Imperial College London



Multivariate analysis of genetic data — uncovering spatial structures —

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Outline

Introduction

Testing spatial structures
Moran's Index
Mantel's correlation

Multivariate analysis of spatial patterns

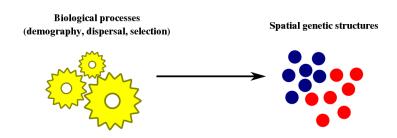
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Moran's Index

From processes to structures

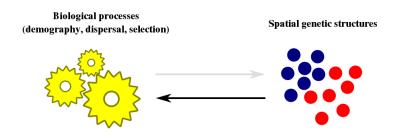
Genetic structure: non-random distribution of genetic diversity.



Identify structures to infer processes.

From processes to structures

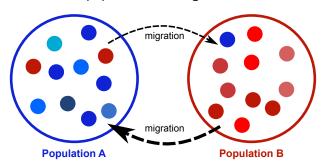
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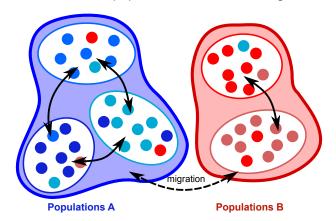
Island model

Reproduction within populations + migration.



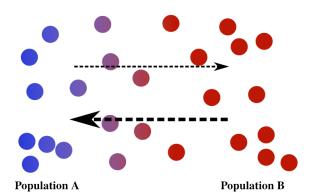
Hierarchical island model

Reproduction within subpopulations + stratified migration.



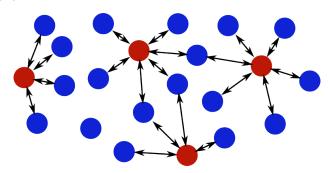
Isolation by distance (IBD)

Reproduction between neighbours → 'diffusion' of genes



Inbreeding avoidance

Mating with individuals from another population \rightarrow 'repulsion' structure



- island / hierarchical island model: patches of related genotypes
- isolation by distance (IBD): clines of genetic differentiation
- inbreeding avoidance: repulsion structure

⇒ Genetic processes often create spatial structures. **How can we reveal them?**

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Spatial autocorrelation

Definitions:

- *in general*: values of a variable non independent from the corresponding spatial locations
- in genetics: genetic distance is correlated to spatial distance

Two types of spatial autocorrelation:

- positive: closer individuals are more similar than at random
- negative: closer individuals are more dissimilar than at random

Spatial autocorrelation

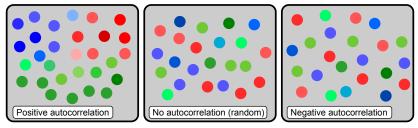
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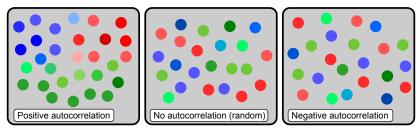
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Spatial autocorrelation: illustration



How do we measure spatial autocorrelation?

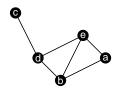
Spatial autocorrelation: illustration



How do we measure spatial autocorrelation?

From spatial coordinates to spatial weights

Matrix of spatial weights L



Row i: uniform weights for neighbours of i.

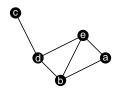
	а	b	С	d	€
а	0.000	0.500	0.000	0.000	0.500
Ь	0.333	0.000	0.000	0.333	0.333
С	0.000	0.000	0.000	1.000	0.000
d	0.000	0.333	0.333	0.000	0.333
е	0.333	0.333	0.000	0.333	0.000

Let x be a variable with one value at each location.

The lag vector Lx computes mean values of neighbours.

From spatial coordinates to spatial weights

Matrix of spatial weights ${f L}$



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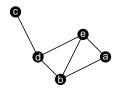
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а	0.000	0.500	0.000	0.000	0.500
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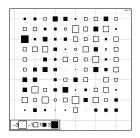
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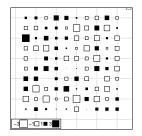
Lag vector:

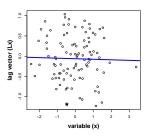
Random:



Lag vector :

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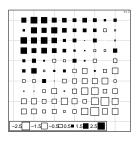


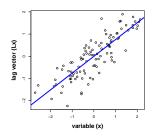


	Df	Sum Sq	Mean Sq	F value	Pr(>F)
X	1	0.02	0.02	0.06	0.8081
Residuals	98	31.53	0.32		

Lag vector :

Positive autocorrelation:

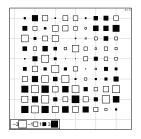




	Df	Sum Sq	Mean Sq	F value	Pr(>F)
xG	1	65.91	65.91	245.69	0.0000
Residuals	98	26.29	0.27		

Lag vector:

Negative autocorrelation:



variable (x)

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
xL	1	87.56	87.56	77.80	0.0000
Residuals	98	110.29	1.13		

Moran's index: definition

Moran's I:

$$I(\mathbf{x}) = \frac{\mathbf{x}^T \mathbf{L} \mathbf{x}}{n} \frac{1}{\mathsf{var}(\mathbf{x})}$$

where:

- $\mathbf{x} \in \mathbb{R}^n$: a centred variable (e.g. allele frequency, PC)
- L : matrix of spatial weights $(n \times n)$
- Lx : lag vector
- $I_0 = \frac{-1}{n-1} \approx 0$: null value (no autocorrelation, i.e. random spatial distribution)
- \Rightarrow Moran's I varies like $\langle \mathbf{x}, \mathbf{L} \mathbf{x} \rangle$.

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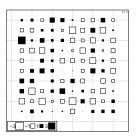
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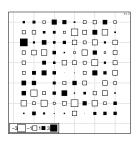
Random:

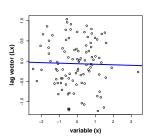


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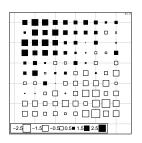
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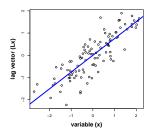


Moran's I: $I(\mathbf{x}) \approx I_0$

Positive autocorrelation:



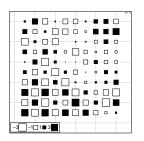
Lag vector:



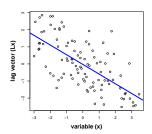
Moran's I:

$$I(\mathbf{x}) > I_0$$

Negative autocorrelation:



Lag vector :



Moran's I: $I(\mathbf{x}) < I_0$

- ullet compute I from the data
- permute randomly the locations to get a value of I under H_0 :
 "x is distributed at random across space."
- repeat this operation a large number of times to obtain a reference distribution of I under H_0
- compare initial value to the reference distribution to get a p-value.

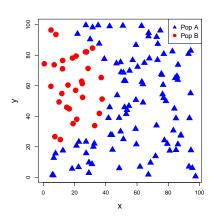
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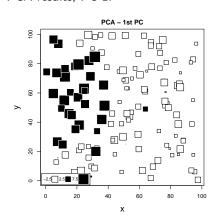
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Application: testing spatial structures in principal components

Data (2 population, island model):

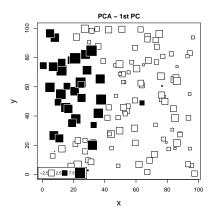


PCA results, PC 1:

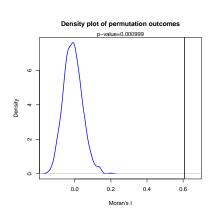


Application: testing spatial structures in principal components

PCA results, PC 1:



Moran's I test of PC1:



- Moran's I is univariate
- solution: test a few principal components
- problems:
 - does not use all the genetic information
 - which PC to test?
 - correction for multiple testing
- ⇒ need for multivariate tests

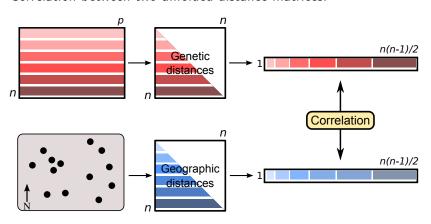
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Mantel's correlation: rationale

Correlation between two unfolded distance matrices.



Mantel's correlation: definition

Notations:

- $\mathbf{X} = [x_{ij}]$ ($\mathbf{X} \in \mathbb{R}^{n \times n}$): genetic distances
- $\mathbf{Y} = [y_{ij}] \ (\mathbf{Y} \in \mathbb{R}^{n \times n})$: geographic distances
- \bar{x} , \bar{y} : means of x and y (excepting diagonals)
- s_x , s_y : standard deviation of x and y (excepting diagonals)

Original definition (unstandardized):

$$z_{\mathsf{M}} = \sum_{i=1}^{n-1} \sum_{j=i+1}^{n} x_{ij} y_{ij}$$

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Standardized coefficient (true correlation):

$$r_{\mathsf{M}} = \frac{1}{d-1} \sum_{i=1}^{n-1} \sum_{j=i+1}^{n} \left(\frac{x_{ij} - \bar{x}}{s_x} \right) \left(\frac{y_{ij} - \bar{y}}{s_y} \right)$$

- ullet compute $z_{f M}$ or $r_{f M}$ from the data
- permute randomly the rows and columns of one matrix, recompute the test statistic (i.e., under H_0 : "no correlation")
- repeat this operation many times to generate a reference distribution
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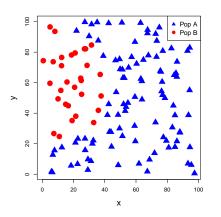
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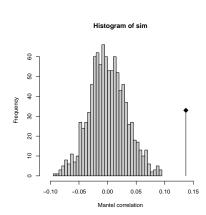
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Mantel test:



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Mapping principal components

Maps of the three first principal components of PCA.



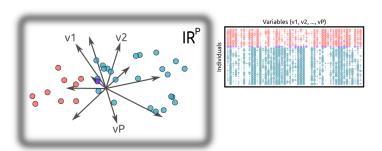
Are we actually looking for spatial patterns here?

Mapping principal components

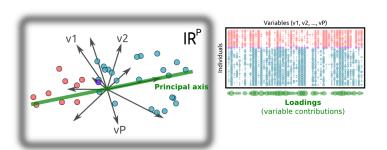
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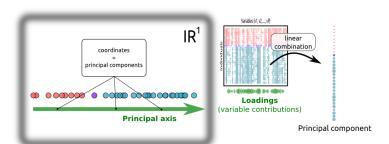
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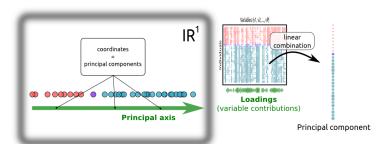
Principal components with maximum total variance.



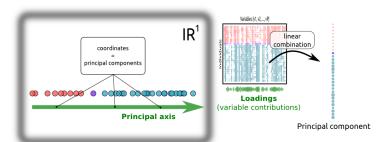
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- they may reveal obvious spatial structures, but overlook finer patterns
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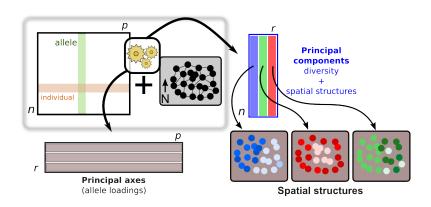
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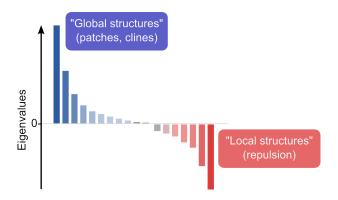
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Spatial Principal Component Analysis (sPCA): outputs



Global and local structures

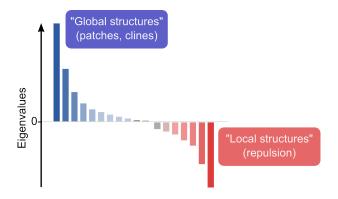
Unlike other multivariate methods, sPCA has **positive** and **negative** eigenvalues



How do we get these in practice?

Global and local structures

Unlike other multivariate methods, sPCA has **positive** and **negative** eigenvalues



How do we get these in practice?

Time to get your hands dirty (one last time)!



The pdf of the practical is online:

http://adegenet.r-forge.r-project.org/

or

Google o adegenet o documents o "Workshop Barcelona, October 2015"