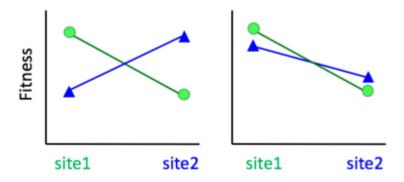
Genome-scans Landscape genomics

Claire Mérot & Anna Tigano Physalia Courses

Basic principle: Local adaptation

Geographic heterogeneity in environment

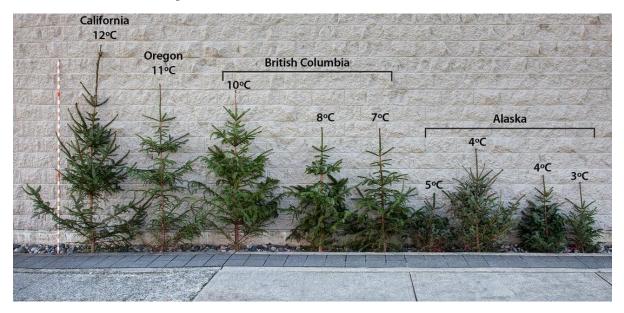
Local populations have been selected by local ecological conditions



Basic principle: Local adaptation

Common garden experiment

- ⇒ local adaptation has heritable genetic basis
- ⇒ phenotypes related to local adaptation



Provenance variation in 8-year-old *Picea sitchensis* from across the species range grown in a common garden in Vancouver, BC, Canada

Aitken et al, 2015 Evol.App https://www.onlinelibrary.wiley.com/doi/10.1111/eva.12293

- ⇒ This may be visible as « signature of selection »
- ⇒ We expect differences of allelic frequency between sites

Basic principle: Local adaptation

Can we use genomic data to understand the genetic basis of local adaptation?

Can we find the loci contributing to divergence between populations?

Can we find the loci possibly associated with relevant traits or relevant ecological variables?

Genome-scan for local adaptation

Approach 1: detect outliers of divergence

-> Search for unexpected patterns in allele frequencies accross the genome

Approach 2: detect associations with environment/phenotype

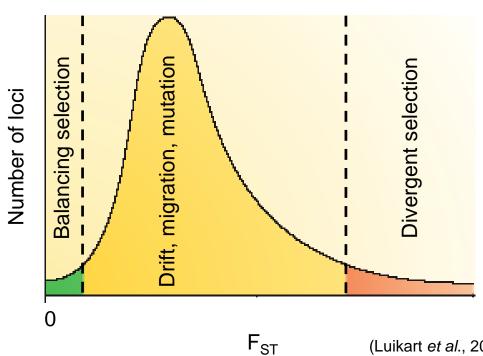
-> Search for correlations between allelic frequencies and variables

Fst statistics

- A measure of differentiation between populations relatively to intra-population diversity

Fst=1: complete fixation of the alleles in each population

Fst=0 : same allelic frequencies



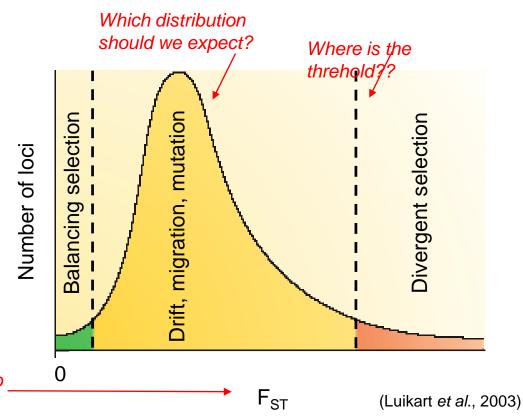
(Luikart et al., 2003)

Fst statistics

- A measure of differentiation between populations relatively to intra-population diversity

Fst=1: complete fixation of the alleles in each population

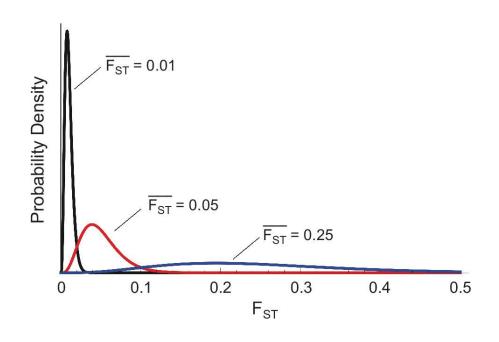
Fst=0 : same allelic frequencies



Which pair to compare?

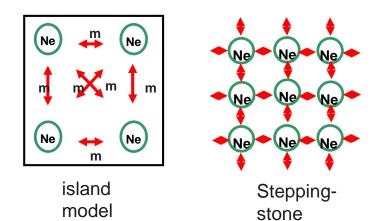
Problem 1: Population structure

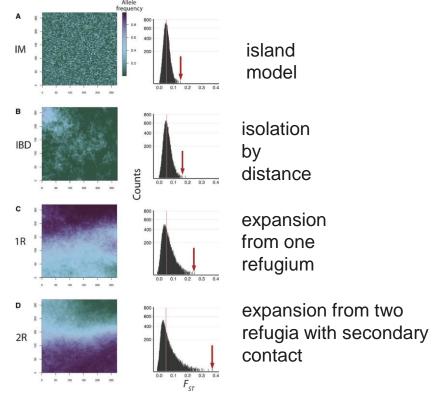
When the average level of differentiation is high,, the variance in $F_{\rm ST}$ values among loci increases with average $F_{\rm ST}$, which makes detection of outlier loci difficult for highly differentiated populations



Problem 2: Demography

The distribution of FST will depend on the real demography of the populations





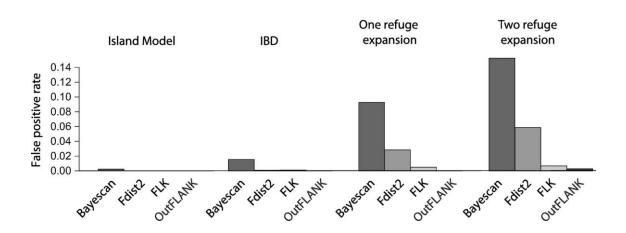
Landscapes of frequency and FST for an outlier neutral locus at the end of the simulation (75 random samples)

1st solution: assume a model of dispersion and demography (Fdist, Bayescan)

2nd solution: try to estimate a neutral model from data

- -> covariance matrix between pop (Baypass/Bayenv2, FLK)
- -> X² distribution of pruned SNPs (OutFLANK)

Many « false positive »...



Or capturing effects unlinked to selection (sampling bias, unaccounted structure, hybrididation)...

Problem 3: Sample size

The power to detect meaningful differences based on genome-scan will depend on the number of populations, and the number of samples within a population

Table 2: Power for the one-refuge case analyzed with OutFLANK as a function of sample sizes, based on loci with $s_L = 0.01$, using >1,700 tests per case with a q value threshold of 5%

	8 1			
No. individuals	5	10	20	40
per population	populations	populations	populations	populations
5	0	.09	.52	.84
10	.10	.56	.82	.94
20	.37	.75	.90	.95
40	.55	.81	.94	.97

Few samples per pop -> stochasticity in allelic frequency

Note: Parameters otherwise similar to those in figure 2.

Few populations

-> low statistical power

Problem 4: Background selection

Fst is a relative measure of variation among populations. Low heterozygosity can inflate Fst values even for small differences in allelic frequencies...

For e.g. background selection (negative selection in regions of low-recombination)

-> Should we use other measures: dxy? (but lower power for early stages of divergence)

AFD: Allelic frequency differences?

(Berner 2019, Genes 10.3390/genes10040308)

-> but the effect of background selection on Fst may not be that bad for populations <u>connected</u> <u>by high gene flow</u>.

See Matthey-Doret & Whitlock 2019 MolEcol https://onlinelibrary.wiley.com/doi/abs/10.1111/mec.15197

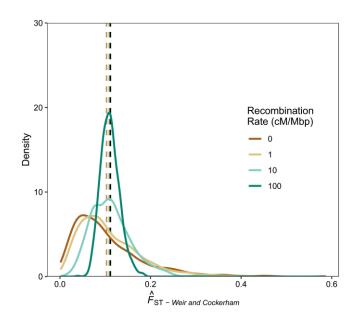
Problem 5: Recombination

Even without selection, Fst variance is expected to be higher in low-recombination regions...

-> Should we use a different threshold depending on recombination??

See Booker, Yeaman & Whitlock 2020 MolEcol https://onlinelibrary.wiley.com/doi/abs/10.1111/mec.15501

A field in development with upcoming whole-genome data...



But the main problem is: how can we interpret the results?

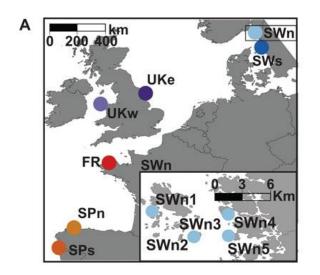
Outliers tests can be useful to remove "putatively-adaptive" loci from analysis of neutral structure/demography...

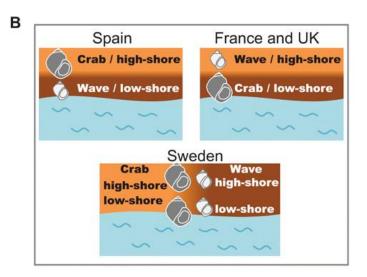
->But do we want to avoid taking into account adaptation or specific loci showing traces of differentiation?!

What does outliers of differentiation means?

-> depend on the study design/ecological information...

Pairs of populations: contrast Fst



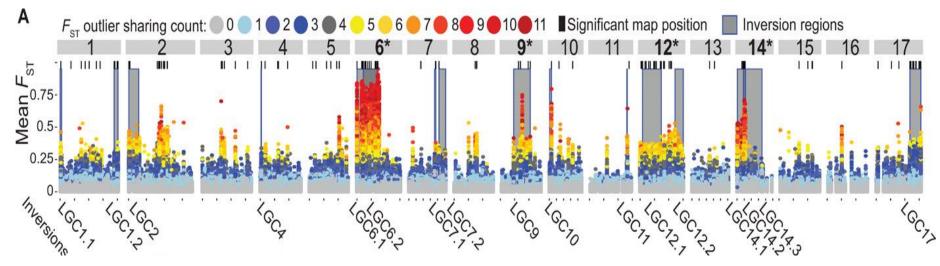


Outliers in pairs of *Littorina* ecotypes Crab vs Wave at different localities accross Europe...



Morales et al, 2019 Science advances https://doi.org/10.1126/sciadv.aav9963

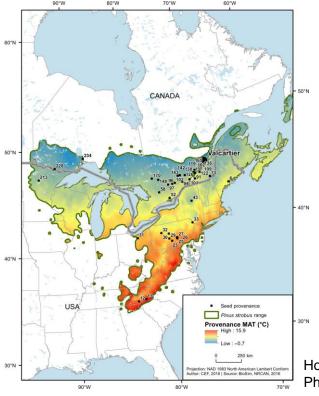
Pairs of populations: contrast Fst





Fst Crab vs Wave at different localities accross Europe: outlier sharing

Along a cline, accross a geogaphic gradient



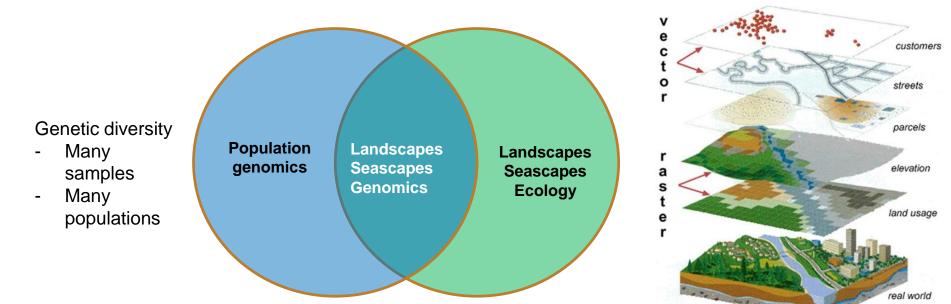
Ecological information

Genetic information

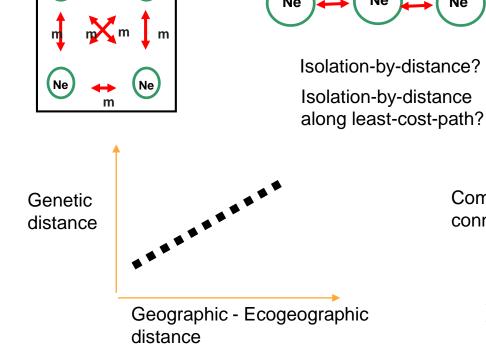
⇒ hypothesis: populations experiencing the same selective pressures (same environment) will be less differentiated then populations experiencing different ecological conditions at adaptive loci

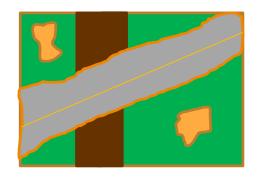
Housset, 2018, New Phytologist

Lanscapes or seascapes genomics contribute to understand both neutral and adaptive genetic variation

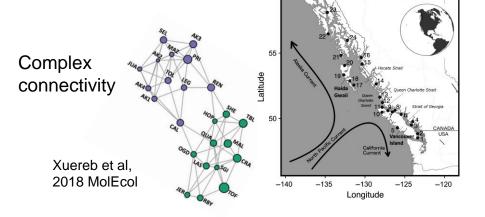


a better understanding of gene flow & connectivity

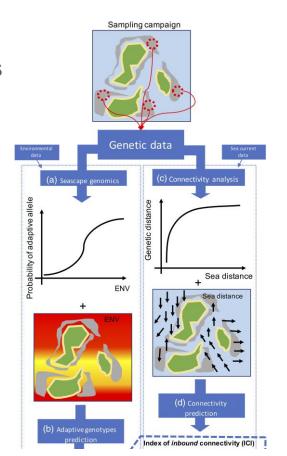


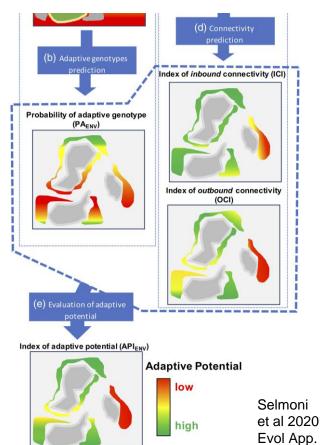


Isolation-by-resistance?

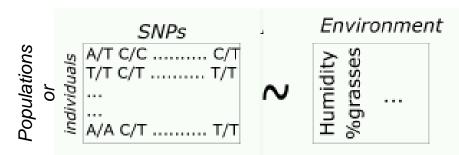


- Analyzing adaptation to known ecological variables
- ⇒ A better sense of why populations are differentiated?
- ⇒ Predictions for models of adaptation, response to climate change, etc





Environmental associations methods



Univariate methods: locus by locus
 (correlation freq/env, LFMM, Bayenv/Baypass, etc)

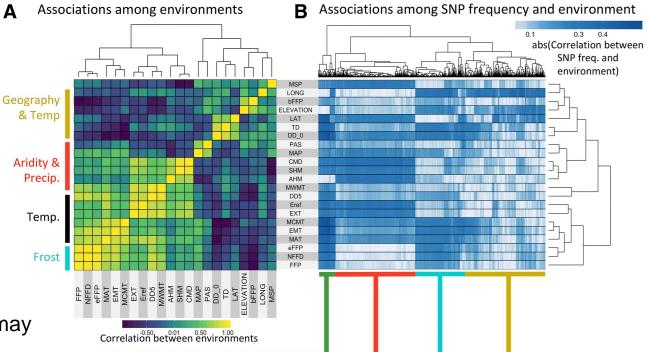
Genome scan methods against more complex models: when and how much should we trust them? (Villemereuil et al, 2014 MolEcol)

 Multivariate methods (redundancy analysis RDA)

Comparing methods for detecting multilocus adaptation with multivariate genotype—environment associations (Forester et al 2018, MolEcol)

Univariate associations:

Which locus is associated with which environmental variable? -> Spearman's correlation

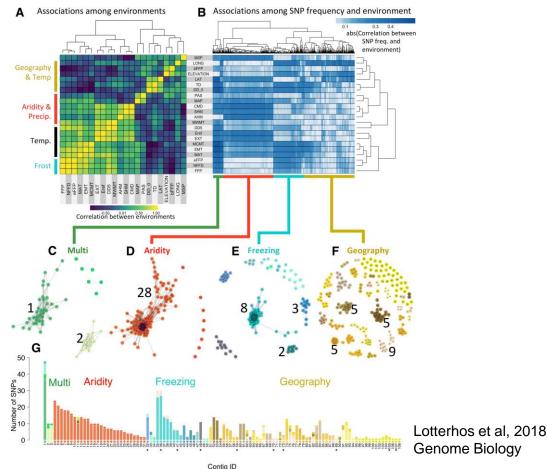


Caution:

- environmental variables may be correlated!
- SNPs may be in physical LD

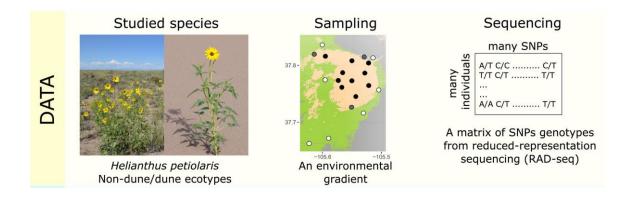
Univariate associations:

Modular group of adaptive loci to different axis of environmental variation



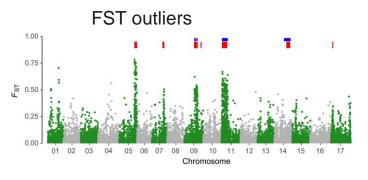
Univariate associations:

Bayenv2 - Baypass

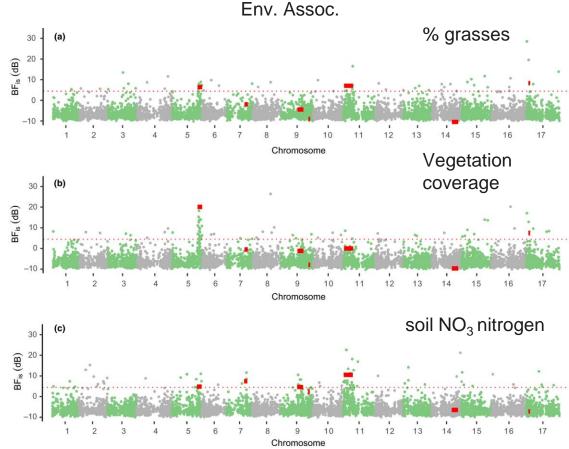


Huang et al, 2020 MolEcol

Univariate associations: Bayenv2 – Baypass

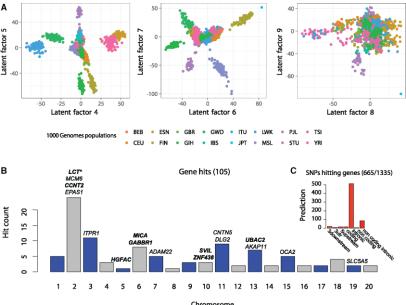


Huang et al, 2020 MolEcol



- Univariate associations: LFMM (latent factor mixed model)
- Detects correlations between environmental and genetic variation while simultaneously inferring background levels of population structure
- Residual population structure is introduced via unobserved K (latent) factors
- Latent factors represent demographic history, IBD, hidden substructure
- Lfmm2: New & faster version which can works on SNP or methylation matrix

Caye et al, 2019, Molecular Biology and Evolution https://doi.org/10.1093/molbev/msz008

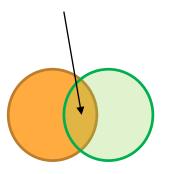


Human GEA study. Association study based on genomic data from the 1000 Genomes Project database and climatic ...

Univariate associations:

Recommendation:

- intersect of several methods
- ⇒ More likely to be strong candidates for adaptation
- ⇒ Reduce false positive but may also miss variants with less signal...
- Controlling false discovery rate
- Correct for population structure
- ⇒ An open debate?
- ⇒ Likely depends on the system: high gene flow? IBD? Geography correlated with environmental variation?

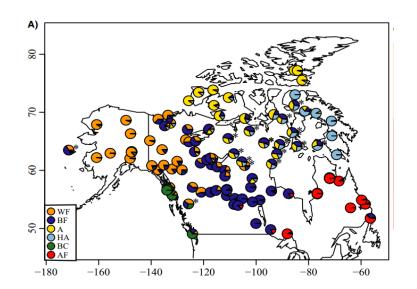


Controlling false discoveries in genome scans for selection

François et al, 2015, Molecular Ecology https://doi.org/10.1111/mec.13513

RDA

Multivariate associations:

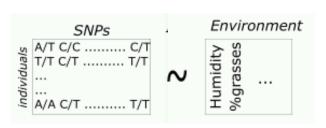




species

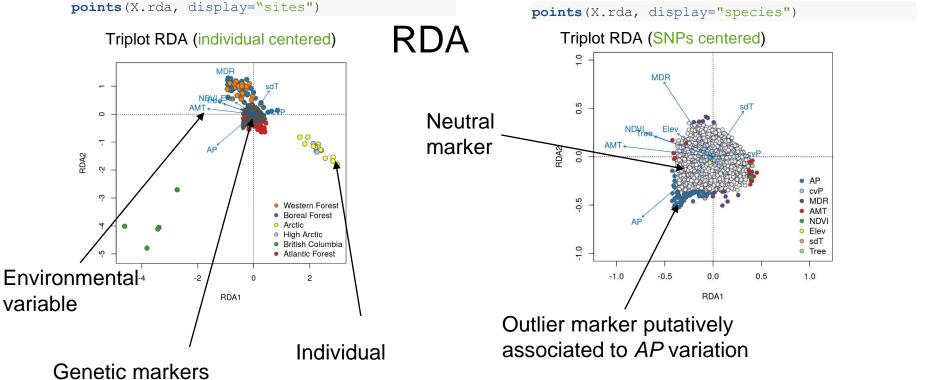
sites

In community ecology (package vegan!)



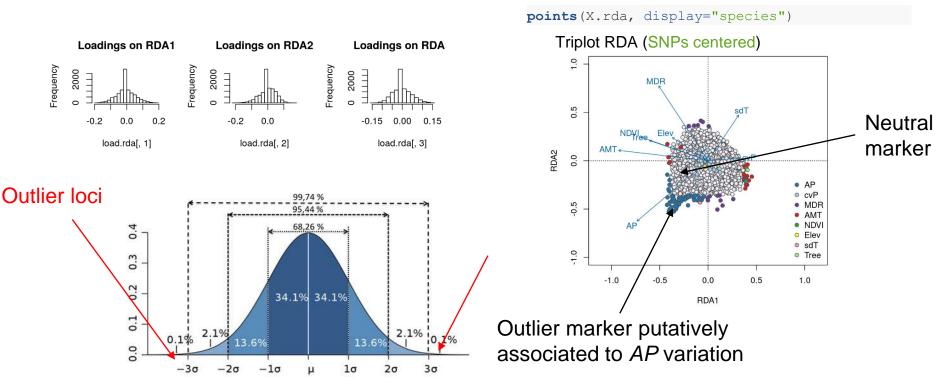
94 wolves42 597 SNPs

Forester et al 2018 Mol Ecol



Forester et al 2018 Mol Ecol

Use the contribution of genetic markers along the different axis to detect putatively-selected loci



Forester et al 2018 Mol Ecol

Use the contribution of genetic markers along the different axis to detect putatively-selected loci

A super good vignette to understand and do Rda analysis:

https://popgen.nescent.org/2018-03-27 RDA GEA.html

Population Genetics in R

Users ▼ Package Developers ▼ Contribute! ▼ Useful Links

Detecting multilocus adaptation using Redundancy Analysis (RDA)

- Introduction
- Assumptions
- Data & packages
- Analysis
- Conclusions
- Contributors
- References
- Session Information

Introduction

The purpose of this vignette is to illustrate the use of Redundancy Analysis (RDA) as a genotype-environment association (GEA) method to detect loci under selection (Forester et al., 2018). RDA is a multivariate ordination technique that can be used to analyze many loci and environmental predictors simultaneously. RDA determines how groups of loci covary in response to the multivariate environment, and can detect processes that result in weak, multilocus molecular signatures (Rellstab et al., 2015; Forester et al., 2018).

RDA is a two-step analysis in which genetic and environmental data are analyzed using multivariate linear regression, producing a matrix of fitted values. Then PCA of the fitted values is used to produce canonical axes, which are linear combinations of the predictors (Legendre & Legendre, 2012). RDA can be used to analyze genomic data derived from both individual and population-based sampling designs.

Assumptions

RDA is a linear model and so assumes a linear dependence between the response variables (genotypes) and the explanatory variables (environmental predictors). Additional detail can be found in Legendre & Legendre (2012). We also recommend Borcard et al. (2011) for details on the implementation and interpretation of RDA using the vegan package (Oksanen et al, 2017).

Contributors

- · Brenna R. Forester (Author)
- Martin Laporte (reviewer)
- Stéphanie Manel (reviewer)

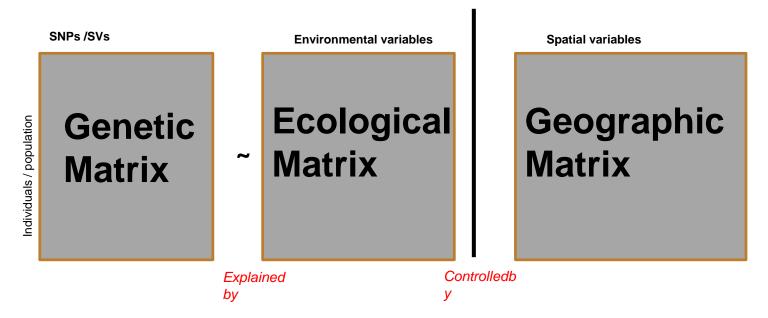
Multi-locus

-> Polygenic adaptation possible to detect?

Multi-variable analysis -> realistic environmental caracterisation

Very fast + Global information:

- Which variables explain genetic variance?
- Correction possible by population/geographic structure



https://doi.org/10.1016/B978-0-444-53868-0.50014-9

RDA

Spatial-eigen vectors are a way to reduce a distance matrix between samples/populations

- -> not necessarily neutral
- -> describe different possible spatial combination

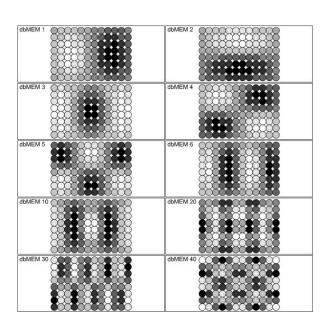






Latitude + Longitude or Spatial eigenvectors

= db-MEM



More information: Legendre & Legendre

https://doi.org/10.1016/B978-0-444-53868-0.50014-9

A must-read:

Finding the Genomic Basis of Local Adaptation: Pitfalls, Practical Solutions, and Future Directions
Hoban et al 2016 Am Nat
https://www.journals.uchicago.edu/doi/full/10.1086/688018