```
knitr::opts_chunk$set(echo = TRUE)
load("lattice.RData")
library(ggplot2)
## Warning: package 'ggplot2' was built under R version 4.1.2
## Warning in as.POSIX1t.POSIXct(Sys.time()): unable to identify current timezone 'H':
## please set environment variable 'TZ'
library(devtools)
## Warning: package 'devtools' was built under R version 4.1.2
## Loading required package: usethis
## Warning: package 'usethis' was built under R version 4.1.2
library(ade4)
## Warning: package 'ade4' was built under R version 4.1.2
library(ggbiplot)
## Loading required package: plyr
## Warning: package 'plyr' was built under R version 4.1.2
## Attaching package: 'plyr'
## The following object is masked _by_ '.GlobalEnv':
##
##
       ozone
## Loading required package: scales
## Warning: package 'scales' was built under R version 4.1.2
## Loading required package: grid
library(viridis)
## Warning: package 'viridis' was built under R version 4.1.2
## Loading required package: viridisLite
## Warning: package 'viridisLite' was built under R version 4.1.2
```

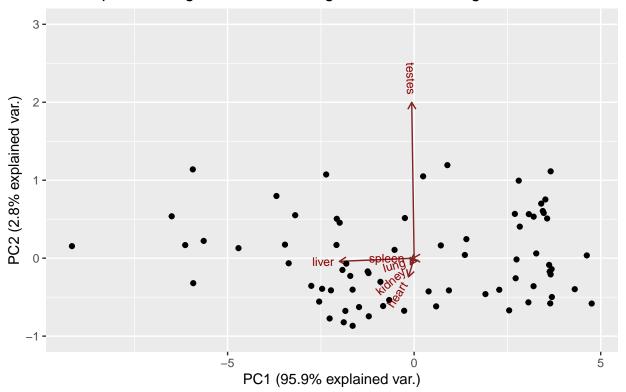
```
##
## Attaching package: 'viridis'
## The following object is masked from 'package:scales':
##
##
      viridis_pal
library(tidyverse)
## Warning: package 'tidyverse' was built under R version 4.1.2
## -- Attaching packages ------ 1.3.1 --
## v tibble 3.1.6
                    v dplyr 1.0.7
## v tidyr 1.1.4 v stringr 1.4.0
          2.1.1
## v readr
                    v forcats 0.5.1
          0.3.4
## v purrr
## Warning: package 'tibble' was built under R version 4.1.2
## Warning: package 'tidyr' was built under R version 4.1.2
## Warning: package 'readr' was built under R version 4.1.2
## Warning: package 'purrr' was built under R version 4.1.2
## Warning: package 'dplyr' was built under R version 4.1.2
## Warning: package 'stringr' was built under R version 4.1.2
## Warning: package 'forcats' was built under R version 4.1.2
## -- Conflicts ----- tidyverse_conflicts() --
## x dplyr::arrange()
                       masks plyr::arrange()
## x readr::col_factor() masks scales::col_factor()
## x purrr::compact()
                       masks plyr::compact()
## x dplyr::count()
                       masks plyr::count()
## x purrr::discard()
                       masks scales::discard()
## x dplyr::failwith()
                      masks plyr::failwith()
## x dplyr::filter()
                       masks stats::filter()
## x dplyr::id()
                       masks plyr::id()
## x dplyr::lag()
                       masks stats::lag()
## x dplyr::mutate()
                       masks plyr::mutate()
## x dplyr::rename()
                       masks plyr::rename()
## x dplyr::summarise() masks plyr::summarise()
## x dplyr::summarize() masks plyr::summarize()
library(dplyr)
library(ggrepel)
```

 $\mbox{\tt \#\#}$  Warning: package 'ggrepel' was built under R version 4.1.2

```
head(hamster,5)
##
        lung
                heart
                         liver
                                   spleen kidney
                                                     testes
## 1 0.969500 1.352002 8.185496 0.22550020 1.348003 0.3415002
## 2 1.141003 1.499403 8.544507 0.27049980 1.322103 0.4170000
## 3 0.872000 1.268503 6.823009 0.04999999 0.897001 1.2200020
## 4 0.739900 1.270203 9.067509 0.11299980 1.278003 0.8145000
## 5 1.139302 1.496203 9.621993 0.11980000 1.415003 1.1568030
ham_noscale = prcomp(hamster, scale. = FALSE)
ham_noscale$rotation[,1:2]
##
                 PC1
                              PC2
## lung
         -0.04844187 -1.400244e-02
## heart -0.07583873 -1.194474e-01
## liver -0.99409891 -2.013637e-02
## spleen -0.01213091 -1.658741e-05
## kidney -0.04990582 -4.195610e-02
## testes -0.03211681 9.916504e-01
ham_noscale2 = prcomp(hamster, scale. = TRUE)
ham noscale2$rotation[,1:2]
##
                 PC1
                             PC2
         -0.40885549 -0.05704641
## lung
## heart -0.46129490 -0.20800130
## liver -0.48564405 0.08125510
## spleen -0.41489143 0.09751789
## kidney -0.45260553 -0.09380103
## testes -0.08489638 0.96362358
summary(hamster)
##
        lung
                        heart
                                        liver
                                                         spleen
## Min. :0.3216
                   Min.
                          :0.4134 Min. : 1.626
                                                     Min.
                                                            :0.0293
  1st Qu.:0.6220
                    1st Qu.:0.8039
                                    1st Qu.: 3.262
                                                     1st Qu.:0.0657
## Median :0.8416
                   Median :1.0690 Median : 6.439
                                                     Median :0.0896
## Mean
         :0.8457
                    Mean
                         :1.0581
                                    Mean : 6.299
                                                     Mean
                                                          :0.1035
##
   3rd Qu.:1.0260
                    3rd Qu.:1.2763
                                    3rd Qu.: 8.307
                                                     3rd Qu.:0.1283
##
   Max.
         :1.6521
                    Max.
                         :1.8390
                                    Max. :15.492
                                                     Max. :0.2959
##
                        testes
       kidney
## Min.
         :0.8199
                    Min. :0.3082
                    1st Qu.:0.6387
  1st Qu.:1.0751
##
## Median :1.2291
                   Median :0.9761
## Mean :1.2256
                   Mean :1.0900
## 3rd Qu.:1.3642
                    3rd Qu.:1.4862
## Max. :1.7381
                    Max. :2.4244
```

```
## scale = 0 means form biplot
ggbiplot(ham_noscale, scale = 0) + ggtitle("PCA Biplot for weights of hamster organs without scaling")
```

#### PCA Biplot for weights of hamster organs without scaling

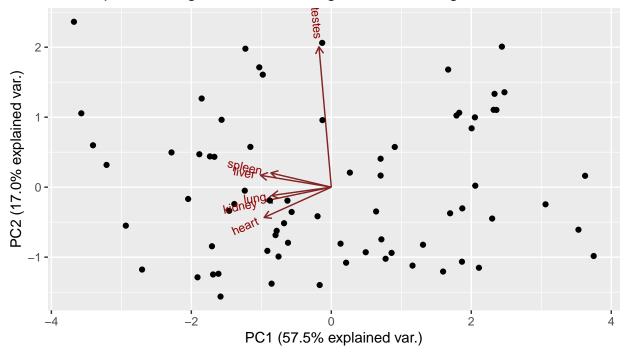


#### ## Q.1)

- 1. For the above plot, we applied PCA on the hamster dataset without scaling the data. We see that all variables are on different scales.
- 2. The first principal component i.e. PC1 shows an explained variance of almost 96%.
- 3. However we see that Testes shows the most variation followed by liver.
- 4. Since we are not scaling the features, we won't be able to take the variation of other features like kidney, heart, spleen and lung.
- 5. This results in capturing most of the variation from Testes and Liver. We can solve this problem if we scale all features.

```
## scale = 0 means form biplot
ggbiplot(ham_noscale2, scale = 0) + ggtitle("PCA Biplot for weights of hamster organs with scaling")
```

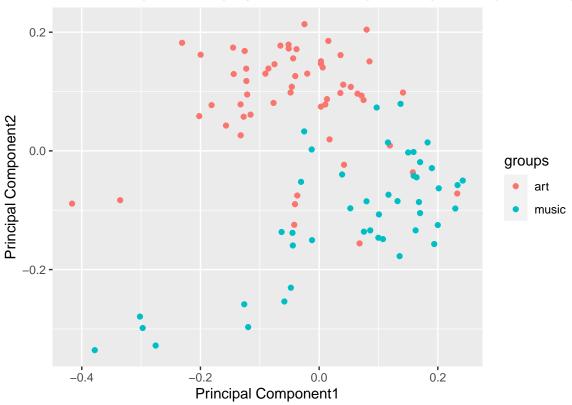
## PCA Biplot for weights of hamster organs with scaling



6. In the above PCA biplot we applied PCA on the scaled data for all organ weights. 7. Compared to the earlier plot wherein we were capturing variation from only two features, here we can capture variation of all the features using two principal components 8. Thus we are able to capture variance from all features or dimensions by projecting it on principal components.

# Q.2)

## Plot of Sample Points projected on Principal Components(Unscaled)

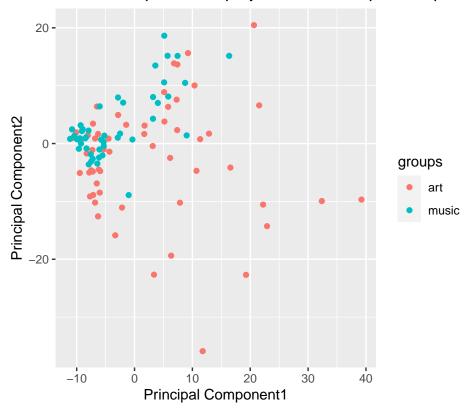


1. In the above PCA plot for sample points, we did not apply scaling on the features. 2. We see that we are able to distinguish art and music articles because of the variation of the features from the data. that is captured by our two main principal components 3. We have differentiated the art and music articles on the basis of seperate colors to represent these clusters.

```
nyt_noscale2 = prcomp(nyt_pca1 , scale. = TRUE)
```

ggbiplot(nyt\_noscale2, scale = 0, var.axes = FALSE, groups = nyt\_pca\$class.labels)+ ggtitle("Plot of Sar xlab("Principal Component1") + ylab("Principal Component2")

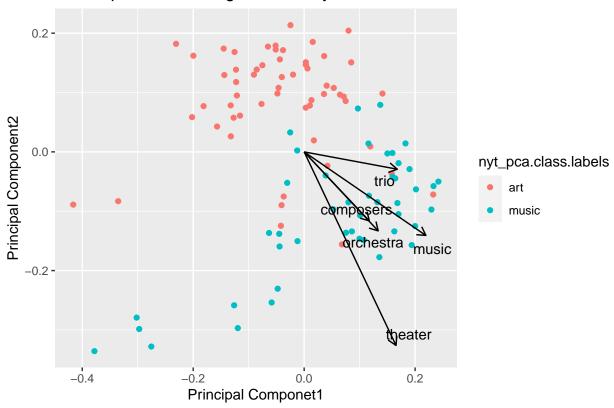
#### Plot of Sample Points projected on Principal Components(Scaled



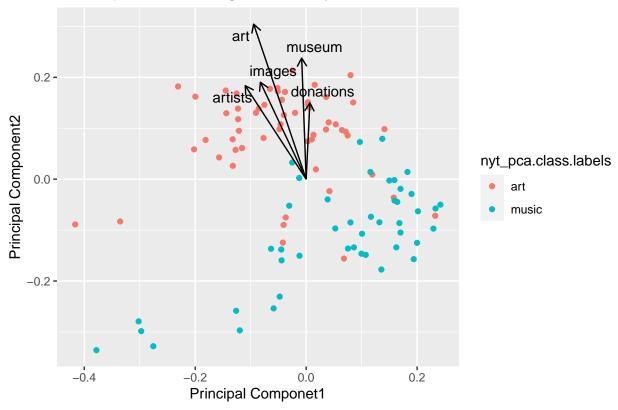
- 4. In the above PCA plot for sample points, we applied scaling of the features.
- 5. Here we see that the clusters for art and music articles are focussed towards same location and as a result we are not able to differentiate between the two article classes.
- 6. Since the data is already normalized there is no need to apply normalization the second time to the data
- 7. Thus the scale is same for all features and therefore using PCA on unscaled features is more beneficial.
- 8. We can say that using our main two principal components, we are able to capture variation in features with respect to art and music articles.

#### Q.3)

## PCA Biplot with loadings ordered by PC1







We created two biplots. One biplot was used for plotting largest loadings according to principal component1 i.e PC1 and other biplot was used for plotting largest loadings according to principal component2 i.e PC2

- 1. From the plot of largest loadings with respect to PC1, we see that variation of features that are most commonly found in music articles is captured by PC1.
- 2. From the plot of largest loadings with respect to PC2, we see that variation of features that are most commonly found in art articles is captured by PC2.
- 3. From this, we can conclude that PC1 and PC2 capture variation from only a particular type of article class at a time. So music and art articles must be having some different words.