





Joint research on the Lake Atnsjøen ecosystem changes – assumptions and methodology

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- Project: FSS/2013/IIC/W/0022, "Reconstruction of the Environmental Changes and Monitoring - Tools for Planning the Sustainable Development of the Lake Ecosystem"
- Project duration: 01.03.2014 29.02.2016
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- External cooperation: dr L. Nevalainen, dr T. Luoto, M. Oksman







The **main aim** of the project is to exchange knowledge between the partners on the possibilities of using results of the research on lakes' history (**paleolimnology**) and contemporary **environmental monitoring**. Our study site is **Lake Atnsjøen** located in eastern part of Southern Norway, near Rondane National Park.

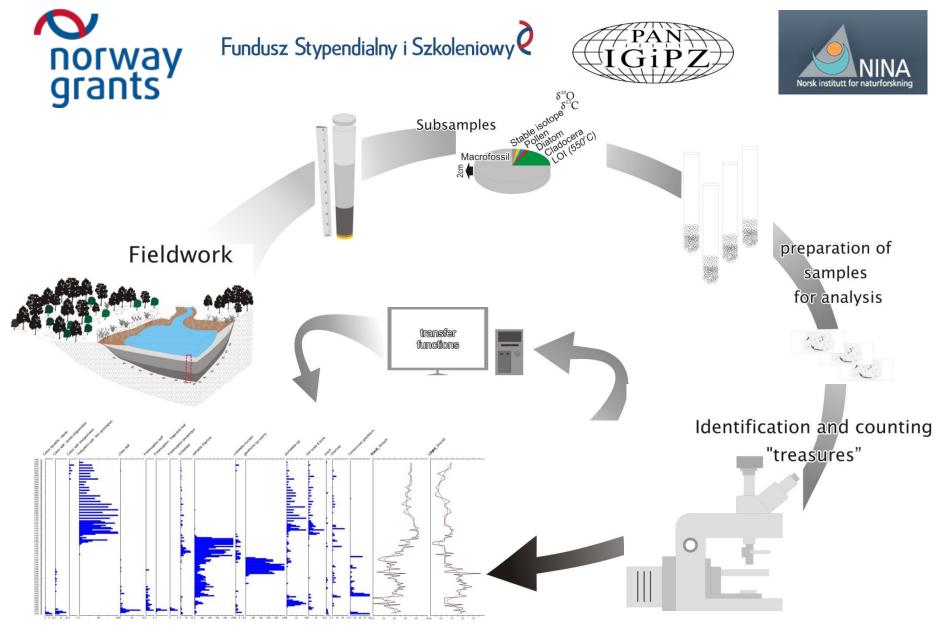






Paleolimnology is a scientific subdiscipline which deals with reconstructing the paleoenvironments of inland waters and especially changes associated with climatic fluctuations, human impacts (e.g., eutrophication, or acidification).

Paleolimnological studies are based on analyses of **lake sediment cores**, including the physical, chemical and mineralogical properties of sediments, and diverse biological records (e.g., fossil diatoms, Cladocera, ostracodes, molluscs, pollen, pigments, or chironomids).



Construction diagram and interpretation









The age of Lake Atna sediments was detrmined using standard dating methods: ²¹⁰Pb and ¹⁴C. The results of dating allowed to build depth scale into age scale.

The history of the lake environment and catchment changes were revealed with the use of various paleolimnological methods:

- Subfossil Cladocera and ephippia analysis
- Palinology
- Macrofossil analysis
- Chironomid analysis
- Diatom analysis
- Chemical Composition of the sediment the Environmental Changes and Monitoring - Tools for Planning the Sustainable Development of the

Lake Ecosystem"

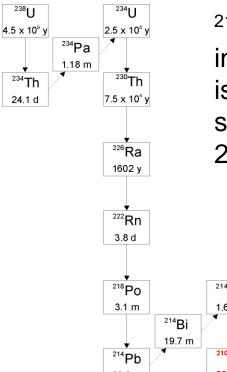




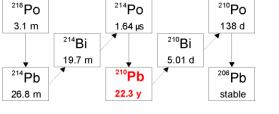




²¹⁰Pb dating



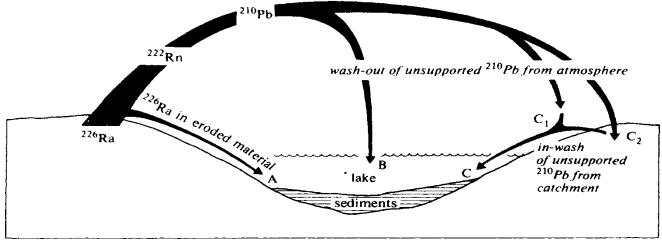
²¹⁰Pb occurs naturally as one of the radionuclides in the ²³⁸U decay series. Disintegration of the isotope ²²⁶Ra yields ²²²Rn which decays through a series of short-lived isotopes to ²¹⁰Pb (half-life 22.26 years).



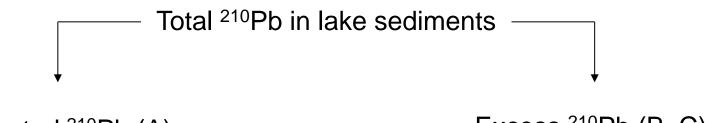








Håkanson and Jansson (2002)



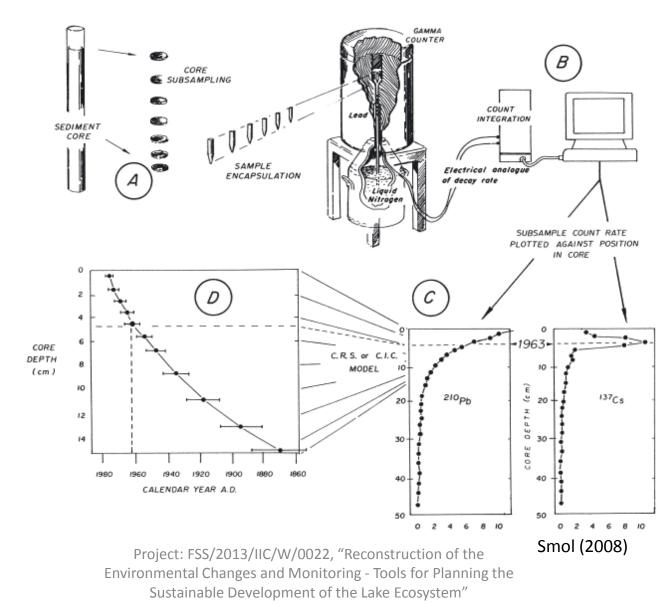
Supported ²¹⁰Pb (A)

Excess ²¹⁰Pb (B, C)















What can be dated with ²¹⁰Pb?

- 1. Glacier ice
- 2. Lake sediments
- 3. Marine sediments
- 4. Peat
- 5. Alluvial sediments

Analytical techniques:

- 1. Alpha spectrometry
- 2. Gamma spectrometry



Limitations:

- 1. Only undisturbed sediments
- 2. Time range 120-150 years
- 3. Uncertainty increases with age
- 4. Dates >100 years risky!!!
- 5. Different calculation models can produce different results
- 6. Need for independent validation!







Laboratory techniques

Alpha spectrometry

Prons:

- 1. Small mass of dry sediment required
- 2. Low detection limits
- 3. High measurement precision
- 4. Reasonable counting time

<u>Cons:</u>

- 1. Wet digestion necessary
- 2. Destructive technique
- 3. Only ²¹⁰Pb measured
- 4. Isotope equilibrium in almost 2 years

Gamma spectrometry

Prons:

- 1. No chemical treatment
- 2. Non-destructive measurements
- 3. Other isotopes measured simultanously
- 4. Isotope equilibrium in 3 weeks

<u>Cons:</u>

- 1. More sediment necessary
- 2. Longer measurement time
- 3. Higher detection limits
- 4. Lower precision

Combination of both methods possible







Cladocera analysis

The Cladocera (water fleas) are major component of the microcrustacean fauna of freshwater lakes. They live close to the shore, among the plants in littoral zone and also offshore in pelagic zone. Some species are herbivores (plant eaters) while some are carnivores (animal eaters). During most of the growing season, Cladocera reproducee via unfertilized female-bearing eggs. If the environmental conditions deteriorate (e.g., overcrowding, limited food availability, temperature exceeds limits, oxygen depletion), males are produced parthenogenetically and efippia are generated sexually (gamogenesis).







Cladocera

- 620 species discovered so far, ~90 live in Europe
- 0,2 18mm size
- Live in pelagic, litoral, benthic zones
- Reproduce by parthenogenesis and sexually
- Cladocera produce dispousal eggs (efipia) that preserve in the sediments
- Remains of \approx 70 species preserve in the sediments
- Efipia- dispousal eggs







Subfossil Cladocera analysis allows to reconstruct:

- Trophy changes
- Acidification changes
- Climate temperature is an important factor controlling Cladocera distribution
- Lake level fluctuations







Subfossil Cladocera analysis – laboratory method

Material for subfossil Cladocera analysis is always prepared according to standard procedure described in Frey (1986). Samples (1 cm³ of fresh sediment each) are treated with hot 10% KOH for 20 min using a magnetic stirrer in order to deflocculate the material, sieved through 33-µm mesh and diluted in 10 cm³ distilled water. Slides are prepared from 0.1ml of each sample and examined with a microscope (Olympus BX41).

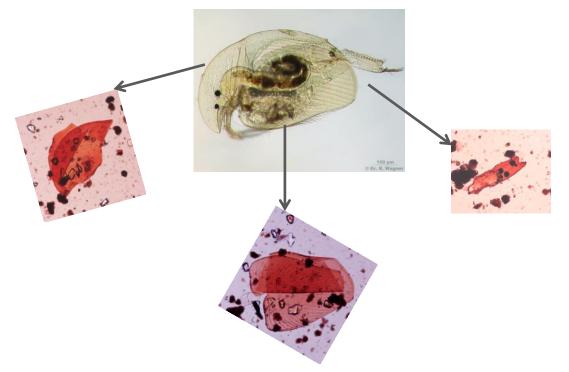








Picture of living Cladcocera specimen (Acroperus harpae)



Pictures of subfossil remains of *Acroperus harpae* that preserve in the lake sediments



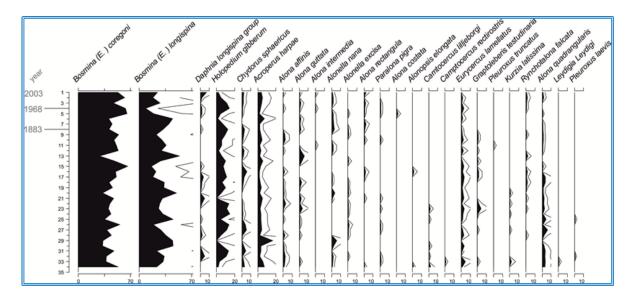




Subfossil Cladocera analysis - results

Results are presented on the stratigraphic diagrams of percantage share of species, absolute abundances and species fluxes, created by means of specialised software. One of the most popular is C2 by Juggins:

(https://www.staff.ncl.ac.uk/stephen.juggins/software/C2Home.htm)









Pollen analysis – Palynology

Palynology studies composition of pollen grains and spores that accumulated in lake sediments. The analysis allows to reconstruct past vegetation changes in the area, which are often due to climate changes or human activity. Pollen grains and spores are extremely resistant to decay.







Pollen analysis

Material for pollen analysis is always prepared according to standard procedure, described in Berglund and Ralska-Jasiewiczowa (1986). All sporomorphs are identified and counted until a minimum of 500 pollen grains of trees and shrubs (arboreal pollen, AP). Pollen grains of all herbaceous species (except the local aquatic and telmatic plants) are counted as non-arborescent pollen (NAP), and the sum of AP and NAP is considered as 100%. The pollen distribution diagrams are drawn by the means of specialised software, for example Tilia2 and Tilia-Graph (Grimm,1992). Basing on the changes in the share of AP and NAP local pollen assemblage zones are divided.







Classical pollen diagram prepared by Ulf Hafsten (1957) with modification

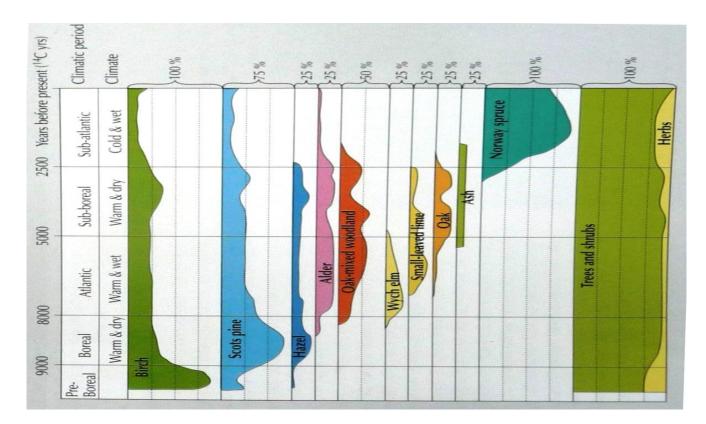


Figure from "National Atlas of Norway. Vegetation" Asbjørn MOEN. 1999



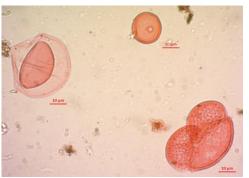




Example of pollen grains from Lake Atnsjøen



Carex t..



Isoëtes lacustris, Poaceae, Pinus



Juniperus communis



Picea abies







Macrofossils analysis

Macrofossil are plant and animal parts visible with a bare eye or under the binocular. In the lake sediments preserve well:

- Plant remains such as: seeds + fruits (orchids to coconuts), cones, Sporangia, leaves + buds, needles, wood
- Remains of different animals such as: fish, oribatida, Cladocera ephippia etc.

Macrofossil analysis provides record of <u>local</u> vegetation and helps to reconstruct:

- Water level fluctuations (paleohydrology)
- Lake trophy changes
- Climate changes







Macrofossil analysis

The samples for the plant macrofossil analysis are rinsed in warm water on 0.125 mm mesh size sieves. Macrofossils are diagnosed with the use of a stereoscopic microscope and a light microscope. Species determination of individual fossils is performed with the help of the appropriate keys (Berggren 1968, 1981; Tobolski 2000; Velichkevich and Zastawniak 2006, 2009). The results are usually presented in the form of stratigraphic diagrams with the use of a program - C2 (Juggins, 2003). The generative parts are summarized in the diagrams in absolute numbers (seeds, fruits), while the vegetative fragments of vascular plants as well as *Sphganum* mosses were visually determined in percentages of the total volume of the sediment samples.

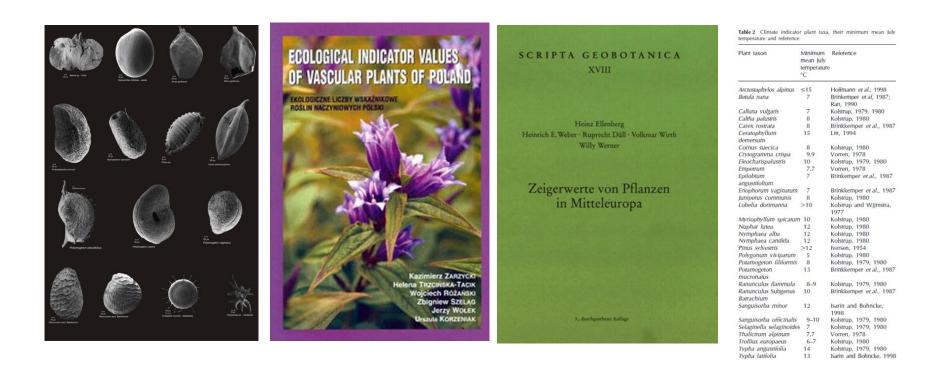








Main books used in interpretation of macrofossils data:









Macrofossils from Lake Atna





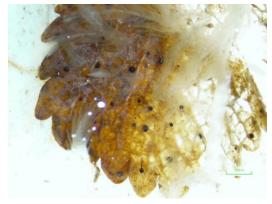
http://www.butbn.cas.cz/en/ecology/staff/martina-ctvrtlikova





Sphagnum sp.

Isoëtes lacustris



Betula nana







Chemical composition of the sediments helps to reconstruct:

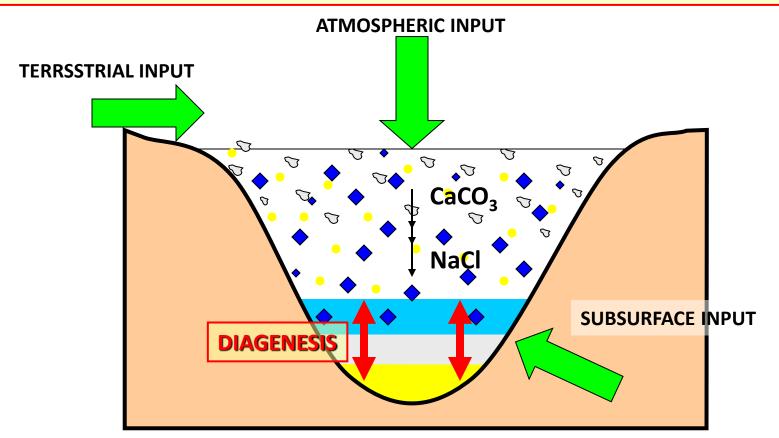
- 1) Rate of denudation (SiO_{2ter}, Ti, Al, AM, AEM)
- 2) Intensity of bioproductivity (OM, SiO_{2biog}, P, carbonates)
- 3) Red-ox/oxygenation (Fe/Mn, Cu/Zn, S, pristane/phytane)
- 4) Salinity (Mg/Ca, Sr/Ca, Na/K, B, TOC/S)
- 5) Anthropopressure (heavy metals, FAP, BC, PAH)







BASIC ASSUMPTION OF CHEMICAL ANALYSIS OF LAKE SEDIMENTS: EQUILIBRIUM BETWEEN THE COMPOSITION OF WATER AND SEDIMENTS









Lake sediments – geochemical perspective

• <u>Endogenic</u> – formed in lake water column

INDICATE PHYSICAL AND CHEMICAL CONDITIONS IN THE LAKE

• <u>Allogenic</u> – delivered from the catchment

INDICATE PHYSICAL PROCESSES IN THE CATCHMENT

<u>Diagenetic</u> – formed within the sediment

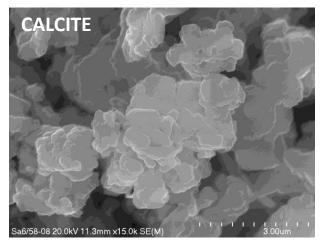
NOISE

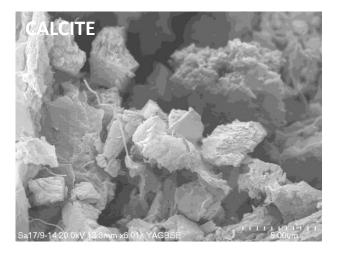


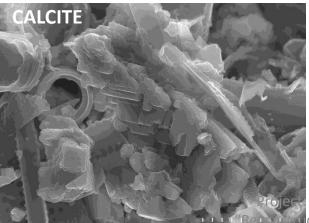




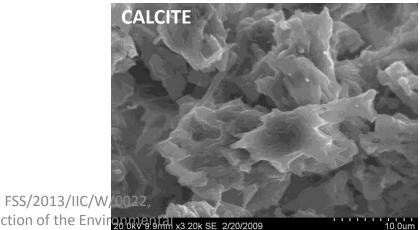
Endogenic components: Carbonates







Sa18/7-09 20.0kV 10.9mm x4.00k SE(M)



Changes and Monitoring - Tools for Phot. M. Woszczyk Planning the Sustainable Development of

Phot. K. Apolinarska







Genesis of carbonates

- 1) Leaching of soils
- 2) Carbonate sceletons
- 3) Incrustations (e.g. charophyts)
- 4) Evaporation
- 5) Mixing precipitation
- 6) Microbial activity



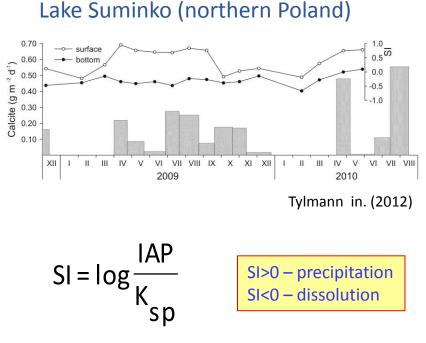




Genesis of carbonates: activity of phytoplankton

Lake Lucieńskie (central Poland)





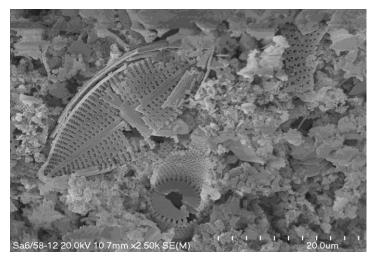






Endogenic components: biogenic silica

- Sponge spicules
- Phytoliths
- Chrysophycae cysts
- Diatom frustules



(phot. M. Woszczyk)







Endogenic components: organic matter

Genesis of organic matter:

- Biomass production
- Oxygen deficiency in water
- Great water depth







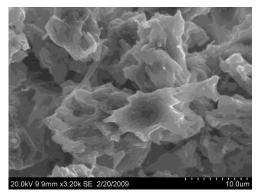
Allogenic components

Terrigenous silica



Fot. M. Woszczyk

Detrital calcite



Fot. K. Apolinarska

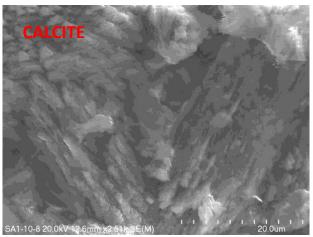


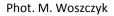


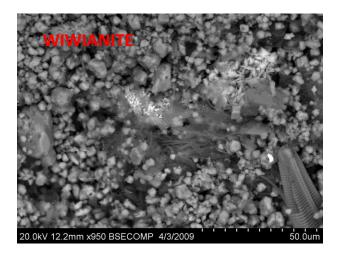


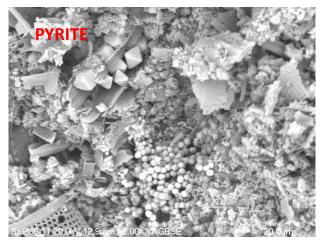
Diagenetic components











Phot. K. Apolinarska







