LDA flu test

```
library(MASS)
library(ggplot2)
library(scales)
flu virus all test set 1
# label and seq data
uniflna_test1 <-
  read.csv("/home/brian/Documents/data_mining/flu_project/uniflna_test1.csv", na.strings="")
# comparisons from sourmash
uniflna_test1_cmp <-
  read.csv("/home/brian/Documents/data_mining/flu_project/uniflna_test1_cmp.csv")
# Label the rows
rownames(uniflna_test1_cmp) <- colnames(uniflna_test1_cmp)</pre>
# add gene column
uniflna test1 wgs <- uniflna test1 cmp
uniflna_test1_wgs$gene <- uniflna_test1$gene
uniflna_test1_wgs$segment <- factor(uniflna_test1$segment)
# Transform for plotting
uniflna_test1_cmp_mat <- as.matrix(uniflna_test1_cmp)</pre>
lda <- lda(gene ~ .,
           uniflna_test1_wgs[,-1002])
## Warning in lda.default(x, grouping, ...): variables are collinear
prop.lda = lda$svd^2/sum(lda$svd^2)
plda <- predict(object = lda,</pre>
                newdata = uniflna_test1_wgs)
dataset = data.frame(gene = uniflna_test1_wgs[,1001],
                     1da = plda$x)
p1 <- ggplot(dataset) + geom_point(aes(lda.LD1, lda.LD2, colour = gene), size=2, alpha=0.2) +
  #theme_minimal() +
  labs(x = paste("LD1 (", percent(prop.lda[1]), ")", sep=""),
       y = paste("LD2 (", percent(prop.lda[2]), ")", sep="")) +
  guides(colour = guide_legend(override.aes = list(alpha = 1)))
# https://stackoverflow.com/questions/5290003/how-to-set-legend-alpha-with-qqplot2
# ggsave("gene_LDA_plot.png", plot = p1, width = 7, height = 4, dpi = 200)
р1
```



