# STAT406- Methods of Statistical Learning Lecture 21

Matias Salibian-Barrera

UBC - Sep / Dec 2017

## Unsupervised learning

- Unsupervised ≠ Supervised
- High-"density" regions (w/o model)
- Agglomerative / hierarchical methods
- High-"density" regions (with a model)EM-algorithm
- Dimension reduction (PCA, MDS, etc.)

## Clustering - Problem

Data: p features / variables per "unit"

$$\mathbf{X} = \begin{pmatrix} X_1 \\ X_2 \\ \vdots \\ X_p \end{pmatrix}$$

•  $\mathbf{X}_1, \ldots, \mathbf{X}_n$ 

- Goal: find regions where X<sub>i</sub>'s are "clustered"
- Goal: find regions where P(X) is relatively high
- These regions are sometimes modeled

- Lower dimensional subspaces (linear manifolds)
  - **Principal Components**
- Convex regions with high P(X)
   K-means / K-medoids Hierarchical methods

- Intrinsically different from classification
- There is no clear performance measure to evaluate "success"
- Hence the name: "unsupervised learning"

#### Example 1 - 9 Breweries - 26 attributes

```
> a
   [,1] [,2] [,3] [,4] [,5] [,6] [,7] [,8] [,9] [...]
V1 3.51 3.41 3.20 2.73 2.35 3.03 2.21 3.91 3.07 [...]
V2 4.43 4.05 3.66 5.25 3.88 4.23 3.27 2.71 4.08 [...]
V3 4.76 3.42 4.22 2.44 4.18 2.47 3.67 4.59 4.74 [...]
V4 3.68 3.78 3.07 2.75 2.78 3.12 2.49 3.91 3.34 [...]
V5 4.77 1.04 3.86 5.28 3.86 4.24 3.40 4.23 4.23 [...]
[\ldots]
   [,15] [,16] [,17] [,18] [,19] [,20] [,21] [,22] [...]
V1 3.07 3.45 2.53 3.12 2.93 2.24 2.41 3.32 [...]
V2 3.82 4.29 4.71 3.58 3.27
                                3.11 3.14 3.74 [...]
                                4.12 3.43 4.32 [...]
V3
   4.17 4.44
              4.53 4.10 4.13
V4 3.21 3.74 2.83 3.14 2.80 2.39 2.40 3.32 [...]
V5 3.94 4.47 4.83 3.82 3.46
                                 3.39 3.22 4.01 [...]
[...]
```

$$\mathbf{X}_1, \ \mathbf{X}_2, \ \dots \ \mathbf{X}_9 \in \mathbb{R}^{26}$$

Do they appear grouped / clustered?

#### **UN Votes**

- From http://hdl.handle.net/1902.1/12379
- UN, founded 1946, 193 members
- "important" votes (U.S. State Department)
- Votes: Yes (1), Abstain (2), No (3),
   Absent (8), Not a Member (9)
- 368 important votes, 77 countries voted > 95% of these

#### **UN Votes**

- Do voting patterns reflect political alignments?
- Do countries vote along known political blocks?
- Data: X<sub>i</sub>: votes for country i

$$X_i \in \mathbb{R}^{368}$$
,  $i = 1, ..., 77$  (countries)

What groups are there?

## Cancer example

- From [HTF09], details in script
- Gene expression for 64 samples
- There are 6830 genes
- $X_1, X_2, \ldots, X_{64} \in \mathbb{R}^{6830}$
- We know tissue type for ea. sample
- Really: "feature selection"

- Look for convex sets of relative high density
- The number of sets K is specified a priori (but we'll come back to this)
- Since "high density" is related to "closeness"

$$\min \sum_{r=1}^{\mathbf{K}} \sum_{i,j \in \mathcal{C}_r} d^2 \left( \mathbf{X}_i, \mathbf{X}_j \right)$$

minimize over all partitions  $C_1, \ldots, C_K$ 

#### Note that

$$\sum_{i=1}^{n} \sum_{j=1}^{n} d^{2} \left( \mathbf{X}_{i}, \mathbf{X}_{j} \right) = \sum_{r=1}^{K} \sum_{i \in \mathcal{C}_{r}} \sum_{j=1}^{n} d^{2} \left( \mathbf{X}_{i}, \mathbf{X}_{j} \right)$$

$$= \sum_{r=1}^{K} \sum_{i \in \mathcal{C}_{r}} \left[ \sum_{j \in \mathcal{C}_{r}} d^{2} \left( \mathbf{X}_{i}, \mathbf{X}_{j} \right) + \sum_{j \notin \mathcal{C}_{r}} d^{2} \left( \mathbf{X}_{i}, \mathbf{X}_{j} \right) \right]$$

$$\sum_{r=1}^{K} \sum_{i,j \in \mathcal{C}_{r}} d^{2} \left( \mathbf{X}_{i}, \mathbf{X}_{j} \right) + \sum_{r=1}^{K} \sum_{i \in \mathcal{C}_{r}} \sum_{j \notin \mathcal{C}_{r}} d^{2} \left( \mathbf{X}_{i}, \mathbf{X}_{j} \right)$$

$$T = W + B$$

When 
$$d^{2}(\mathbf{X}_{i}, \mathbf{X}_{j}) = \|\mathbf{X}_{i} - \mathbf{X}_{j}\|^{2}$$

$$W = \sum_{r=1}^{K} \sum_{i,j \in \mathcal{C}_{r}} \|\mathbf{X}_{i} - \mathbf{X}_{j}\|^{2} = \sum_{r=1}^{K} \sum_{i \in \mathcal{C}_{r}} \|\mathbf{X}_{i} - \bar{\mathbf{X}}_{r}\|^{2}$$

• Given  $C_1, C_2, \ldots, C_K$ , assign  $X_i$  to the cluster  $C_i$  with closest mean

$$\mathbf{X}_{i} \leftarrow \arg\min_{1 \leq i \leq \mathcal{K}} \left\| \mathbf{X}_{i} - \bar{\mathbf{X}}_{j} \right\|^{2}$$

Note that

$$ar{\mathbf{X}}_r = \hat{\boldsymbol{\mu}}_r = \arg\min_{\boldsymbol{\mu}} \sum_{i \in \mathcal{C}_r} \|\mathbf{X}_i - \boldsymbol{\mu}\|^2$$

• Given  $\hat{\mu}_1, \ldots, \hat{\mu}_K$ 

$$\min_{\mathcal{C}_1, \dots, \mathcal{C}_k} \sum_{r=1}^k \sum_{i \in \mathcal{C}_r} \|\mathbf{x}_i - \hat{\boldsymbol{\mu}}_r\|^2$$

is attained with

$$\mathbf{X}_i \leftarrow \arg\min_{1 \leq i \leq K} \left\| \mathbf{X}_i - \hat{\boldsymbol{\mu}}_j \right\|^2$$

• And, given  $C_1, \ldots, C_K$ 

$$\min_{\hat{\boldsymbol{\mu}}_1, \dots, \hat{\boldsymbol{\mu}}_K} \sum_{r=1}^k \sum_{i \in \mathcal{C}_r} \|\mathbf{x}_i - \hat{\boldsymbol{\mu}}_r\|^2$$

is attained with

$$\hat{\boldsymbol{\mu}}_r \leftarrow \bar{\mathbf{X}}_r = \frac{1}{n_r} \sum_{i \in \mathcal{C}} \mathbf{x}_i$$

This suggests a simple iterative (and greedy) algorithm.

#### Remarks

- Algorithm is greedy
- Answer depends on the initial configuration
- It needs to be started from many initial configurations

#### Cancer data

```
> set.seed(31)
> nci.km <- kmeans(nci, centers=8)</pre>
> table(nci.km$cluster)
 1 2 3 4 5 6 7 8
8 6 6 14 3 8 4 15
> set.seed(311)
> nci.km <- kmeans(nci, centers=8)</pre>
> table(nci.km$cluster)
 1 2 3 4 5 6 7 8
4 12 6 9 4 8 19 2
```

Need **more** starting points...

```
> set.seed (31)
> nci.km <- kmeans(nci, centers=8, iter.max = 5000,
  nstart=1000)
> table(nci.km$cluster)
 1 2 3 4 5 6 7 8
3 8 5 14 6 15 9 4
> set.seed(311)
> nci.km <- kmeans(nci, centers=8, iter.max = 5000,
  nstart=1000)
> table(nci.km$cluster)
 1 2 3 4 5 6 7 8
4 5 8 9 14 3 15 6
```

#### These clusters are the same

Tissue	$C_1$	$C_2$	<i>C</i> <sub>3</sub>	<i>C</i> <sub>4</sub>	<i>C</i> <sub>5</sub>	<i>C</i> <sub>6</sub>	<i>C</i> <sub>7</sub>	<i>C</i> <sub>8</sub>
LEUKEMIA BREAST RENAL COLON PROSTATE	1	2	5	1 2	1 <b>8</b>	6	2	2
MELANOMA OVARIAN NSCLC OTHER	2	5		<b>5 6</b>	1 1 3 1		7	2
• =	_				_		_	_

(C) Matias Salibian-Barrera, 2017. All rights reserved. Cannot be copied, re-used, or edited.

## **UN Votes example**

Not all countries voted each time

```
> dim(X)
[1] 368 77
> sum( complete.cases(X) )
[1] 145
```

Only use resolutions with full votes

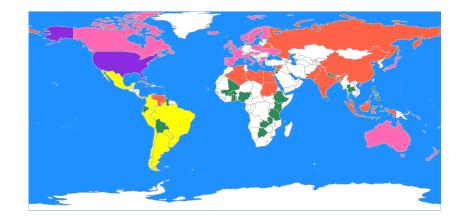
```
X2 <- X[complete.cases(X),]</pre>
```

• Use kmeans in R

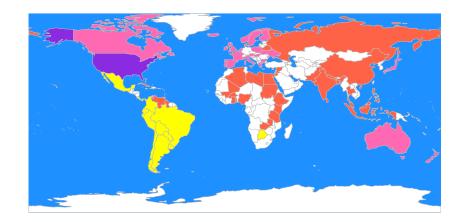
## **UN Votes example**

```
> set.seed(123)
> b <- kmeans(t(X2), centers=5,
                   iter.max=20, nstart=1)
> table(b$cluster)
1 2 3 4 5
18 2 7 19 31
> b <- kmeans(t(X2), centers=5,
                   iter.max=20, nstart=1)
> table(b$cluster)
1 2 3 4 5
27 12 13 7 18
```

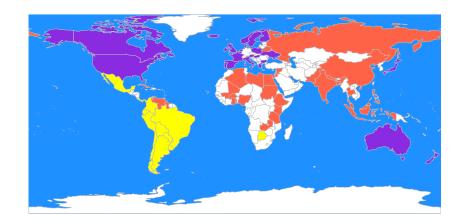
## UN Votes example - K=5



## UN Votes example - K=4



## UN Votes example - K=3



#### K-means++

- A cleverly chosen set of initial centres
- K-means++
  - Pick a centre c<sub>1</sub> at random (from data)
  - Then for j in 2:k
    - Compute weights

$$w_i = \min \left( d^2(\mathbf{x}_i, \mathbf{c}_1), \dots, d^2(\mathbf{x}_i, \mathbf{c}_{j-1}) \right), \quad i = 1, \dots, n$$

- ▶ Pick next centre  $\mathbf{c}_i$  from data with prob  $\propto d_i$
- Implemented in flexclust::kcca

## Choosing *K*

For each cluster  $C_r$ , let

$$W\left(\mathcal{C}_{r}\right) = \sum_{i,j \in \mathcal{C}_{r}} d^{2}\left(\mathbf{X}_{i}, \mathbf{X}_{j}\right) \qquad r = 1, \dots, \mathbf{K}$$

and

$$W_{\mathbf{K}} = \sum_{i=1}^{\mathbf{K}} W(\mathcal{C}_r)$$

## Choosing *K*

- Note that selecting K to minimize W<sub>K</sub> does not generally work
- W<sub>K</sub> typically decreases with K
- A simple example follows

## Selecting the number *K* of clusters

For each cluster  $C_r$ , let

$$W(C_r) = \sum_{i,i \in C_r} d(\mathbf{X}_i, \mathbf{X}_j)$$
  $r = 1, ..., \mathbf{K}$ 

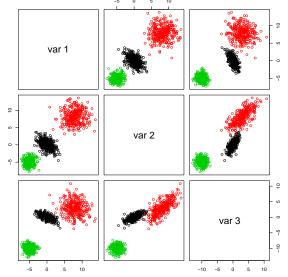
and

$$W_{\mathbf{K}} = \sum_{r=1}^{\mathbf{K}} W(\mathcal{C}_r)$$

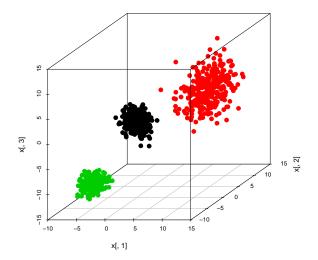
## Selecting the number *K* of clusters

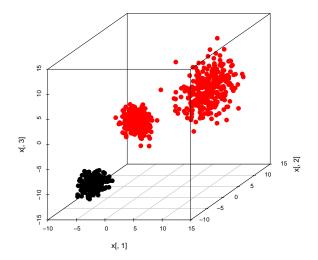
- Note that selecting K to minimize W<sub>K</sub> does not generally work
- W<sub>K</sub> typically decreases with K
- A simple example follows

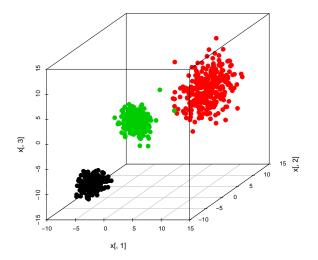
## Pairs plot - Easy case

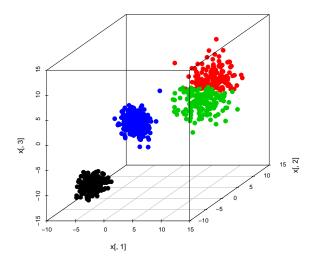


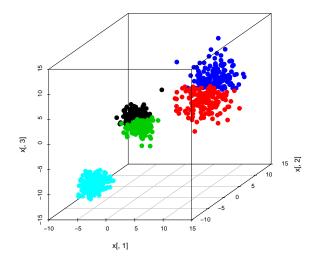
## Pairs plot - Easy case

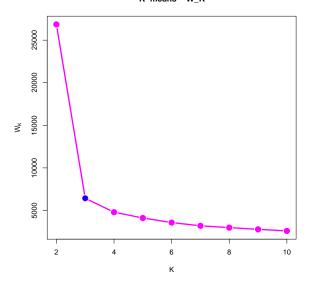




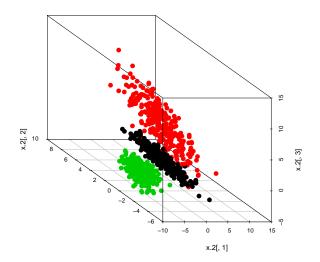


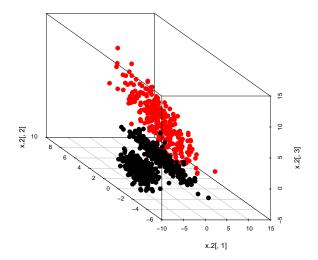


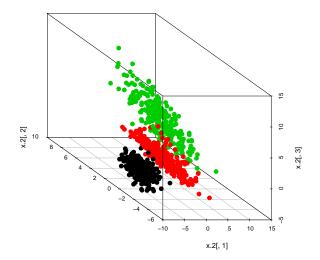


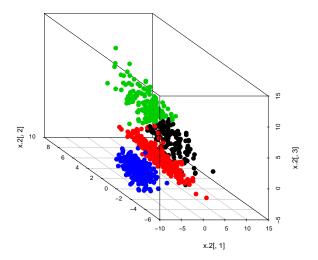


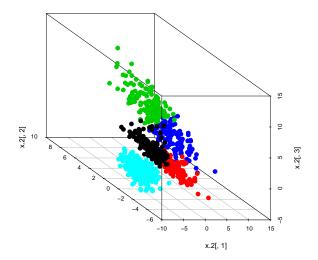
## Pairs plot - K-means

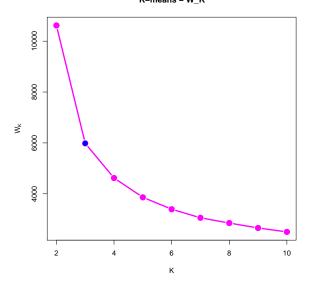












#### **GAP Statistic**

GAP Statistic (Tibshirani, Walther and Hastie, 2001)

Consider

$$G(\mathbf{K}) = E[\log(W_{\mathbf{K}})] - \log(W_{\mathbf{K}})$$

where  $E[\log(W_{\mathbf{K}})]$  is the expected value under a certain reference distribution

## Clest algorithm

#### **Clest algorithm**

Idea - select the value of **K** that produces classes that are best predicted by your favourite classification method.

Dudoit, Fridlyand, 2002, A prediction-based resapmling method for estimating the number of clusters in a dataset, Genome Biology **3(7)**: research0036.1 - 0036.21

## Other approaches to select **K**

Dudoit, Fridlyand, 2003, Bagging to improve the accuracy of a clustering procedure, Bioinformatics, **19**, 1090-1099

#### Note that in K-means

- We used  $d^2(\mathbf{X}_i, \mathbf{X}_j) = \|\mathbf{X}_i \mathbf{X}_j\|^2$
- The cluster "centers" may not be actual observations
- Need to manipulate the "features" (X<sub>i</sub>)
- Can we use different distance measures?
- Can we work with the dissimilarities only?