

A (short) introduction to ordination with the vegan package

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	Raw data	Transformed data	Unimodal	Distance-based	Model-based
Unconstrained	PCA		CA, DCA	PCoA, NMDS	MM, LVM
Constrained	RDA	tb-PCA tb-RDA	CCA	db-RDA	CAO, CQO
Other				Permanova, Dispersion	manyglm

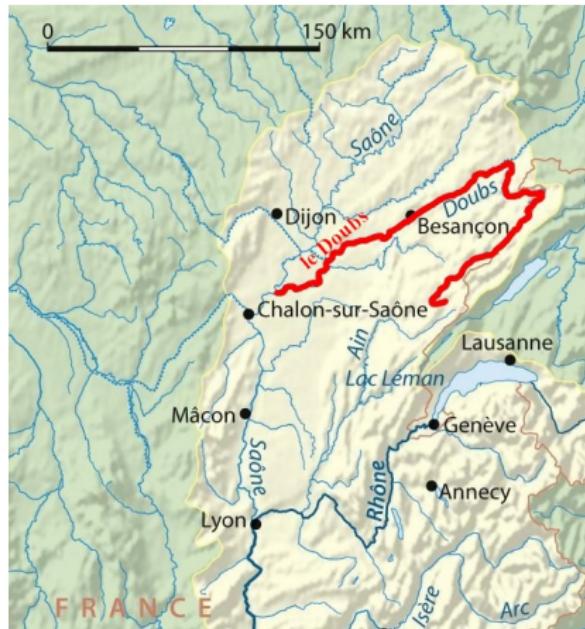
(Nearly) no maths today ;)



Datasets

Demonstration: Doubs river fish communities

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- ▶ Fish communities
- ▶ 30 sites along the Doubs River

Verneaux, J. (1973) Cours d'eau de Franche-Comte (Massif du Jura). Recherches écologiques sur le réseau hydrographique du Doubs. Essai de biotypologie. These d'état, Besançon. 1–257.

Datasets



Unconstrained Ordination

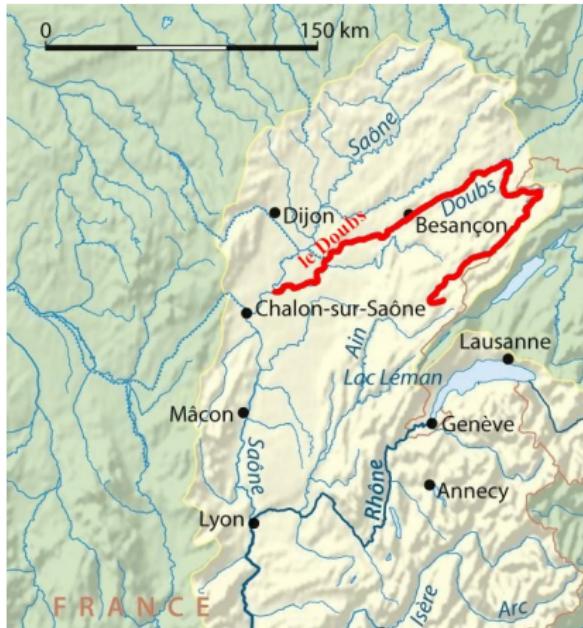


Constrained Ordination



Model diagnostics / testing





- ▶ Fish communities
 - ▶ 30 sites along the Doubs River

Questions

- ▶ How does fish composition change downstream?
 - ▶ Environmental drivers?

Verneaux, J. (1973) Cours d'eau de Franche-Comte (Massif du Jura). Recherches ecologiques sur le reseau hydrographique du Doubs. Essai de biotypologie. These d'etat, Besancon. 1-257.

```
Dabu <- read.table('doubtsAbu.csv', sep = ',', header = TRUE)
Denv <- read.table('doubtsEnv.csv', sep = ',', header = TRUE)
Dspa <- read.table('doubtsSpa.csv', sep = ',', header = TRUE)
```

dim(Dabu)

[1] 30 27

30 sites, 27 taxa

```
head(Dabu[, 1:18])
```

	CHA	TRU	VAI	LOC	OMB	BLA	HOT	TOX	VAN	CHE	BAR	SPI	GOU	BRO	PER	BOU	PSO	ROT
1	0	3	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
2	0	5	4	3	0	0	0	0	0	0	0	0	0	0	0	0	0	
3	0	5	5	5	0	0	0	0	0	0	0	0	0	1	0	0	0	
4	0	4	5	5	0	0	0	0	0	1	0	0	1	2	2	0	0	
5	0	2	3	2	0	0	0	0	5	2	0	0	2	4	4	0	0	
6	0	3	4	5	0	0	0	0	1	2	0	0	1	1	1	0	0	

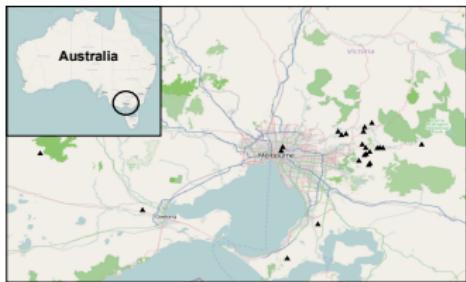
```
# Dimension and first rows of Environmental data  
dim(Denv)
```

[1] 30 11

30 sites, 11 variables

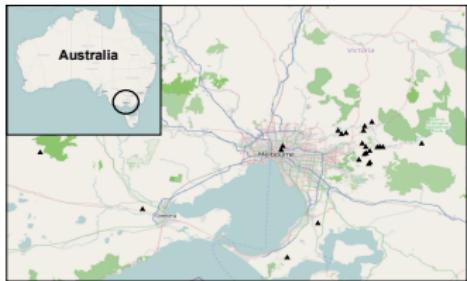
head(Deny)

	das	alt	pen	deb	pH	dur	pho	nit	amm	oxy	dbo
1	0.3	934	48.0	0.84	7.9	45	0.01	0.20	0.00	12.2	2.7
2	2.2	932	3.0	1.00	8.0	40	0.02	0.20	0.10	10.3	1.9
3	10.2	914	3.7	1.80	8.3	52	0.05	0.22	0.05	10.5	3.5
4	18.5	854	3.2	2.53	8.0	72	0.10	0.21	0.00	11.0	1.3
5	21.5	849	2.3	2.64	8.1	84	0.38	0.52	0.20	8.0	6.2
6	32.4	846	3.2	2.86	7.9	60	0.20	0.15	0.00	10.2	5.3



- ▶ Macrovertebrate communities
 - ▶ 24 sites
 - ▶ covering a salinity and toxicity gradient

The dataset is published in: Szöcs, E., Kefford, B.J., Schäfer, R.B., 2012. Is there an interaction of the effects of salinity and pesticides on the community structure of macroinvertebrates? *Science of the Total Environment* 437, 121–126.



- ▶ Macroinvertebrate communities
 - ▶ 24 sites
 - ▶ covering a salinity and toxicity gradient

Questions:

- ▶ Interaction between salinization and pesticides?
 - ▶ Which species are affected?
 - ▶ Other influences?

The dataset is published in: Szöcs, E., Kefford, B.J., Schäfer, R.B., 2012. Is there an interaction of the effects of salinity and pesticides on the community structure of macroinvertebrates? *Science of the Total Environment* 437, 121–126.

Exercise: Salinization and Pesticides

```
# setwd('3-Ordination/data/')
abu <- read.table('melbourneAbu.csv', sep = ';', header = TRUE)
env <- read.table('melbourneEnv.csv', sep = ';', header = TRUE)
```

```
# dimensions of data.frame  
dim(env)
```

[1] 24 23

dim(abu)

[1] 24 76

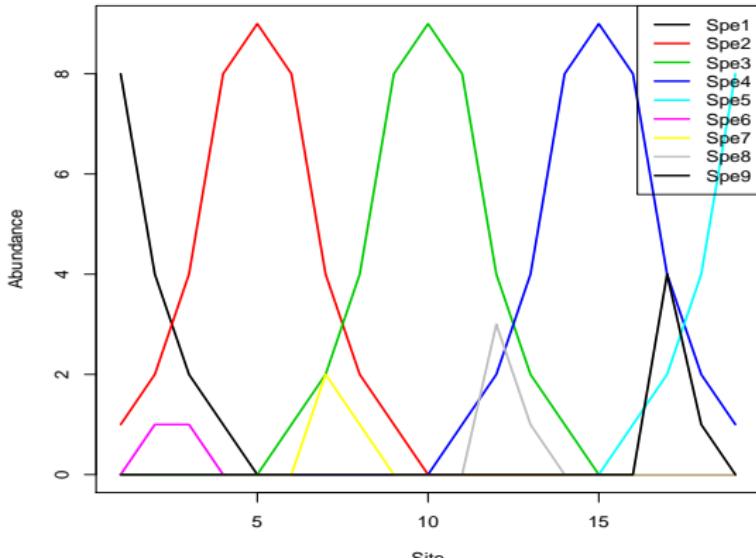
24 sites, 22 environmental variables, 75 taxa

```
head(env[ , 1:10])
```

ID	T	pH	oxygen	Depth	maxwidth	minwidth	rifperc	poolperc	Bedrock	
1	1-11	16.8	7.67	80.1	0.9	15	12.0	0	100	0
2	2-11	16.5	7.29	83.0	0.9	30	15.0	0	100	0
3	3-11	17.3	7.20	77.9	0.4	4	2.5	0	100	0
4	4-11	15.6	7.84	72.0	0.7	8	2.5	0	100	0
5	5-11	17.2	6.97	69.9	0.9	7	4.0	0	100	0
6	6-11	15.5	7.26	80.0	0.2	3	2.0	5	95	0

Exercise: Dummy abundances

```
# Load dummy data
dummy <- read.table('dummydata.csv', header = TRUE, sep = ';')
# plot dummy data
matplot(dummy[ , -1], type = 'l', xlab = 'Site', ylab = 'Abundance',
        lty = 'solid', lwd = 2, col = 1:9)
legend('topright', legend = colnames(dummy)[-1],
       col = 1:9, lty = 'solid', lwd = 2)
```



Unconstrained Ordination

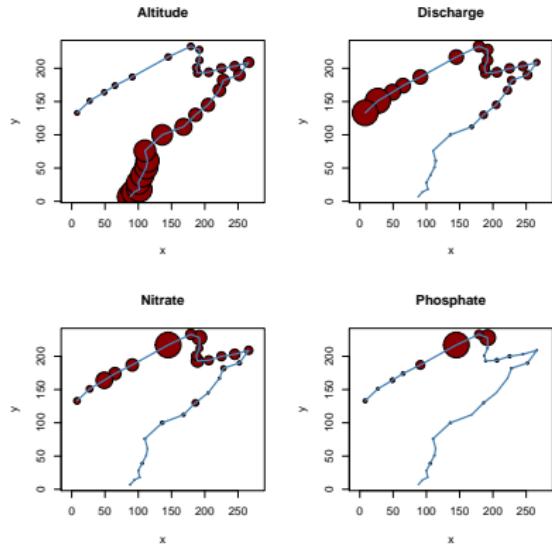
Principal Components Analysis (PCA)

Principal coordinates analysis (PCoA)

Nonmetric Multidimensional Scaling (NMDS)

Principal Components Analysis (PCA) — Why?

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- ▶ 11 variables

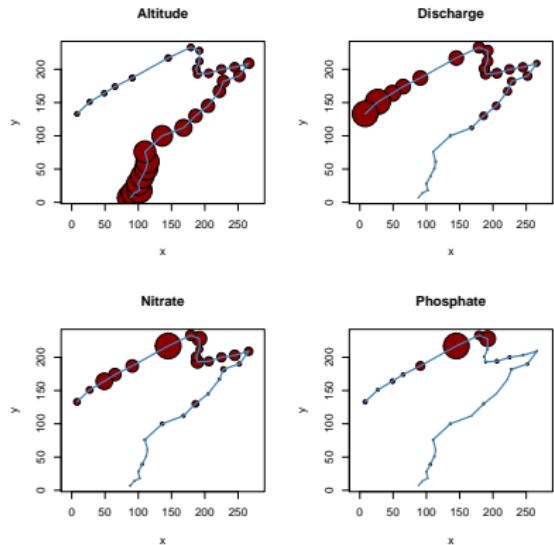
Questions:

- ▶ Which variables are correlated?
- ▶ Which sites have similar conditions?
- ▶ How do conditions change downstream?

Solutions?

Principal Components Analysis (PCA) — Why?

11 / 53



- ▶ 11 variables

Questions:

- ▶ Which variables are correlated?
- ▶ Which sites have similar conditions?
- ▶ How do conditions change downstream?

Solutions?

- ▶ pairwise comparisons
- ▶ 3D possible
- ▶ more than 3 dimensions?

Datasets
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Unconstrained Ordination
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Constrained Ordination
oooooooooooo

Model diagnostics / testing
oooooooooooo

- ▶ *"Look from another angle on the data"*
- ▶ PCA is just a rotation of the coordinate system
- ▶ The rotation is done so that the first axis contains as much variation as possible
- ▶ Second axis than most of remaining variation

Short Demo.

- ▶ *"Look from another angle on the data"*
- ▶ PCA is just a rotation of the coordinate system
- ▶ The rotation is done so that the first axis contains as much variation as possible
- ▶ Second axis than most of remaining variation

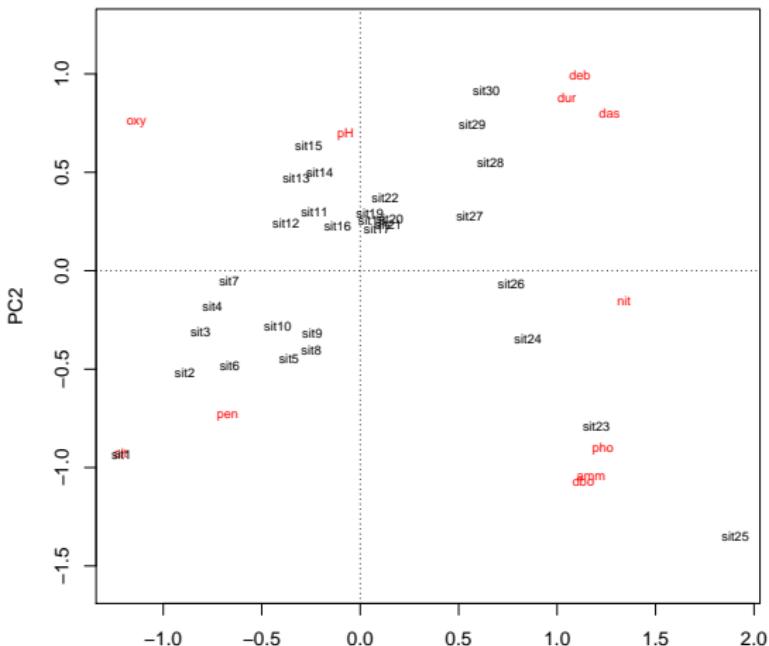
Short Demo.

Maths:

- ▶ The covariance (or correlation) matrix is decomposed into its Eigenvectors and Eigenvalues.
- ▶ The Eigenvectors give the rotation needed
- ▶ The Eigenvalues stretch the axes

Principal Components Analysis (PCA) — How?

```
require(vegan)  
PCA <- rda(Denv, scale = TRUE)  
plot(PCA, scaling = 3)
```



Datasets

Unconstrained Ordination

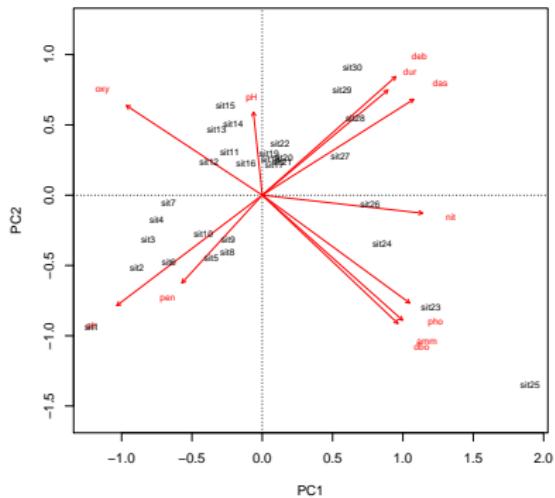
PC1 Constrained Ordination

Model diagnostics / testing

Principal Components Analysis (PCA) — Interpretation? (I)

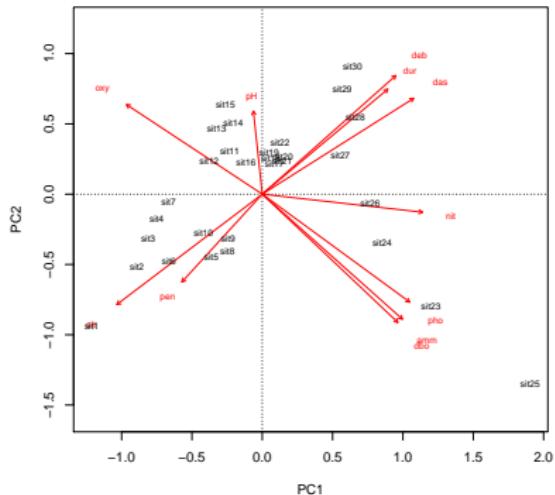
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```
biplot(PCA, cex = 5, scaling = 3)
```

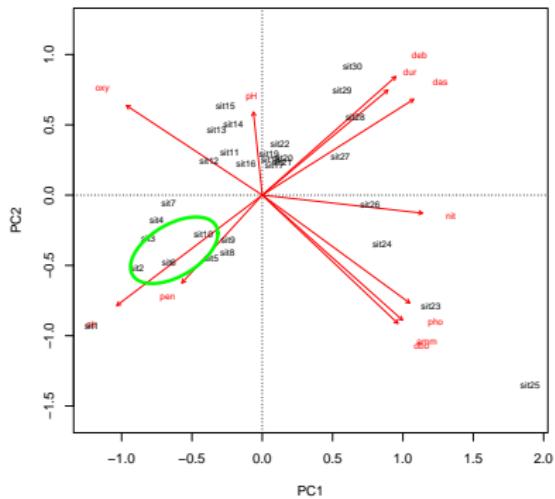


- ▶ angle between variables **approx.** their correlation
- ▶ distance between sites **approx.** their euclidean distance
- ▶ projecting a site on a variable **approx.** the relative value

```
biplot(PCA, cex = 5, scaling = 3)
```



- ▶ angle between variables **approx.** their correlation
 - ▶ distance between sites **approx.** their euclidean distance
 - ▶ projecting a site on a variable **approx.** the relative value
-
- ▶ scaling = 1 - to interpret (only) distances between sites
 - ▶ scaling = 2 - to interpret (only) correlations between variables



- ▶ high altitude + slope, low discharge

Datasets

Unconstrained Ordination



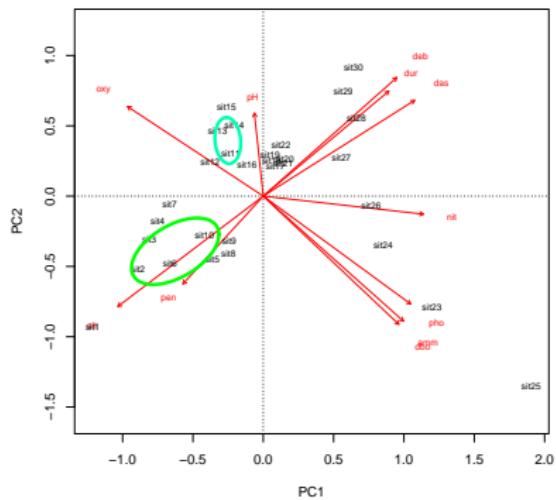
Constrained Ordination



Model diagnostics / testing



Principal Components Analysis (PCA) — Interpretation? (II)



- ▶ high altitude + slope, low discharge
 - ▶ high oxygen, low nutrient

Datasets

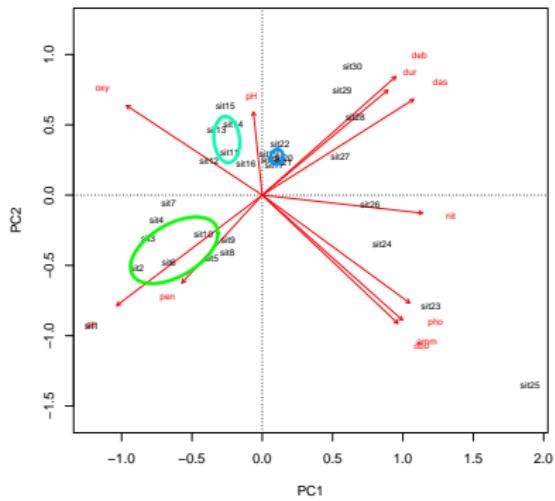
Unconstrained Ordination

Constrained Ordination



Model diagnostics / testing

Principal Components Analysis (PCA) — Interpretation? (II)



- ▶ high altitude + slope, low discharge
 - ▶ high oxygen, low nutrient
 - ▶ intermediate

Datasets

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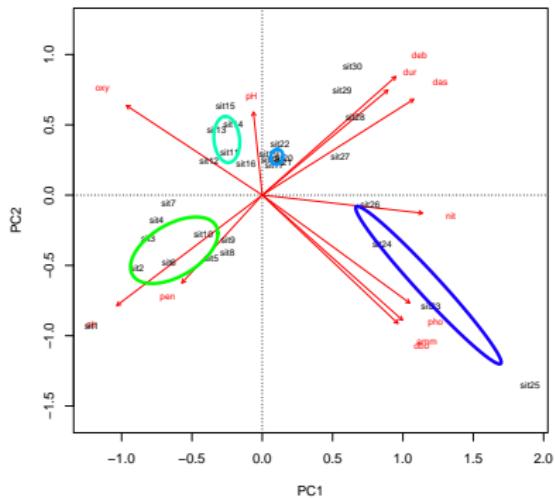
Unconstrained Ordination

Constrained Ordination

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Model diagnostics / testing

Principal Components Analysis (PCA) — Interpretation? (II)



- ▶ high altitude + slope, low discharge
 - ▶ high oxygen, low nutrient
 - ▶ intermediate
 - ▶ nutrient rich

Datasets

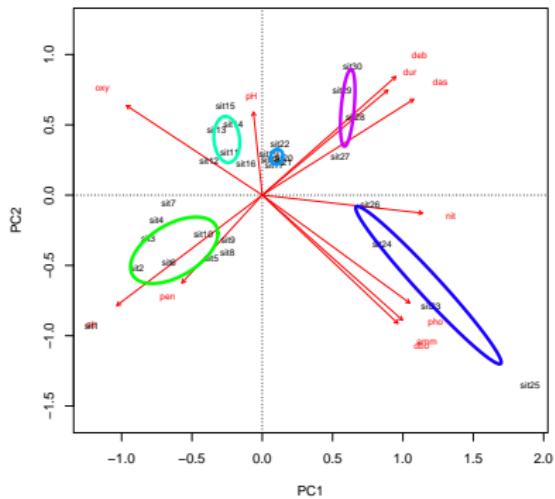
Unconstrained Ordination

Constrained Ordination



Model diagnostics / testing

Principal Components Analysis (PCA) — Interpretation? (II)



- ▶ high altitude + slope, low discharge
 - ▶ high oxygen, low nutrient
 - ▶ intermediate
 - ▶ nutrient rich
 - ▶ high discharge, low altitude, medium nutrient

Datasets

Unconstrained Ordination

Constrained Ordination

Model diagnostics / testing

Principal Components Analysis (PCA) — Interpretation? (III)

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```
summary(PCA, display = NULL, scaling = 3)
```

Call:
rda(X = Denv, scale = TRUE)

Partitioning of correlations:
Inertia Proportion
Total 11 1
Unconstrained 11 1

Eigenvalues, and their contribution to the correlations

Importance of components:

	PC1	PC2	PC3	PC4	PC5	PC6	PC7
Eigenvalue	5.9687	2.1638	1.06516	0.73873	0.40027	0.33565	0.1727
Proportion Explained	0.5426	0.1967	0.09683	0.06716	0.03639	0.03051	0.0157
Cumulative Proportion	0.5426	0.7393	0.83616	0.90331	0.93970	0.97022	0.9859
	PC8	PC9	PC10	PC11			
Eigenvalue	0.10821	0.02368	0.01707	0.005993			
Proportion Explained	0.00984	0.00215	0.00155	0.000540			
Cumulative Proportion	0.99575	0.99790	0.99946	1.000000			

Scaling 3 for species and site scores

- * Both sites and species are scaled proportional to eigenvalues on all dimensions
- * General scaling constant of scores:

Your turn!

Load the Melbourne dataset (only environmental variables).

Exclude the variables ID, logCond and logmaxTU.

Perform a PCA.

Which variables are correlated?

How much variance is explained by the first 2 axes?

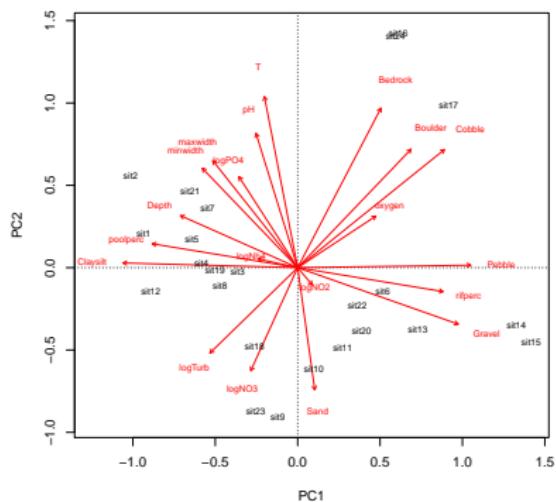
How could the two PCA axes be interpreted?

Exercise

```
take <- env[ , !names(env) %in% c('ID', 'logCond', 'logmaxTU')]
PCA <- rda(take, scale = TRUE)
cumsum(PCA$CA$eig / PCA$tot.chi)[1:2]
```

PC1	PC2
0.2839873	0.4537452

```
biplot(PCA, scaling = 3)
```



- ▶ multiple variables interrelated
- ▶ 1st axis can be interpreted as *hydrological gradient*
- ▶ 2nd axis can be interpreted as *chemistry gradient*

Datasets
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Unconstrained Ordination
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Constrained Ordination
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Model diagnostics / testing
oooooooooooooooo

Question:

- ▶ How is diversity related to salinity, pesticides and other variables?

Datasets

Unconstrained Ordination

Constrained Ordination

Model diagnostics / testing

Question:

- How is diversity related to salinity, pesticides and other variables?

Problem:

- ▶ Only 24 sites
 - ▶ but 22 (potentially correlated) explanatory variables
 - ▶ strong hypotheses about salinity and pesticides

A Solution:

Question:

- ▶ How is diversity related to salinity, pesticides and other variables?

Problem:

- ▶ Only 24 sites
- ▶ but 22 (potentially correlated) explanatory variables
- ▶ strong hypotheses about salinity and pesticides

A Solution:

- ▶ Reduce the number of variables to *Principal Components*
- ▶ regress these

Excursus — principal component regression (PCR)

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```
# calculate shannon diversity index
div <- diversity(abu[, -1], index = 'shannon')
pc <- scores(PCA, choices = c(1, 2), scaling = 1, display = 'sites')
model_data <- data.frame(div, pc, logCond = env$logCond, logmaxTU = env$logmaxTU)
model <- lm(div ~ PC1 + PC2 + logCond + logmaxTU, data = model_data)
summary(model)
```

Call:

```
lm(formula = div ~ PC1 + PC2 + logCond + logmaxTU, data = model_data)
```

Residuals:

Min	1Q	Median	3Q	Max
-0.64415	-0.15688	0.02063	0.18219	0.57929

Coefficients:

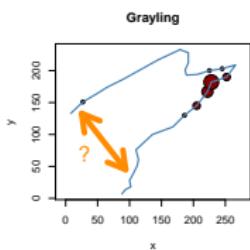
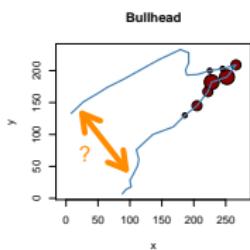
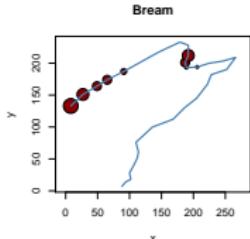
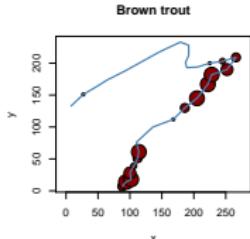
	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	1.83079	0.43429	4.216	0.000468 ***
PC1	0.01971	0.16691	0.118	0.907262
PC2	0.02192	0.19570	0.112	0.911996
logCond	-0.20942	0.13050	-1.605	0.125049
logmaxTU	-0.12572	0.07316	-1.718	0.101994

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 0.3645 on 19 degrees of freedom

Multiple R-squared: 0.2682, Adjusted R-squared: 0.1141

F-statistic: 1.741 on 4 and 19 DF, p-value: 0.1827



- ▶ Species may be absent due to different factors (too high flow, too saline, etc.)
 - ▶ *Absence* contains less information than *Presence*
 - ▶ PCA preserves the euclidean distance between sites
 - ▶ Need another measure of similarity for (raw) abundances

Datasets

Unconstrained Ordination

Constrained Ordination

Model diagnostics / testing

Dissimilarity measures — Species abundance paradox

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	Spe1	Spe2	Spe3
sit1	0	4	8
sit2	0	1	1
sit3	1	0	0

```
vegdist(mat, method = 'euclidean')  
  
      sit1      sit2  
sit2 7.615773  
sit3 9.000000 1.732051
```

```
vegdist(mat, method = 'bray')  
  
      sit1      sit2  
sit2 0.7142857  
sit3 1.0000000 1.0000000
```

from: Anderson, M.J., Crist, T.O., et al. , 2011. Navigating the multiple meanings of beta diversity: a roadmap for the practicing ecologist. *Ecology Letters* 14, 19–28.

Dissimilarity measures — Species abundance paradox

	Spe1	Spe2	Spe3
sit1	0	4	8
sit2	0	1	1
sit3	1	0	0

```
vegdist(mat, method = 'euclidean')
```

```
      sit1      sit2
sit2 7.615773
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```

```
vegdist(mat, method = 'bray')
```

```
      sit1      sit2
sit2 0.7142857
sit3 1.0000000 1.0000000
```

Abundances included
(or cover, biomass, etc.)

Exclude joint absences

Include joint absences

Chi-squared
Hellinger, Chord
Euclidean on proportions
Bray-Curtis, fourth root
Bray-Curtis, square root
Bray-Curtis, raw data
Modified Gower, log10
Modified Gower, log5
Modified Gower, log2

Euclidean, log(y+1)
Euclidean, raw data

Emphasis:

Rare species

Proportions

Composition + relative abund.

Composition

log abundance

Log abundance

Raw abundance

from: Anderson, M.J., Crist, T.O., et al. , 2011. Navigating the multiple meanings of beta diversity: a roadmap for the practicing ecologist. Ecology Letters 14, 19–28.

- ▶ Works on distance matrices
- ▶ Species can be added as *weighted averages*
- ▶ Eigenvalue based
- ▶ PCoA with euclidean distance == PCA

Datasets
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Unconstrained Ordination
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Constrained Ordination
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Model diagnostics / testing
oooooooooooo

Principal coordinates analysis (PCoA)

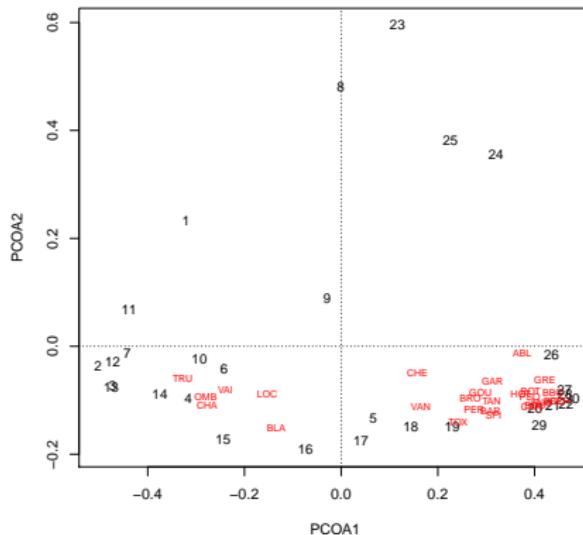
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```
# Distance matrix
Dabu_dist <- vegdist(Dabu, method = 'bray')

# PCoA
PCOA <- cmdscale(Dabu_dist, eig = TRUE)

# Create plot
plot(PCOA$points, type = 'n',
      xlab = 'PCOA1', ylab = 'PCOA2')
text(PCOA$points,
     labels = rownames(Dabu), cex = 0.9)
abline(h = 0, lty = 'dotted')
abline(v = 0, lty = 'dotted')
# Add species as weighted averages
wa <- wascores(PCOA$points, Dabu)
text(wa, labels = colnames(Dabu),
     col = 'red', cex = 0.7)
```

```
# explained variance
(PCOA$eig / sum(PCOA$eig))[1:2] * 100
[1] 49.24914 15.95758
```



Datasets
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Unconstrained Ordination
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Constrained Ordination
oooooooooooo

Model diagnostics / testing
oooooooooooo

- ▶ Similar to PCoA
- ▶ Does not preserve exact distances between objects
- ▶ Possibly better representation in low dimensions
- ▶ **Not** eigenvalue based, iterative algorithm
- ▶ Axes have no meaning, just the relative distances

Datasets
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Unconstrained Ordination
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Constrained Ordination
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Model diagnostics / testing
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Nonmetric Multidimensional Scaling (NMDS)

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```
# Distance matrix
Dabu_0 <- Dabu[!rowSums(Dabu) == 0, ]
Dabu_dist <- vegdist(Dabu_0, method = 'bray')

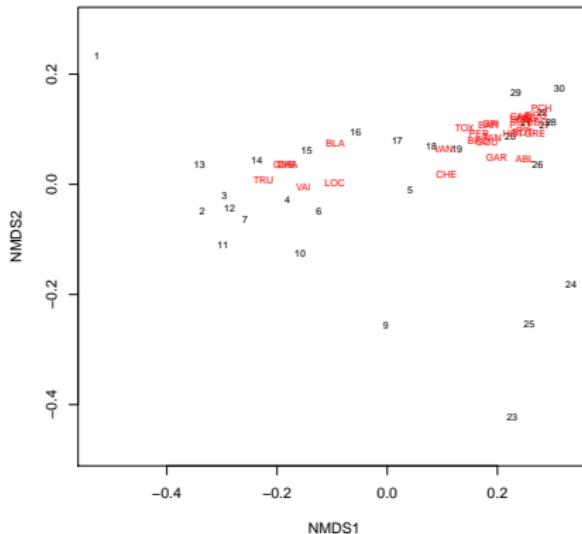
# NMDS
NMDS <- metaMDS(Dabu_dist, k = 2, trace = 0)

# Plot
plot(NMDS, type = 't')

# Add species as weighted averages
wa <- wascores(NMDS$points, Dabu_0)
text(wa, labels = colnames(Dabu),
     col = 'red', cex = 0.7)
```

```
# Stress value
NMDS$stress

[1] 0.07376347
```



Your turn!

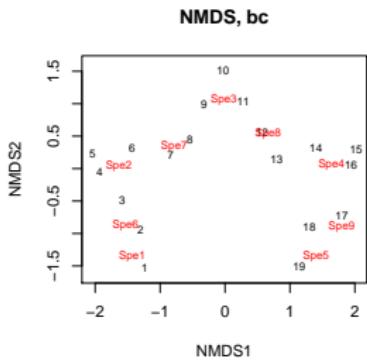
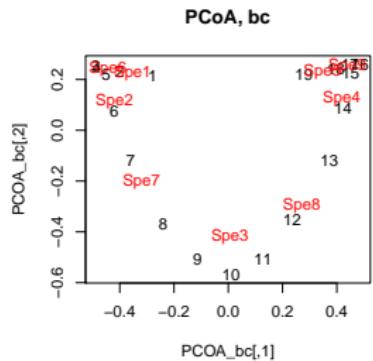
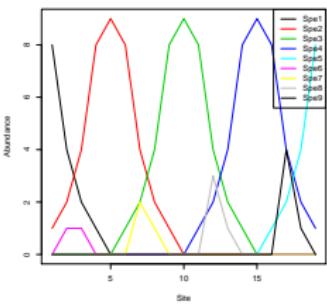
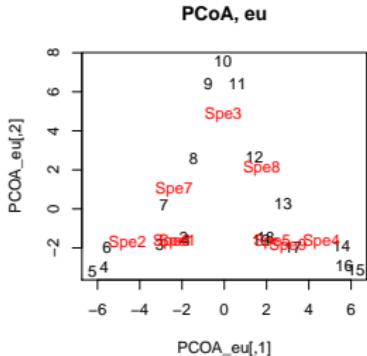
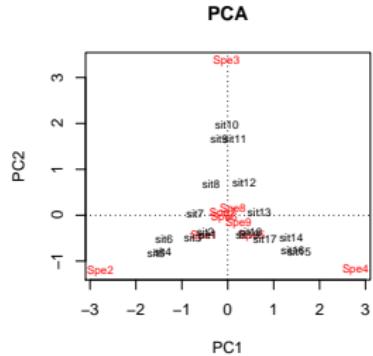
Using the artificial dummy dataset.

Run:

1. PCA
2. PCoA with euclidean distance
3. PCoA with Bray-Curtis dissimilarity
4. NMDS with Bray-Curtis dissimilarity

What are the differences between ordinations?
Which represent better the underlying gradient?

Exercise



Datasets

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Unconstrained Ordination

Constrained Ordination

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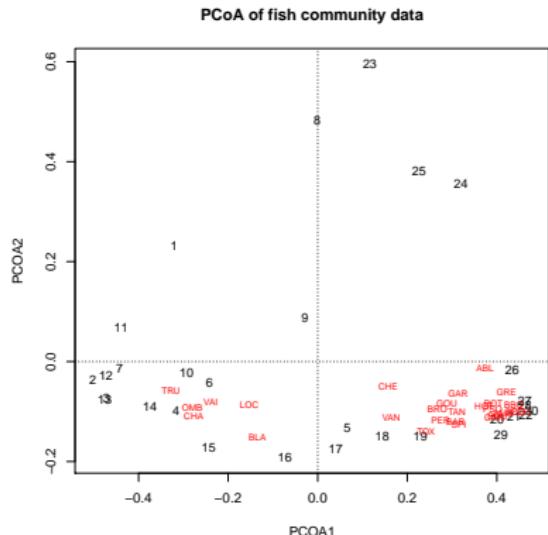
Model diagnostics / testing

- This ordination is **only** driven by fish community data

Question:

- How can we interpret the gradients in community composition?

A solution:



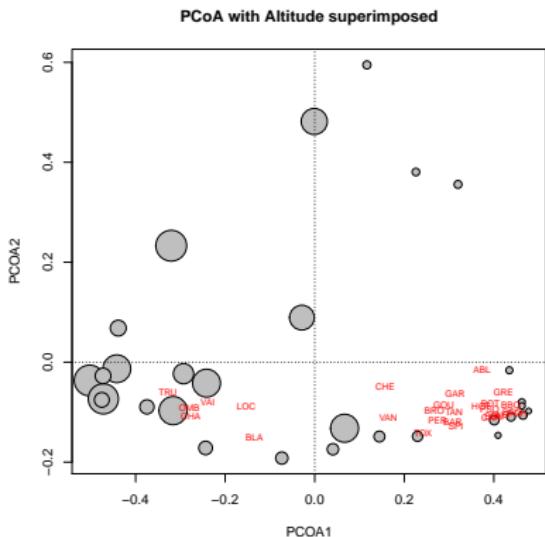
- ▶ This ordination is **only** driven by fish community data

Question:

- ▶ How can we interpret the gradients in community composition?

A solution:

- ▶ Superimpose environmental variables

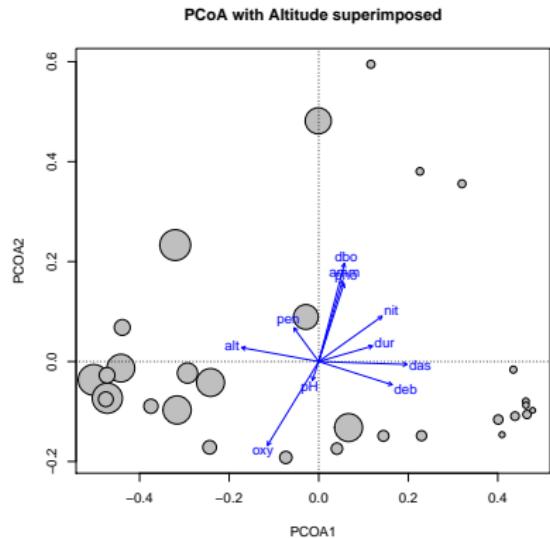


Fit environmental variables to ordination (II)

```
# PCoA of fish community data
plot(PCoA$points,
     xlab = 'PCoA1', ylab = 'PCoA2',
     cex = 5*Denv$alt / max(Denv$alt),
     main = 'PCoA with Altitude',
     bg = 'grey75', pch = 21)
abline(h = 0 , lty = 'dotted')
abline(v = 0 , lty = 'dotted')

# Fit Altitude to site-scores
ef <- envfit(PCoA, Denv)
plot(ef)
ef # summary

# Fit GAM
ordisurf(PCoA, Denv$alt, add = TRUE)
```



- ▶ Post hoc method
- ▶ non-linearity?

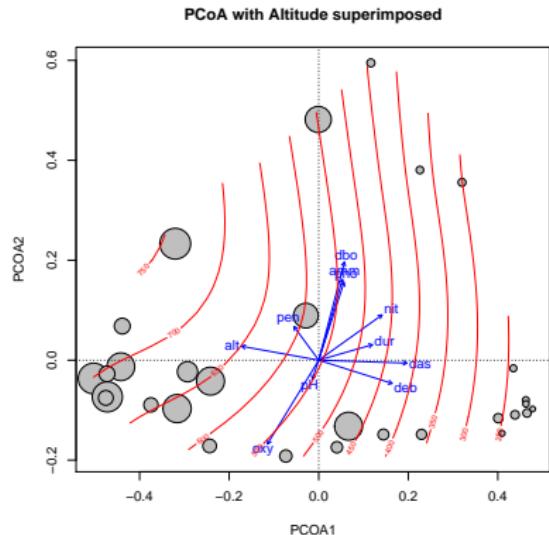
Fit environmental variables to ordination (II)

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```
# PCoA of fish community data
plot(PCOA$points,
     xlab = 'PCOA1', ylab = 'PCOA2',
     cex = 5*Denv$alt / max(Denv$alt),
     main = 'PCoA with Altitude',
     bg = 'grey75', pch = 21)
abline(h = 0 , lty = 'dotted')
abline(v = 0 , lty = 'dotted')

# Fit Altitude to site-scores
ef <- envfit(PCOA, Denv)
plot(ef)
ef # summary

# Fit GAM
ordisurf(PCOA, Denv$alt, add = TRUE)
```



- ▶ Post hoc method
- ▶ non-linearity?
- ▶ be careful with `summary`
- ▶ Constrained ordination a better alternative

Datasets
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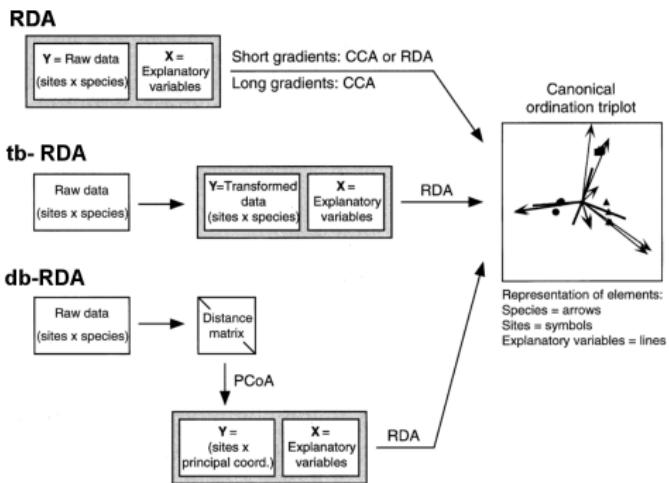
Unconstrained Ordination
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Constrained Ordination
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Model diagnostics / testing
oooooooooooo

Constrained Ordination

- ▶ Redundancy analysis (RDA)
- ▶ Transformation-based RDA (tb-RDA)
- ▶ Distance-based RDA (db-RDA)



Adapted from: Legendre, P., Gallagher, E.D., 2001. Ecologically meaningful transformations for ordination of species data. *Oecologia* 129, 271–280.

- ▶ Associates both environmental and community data at once
- ▶ Combination of regression and PCA:
 1. Regress explanatory variables on community data
 2. Run PCA on fitted values
- ▶ Can test hypothesis about relationships

Datasets
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Unconstrained Ordination
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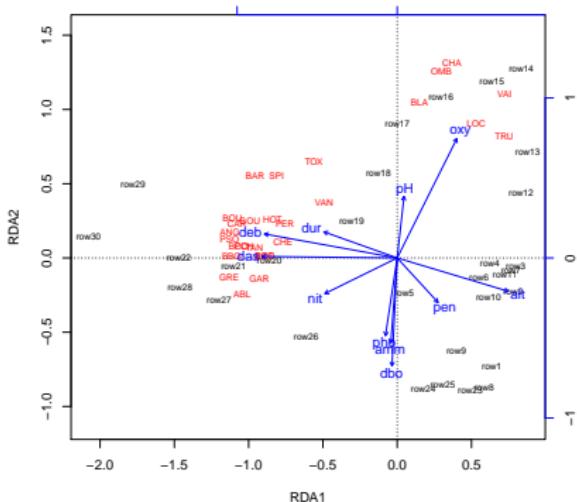
Constrained Ordination
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Model diagnostics / testing
oooooooooooo

```
RDA <- rda(Dabu ~ ., data = Denv,
             scale = TRUE)
plot(RDA, scaling = 3)
```

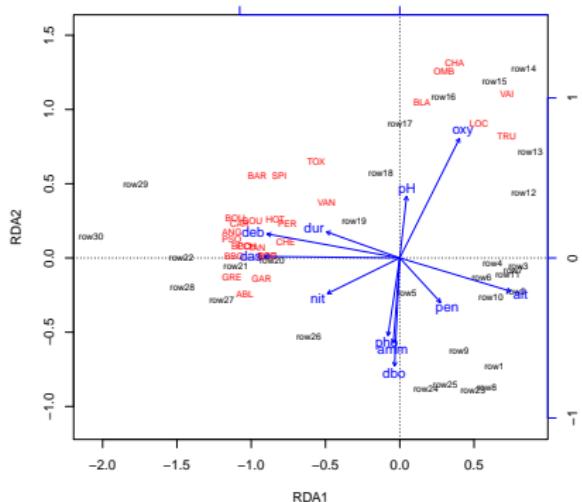
► Formula interface

- Left side: Response matrix
- Right side: Response variables from Denv



Redundancy analysis (RDA) — Interpretation? (I)

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- ▶ Similar to PCA
- ▶ Projecting a site on response or explanatory variable **approx.** the value
- ▶ angles between response and expl. variables **approx.** their correlations

```
...
```

Partitioning of correlations:

	Inertia	Proportion
Total	27.000	1.0000
Constrained	20.177	0.7473
Unconstrained	6.823	0.2527

Eigenvalues, and their contribution to the correlations

Importance of components:

	RDA1	RDA2	RDA3	RDA4	RDA5	RDA6	RDA7
Eigenvalue	14.714	2.6433	1.1341	0.76821	0.33807	0.28135	0.09356
Proportion Explained	0.545	0.0979	0.0420	0.02845	0.01252	0.01042	0.00347
Cumulative Proportion	0.545	0.6429	0.6849	0.71333	0.72585	0.73627	0.73974
	RDA8	RDA9	RDA10	RDA11	PC1	PC2	
Eigenvalue	0.08411	0.07592	0.02314	0.02129	2.44703	1.4094	
Proportion Explained	0.00312	0.00281	0.00086	0.00079	0.09063	0.0522	
Cumulative Proportion	0.74285	0.74566	0.74652	0.74731	0.83794	0.8901	

```
...
```

- RDA (as PCA) preserves the euclidean distance.

What about the species abundance paradox?

Legendre, P., Gallagher, E.D., 2001. Ecologically meaningful transformations for ordination of species data. *Oecologia* 129, 271–280.

Datasets
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Unconstrained Ordination
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Constrained Ordination
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Model diagnostics / testing
oooooooooooo

- ▶ RDA (as PCA) preserves the euclidean distance.

What about the species abundance paradox?

- ▶ Can transform data to use with euclidean distance
 - ▶ Remove differences in total abundance, while keeping the variations relative abundance
- ▶ Chord and Hellinger transformations useful.

Hellinger:

$$y'_{ij} = \sqrt{\frac{y_{ij}}{\sum_{j=1}^p y_{ij}}}$$

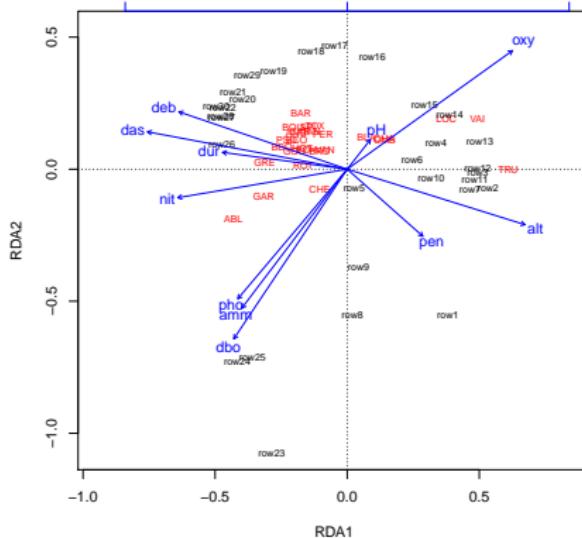
```
mat
  Spe1  Spe2  Spe3
sit1    0     4     8
sit2    0     1     1
sit3    1     0     0

decostand(mat, 'hellinger')

  Spe1      Spe2      Spe3
sit1    0 0.5773503 0.8164966
sit2    0 0.7071068 0.7071068
sit3    1 0.0000000 0.0000000
attr(,"decostand")
[1] "hellinger"
```

Legendre, P., Gallagher, E.D., 2001. Ecologically meaningful transformations for ordination of species data. Oecologia 129, 271–280.

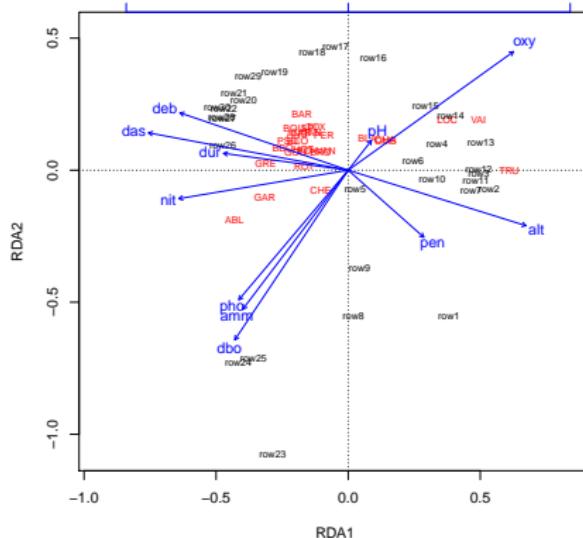
```
# Hellinger transformation  
Dabu_h <- decostand(Dabu, 'hellinger')  
# RDA on Hellinger transformed abundances  
tbRDA <- rda(Dabu_h ~ ., data = Denv)  
  
# Plot  
plot(tbRDA, type = 't')
```



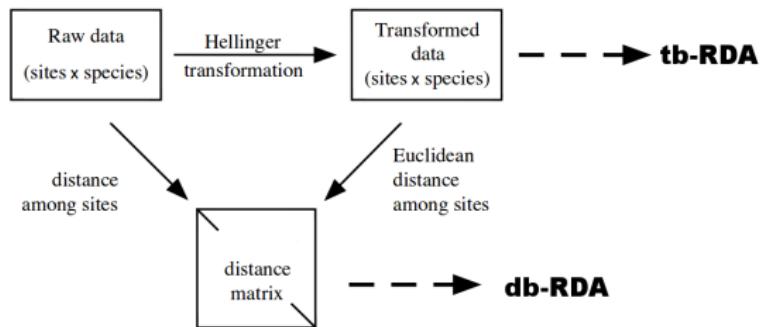
```
# Hellinger transformation
Dabu_h <- decostand(Dabu, 'hellinger')
# RDA on Hellinger transformed abundances
tbRDA <- rda(Dabu_h ~ ., data = Denv)

# Plot
plot(tbRDA, type = 't')
```

- ▶ alt, oxy and nutrients important
- ▶ Trout (TRU) and minow (VAI) found at high sites with high oxygen
- ▶ Bleak (ABL) is found at low oxygen and high nutrients
- ▶ Other species in similar environments



- db-RDA is a related method
- hellinger transformation can also be expressed as distance matrix
- *Constrained PCoA*
- Can use every distance metric



modified from: Legendre, P., Gallagher, E.D., 2001. Ecologically meaningful transformations for ordination of species data. *Oecologia* 129, 271–280.

distance-based RDA

```
# dbRDA
dbRDA <- capscale(Dabu ~ ., data = Denv,
                     distance = 'bray')
plot(dbRDA, type = 't')
```

```
summary(dbRDA)
```

...

Partitioning of squared Bray distance:

Inertia Proportion

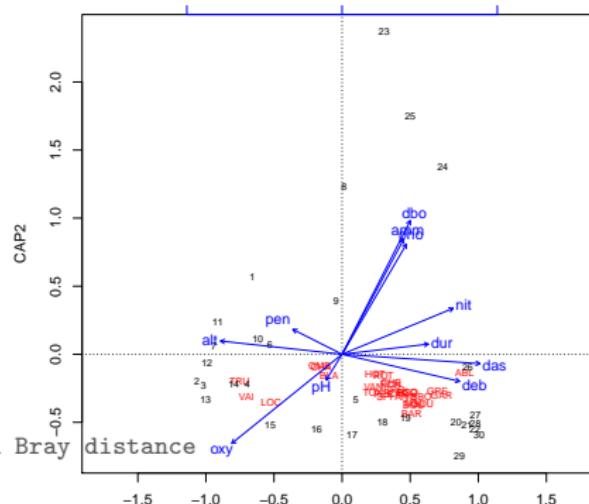
Total	7.802	1.0000
Constrained	5.715	0.7325
Unconstrained	2.087	0.2675

Eigenvalues, and their contribution to the squared Bray distance

Importance of components:

	CAP1	CAP2	CAP3	CAP4	CAP5	CAP6	CAP1
Eigenvalue	3.2627	1.0283	0.53633	0.37863	0.24913	0.10755	
Proportion Explained	0.4182	0.1318	0.06874	0.04853	0.03193	0.01379	
Cumulative Proportion	0.4182	0.5500	0.61875	0.66729	0.69922	0.71300	

...



Your turn!

Using the artificial dummy dataset and Site as only constraining variable

Run:

1. RDA
2. tbRDA (Hellinger)
3. dbRDA with Bray-Curtis
4. dbRDA with $x^{0.25}$ transformed abundances and Bray-Curtis

What ordination presents best the gradient?

What method explains most of variance?

See Demo.

Datasets
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Unconstrained Ordination
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Constrained Ordination
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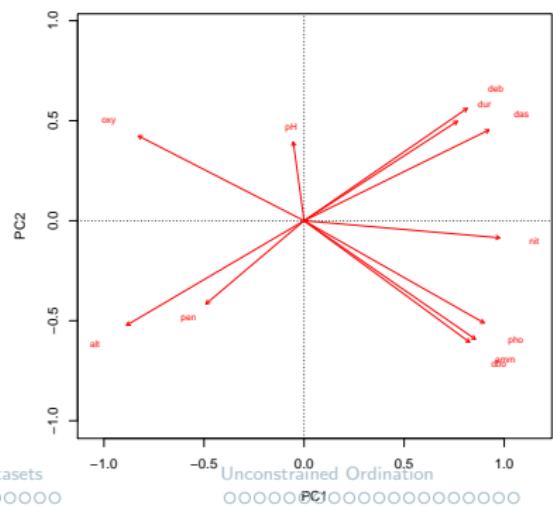
Model diagnostics / testing
oooooooooooo

Model diagnostics and testing

Collinearity

- RDA et al. are *regression methods* - everything you know applies also here
- Collinearity of predictors may lead to wrong conclusions.
- Many methods available (see ref.). Additional: *Use your ecological knowledge!*

```
# visual inspection via PCA
biplot(rda(Denv, scale = TRUE),
       display = 'species', scaling = 2)
```



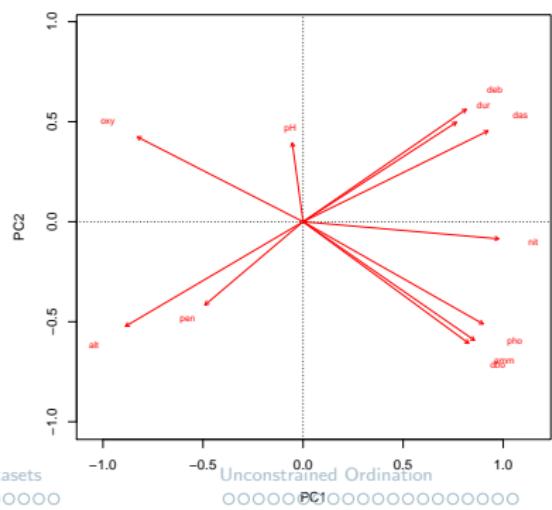
Constrained Ordination
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Model diagnostics / testing
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Collinearity

- RDA et al. are *regression methods* - everything you know applies also here
- Collinearity of predictors may lead to wrong conclusions.
- Many methods available (see ref.). Additional: *Use your ecological knowledge!*

```
# visual inspection via PCA
biplot(rda(Denv, scale = TRUE),
       display = 'species', scaling = 2)
```



```
# RDA on Hellinger transformed abundances
tbRDA <- rda(decostand(Dabu, "hellinger") ~ .,
# Variance inflation factors
vif.cca(tbRDA)
```

	das	alt	pen	deb
104.008842	39.589948	2.023986	36.048637	
pho	nit	amm	oxy	
25.329286	17.284287	31.099024	12.963327	17

- many different cutoffs (see 1st part)
- very high values ($\gg 20$)
- strong collinearity

- ▶ cumulative variance explained by constraints
- ▶ available for *sites* or *species*
- ▶ `summarize = TRUE` gives the accumulated total variance (= last column)

```
# GOF for each species on  
goodness(tbRDA)[ , 1:3]
```

	RDA1	RDA2	RDA3
CHA	0.16597466	0.28222282	0.72531373
TRU	0.68728597	0.68728738	0.75205565
VAI	0.66278142	0.76121936	0.77215444
LOC	0.50167024	0.63319561	0.63433575
OMB	0.16050386	0.25551654	0.64853919
BLA	0.04766735	0.17402885	0.47152162
...			

```
# total var explained by constraints  
goodness(tbRDA, summarize = TRUE)[1:6]
```

	CHA	TRU	VAI	LOC	OMB	BLA
	0.8010478	0.8798422	0.8676846	0.8223702	0.7889173	0.5669931

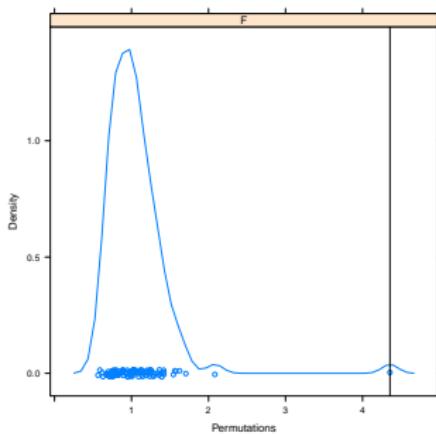
- ▶ can decompose variance into constrained (CCA) and unconstrained (CA) parts

```
# variance explained by constrained (CCA) and unconstrained (CA) axes  
inertcomp(tbRDA, proportional = TRUE)
```

	CCA	CA
CHA	0.8010478	0.1989522
TRU	0.8798422	0.1201578
VAI	0.8676846	0.1323154
LOC	0.8223702	0.1776298
OMB	0.7889173	0.2110827
BLA	0.5669931	0.4330069
...		

- ▶ Cannot use parametric test theory
- ▶ Use of *permutation tests* :
 1. Shuffle the data (H_0 : No effect)
 2. Fit model to shuffled data
 3. Compute (pseudo-)F statistic for each model (Null distribution if H_0 is true)
 - 4.

$$p = \frac{(\text{No. of } F_{\text{perm}} \geq F) + 1}{\text{Total No. of } F_{\text{perm}} + 1}$$



A number of different test can be applied to RDA:

- ▶ Test overall significance of model
- ▶ Test RDA axes
- ▶ Test terms:
 - ▶ sequential
 - ▶ marginal

I omit some variables due to collinearity:

```
# RDA on Hellinger transformed abundances
tbRDA <- rda(decostand(Dabu, 'hellinger') ~ alt + oxy + pH + nit + pho,
               data = Denv)
vif.cca(tbRDA)

      alt      oxy      pH      nit      pho
2.929759 2.198671 1.047617 6.577783 4.164368
```

```
# Tests if the overall model is significant
anova(tbRDA)
```

Permutation test for rda under reduced model

Permutation: free

Number of permutations: 999

```
Model: rda(formula = decostand(Dabu, "hellinger") ~ alt + oxy + pH + nit + pho, data = Denv)
        Df Variance     F Pr(>F)
Model      5  0.28776 6.4372  0.001 ***
Residual 24  0.21458
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
# Tests RDA axes
anova(tbRDA, by = 'axis')

Permutation test for rda under reduced model
Marginal tests for axes
Permutation: free
Number of permutations: 999

Model: rda(formula = decostand(Dabu, "hellinger") ~ alt + oxy + pH + nit + pho, data = Dent)
        Df Variance      F Pr(>F)
RDA1     1 0.212483 23.7658  0.001 ***
RDA2     1 0.046671  5.2201  0.001 ***
RDA3     1 0.020291  2.2695  0.029 *
RDA4     1 0.006385  0.7141  0.675
RDA5     1 0.001935  0.2165  0.998
Residual 24 0.214578
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Legendre, P., Oksanen, J. and ter Braak, C.J.F. (2011). Testing the significance of canonical axes in redundancy analysis. Methods in Ecology and Evolution 2, 269–277.

- ▶ Variables are tested in the order they were specified (first to last)
- ▶ Test of additional variance explained by adding the variable to the model
- ▶ Order matters!

```
# Tests RDA axes
anova(tbRDA, by = 'terms')

Permutation test for rda under reduced model
Terms added sequentially (first to last)
Permutation: free
Number of permutations: 999

Model: rda(formula = decostand(Dabu, "hellinger") ~ alt + oxy + pH + nit + pho, data = Den)
        Df Variance      F Pr(>F)
alt      1 0.153953 17.2193 0.001 ***
oxy      1 0.085314  9.5422 0.001 ***
pH       1 0.004285  0.4793 0.814
nit      1 0.007642  0.8547 0.489
pho      1 0.036571  4.0904 0.006 **
Residual 24 0.214578
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

- ▶ Test explained variance of variable when all other variable are included in the model
- ▶ Order has no influence

```
# Tests RDA axes
anova(tbRDA, by = 'margin')

Permutation test for rda under reduced model
Marginal effects of terms
Permutation: free
Number of permutations: 999

Model: rda(formula = decostand(Dabu, "hellinger") ~ alt + oxy + pH + nit + pho, data = Dent)
        Df Variance      F Pr(>F)
alt      1 0.031774 3.5539  0.020 *
oxy      1 0.067640 7.5653  0.001 ***
pH       1 0.003905 0.4367  0.862
nit      1 0.011760 1.3153  0.238
pho      1 0.036571 4.0904  0.010 **
Residual 24 0.214578
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Multivariate topics not covered here:

- ▶ Model selection (be careful with automatic methods!)
- ▶ Distance-based hypothesis testing ((PER-)MANOVA, SIMPER, ANOSIM)
- ▶ Dispersion measures (β -Diversity, Functional diversity)
- ▶ Consensus RDA, RLQ (traits), ...
- ▶ Model-based multivariate framework (See work of David Warton et al.)
- ▶ manymore