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### Population substructure

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- Population substructure
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Metric MDS

- Genotype frequencies and allele frequencies vary over human populations.
- If data is a mixture of individuals from different populations, spurious associations may result.
- If the subpopulations are known then
  - a stratified analysis may be more adequate
  - account for population substructure by defining a covariate
- How to detect population substructure?

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### Consequences of population substructure

- Population substructure can influence many types of analysis in statistical genetics.
- It can affect tests for HWE.
- It can affect tests for LD.
- It can affect marker-trait association tests.

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Let there be two populations, and consider one polymorphism. The polymorphism has allele frequency  $p_1 = 0.3$  in the first population, and allele frequency  $p_2 = 0.8$  in the second population. Let there be 300 individuals in each population ( $n_1 = n_2 = 300$ ). We assume Hardy-Weinberg equilibrium within each population. Then

Pop 1	A	В	
Α	$300 \cdot 0.3^2 = 27$	$300 \cdot 0.3 \cdot 0.7 = 63$	90
В	$300 \cdot 0.3 \cdot 0.7 = 63$	$300 \cdot 0.7^2 = 147$	210
	90	210	300

Pop 2	A	В	
Α	$300 \cdot 0.8^2 = 192$	$300 \cdot 0.2 \cdot 0.8 = 48$	240
В	$300 \cdot 0.2 \cdot 0.8 = 48$	$300 \cdot 0.2^2 = 12$	60
	240	60	300

Joint	A	В	
A	27 + 192 = 219	63 + 48 = 111	330
В	63 + 48 = 111	147 + 12 = 159	270
	330	270	600

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```
library(HardyWeinberg)
> x1
[1] 27 126 147
> out1 <- HWChisq(x1,cc=0,verbose=TRUE)
Chi-square test for Hardy-Weinberg equilibrium
Chi2 = 5.962667e-30 p-value = 1 D = 0
> x2
[1] 192 96 12
> out2 <- HWChisq(x2,cc=0,verbose=TRUE)
Chi-square test for Hardy-Weinberg equilibrium
Chi2 = 0 p-value = 1 D = 0
> x3 <- x1+x2
> x4 < - x3/2
> v4
[1] 109.5 111.0 79.5
> out4 <- HWChisq(x4,cc=0,verbose=TRUE)
Chi-square test for Hardy-Weinberg equilibrium
Chi2 = 19.1307 p-value = 1.220655e-05 D = -18.75
```

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Let there be two populations, and consider two polymorphisms, A/a and B/b. In the first population we have  $p_A = 0.7$  and  $p_B = 0.6$ . In the second population we have  $p_A = 0.3$  and  $p_B = 0.9$ . Let there be 100 individuals (200 haplotypes) in each population ( $n_1 = n_2 = 100$ ). We assume linkage equilibrium within each population. Then

Pop 1	В	b	
A	$200 \cdot 0.7 \cdot 0.6 = 84$	$200 \cdot 0.7 \cdot 0.4 = 56$	140
a	$200 \cdot 0.3 \cdot 0.6 = 36$	$200 \cdot 0.3 \cdot 0.4 = 24$	60
	120	80	200

Pop 2	В	b	
A	$200 \cdot 0.3 \cdot 0.9 = 54$	$200 \cdot 0.3 \cdot 0.1 = 6$	60
а	$200 \cdot 0.7 \cdot 0.9 = 126$	$200 \cdot 0.7 \cdot 0.1 = 14$	140
	180	20	200

Joint	В	b	
Α	84 + 54 = 138	56 + 6 = 62	200
a	36 + 126 = 162	24 + 14 = 38	200
	300	100	400

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```
> X1
     [,1] [,2]
[1,]
      84
           56
[2.]
      36 24
> out <- chisq.test(X1,correct=FALSE)
> print(out)
Pearson's Chi-squared test
data: X1
X-squared = 0, df = 1, p-value = 1
> X2
     [,1] [,2]
[1.] 54 6
[2,] 126 14
> out <- chisq.test(X2,correct=FALSE)
> print(out)
Pearson's Chi-squared test
data: X2
X-squared = 0, df = 1, p-value = 1
```

### Chi-square tests

```
> X4 <- (X1+X2)/2
> X4
    [.1] [.2]
[1,] 69 31
[2,] 81 19
> out <- chisq.test(X4,correct=FALSE)
> print(out)
Pearson's Chi-squared test
data: X4
X-squared = 3.84, df = 1, p-value = 0.05004
```

#### How to detect substructure

- Principal component analysis of the marker data
- Multidimensional scaling of distance matrix computed from the marker data
- ..
- In the remainder of this module we focus on MDS.

### Multidimensional scaling

Population substructure

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#### Objective

On the basis of information regarding the distances (or similarities) of n objects, construct a configuration of n points in a low-dimensional space (a **map**).

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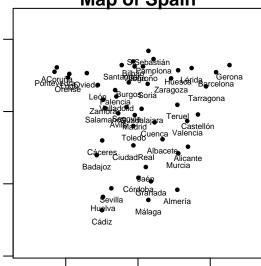
Population substructure

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	Albacete	Alicante	Almer	Avila	Badajoz	Barcelona	Bilbao	Burgos	
Albacete	0	171	369	366	525	540	646	488	
Alicante	171	0	294	537	696	515	817	659	
Almer	369	294	0	663	604	809	958	800	
Avila	366	537	663	0	318	717	401	243	
Badajoz	525	696	604	318	0	1022	694	536	
Barcelona	540	515	809	717	1022	0	620	583	
Bilbao	646	817	958	401	694	620	0	158	
Burgos	488	659	800	243	536	583	158	0	
	•								
:		:	:	:	:	:	:	:	

Download SpainDist.dat

## Map of Spain



Population substructure

### Some basic terminology

#### Terminology

Population substructure

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- proximity
- similarity (s<sub>rs</sub>)
- dissimilarity or distance (d<sub>rs</sub>)

A similarity measure satisfies:

- s(A, B) = s(B, A)
- s(A, B) > 0
- s(A, B) increases as the similarity between A and B increases

A distance measure,  $\delta(A, B)$  satisfies:

- $\bullet$   $\delta(A,B) = \delta(B,A)$
- $\bullet$   $\delta(A,B)>0$
- $\delta(A,A) = 0$

The distance function  $\delta(A, B)$  called a **metric** if also

- $\bullet$   $\delta(A,B)=0$  iff A=B
- the triangle inequality holds:  $\delta(A, B) \leq \delta(A, C) + \delta(C, B)$ .

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### Some dissimilarity measures (quantitative data)

Euclidean distance:

$$\delta_{rs} = \sqrt{(\mathbf{x}_r - \mathbf{x}_s)'(\mathbf{x}_r - \mathbf{x}_s)} = \left\{ \sum_{i=1}^p (x_{ri} - x_{si})^2 \right\}^{\frac{1}{2}}$$

Mahalanobis distance:

$$\delta_{rs} = \left\{ (\mathbf{x}_r - \mathbf{x}_s)' \mathbf{S}^{-1} (\mathbf{x}_r - \mathbf{x}_s) \right\}^{\frac{1}{2}}$$

Minkowski distance

$$\delta_{rs} = \left\{ \sum_{i=1}^{p} |x_{ri} - x_{si}|^{\lambda} \right\}^{\frac{1}{\lambda}}$$

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- In metric MDS, the configuration of points is directly obtained from the distances.
- In non-metric MDS, only the rank order of the distances is important.
- $d_{rs} \approx \delta_{rs}$ : Classical scaling.
- $d_{rs} \approx f(\delta_{rs})$  with  $f(\delta_{rs}) = \alpha + \beta \delta_{rs}$ : Metric scaling.
- $d_{rs} \approx f(\delta_{rs})$  with  $f(\delta_{rs})$  arbitrary, monotone: Non-metric scaling.

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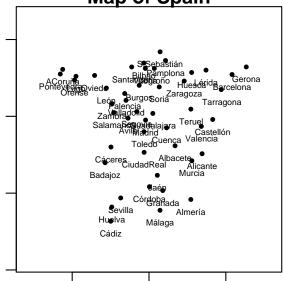
Population substructure

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- Also known as: classical scaling, principal coordinate analysis (PCO).
- Given n objects with dissimiliarities  $(\delta_{rs})$  find a set of points in Euclidean space such that  $d_{rs} \approx \delta_{rs}$ .
- Classical application: given a distance matrix (in km or in travel time) between cities, contruct a map of the cities.

	Albacete	Alicante	Almer	Avila	Badajoz	Barcelona	Bilbao	Burgos	
Albacete	0	171	369	366	525	540	646	488	
Alicante	171	0	294	537	696	515	817	659	
Almer	369	294	0	663	604	809	958	800	
Avila	366	537	663	0	318	717	401	243	
Badajoz	525	696	604	318	0	1022	694	536	
Barcelona	540	515	809	717	1022	0	620	583	
Bilbao	646	817	958	401	694	620	0	158	
Burgos	488	659	800	243	536	583	158	0	
:			:	:	:			1	

Map of Spain



## Theory (1)

Population substructure

Let X be the matrix of coordinates with the solution.  $\mathbf{x}_r, \mathbf{x}_s$  two rows of  $\mathbf{X}$ .

$$d_{rs}^2 = (\mathbf{x}_r - \mathbf{x}_s)'(\mathbf{x}_r - \mathbf{x}_s)$$

Let **B** be the inner product matrix with

$$b_{rs} = \mathbf{x}_r' \mathbf{x}_s$$

Assume the solution to be centered at the origin:

$$\sum_{r=1}^{n} x_{ri} = 0$$

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$$d_{rs}^2 = \mathbf{x}_r' \mathbf{x}_r + \mathbf{x}_s' \mathbf{x}_s - 2\mathbf{x}_r' \mathbf{x}_s$$

$$\frac{1}{n}\sum_{r=1}^{n}d_{rs}^{2}=\frac{1}{n}\sum_{r=1}^{n}\mathbf{x}_{r}'\mathbf{x}_{r}+\mathbf{x}_{s}'\mathbf{x}_{s}$$

$$\frac{1}{n}\sum_{s=1}^n d_{rs}^2 = \mathbf{x}_r'\mathbf{x}_r + \frac{1}{n}\sum_{s=1}^n \mathbf{x}_s'\mathbf{x}_s$$

$$\frac{1}{n^2} \sum_{r=1}^{n} \sum_{s=1}^{n} d_{rs}^2 = \frac{2}{n} \sum_{r=1}^{n} \mathbf{x}_r' \mathbf{x}_r$$

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Let 
$$b_{rs} = \mathbf{x}_{r}' \mathbf{x}_{s} = -\frac{1}{2} (d_{rs}^{2} - \mathbf{x}_{r}' \mathbf{x}_{r} - \mathbf{x}_{s}' \mathbf{x}_{s})$$

$$b_{rs} = -\frac{1}{2} \left( d_{rs}^2 - \frac{1}{n} \sum_{s=1}^n d_{rs}^2 - \frac{1}{n} \sum_{r=1}^n d_{rs}^2 + \frac{1}{n^2} \sum_{r=1}^n \sum_{s=1}^n d_{rs}^2 \right).$$

We define  $a_{rs} = -\frac{1}{2}d_{rs}^2$  so that  $b_{rs} = a_{rs} - a_{r.} - a_{.s} + a_{.s}$  and build matrix **A** 

$$B = HAH \quad H = I - \frac{1}{n}11'$$

and

$$\mathbf{B} = \mathbf{X}\mathbf{X}'$$

We wish to approximate **B** in a low dimensional space.

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## Theory (4) Spectral Decomposition

Let **B** be any  $k \times k$  symmetric matrix we want to approximate

$$\mathbf{B} = \mathbf{V}\mathbf{D}_{\lambda}\mathbf{V}' = \sum_{i=1}^{k} \lambda_{i}\mathbf{v}_{i}\mathbf{v}_{i}'$$

with  $\mathbf{D}_{\lambda} = \operatorname{diag}(\lambda_1, \dots, \lambda_k)$  and  $\mathbf{V} = [\mathbf{v}_i, \dots, \mathbf{v}_k]$ 

$$ilde{\mathbf{B}} = \mathbf{V}_{(,1:k)} \mathbf{D}_{\lambda(1:k,1:k)} \mathbf{V}_{(,1:k)}'$$

gives the rank k least squares approximation to  $\mathbf{B}$ 

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$$\mathbf{B} = \mathbf{X}\mathbf{X}' = \mathbf{V}\mathbf{D}_{\lambda}\mathbf{V}'$$

The coordinates of the solution are obtained as:

$$\mathbf{X} = \mathbf{V} \mathbf{D}_{\lambda}^{rac{1}{2}}$$

Note: there will always be at least one eigenvalue equal to zero.

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- Compute a distance or dissimilarity matrix.
- Compute  $[a_{rs}] = -\frac{1}{2}\delta_{rs}^2$
- Double center A to obtain B = HAH
- Compute eigenvalues and eigen vectors of B
- Compute the solution as  $\mathbf{X} = \mathbf{V} \mathbf{D}_{\lambda}^{\frac{1}{2}}$

# How well do we manage to approximate the distance matrix?

$$\frac{\sum_{i=1}^{P} \lambda_i}{\sum_{i=1}^{n-1} \lambda_i}$$

If **B** is not positive semi-definite:

$$\frac{\sum_{i=1}^{P} \lambda_i}{\sum_{i=1}^{n-1} |\lambda_i|} \quad \text{or} \quad \frac{\sum_{i=1}^{P} \lambda}{\sum_{\lambda_i > 0} \lambda_i}$$

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#### chidean Distance matrix

#### Definition

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A distance matrix **D** is called **Euclidean** if there exists a configuration of points in Euclidean space whose interpoint distances are given by **D**. That is, for some p there exists points  $\mathbf{x}_1, \mathbf{x}_2, \ldots, \mathbf{x}_n$  such that  $d_{rs}^2 = (\mathbf{x}_r - \mathbf{x}_s)'(\mathbf{x}_r - \mathbf{x}_s)$ .

#### Theorem

A distance matrix  $\mathbf{D}$  is Euclidean if and only if  $\mathbf{B}$  (=  $\mathbf{HAH}$ , as previously defined) is positive semi definite.

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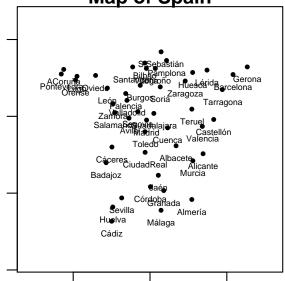
- Sometimes data are given in the form of similarities ( $c_{rs}$ ).
- A similarity matrix **C** has  $c_{rs} = c_{sr}$  and  $c_{rs} \leq c_{rr}$ .
- Similarities can be transformed into distances with the transformation  $d_{rs} = \sqrt{c_{rr} - 2c_{rs} + c_{ss}}$
- If C is psd, then the obtained distance matrix will be Euclidean.

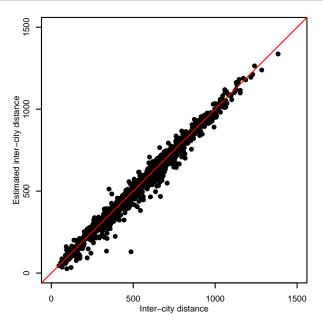
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### R code for classical scaling

```
> ev <- mds.out$eig
> gof <- mds.out$GOF
> print(round(ev,digits=2))
 [1] 4419357.73 3710242.86
                                                    215904.45
                             523390.06
                                         222914.52
                                                                143955.45
 [7]
      128021.63
                 103602.38
                              92361.07
                                          77669.80
                                                     67866.94
                                                                 55724.33
Γ137
       51347.16
                  38327.38
                              32347.58
                                          29609.07
                                                     18785.64
                                                                 14974.46
Γ197
        9473.34
                   9317.99
                               6911.58
                                           4219.73
                                                      1459.24
                                                                   105.43
[25]
           0.00
                   -854.58
                              -3724.49
                                          -4557.54
                                                     -5306.92
                                                                 -8958.67
Γ317
                             -16867.79
      -11879.05
                 -15217.83
                                         -24417.22
                                                    -34120.67
                                                                -43608.19
[37]
      -50334.85
                 -63916.60
                             -77134.54
                                         -80754.15
                                                    -91612.38
                                                                -97422.06
[43] -120383.81 -125973.49 -179445.66 -253056.31 -340735.97
> print(round(gof,digits=4))
[1] 0.8581 1.0000
```

Map of Spain





• STRESS = 
$$\sqrt{\frac{\sum_{r \neq s}^{n} (f(\delta_{rs}) - d_{rs})^{2}}{\sum_{r \neq s} d_{rs}^{2}}}$$

- $stress(\Delta, \hat{\mathbf{X}}) = \min_{all\mathbf{X}} stress(\Delta, \mathbf{X})$
- We minimize the objective function numerically, starting from an initial configuration.

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- Choose a distance measure (e.g.  $\delta_{rs} = \left\{\sum_{i=1}^p |x_{ri} x_{si}|^{\lambda}\right\}^{\frac{1}{\lambda}}$ )
- Choose a monotone transformation f
- Choose an algorithm to minimize Stress.

#### Global versus local minima

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- Use different initial configurations
- Compare stress over 1,2,3,... dimensional solutions

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### Diagnostics

- Scatter plot of  $\delta_{rs}$  versus  $d_{rs}$
- Plot stress versus number of dimensions
- Degeneracy (many points with the same  $d_{rs}$ )
- Compute residuals  $(d_{rs} f(\delta_{rs}))$

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Population substructure

## R code for non-metric MDS

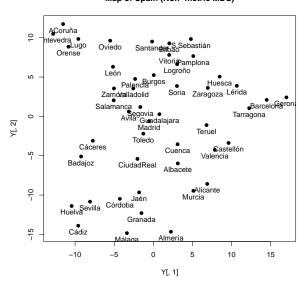
```
> init <- scale(matrix(runif(n*2),ncol=2),scale=FALSE)
> nmmds.out <- isoMDS(Spain,y=init,k=2)
initial value 41.659041
iter
       5 value 40.219780
iter 10 value 37,286307
iter 15 value 30.177635
iter 20 value 22.661686
iter 25 value 14.483317
iter 30 value 10.703962
iter 35 value 7.756514
iter 40 value 6.116380
iter 45 value 5.360785
iter 50 value 5 145884
final value 5.145884
stopped after 50 iterations
```

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Population substructure

```
> nmmds.out <- isoMDS(Spain, y=init, k=2, maxit=100)
initial value 41.659041
       5 value 40.219780
iter
iter 10 value 37,286307
iter 15 value 30.177635
iter 20 value 22 661686
iter 25 value 14.483317
iter 30 value 10.703962
iter 35 value 7.756514
iter 40 value 6.116380
iter 45 value 5.360785
iter 50 value 5.145884
iter 55 value 5.088756
final value 5.057439
converged
> Y <- nmmds.out$points
> nmmds2.out <- isoMDS(Spain,y=X2,k=2) # PCO solution as initial configuration
initial value 6.252429
final value 6.252214
converged
> Y2 <- nmmds2.out$points
> plot(Y[,2],Y[,1],pch=19)
> text(Y[,2], Y[,1], rownames(Spain), cex=0.5,pos=1)
```

### Map of Spain (Non-metric MDS)

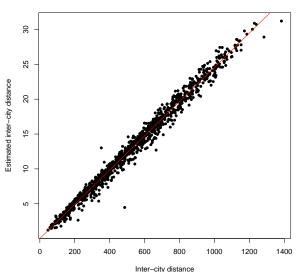


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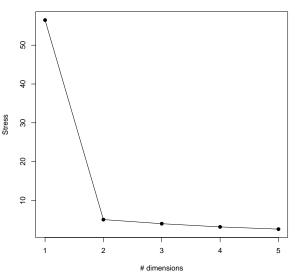
# Diagnostics non-metric MDS

#### Non-metric MDS

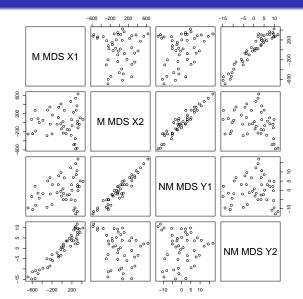


## Diagnostics non-metric MDS

#### Stress versus dimensionality



### Relation metric MDS and non-metric MDS solutions



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## Correlation solutions MDS versus non-metric MDS

	М	MDS	Х1	М	MDS	Х2	NM	MDS	Y1	NM	MDS	Y2
M MDS X1		1.	.00		0	.00		0.	.31		0	. 96
M MDS X2		0	.00		1	.00		0.	. 95		-0	. 29
NM MDS Y1		0	.31		0	. 95		1.	.00		0	. 02
NM MDS Y2		0	.96		-0	. 29		0.	.02		1	.00

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## MDS for genetic data

- There is a rich literature on how to measure genetic distance
- The allele sharing distance is an often used measure

i∖j	AA	AB	BB
AA	2	1	0
AB	1	2	1
BB	0	1	2

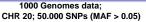
$i \setminus j$	AA	AB	ВВ
AA	0	1	2
AB	1	0	1
BB	2	1	0

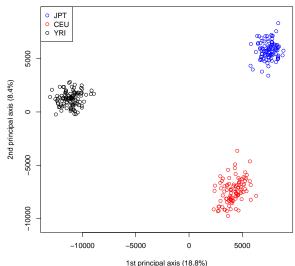
- Let  $x_{ijk}$  be the number of shared alleles of individual i and j for variant k
- $d_{ijk} = 2 x_{ijk}$
- Often scaled by multiplying by  $\frac{1}{2}$
- Typically averaged over K genetic variants:

$$d_{ij} = \frac{1}{K} \sum_{k=1}^{K} d_{ijk}$$

• The so obtained  $D = [d_{ij}]$  is used as input for MDS.

#### MDS with genetic data (CEU, JPT and YRI samples from 1,000 Genomes)





Dim	λ	%	% Cum.
1	4.04	0.14	0.14
2	1.64	0.06	0.20
3	1.43	0.05	0.25
4	0.98	0.04	0.29
5	0.98	0.04	0.32
6	0.78	0.03	0.35
7	0.74	0.03	0.37
8	0.69	0.02	0.40
9	0.65	0.02	0.42
	0.03	0.02	0.42
62	0.14	0.00	0.98
63	0.13	0.00	0.99
64	0.13	0.00	0.99
65	0.13	0.00	1.00
66	0.12	0.00	1.00
67	0.12	0.00	1.01
68	0.11	0.00	1.01
69	0.11	0.00	1.01
121	0.01	0.00	1.11
122 123	0.01	0.00	1.11
	0.00	0.00	1.11
124 125	0.00	0.00	1.11
125	0.00	0.00	1.11
127	-0.00	-0.00	1.11
128	-0.00	-0.00	1.11
129	-0.00	-0.00	1.11
130	-0.01	-0.00	1.11
200	-0.08	-0.00	1.01
201	-0.08	-0.00	1.01
202	-0.09	-0.00	1.00
203	-0.10	-0.00	1.00

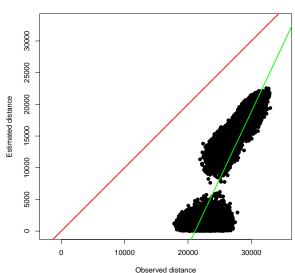
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 Metric MDS
 Non-metric MDS
 Example
 Computer exercise

 00000000000000
 00000000000
 00

## Goodness of fit

R2 0.81



Population substructure

- Load the database CHBChr2-200.rda
- Convert the genotype data into and  $n \times n$  distance matrix.
- Produce a map of the individuals by metric multidimensional scaling. Is there evidence for the existence of groups?
- Make a graph of the fitted against the observed distances, and comment on the results.
- Produce a map of the individuals by non-metric multidimensional scaling. Are the results comparable not those obtained by metric MDS? Is there evidence for the existence of groups?

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