

class08_mini_project

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Unsupervised LEarning Analysis of Human Breast Cancer Cells.

Our data

```
# Save your input data file into your Project directory
fna.data <- "WisconsinCancer.csv"

# Complete the following code to input the data and store as wisc.df
wisc.df <- read.csv(fna.data, row.names=1)
head(wisc.df)
```

	diagnosis	radius_mean	texture_mean	perimeter_mean	area_mean
842302	M	17.99	10.38	122.80	1001.0
842517	M	20.57	17.77	132.90	1326.0
84300903	M	19.69	21.25	130.00	1203.0
84348301	M	11.42	20.38	77.58	386.1
84358402	M	20.29	14.34	135.10	1297.0
843786	M	12.45	15.70	82.57	477.1

	smoothness_mean	compactness_mean	concavity_mean	concave.points_mean
842302	0.11840	0.27760	0.3001	0.14710
842517	0.08474	0.07864	0.0869	0.07017
84300903	0.10960	0.15990	0.1974	0.12790
84348301	0.14250	0.28390	0.2414	0.10520
84358402	0.10030	0.13280	0.1980	0.10430
843786	0.12780	0.17000	0.1578	0.08089

	symmetry_mean	fractal_dimension_mean	radius_se	texture_se	perimeter_se
842302	0.2419	0.07871	1.0950	0.9053	8.589
842517	0.1812	0.05667	0.5435	0.7339	3.398
84300903	0.2069	0.05999	0.7456	0.7869	4.585
84348301	0.2597	0.09744	0.4956	1.1560	3.445
84358402	0.1809	0.05883	0.7572	0.7813	5.438
843786	0.2087	0.07613	0.3345	0.8902	2.217

	area_se	smoothness_se	compactness_se	concavity_se	concave.points_se
842302	153.40	0.006399	0.04904	0.05373	0.01587
842517	74.08	0.005225	0.01308	0.01860	0.01340
84300903	94.03	0.006150	0.04006	0.03832	0.02058
84348301	27.23	0.009110	0.07458	0.05661	0.01867
84358402	94.44	0.011490	0.02461	0.05688	0.01885
843786	27.19	0.007510	0.03345	0.03672	0.01137

	symmetry_se	fractal_dimension_se	radius_worst	texture_worst
842302	0.03003		0.006193	25.38
842517	0.01389		0.003532	24.99
84300903	0.02250		0.004571	23.57
84348301	0.05963		0.009208	14.91
84358402	0.01756		0.005115	22.54
843786	0.02165		0.005082	15.47

	perimeter_worst	area_worst	smoothness_worst	compactness_worst
842302	184.60	2019.0	0.1622	0.6656
842517	158.80	1956.0	0.1238	0.1866
84300903	152.50	1709.0	0.1444	0.4245
84348301	98.87	567.7	0.2098	0.8663
84358402	152.20	1575.0	0.1374	0.2050
843786	103.40	741.6	0.1791	0.5249

	concavity_worst	concave.points_worst	symmetry_worst
842302	0.7119	0.2654	0.4601
842517	0.2416	0.1860	0.2750
84300903	0.4504	0.2430	0.3613
84348301	0.6869	0.2575	0.6638
84358402	0.4000	0.1625	0.2364
843786	0.5355	0.1741	0.3985

	fractal_dimension_worst
842302	0.11890
842517	0.08902
84300903	0.08758
84348301	0.17300
84358402	0.07678
843786	0.12440

Q1. How many observations/samples/patients/rows?

There are 569 individuals in this dataset.

Q2. What is in the 'diagnosis' column? How many of each type?

```
sum(wisc.df$diagnosis == "M")
```

```
[1] 212
```

```
sum(wisc.df$diagnosis == "B")
```

```
[1] 357
```

Use table.

```
table(wisc.df$diagnosis)
```

```
  B    M  
357 212
```

Q3. How many variables/features in the data are suffixed with `_mean`?

```
length(grep("_mean", colnames(wisc.df), value=TRUE))
```

```
[1] 10
```

Q. How many variables/dimensions we have?

```
ncol(wisc.df)
```

```
[1] 31
```

Save the diagnosis for reference later

```
diagnosis <- as.factor(wisc.df$diagnosis)  
head(diagnosis)
```

```
[1] M M M M M M  
Levels: B M
```

and remove or exclude this (diagnosis) column from any of our analysis.

```
wisc.data <- wisc.df[,-1]
print(head(wisc.data))
```

	radius_mean	texture_mean	perimeter_mean	area_mean	smoothness_mean
842302	17.99	10.38	122.80	1001.0	0.11840
842517	20.57	17.77	132.90	1326.0	0.08474
84300903	19.69	21.25	130.00	1203.0	0.10960
84348301	11.42	20.38	77.58	386.1	0.14250
84358402	20.29	14.34	135.10	1297.0	0.10030
843786	12.45	15.70	82.57	477.1	0.12780

	compactness_mean	concavity_mean	concave.points_mean	symmetry_mean
842302	0.27760	0.3001	0.14710	0.2419
842517	0.07864	0.0869	0.07017	0.1812
84300903	0.15990	0.1974	0.12790	0.2069
84348301	0.28390	0.2414	0.10520	0.2597
84358402	0.13280	0.1980	0.10430	0.1809
843786	0.17000	0.1578	0.08089	0.2087

	fractal_dimension_mean	radius_se	texture_se	perimeter_se	area_se
842302	0.07871	1.0950	0.9053	8.589	153.40
842517	0.05667	0.5435	0.7339	3.398	74.08
84300903	0.05999	0.7456	0.7869	4.585	94.03
84348301	0.09744	0.4956	1.1560	3.445	27.23
84358402	0.05883	0.7572	0.7813	5.438	94.44
843786	0.07613	0.3345	0.8902	2.217	27.19

	smoothness_se	compactness_se	concavity_se	concave.points_se
842302	0.006399	0.04904	0.05373	0.01587
842517	0.005225	0.01308	0.01860	0.01340
84300903	0.006150	0.04006	0.03832	0.02058
84348301	0.009110	0.07458	0.05661	0.01867
84358402	0.011490	0.02461	0.05688	0.01885
843786	0.007510	0.03345	0.03672	0.01137

	symmetry_se	fractal_dimension_se	radius_worst	texture_worst
842302	0.03003	0.006193	25.38	17.33
842517	0.01389	0.003532	24.99	23.41
84300903	0.02250	0.004571	23.57	25.53
84348301	0.05963	0.009208	14.91	26.50
84358402	0.01756	0.005115	22.54	16.67
843786	0.02165	0.005082	15.47	23.75

	perimeter_worst	area_worst	smoothness_worst	compactness_worst
842302	184.60	2019.0	0.1622	0.6656
842517	158.80	1956.0	0.1238	0.1866
84300903	152.50	1709.0	0.1444	0.4245

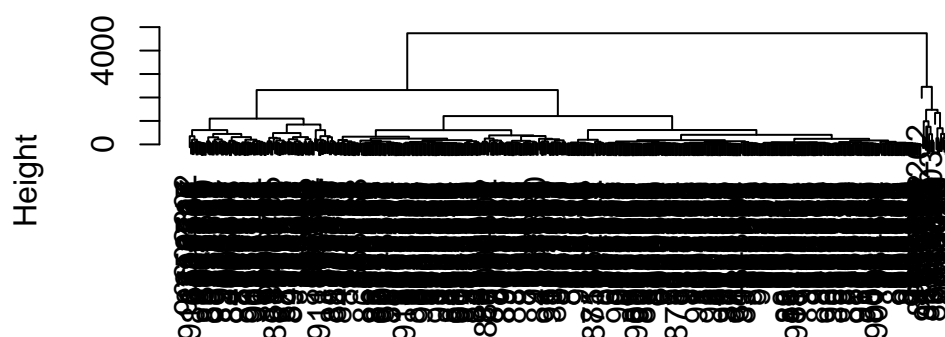
84348301	98.87	567.7	0.2098	0.8663
84358402	152.20	1575.0	0.1374	0.2050
843786	103.40	741.6	0.1791	0.5249
	concavity_worst	concave.points_worst	symmetry_worst	
842302	0.7119	0.2654	0.4601	
842517	0.2416	0.1860	0.2750	
84300903	0.4504	0.2430	0.3613	
84348301	0.6869	0.2575	0.6638	
84358402	0.4000	0.1625	0.2364	
843786	0.5355	0.1741	0.3985	
	fractal_dimension_worst			
842302	0.11890			
842517	0.08902			
84300903	0.08758			
84348301	0.17300			
84358402	0.07678			
843786	0.12440			

Let's try clustering this data:

Hierarchical Clustering with `hclust()`

```
wisc.hc <- hclust(dist(wisc.data))
plot(wisc.hc)
```

Cluster Dendrogram



```
dist(wisc.data)
hclust (*, "complete")
```

Principal Component Analysis

Let's try PCA on this data. Before doing any analysis like we should check if our input data needs to be called first?

side-note: example with mtcars

```
head(mtcars)
```

	mpg	cyl	disp	hp	drat	wt	qsec	vs	am	gear	carb
Mazda RX4	21.0	6	160	110	3.90	2.620	16.46	0	1	4	4
Mazda RX4 Wag	21.0	6	160	110	3.90	2.875	17.02	0	1	4	4
Datsun 710	22.8	4	108	93	3.85	2.320	18.61	1	1	4	1
Hornet 4 Drive	21.4	6	258	110	3.08	3.215	19.44	1	0	3	1
Hornet Sportabout	18.7	8	360	175	3.15	3.440	17.02	0	0	3	2
Valiant	18.1	6	225	105	2.76	3.460	20.22	1	0	3	1

```
apply(mtcars,2,mean)
```

mpg	cyl	disp	hp	drat	wt	qsec
20.090625	6.187500	230.721875	146.687500	3.596563	3.217250	17.848750
vs	am	gear	carb			
0.437500	0.406250	3.687500	2.812500			

```
apply(mtcars,2,sd)
```

mpg	cyl	disp	hp	drat	wt
6.0269481	1.7859216	123.9386938	68.5628685	0.5346787	0.9784574
qsec	vs	am	gear	carb	
1.7869432	0.5040161	0.4989909	0.7378041	1.6152000	

```
#the variance is high, so scaling it helps to bring the values together with less spread s
```

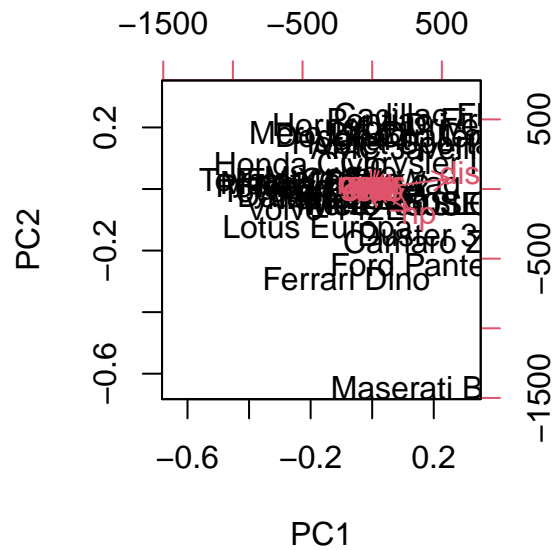
Let's try a PCA on this car dataset.

```
pc<- prcomp(mtcars)
summary(pc)
```

Importance of components:

	PC1	PC2	PC3	PC4	PC5	PC6	PC7
Standard deviation	136.533	38.14808	3.07102	1.30665	0.90649	0.66354	0.3086
Proportion of Variance	0.927	0.07237	0.00047	0.00008	0.00004	0.00002	0.0000
Cumulative Proportion	0.927	0.99937	0.99984	0.99992	0.99996	0.99998	1.0000
	PC8	PC9	PC10	PC11			
Standard deviation	0.286	0.2507	0.2107	0.1984			
Proportion of Variance	0.000	0.0000	0.0000	0.0000			
Cumulative Proportion	1.000	1.0000	1.0000	1.0000			

```
biplot(pc)
```



Scale the mtcars.

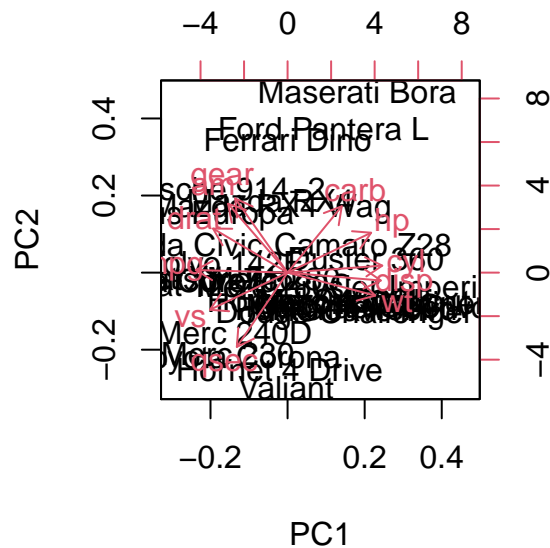
```
pc.scale<- prcomp(mtcars, scale= TRUE)
summary(pc.scale)
```

Importance of components:

	PC1	PC2	PC3	PC4	PC5	PC6	PC7
Standard deviation	2.5707	1.6280	0.79196	0.51923	0.47271	0.46000	0.3678
Proportion of Variance	0.6008	0.2409	0.05702	0.02451	0.02031	0.01924	0.0123
Cumulative Proportion	0.6008	0.8417	0.89873	0.92324	0.94356	0.96279	0.9751

	PC8	PC9	PC10	PC11
Standard deviation	0.35057	0.2776	0.22811	0.1485
Proportion of Variance	0.01117	0.0070	0.00473	0.0020
Cumulative Proportion	0.98626	0.9933	0.99800	1.0000

```
biplot(pc.scale)
```

We want to scale, because some data have different spreads, so we want to scale to reduce that. We can check the plot before scaling and after scaling to check if we need to do that. Scale helps to reduce the SD.

Back to our cancer data set.

Do we need to scale this dataset? Yes, we do because the spread is very different.

```
# Check column means and standard deviations
colMeans(wisc.data)
```

radius_mean	texture_mean	perimeter_mean
1.412729e+01	1.928965e+01	9.196903e+01
area_mean	smoothness_mean	compactness_mean
6.548891e+02	9.636028e-02	1.043410e-01
concavity_mean	concave.points_mean	symmetry_mean
8.879932e-02	4.891915e-02	1.811619e-01
fractal_dimension_mean	radius_se	texture_se
6.279761e-02	4.051721e-01	1.216853e+00
perimeter_se	area_se	smoothness_se
2.866059e+00	4.033708e+01	7.040979e-03

compactness_se	concavity_se	concave.points_se
2.547814e-02	3.189372e-02	1.179614e-02
symmetry_se	fractal_dimension_se	radius_worst
2.054230e-02	3.794904e-03	1.626919e+01
texture_worst	perimeter_worst	area_worst
2.567722e+01	1.072612e+02	8.805831e+02
smoothness_worst	compactness_worst	concavity_worst
1.323686e-01	2.542650e-01	2.721885e-01
concave.points_worst	symmetry_worst	fractal_dimension_worst
1.146062e-01	2.900756e-01	8.394582e-02

```
head(apply(wisc.data,2,sd))
```

radius_mean	texture_mean	perimeter_mean	area_mean
3.52404883	4.30103577	24.29898104	351.91412918
smoothness_mean	compactness_mean		
0.01406413	0.05281276		

```
# Perform PCA on wisc.data by completing the following code
wisc.pr <- prcomp( wisc.data, scale= TRUE)
#wisc.pr
```

How well do the PCS capture the variance in the original data?

```
summary(wisc.pr)
```

Importance of components:

	PC1	PC2	PC3	PC4	PC5	PC6	PC7
Standard deviation	3.6444	2.3857	1.67867	1.40735	1.28403	1.09880	0.82172
Proportion of Variance	0.4427	0.1897	0.09393	0.06602	0.05496	0.04025	0.02251
Cumulative Proportion	0.4427	0.6324	0.72636	0.79239	0.84734	0.88759	0.91010
	PC8	PC9	PC10	PC11	PC12	PC13	PC14
Standard deviation	0.69037	0.6457	0.59219	0.5421	0.51104	0.49128	0.39624
Proportion of Variance	0.01589	0.0139	0.01169	0.0098	0.00871	0.00805	0.00523
Cumulative Proportion	0.92598	0.9399	0.95157	0.9614	0.97007	0.97812	0.98335
	PC15	PC16	PC17	PC18	PC19	PC20	PC21
Standard deviation	0.30681	0.28260	0.24372	0.22939	0.22244	0.17652	0.1731
Proportion of Variance	0.00314	0.00266	0.00198	0.00175	0.00165	0.00104	0.0010
Cumulative Proportion	0.98649	0.98915	0.99113	0.99288	0.99453	0.99557	0.9966
	PC22	PC23	PC24	PC25	PC26	PC27	PC28

Standard deviation	0.16565	0.15602	0.1344	0.12442	0.09043	0.08307	0.03987
Proportion of Variance	0.00091	0.00081	0.0006	0.00052	0.00027	0.00023	0.00005
Cumulative Proportion	0.99749	0.99830	0.9989	0.99942	0.99969	0.99992	0.99997
	PC29	PC30					
Standard deviation	0.02736	0.01153					
Proportion of Variance	0.00002	0.00000					
Cumulative Proportion	1.00000	1.00000					

Q4. From your results, what proportion of the original variance is captured by the first principal components (PC1)?

PC1 captures 44.27%

Q5. How many principal components (PCs) are required to describe at least 70% of the original variance in the data?

PC3 > Q6. How many principal components (PCs) are required to describe at least 90% of the original variance in the data?

PC7

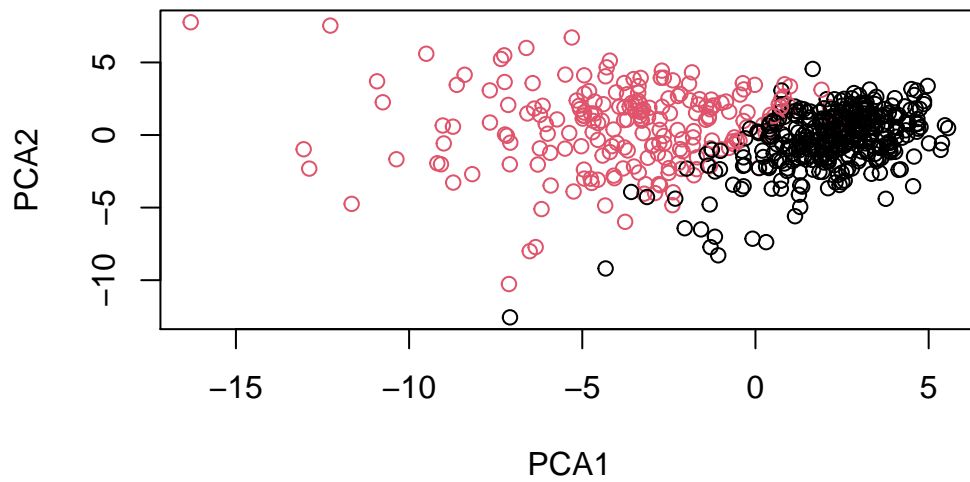
our main PC score plot(aka PC plot, PC1 vs PC2, coordination plot)

```
attributes(wisc.pr)
```

```
$names
[1] "sdev"      "rotation" "center"   "scale"    "x"

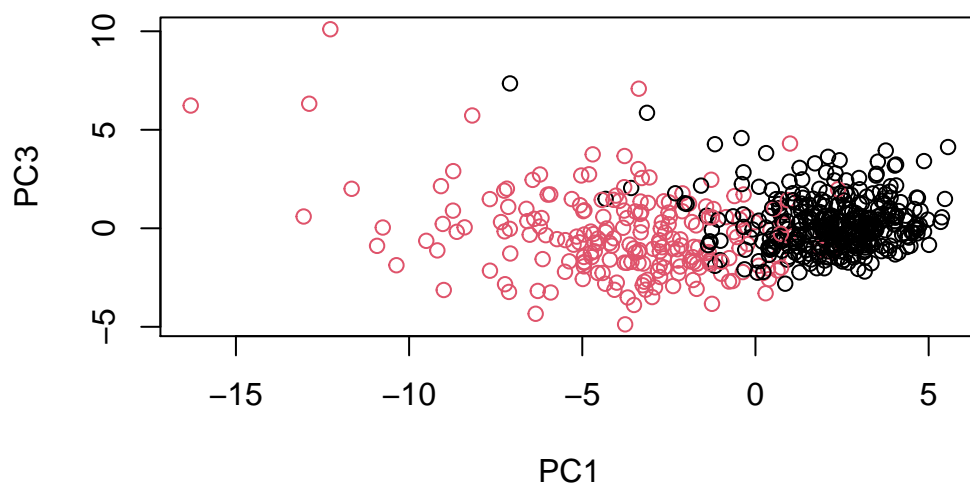
$class
[1] "prcomp"
```

```
biplot(wisc.pr)
```

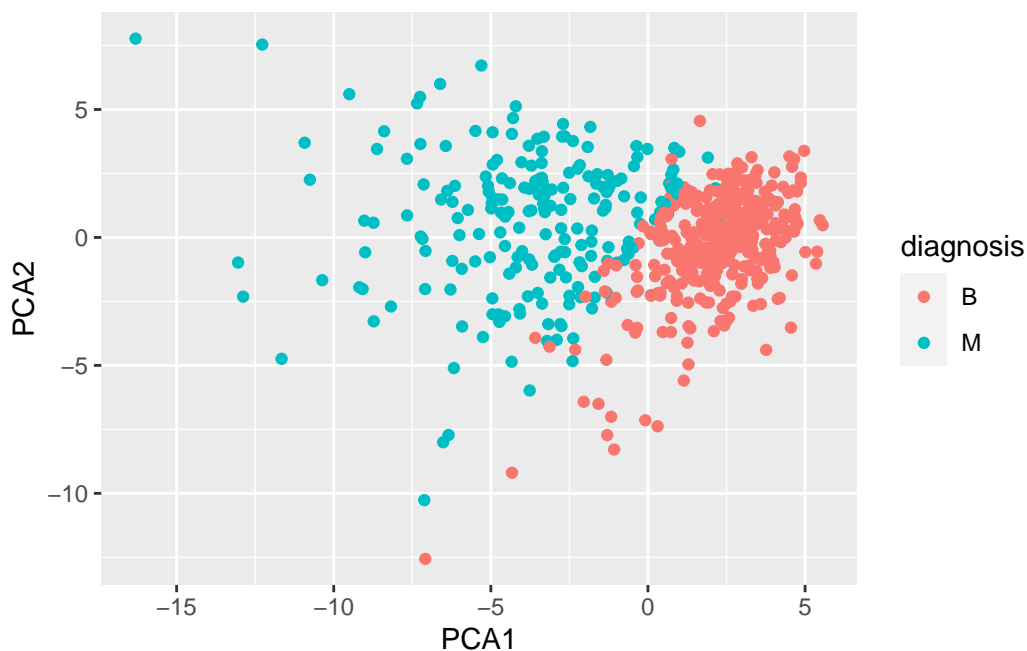
Q8. Q8. Generate a similar plot for principal components 1 and 3. What do you notice about these plots? The variance of the cells lie toward PCA1 (capture by PCA1)

```
plot(wisc.pr$x[, 1], wisc.pr$x[, 3], col = diagnosis,  
     xlab = "PC1", ylab = "PC3")
```



Make a nice ggplot version

```
library(ggplot2)
pc <- as.data.frame(wisc.pr$x)
ggplot(pc, aes(wisc.pr$x[,1], wisc.pr$x[,2], col = diagnosis))+
  geom_point()+
  xlab( "PCA1")+
  ylab( "PCA2")
```



```
v<-summary(wisc.pr)
v$importance[2,]
```

PC1	PC2	PC3	PC4	PC5	PC6	PC7	PC8	PC9	PC10
0.44272	0.18971	0.09393	0.06602	0.05496	0.04025	0.02251	0.01589	0.01390	0.01169
PC11	PC12	PC13	PC14	PC15	PC16	PC17	PC18	PC19	PC20
0.00980	0.00871	0.00805	0.00523	0.00314	0.00266	0.00198	0.00175	0.00165	0.00104
PC21	PC22	PC23	PC24	PC25	PC26	PC27	PC28	PC29	PC30
0.00100	0.00091	0.00081	0.00060	0.00052	0.00027	0.00023	0.00005	0.00002	0.00000

```
# Scale the wisc.data data using the "scale()" function
data.scaled <- scale(wisc.data)
```

Calculate the variance of each principal component by squaring the sdev component of wisc.pr (i.e. `wisc.pr$sdev^2`). Save the result as an object called `pr.var`.

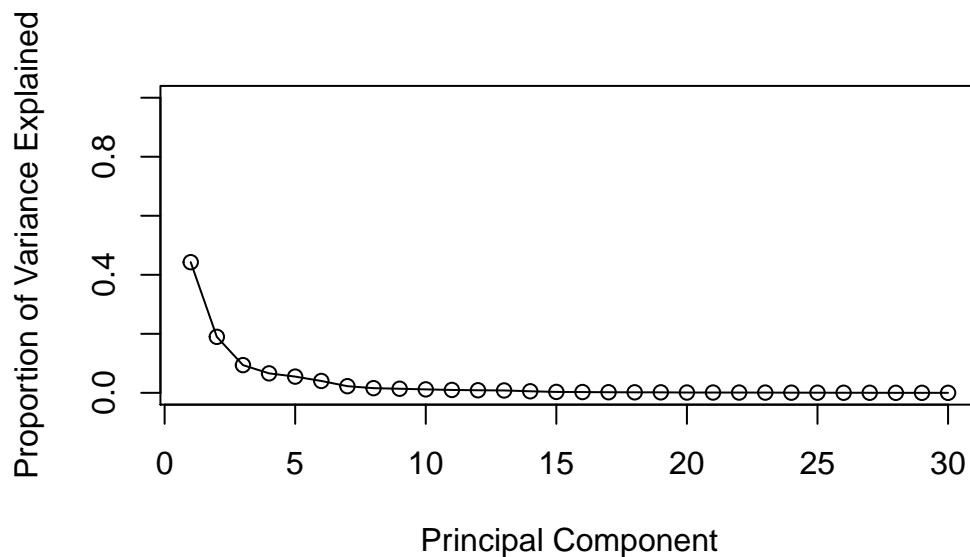
```
# Calculate variance of each component
pr.var <- wisc.pr$sdev^2
head(pr.var)
```

```
[1] 13.281608  5.691355  2.817949  1.980640  1.648731  1.207357
```

Calculate the variance explained by each principal component by dividing by the total variance explained of all principal components. Assign this to a variable called pve and create a plot of variance explained for each principal component.

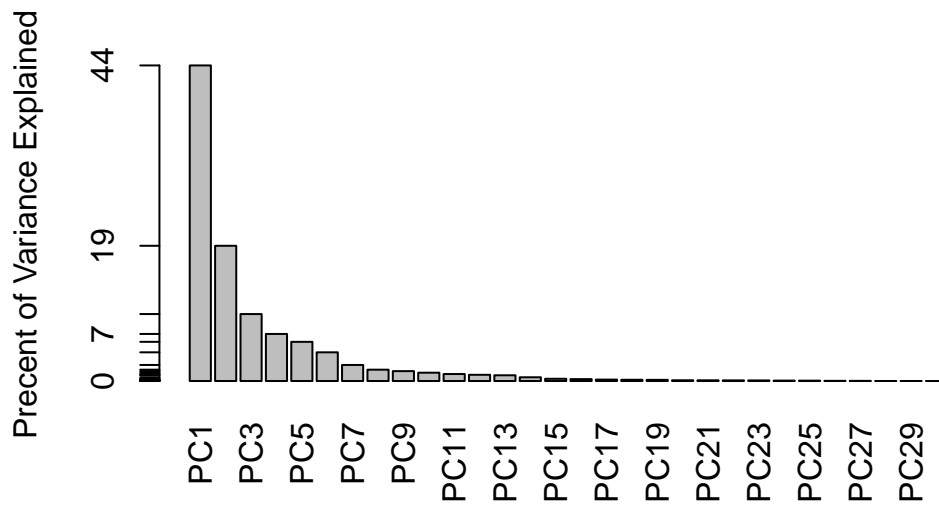
```
# Variance explained by each principal component: pve
pve <- pr.var / sum(pr.var)

# Plot variance explained for each principal component
plot(pve, xlab = "Principal Component",
     ylab = "Proportion of Variance Explained",
     ylim = c(0, 1), type = "o")
```



The elbow is around 0.1.

```
# Alternative scree plot of the same data, note data driven y-axis
barplot(pve, ylab = "Percent of Variance Explained",
       names.arg=paste0("PC",1:length(pve)), las=2, axes = FALSE)
axis(2, at=pve, labels=round(pve,2)*100 )
```

Communicating PCA results: > Q9. For the first principal component, what is the component of the loading vector (i.e. `wisc.pr$rotation[,1]`) for the feature `concave.points_mean`? This tells us how much this original feature contributes to the first PC.

-0.26085376

```
head(wisc.pr$rotation)
```

	PC1	PC2	PC3	PC4	PC5
radius_mean	-0.2189024	0.23385713	-0.008531243	0.04140896	-0.03778635
texture_mean	-0.1037246	0.05970609	0.064549903	-0.60305000	0.04946885
perimeter_mean	-0.2275373	0.21518136	-0.009314220	0.04198310	-0.03737466
area_mean	-0.2209950	0.23107671	0.028699526	0.05343380	-0.01033125
smoothness_mean	-0.1425897	-0.18611302	-0.104291904	0.15938277	0.36508853
compactness_mean	-0.2392854	-0.15189161	-0.074091571	0.03179458	-0.01170397
	PC6	PC7	PC8	PC9	PC10
radius_mean	0.018740790	-0.12408834	0.007452296	-0.223109764	0.09548644
texture_mean	-0.032178837	0.01139954	-0.130674825	0.112699390	0.24093407
perimeter_mean	0.017308445	-0.11447706	0.018687258	-0.223739213	0.08638562
area_mean	-0.001887748	-0.05165343	-0.034673604	-0.195586014	0.07495649
smoothness_mean	-0.286374497	-0.14066899	0.288974575	0.006424722	-0.06929268
compactness_mean	-0.014130949	0.03091850	0.151396350	-0.167841425	0.01293620

	PC11	PC12	PC13	PC14	PC15
radius_mean	-0.04147149	0.05106746	0.01196721	0.059506135	-0.05111877
texture_mean	0.30224340	0.25489642	0.20346133	-0.021560100	-0.10792242
perimeter_mean	-0.01678264	0.03892611	0.04410950	0.048513812	-0.03990294
area_mean	-0.11016964	0.06543751	0.06737574	0.010830829	0.01396691
smoothness_mean	0.13702184	0.31672721	0.04557360	0.445064860	-0.11814336
compactness_mean	0.30800963	-0.10401704	0.22928130	0.008101057	0.23089996
	PC16	PC17	PC18	PC19	PC20
radius_mean	-0.1505839	0.20292425	0.146712338	0.22538466	-0.04969866
texture_mean	-0.1578420	-0.03870612	-0.041102985	0.02978864	-0.24413499
perimeter_mean	-0.1144540	0.19482131	0.158317455	0.23959528	-0.01766501
area_mean	-0.1324480	0.25570576	0.266168105	-0.02732219	-0.09014376
smoothness_mean	-0.2046132	0.16792991	-0.352226802	-0.16456584	0.01710096
compactness_mean	0.1701784	-0.02030771	0.007794138	0.28422236	0.48868633
	PC21	PC22	PC23	PC24	PC25
radius_mean	-0.06857001	-0.07292890	-0.0985526942	-0.18257944	-0.01922650
texture_mean	0.44836947	-0.09480063	-0.0005549975	0.09878679	0.08474593
perimeter_mean	-0.06976904	-0.07516048	-0.0402447050	-0.11664888	0.02701541
area_mean	-0.01844328	-0.09756578	0.0077772734	0.06984834	-0.21004078
smoothness_mean	-0.11949175	-0.06382295	-0.0206657211	0.06869742	0.02895489
compactness_mean	0.19262140	0.09807756	0.0523603957	-0.10413552	0.39662323
	PC26	PC27	PC28	PC29	
radius_mean	-0.12947640	-0.13152667	2.111940e-01	0.211460455	
texture_mean	-0.02455666	-0.01735731	-6.581146e-05	-0.010533934	
perimeter_mean	-0.12525595	-0.11541542	8.433827e-02	0.383826098	
area_mean	0.36272740	0.46661248	-2.725083e-01	-0.422794920	
smoothness_mean	-0.03700369	0.06968992	1.479269e-03	-0.003434667	
compactness_mean	0.26280847	0.09774871	-5.462767e-03	-0.041016774	
	PC30				
radius_mean	0.702414091				
texture_mean	0.000273661				
perimeter_mean	-0.689896968				
area_mean	-0.032947348				
smoothness_mean	-0.004847458				
compactness_mean	0.044674186				

Calculate the (Euclidean) distances between all pairs of observations in the new scaled dataset and assign the result to `data.dist`.

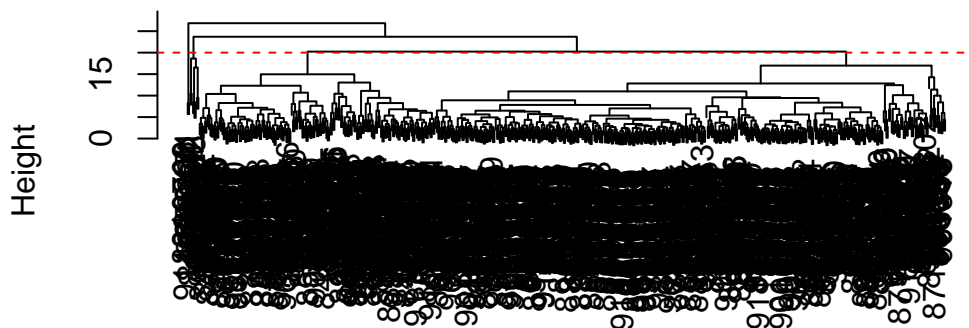
```
data.dist<- dist(data.scaled)
```

Create a hierarchical clustering model using complete linkage. Manually specify the method argument to `hclust()` and assign the results to `wisc.hclust`.

```
wisc.hclust <- hclust(data.dist, method ="complete")
```

```
plot(wisc.hclust)
abline(h=20, col="red", lty=2)
```

Cluster Dendrogram



```
data.dist
hclust (*, "complete")
```

Q10. Using the `plot()` and `abline()` functions, what is the height at which the clustering model has 4 clusters?

H=20

Use `cutree()` to cut the tree so that it has 4 clusters. Assign the output to the variable `wisc.hclust.clusters`.

```
wisc.hclust.clusters <- cutree(wisc.hclust, h=20)
```

We can use the `table()` function to compare the cluster membership to the actual diagnoses.

```
table(wisc.hclust.clusters, diagnosis)
```

```

              diagnosis
wisc.hclust.clusters  B   M
1      12 165
```

2	2	5
3	343	40
4	0	2

Using different methods

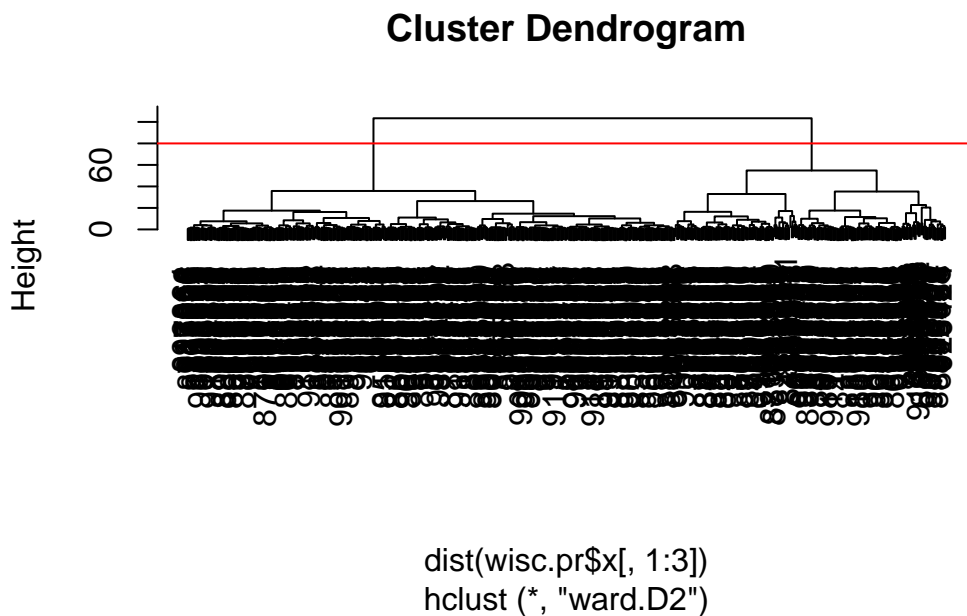
Q12. Which method gives your favorite results for the same data.dist dataset?
Explain your reasoning.

The ward.D2 gives me the better results, as I can distinct the number of clusters clearly.

4. Combining methods

Here we will use the results of PCA as the input to a clustering analysis. We start with 3 PCs.

```
wisc.pr.hclust<- hclust(dist(wisc.pr$x[,1:3]), method ="ward.D2")
plot(wisc.pr.hclust)
abline(h=80, col="red")
```



Q13. How well does the newly created model with four clusters separate out the two diagnoses?

By grps 1 and 2.

```
grps <- cutree(wisc.pr.hclust, h=80)
table(grps)
```

```
grps
  1  2
203 366
```

Q14. How well do the hierarchical clustering models you created in previous sections (i.e. before PCA) do in terms of separating the diagnoses? Again, use the table() function to compare the output of each model (wisc.km\$cluster and wisc.hclust.clusters) with the vector containing the actual diagnoses.

By grps 1 and 2 and diagnosis by B and M.

```
table(grps, diagnosis)
```

```
      diagnosis
grps   B    M
  1  24 179
  2 333  33
```

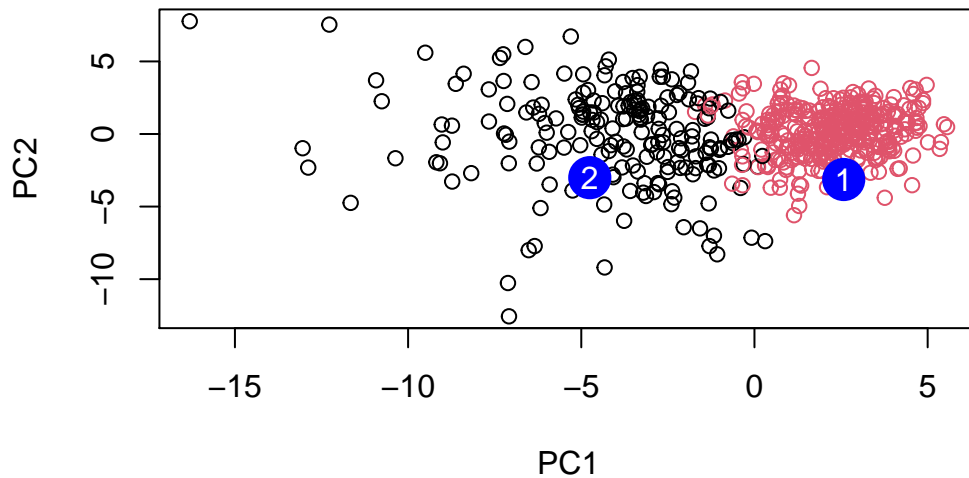
Q14. How well do the hierarchical clustering models you created in previous sections (i.e. before PCA) do in terms of separating the diagnoses? Again, use the table() function to compare the output of each model (wisc.km\$cluster and wisc.hclust.clusters) with the vector containing the actual diagnoses.

Prediction

```
#url <- "new_samples.csv"
url <- "https://tinyurl.com/new-samples-CSV"
new <- read.csv(url)
npc <- predict(wisc.pr, newdata=new)
npc
```

	PC1	PC2	PC3	PC4	PC5	PC6	PC7
[1,]	2.576616	-3.135913	1.3990492	-0.7631950	2.781648	-0.8150185	-0.3959098
[2,]	-4.754928	-3.009033	-0.1660946	-0.6052952	-1.140698	-1.2189945	0.8193031
	PC8	PC9	PC10	PC11	PC12	PC13	PC14
[1,]	-0.2307350	0.1029569	-0.9272861	0.3411457	0.375921	0.1610764	1.187882
[2,]	-0.3307423	0.5281896	-0.4855301	0.7173233	-1.185917	0.5893856	0.303029
	PC15	PC16	PC17	PC18	PC19	PC20	
[1,]	0.3216974	-0.1743616	-0.07875393	-0.11207028	-0.08802955	-0.2495216	
[2,]	0.1299153	0.1448061	-0.40509706	0.06565549	0.25591230	-0.4289500	
	PC21	PC22	PC23	PC24	PC25	PC26	
[1,]	0.1228233	0.09358453	0.08347651	0.1223396	0.02124121	0.078884581	
[2,]	-0.1224776	0.01732146	0.06316631	-0.2338618	-0.20755948	-0.009833238	
	PC27	PC28	PC29	PC30			
[1,]	0.220199544	-0.02946023	-0.015620933	0.005269029			
[2,]	-0.001134152	0.09638361	0.002795349	-0.019015820			

```
g<- as.factor(grps)
plot(wisc.pr$x[,1:2], col=g)
points(npc[,1], npc[,2], col="blue", pch=16, cex=3)
text(npc[,1], npc[,2], c(1,2), col="white")
```



Q16. Which of these new patients should we prioritize for follow up based on your

results?

Patient 2.