

bbe

moldaenke

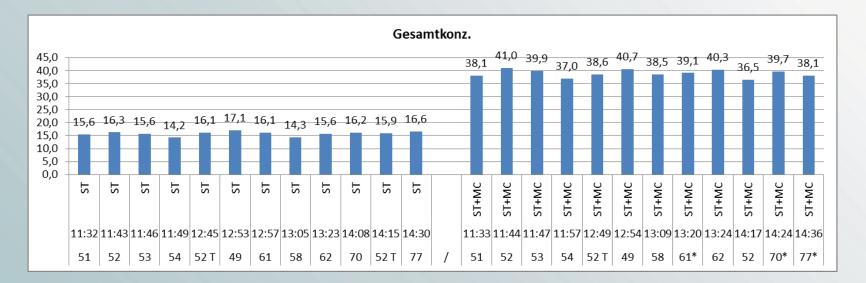
28 FluoroProbes Measuring in Parallel

To what extent can we expect the same results with each probe?

Chr. Moldaenke, bbe Marén Lentz, IGB, Berlin



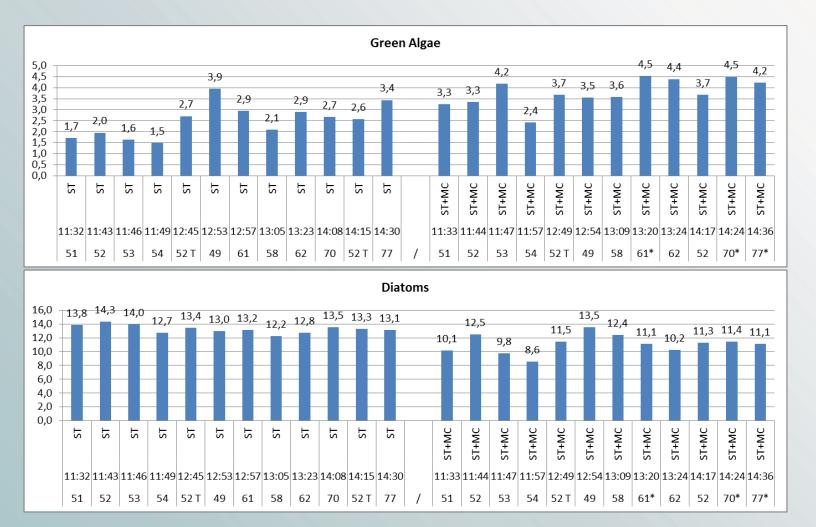
Results not equal enough



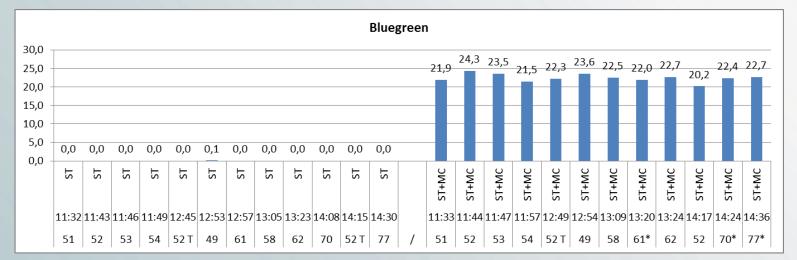
ST= Lake Stechlin ; MC = Microcystis; measurement on 14.03.2012

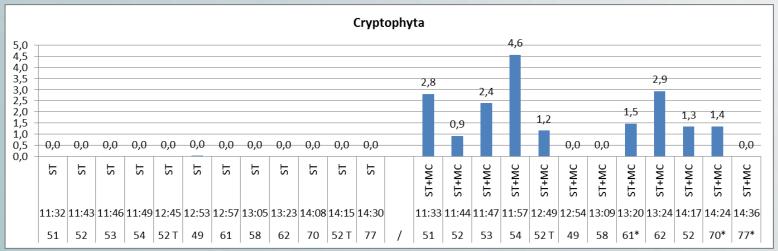
-> Scattering in all parameters not accepted





Page 3





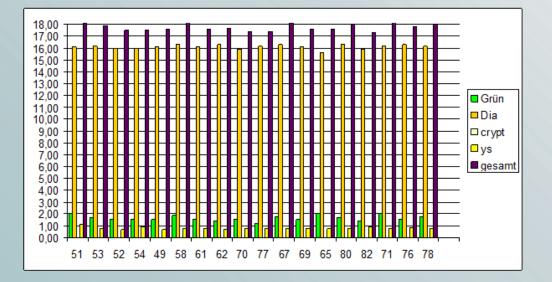
moldaenke

bbe



First steps for better evaluation:

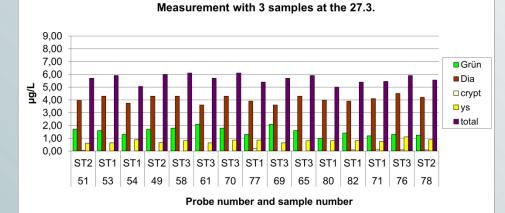
- Determine the offset with bi-distilled water
- Filtrate (0.2-0.45 µm) sample water and re-calibrate the offset for yellow substances
- Re-scale the fingerprints

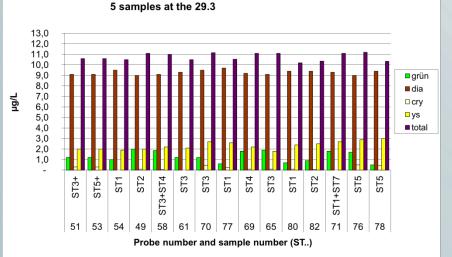


Looks good... But is it the truth?

..only if the settings/fingerprints are valid for other concentrations and distributions as well







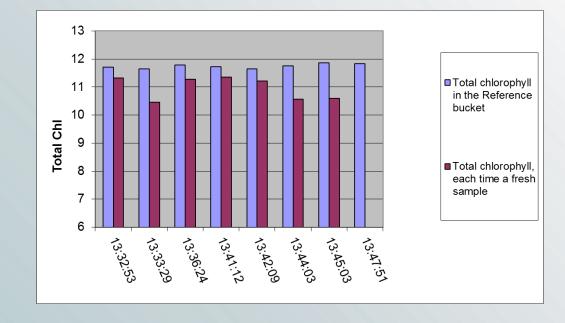
Different concentrations on March 27 and March 29:

Still some scattering:

- 'Induction Kinetics' considered, keep the probe moving
- Also, a warm probe in cold water can produce air bubbles: move the probe

Does the sampling influence the results?





Probes were alternately submersed in a reference bucket, always containing the same sample, and a bucket with samples which were exchanged before each measurement.

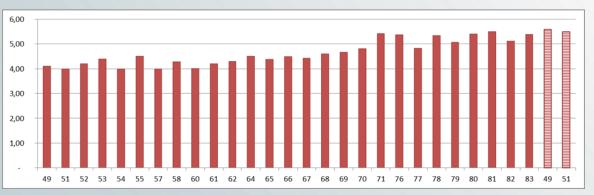
-> sampling is a strong issue

All measurements in one large bucket !?



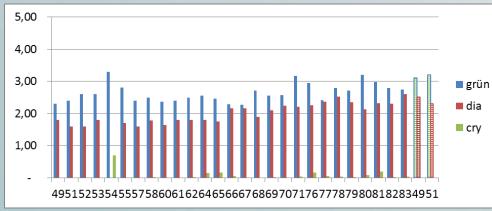
bbe

moldaenke



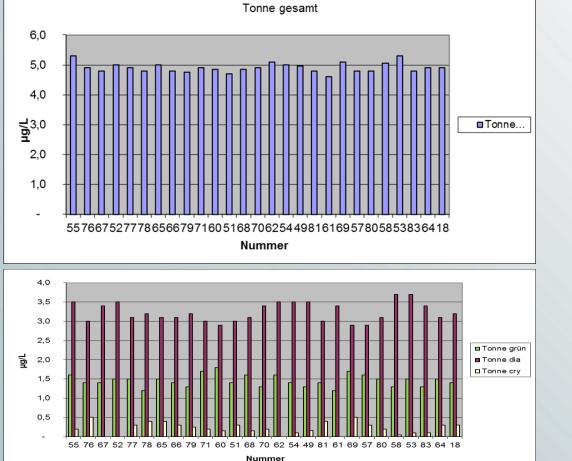
Total chlorophyll of a measurement on May 2. The measurements with the probes 49 and 51 were repeated at the end

-> 28 measurements tend to consume too much time, the algae physiology changes and the results differ



Interesting aspect: diatoms seem to be most affected. <u>Possible reasons</u>: light changes, temperature changes



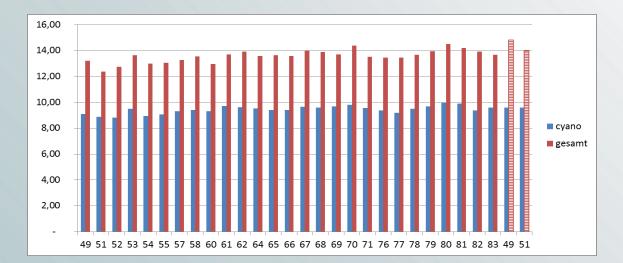


Measurement in a large bucket on April 24

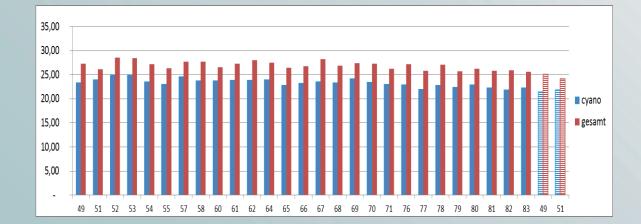
...so far , so good...

Is the variance in the algae class distribution an incorrect calibration or physiology?





Spike test of the sample on May 2, additional 9µg/L cyanobacteria



Spike test of the sample on May 2, additional 21µg/L cyanobacteria

....physiology....



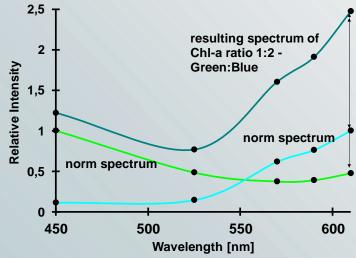
bbe

moldaenke

Spike tests have their own difficulties: cultured algae will be shocked by sample water due to changes in temperature, osmotic pressure etc.

This can induce drifts, oscillations and changes in fingerprints

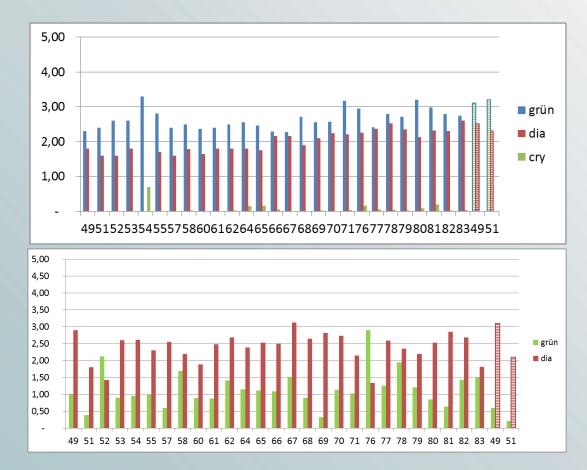
Example using bbe++: Cyanobacteria fingerprint changes





bbe

moldaenke



Spike test of the sample of May 2, additional 9µg/L cyanobacteria, residual algae classes

Spike test of the sample of May 2, additional 21µg/L cyanobacteria, residual algae classes

There are natural limits for the resolution depending on the maximum concentration



moldaenke

It appears that the lowest class scatters the most. This can be up to 5% of the total concentration, resulting in a broad scattering of the class of minimum concentration.

In the case discussed below, the total concentration is about $27\mu g/L$, the lowest class - the green algae - scatter up to $1\mu g/L$, which is about 4%.



bbe

moldaenke

Many Thanks for your Attention