

Analysing Mesocosm Experiments - Principal Response Curves

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Mesocosm studies

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Curves

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Interpretation
Alternatives
How to

- ▶ Between laboratory and field studies (relevance and control)
- ▶ Higher Tier Tests in ecological risk assessment
- ▶ Effects on community structure and functioning



IGB, LakeLab, Lake Stechlin, Brandenburg

Mesocosm data

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How to

General experimental setup:

- ▶ Mesocosms treated with test item
- ▶ Communities sampled before and after exposure
- ▶ Multiple timepoints (recovery?)

This yields:

- ▶ large and complex data sets
- ▶ multiple species
- ▶ many time points

Multivariate techniques are needed:

- ▶ analyse
- ▶ summarise
- ▶ interpret

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Aquatic Ecology 32: 163–178, 1998.
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Multivariate analysis of stress in experimental ecosystems by Principal Response Curves and similarity analysis

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Wageningen, CPRO-DLO, P.O. Box 16, 6700 AA Wageningen, The Netherlands)

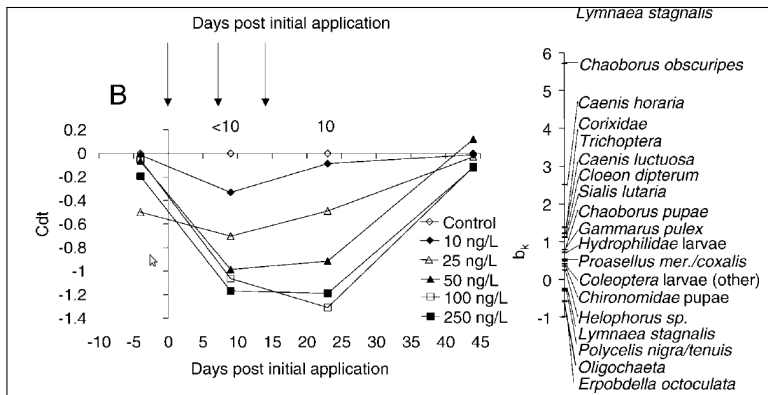


Accepted 4 June 1998

Key words: aquatic, chlorpyrifos, microcosm, nutrients, principal component analysis, similarity index

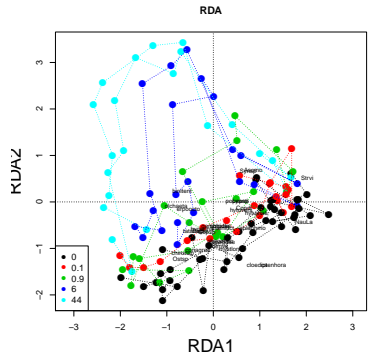
Abstract

Experiments in microcosms and mesocosms, which can be carried out in an advanced tier of risk assessment, usually result in large data sets on the dynamics of biological communities of treated and control cosms. Multi-



RDA:

- ▶ treatment, time , treatment x time
- ▶ cluttered biplot
- ▶ can spot effects, but not obvious
- ▶ effect of time?



RDA?

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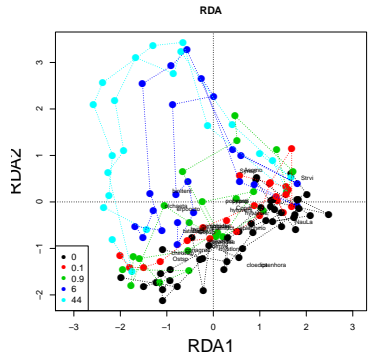
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RDA:

- ▶ treatment, time , treatment x time
- ▶ cluttered biplot
- ▶ can spot effects, but not obvious
- ▶ effect of time?



Need a better way to represent the treatment effects!

partial RDA?

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How to

partial RDA:

- ▶ Not interested in the pure time effect
- ▶ remove time effect (*partial out*)

partial RDA?

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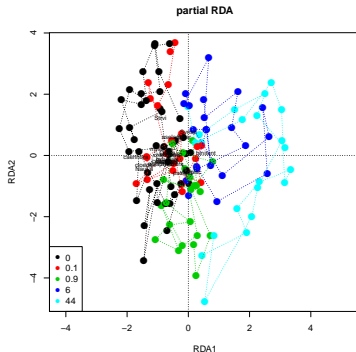
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How to

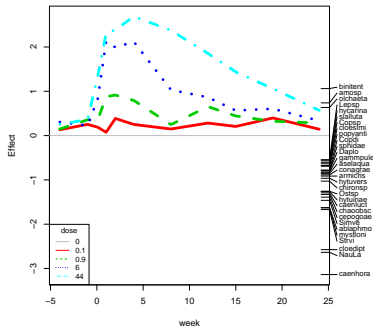
partial RDA:

- ▶ Not interested in the pure time effect
- ▶ remove time effect (*partial out*)
- ▶ treatment + treatment x time on first axis
- ▶ first axis better separates treatments
- ▶ still cluttered
- ▶ time course?



PRC:

- ▶ **1st axis**
- ▶ site scores as contrast to control
- ▶ species scores
- ▶ easy plot
- ▶ easy to interpret



Summary

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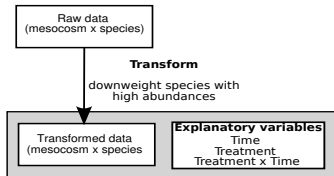
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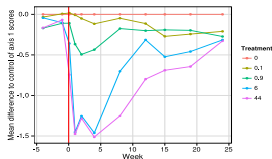
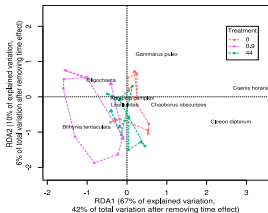
How to



Partial RDA:

- 1) 'partial out' time effect
- 2) Fit RDA model

= axes display effect of Treatment and Treatment x Time



A) Site scores

Site scores on first axis as contrast to control

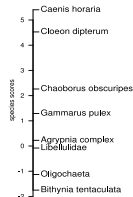
= Effect of treatment along time

B) Species scores

Species scores on first axis

= species responsible for the pattern in A)

C) Significance test



Permutation tests of first axis

= axis displays statistically significant amount of variation

Numerical output

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How to

- ▶ How much variation is displayed on first axis?
- ▶ Significance testing:
 - ▶ of first axis
 - ▶ of predictors (treatment x time)
 - ▶ need to take repeated measures into account
- ▶ Recovery?
 - ▶ separate RDA per time point
- ▶ NOEC?
 - ▶ poorly replicated
 - ▶ cannot use permutations

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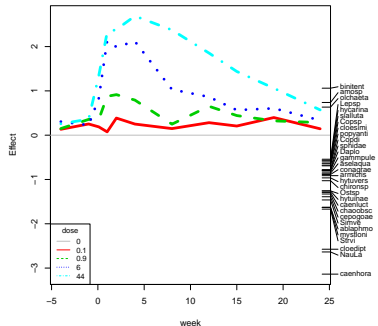
How to

Site scores:

- ▶ Date on **x-axis**
- ▶ deviations in composition between treatments and control on the **y-axis**
- ▶ spot recovery (treatment x time)

Species scores

- ▶ species responsible for the pattern
- ▶ increase / decrease



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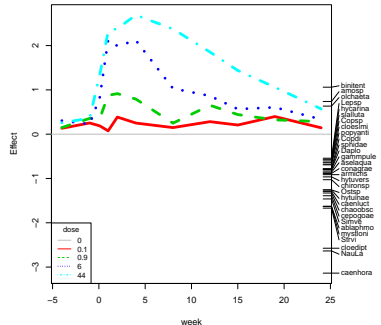
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Caveats

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How to

- ▶ low species scores not necessarily mean low response!
 - ▶ species associated with main pattern
 - ▶ special responses (no recovery)
- ▶ Need to transform abundance data (often $\ln(Ax + 1)$)

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- ▶ (multivariate) GLM (Warton 2012)
 - ▶ new method, can model counts
 - ▶ advantages, but also some disadvantages (e.g. computation time, complexity)
- ▶ Univariate indicators?
 - ▶ SPEAR, diversity, ...
 - ▶ univariate solution to a multivariate problem
 - ▶ what about species responses?

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Lets do it in R!

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*use***R**!