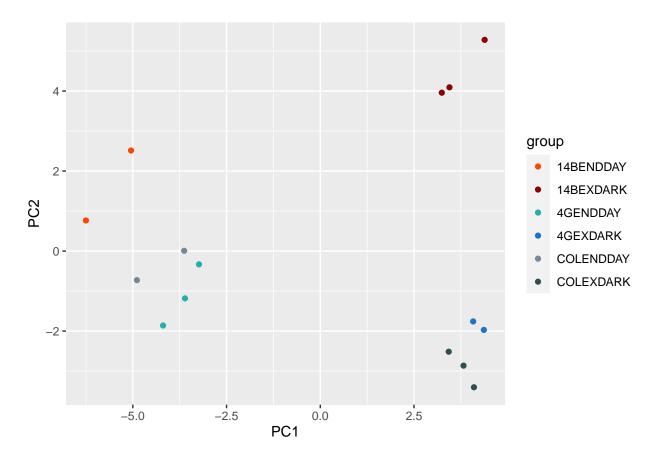
```
## Filtering counts for assigned geneset
lgNorm[rownames(lgNorm) %in% gene_vec,] %>% data.frame(check.names = FALSE) -> lgGo

## PCA on log normalized counts, assigned genes
pca_lggo <- prcomp(t(lgGo))

pca_lggo_dat <- data.frame(pca_lggo$x[,1:2])
pca_lggo_dat$group <- sampleAnnotation[rownames(pca_lggo_dat), "group"]
pca_lggo_dat$sample <- rownames(pca_lggo_dat)

pca_lggo_dat %>% ggplot(aes(x = PC1, y = PC2, color = group, label = sample)) + geom_point() + scale_co
```



```
## Clustered heatmap, assigned genes
library(pheatmap)
heatData <- lgGo -rowMeans(lgGo)
heatData[heatData > 2] = 2
heatData[heatData < -2] = -2

pdf("BIO321G-RNAseq.pdf", width = 24, height = 8)
pheatmap(heatData, color = heatPalette, clustering_method = "average", labels_row = geneNamesAndDescrip dev.off()
## pdf
## 3</pre>
```

```
## Expression stripchart, assigned genes
data.frame(results(dds)[5]) -> pmat

subset(pmat, rownames(pmat) %in% gene_vec) %>% arrange(pvalue) %>% head(9) %>% rownames -> top_9_exp

lgGo[rownames(lgGo) %in% top_9_exp,] -> lll
stripchart321g(data = lll, sampleAnnotation = sampleAnnotation) -> stripgg
stripgg
```

