

Investigating the dynamics of the aquatic community in Oslofjord through time series analysis of eDNA

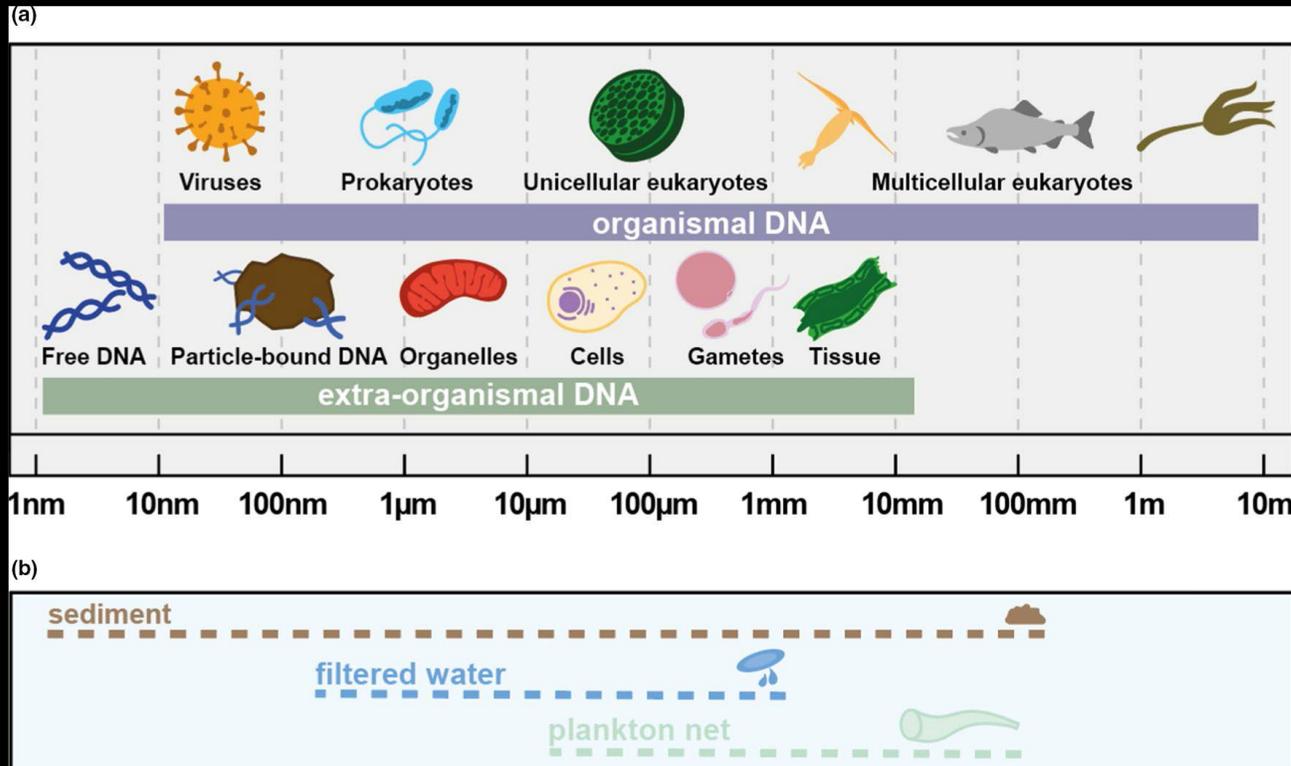
Quentin Mauvisseau

EDGE group (Evolution, eDNA, Genomics and Ethnobotany)

Natural History Museum, University of Oslo



What is eDNA (environmental DNA)?



Rodriguez-Ezpeleta et al. 2021. *Molecular Ecology*

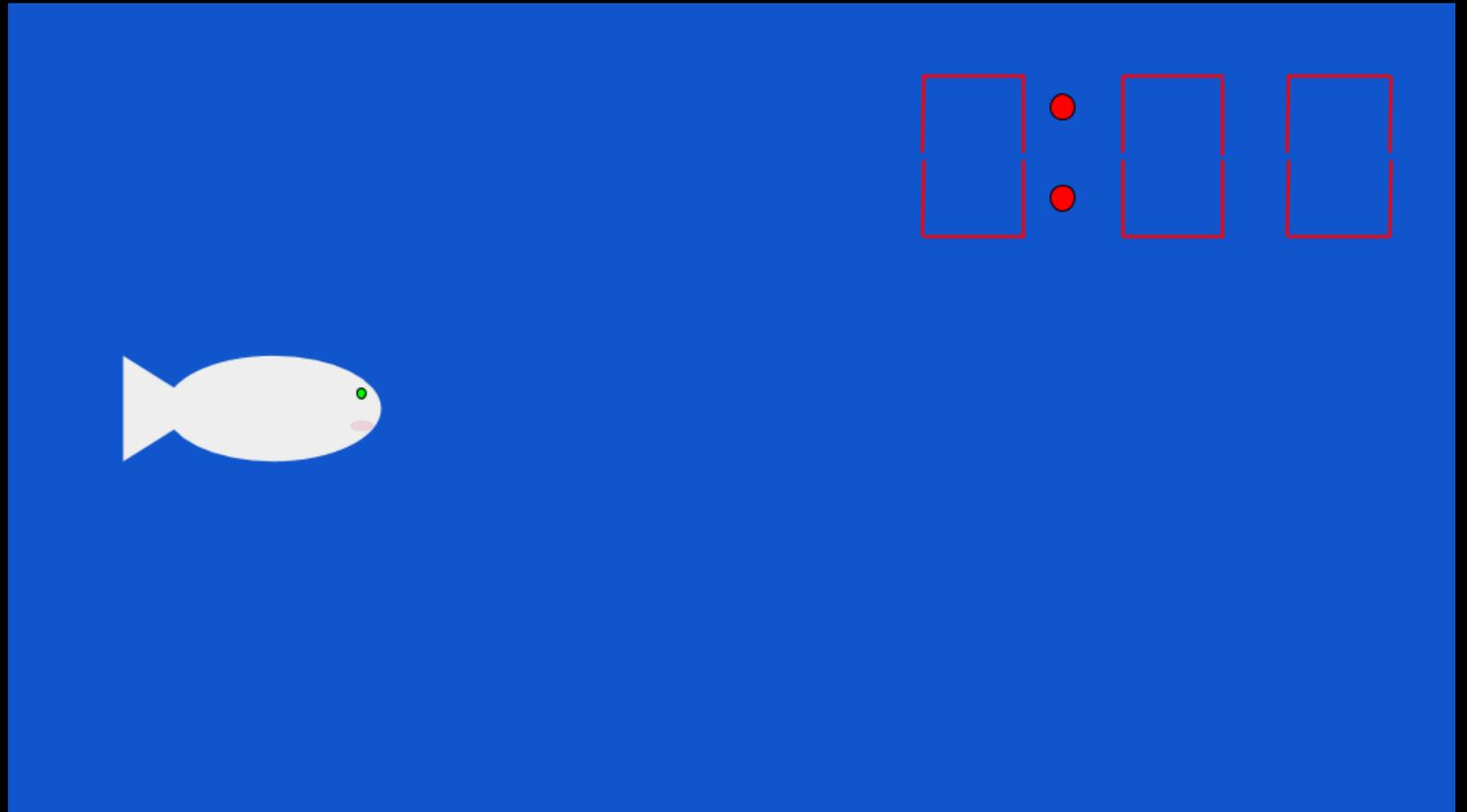
- Free DNA originating from an organism which can be isolated from the habitat of the organism, rather than directly from the organism itself

- eDNA should be used to refer to the total pool of DNA isolated from the environment

- eDNA is composed of organismal and extra organismal DNA

What is eDNA (environmental DNA)?

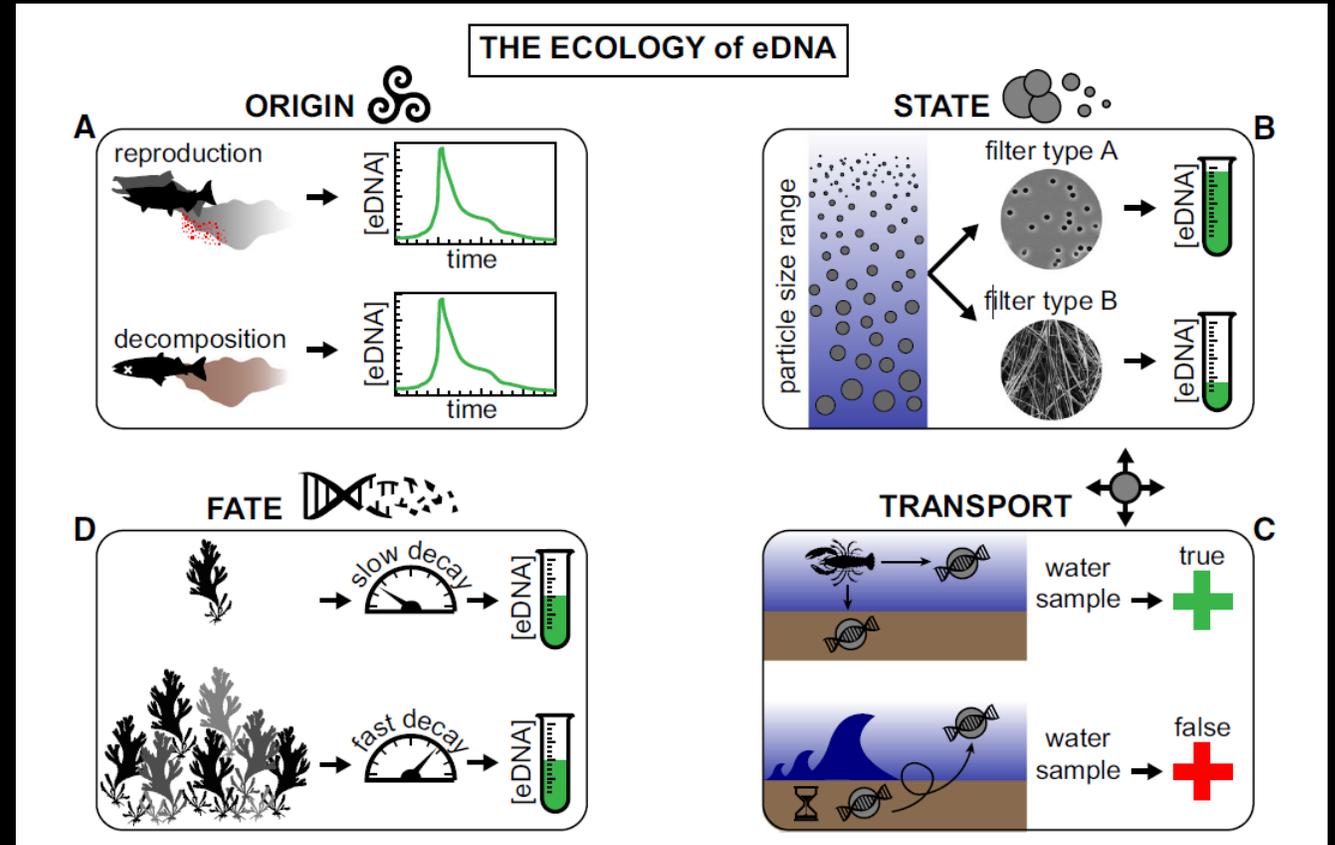
- Limitations
- Persistence
- Abundance
- Sampling technique
- Habitat parameters



False positive and false negative

Limitations of eDNA based methods

- Experimental design
- Weather / seasonality
- Environment / habitat constraints
- Sampling technique
- UV / pH / temperature



Barnes & Turner 2015. *Conservation Genetics*

False positive / negative

Metabarcoding – Benefits

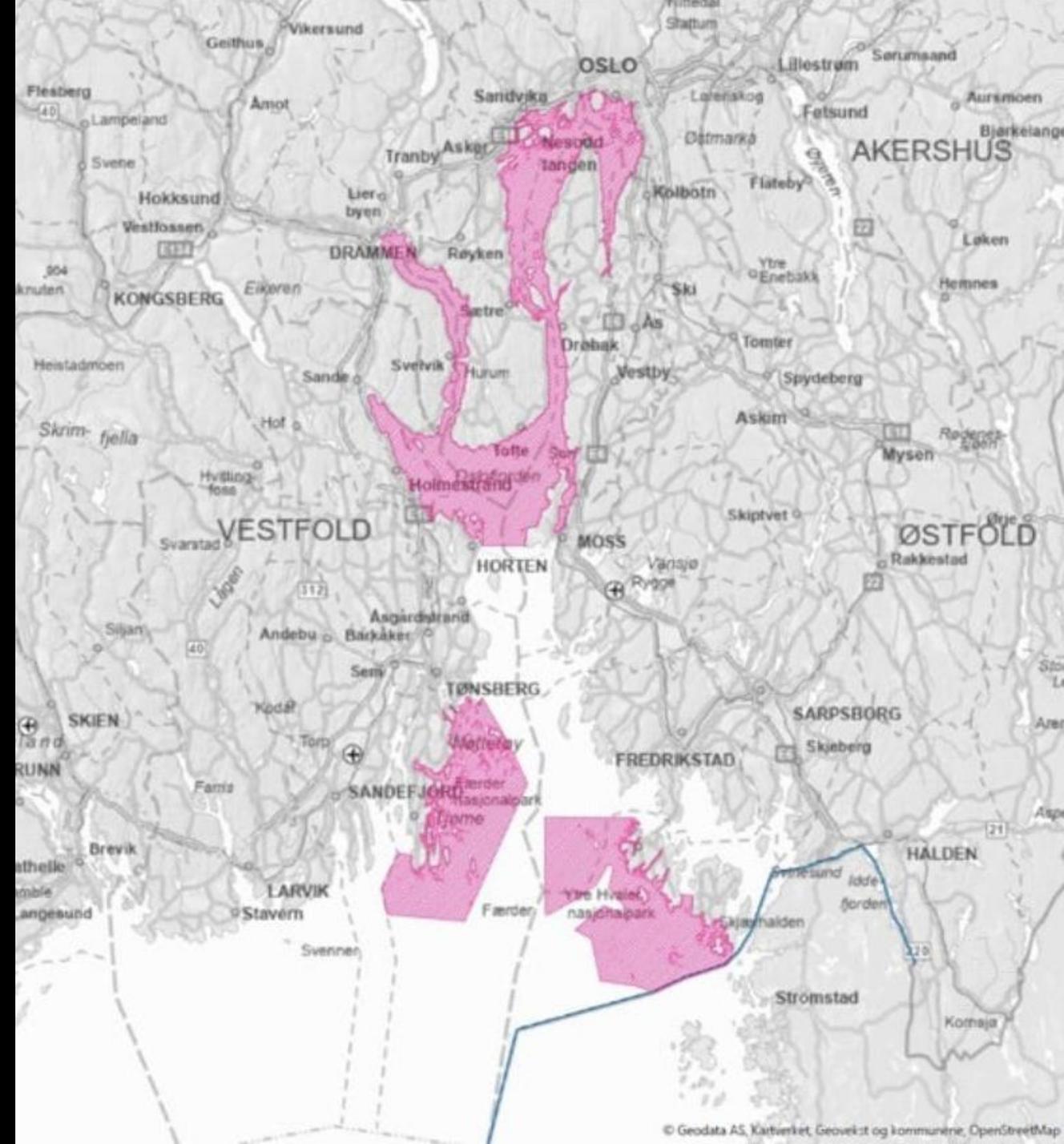
- **Non-destructive and non-invasive**
- **Detect rare, invasive, cryptic and elusive species**
- **Monitor imminent or ongoing environmental changes in remote locations**
- **Easily standardized**
- **Simultaneous assessment of wide range of organisms**

Limitations?

**False
positive/negative**

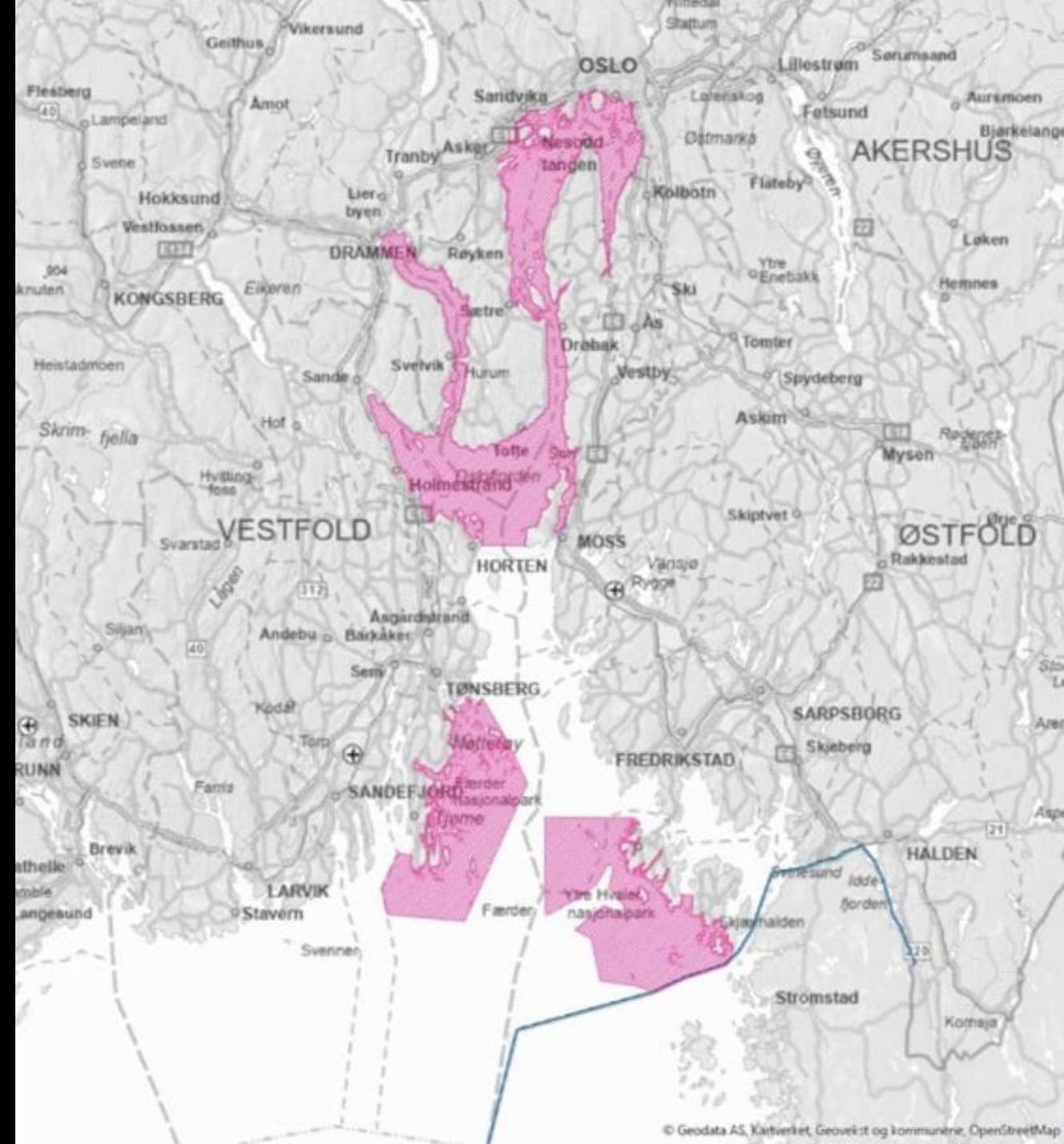
Oslofjord

- Sever threats on the ecosystem
- Fisheries, pollution and runoff from agriculture, sewages
- Fishing ban for cod in 2018: “not had the desired effect”
- Extensive consultation process
- Introduction of zero-fishing zones for 10 years



Oslofjord in Drøbak

- How to monitor the Oslofjord with minimal impacts on the environment?
- Environmental DNA based monitoring?
- Does eDNA time series analysis allow to observe ecosystem shifts?
- Past and ongoing work





OPEN **Harnessing eDNA metabarcoding to investigate fish community composition and its seasonal changes in the Oslo fjord**

Cintia Oliveira Carvalho^{1,2,5}, William Gromstad^{1,5}, Micah Dunthorn¹, Hans Erik Karlsen³, Audun Schröder-Nielsen¹, Jonathan Stuart Ready^{1,2}, Torbjørn Haugaasen⁴, Grete Sørnes³, Hugo de Boer¹ & Quentin Mauvisseau^{1,2}

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REGULAR ARTICLE



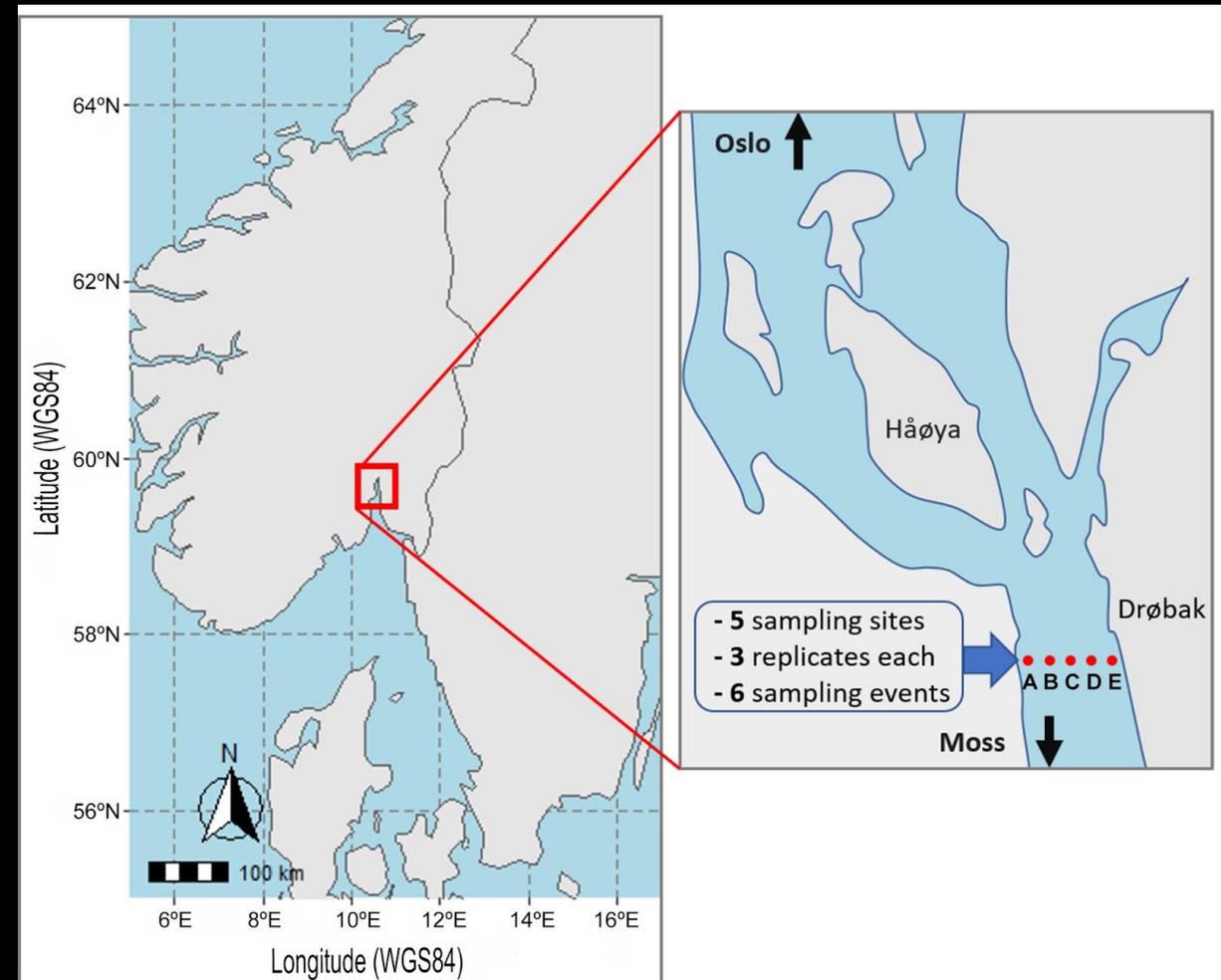
Investigating the dynamics of the aquatic community in Oslofjord through time series analysis of eDNA

Sibusiso Mahlangu¹ | Isabelle Ewers¹ | Cintia Oliveira Carvalho^{1,2} | Audun Schröder-Nielsen¹ | Micah Dunthorn¹ | Hans Erik Karlsen³ | Grete Sørnes³ | Louise Chavarie² | Dag Endresen¹ | Jonathan Stuart Ready⁴ | Hugo J. de Boer¹ | Quentin Mauvisseau¹



eDNA in Drøbak: method

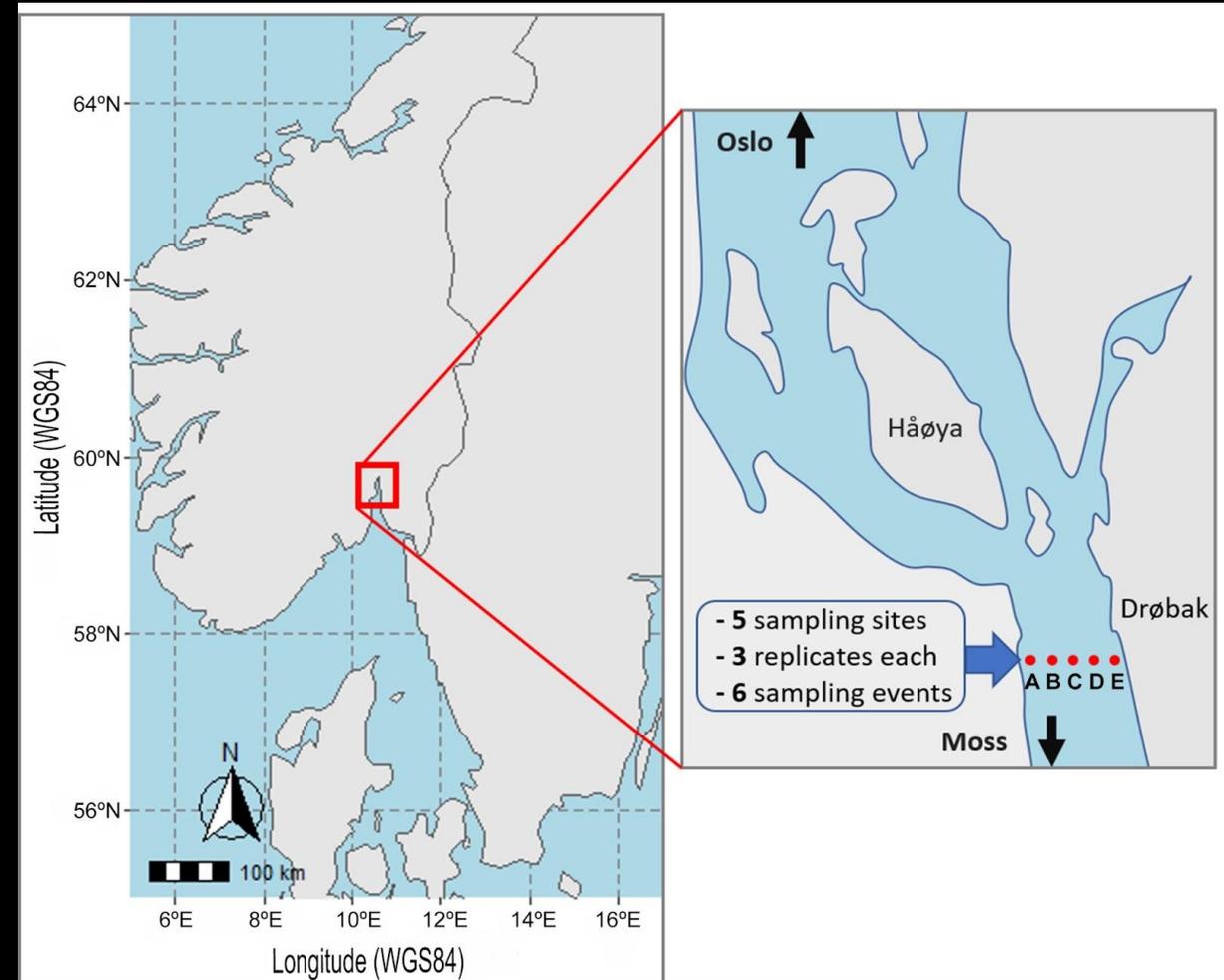
- Oslofjord stretches 100 km from Skagerrak/North Sea to Oslo
- Maximum depth of 258 m
- Inner fjord separated from outer fjord near Drøbak
- Drøbak: shallow sill of about 20 m
- Dramatic environmental decline



Carvalho et al 2024. *Scientific Reports*

eDNA in Drøbak: method

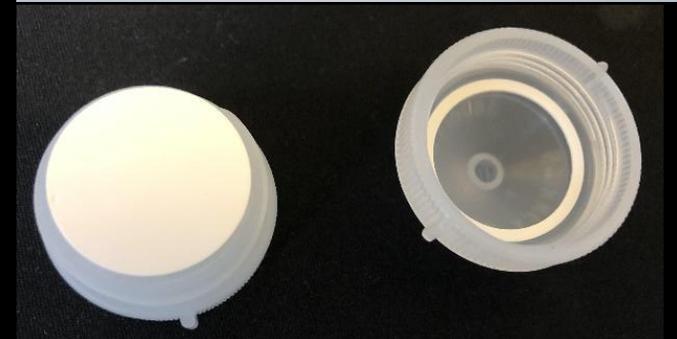
- Two years of sampling: March 2022 to January 2024 (January 2025...)
- 180 eDNA samples, 12 field negative controls
- Sampling every two months
- Other environmental parameters
- MiFish and Elas02 primers. MiSeq. Custom bioinformatics pipelines



Carvalho et al 2024. *Scientific Reports*

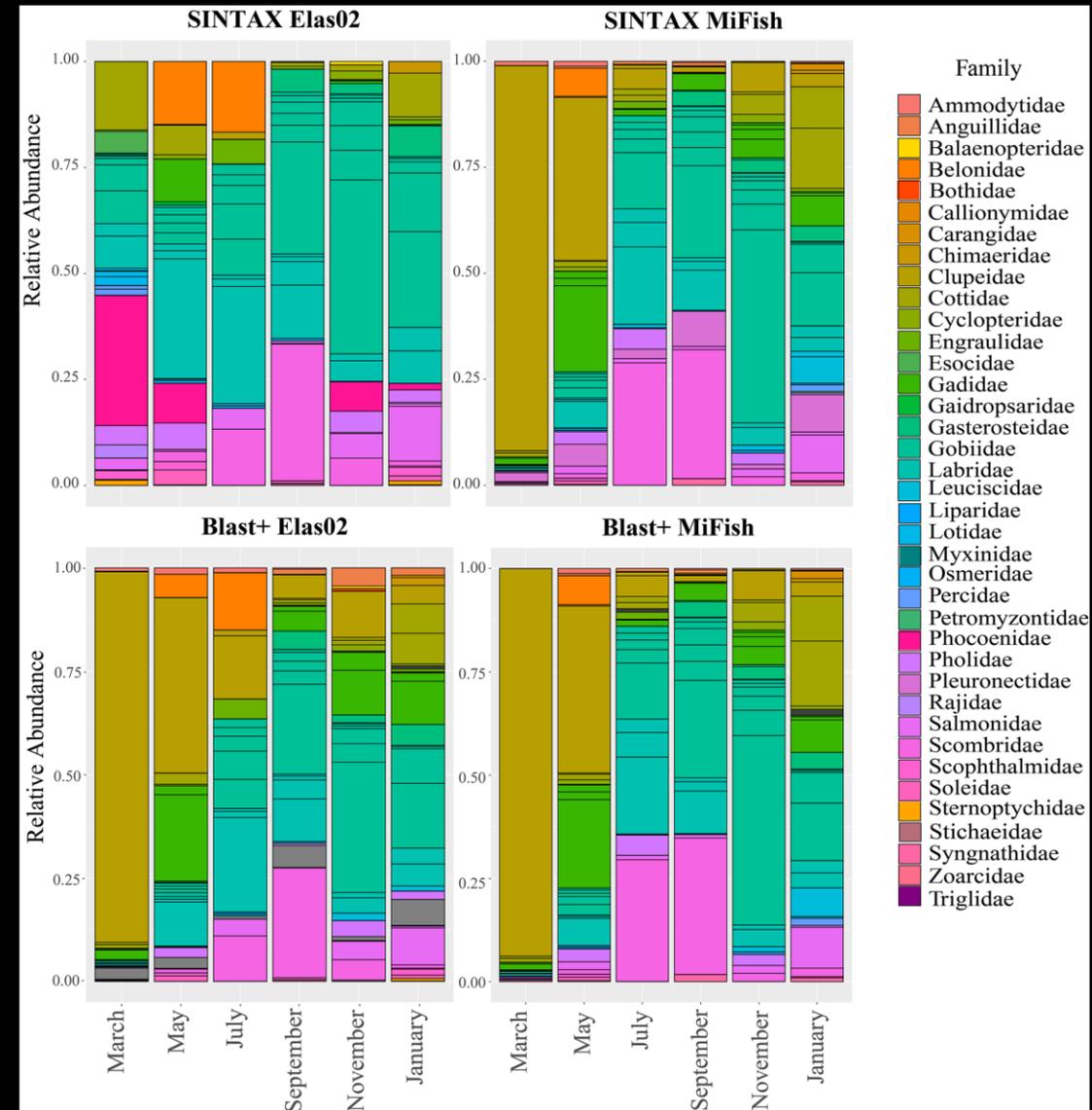
eDNA in Drøbak: method

- Collect water in sterile containers
- Filter holders and sterile membranes
- 0.8 μm pore size and 22 mm diameters
- Freeze samples until DNA extraction and analysis



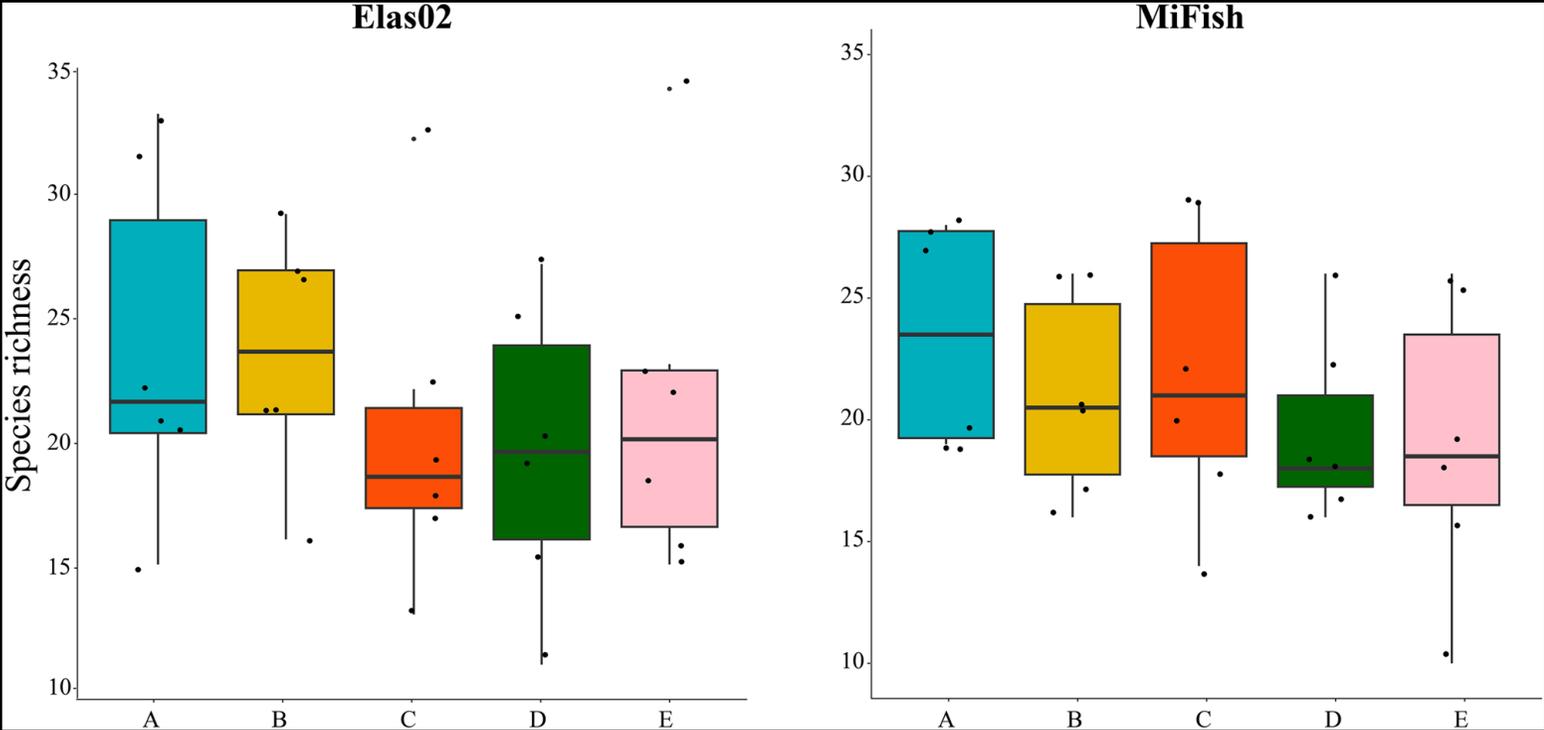
eDNA in Drøbak - year 1

- 63 unique fish species + Sei whale and Harbour porpoise
- 35 species detected with both primers (16 Elas02, and 14 MiFish only)
- Many other birds and mammals detected
- No differences between SINTAX and Blast+ for the taxonomic assignment (using the same database MIDORI2)



eDNA in Drøbak - year 1

No significant difference of species richness across sampling sites for each primers

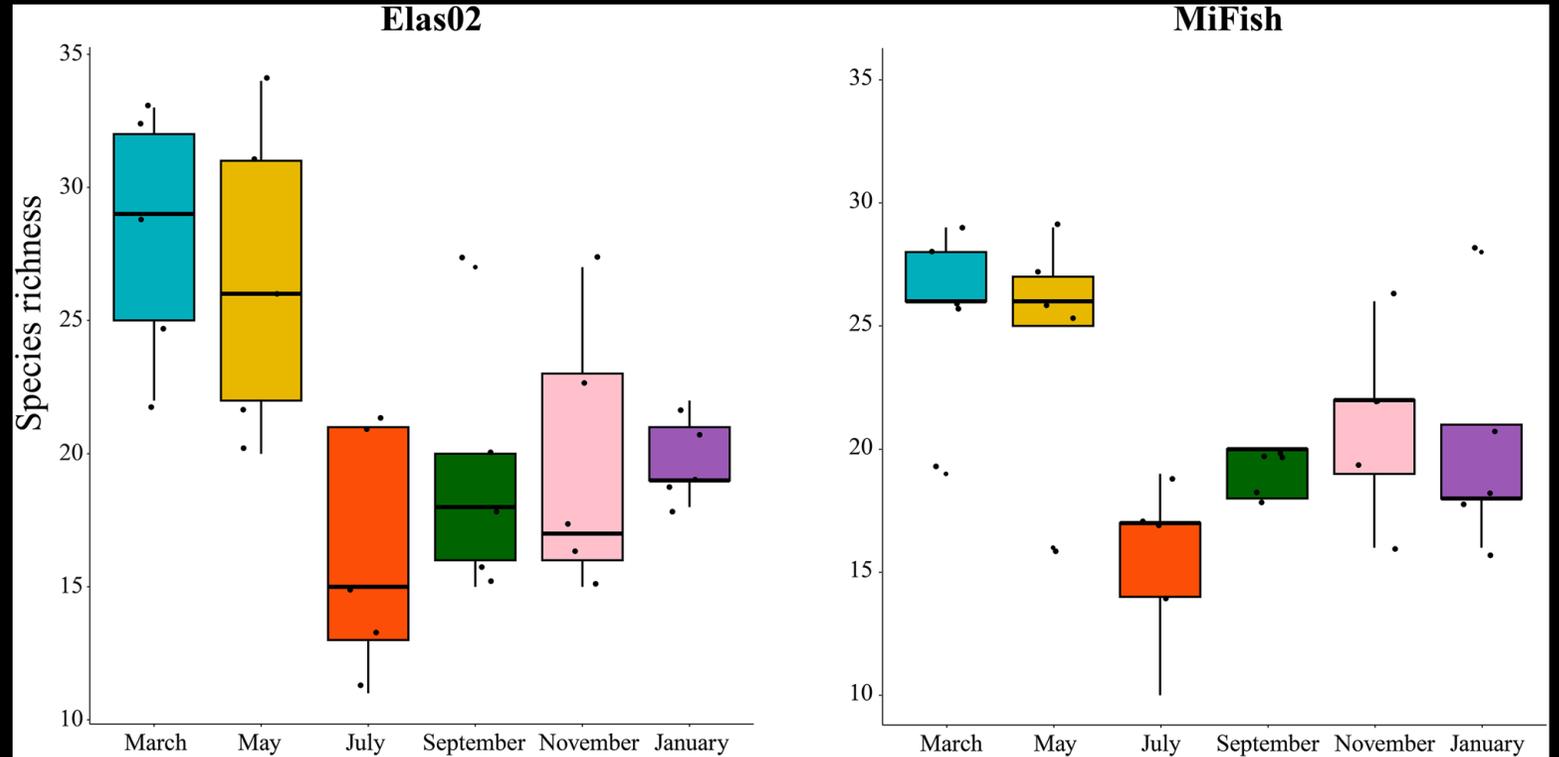


Carvalho et al 2024. *Scientific Reports*

Oslofjord in Drøbak - year 1

Strong seasonal variation for both primer sets

Volume, temperature, conductivity (linked to seasons)

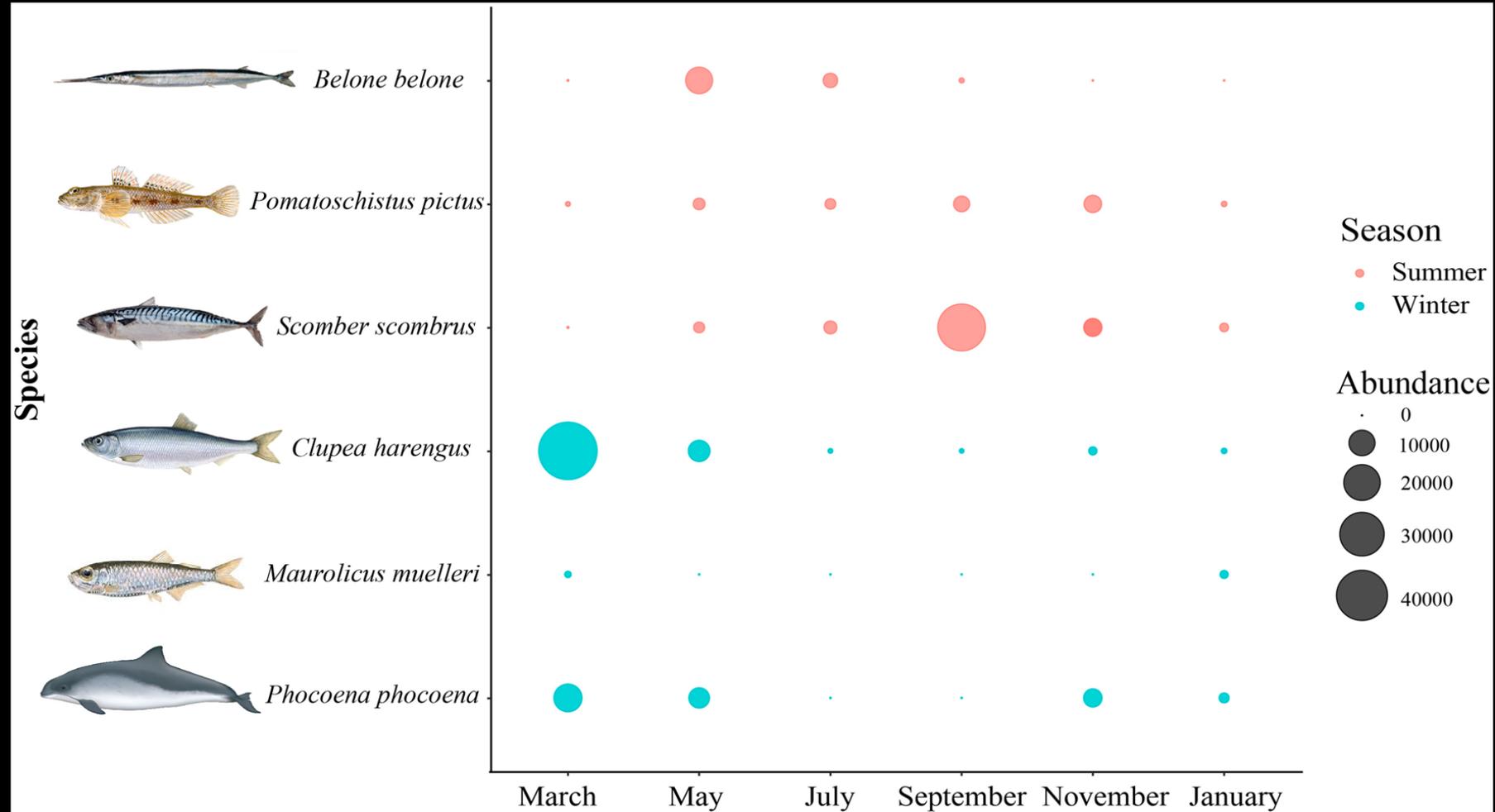


Carvalho et al 2024. *Scientific Reports*

Oslofjord in Drøbak - year 1

Interesting pattern of seasonal changes (read counts as a proxy for relative abundance)

Winter vs summer species



Carvalho et al 2024. *Scientific Reports*

eDNA in Drøbak - year 2

- 63 unique fish species, key species patterns

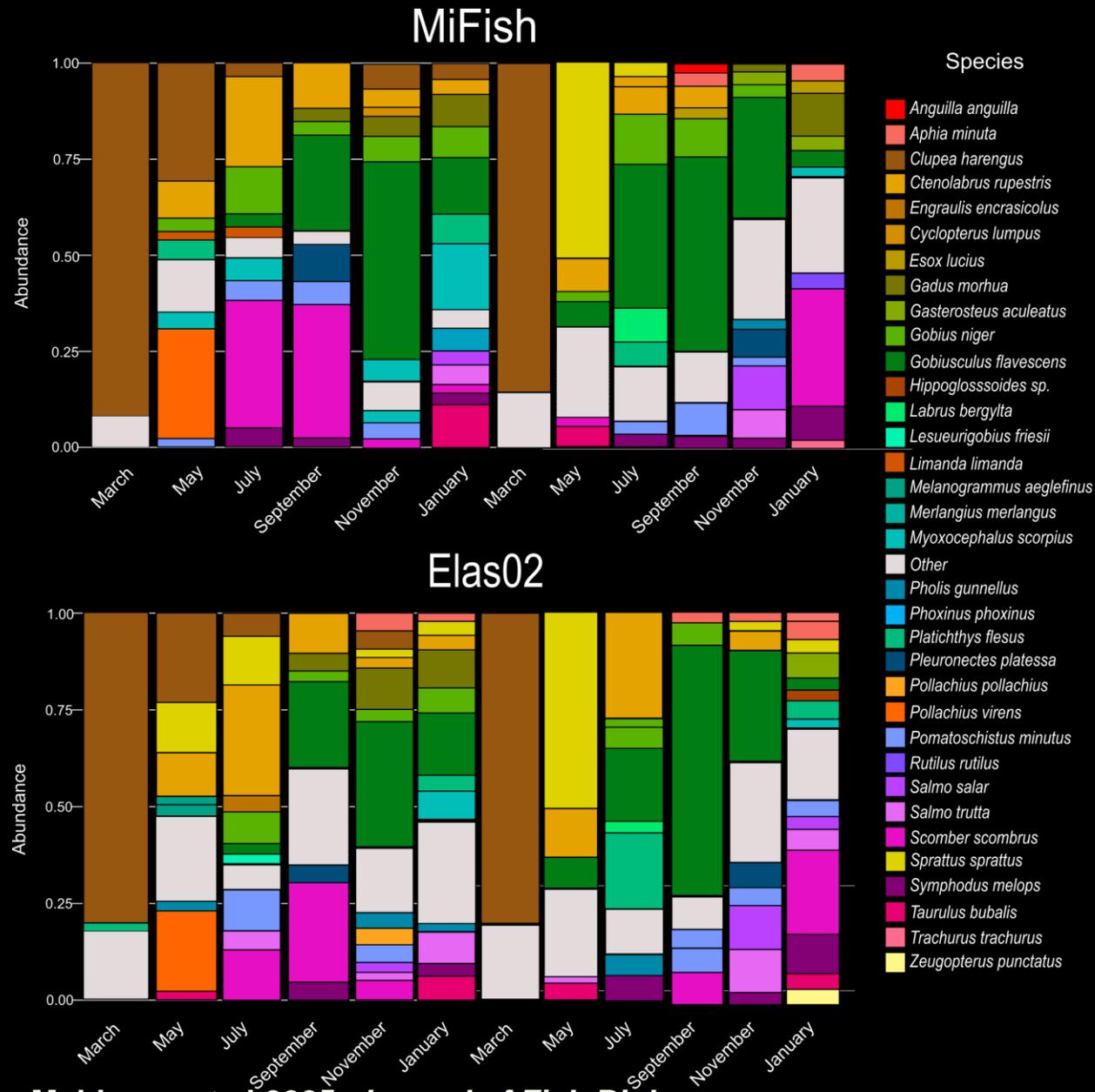
- *Clupea harengus* in March



- *Scomber scombrus* in Summer

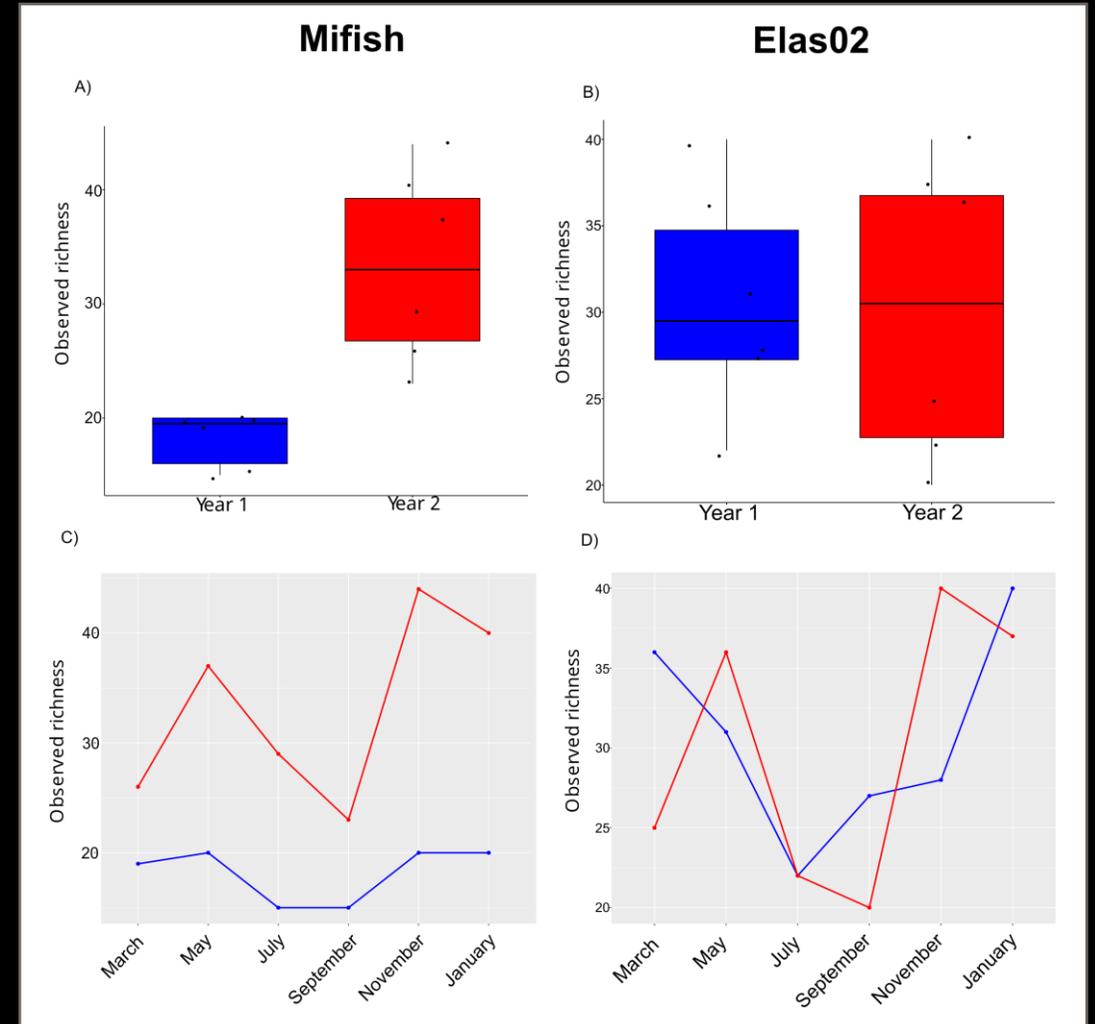


- *Gobiusculus flavescens* in Autumn months

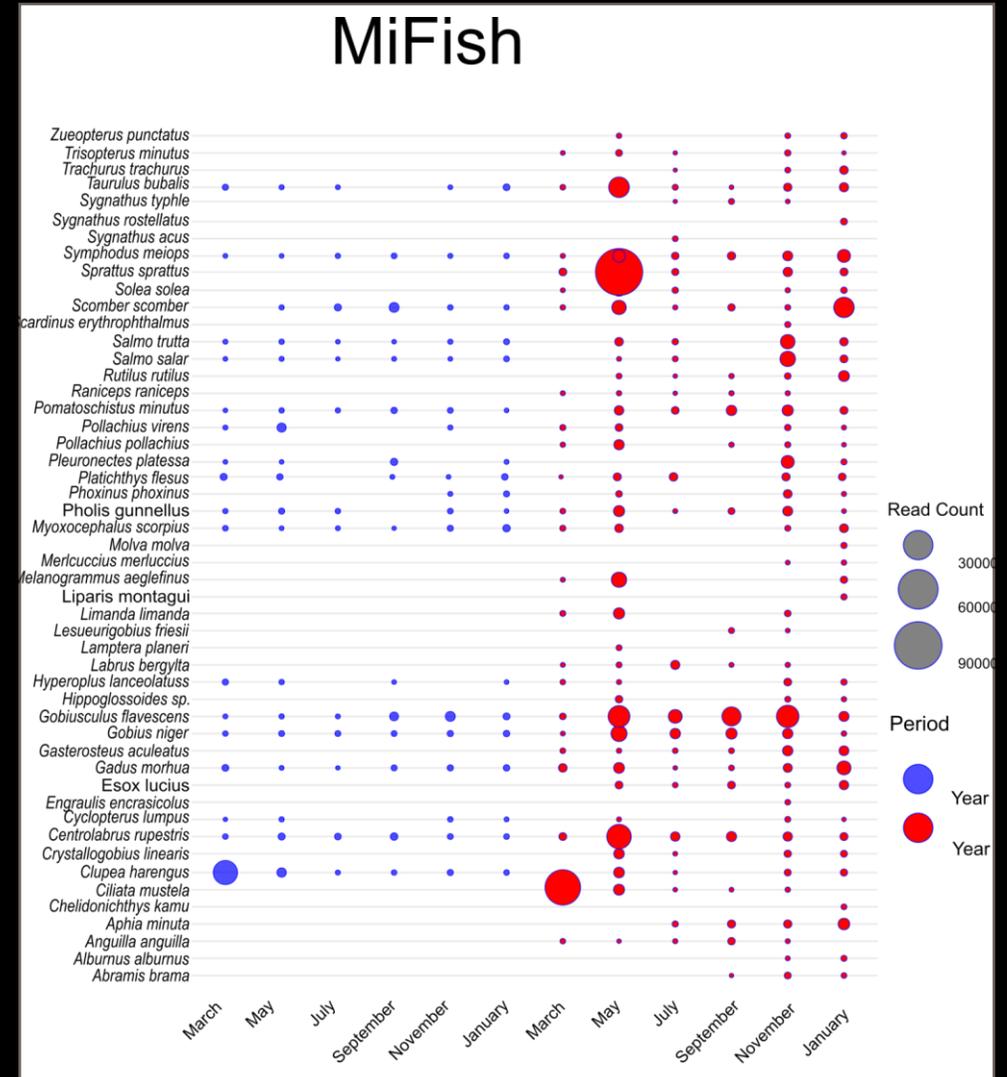
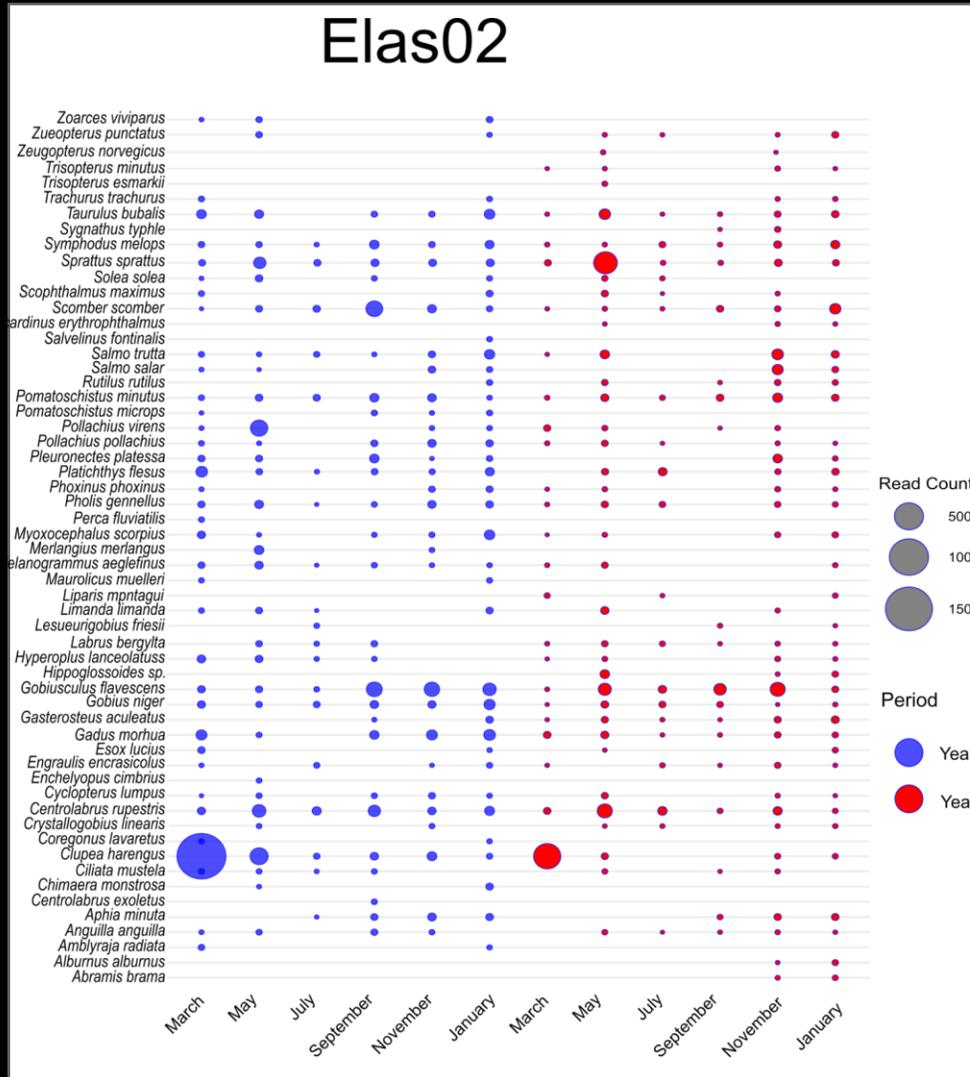


eDNA in Drøbak - year 2

- MiFish: significant increase in species richness in year 2 (linked to library-prep before sequencing)
- Elas02: stable richness across years but turnover in taxa
- Presence/absence data showed significant differences between years (MiFish $p = 0.002$, Elas02 $p = 0.007$)
- Community shifts driven by species turnover, not abundance



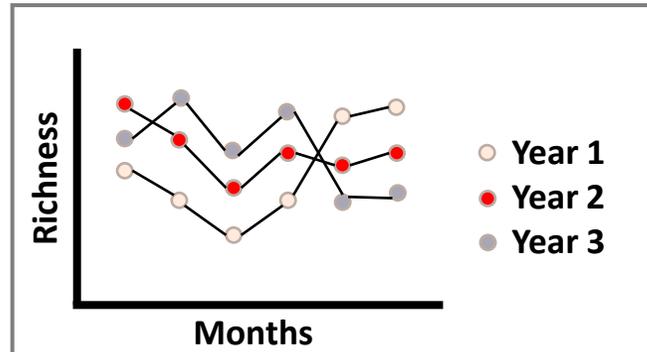
eDNA in Drøbak - year 2



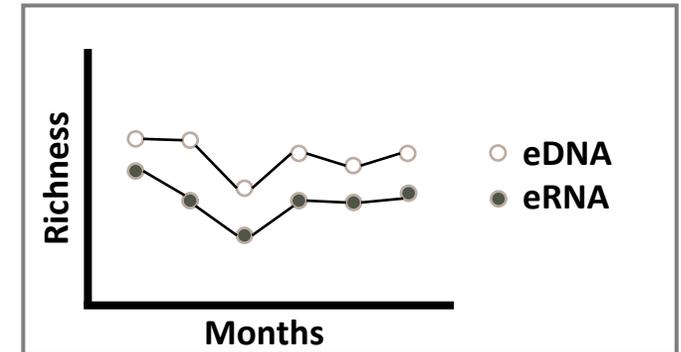
eDNA in Drøbak - year 3 incoming!

- Third year of sampling (until January 2025)
- eDNA following the same protocol as studies 1-2
- eDNA/eRNA using different protocol
- MiFish, Elas02 (and ITS2!)

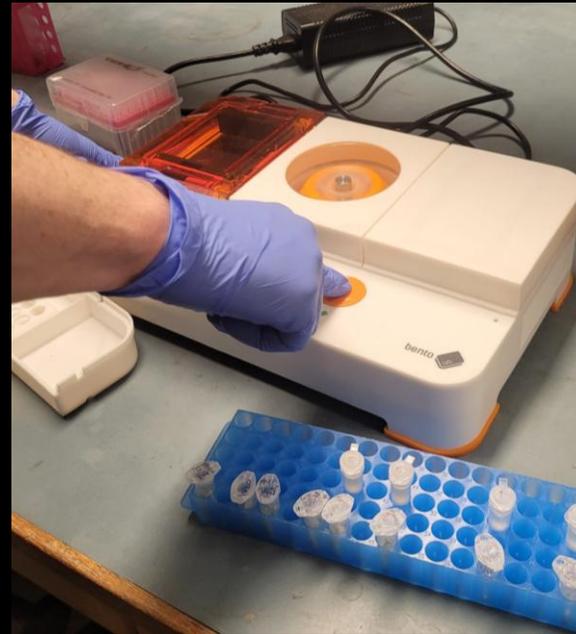
eDNA as a source of temporal ecological data



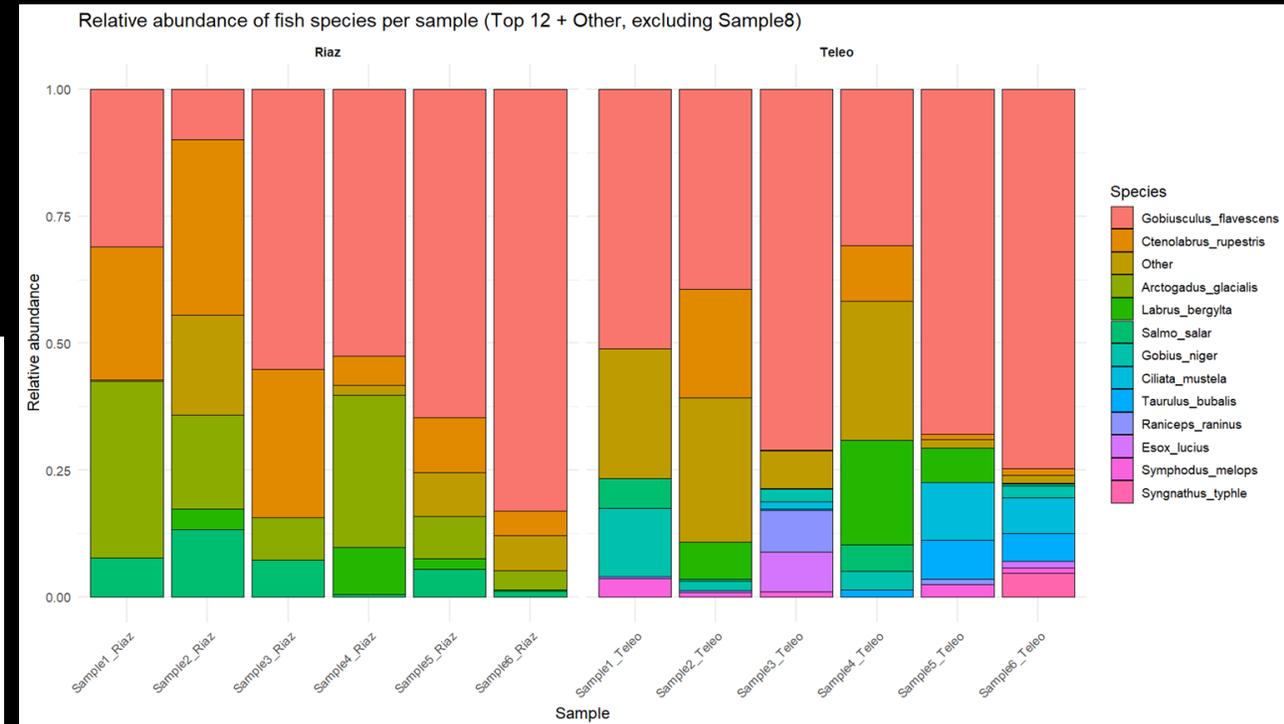
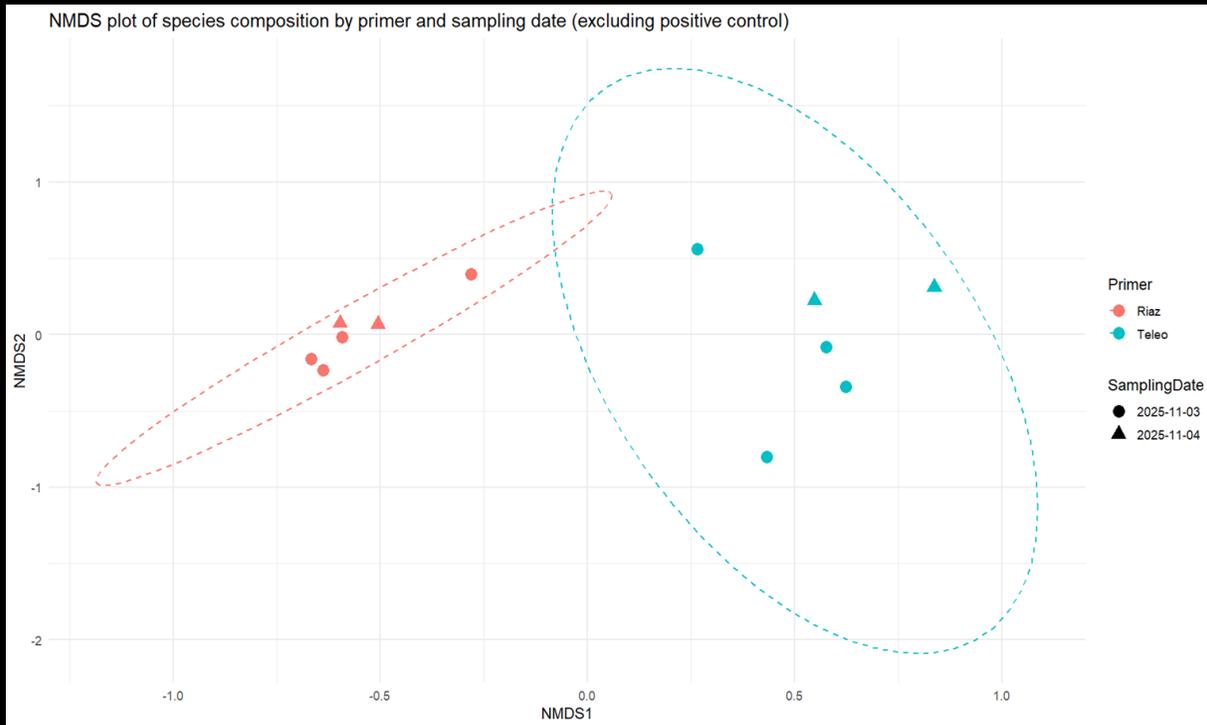
eDNA vs eRNA for time series analysis



What about on-site monitoring?



What about on-site monitoring?



How to increase results reliability?

Environmental DNA

Dedicated to the study and use of environmental DNA for basic and applied sciences

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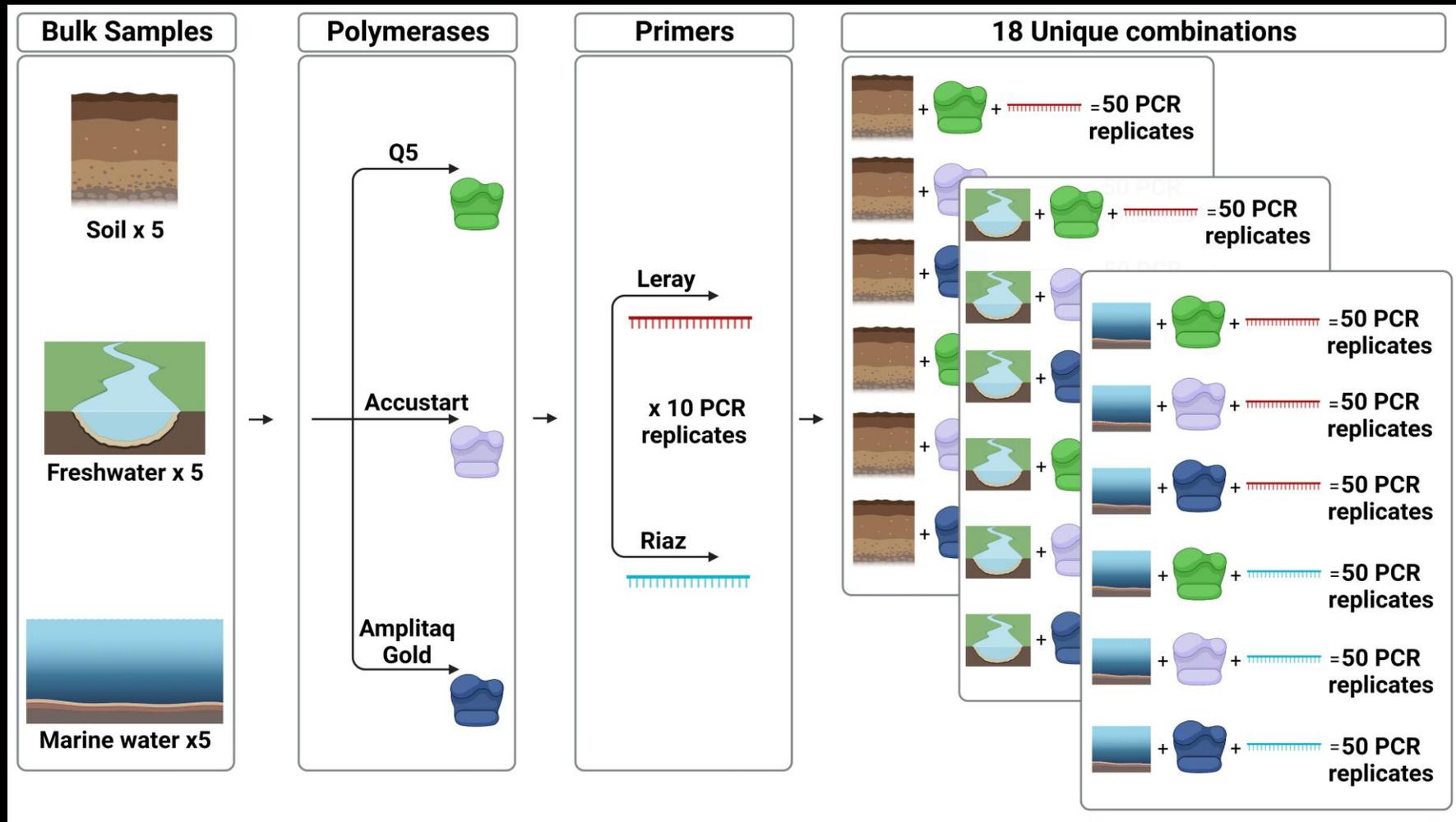
eDNA Replicates, Polymerase and Amplicon Size Impact Inference of Richness Across Habitats

Jarl Andreas Anmarkrud, Fabricio dos Anjos Santa Rosa, Lisbeth Thorbek, Audun Schrøder-Nielsen, Silvana Melo Sviggum, Jonathan Stuart Ready , Hugo J. de Boer, Quentin Mauvisseau 

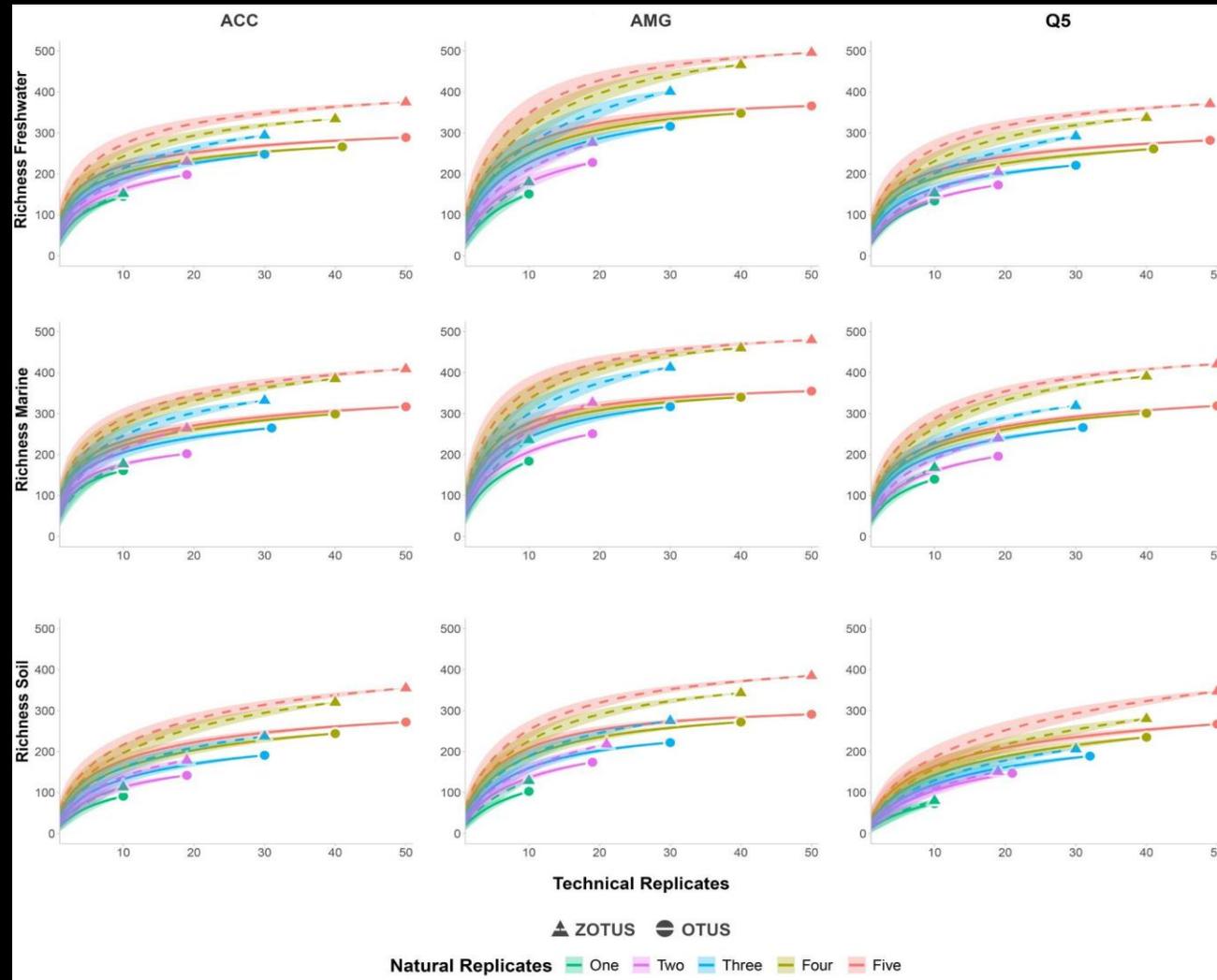
First published: 09 May 2025 | <https://doi.org/10.1002/edn3.70095> | Citations: 3

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How to increase results reliability?



How to increase results reliability?



Thanks!!!



Any questions?

