

Corina Carpentier

***In situ* Biomass Quantification of Microphyto-  
benthos: a First Step Towards an Alternative  
Method for Assessing Stream Eutrophication  
under the WFD**

AquaLife Workshop  
4-6 June  
Kiel, Germany



# Introduction

- ❑ WFD and microphytobenthos in rivers and streams
- ❑ Commonly used methods
  - EN 13946 / EN 14407
  - EN 15708
- ❑ *In situ* fluorescence measurements
- ❑ Nutrient – biomass relationships



# WFD - Microphytobenthos

- ❑ Indicator for eutrophication in rivers and streams
- ❑ Organisms:
  - sessile
  - take up nutrients from water-phase
  - respond rapidly to changing conditions
- ❑ Ecology:
  - support lotic food webs – food source for grazers
  - attenuate current
  - stabilise sediments



# Microphytobenthos Analysis

## □ Biodiversity assessment

- EN 13946 / EN 14407: benthic diatoms
- EN 15708: phytobenthos

## □ Calculation of index values:

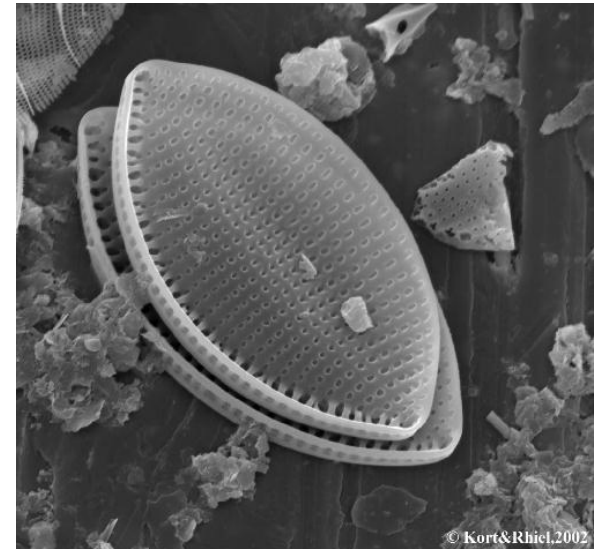
- diversity index: disturbance/stress
- saprobic index: organic pollution

## □ Additionally:

- evenness index: even distribution of abundance

# Species Diversity Assessment

- ❑ Hungary: 6500 diatom species
- ❑ Germany: 1000 phytobenthos species
- ❑ France: 6500 diatom species
- ❑ Poland: 520 diatom species
- ❑ Czech Republic:
  - 252 blue-green algae species
  - 521 diatom species
  - 990 green algae species
  - 750 chromophyte species
  - 324 flagellate species
  - Total: 2,837 species



Diatomee (Kieselalge)

10µm



# WFD Implementation

- Austria:
  - experts at eco-region level required
  - official registry of biological experts: 6
  - high quality guaranteed
- Biodiversity analysis very difficult as *routine* method
- > 120 river basins in Europe
  - minimum frequency: once every 3 years
  - provide “sufficient data for reliable assessment”
  - exclude seasonal and geographical variation



# Alternative Method

- Suitable for routine analysis
- Practical
- Fast
- Affordable



# Sampling Errors



substrate	before ( $\mu\text{g}/\text{cm}^2$ )	after ( $\mu\text{g}/\text{cm}^2$ )	removal
cyanobacteria	1.20	0.45	62.4%
diatoms	0.17	0.15	13.4%

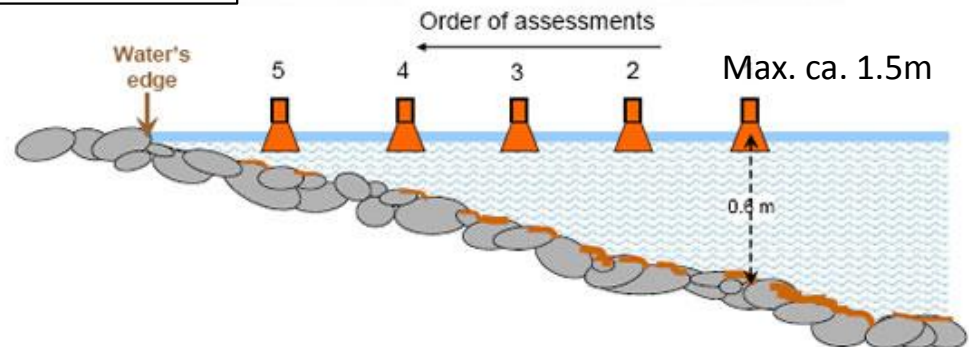
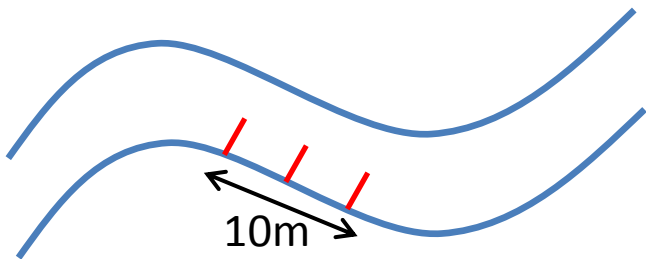
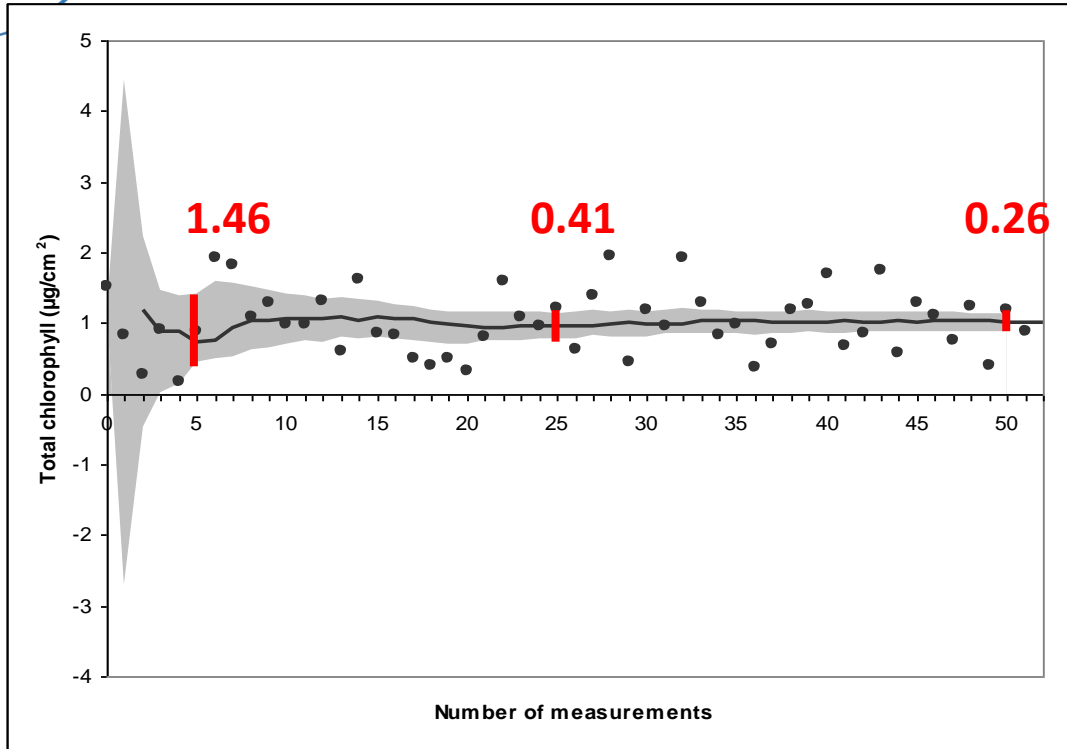




# *In Situ* Fluorescence Method

- ❑ Easy and fast
- ❑ Large transects, in situ measurements
- ❑ General information on all algal groups
- ❑ No detailed information at species level
- ❑ Substratum variability solved by 700nm LED
- ❑ To be studied:
  - dark adaptation?
  - representativeness – patchy distribution

# Representativeness





# Sampling Sites

- Aim of study
- Select sites representative of river-stretch under investigation
- Avoid heavily shaded sites
- Check similar conditions for
  - light
  - current velocity
  - substratum
- Survey units of similar length (10m)
- Consider seasonal run-off

# Microphytobenthos Biomass

- Current velocity
- Frequency of biomass-scouring floods
- Suspended sediment
- Shading
- Substratum type
- Grazing



# Nutrient-Biomass Relationships

## ☐ Lakes:

- nutrient loading
- average residence time

## ☐ Streams and Rivers:

- nutrient supply
- frequency of flood disturbance
- (if both low: grazing)





# Example from Australia/USA

$$B^* = k_1 d_a + k_2 n + c$$

$B^*$  : mean monthly biomass of benthic algae

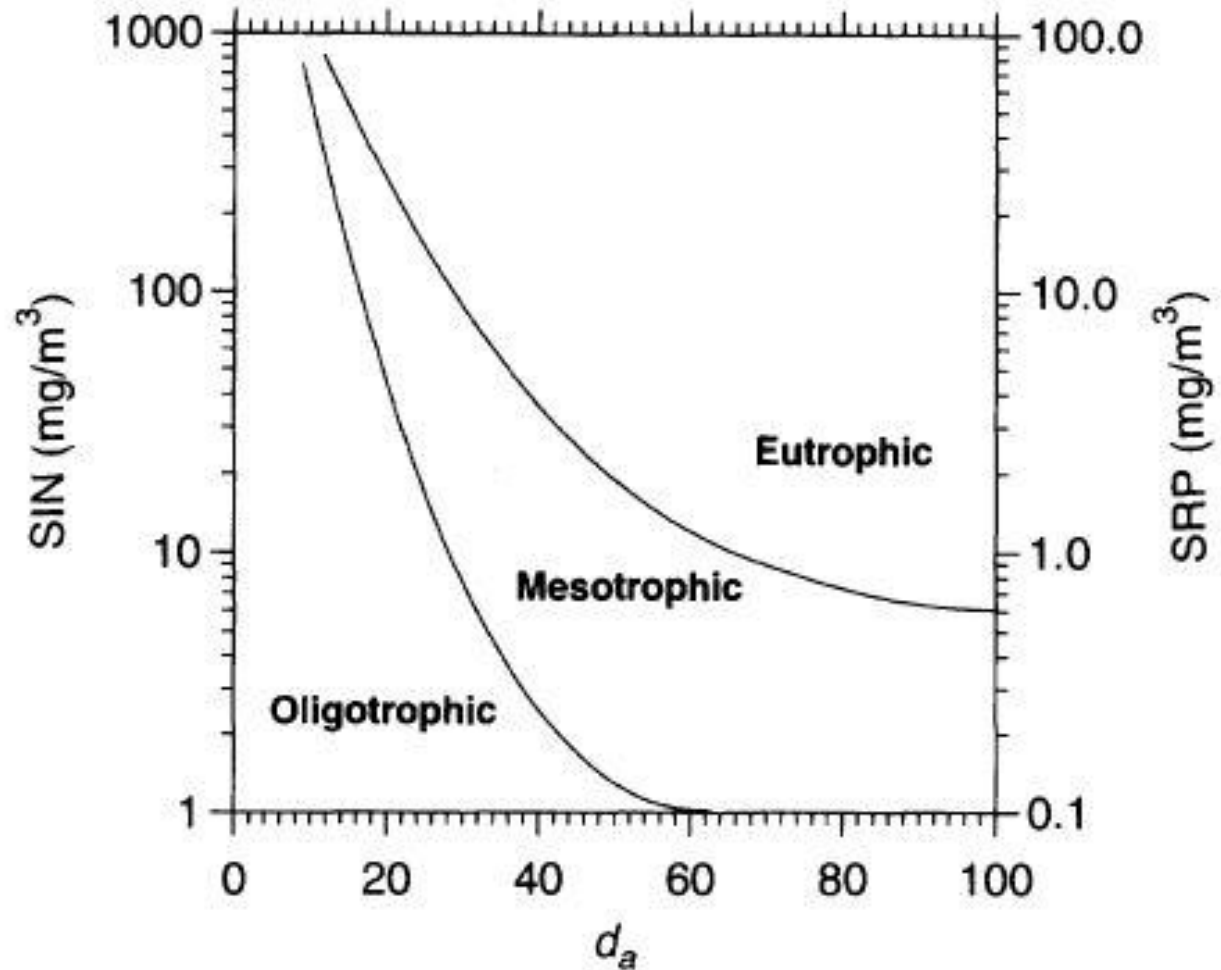
$k_1/k_2$  : coefficients

$d_a$  : number of days available for biomass accrual

$n$  : measure of nutrient supply (mean monthly SRP/SIN)

$c$  : empirical constant

# Conditions resulting in max. phyto**benthos** biomass



O-M:  $6 \mu\text{g}/\text{cm}^2$

M-E:  $20 \mu\text{g}/\text{cm}^2$



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