BIMM 143

Unsupervised Learning

Lecture 8

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http://thegrantlab.org/bimm143

Recap of Lecture 7

- Reviewed the major steps for R function writing

- Introduced the CRAN & Bioconductor repositories for packages that extend R functionality

- Explored installing and using the ggplot2, bio3d, rgl, rentrez, igraph, blogdown, shiny, and circlize packages

- Discussed how to judge the utility, usability and development status of R packages.

[Muddy Point Feedback Link]

Today's Menu

- Introduction to machine learning
  - <u>Unsupervised</u>, supervised and reinforcement learning

- Clustering
  - K-means clustering
  - Hierarchical clustering
  - Heatmap representations

- Dimensionality reduction, visualization and 'structure' analysis
  - Principal Component Analysis (PCA)

- Hands-on application to cell classification

Types of machine learning

- <u>Unsupervised learning</u>
  - ➡ Finding structure in unlabeled data

- <u>Supervised learning</u>
  - ➡ Making predictions based on labeled data
  - ➡ Predictions like regression or classification

- <u>Reinforcement learning</u>
  - ➡ Making decisions based on past experience

[Muddy Point Feedback Link]

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k-means clustering algorithm

- Breaks observations into *k* pre-defined number of clusters
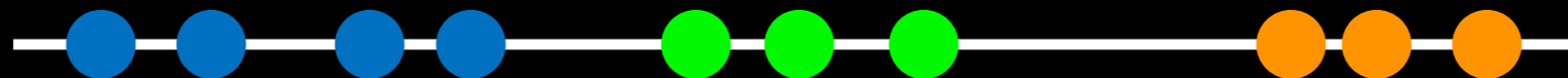
- You define *k* the number of clusters!

k-means clustering algorithm

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- You define *k* the number of clusters!
  - ➡ Imagine you had data that you could plot along a line and you knew you had to put them into *k*=3 "clusters" (e.g. data from three types of tumor cells)

k-means clustering algorithm

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Here your eyes can clearly see 3 natural groupings

k-means clustering algorithm

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- You define *k* the number of clusters!
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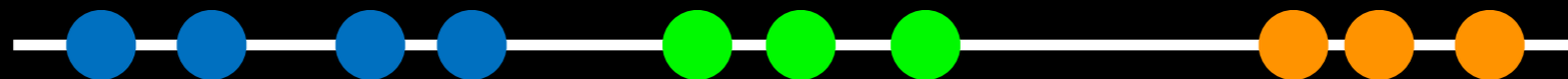
Here your eyes can clearly see 3 natural groupings
How does k-means attempt to define this grouping?
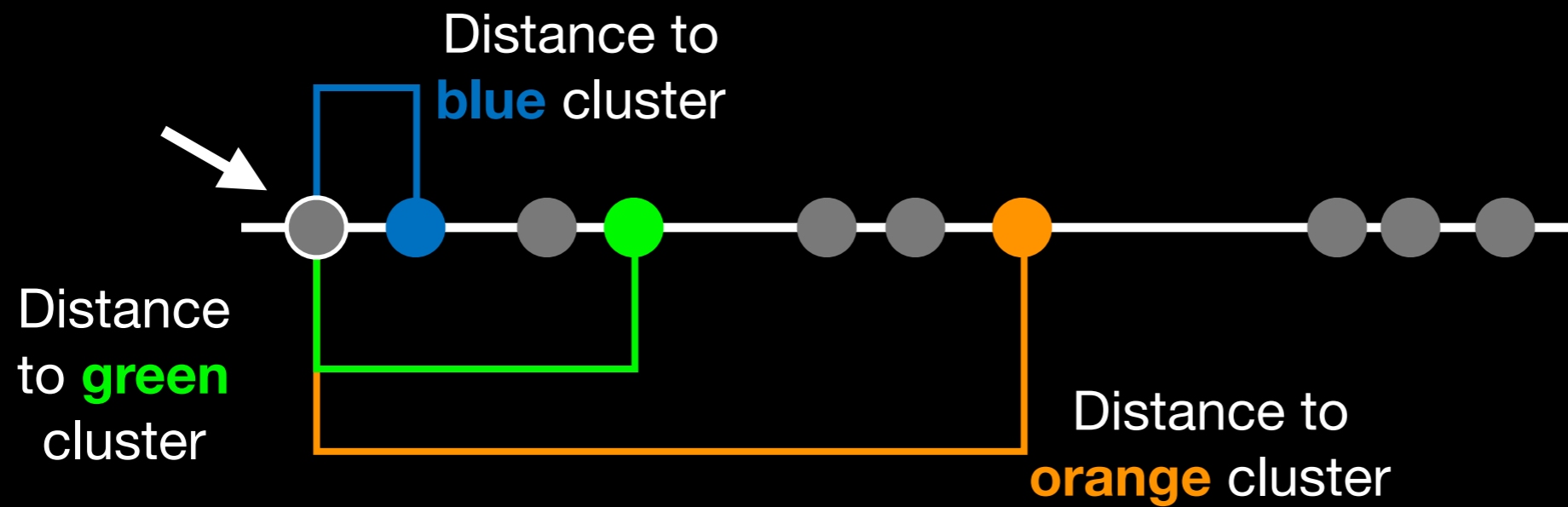
Step 1.
Select *k* (the number of clusters)
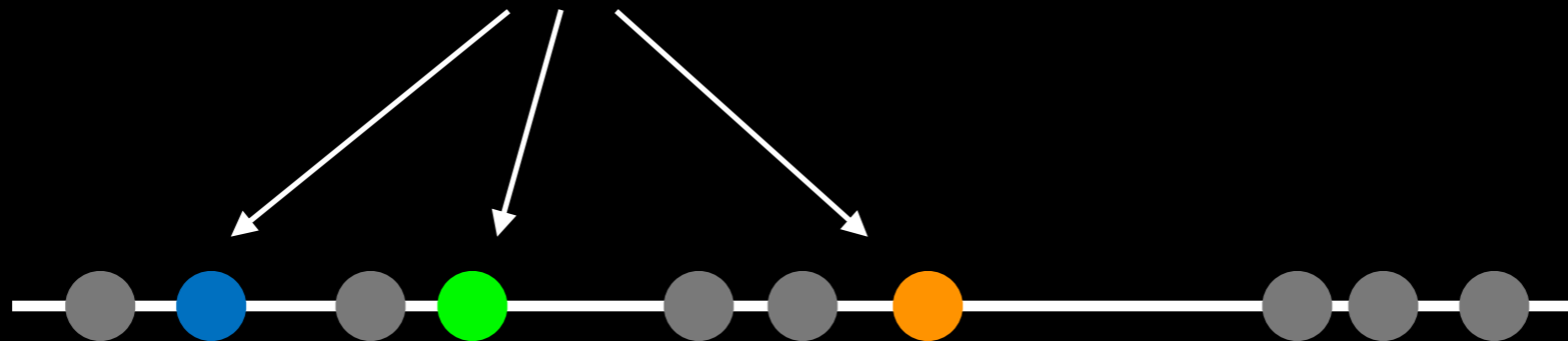
Step 3.

Measure distance between the 1st point and the *k=3* initial clusters

Step 4.

Assign the 1st point to the nearest cluster

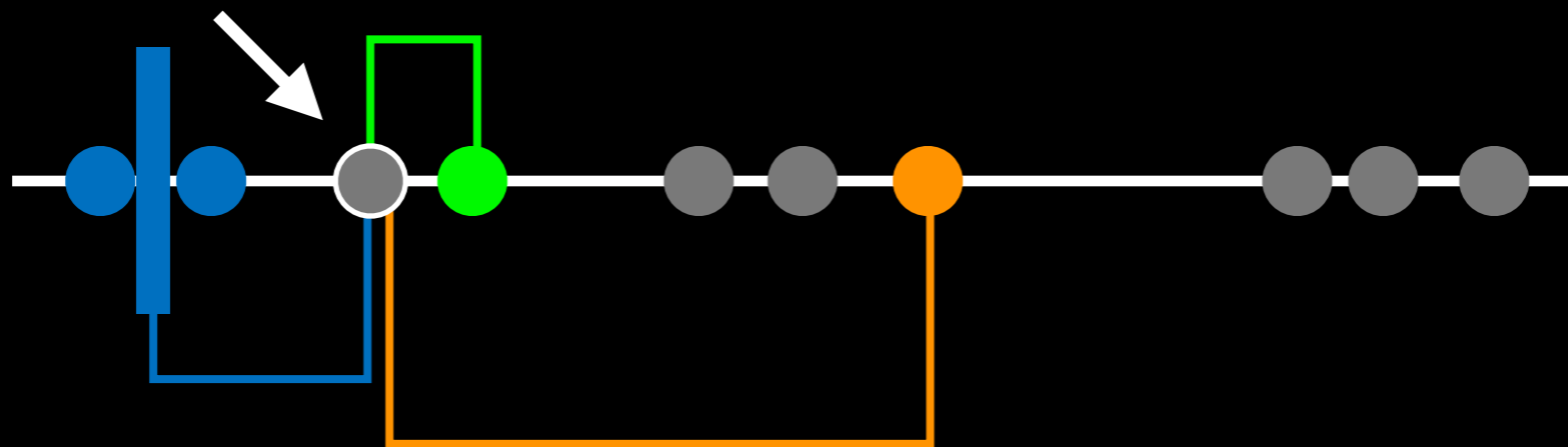# Step 5.

Update cluster centers

Calculate the mean value for the blue cluster including the new point
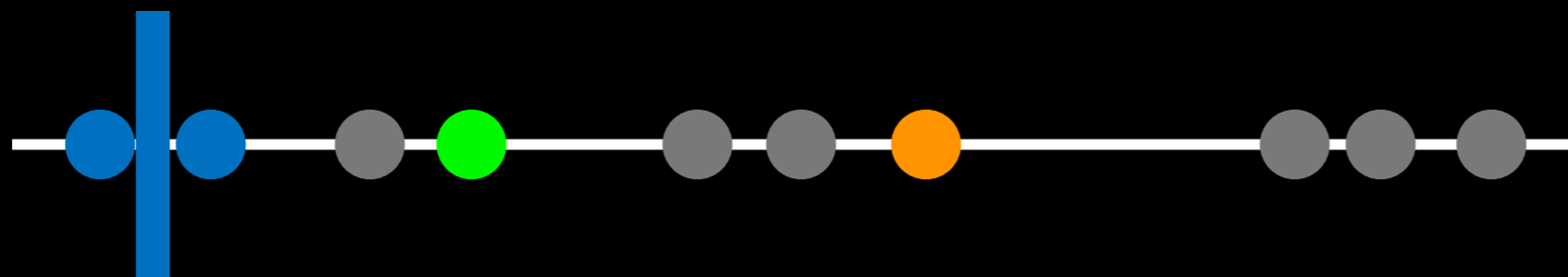
Step 6.

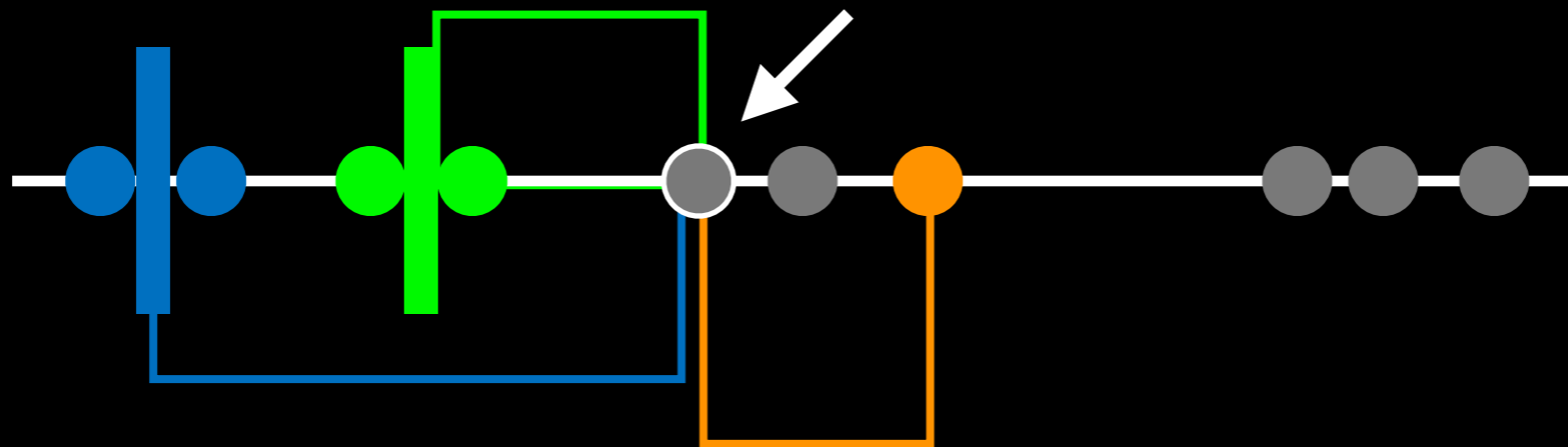Assign next point to closest cluster

Use updated cluster centers for distance calculation

Step 7.

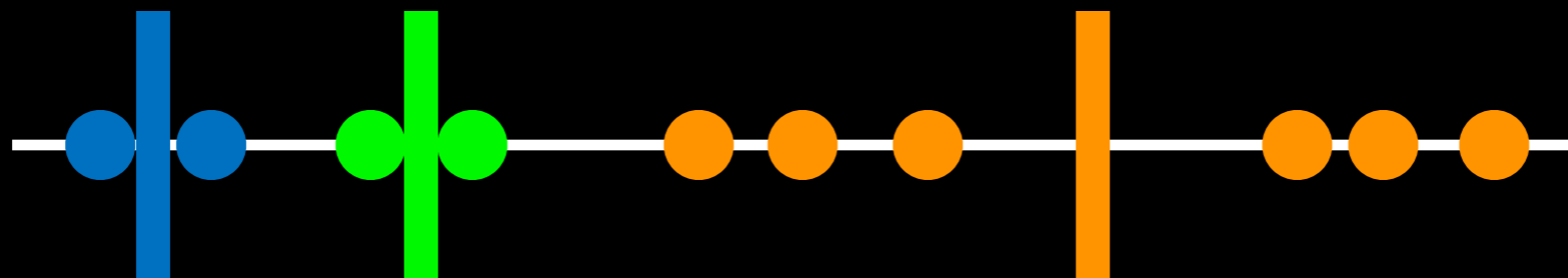Update cluster centers and move to next point

Use updated cluster centers for distance calculation

Step 8.
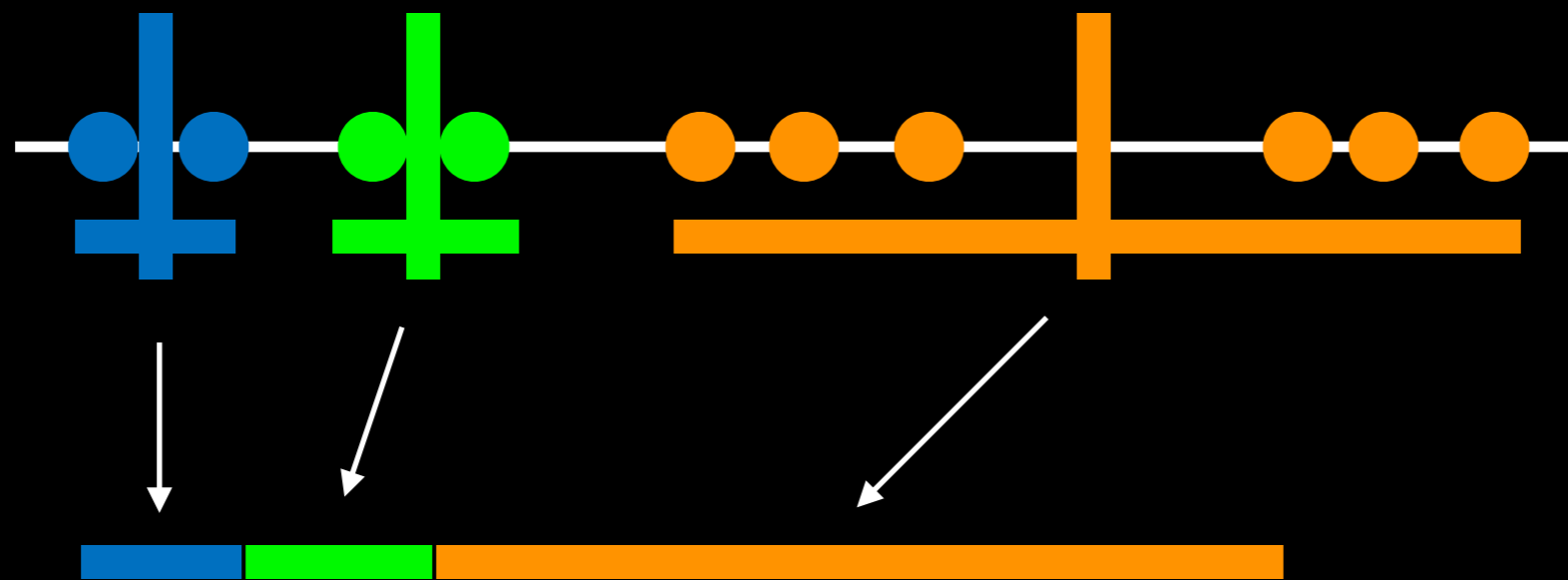Repeat for each point
Each time updating cluster centers

Step 9.

Assess the quality of the clustering by adding up the variation within each cluster
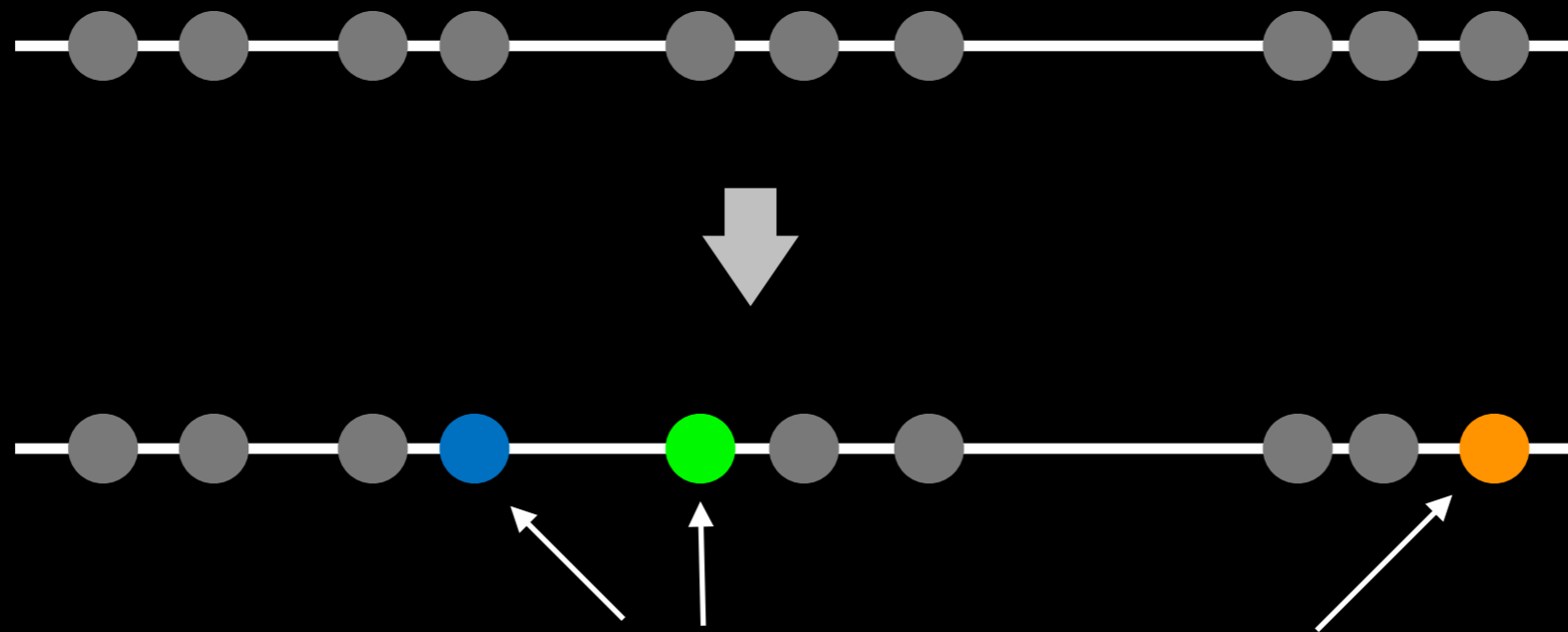
The total variation within clusters

K-means keeps track of these clusters and their total **variance** and then does the whole thing over again with different starting points

Step 10.
Repeat with different starting points
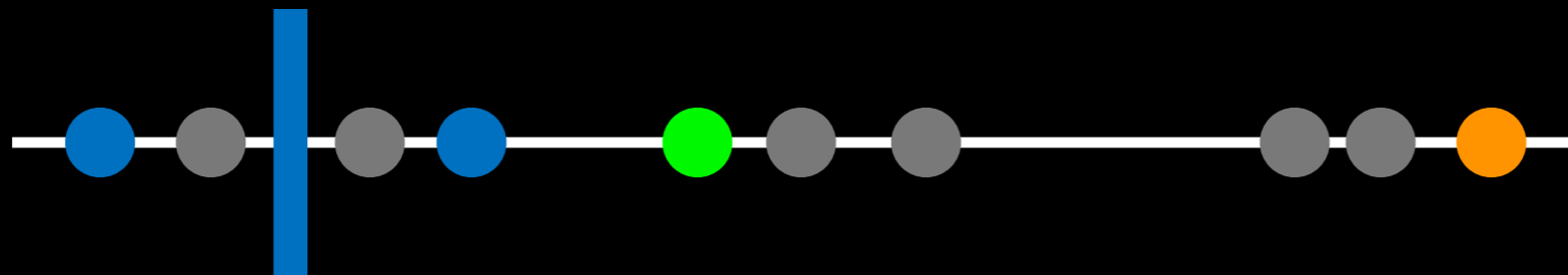Back to the beginning and do all steps over again…
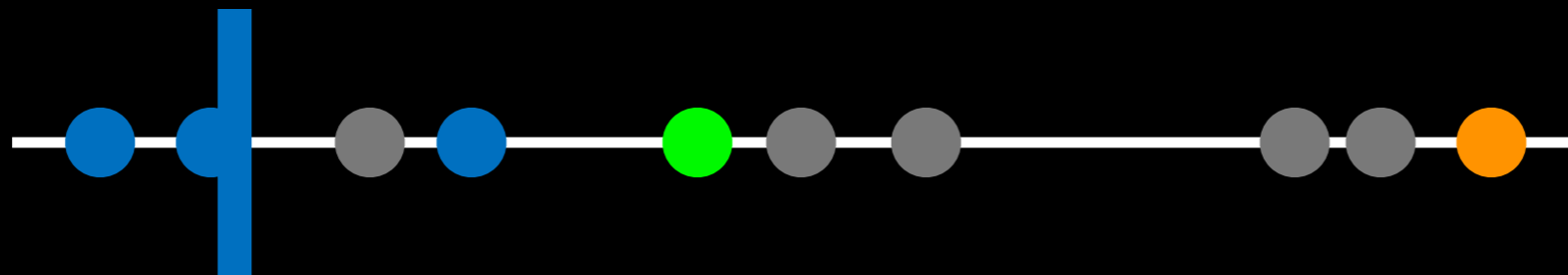

Pick new points as "initial" clusters

…Pick *k*=3 initial clusters and add the remaining points to the cluster with the nearest mean, recalculating the mean each time a new point is added…

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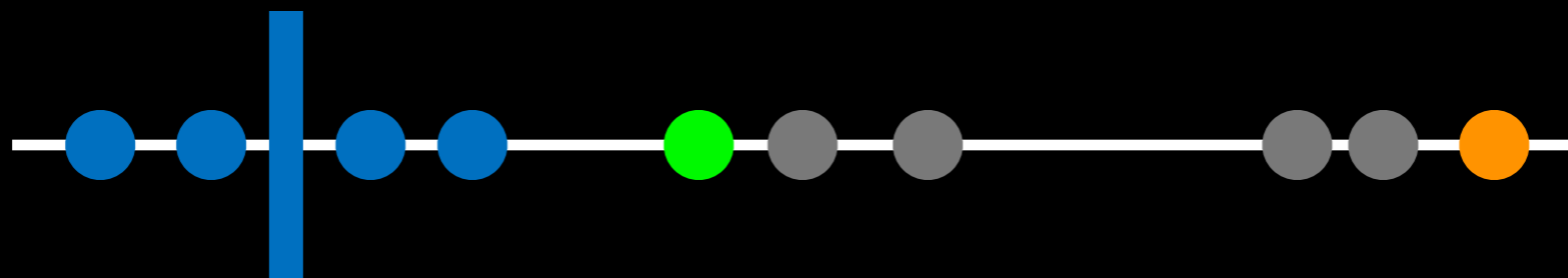
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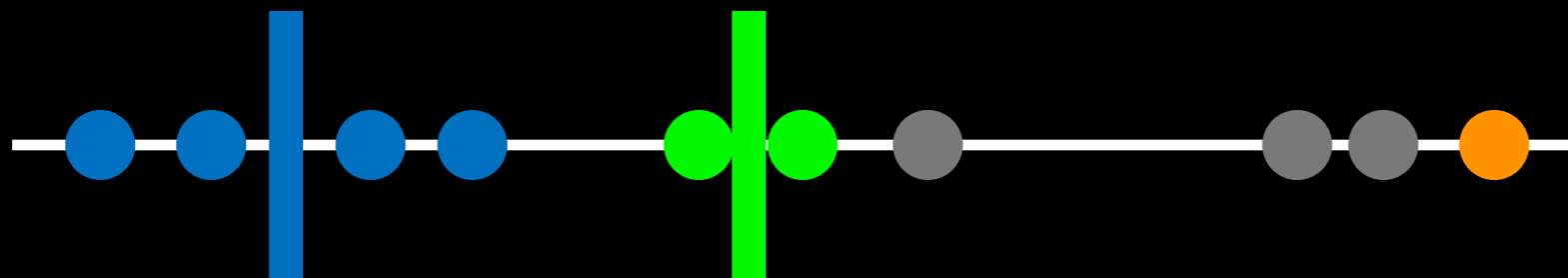
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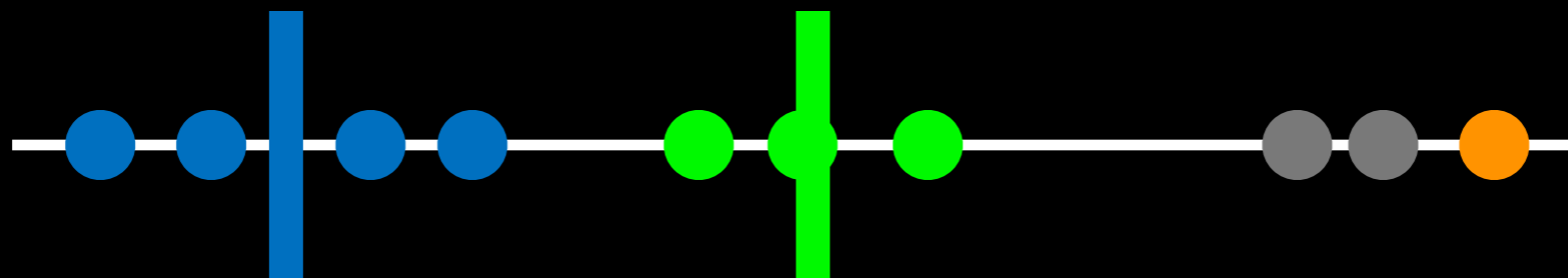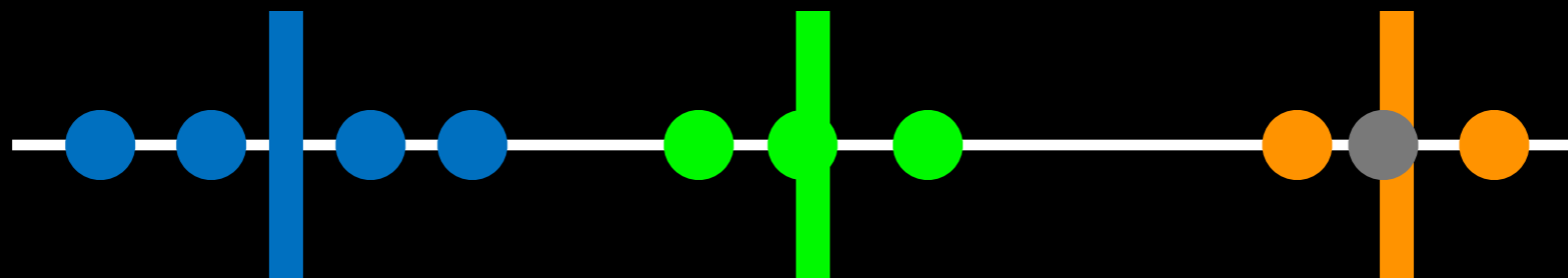
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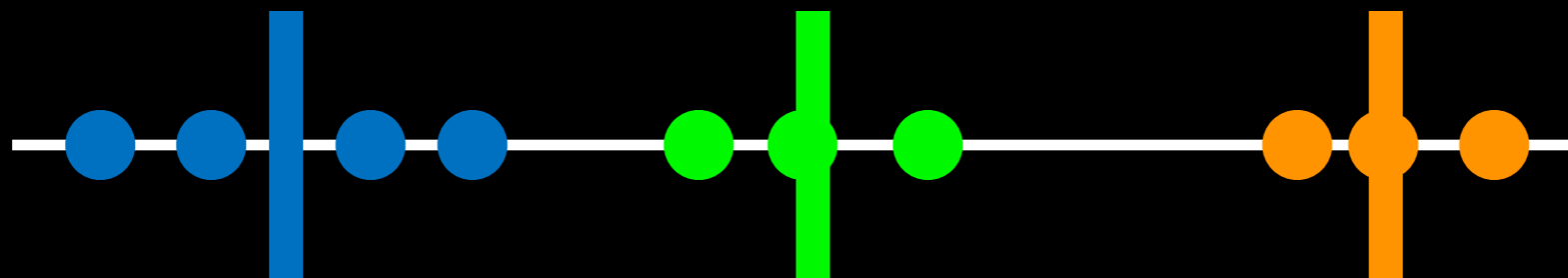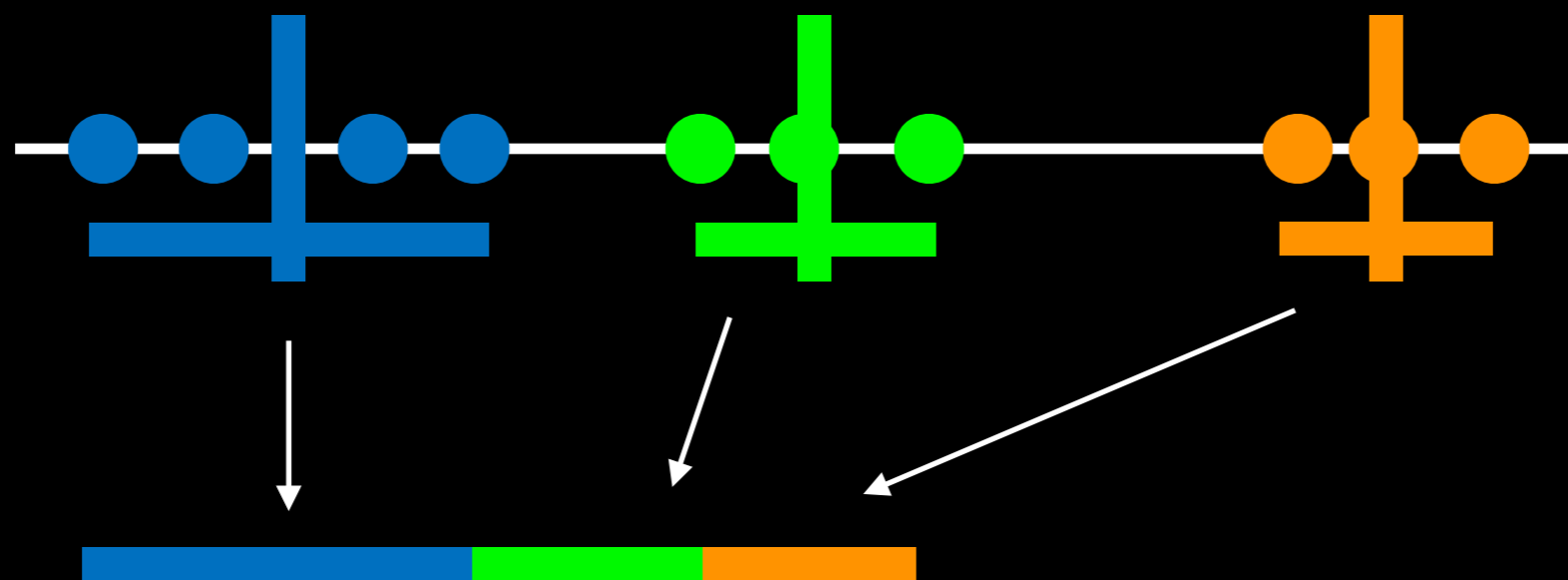
…Pick *k*=3 initial clusters and add the remaining points to the cluster with the nearest mean, recalculating the mean each time a new point is added…

Now the data are all assigned to clusters, we again sum the variation within each cluster

The total variation within clusters

Step 10.
Repeat again with different starting points

After several iterations k-means clustering knows it has the _best clustering so-far_ based on the smallest total variation with clusters.

However, it does not know if it has found the _best overall_. So it will perform several more iterations with different starting points…

Iteration 1

Iteration 2　　　　　　　**The winner!**

Iteration 3

Iteration 4
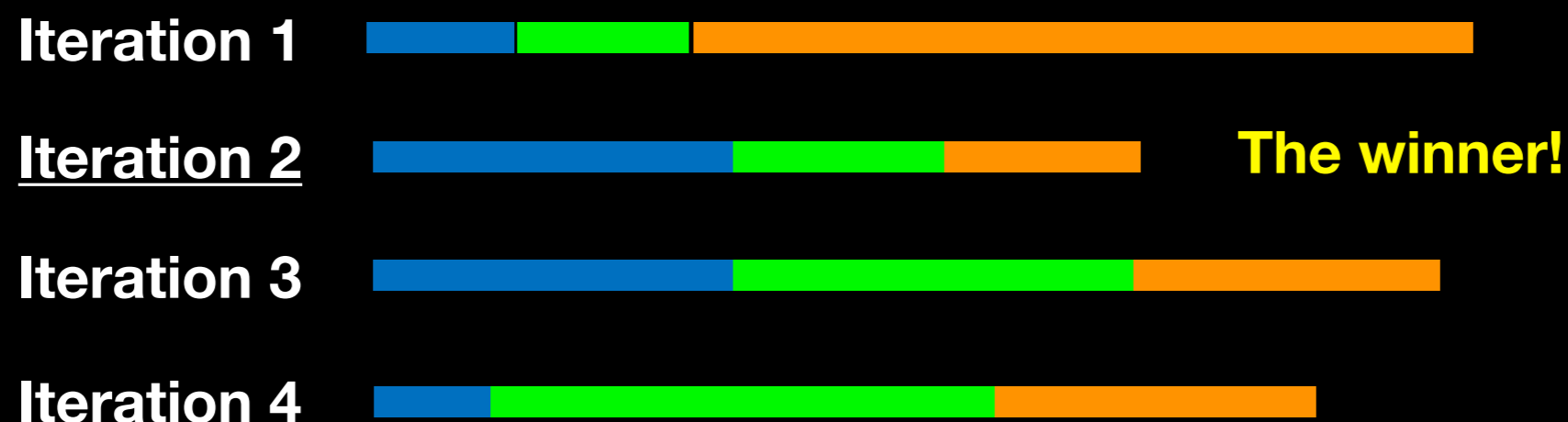
The total variation within clusters

What if we have more dimensions?
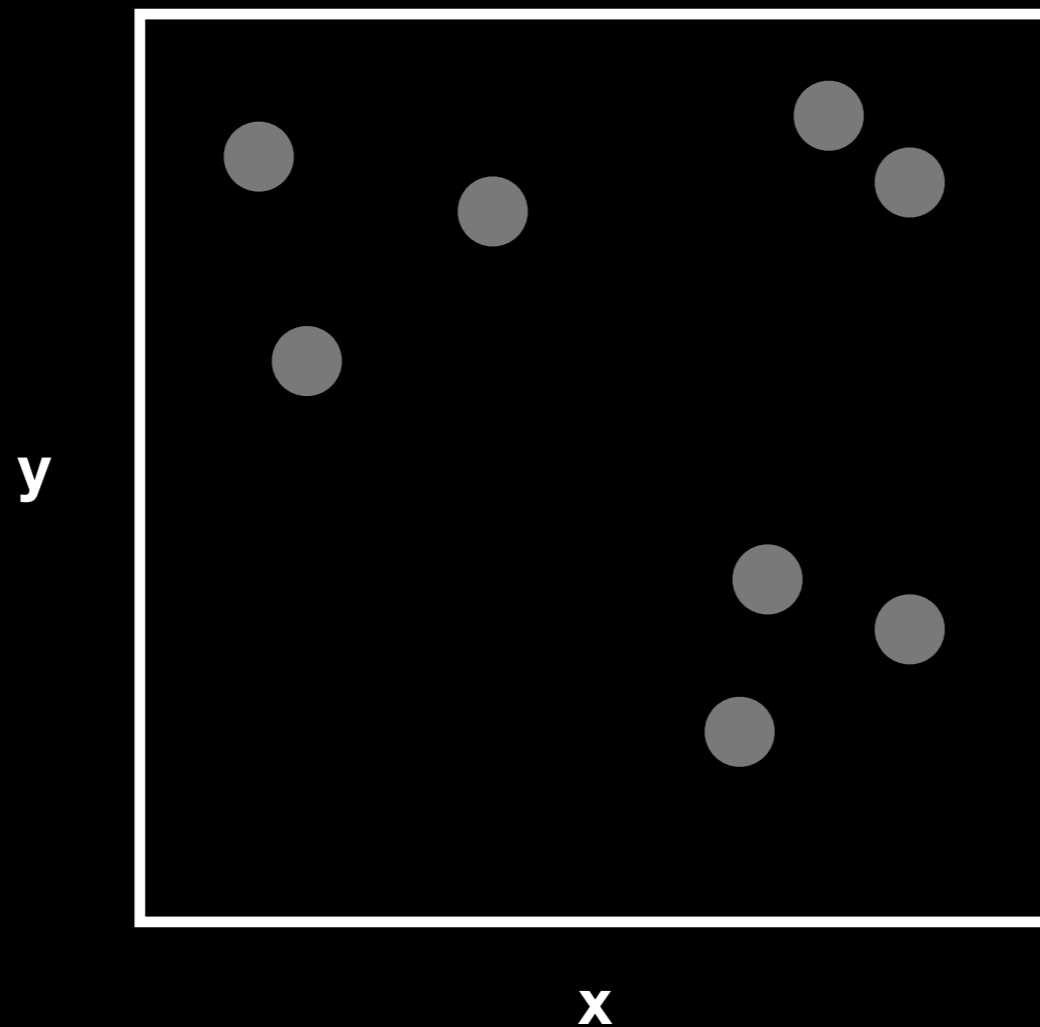
Just like before, we pick 3 random points…
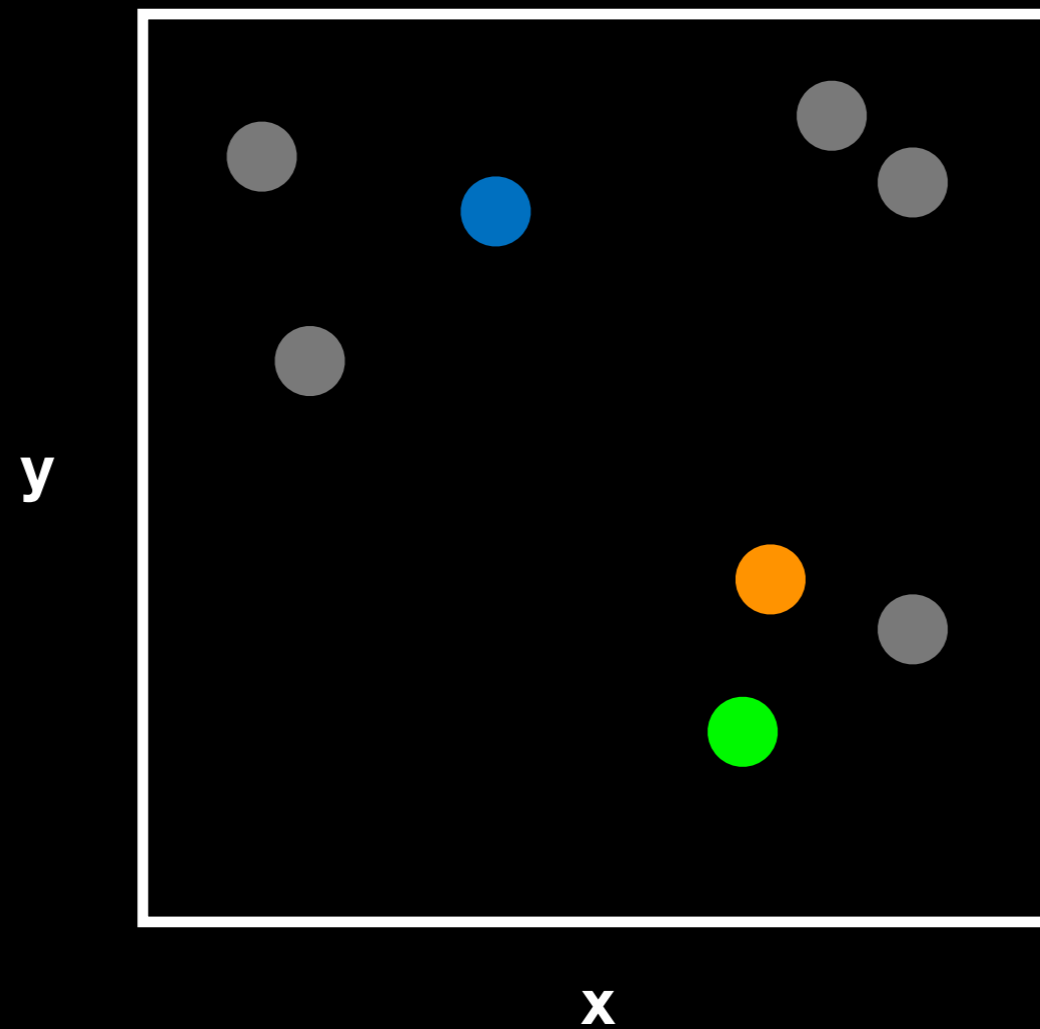
…and use the Euclidean distance.
In 2 dimensions the Euclidean distance is the same as the
Pythagorean theorem d = sqrt(x^2 + y^2)

…assign point to nearest cluster and update cluster center *
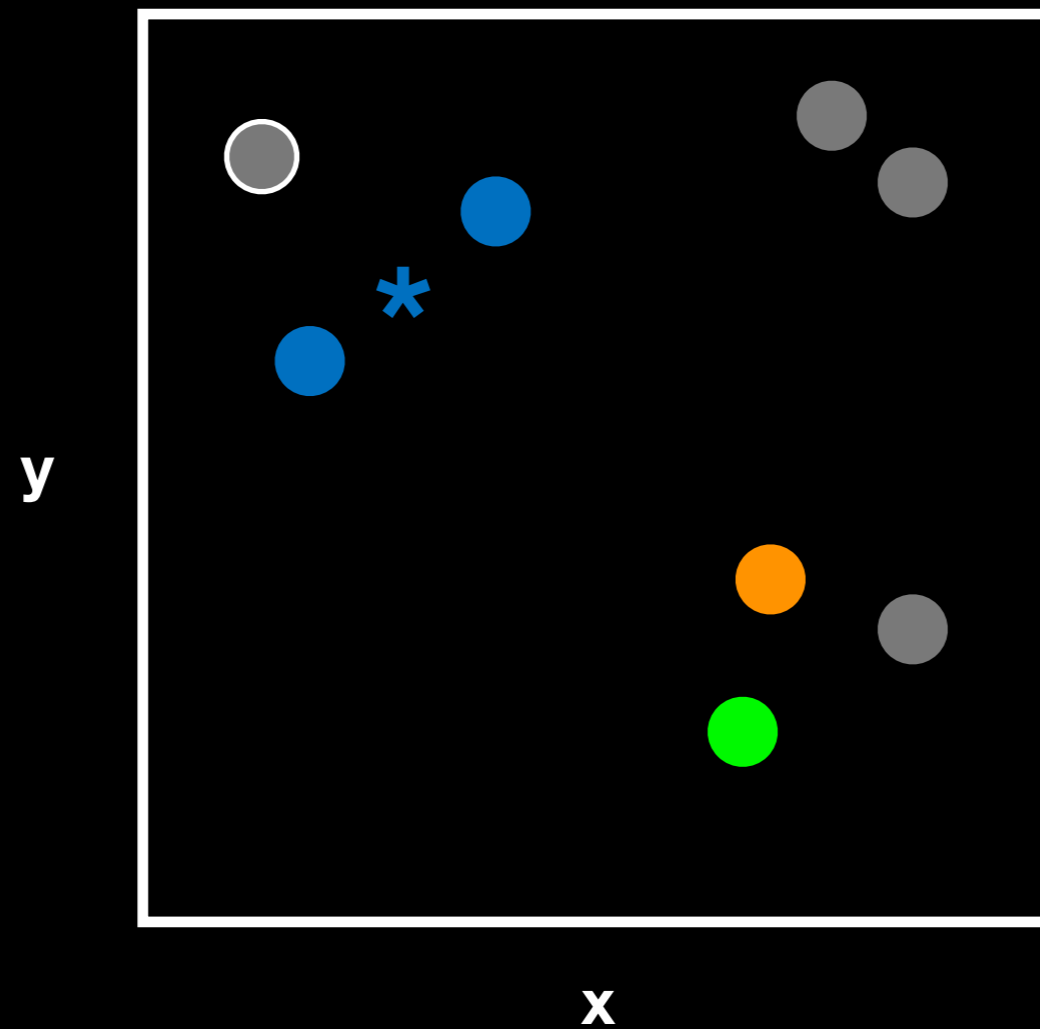
…and continue
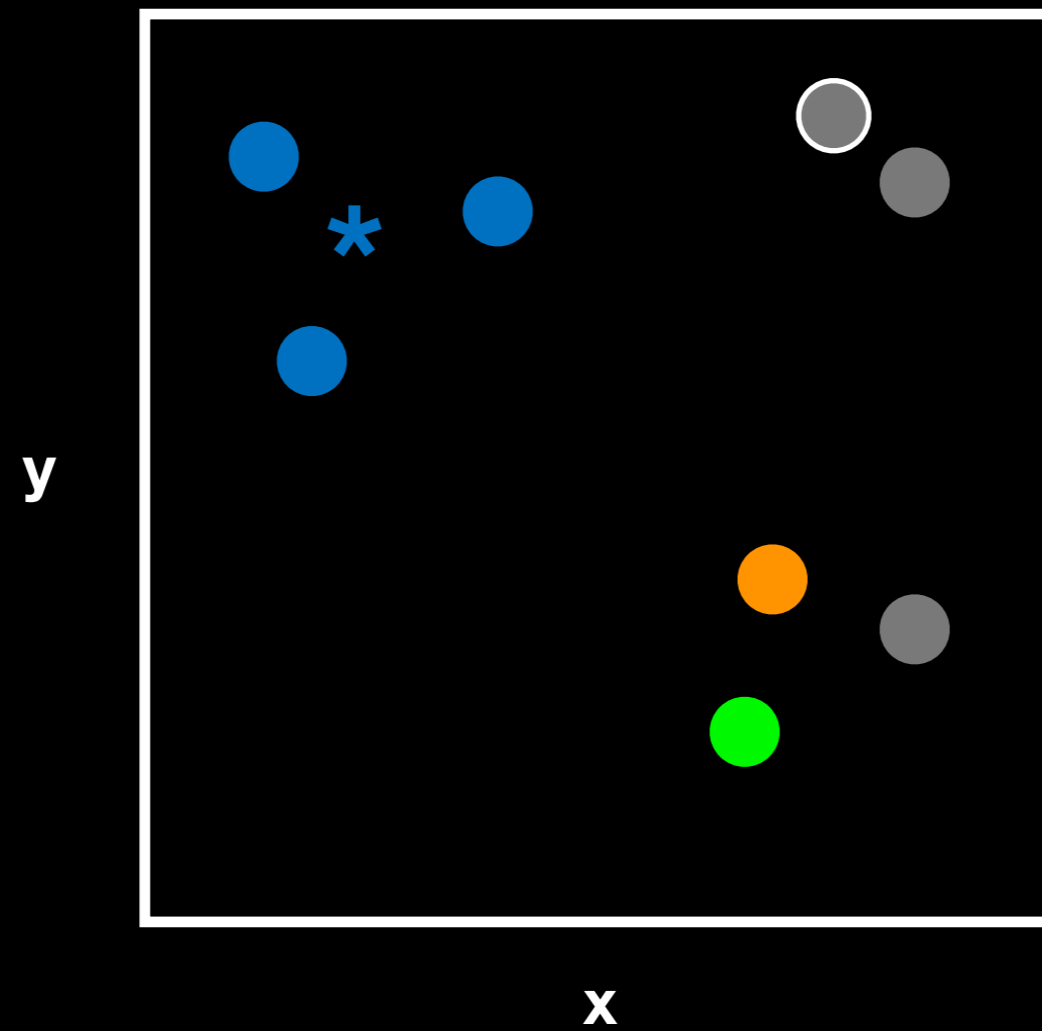
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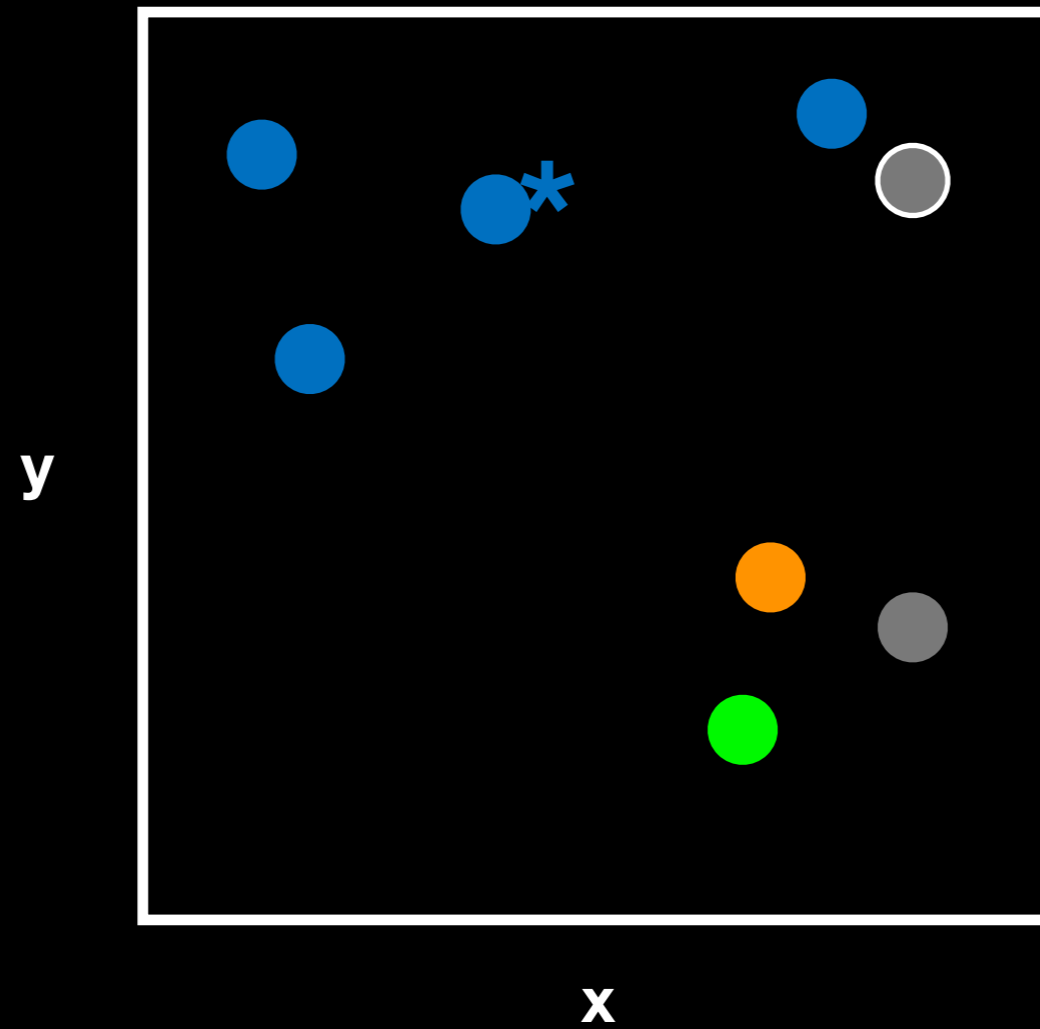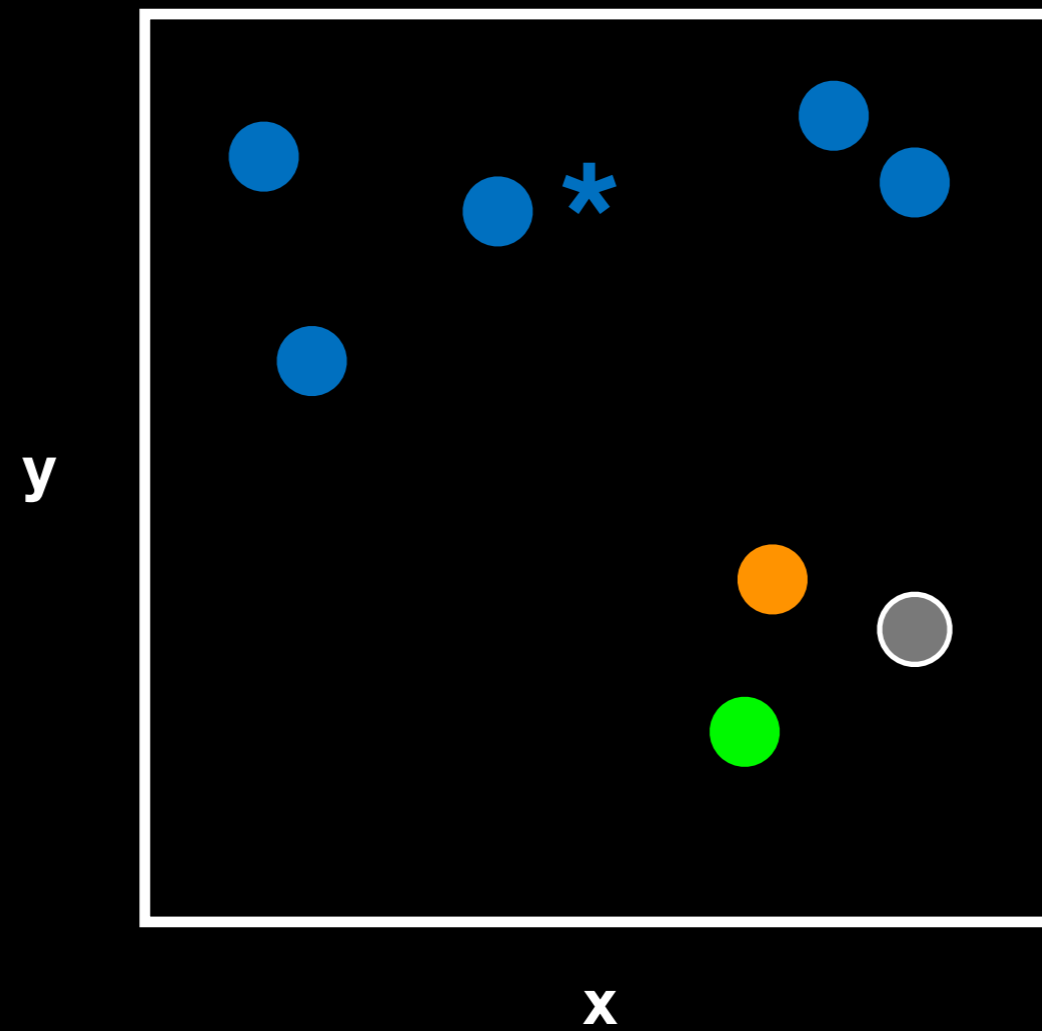
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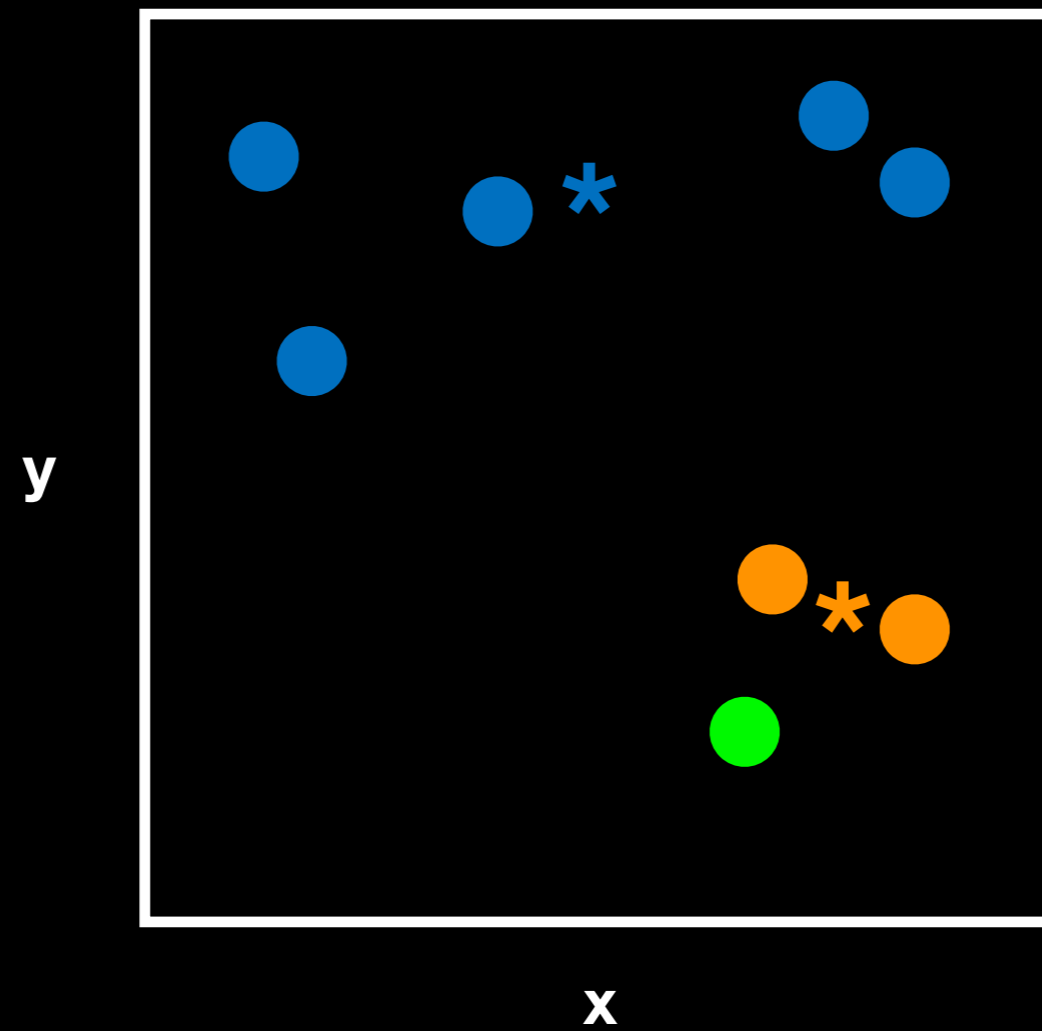
Again we have to use a number of different starting conditions before deciding on a good clustering!
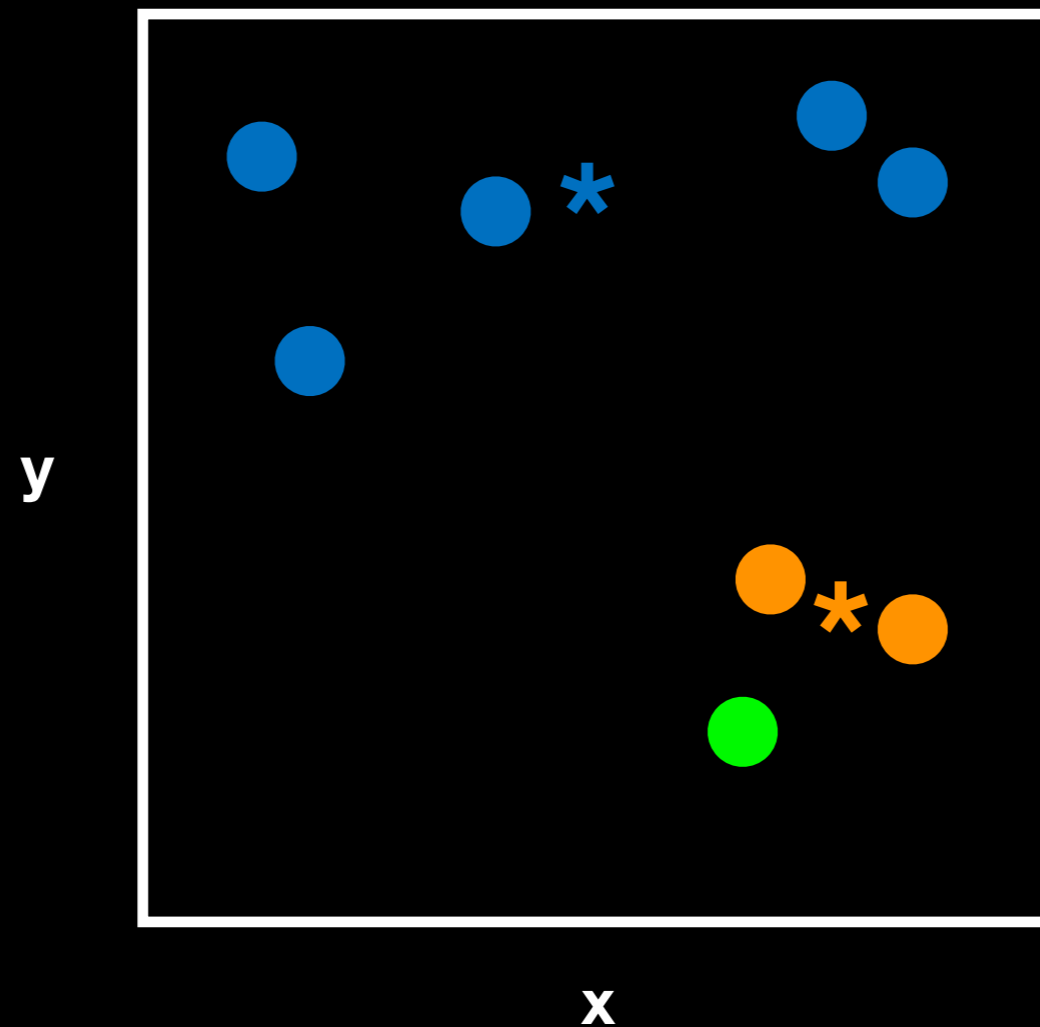
What if we have even more dimensions?

Cell Samples

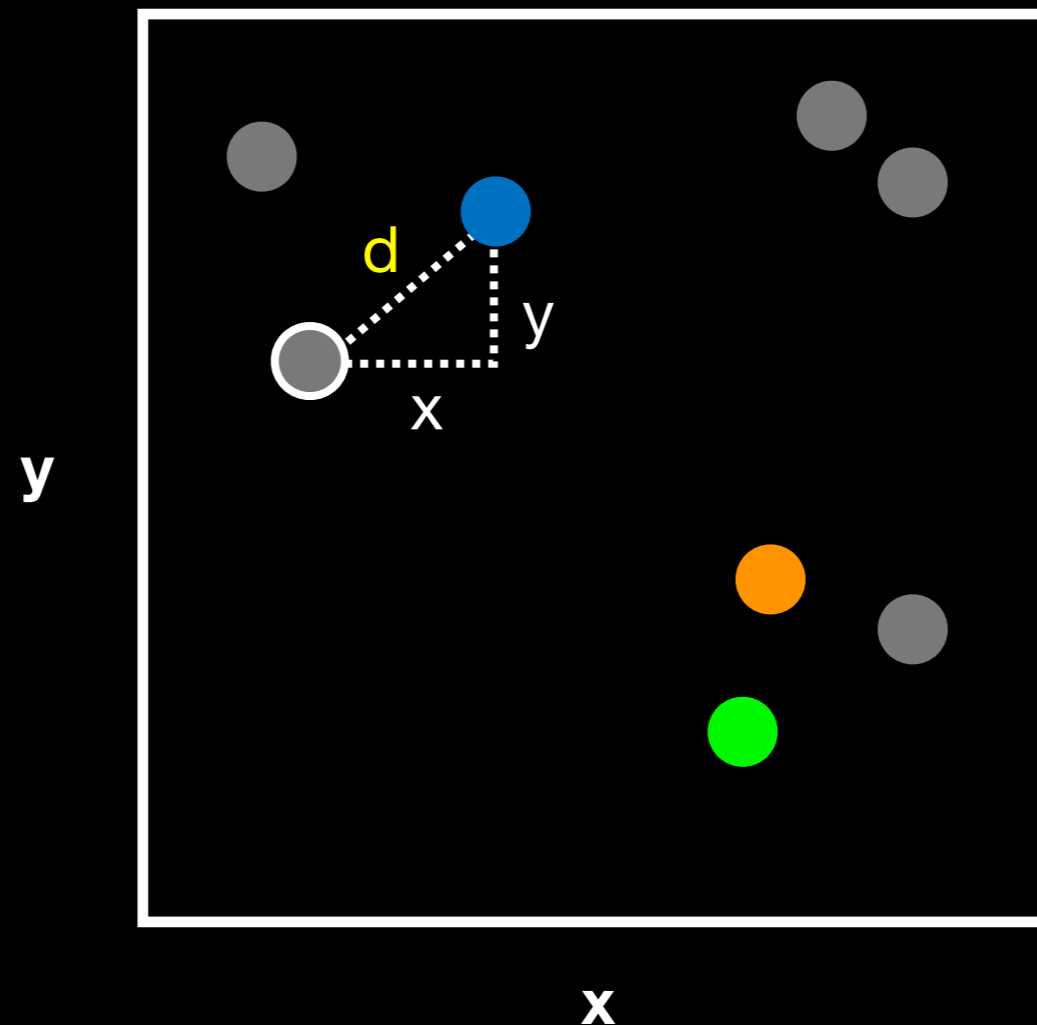|	#1	#2	#3
Gene 1	12	6	-13
Gene 2	-7	13	10
Gene 3	8	6	-9
Gene 4	9	5	-11
Gene 5	-3	1	6
Gene 6	10	4	-8

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	x	**y**	**z**

We could simply plot them by relabeling the cell samples as **x**, **y**, and **z** (i.e. a 3D plot)

What if we have even more dimensions?

Cell Samples

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x   **y**   **z**

…and go through exactly the same procedure with initial cluster assignment followed by distance calculation etc…

$$d = sqrt(x^2 + y^2 + z^2)$$

Cell Samples

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Of course we don't actually need to plot anything.

We can just calculate the Euclidean distance along any number of dimensions and perform our k-means clustering in the same way.

k-means in R

```
# k-means algorithm with 3 centers, run 20 times
kmeans(x, centers= 3, nstart= 20)
```

- Input **x** is a numeric matrix, or data.frame, with one observation per row, one feature per column

- k-means has a random component

- Run algorithm multiple times to improve odds of the best model
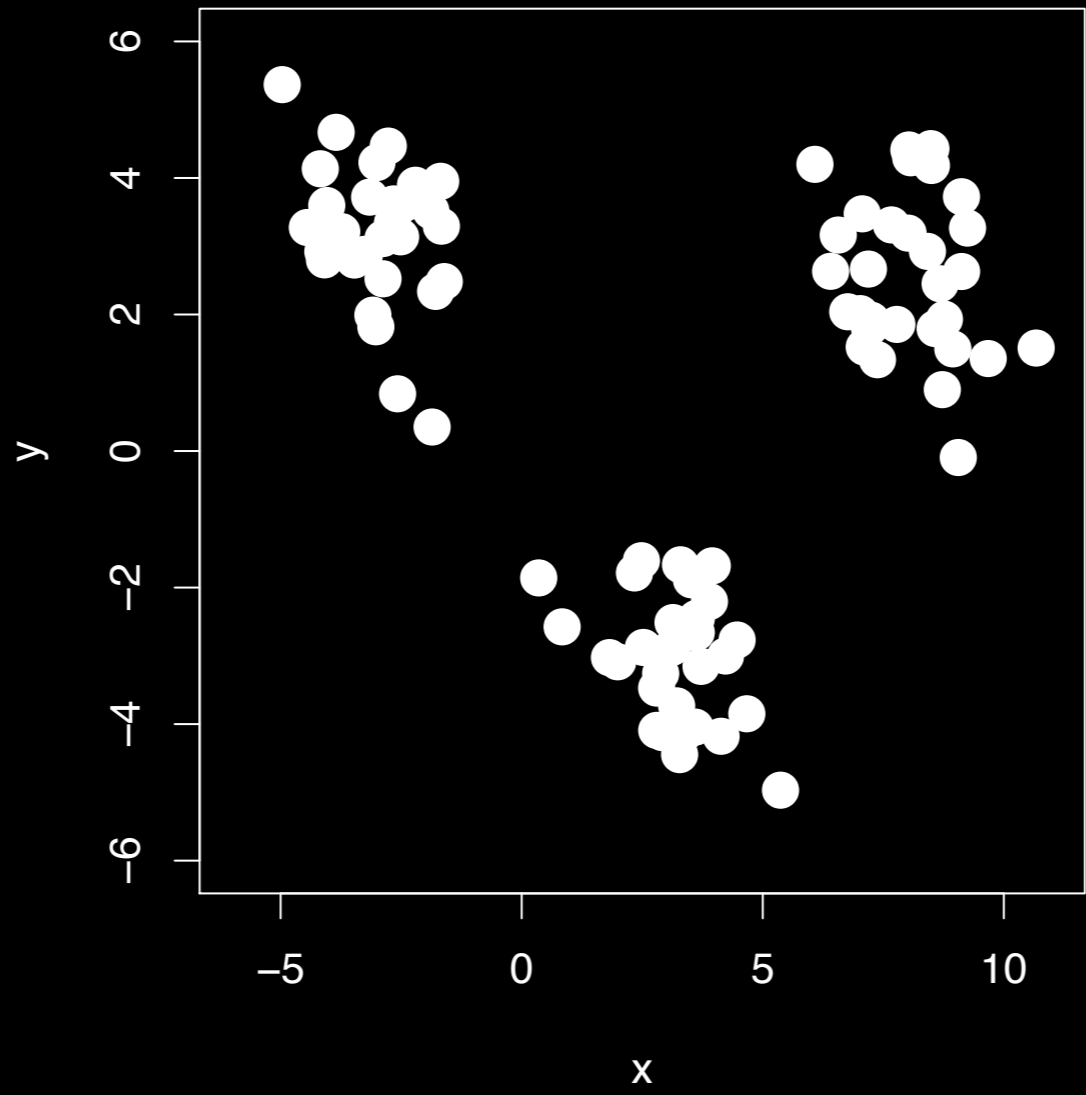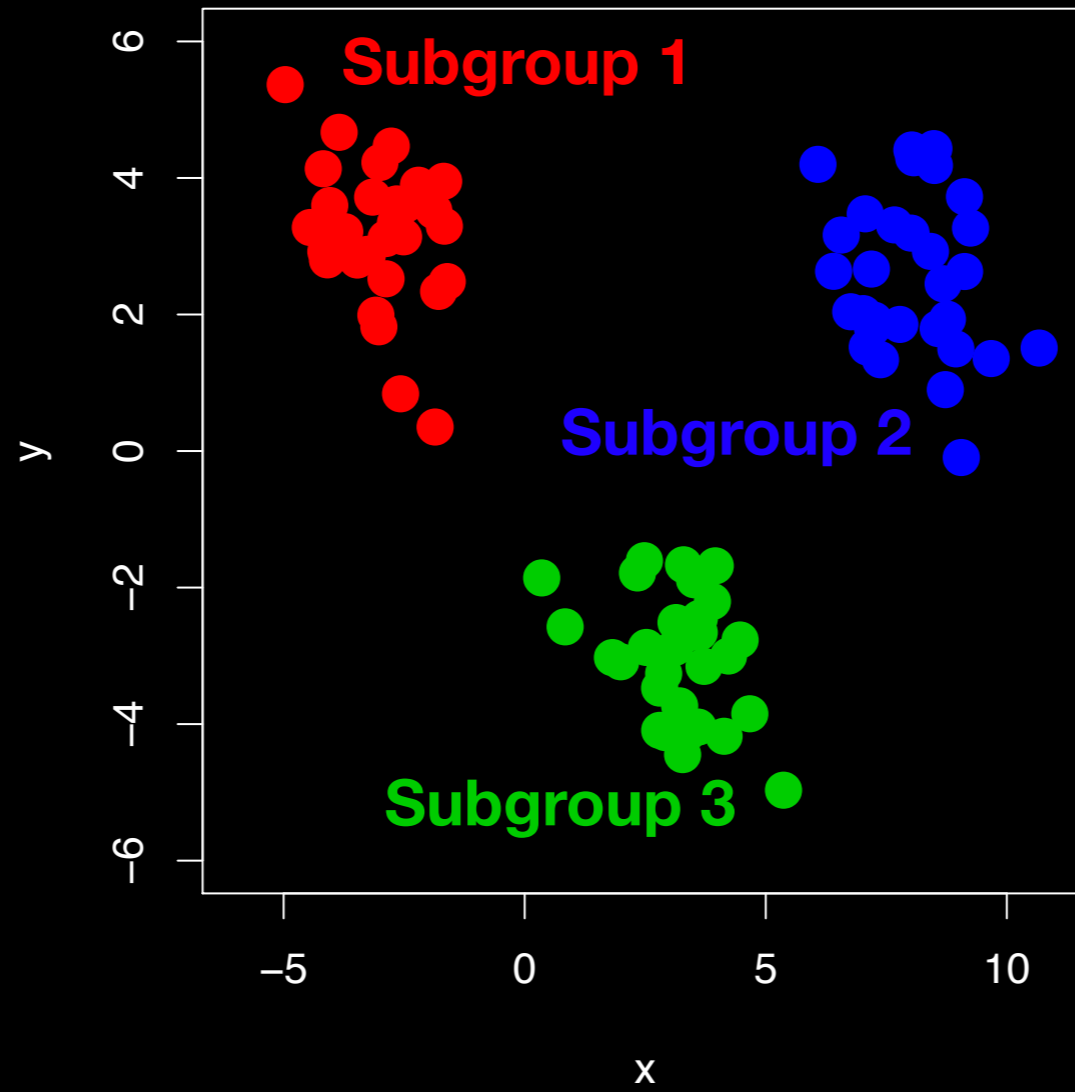
Model selection

- Recall k-means has a random component

- Best outcome is based on total within cluster sum of squares:
  - ➡ For each cluster
    - For each observation in the cluster
      - Determine squared distance from observation to cluster center
  - ➡ Sum all of them together

Model selection

```
# k-means algorithm with 5 centers, run 20 times
kmeans(x, centers=5, nstart=20)
```

- Running algorithm multiple times (i.e. setting nstart) helps find the global minimum total within cluster sum of squares

- Increasing the default value of **nstart** is often sensible

Total Within SS = 615.340155087286   Total Within SS = 533.574674546762
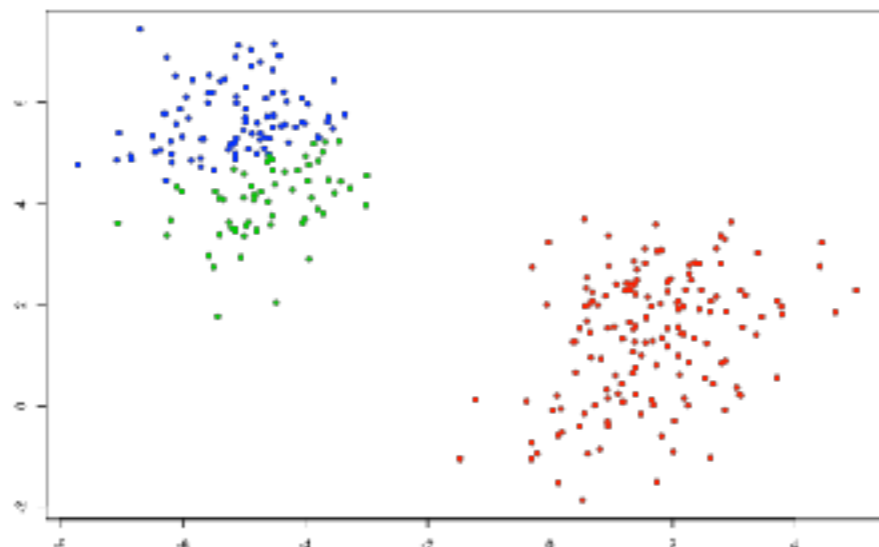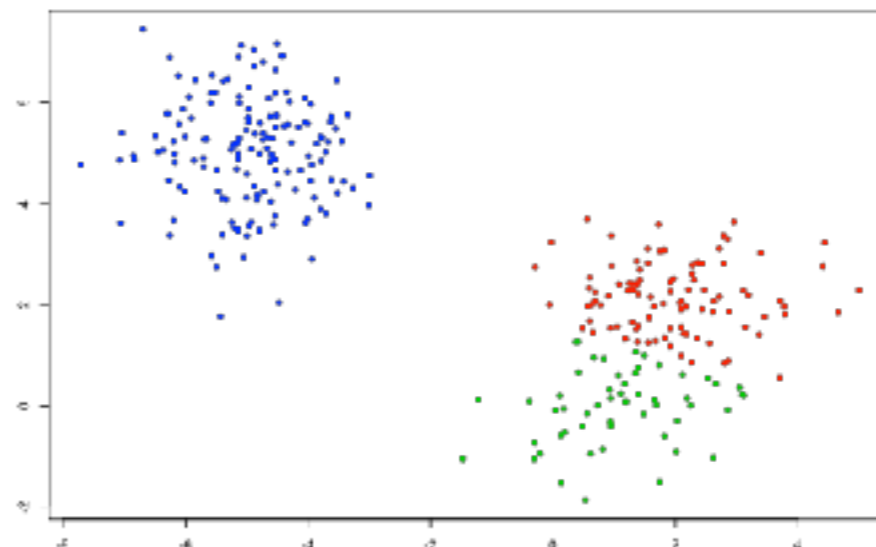
Total Within SS = 615.340155087286   Total Within SS = 533.630315419879

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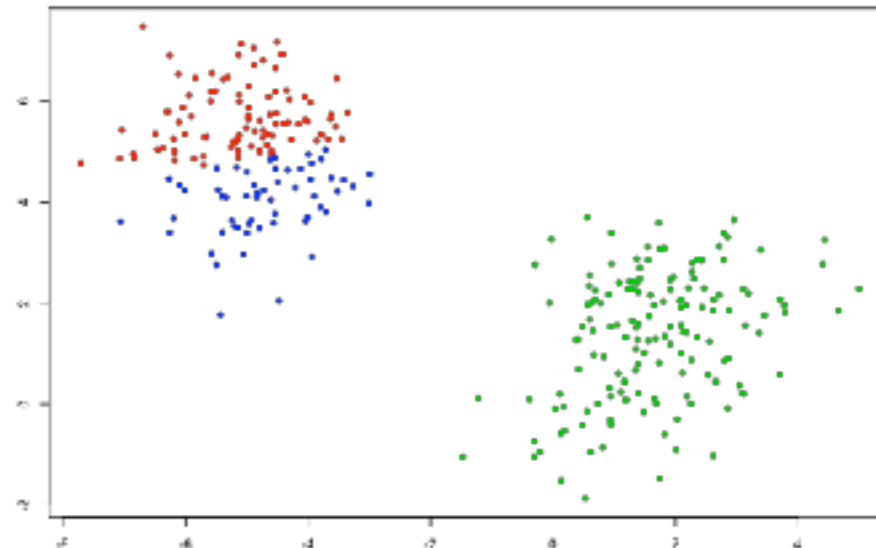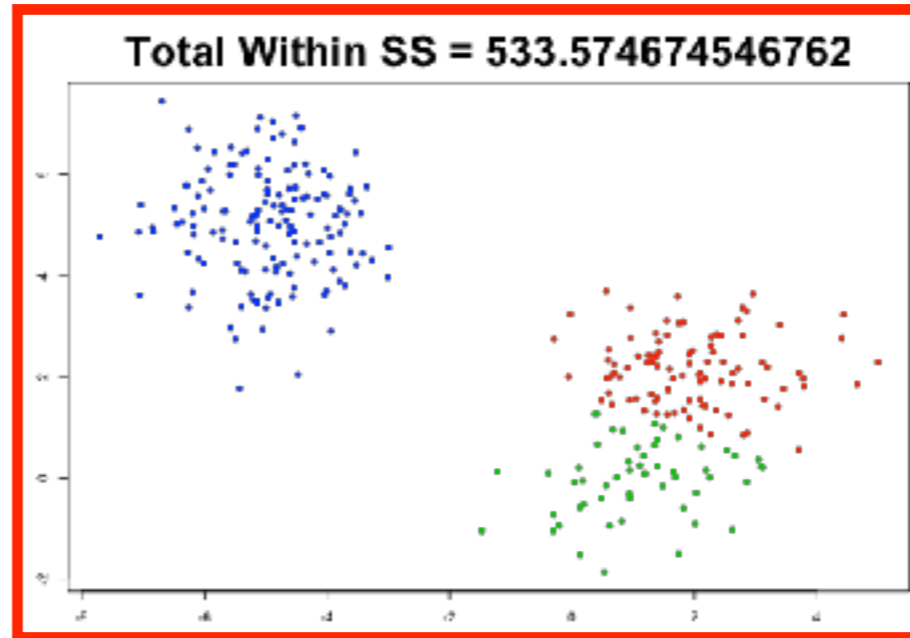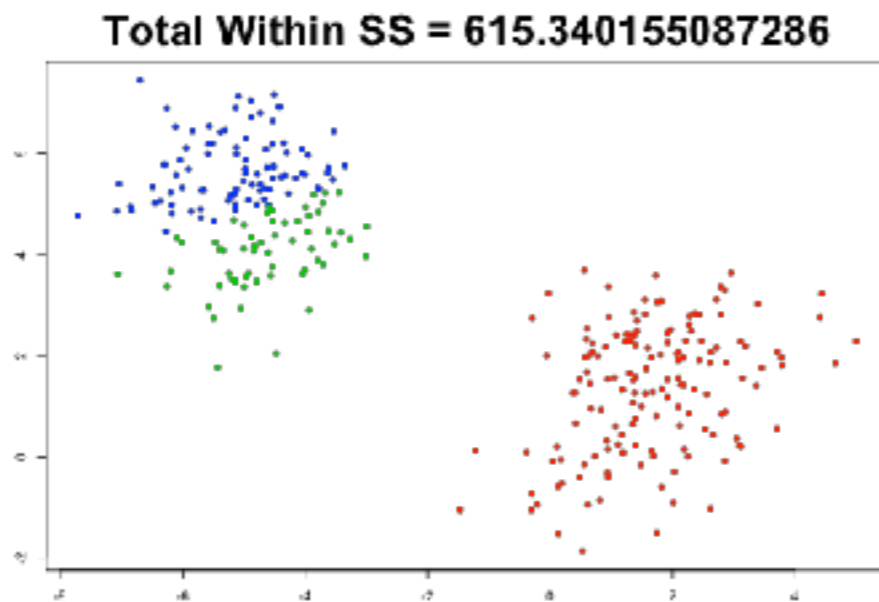
Winner has the smallest within cluster SS

Total Within SS = 615.340155087286

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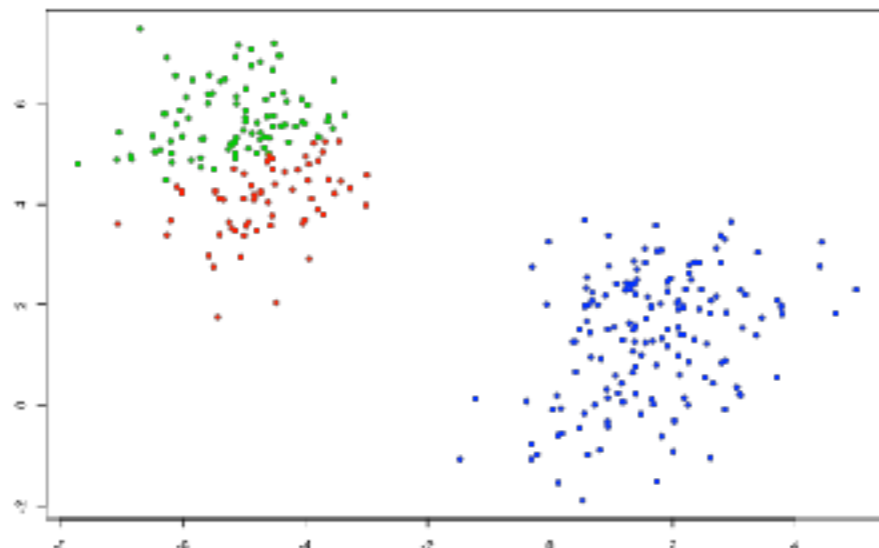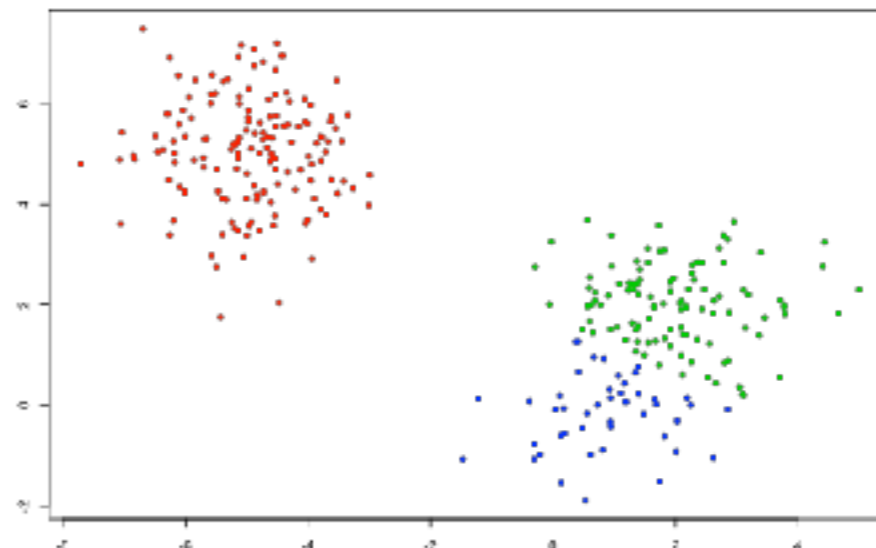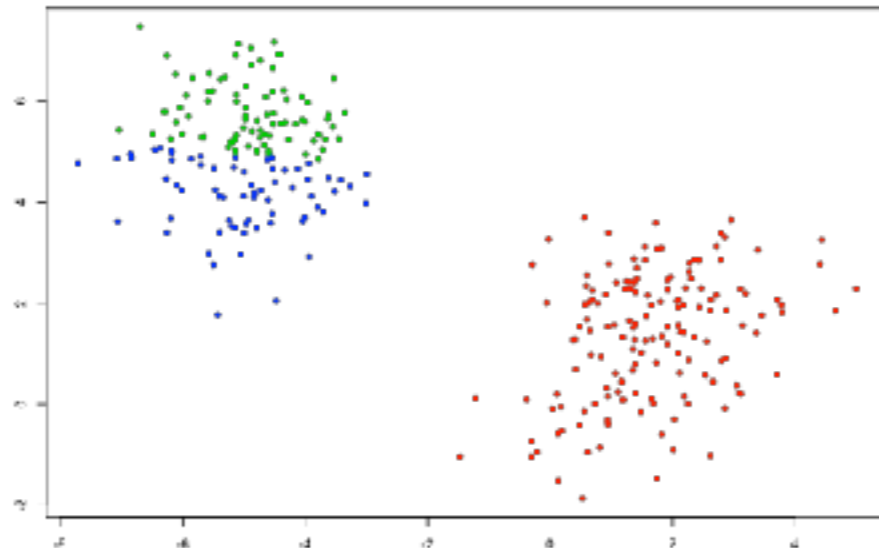
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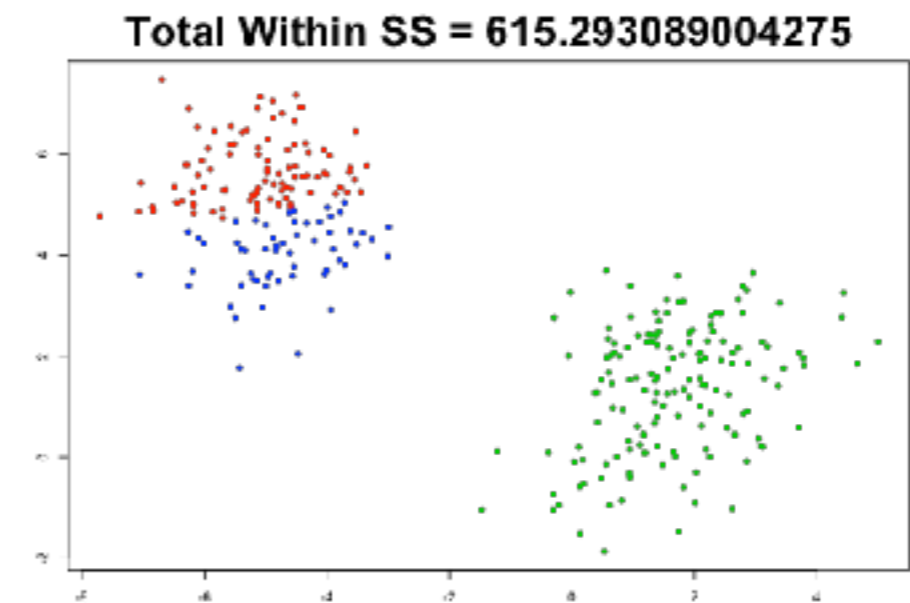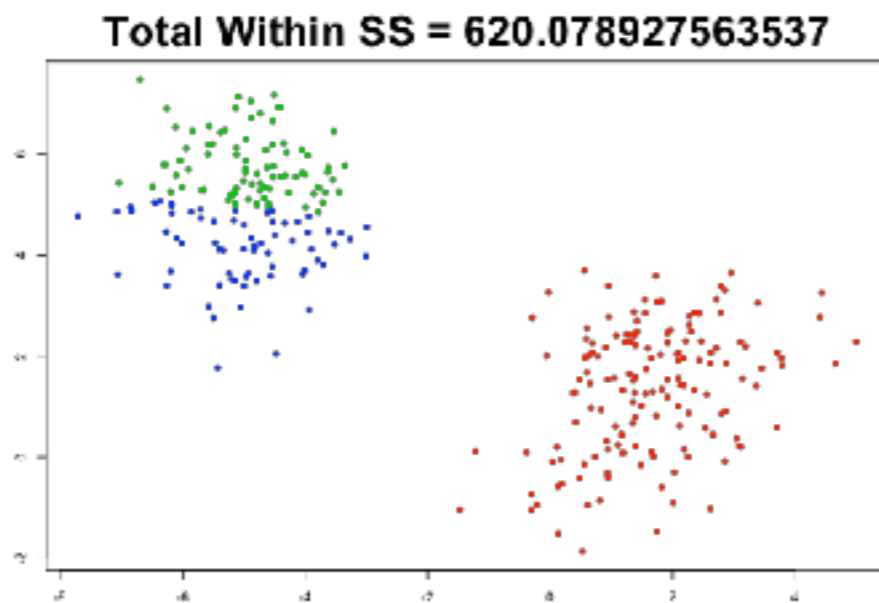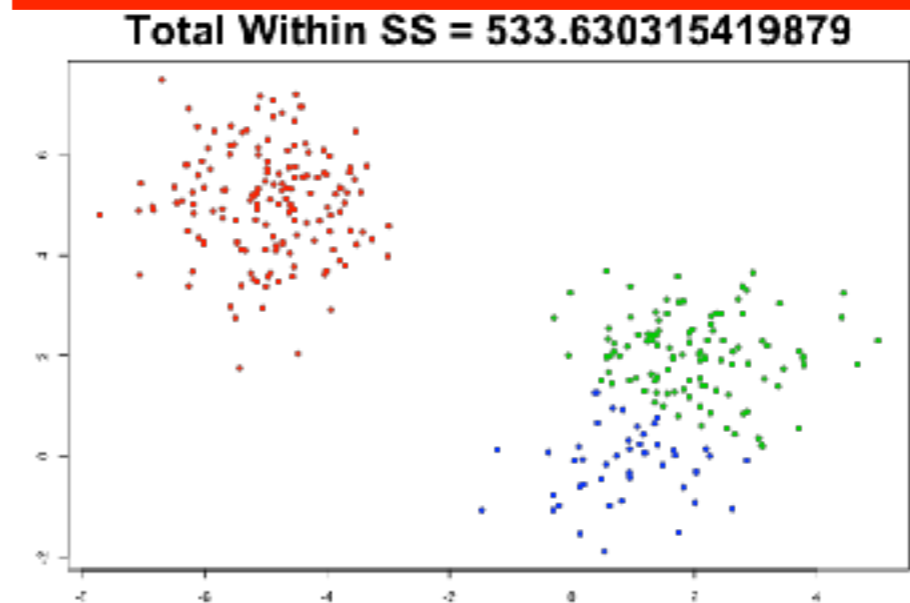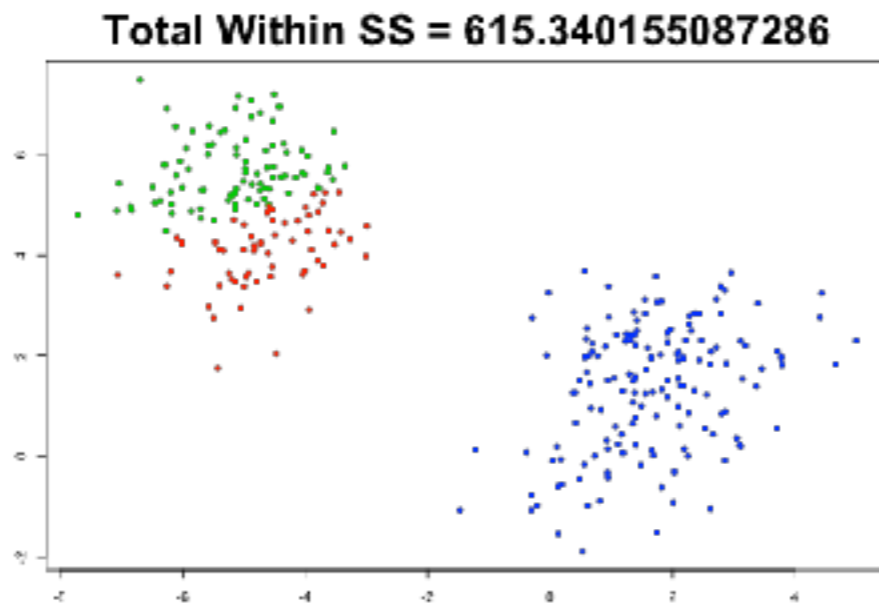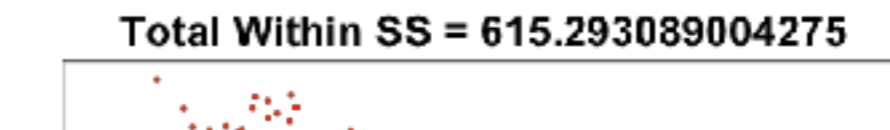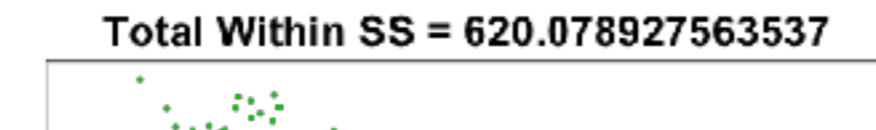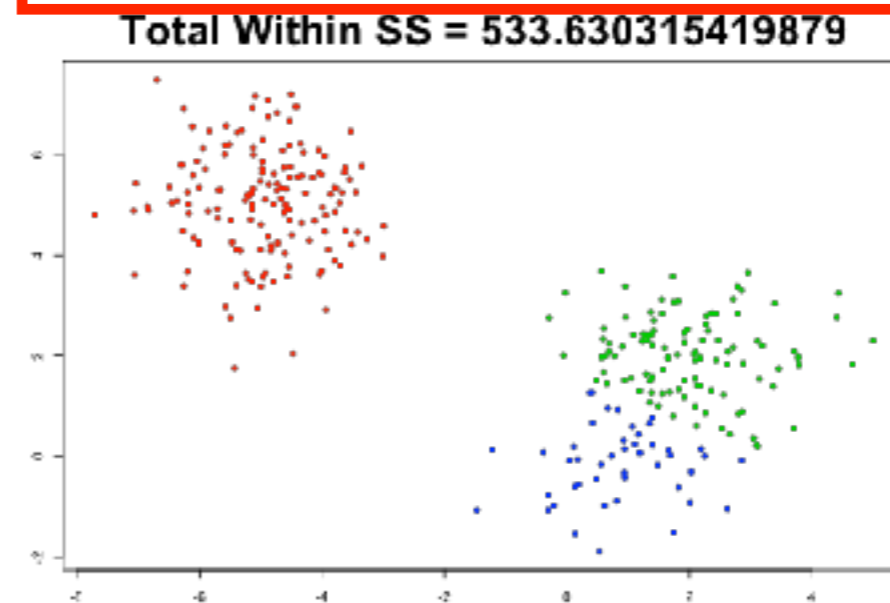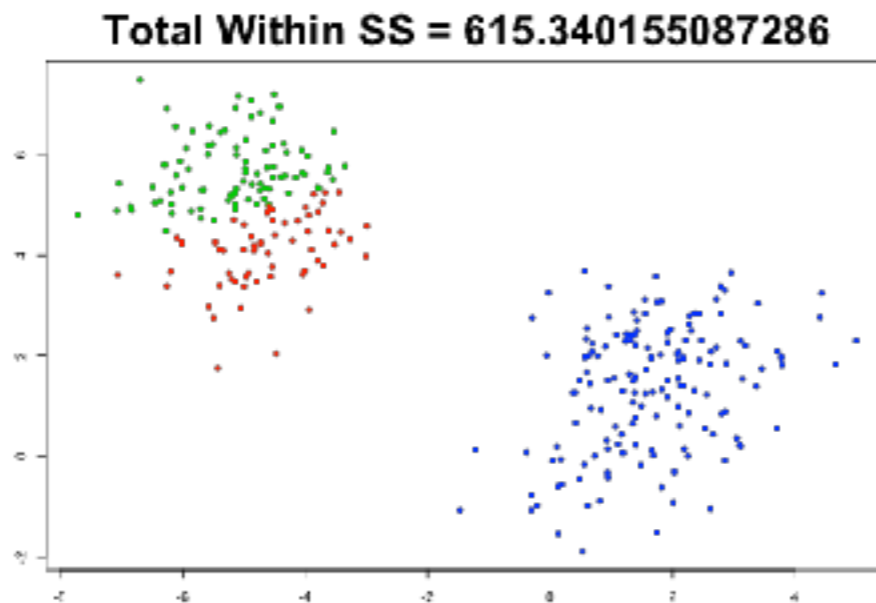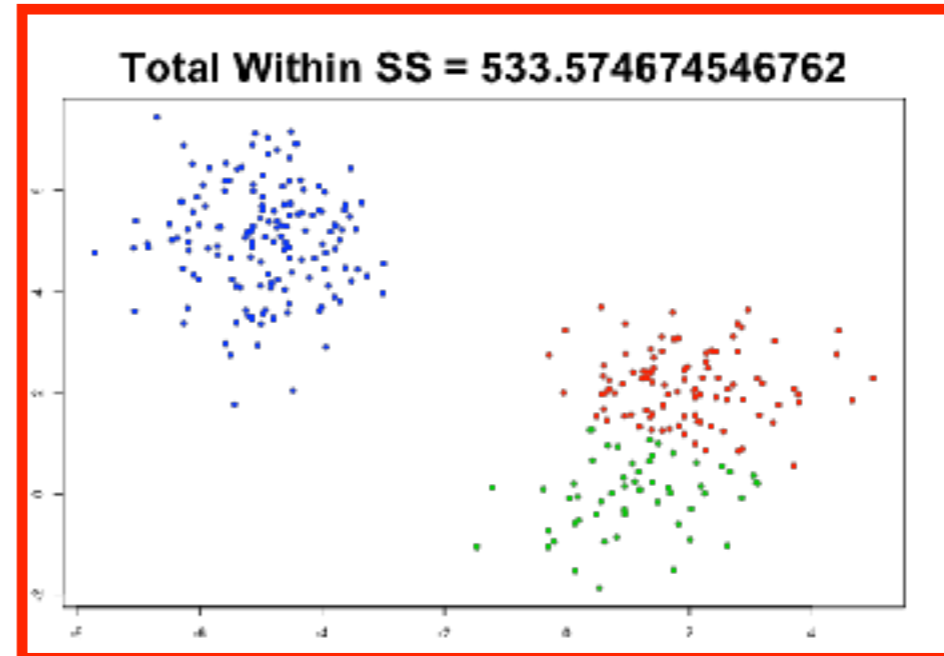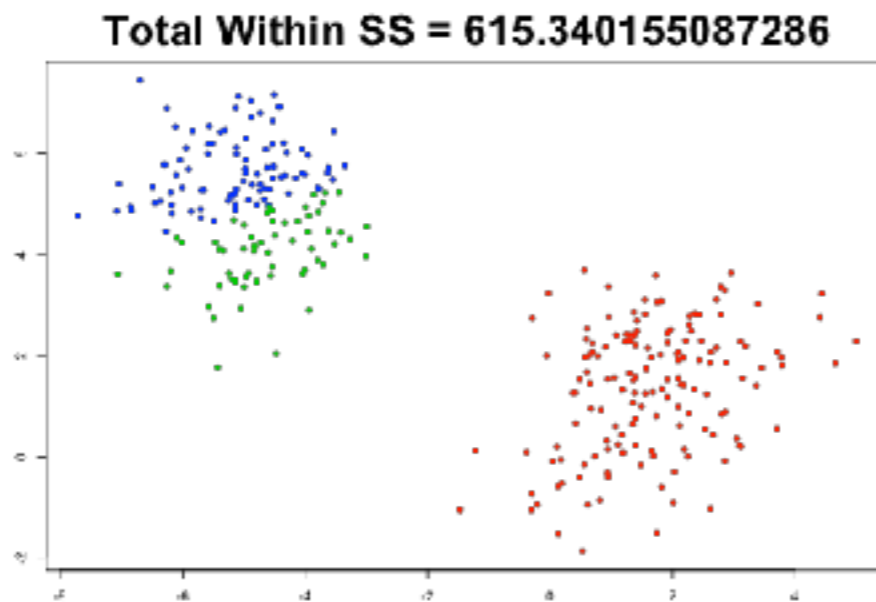
Note. k-means will always give you the renumber of clusters you request! (Here for example, k=2 may be better but we asked for k=3)
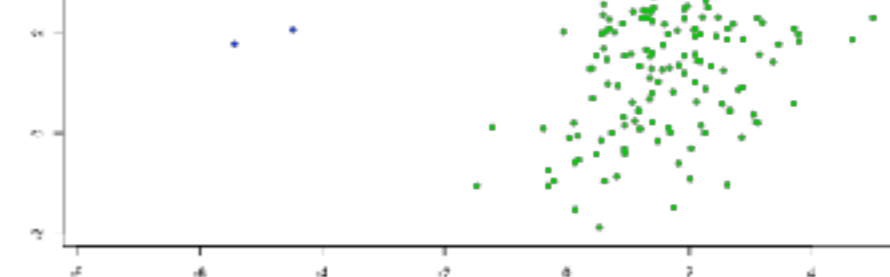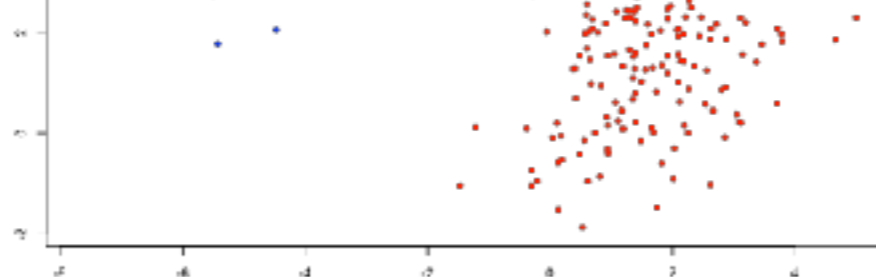
Determining number of clusters

Scree plot

"Elbow"

Total within SS

Number of clusters (k)
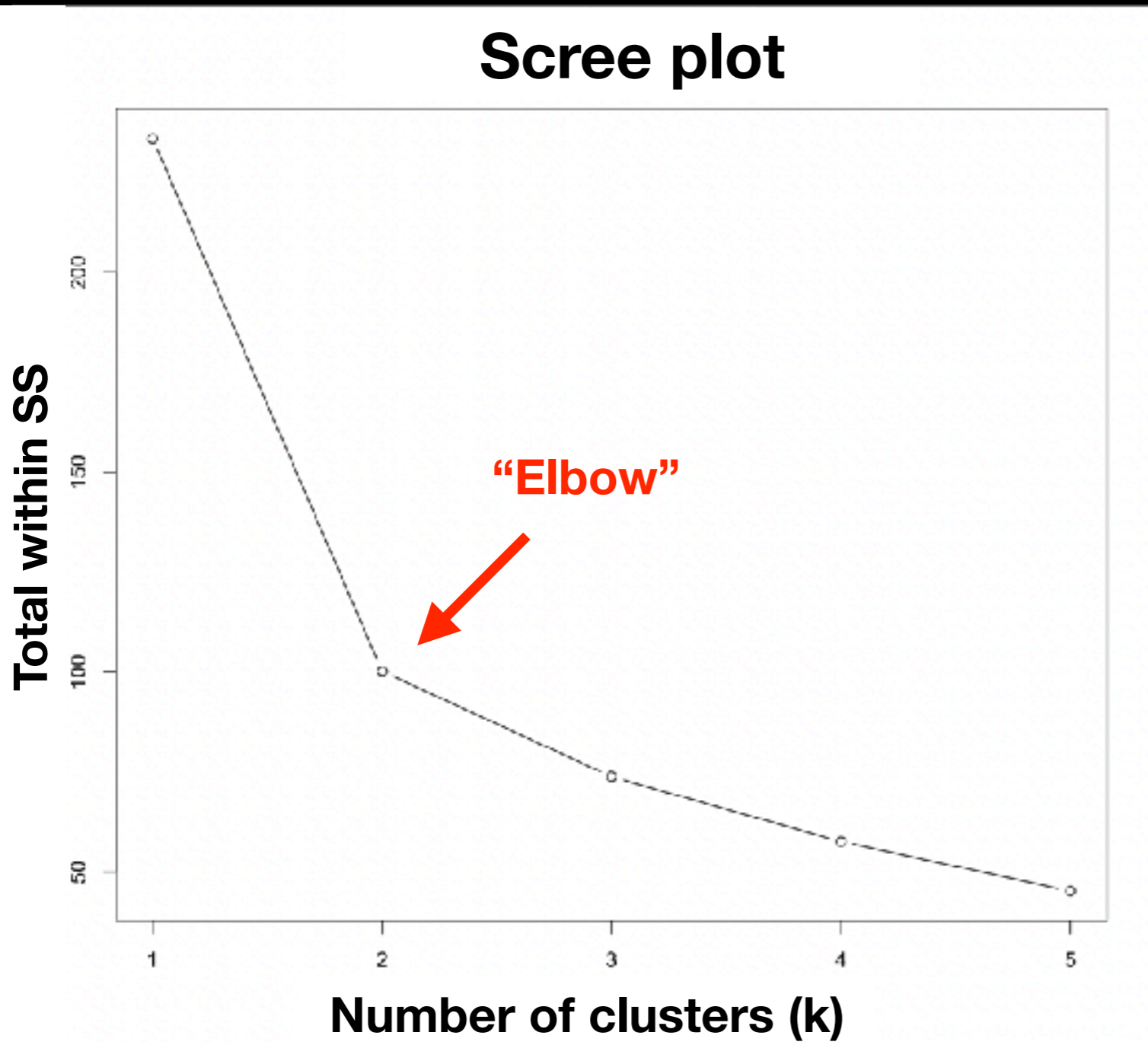
Trial and error is not the best approach

Systematically try a range of different k values and plot a "scree plot".

Here there is a large reduction in SS with **k=2** but after that the values do not go down as quickly!

Your Turn!

```r
# Generate some example data for clustering
tmp <- c(rnorm(30,-3), rnorm(30,3))
x <- cbind(x=tmp, y=rev(tmp))

plot(x)

# Use the kmeans() function setting k to 2 and nstart=20

# Inspect/print the results

# Q. How many points are in each cluster?
# Q. What 'component' of your result object details
#       - cluster size?
#       - cluster assignment/membership?
#       - cluster center?


# Plot x colored by the kmeans cluster assignment and
#       add cluster centers as blue points
# Q. Repeat for k=3, which one has the better total SS?
```

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- Hands-on application to cell classification

Hierarchical clustering

- Number of clusters is not known ahead of time

- Two kinds of hierarchical clustering:
  - ➡ <u>bottom-up</u>
  - ➡ top-down

Hierarchical clustering

Simple example:

5 clusters: Each point starts as it's own "cluster"!
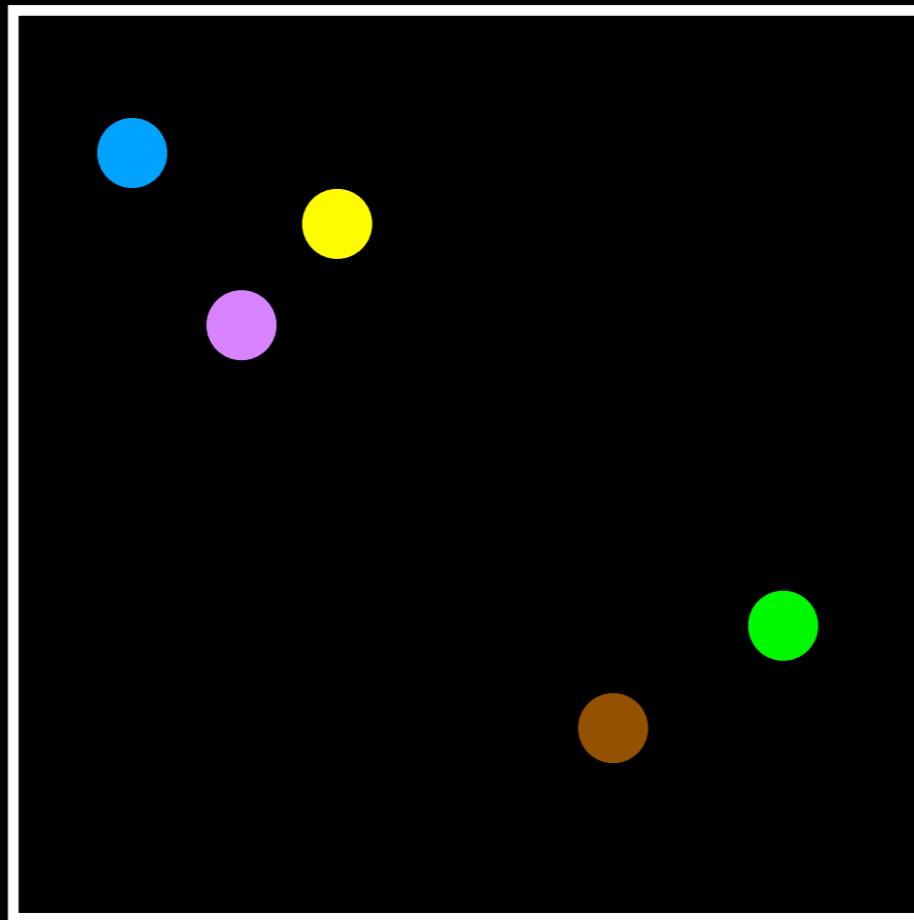
Hierarchical clustering

4 clusters
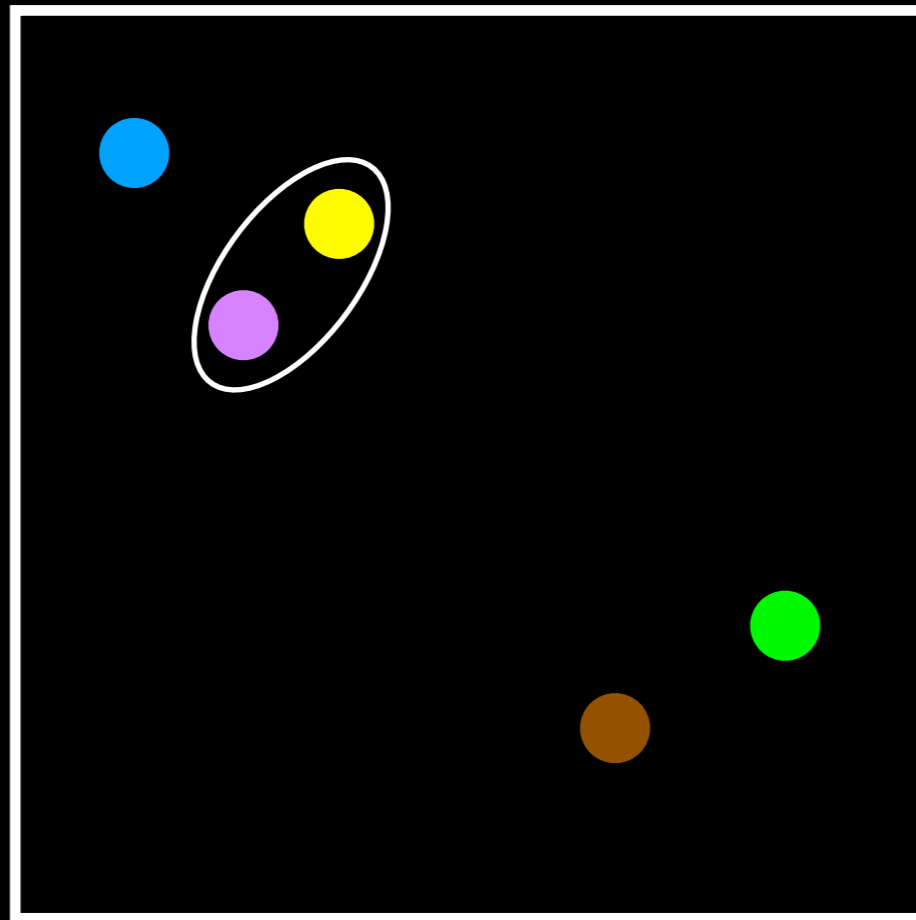
Hierarchical clustering

3 clusters
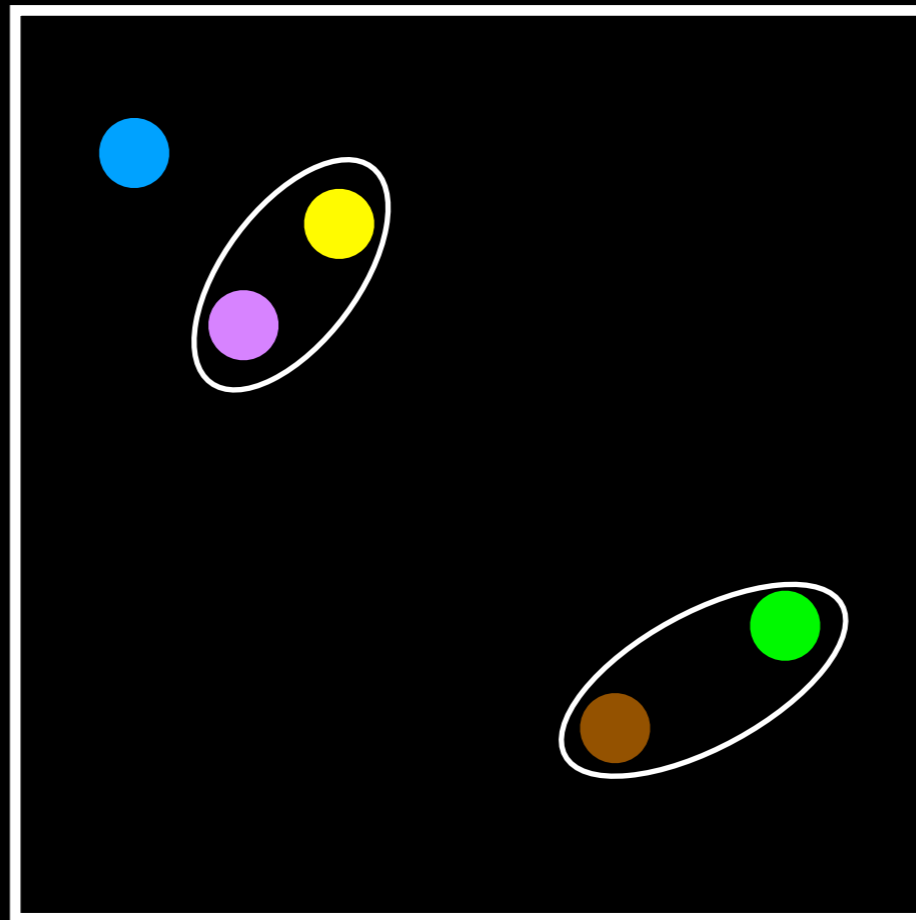
Hierarchical clustering

2 clusters
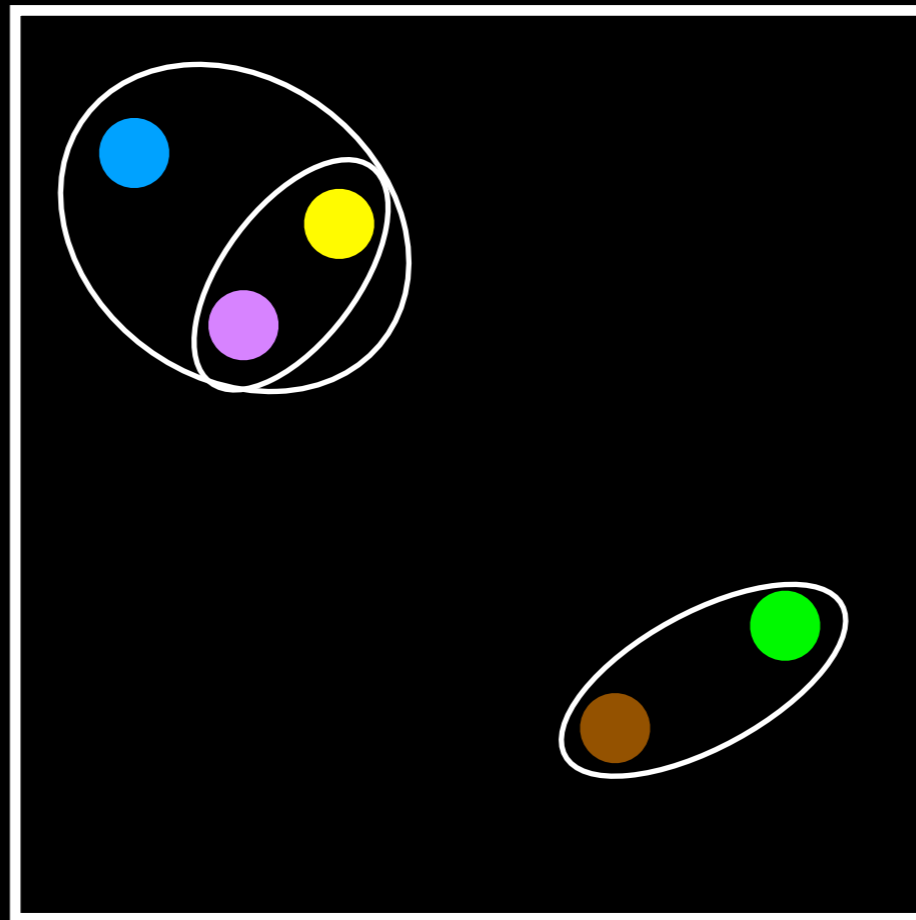
Hierarchical clustering

End: 1 cluster
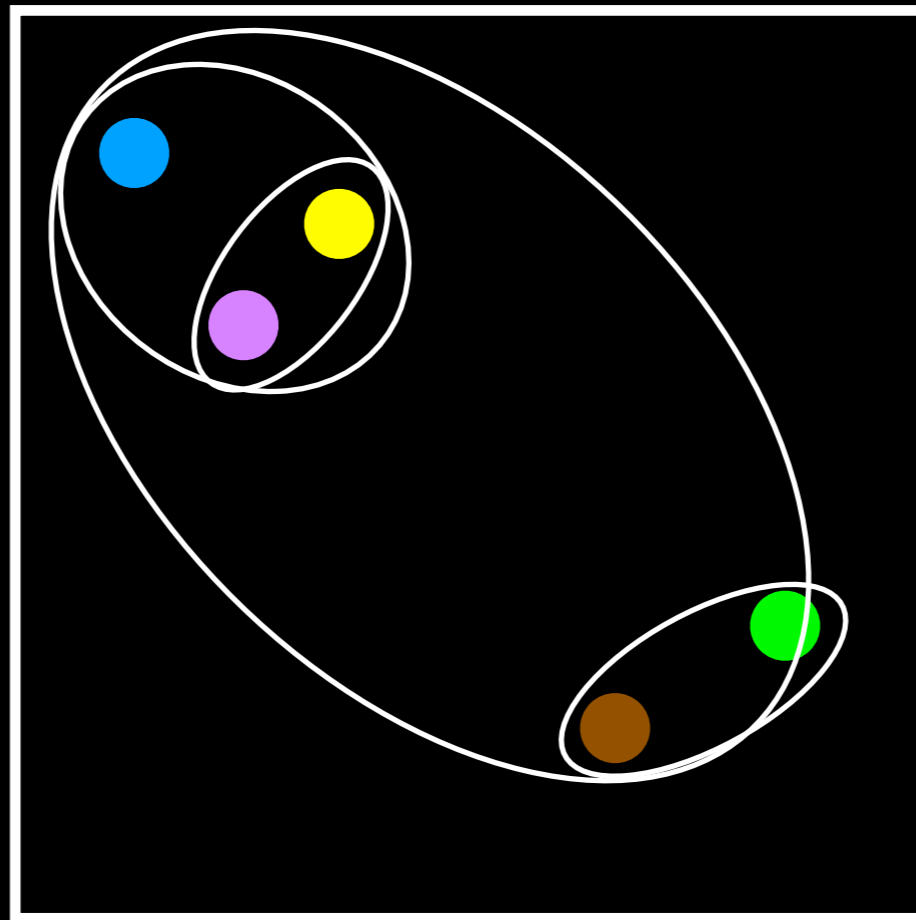
Hierarchical clustering in R

```
# First we need to calculate point (dis)similarity
#   as the Euclidean distance between observations
dist_matrix <- dist(x)

# The hclust() function returns a hierarchical
#   clustering model
hc <- hclust(d = dist_matrix)

# the print method is not so useful here
hc

Call:
hclust(d = dist_matrix)

Cluster method   : complete
Distance         : euclidean
Number of objects: 60
```

Lets have a closer look…

```r
# Our input is a distance matrix from the dist()
# function. Lets make sure we understand it first
dist_matrix <- dist(x)

dim(dist_matrix)
NULL

View( as.matrix(dist_matrix) )

dim(x)
[1] 60  2

dim( as.matrix(dist_matrix) )
[1] 60 60

# Note. symmetrical pairwise distance matrix
```

Interpreting results

```
# Create hierarchical cluster model: hc
hc <- hclust(dist(x))



# We can plot the results as a dendrogram
plot(hc)

# What do you notice?
# Does the dendrogram
# make sense based on
# your knoweledge of x?
```
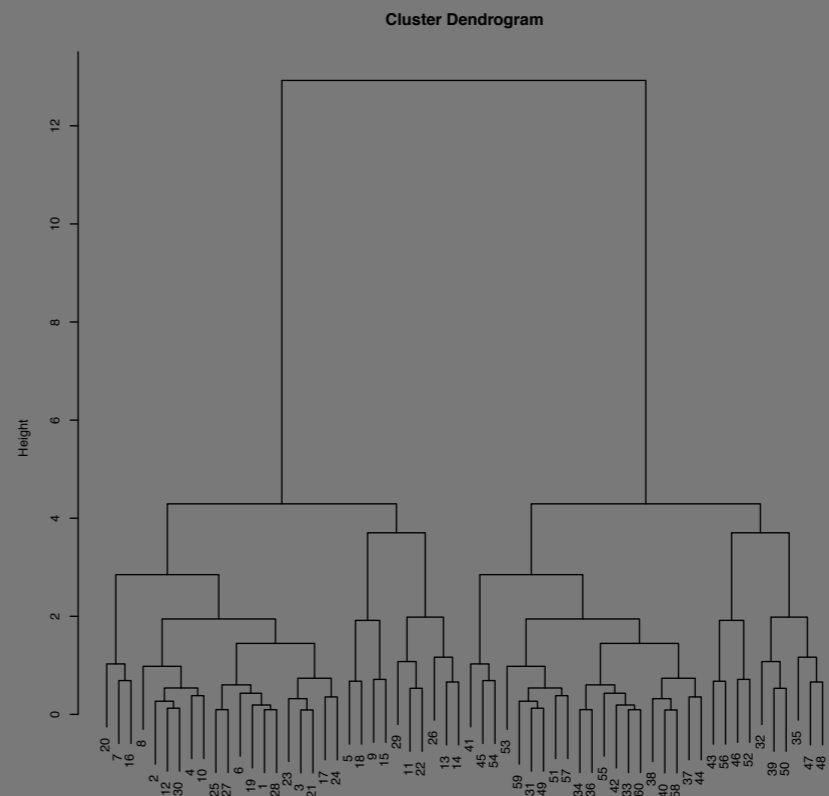

Cluster Dendrogram

Dendrogram

- Tree shaped structure used to interpret hierarchical clustering models

Dendrogram

- Tree shaped structure used to interpret hierarchical clustering models
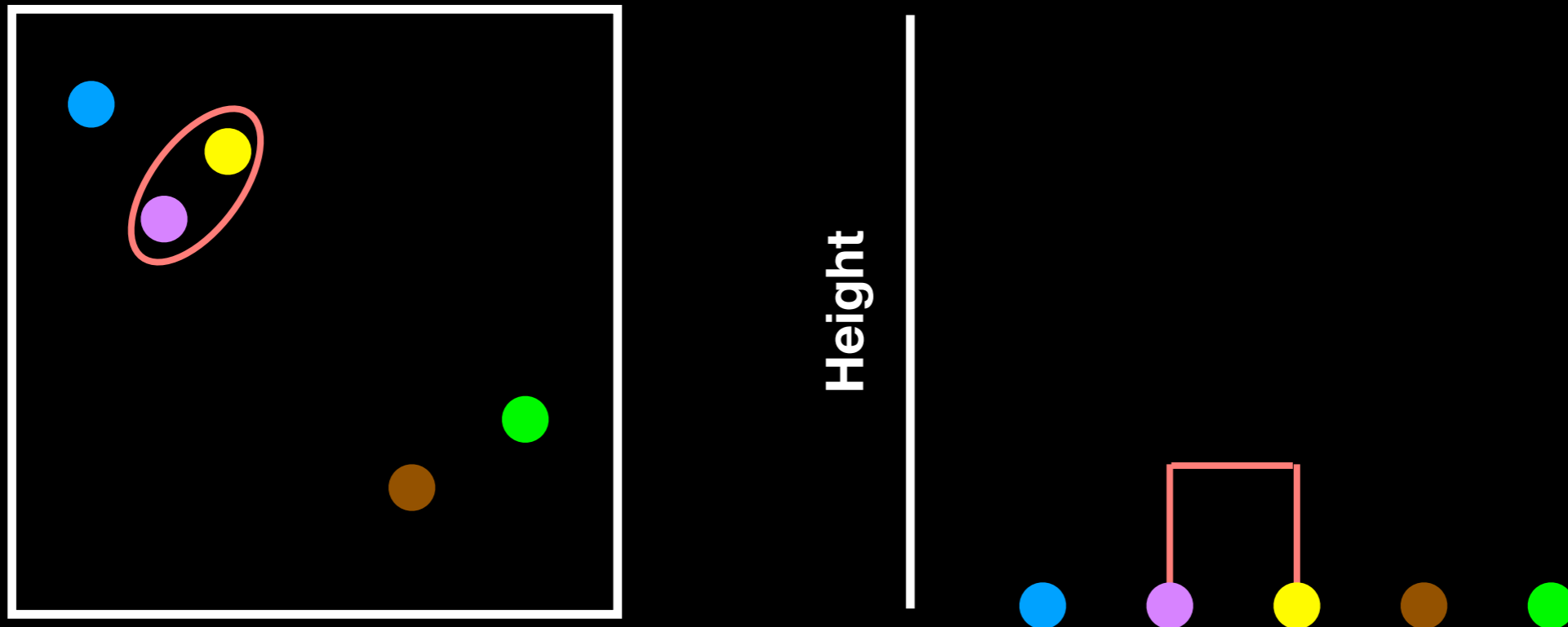
Dendrogram

- Tree shaped structure used to interpret hierarchical clustering models
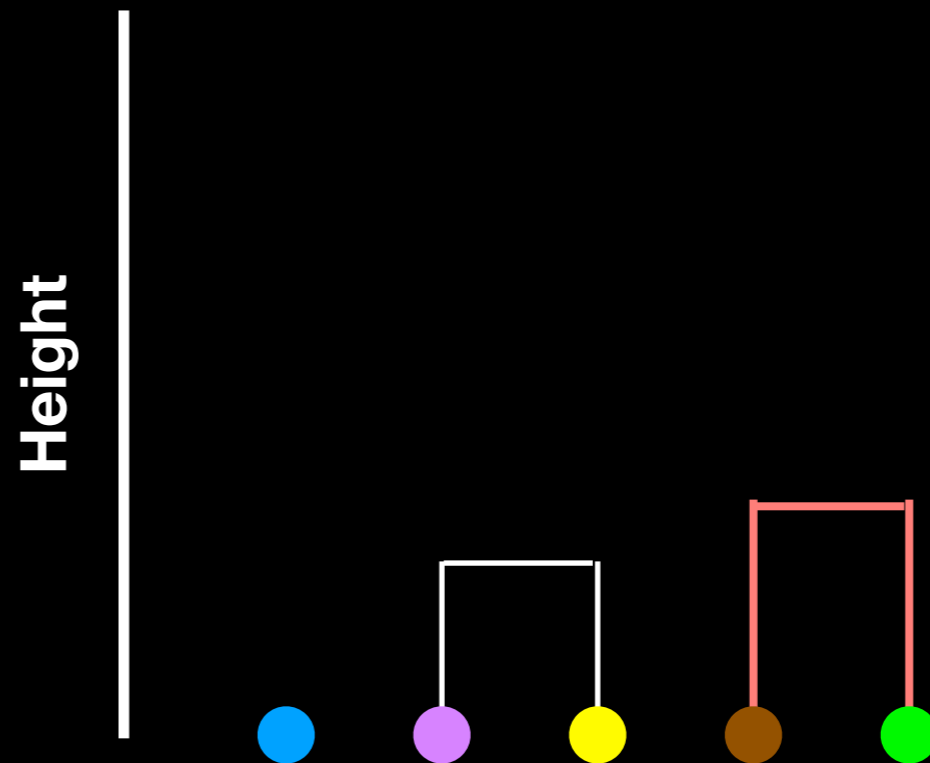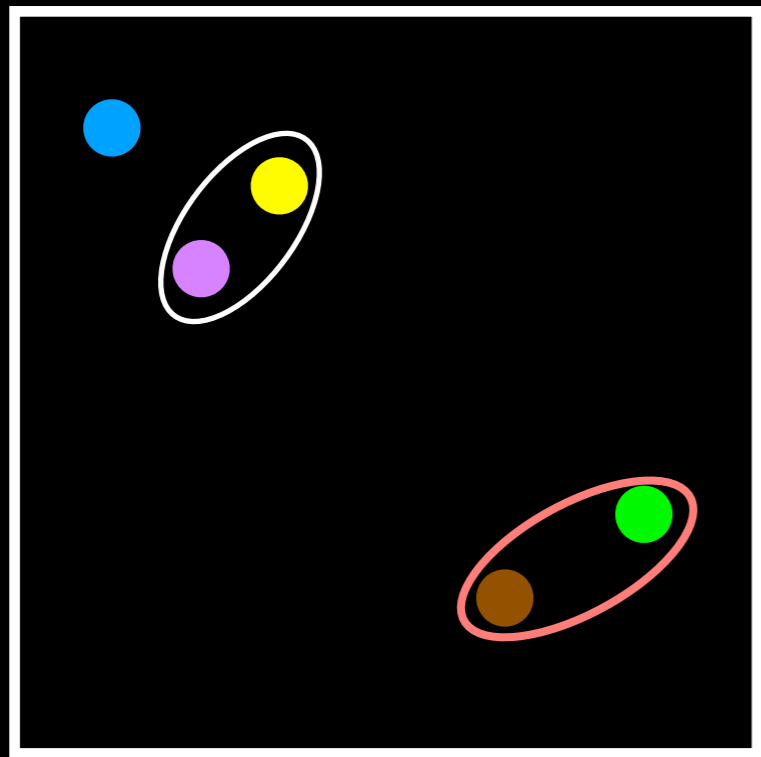
Dendrogram

- Tree shaped structure used to interpret hierarchical clustering models
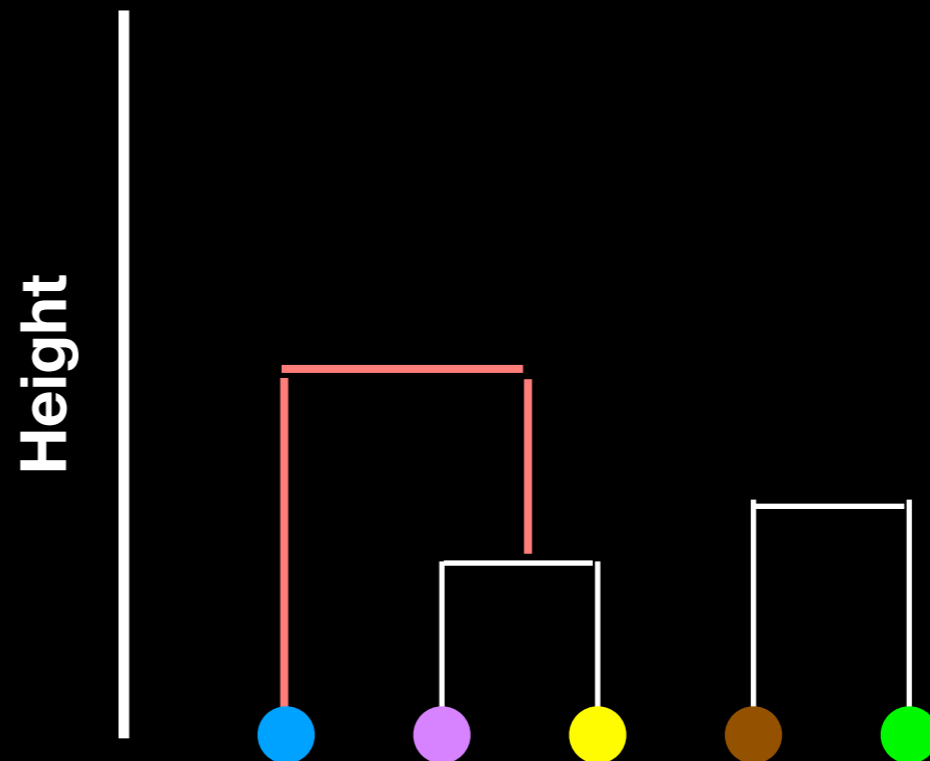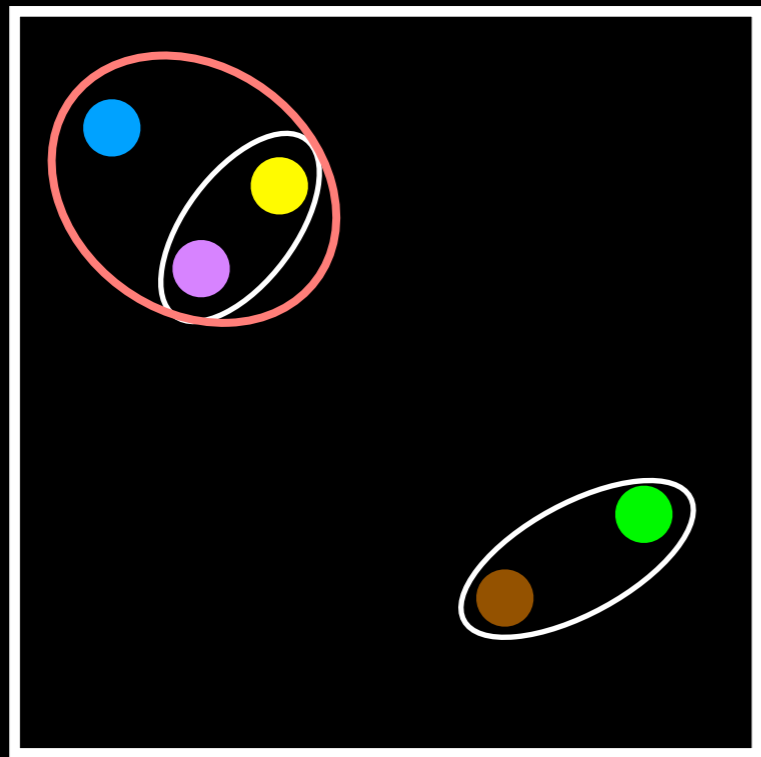
Dendrogram

- Tree shaped structure used to interpret hierarchical clustering models

Dendrogram

- Tree shaped structure used to interpret hierarchical clustering models
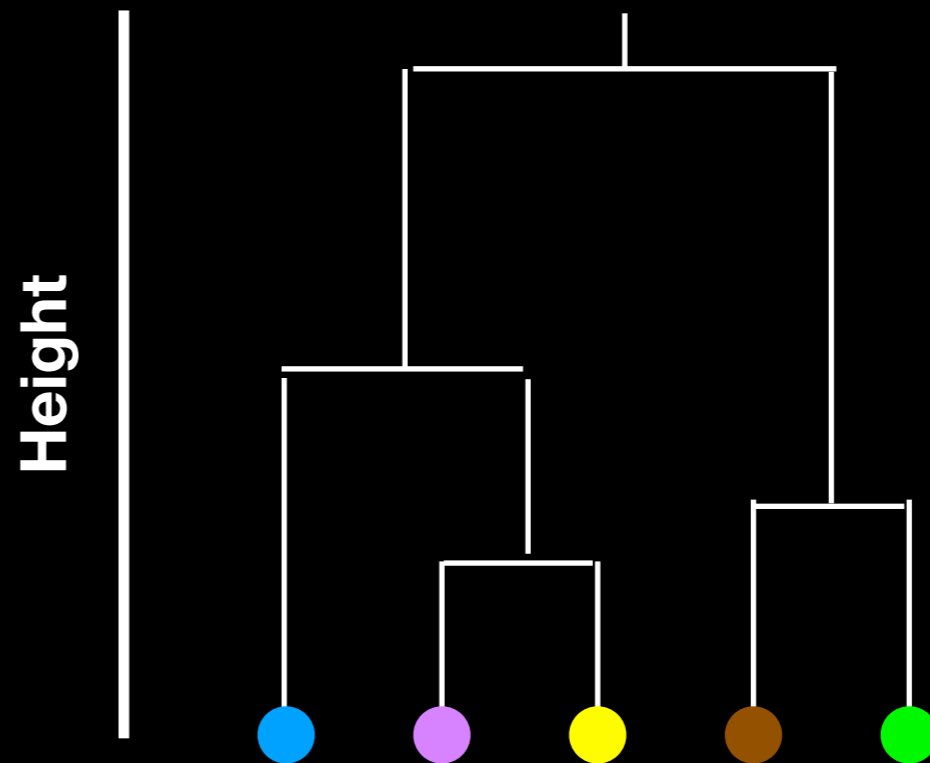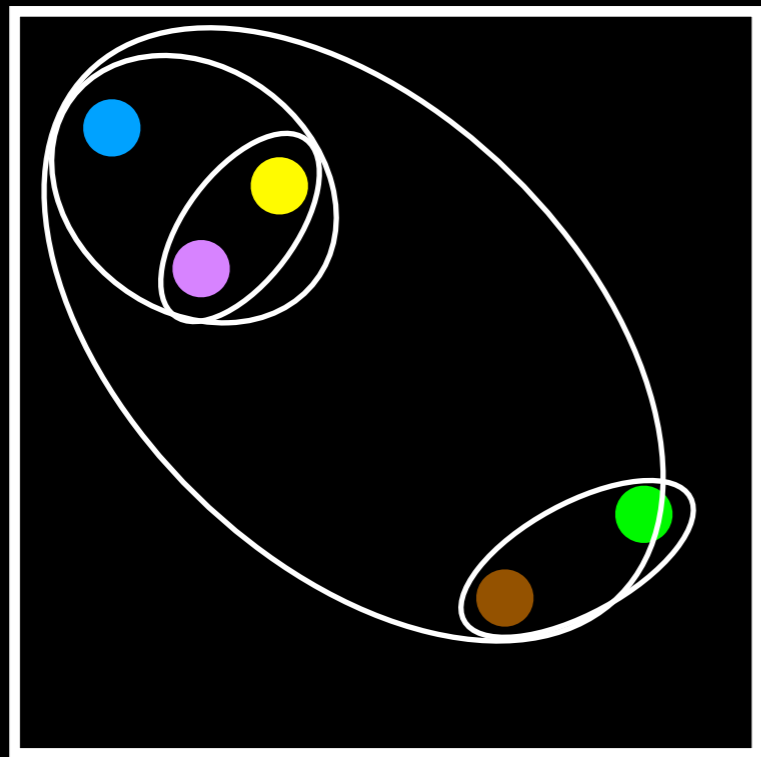
Dendrogram plotting in R
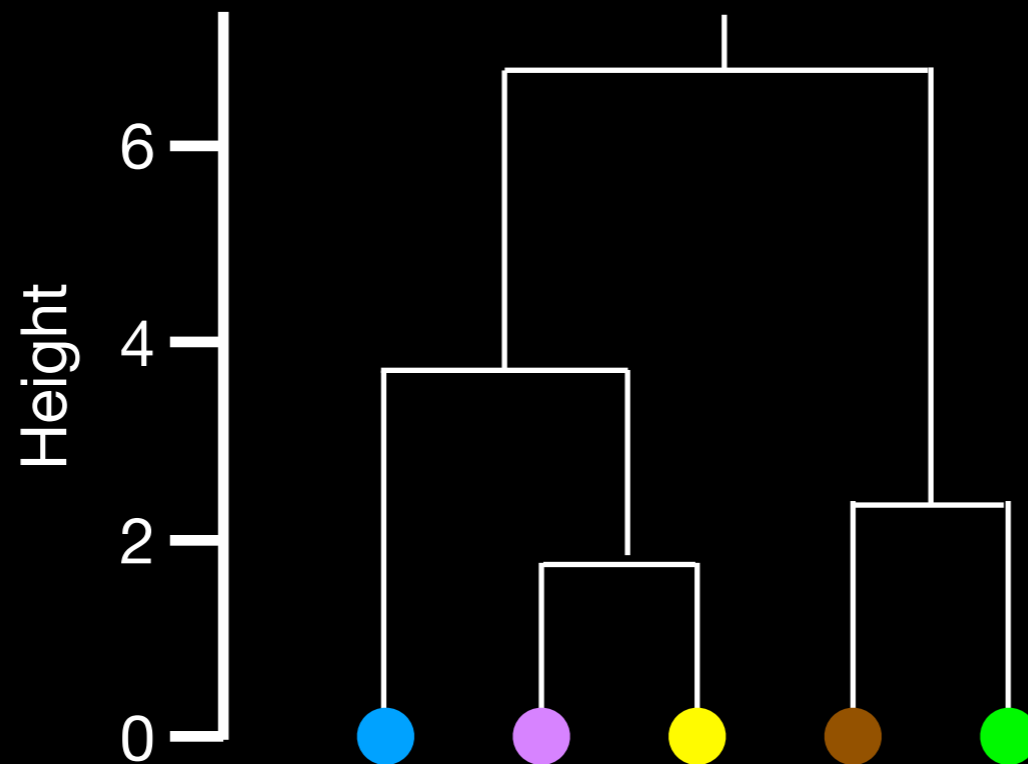
```
# Draws a dendrogram
plot(hc)
```

Dendrogram plotting in R
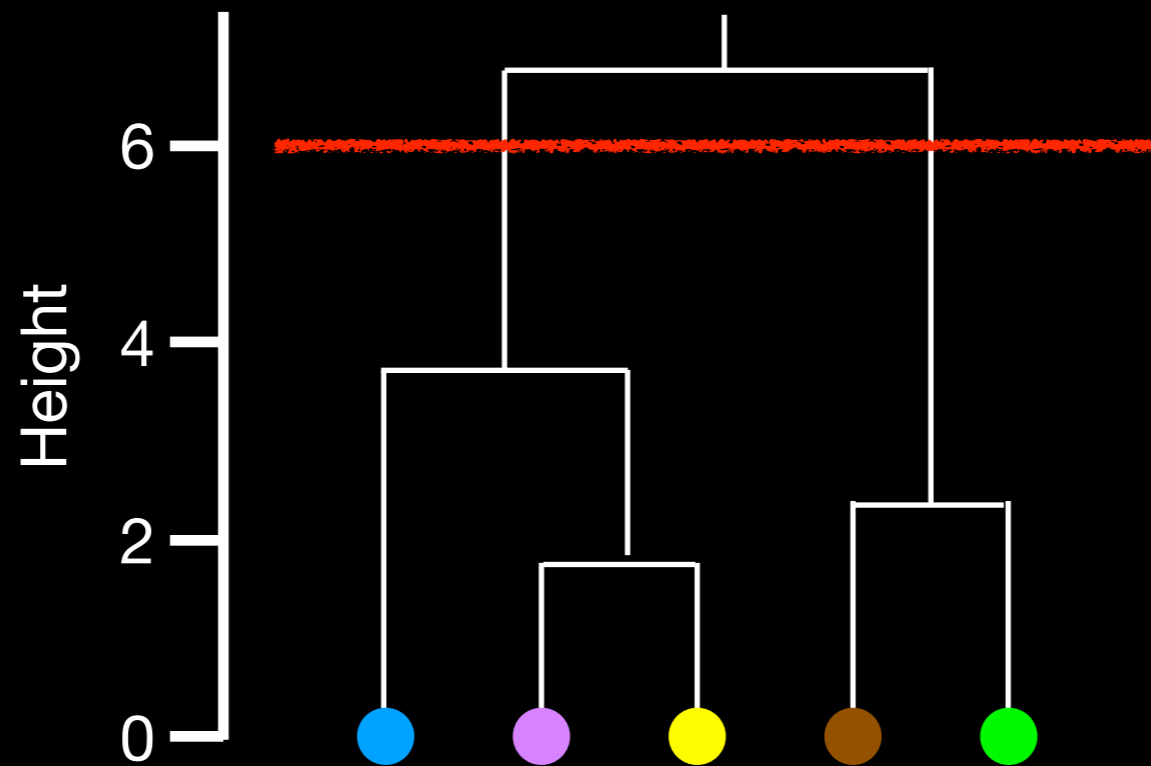
```
# Draws a dendrogram
plot(hc)
abline(h=6, col="red")
```

Dendrogram plotting in R

```r
# Draws a dendrogram
plot(hc)
abline(h=6, col="red")
```
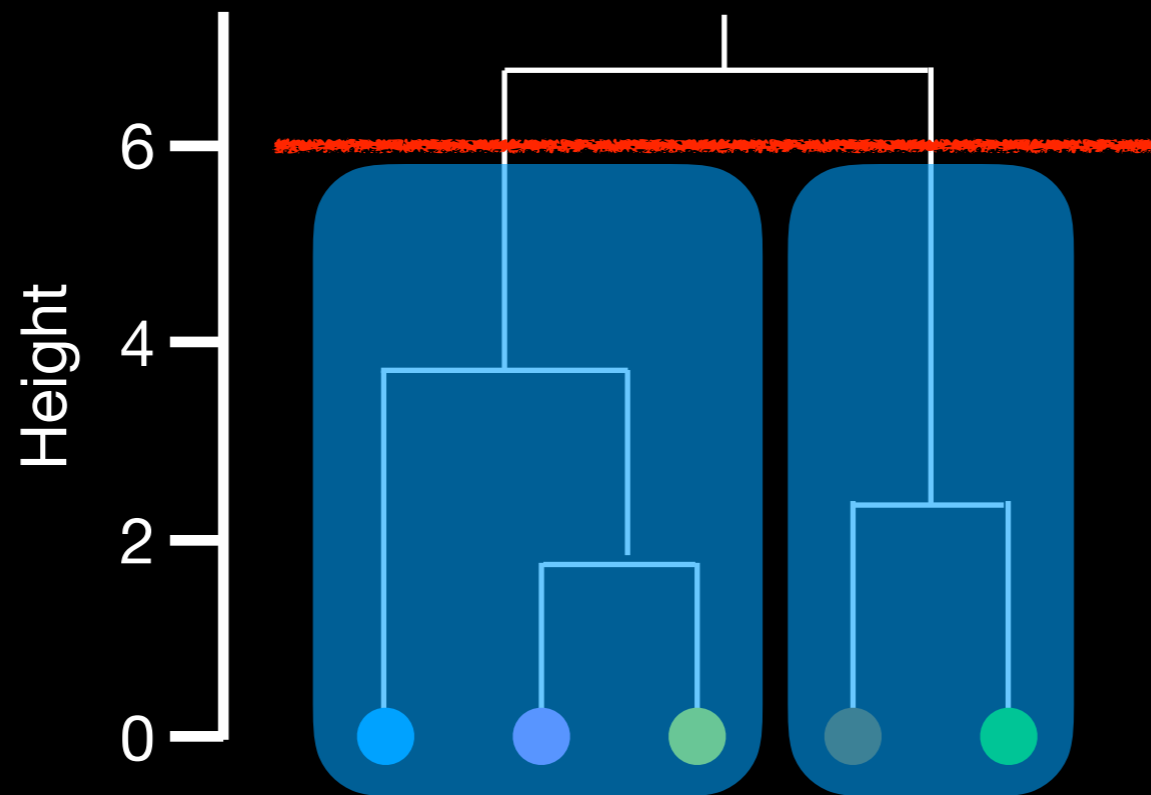
Dendrogram plotting in R

```
# Draws a dendrogram
plot(hc)
abline(h=6, col="red")
cutree(hc, h=6) # Cut by height h
[1] 1,1,1,2,2
```
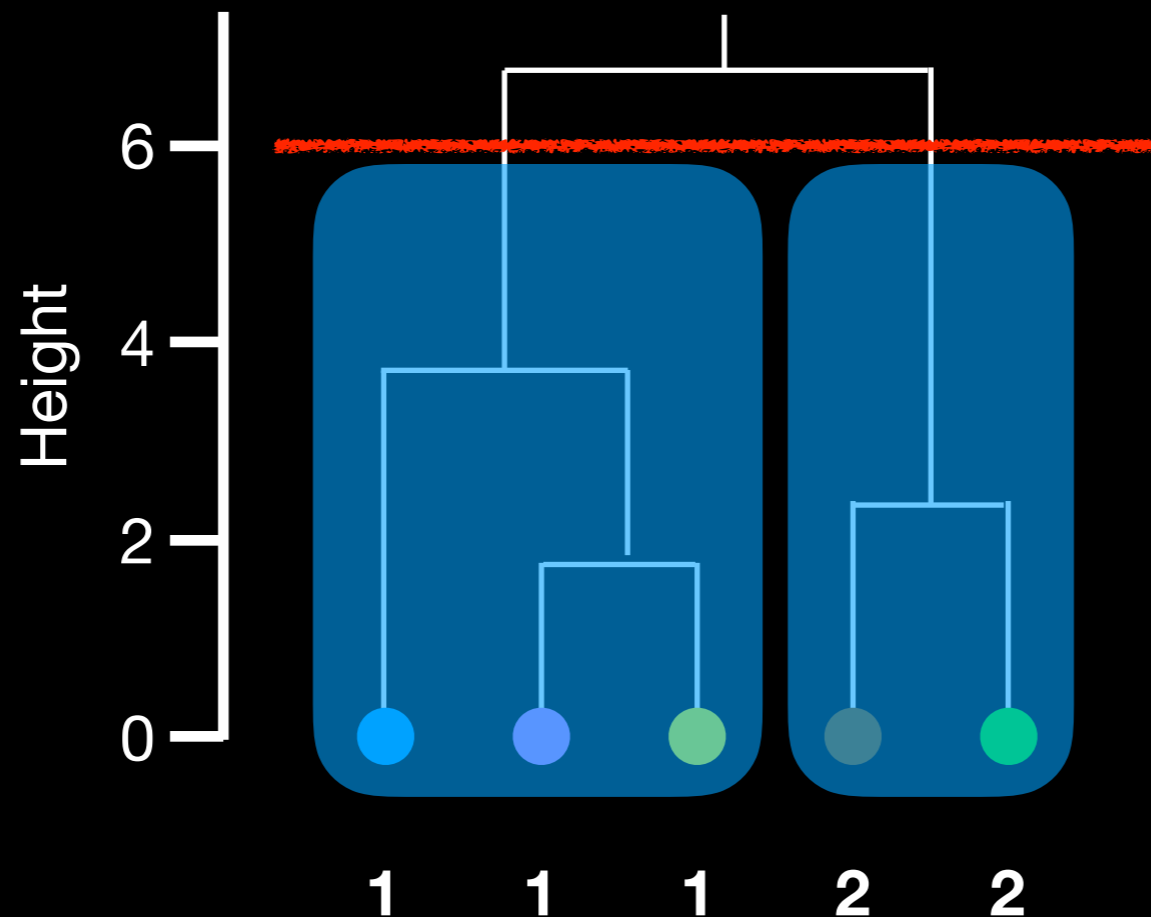
Dendrogram plotting in R

```
# Draws a dendrogram
plot(hc)
abline(h=6, col="red")
cutree(hc, k=2 ) # Cut into k grps
[1] 1,1,1,2,2
```
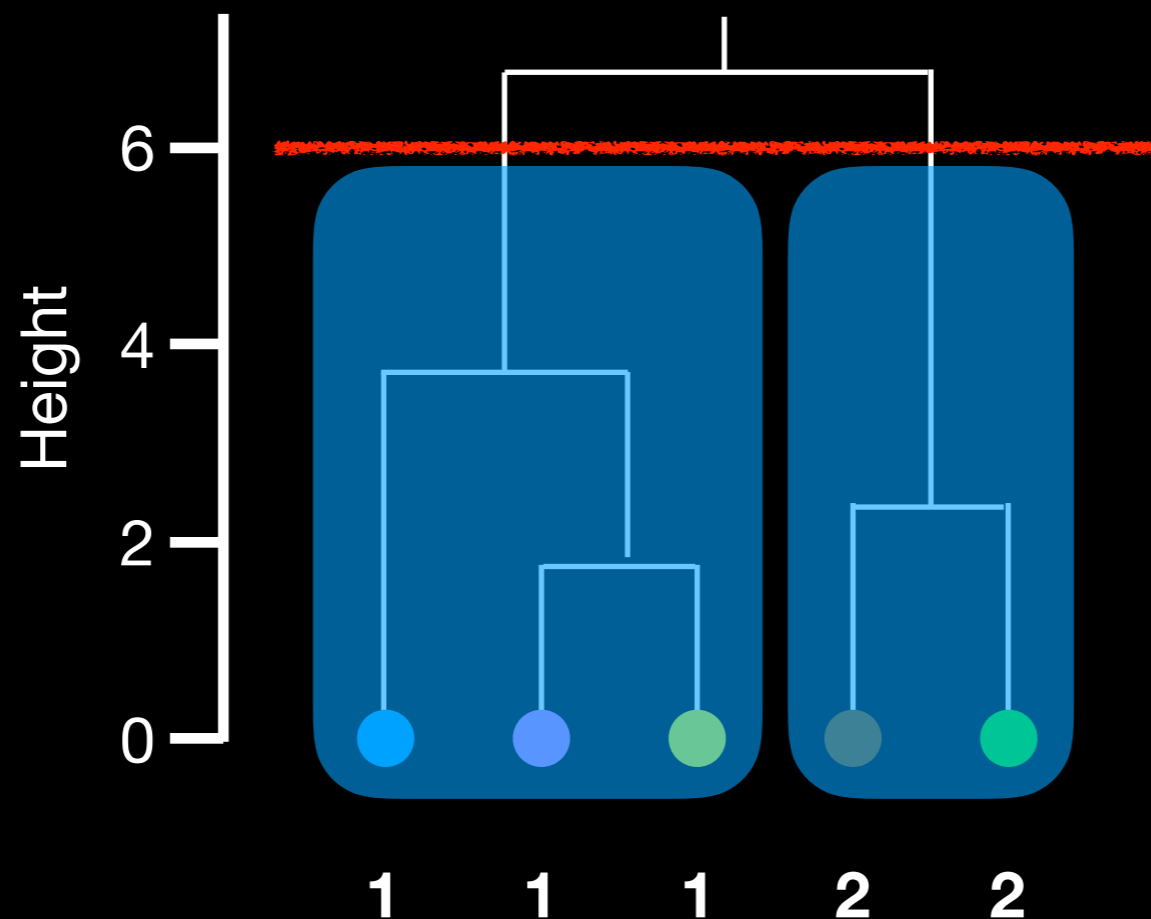
Linking clusters in hierarchical clustering

- How is distance between clusters determined?

- There are four main methods to determine which cluster should be linked:

  ➡ **Complete**: pairwise similarity between all observations in cluster 1 and cluster 2, and uses largest of similarities

  ➡ **Single**: same as above but uses smallest of similarities

  ➡ **Average**: same as above but uses average of similarities

  ➡ **Centroid**: finds centroid of cluster 1 and centroid of cluster 2, and uses similarity between two centroids

Linking methods: complete and average
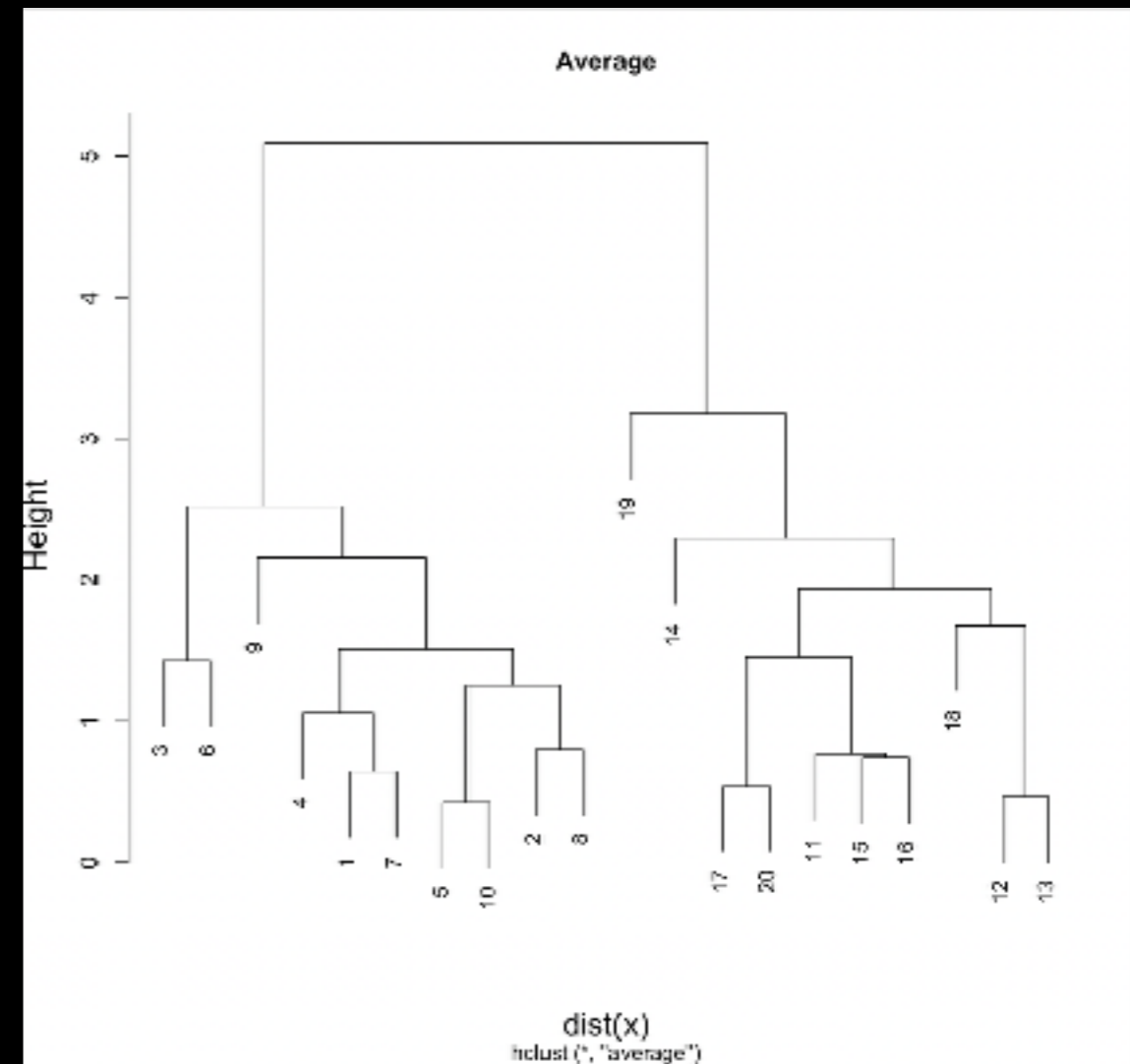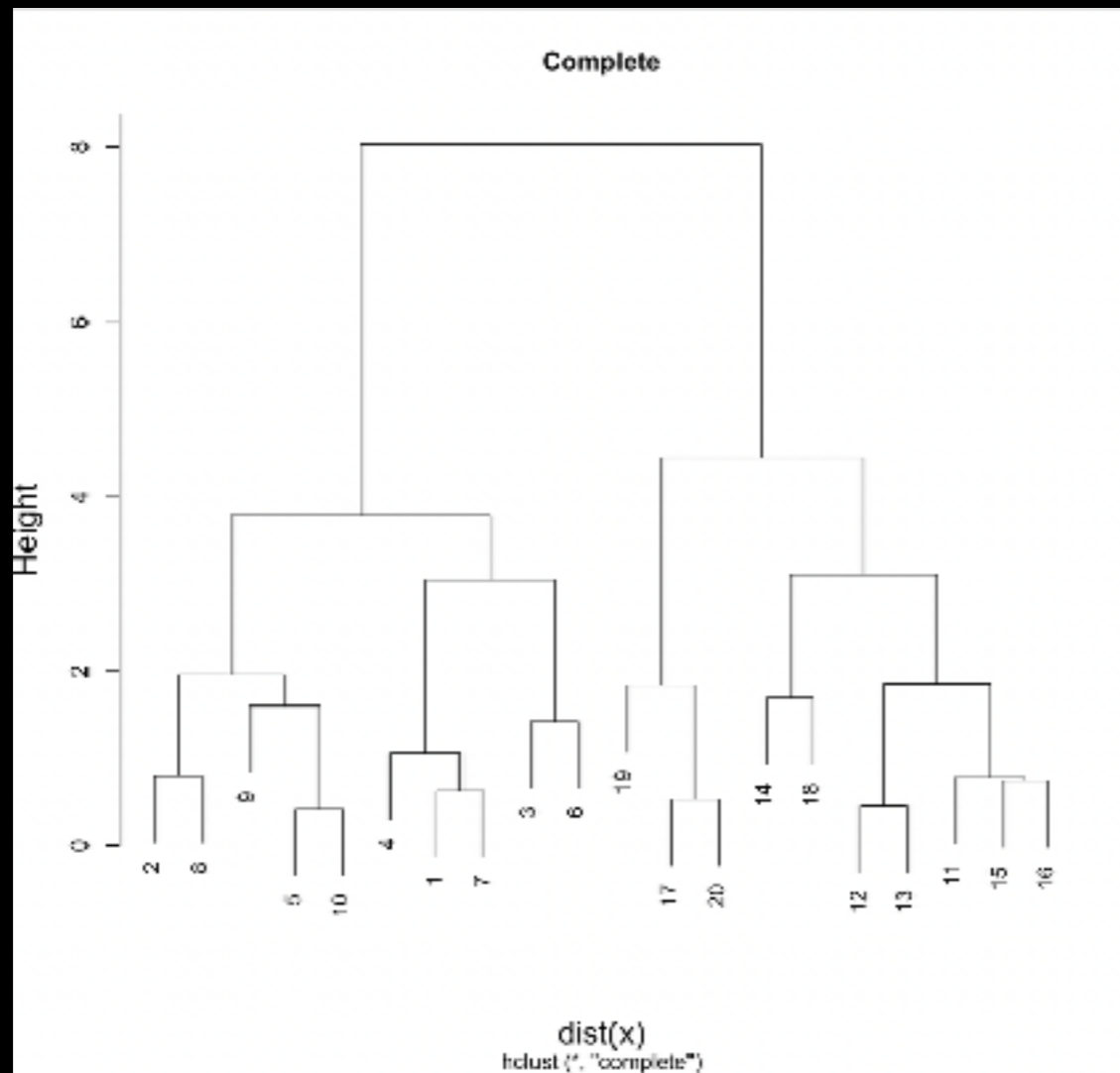
Linking method: single
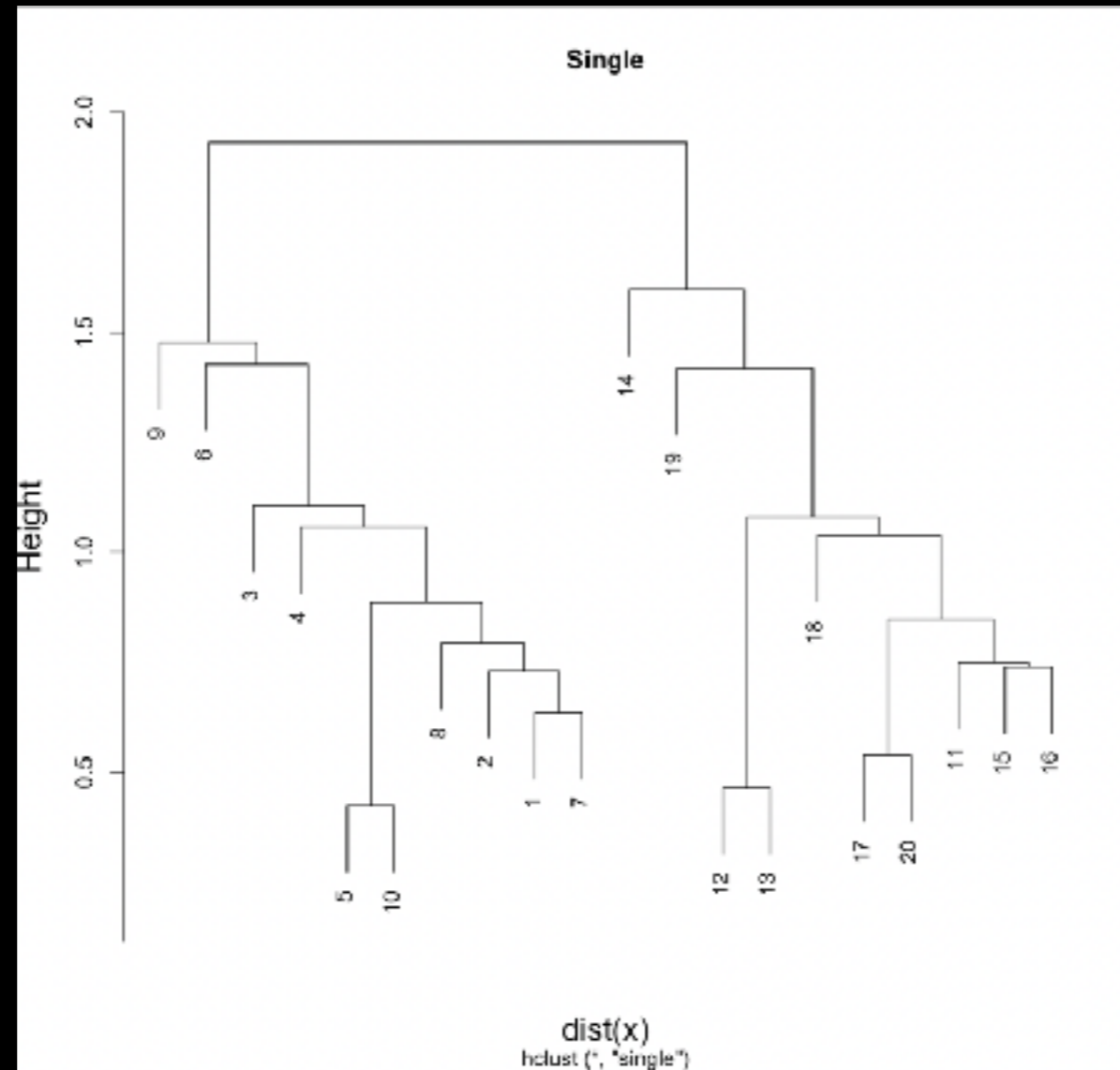
Linking method: centroid
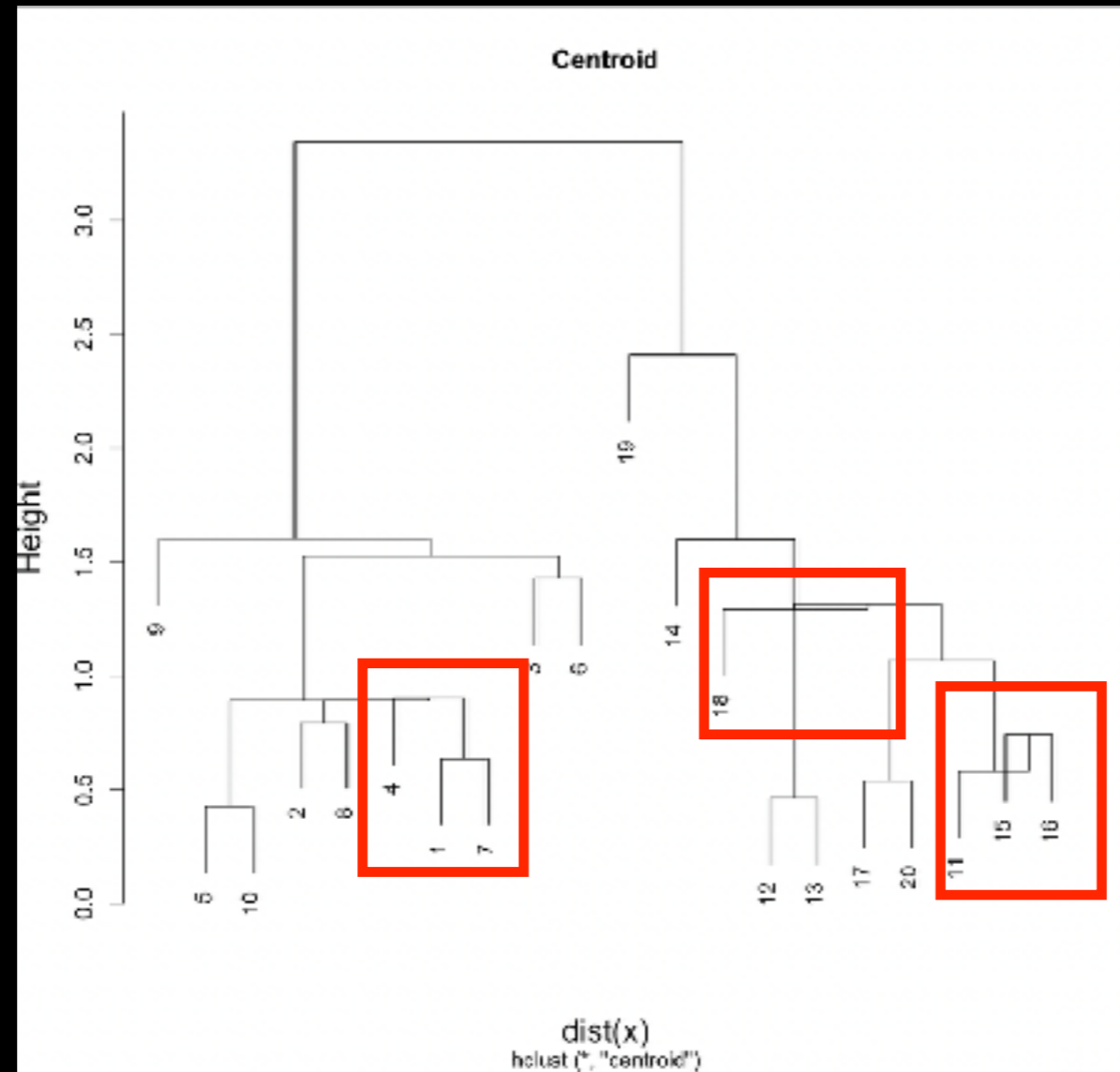
Linkage in R

```r
# Using different hierarchical clustering methods
hc.complete <- hclust(d, method="complete")

hc.average  <- hclust(d, method="average")

hc.single   <- hclust(d, method="single")
```

Side-Note: Practical matters

- Data on different scales can cause undesirable results in clustering methods

- Solution is to scale data so that features have same mean and standard deviation
  - ➡ Subtract mean of a feature from all observations
  - ➡ Divide each feature by the standard deviation of the feature
  - ➡ Normalized features have a mean of zero and a standard deviation of one

Side-Note: Practical matters

```
# Check if scaling is necessary
colMeans(x)    # x is a data matrix
[1] -0.1337828  0.0594019

apply(x, 2, sd)
[1] 1.974376 2.112357

# New matrix with col mean of 0 and sd of 1
scaled_x <- scale(x)

colMeans(scaled_x)
[1] 2.775558e-17 3.330669e-17

apply(scaled_x, 2, sd)
[1] 1 1
```

Your Turn!

```r
# Step 1. Generate some example data for clustering
x <- rbind(
  matrix(rnorm(100, mean=0, sd = 0.3), ncol = 2),    # c1
  matrix(rnorm(100, mean = 1, sd = 0.3), ncol = 2), # c2
  matrix(c(rnorm(50, mean = 1, sd = 0.3),            # c3
           rnorm(50, mean = 0, sd = 0.3)), ncol = 2))

colnames(x) <- c("x", "y")

# Step 2. Plot the data without clustering
plot(x)

# Step 3. Generate colors for known clusters
#         (just so we can compare to hclust results)
col <- as.factor( rep(c("c1","c2","c3"), each=50) )

plot(x, col=col)

# Q. Use the dist(), hclust(), plot() and cutree()
#      functions to return 2 and 3 clusters
# Q. How does this compare to your known 'col' groups?
```

Today's Menu

- Introduction to machine learning
  - <u>Unsupervised</u>, supervised and reinforcement learning

- Clustering
  - K-means clustering
  - Hierarchical clustering
  - Heatmap representations

- Dimensionality reduction, visualization and 'structure' analysis
  - Principal Component Analysis (PCA)

- Hands-on application to cell classification

Bunch of "normal" cells
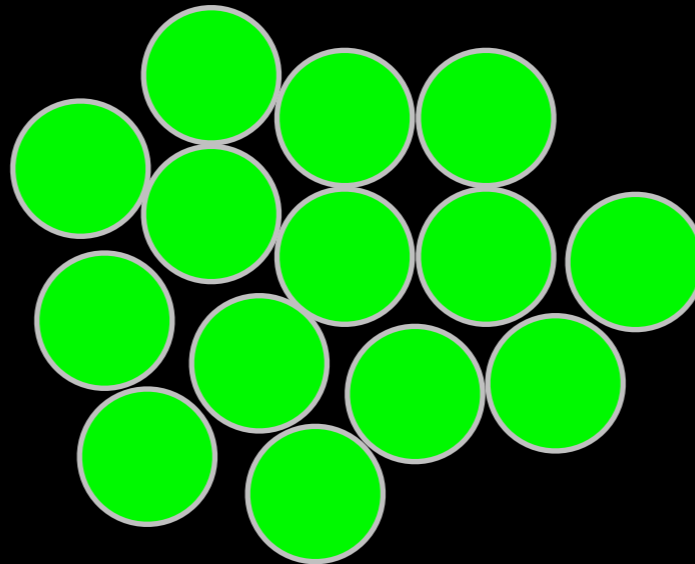
Even though they look the same we suspect that there are differences…

These might be
one cell type...
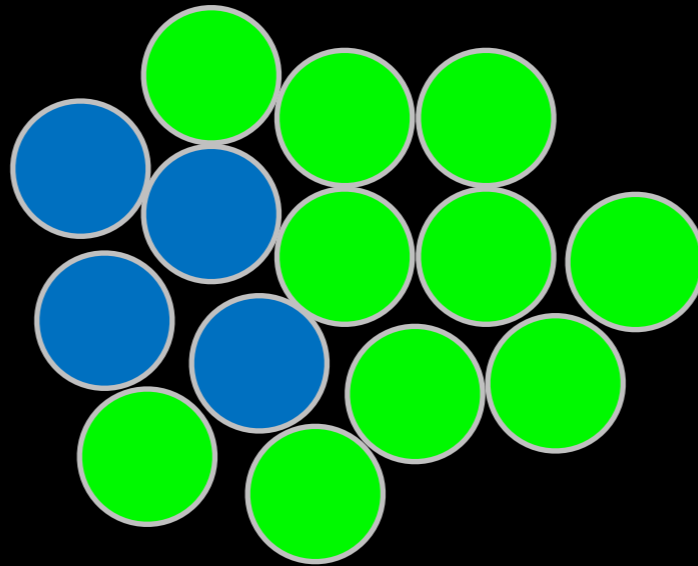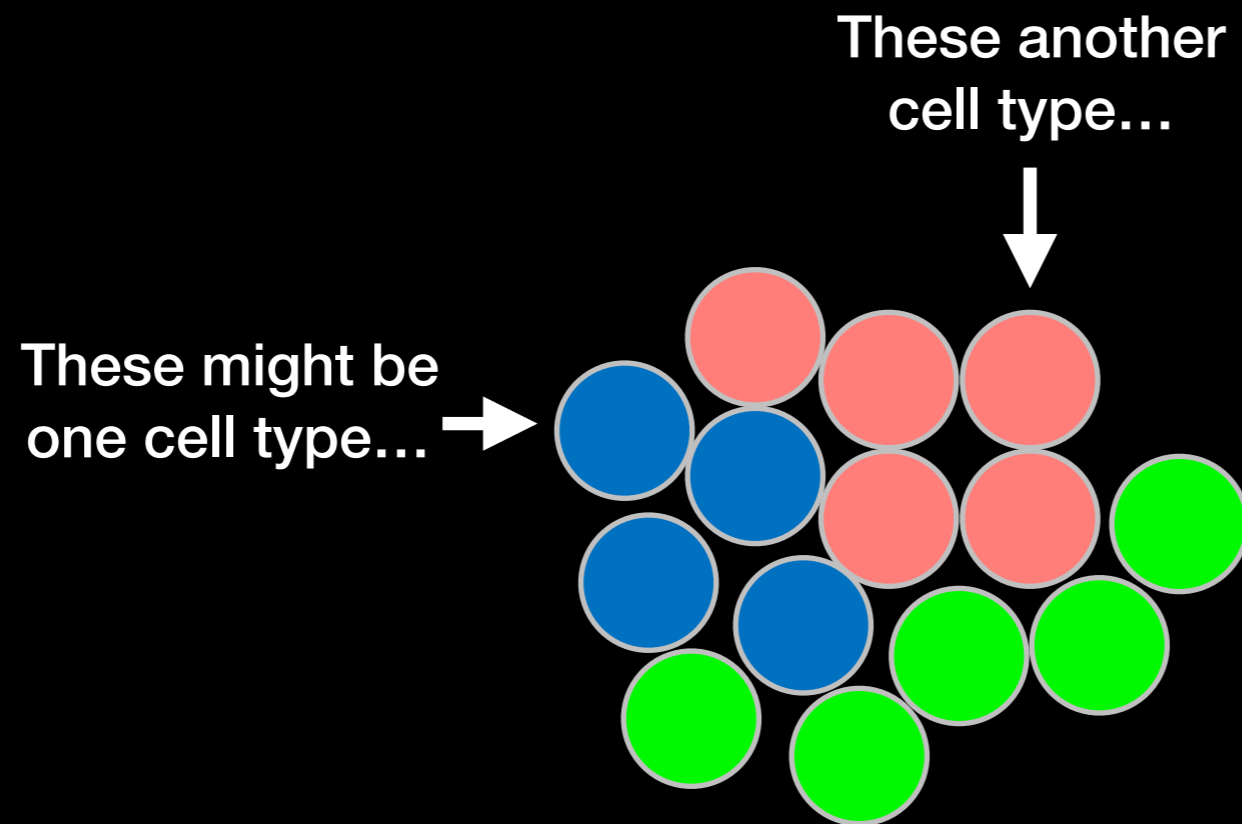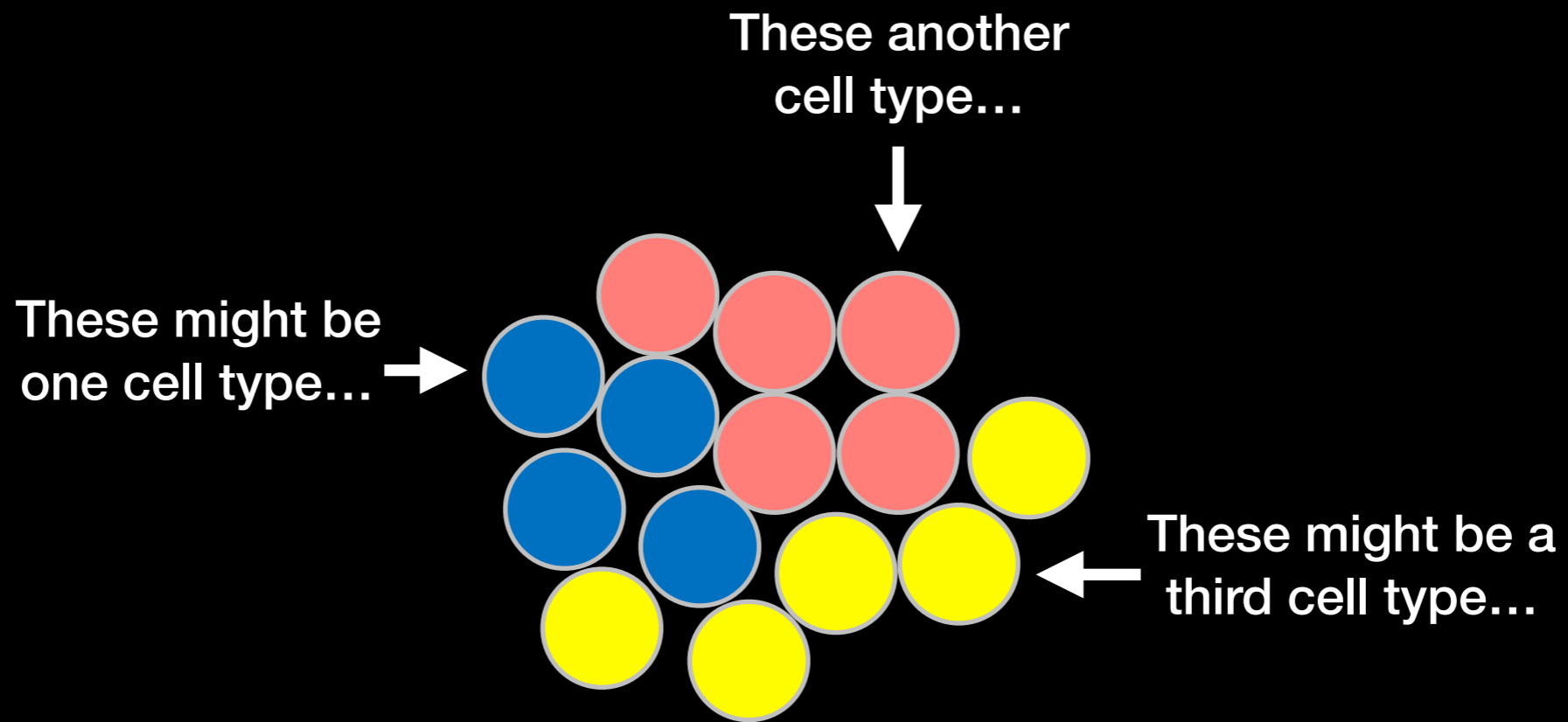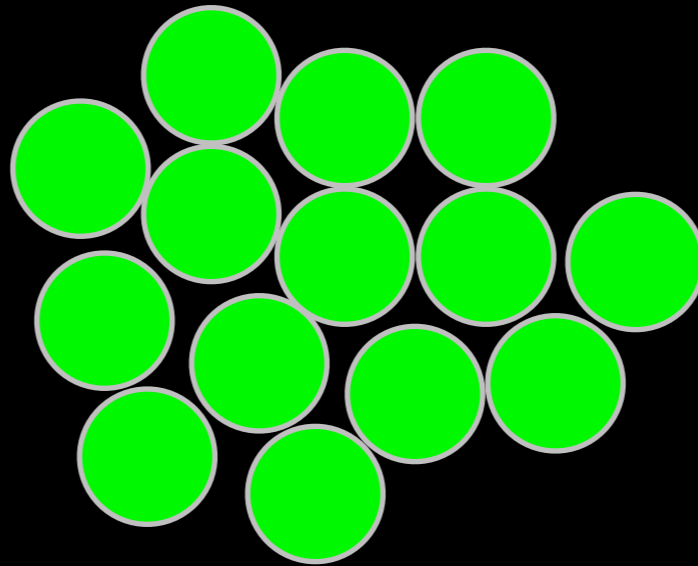
Unfortunately we can't observe
the differences visually
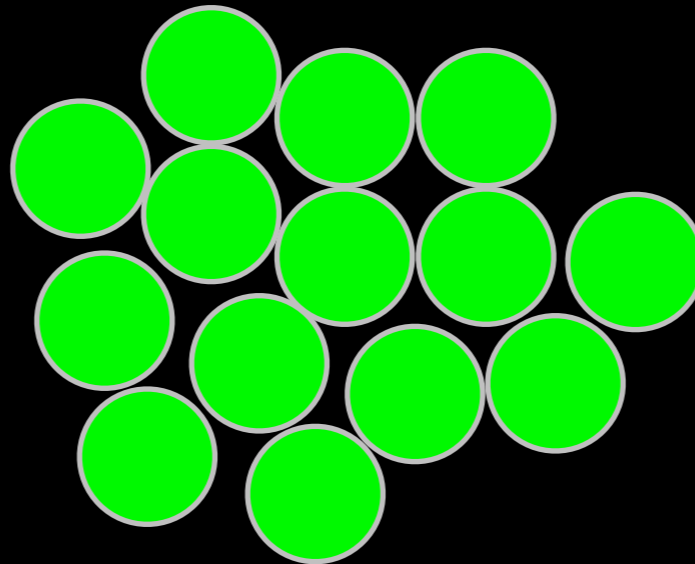
Unfortunately we can't observe
the differences visually

So we sequence the mRNA in each
cell to identify which genes are
active and at what levels.

Here is the data…

	Cell1	Cell2	Cell3	Cell4	…
Gene1	3	0.25	2.8	0.1	…
Gene2	2.9	0.8	2.2	1.8	…
Gene3	2.2	1	1.5	3.2	…
Gene4	2	1.4	2	0.3	…
Gene5	1.3	1.6	1.6	0	…
Gene6	1.5	2	2.1	3	…
Gene7	1.1	2.2	1.2	2.8	…
Gene8	1	2.7	0.9	0.3	…
Gene9	0.4	3	0.6	0.1	…

Each column shows how much each
gene is transcribed in each cell

Here is the data...

	Cell1	Cell2	Cell3	Cell4	...
Gene1	3	0.25	2.8	0.1	...
Gene2	2.9	0.8	2.2	1.8	...
Gene3	2.2	1	1.5	3.2	...
Gene4	2	1.4	2	0.3	...
Gene5	1.3	1.6	1.6	0	...
Gene6	1.5	2	2.1	3	...
Gene7	1.1	2.2	1.2	2.8	...
Gene8	1	2.7	0.9	0.3	...
Gene9	0.4	3	0.6	0.1	...

For now lets consider
only two cells

	Cell1	Cell2
Gene1	3	0.25
Gene2	2.9	0.8
Gene3	2.2	1
Gene4	2	1.4
Gene5	1.3	1.6
Gene6	1.5	2
Gene7	1.1	2.2
Gene8	1	2.7
Gene9	0.4	3

We have just 2 cells so we can plot the measurements for each gene
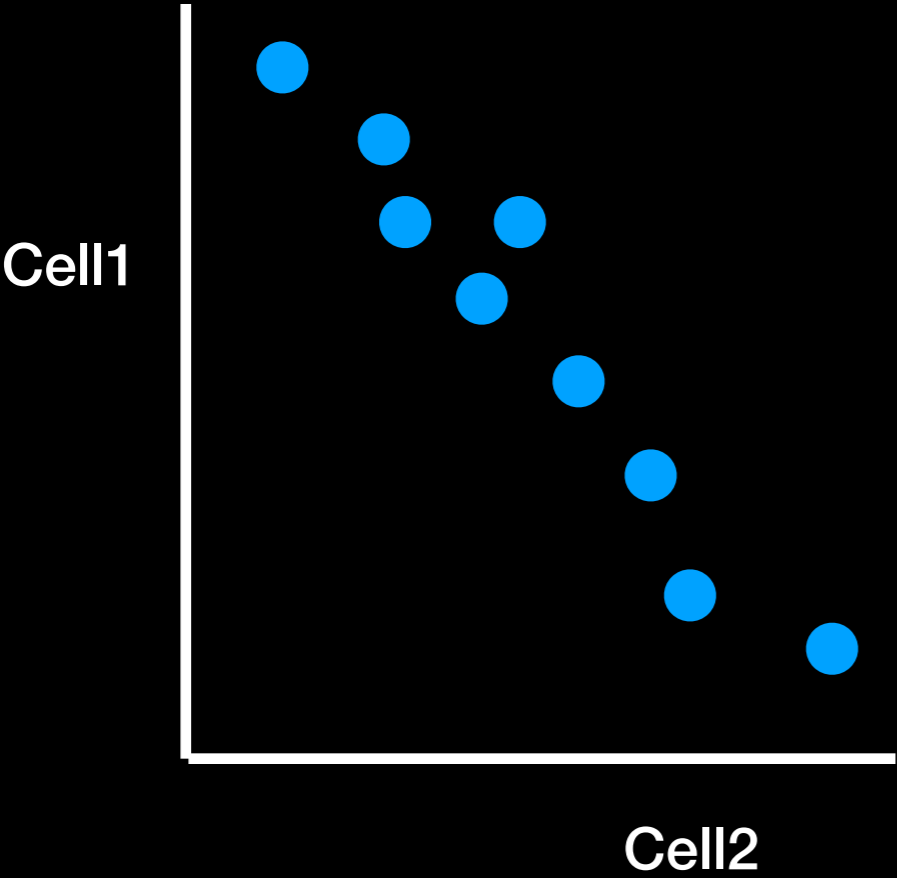
	Cell1	Cell2
Gene1	3	0.25
Gene2	2.9	0.8
Gene3	2.2	1
Gene4	2	1.4
Gene5	1.3	1.6
Gene6	1.5	2
Gene7	1.1	2.2
Gene8	1	2.7
Gene9	0.4	3

This gene (Gene1) is highly transcribed in Cell1 and lowly transcribed in Cell2...
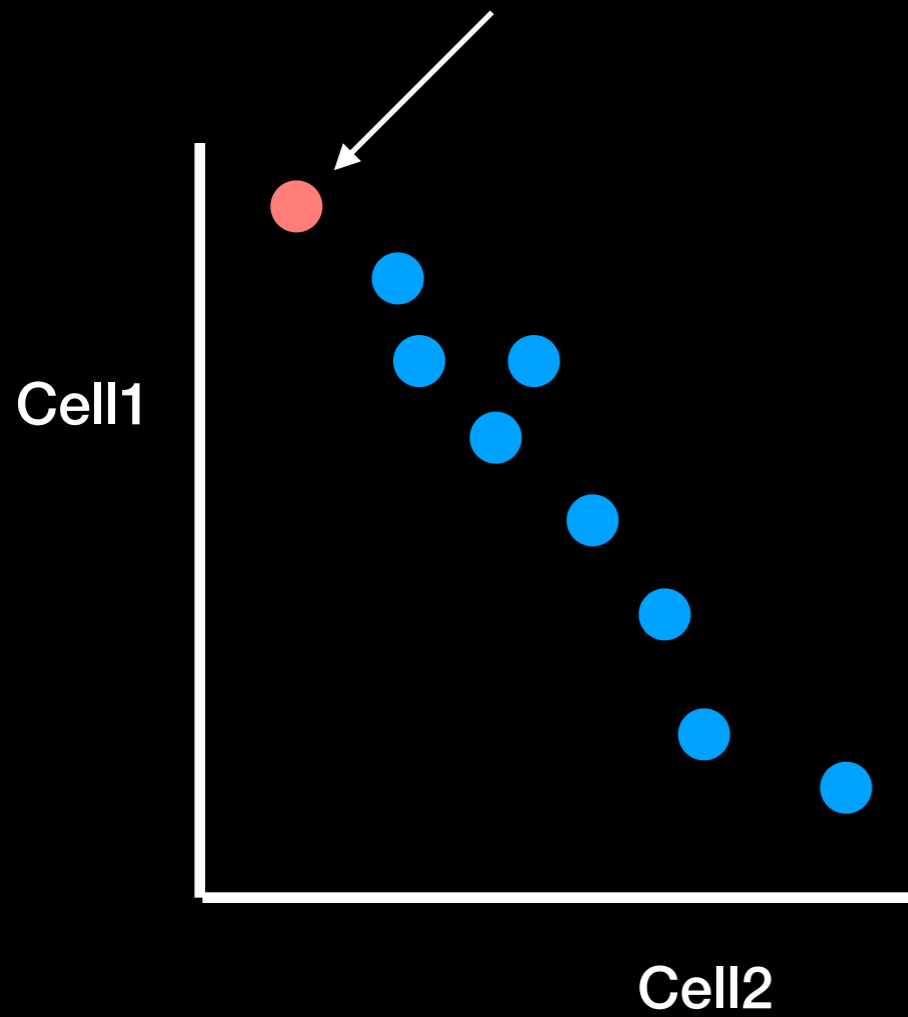
Cell1

Cell2

	Cell1	Cell2
Gene1	3	0.25
Gene2	2.9	0.8
Gene3	2.2	1
Gene4	2	1.4
Gene5	1.3	1.6
Gene6	1.5	2
Gene7	1.1	2.2
Gene8	1	2.7
Gene9	0.4	3

This gene (Gene9) is lowly transcribed in Cell1 and highly transcribed in Cell2…
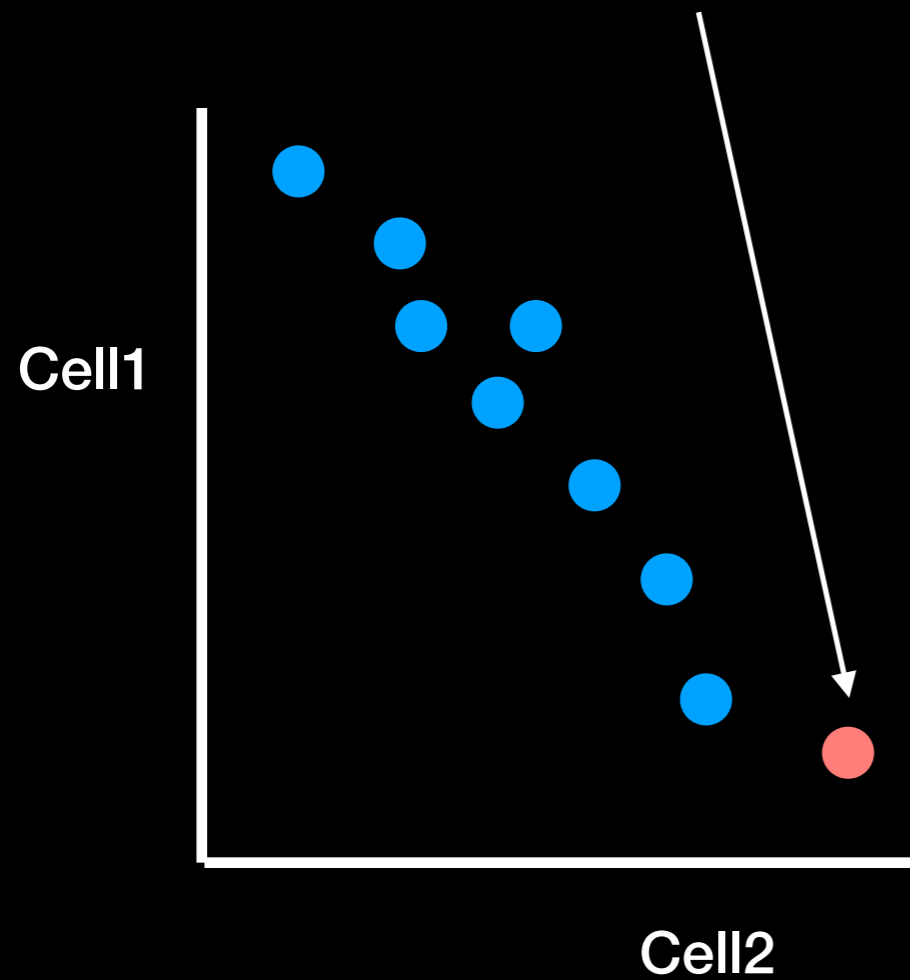
Cell1

Cell2

	Cell1	Cell2
Gene1	3	0.25
Gene2	2.9	0.8
Gene3	2.2	1
Gene4	2	1.4
Gene5	1.3	1.6
Gene6	1.5	2
Gene7	1.1	2.2
Gene8	1	2.7
Gene9	0.4	3

In generel, Cell1 and Cell2 have an **inverse correlation**.

This suggests that they may be two different types of cells as they are using different genes
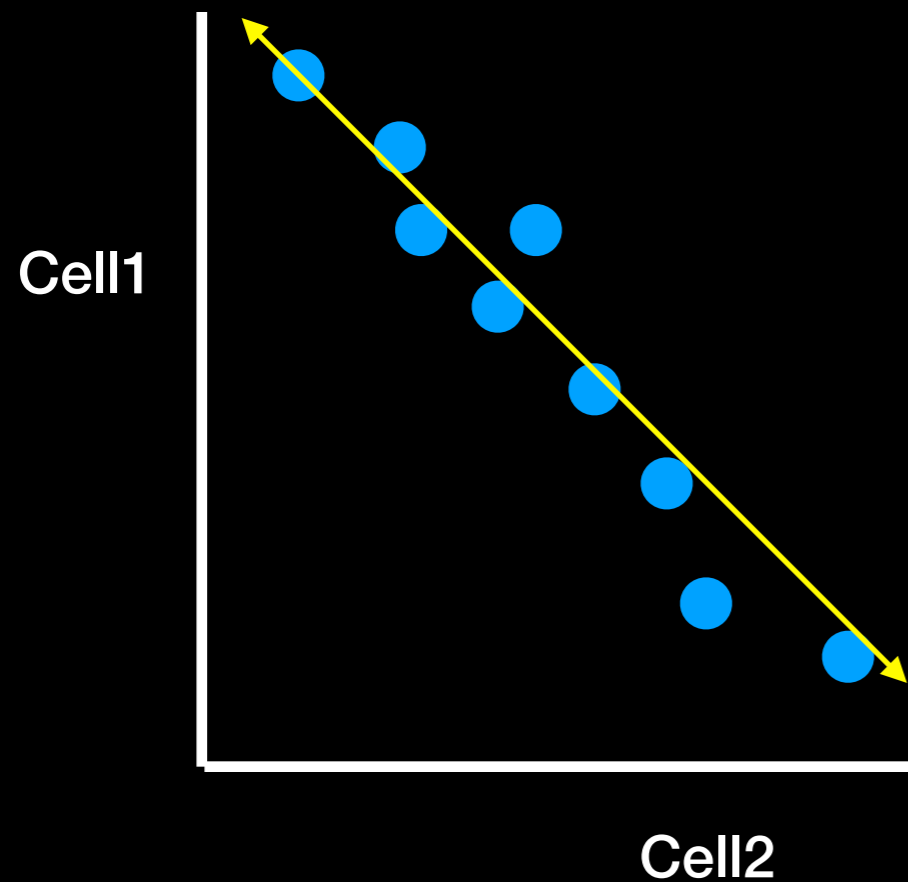
	Cell1	Cell2
Gene1	3	0.25
Gene2	2.9	0.8
Gene3	2.2	1
Gene4	2	1.4
Gene5	1.3	1.6
Gene6	1.5	2
Gene7	1.1	2.2
Gene8	1	2.7
Gene9	0.4	3

Now lets imagine
there are three cells

	Cell1	Cell2	Cell3
Gene1	3	0.25	2.8
Gene2	2.9	0.8	2.2
Gene3	2.2	1	1.5
Gene4	2	1.4	2
Gene5	1.3	1.6	1.6
Gene6	1.5	2	2.1
Gene7	1.1	2.2	1.2
Gene8	1	2.7	0.9
Gene9	0.4	3	0.6

We have already seen how we can plot the first two cells to see how closely related they are
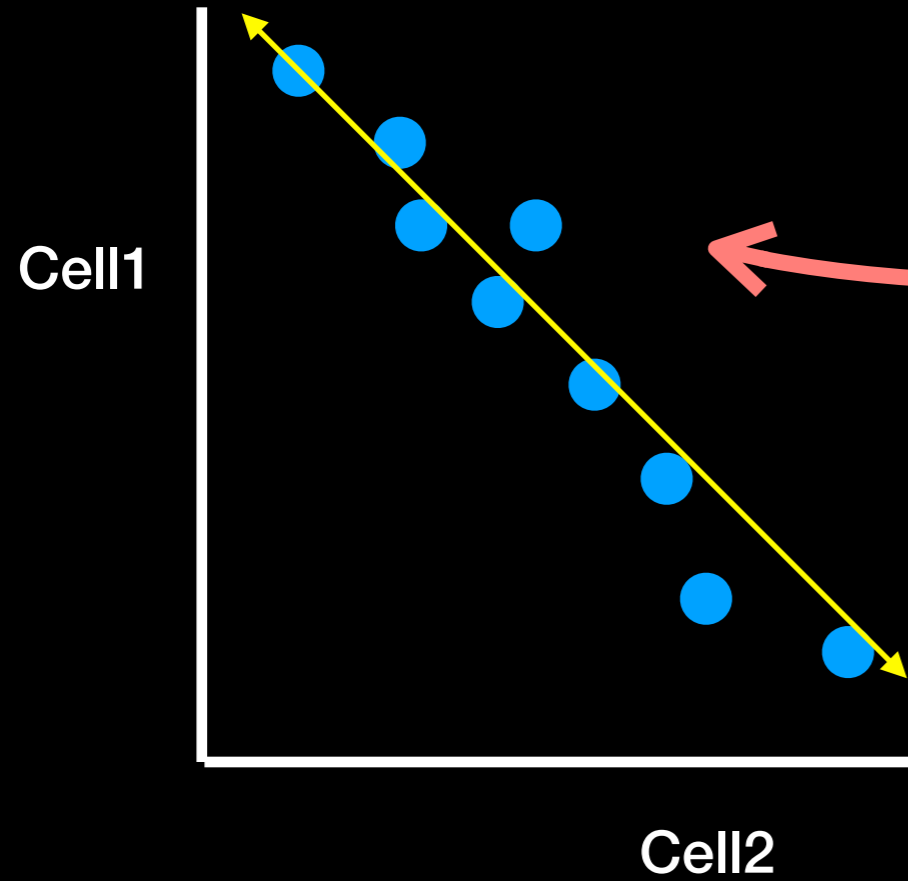
	Cell1	Cell2	Cell3
Gene1	3	0.25	2.8
Gene2	2.9	0.8	2.2
Gene3	2.2	1	1.5
Gene4	2	1.4	2
Gene5	1.3	1.6	1.6
Gene6	1.5	2	2.1
Gene7	1.1	2.2	1.2
Gene8	1	2.7	0.9
Gene9	0.4	3	0.6
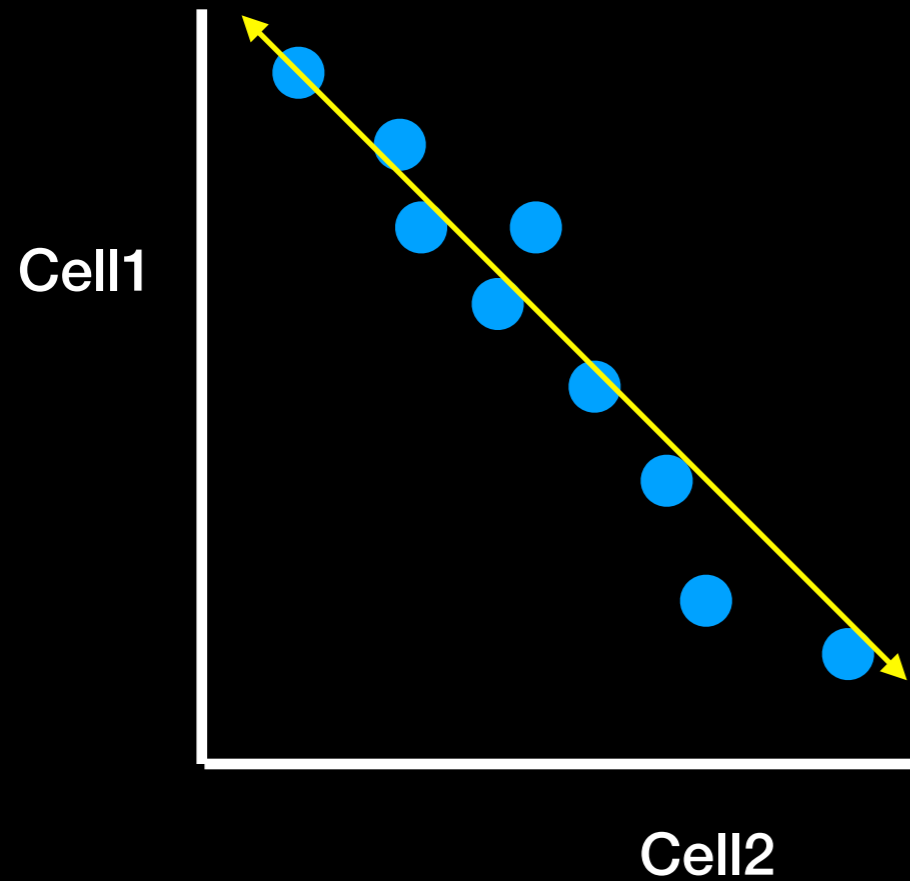
Now we can also compare
Cell1 to Cell3
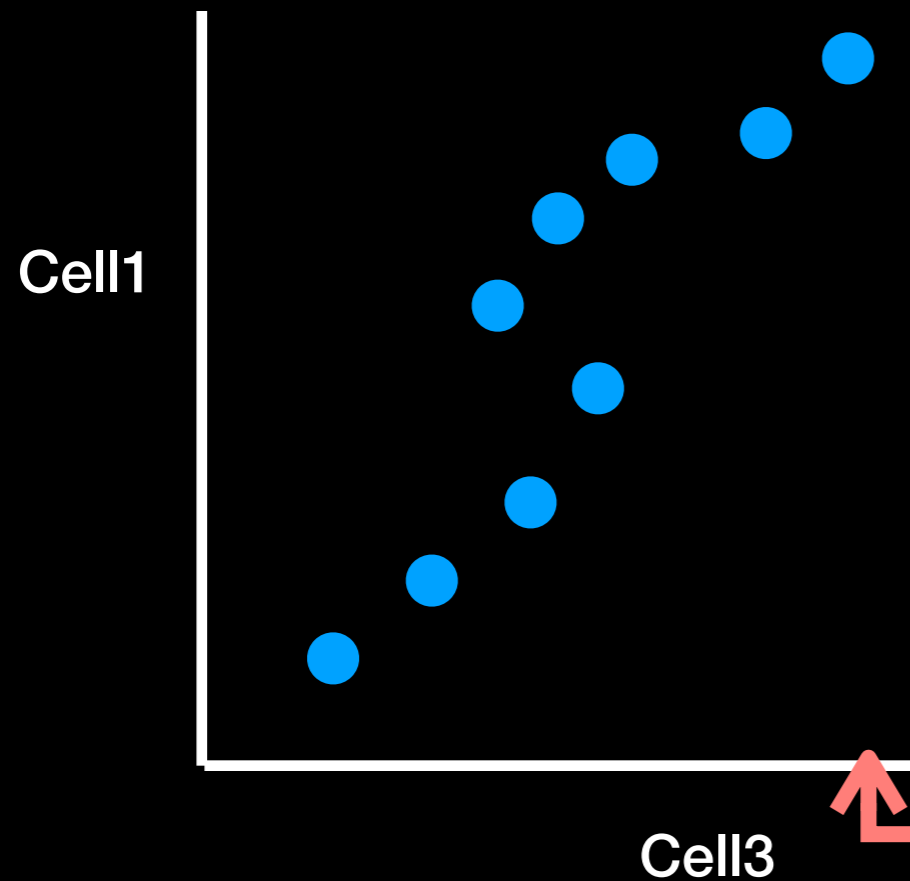
	Cell1	Cell2	Cell3
Gene1	3	0.25	2.8
Gene2	2.9	0.8	2.2
Gene3	2.2	1	1.5
Gene4	2	1.4	2
Gene5	1.3	1.6	1.6
Gene6	1.5	2	2.1
Gene7	1.1	2.2	1.2
Gene8	1	2.7	0.9
Gene9	0.4	3	0.6
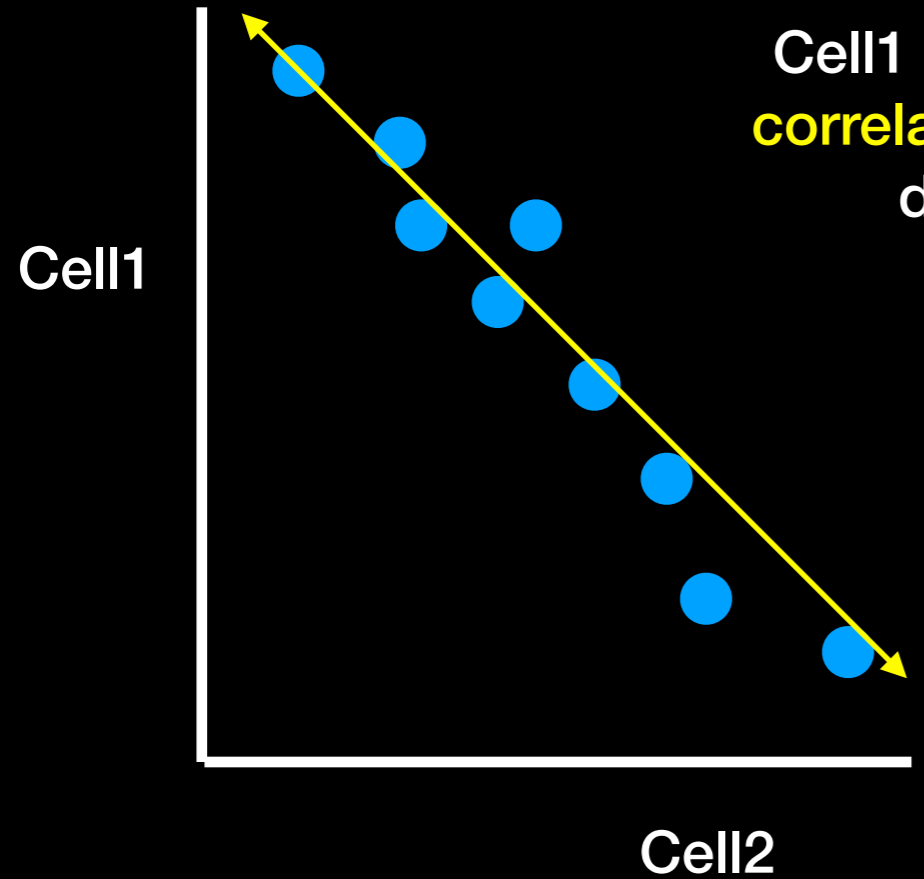
Cell1 and Cell3 are positively correlated suggesting they are doing similar things
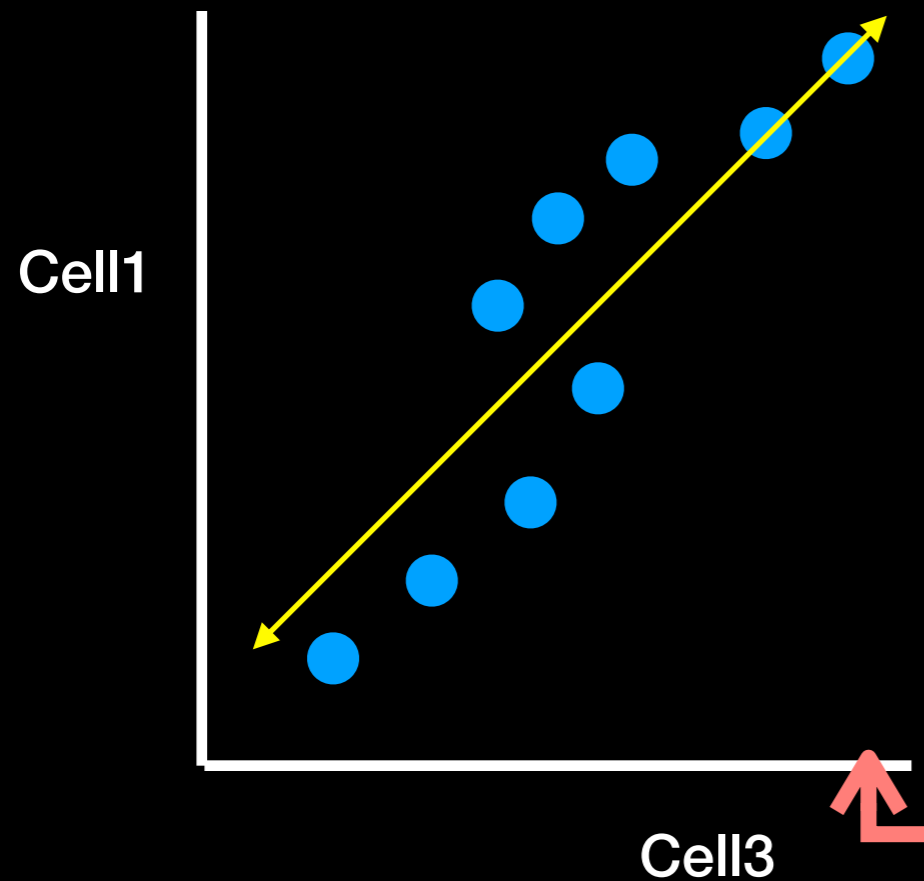
	Cell1	Cell2	Cell3
Gene1	3	0.25	2.8
Gene2	2.9	0.8	2.2
Gene3	2.2	1	1.5
Gene4	2	1.4	2
Gene5	1.3	1.6	1.6
Gene6	1.5	2	2.1
Gene7	1.1	2.2	1.2
Gene8	1	2.7	0.9
Gene9	0.4	3	0.6

Cell1
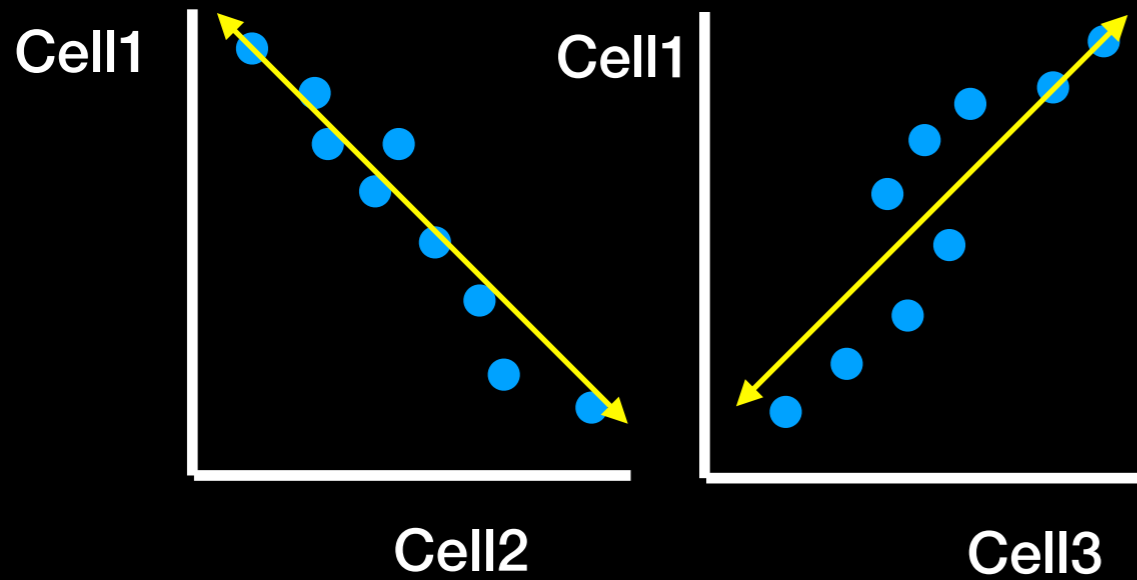
Cell2

Cell1

Cell3

We can also compare Cell2 to Cell3…

The inverse correlation suggests that Cell2 is doing something different from Cell3

	Cell1	Cell2	Cell3
Gene1	3	0.25	2.8
Gene2	2.9	0.8	2.2
Gene3	2.2	1	1.5
Gene4	2	1.4	2
Gene5	1.3	1.6	1.6
Gene6	1.5	2	2.1
Gene7	1.1	2.2	1.2
Gene8	1	2.7	0.9
Gene9	0.4	3	0.6
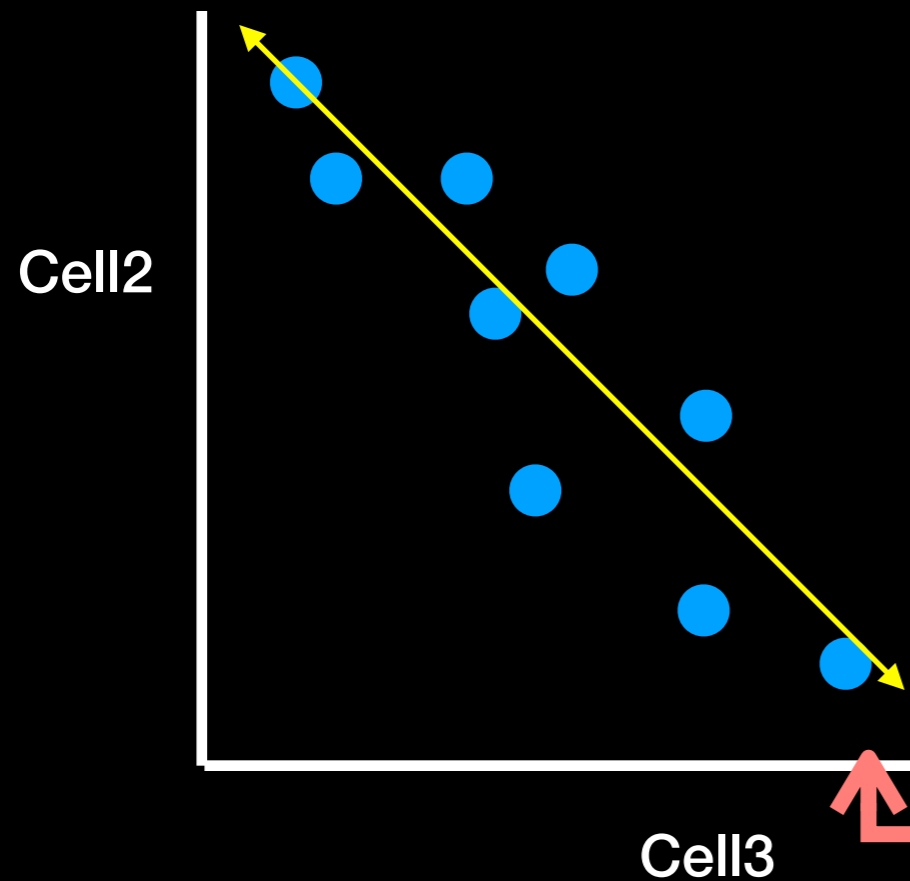
Alternatively, we could try to plot all 3 cells at once on a 3-dimensional graph.
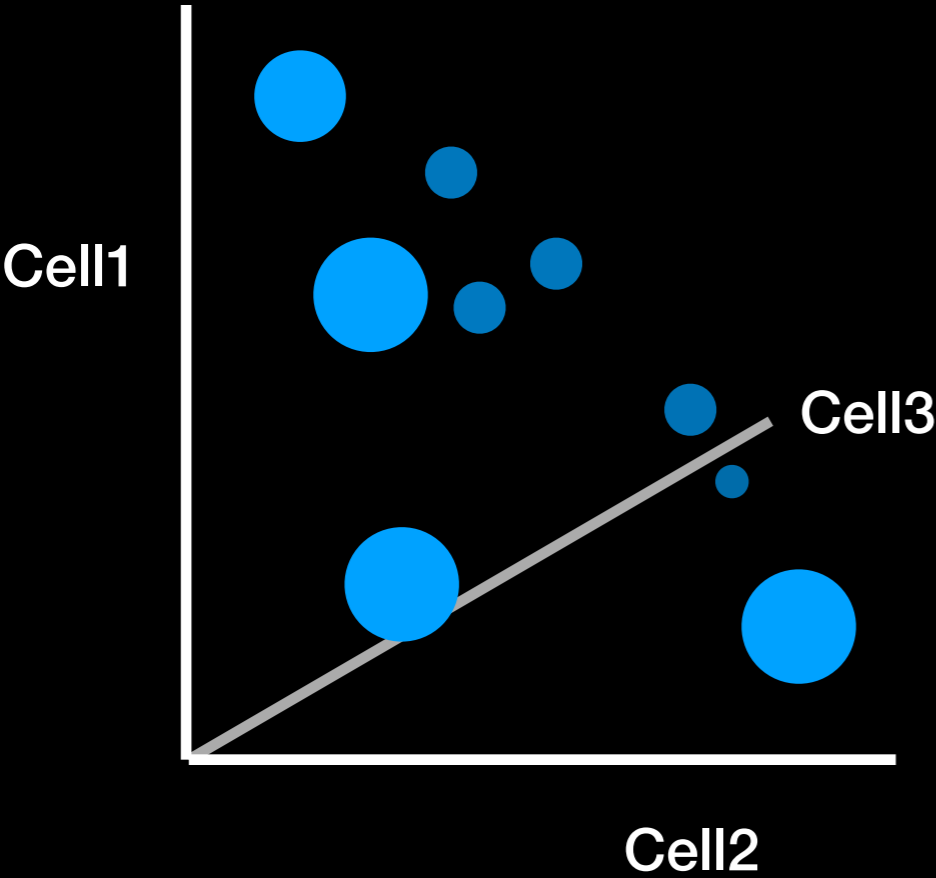
	Cell1	Cell2	Cell3
Gene1	3	0.25	2.8
Gene2	2.9	0.8	2.2
Gene3	2.2	1	1.5
Gene4	2	1.4	2
Gene5	1.3	1.6	1.6
Gene6	1.5	2	2.1
Gene7	1.1	2.2	1.2
Gene8	1	2.7	0.9
Gene9	0.4	3	0.6

But what if we have 4 or more Cells?

	Cell1	Cell2	Cell3	Cell4	...
Gene1	3	0.25	2.8	0.1	...
Gene2	2.9	0.8	2.2	1.8	...
Gene3	2.2	1	1.5	3.2	...
Gene4	2	1.4	2	0.3	...
Gene5	1.3	1.6	1.6	0	...
Gene6	1.5	2	2.1	3	...
Gene7	1.1	2.2	1.2	2.8	...
Gene8	1	2.7	0.9	0.3	...
Gene9	0.4	3	0.6	0.1	...

Draw lots of 2 cell plots and try to make sense of them all?
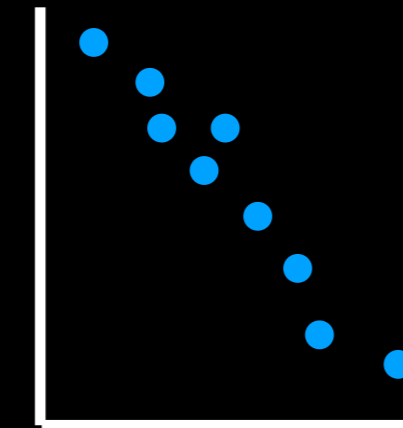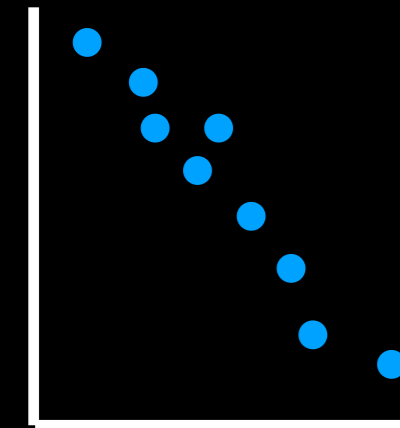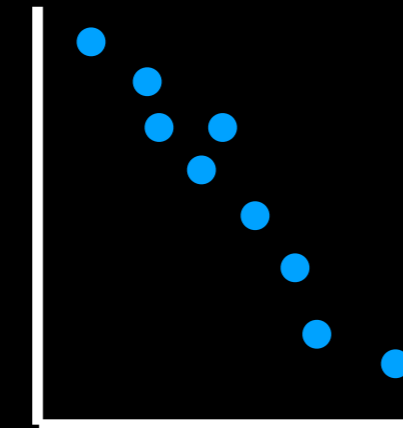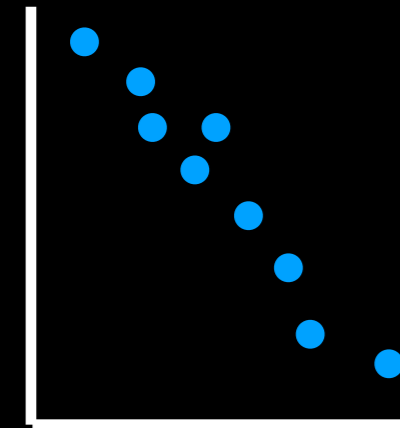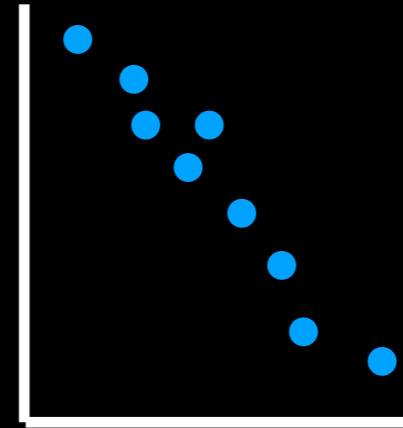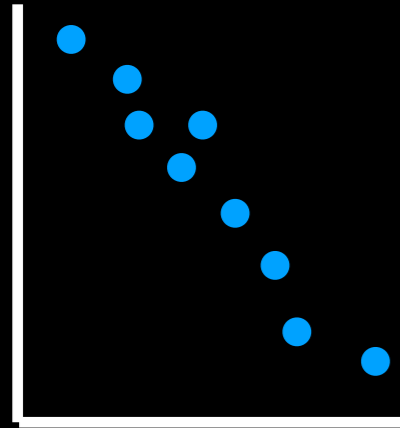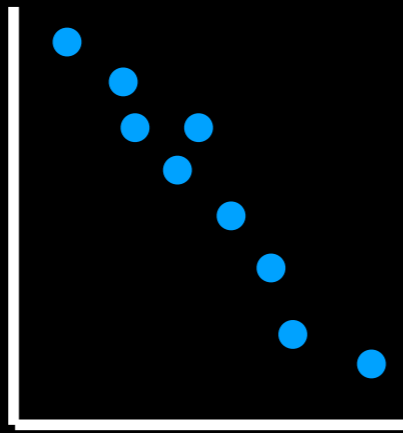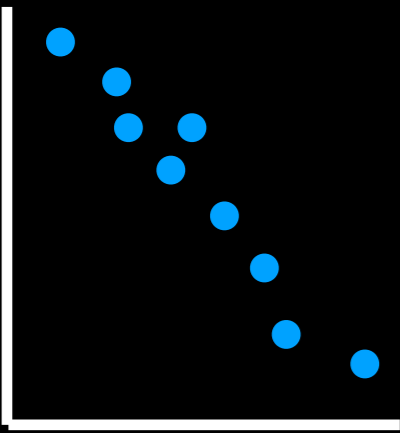
	Cell1	Cell2	Cell3	Cell4	...
Gene1	3	0.25	2.8	0.1	...
Gene2	2.9	0.8	2.2	1.8	...
Gene3	2.2	1	1.5	3.2	...
Gene4	2	1.4	2	0.3	...
Gene5	1.3	1.6	1.6	0	...
Gene6	1.5	2	2.1	3	...
Gene7	1.1	2.2	1.2	2.8	...
Gene8	1	2.7	0.9	0.3	...
Gene9	0.4	3	0.6	0.1	...

Or draw some crazy graph that has an axis for each cell and makes or brains hurt!
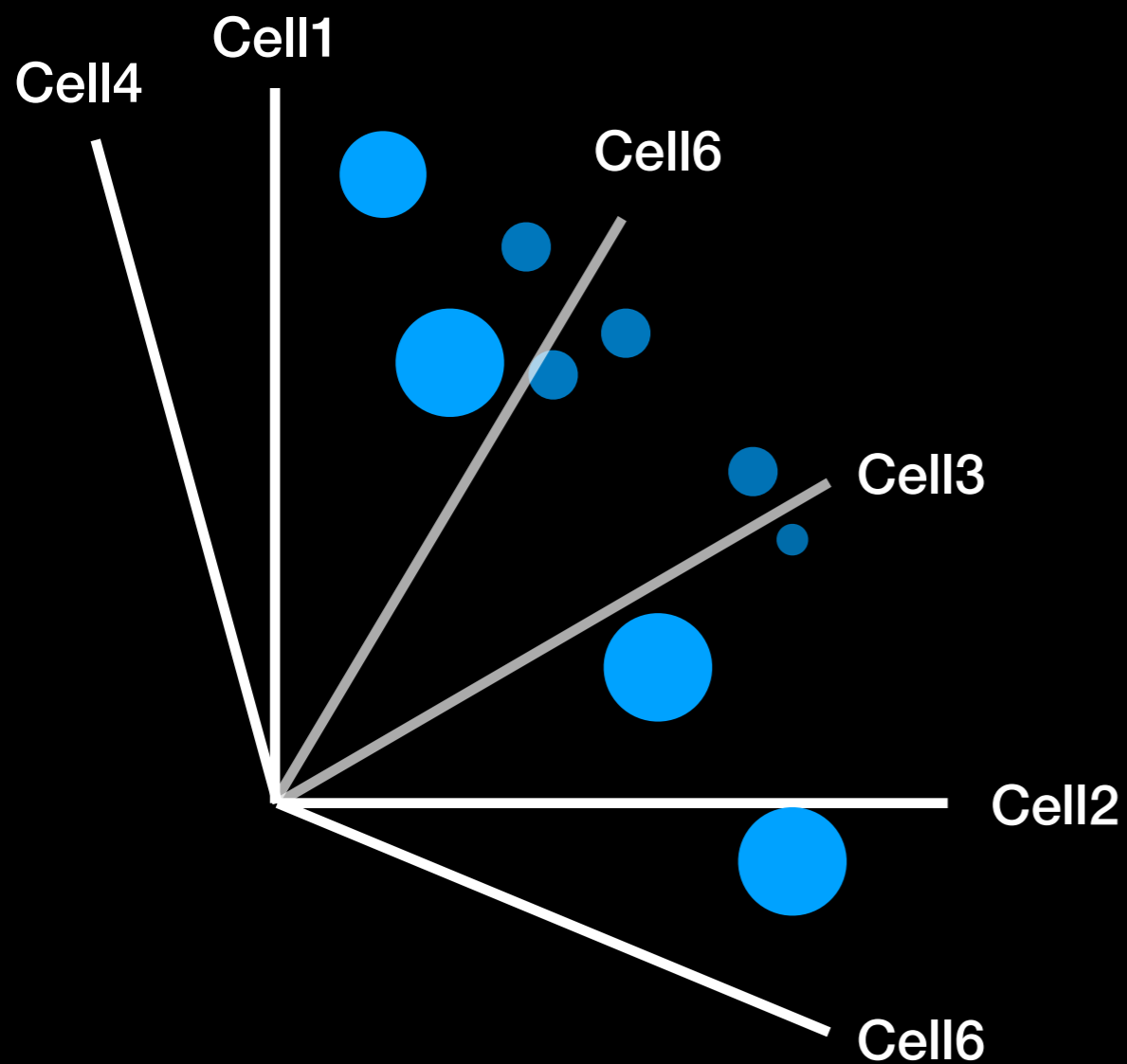
	Cell1	Cell2	Cell3	Cell4	...
Gene1	3	0.25	2.8	0.1	...
Gene2	2.9	0.8	2.2	1.8	...
Gene3	2.2	1	1.5	3.2	...
Gene4	2	1.4	2	0.3	...
Gene5	1.3	1.6	1.6	0	...
Gene6	1.5	2	2.1	3	...
Gene7	1.1	2.2	1.2	2.8	...
Gene8	1	2.7	0.9	0.3	...
Gene9	0.4	3	0.6	0.1	...

	Cell1	Cell2	Cell3	Cell4	...
Gene1	3	0.25	2.8	0.1	...
Gene2	2.9	0.8	2.2	1.8	...
Gene3	2.2	1	1.5	3.2	...
Gene4	2	1.4	2	0.3	...
Gene5	1.3	1.6	1.6	0	...
Gene6	1.5	2	2.1	3	...
Gene7	1.1	2.2	1.2	2.8	...
Gene8	1	2.7	0.9	0.3	...
Gene9	0.4	3	0.6	0.1	...

Cell4

Cell1

Cell6

Cell3

Cell2

Cell6

Enter Principal Component Analysis (PCA)

PC2

PC1
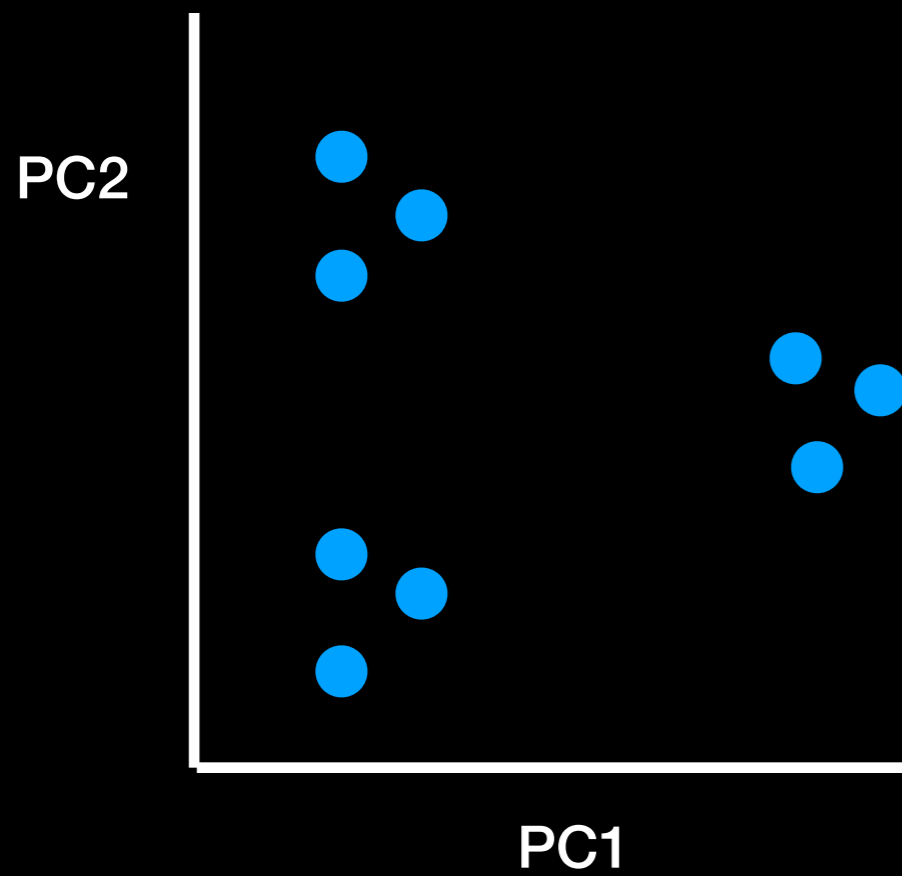
	Cell1	Cell2	Cell3	Cell4	…
Gene1	3	0.25	2.8	0.1	…
Gene2	2.9	0.8	2.2	1.8	…
Gene3	2.2	1	1.5	3.2	…
Gene4	2	1.4	2	0.3	…
Gene5	1.3	1.6	1.6	0	…
Gene6	1.5	2	2.1	3	…
Gene7	1.1	2.2	1.2	2.8	…
Gene8	1	2.7	0.9	0.3	…
Gene9	0.4	3	0.6	0.1	…

PCA converts the correlations (or lack there of) among all cells into a representation we can more readily interpret (e.g. a 2D graph!)
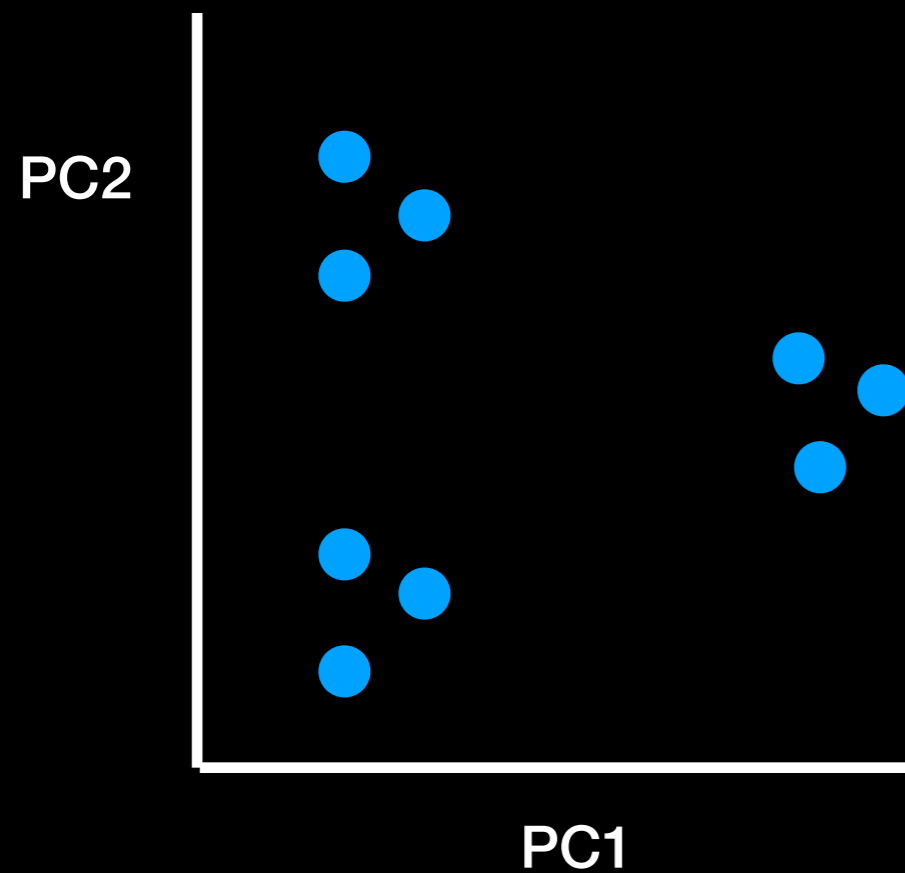
	Cell1	Cell2	Cell3	Cell4	...
Gene1	3	0.25	2.8	0.1	...
Gene2	2.9	0.8	2.2	1.8	...
Gene3	2.2	1	1.5	3.2	...
Gene4	2	1.4	2	0.3	...
Gene5	1.3	1.6	1.6	0	...
Gene6	1.5	2	2.1	3	...
Gene7	1.1	2.2	1.2	2.8	...
Gene8	1	2.7	0.9	0.3	...
Gene9	0.4	3	0.6	0.1	...
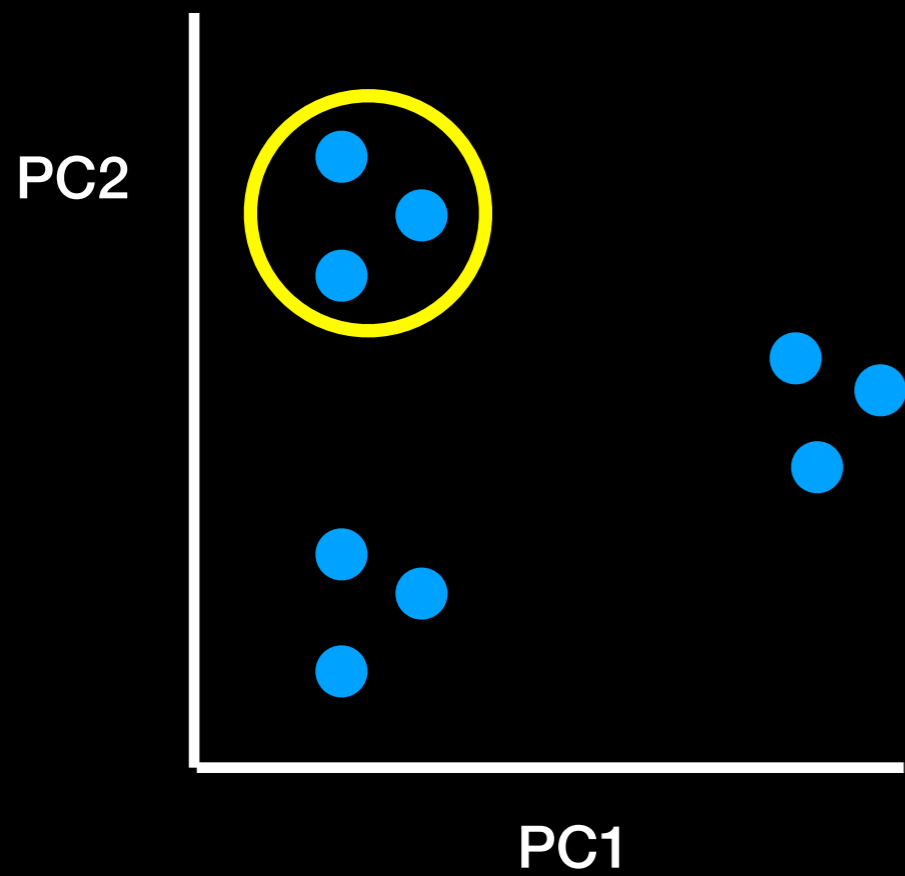
Cells that are highly correlated cluster together

	Cell1	Cell2	Cell3	Cell4	...
Gene1	3	0.25	2.8	0.1	...
Gene2	2.9	0.8	2.2	1.8	...
Gene3	2.2	1	1.5	3.2	...
Gene4	2	1.4	2	0.3	...
Gene5	1.3	1.6	1.6	0	...
Gene6	1.5	2	2.1	3	...
Gene7	1.1	2.2	1.2	2.8	...
Gene8	1	2.7	0.9	0.3	...
Gene9	0.4	3	0.6	0.1	...

Cells that are highly correlated cluster together

PC2

PC1
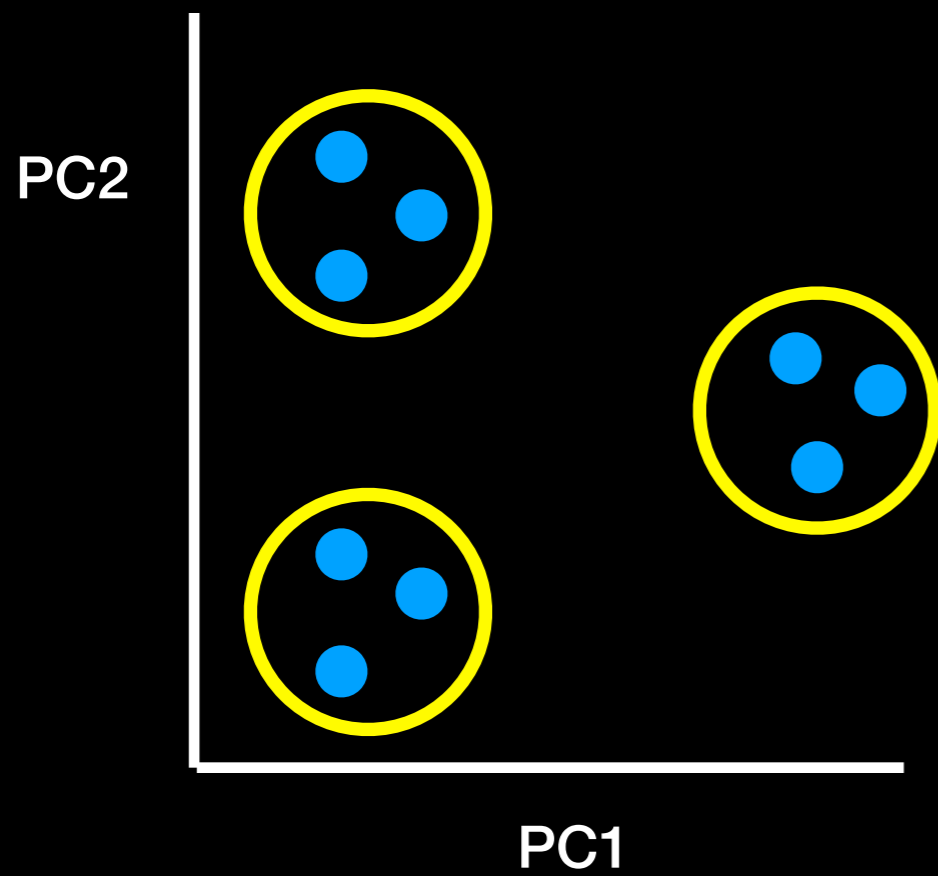
	Cell1	Cell2	Cell3	Cell4	...
Gene1	3	0.25	2.8	0.1	...
Gene2	2.9	0.8	2.2	1.8	...
Gene3	2.2	1	1.5	3.2	...
Gene4	2	1.4	2	0.3	...
Gene5	1.3	1.6	1.6	0	...
Gene6	1.5	2	2.1	3	...
Gene7	1.1	2.2	1.2	2.8	...
Gene8	1	2.7	0.9	0.3	...
Gene9	0.4	3	0.6	0.1	...

To make the clusters easier to see we can color code them…
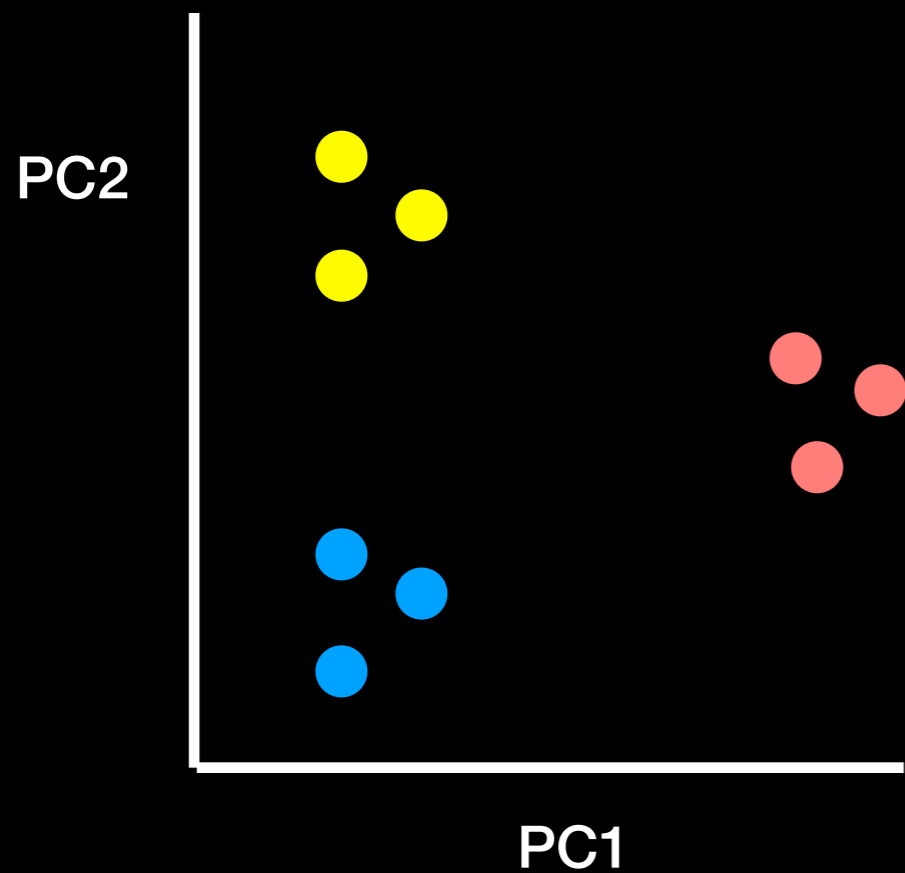
	Cell1	Cell2	Cell3	Cell4	…
Gene1	3	0.25	2.8	0.1	…
Gene2	2.9	0.8	2.2	1.8	…
Gene3	2.2	1	1.5	3.2	…
Gene4	2	1.4	2	0.3	…
Gene5	1.3	1.6	1.6	0	…
Gene6	1.5	2	2.1	3	…
Gene7	1.1	2.2	1.2	2.8	…
Gene8	1	2.7	0.9	0.3	…
Gene9	0.4	3	0.6	0.1	…

Once we have identified the clusters from our PCA results, we can go back to or original cells…
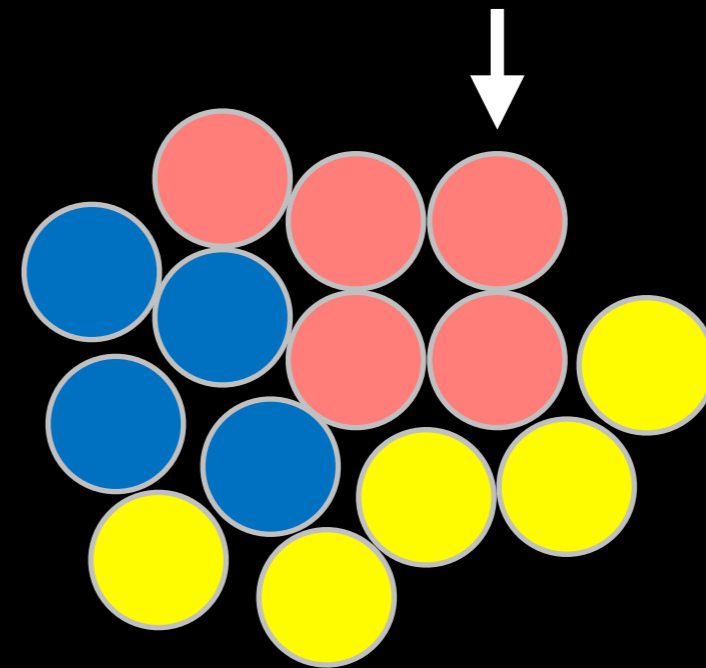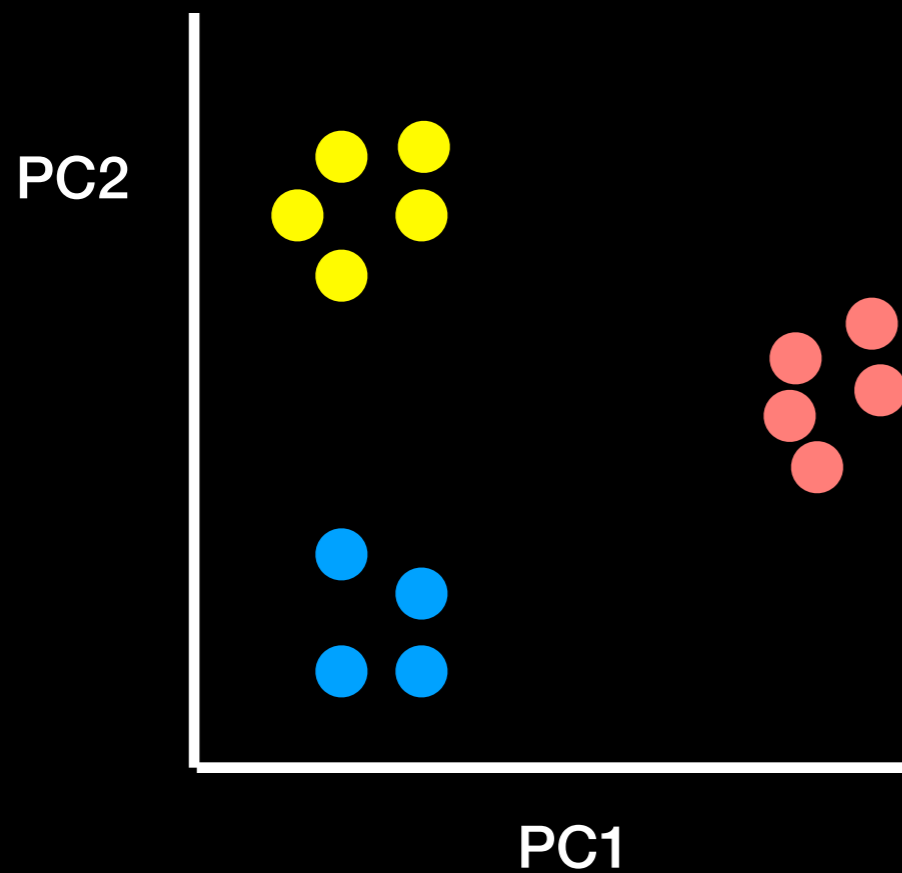
Some key points:

The PCs (i.e. new plot axis) are ranked by their importance

So PC1 is more important than PC2 which in turn is more important than PC3 etc.

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The PCs (i.e. new plot axis) are ranked by their importance

So PC1 is more important than PC2 which in turn is more important than PC3 etc.

So the red and blue cluster are more dissimilar than the yellow and blue clusters

PC2
(11%)

PC1 (44%)
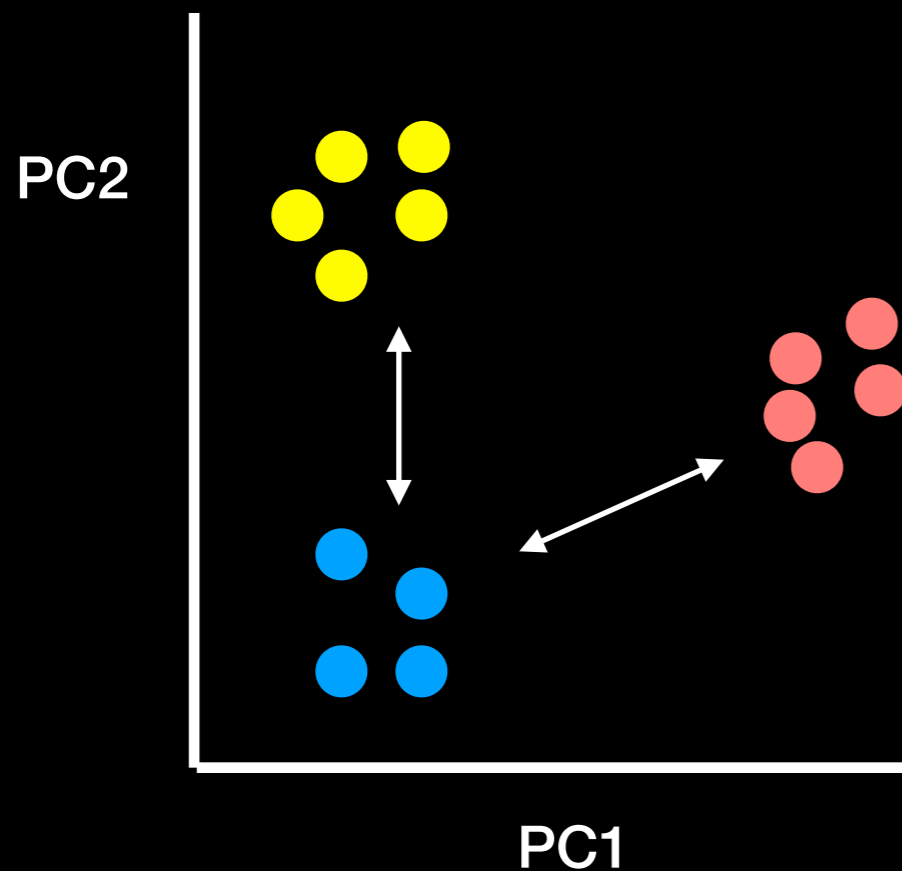
Some key points:

The PCs (i.e. new plot axis) are ranked by their importance

So PC1 is more important than PC2 which in turn is more important than PC3 etc.

So the red and blue cluster are more dissimilar than the yellow and blue clusters

The PCs (i.e. new plot axis) are ranked by the amount of variance in the original data (i.e. gene expression values) that they "capture"

Some key points:

The PCs (i.e. new plot axis) are ranked by their importance

So PC1 is more important than PC2 which in turn is more important than PC3 etc.

So the red and blue cluster are more dissimilar than the yellow and blue clusters

The PCs (i.e. new plot axis) are ranked by the amount of variance in the original data (i.e. gene expression values) that they "capture"

In this example PC1 'captures' 4x more of the original variance than PC2 (44/11 = 4)
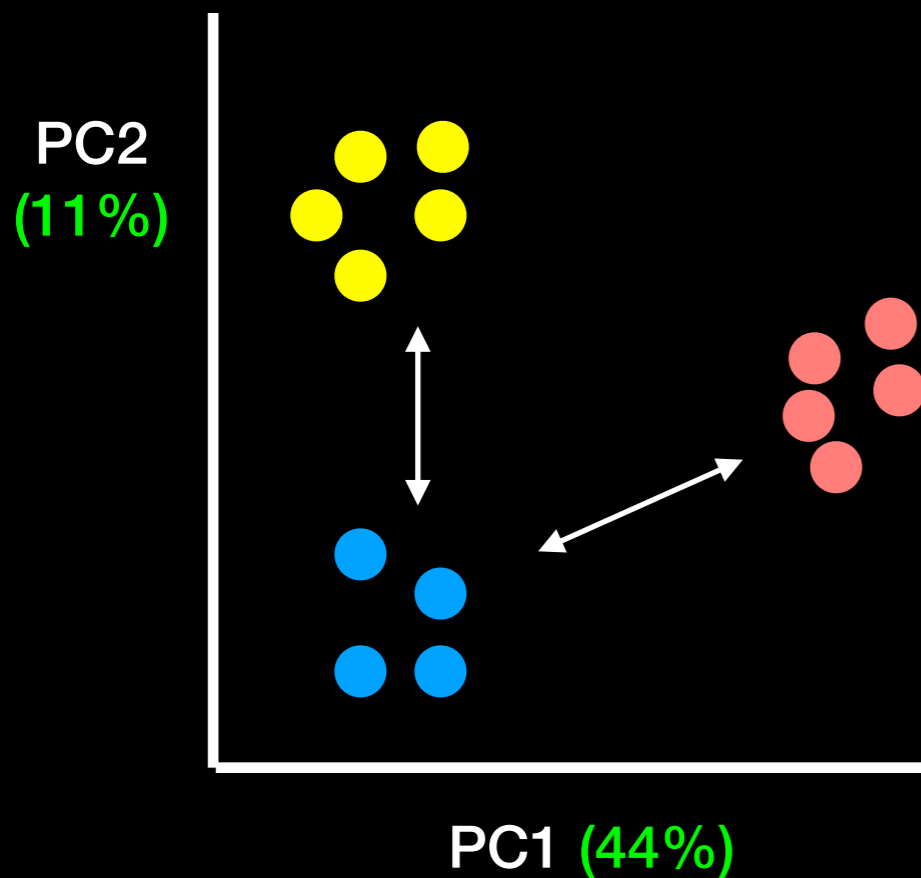
- We actually get two main things out of a typical PCA

  - The new axis (called PCs or Eigenvectors) and

  - Eigenvalues that detail the amount of variance captured by each PC
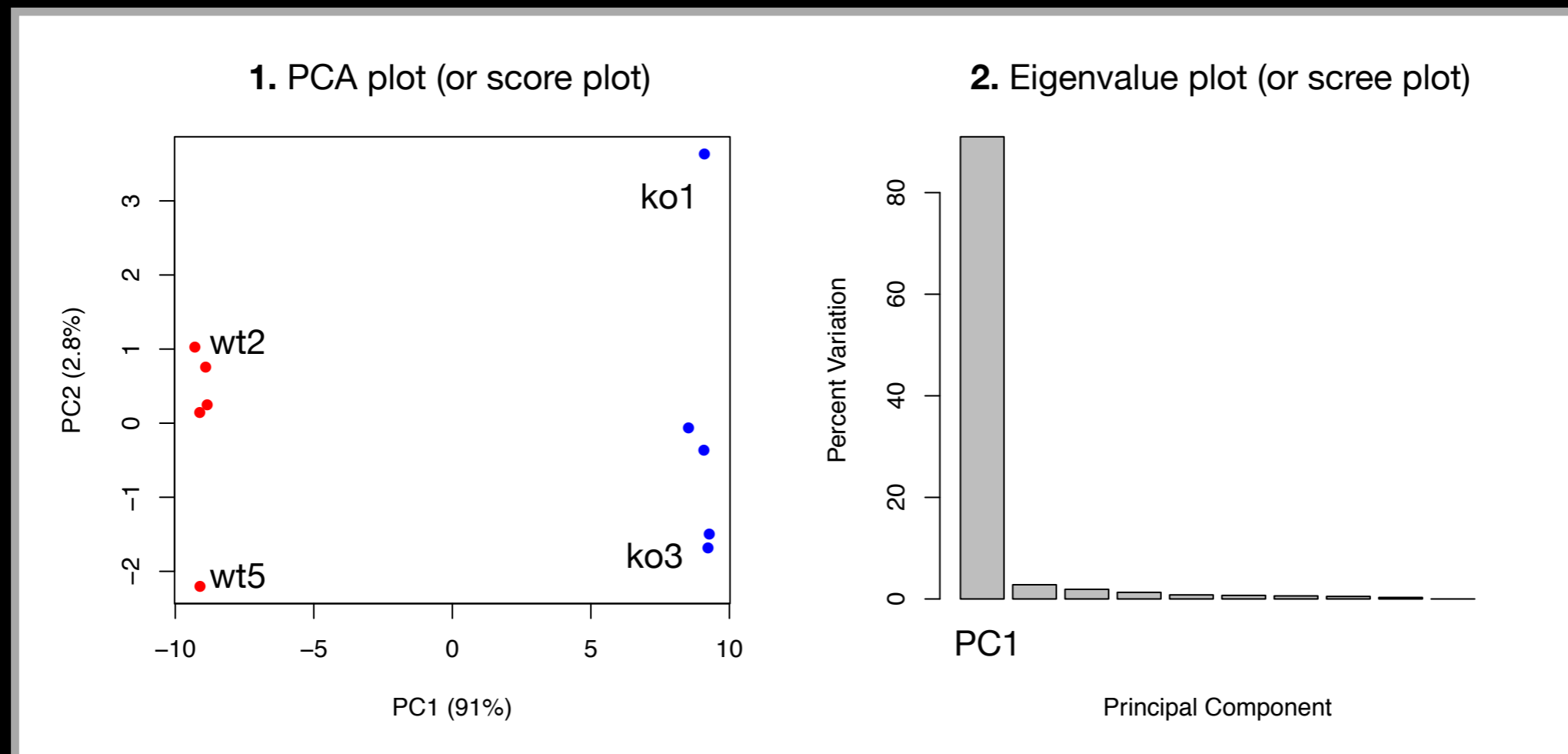
- Another cool thing we can get out of PCA is a quantitive report on how the original variables contributed to each PC

  - In other words, which were the most important genes that lead to the observed clustering in PC-space

  - These are often called the loadings and we can plot them to see which are the most important genes for the observed separation as well as outputting ranked lists of genes that act to discriminate the samples

```
   gene64      gene39
0.1047968   0.1047629
```

```
    gene7      gene65
-0.1047629  -0.1047443
```

Hands-on time!

http://setosa.io/ev/principal-component-analysis/

Outline: How to do PCA in R

- How to use the **prcomp()** function to do PCA.

- How to draw and interpret PCA plots

- How to determine how much variation each principal component accounts for and the the "intrinsic dimensionality" useful for further analysis

- How to examine the **loadings** (or loading scores) to determine what variables have the largest effect on the graph and are thus important for discriminating samples.

- First lets generate some example data to work with.

```
## Initialize a blank 100 row by 10 column matrix
mydata <- matrix(nrow=100, ncol=10)
```

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## Initialize a blank 100 row by 10 column matrix
mydata <- matrix(nrow=100, ncol=10)

## Lets label the rows gene1, gene2 etc. to gene100
rownames(mydata) <- paste("gene", 1:100, sep="")
```

- First lets generate some example data to work with.

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## Initialize a blank 100 row by 10 column matrix
mydata <- matrix(nrow=100, ncol=10)

## Lets label the rows gene1, gene2 etc. to gene100
rownames(mydata) <- paste("gene", 1:100, sep="")

## Lets label the first 5 columns wt1, wt2, wt3, wt4 and wt5
##   and the last 5 ko1, ko2 etc. to ko5 (for "knock-out")
colnames(mydata) <- c( paste("wt", 1:5, sep=""),
                       paste("ko", 1:5, sep="") )
```

- First lets generate some example data to work with.

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## Initialize a blank 100 row by 10 column matrix
mydata <- matrix(nrow=100, ncol=10)

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rownames(mydata) <- paste("gene", 1:100, sep="")

## Lets label the first 5 columns wt1, wt2, wt3, wt4 and wt5
##   and the last 5 ko1, ko2 etc. to ko5 (for "knock-out")
colnames(mydata) <- c( paste("wt", 1:5, sep=""),
                       paste("ko", 1:5, sep="") )

## Fill in some fake read counts
for(i in 1:nrow(mydata)) {
  wt.values <- rpois(5, lambda=sample(x=10:1000, size=1))
  ko.values <- rpois(5, lambda=sample(x=10:1000, size=1))

  mydata[i,] <- c(wt.values, ko.values)
}

head(mydata)
```

- **NOTE**: the samples are columns, and the genes are rows!

```r
mydata <- matrix(nrow=100, ncol=10)

rownames(mydata) <- paste("gene", 1:100, sep="")

colnames(mydata) <- c( paste("wt", 1:5, sep=""),
                       paste("ko", 1:5, sep="") )

for(i in 1:nrow(mydata)) {
  wt.values <- rpois(5, lambda=sample(x=10:1000, size=1))
  ko.values <- rpois(5, lambda=sample(x=10:1000, size=1))
  mydata[i,] <- c(wt.values, ko.values)
}

head(mydata)
```

```
       wt1 wt2 wt3 wt4 wt5 ko1 ko2 ko3 ko4 ko5
gene1  147 171 160 175 187  63  57  58  55  59
gene2  151 134 148 126 134 838 831 894 847 830
gene3  702 672 653 681 701 593 579 644 596 610
gene4  319 297 310 296 304 754 807 734 750 774
```

- Now we have our data we call **prcomp()** to do PCA

  - NOTE: **prcomp()** expects the samples to be rows and genes to be columns so we need to first transpose the matrix with the **t()** function!

```
## lets do PCA
pca <- prcomp(t(mydata), scale=TRUE)
```

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```r
## lets do PCA
pca <- prcomp(t(mydata), scale=TRUE)

## See what is returned by the prcomp() function
attributes(pca)

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

- The returned pca$x here contains the principal components (PCs) for drawing our first graph.

  - Here we will take the first two columns in pca$x (corresponding to PC1 and PC2) to draw a 2-D plot

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## A basic PC1 vs PC2 2-D plot
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- Looks interesting with a nice separation of samples into two groups of 5 samples each

  - Now we can use the square of pca$sdev , which stands for "standard deviation", to calculate how much variation in the original data each PC accounts for

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## A basic PC1 vs PC2 2-D plot
plot(pca$x[,1], pca$x[,2])

## Variance captured per PC
pca.var <- pca$sdev^2
```

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## Precent variance is often more informative to look at
pca.var <- pca$sdev^2
pca.var.per <- round(pca.var/sum(pca.var)*100, 1)
```

- Looks interesting with a nice separation of samples into two groups of 5 samples each

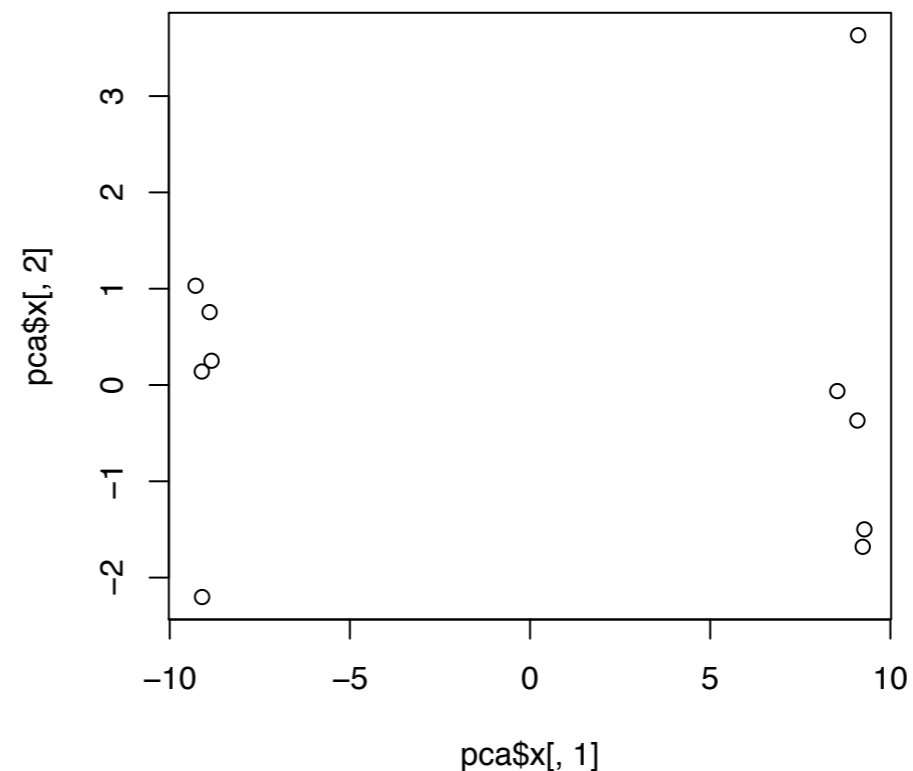 - Now we can use the square of pca$sdev , which stands for "standard deviation", to calculate how much variation in the original data each PC accounts for

```r
## lets do PCA
pca <- prcomp(t(mydata), scale=TRUE)

## A basic PC1 vs PC2 2-D plot
plot(pca$x[,1], pca$x[,2])

## Precent variance is often more informative to look at
pca.var <- pca$sdev^2
pca.var.per <- round(pca.var/sum(pca.var)*100, 1)

pca.var.per

[1] 91.0  2.8  1.9  1.3  0.8  0.7  0.6  0.5  0.3  0.0
```
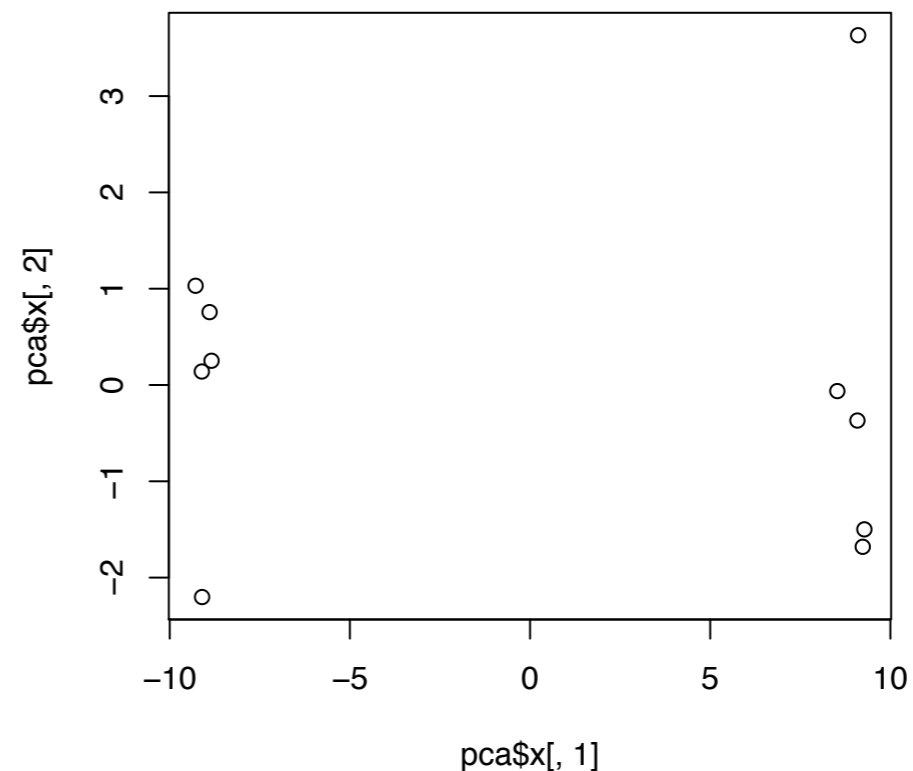
- Looks interesting with a nice separation of samples into two groups of 5 samples each

  - Now we can use the square of pca$sdev , which stands for "standard deviation", to calculate how much variation in the original data each PC accounts for

```
pca.var <- pca$sdev^2
pca.var.per <- round(pca.var/sum(pca.var)*100, 1)

barplot(pca.var.per, main="Scree Plot",
        xlab="Principal Component", ylab="Percent Variation")
```

- From the "scree plot" it is clear that **PC1** accounts for almost all of the variation in the data!

```
pca.var <- pca$sdev^2
pca.var.per <- round(pca.var/sum(pca.var)*100, 1)

barplot(pca.var.per, main="Scree Plot",
        xlab="Principal Component", ylab="Percent Variation")
```
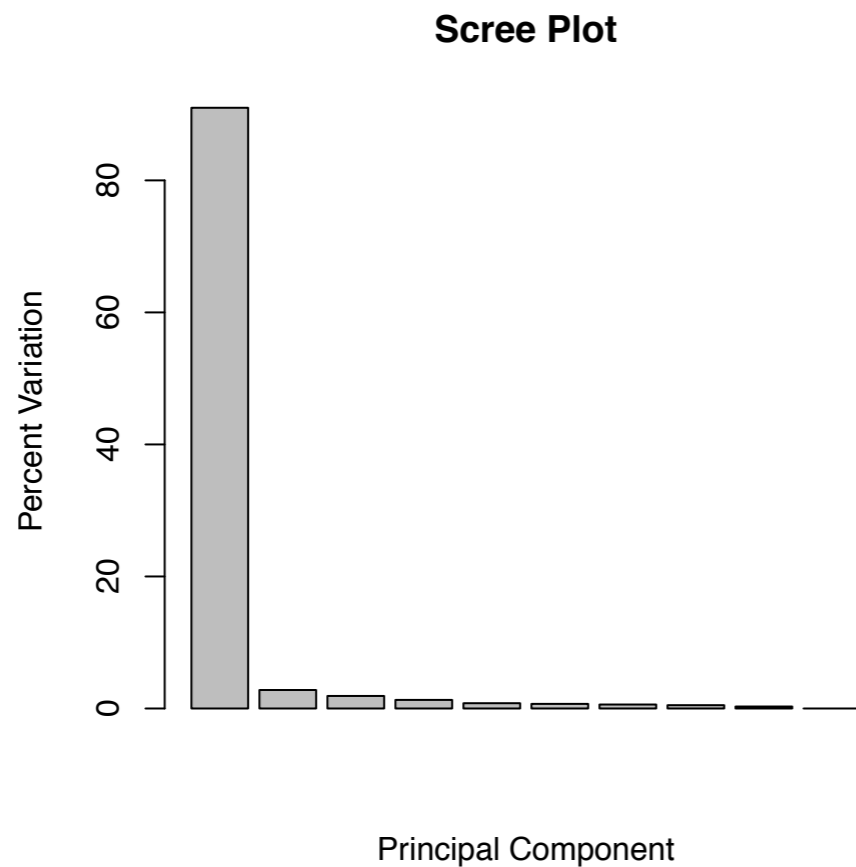

Scree Plot

- Which means there are big differences between these two groups that are separated along the PC1 axis…

```
pca.var <- pca$sdev^2
pca.var.per <- round(pca.var/sum(pca.var)*100, 1)

barplot(pca.var.per, main="Scree Plot",
        xlab="Principal Component", ylab="Percent Variation")
```
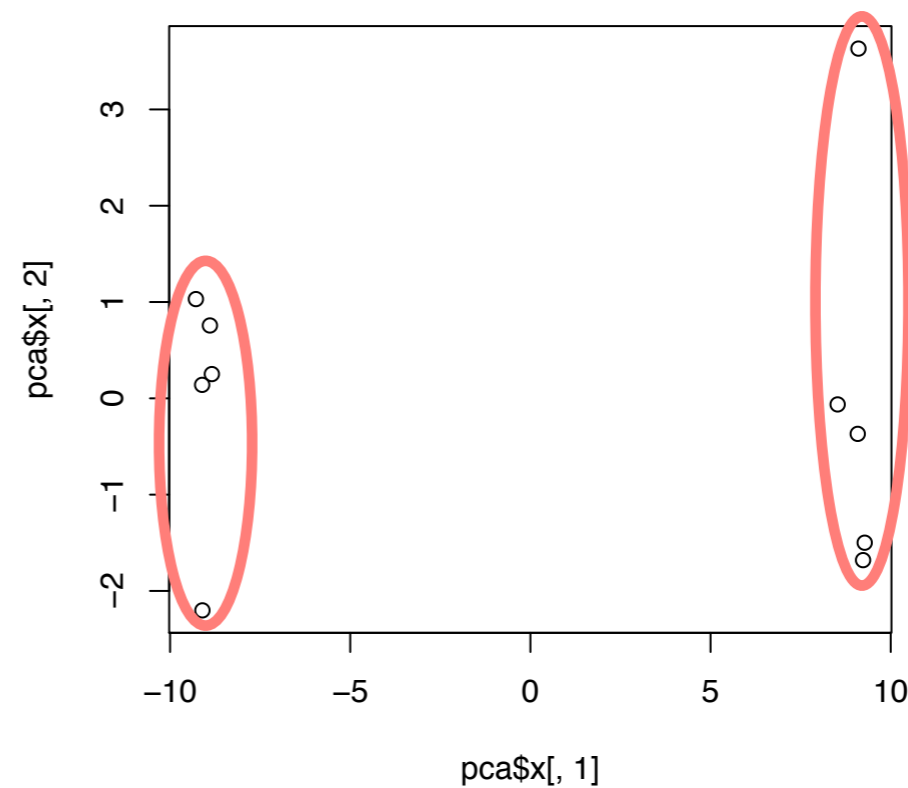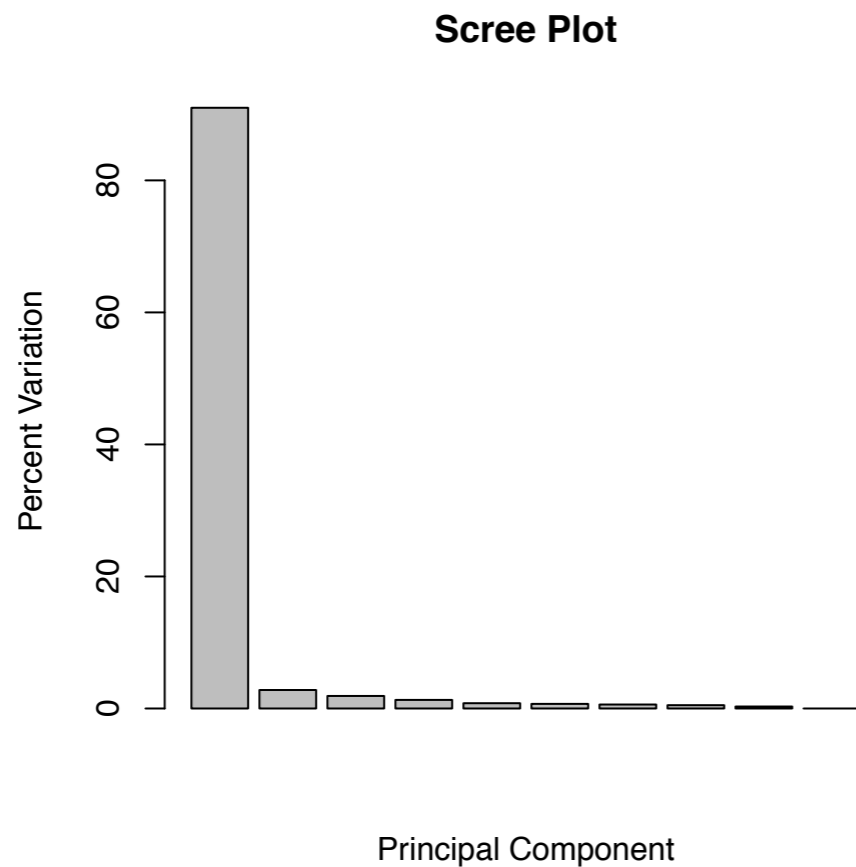
- Lets make our plot a bit more useful…

```
## A vector of colors for wt and ko samples
colvec <- colnames(mydata)
colvec[grep("wt", colvec)] <- "red"
colvec[grep("ko", colvec)] <- "blue"

plot(pca$x[,1], pca$x[,2], col=colvec, pch=16,
     xlab=paste0("PC1 (", pca.var.per[1], "%)"),
     ylab=paste0("PC2 (", pca.var.per[2], "%)"))
```
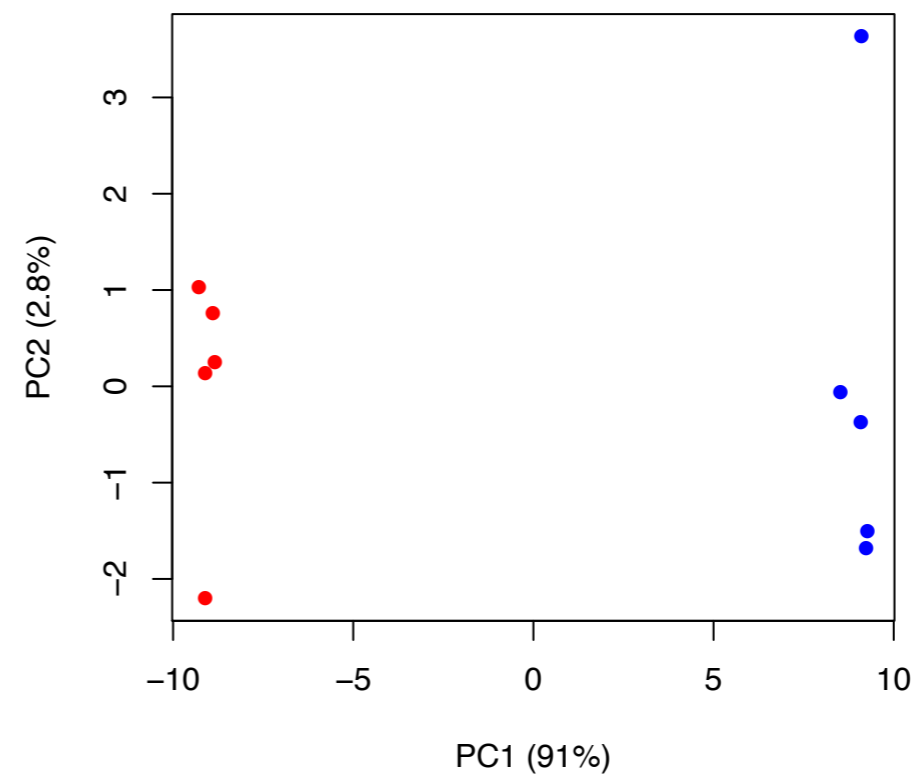
- And add some labels…

```
plot(pca$x[,1], pca$x[,2], col=colvec, pch=16,
     xlab=paste0("PC1 (", pca.var.per[1], "%)"),
     ylab=paste0("PC2 (", pca.var.per[2], "%)"))

## Click to identify which sample is which
identify(pca$x[,1], pca$x[,2], labels=colnames(mydata))

## Press ESC to exit…
```
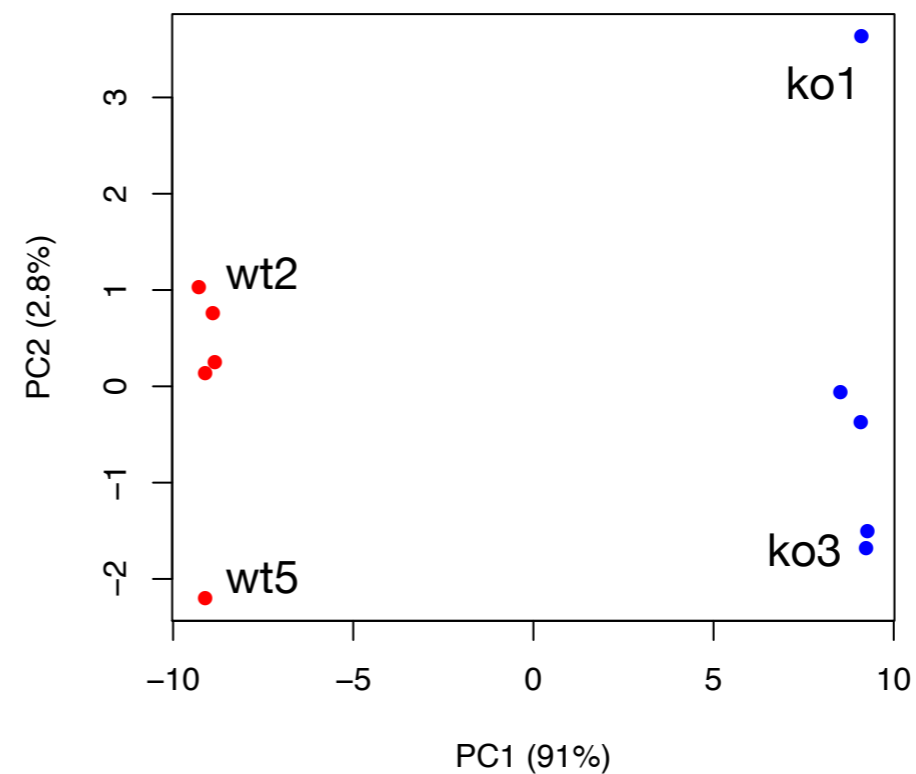
- Finally, lets look at how to use the loading scores to determine which genes have the largest effect on where samples are plotted in the PCA plot

  - The prcomp() function calls loading scores $rotation

```
## Lets focus on PC1 as it accounts for > 90% of variance
loading_scores <- pca$rotation[,1]
```

- Finally, lets look at how to use the loading scores to determine which genes have the largest effect on where samples are plotted in the PCA plot

  - The prcomp() function calls loading scores $rotation

```
## Lets focus on PC1 as it accounts for > 90% of variance
loading_scores <- pca$rotation[,1]

summary(loading_scores)
     Min.   1st Qu.    Median      Mean   3rd Qu.      Max.
-0.104763 -0.104276 -0.068784 -0.005656  0.103926  0.104797

## We are interested in the magnitudes of both plus
##   and minus contributing genes
gene_scores <- abs(loading_scores)
```

- Finally, lets look at how to use the loading scores to determine which genes have the largest effect on where samples are plotted in the PCA plot

  - The prcomp() function calls loading scores $rotation

```
loading_scores <- pca$rotation[,1]

gene_scores <- abs(loading_scores)

## Sort by magnitudes from high to low
gene_score_ranked <- sort(gene_scores, decreasing=TRUE)
```

- Finally, lets look at how to use the loading scores to determine which genes have the largest effect on where samples are plotted in the PCA plot

  - The prcomp() function calls loading scores $rotation

```r
loading_scores <- pca$rotation[,1]

gene_scores <- abs(loading_scores)

## Sort by magnitudes from high to low
gene_score_ranked <- sort(gene_scores, decreasing=TRUE)

## Find the names of the top 5 genes
top_5_genes <- names(gene_score_ranked[1:5])
```

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## Find the names of the top 5 genes
top_5_genes <- names(gene_score_ranked[1:5])

## Show the scores (with +/- sign)
pca$rotation[top_5_genes,1]
```

- Here we see genes with the largest positive loading scores that effectively 'push' the "ko" samples to the right positive side of the plot.

- And the genes with high negative scores that push "wt" samples to the left side of the plot.

```
loading_scores <- pca$rotation[,1]

gene_scores <- abs(loading_scores)

## Sort by magnitudes from high to low
gene_score_ranked <- sort(gene_scores, decreasing=TRUE)

## Find the names of the top 5 genes
top_5_genes <- names(gene_score_ranked[1:5])

## Show the scores (with +/- sign)
pca$rotation[top_5_genes,1]
    gene64       gene39        gene7       gene60       gene65
 0.1047968    0.1047629   -0.1047629    0.1047601   -0.1047443
```
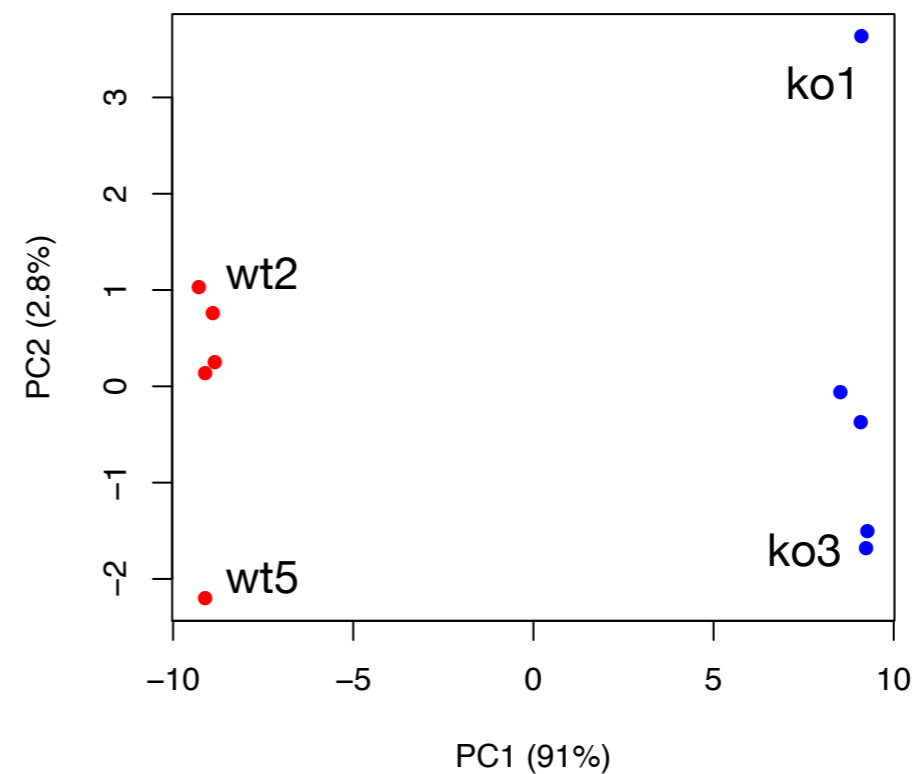
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- And the genes with high negative scores that push "wt" samples to the left side of the plot.

PCA Recap

- PCA is classic "multivariate statistical technique" used to reduce the dimensionality of a complex data set to a more manageable number (typically 2D or 3D)

- For a matrix of m genes x n samples, we mean center (i.e. subtract the sample mean from each sample column), optionally rescale the values for each sample column, then calculate a new covariance matrix of size n x n

- We finally diagonalize the covariance matrix to yield our n Eigenvectors (called principal components or PCs) and n Eigenvalues.

- The top PCs (with largest Eigenvalues) retain the essential features of the original data and represent a useful subspace for further analysis (e.g. visualization, clustering, feature extraction, outlier detection etc…)

PCA objectives in a nutshell

- to reduce dimensionality

- to visualize multidimensional data

- to choose the most useful variables (features)

- to identify groupings of objects (e.g. genes/samples)

- to identify outliers

Your turn!

Perform a PCA on the UK foods dataset

UK_foods.csv

Input: read, View/head,
PCA: prcomp,
Plots: PCA plot
scree plot,
loadings plot.

https://bioboot.github.io/bimm143_S18/class-material/UK_food_pca/

[Muddy Point Feedback Link]

Practical issues with PCA

- Scaling the data

- Missing values:

Scaling

```
> data(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

# Means and standard deviations vary a lot
> round(colMeans(mtcars), 2)
    mpg     cyl    disp      hp    drat      wt    qsec      vs      am    gear    carb
  20.09    6.19  230.72  146.69    3.60    3.22   17.85    0.44    0.41    3.69    2.81
> round(apply(mtcars, 2, sd), 2)
    mpg     cyl    disp      hp    drat      wt    qsec      vs      am    gear    carb
   6.03    1.79  123.94   68.56    0.53    0.98    1.79    0.50    0.50    0.74    1.62
```
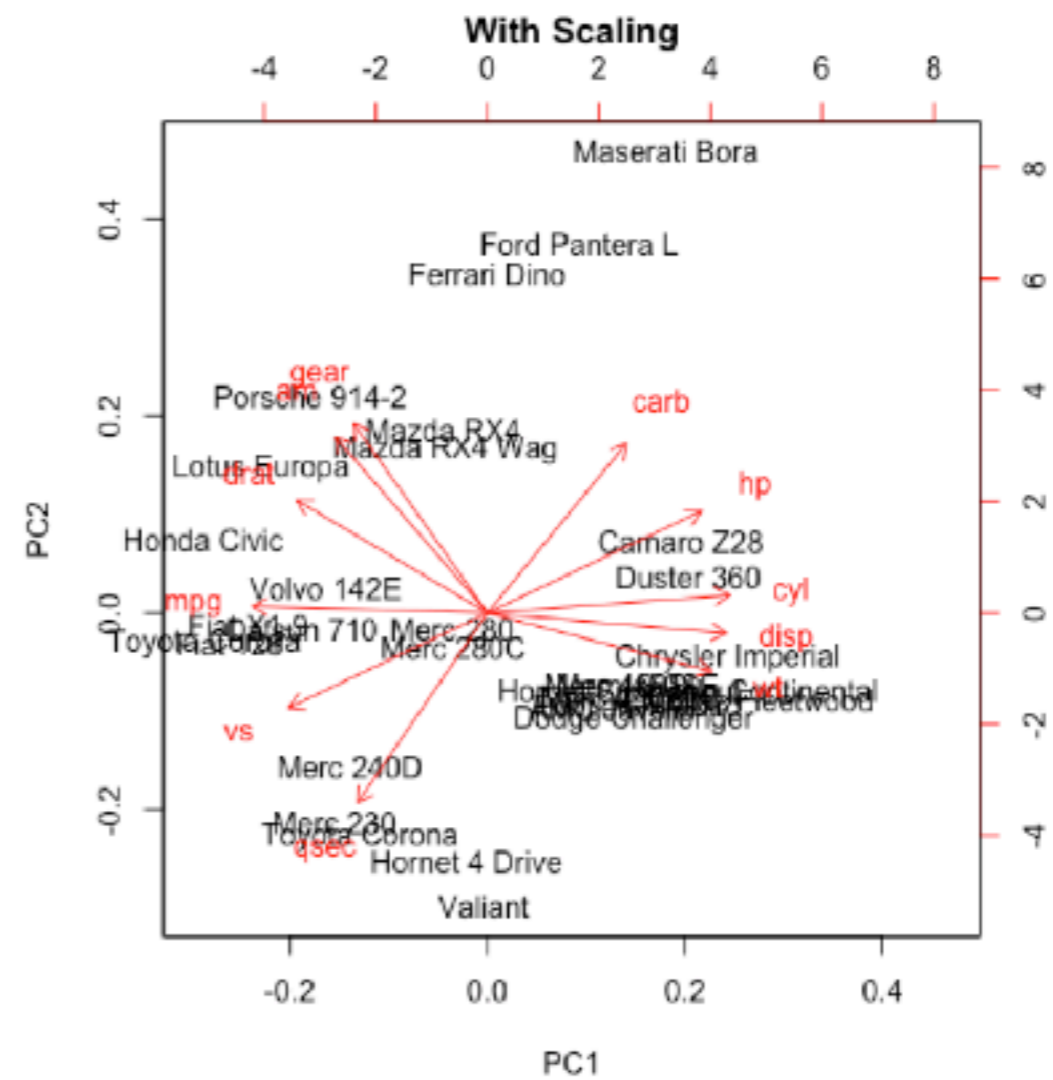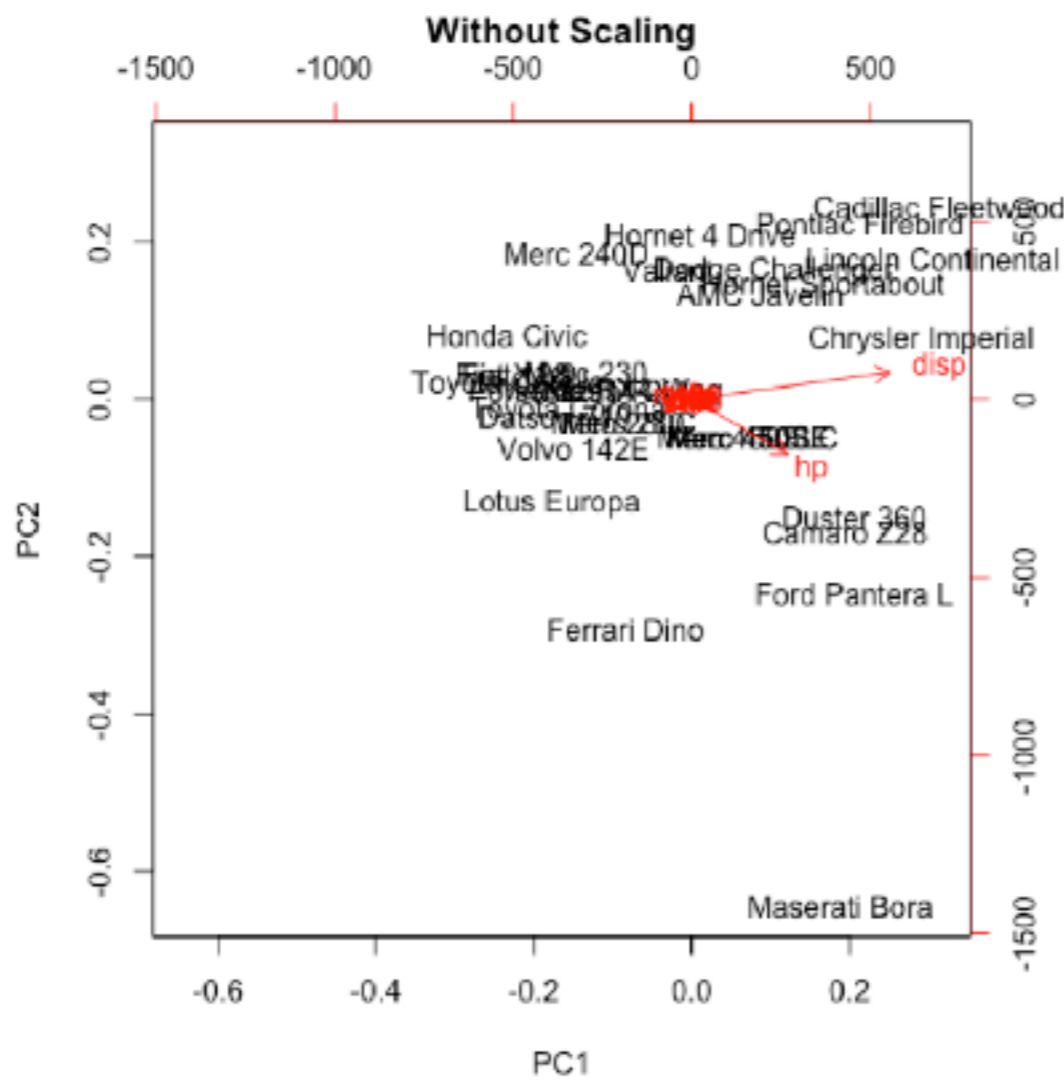
Scaling

Practical issues with PCA

- Scaling the data

- Missing values:
  - ➡ **Drop observations with missing values**
  - ➡ Impute / estimate missing values