



# Monitoring UK lake fish communities using eDNA

---

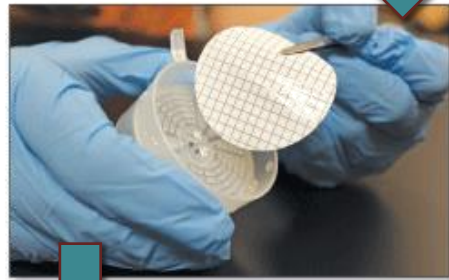
BERND HÄNFLING, NIGEL WILLBY, LORI LAWSON-HANDLEY, DAN READ, IAN WINFIELD *ET AL.*

# Environmental DNA (eDNA):

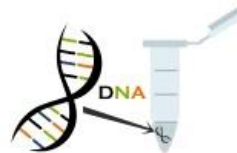


Cells, free-floating DNA  
(from sloughed skin cells  
faeces/urine, gametes,  
decaying matter)

Water sampled & filtered  
(0.5-2L samples)

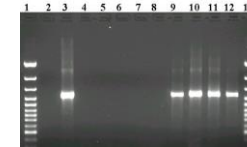


Extract DNA



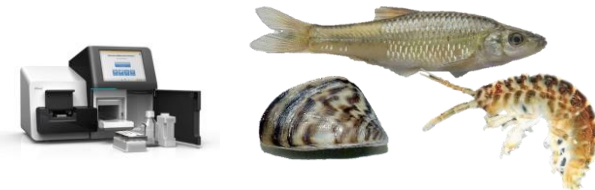
## Targeted detection:

- Species-specific primers
- Standard or qPCR



## Whole community:

- eDNA metabarcoding
- Conserved primers
- Next generation sequencing





Centre for Ecology & Hydrology  
NATURAL ENVIRONMENT RESEARCH COUNCIL



Environment Agency



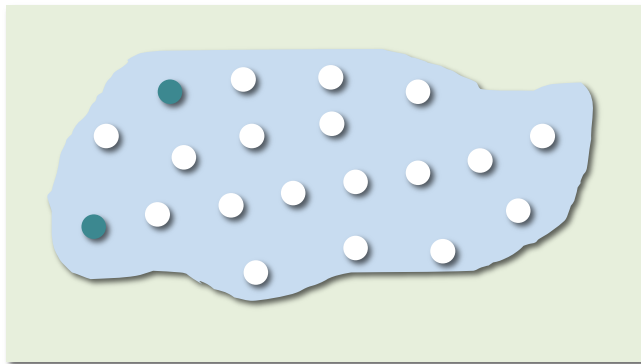
Scottish Environment Protection Agency

# Lake fish monitoring with eDNA metabarcoding

Hänfling et al. (2016) *Molecular Ecology*, **25**, 3101-3119

Li et al. (2019) *Journal of Applied Ecology*, **56**, 1232-1244

Lawson-Handley et al (2019), *Environmental DNA* **1**, 24-36



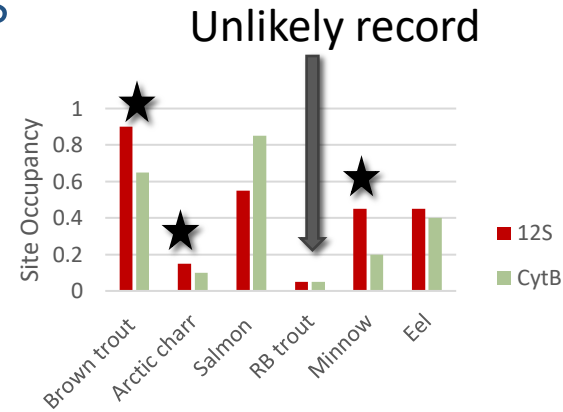
# eDNA metabarcoding has low false negative rates

Highland lakes  
e.g. Scottish Lochs,

Low alkalinity  
oligotrophic  
Deep



Loch Osgaig

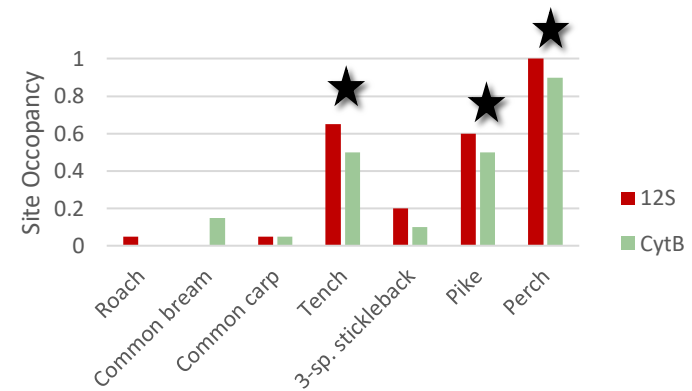


Lowland lakes  
e.g. Cheshire Meres

Medium alkalinity  
mesotrophic  
shallow

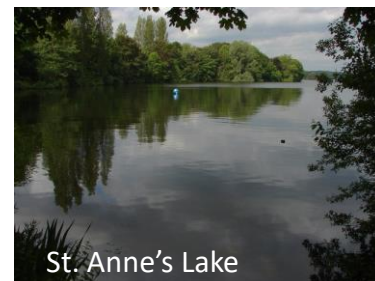


Chapel Mere

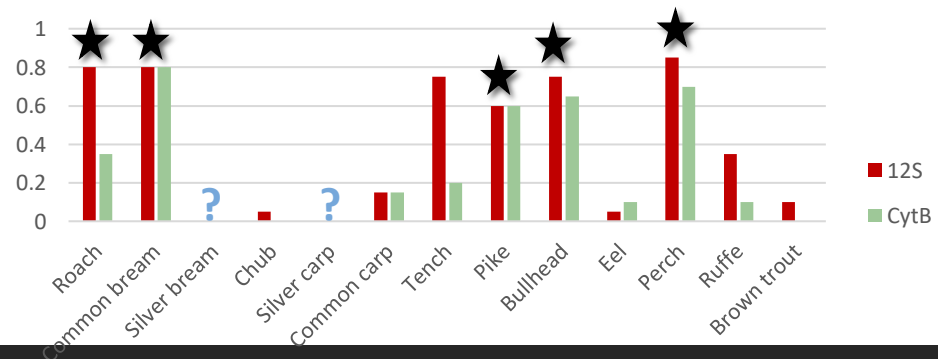


Lowland lakes  
e.g. Thames Flood Plain

Low alkalinity  
Very shallow  
eutrophic



St. Anne's Lake



? = species detected at > 2 sites but at low abundance (below threshold)

★ = species detected in netting surveys

# Implausible detections, “false positives”

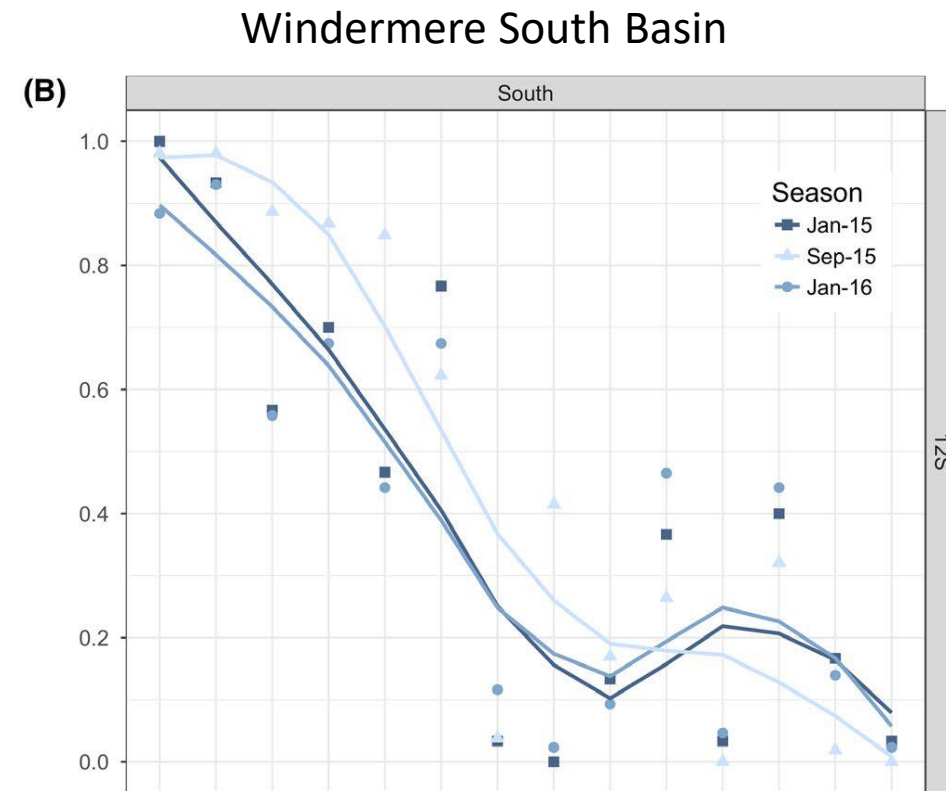
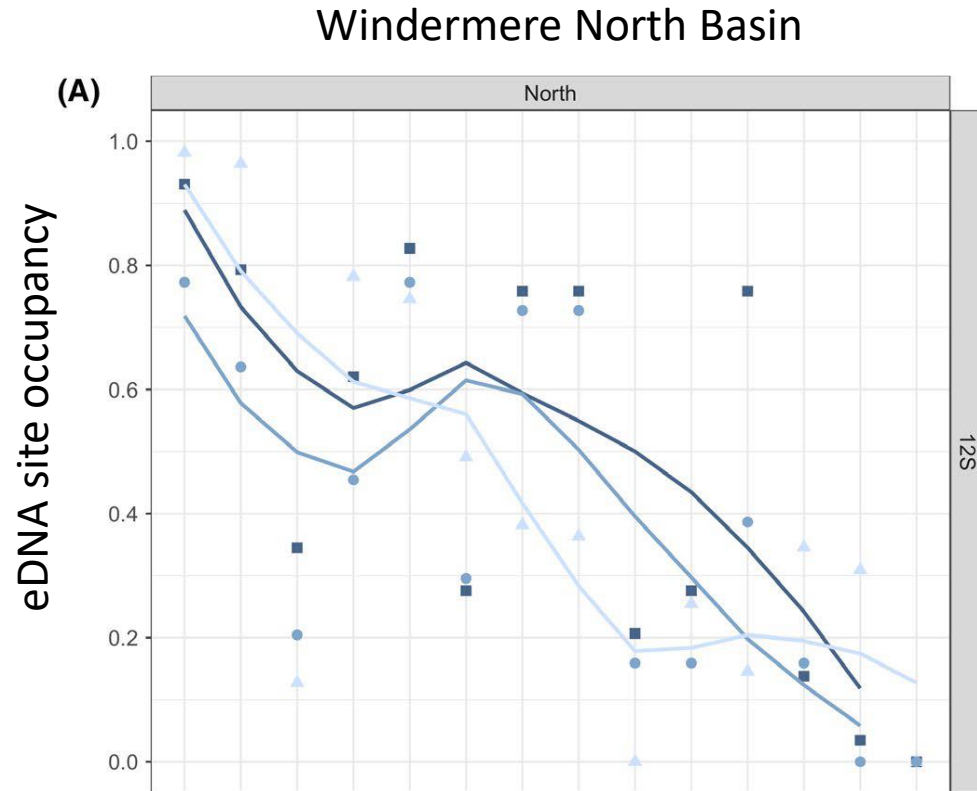
---

- Contamination during sampling
- Contamination in the lab
- Tag-jumping during sequencing
- Bioinformatics/reference data base
- DNA contamination of the environment



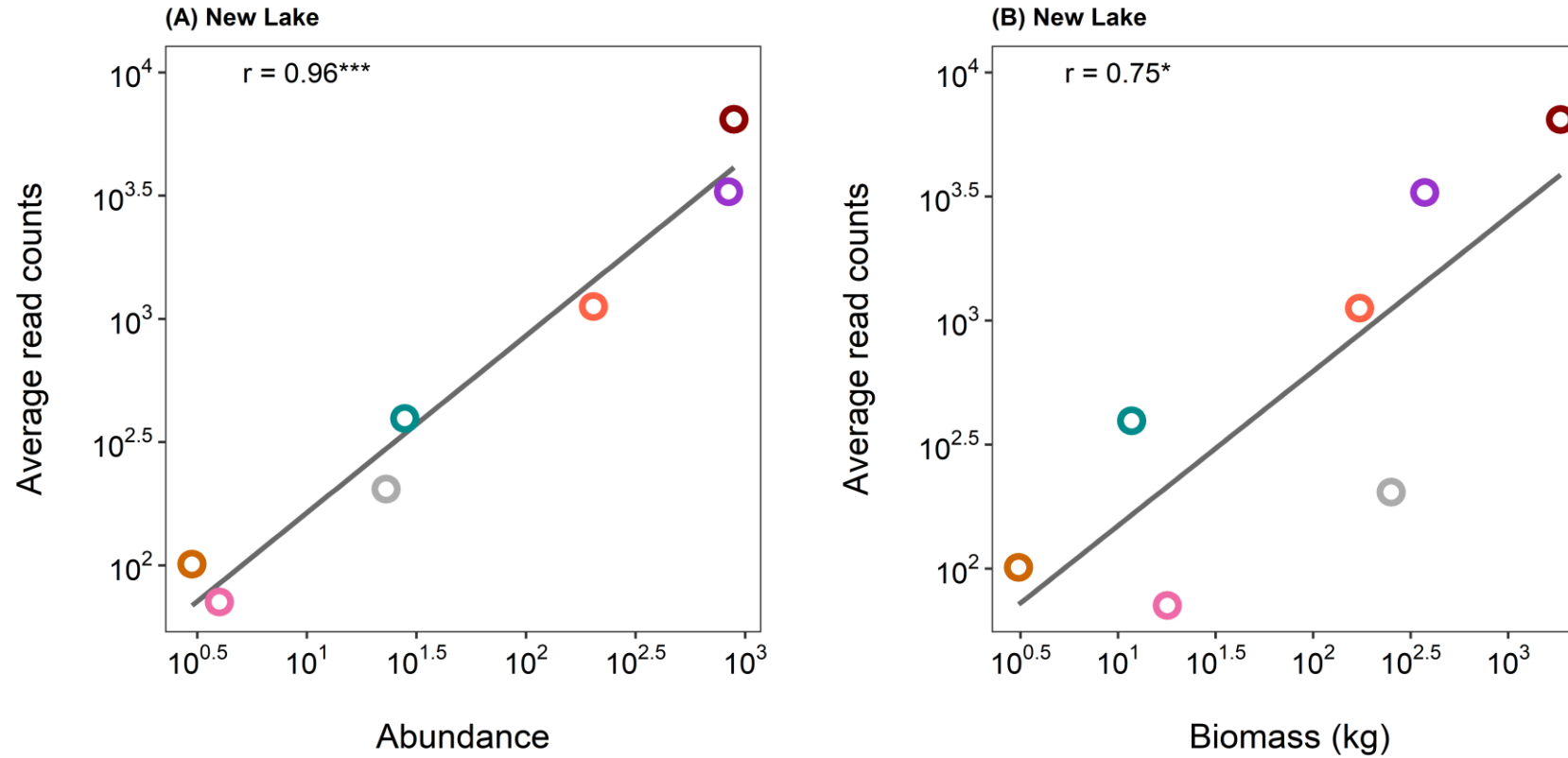
< 2% of all records in our data set

# eDNA abundance reflects relative fish abundance (e.g. Windermere)



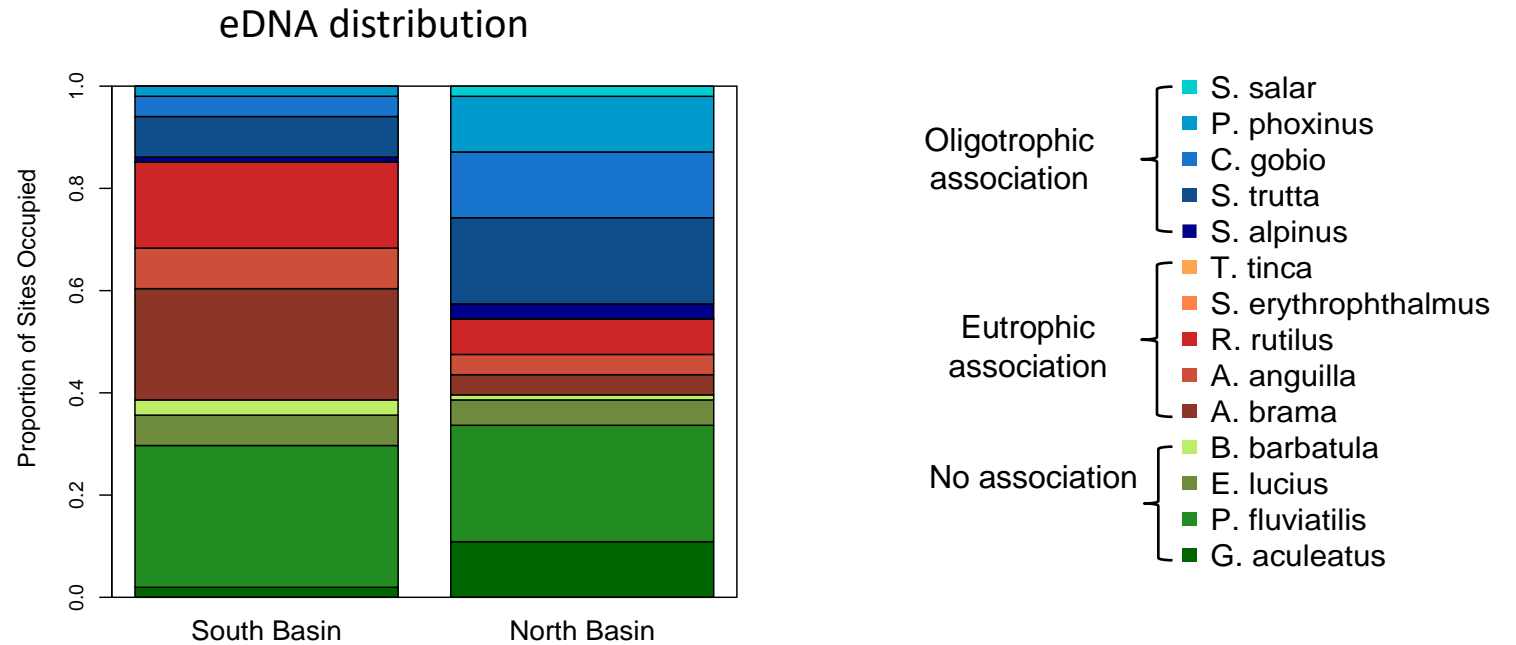
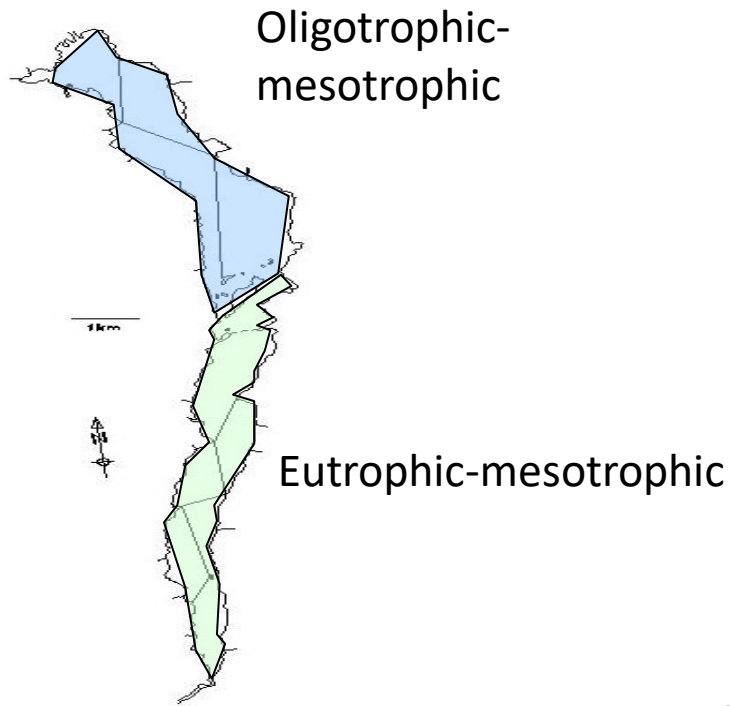
Species abundance rank based on long-term data

# eDNA site abundance reflects relative fish abundance (drained fishing lake)



- Abramis brama x Rutilus rutilus*
- Barbus barbus*
- Ctenopharyngodon idella*
- Perca fluviatilis*
- Scardinius erythrophthalmus*
- Silurus glanis*
- Squalius cephalus*
- Tinca tinca*

# eDNA distribution reflects lake ecology



North basin contains a higher proportion of oligotrophic species



# eDNA provides an accurate description of expected lake fish communities

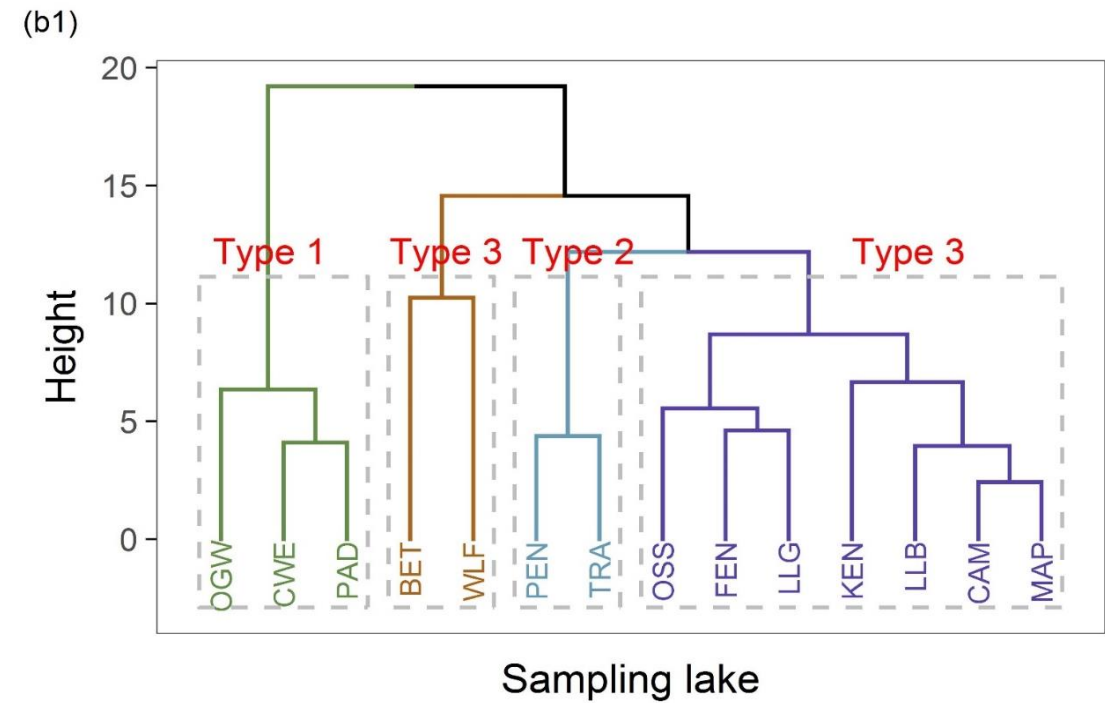
## eDNA metabarcoding of 14 Lakes in Wales and NW England

Pre-classified in 3 types

Type 1: Low alkaline deep upland lakes

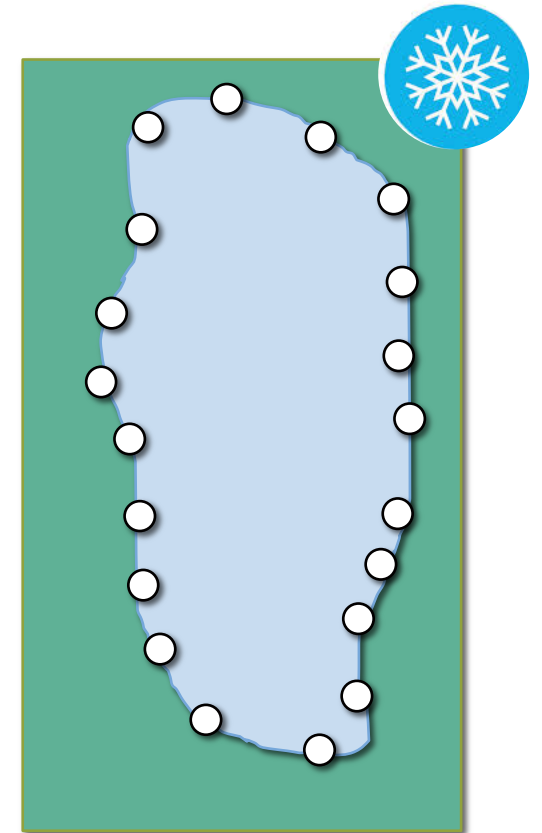
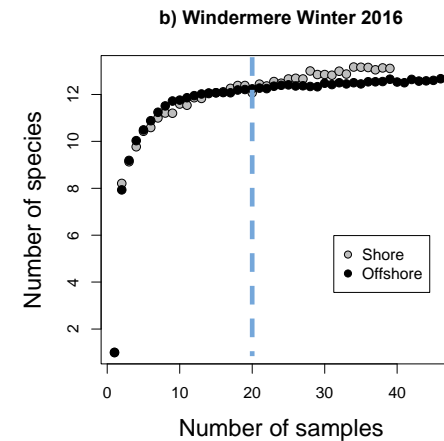
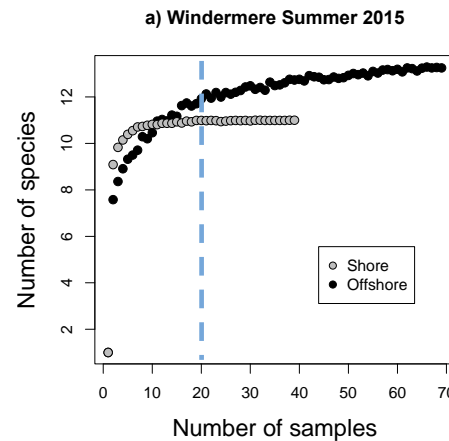
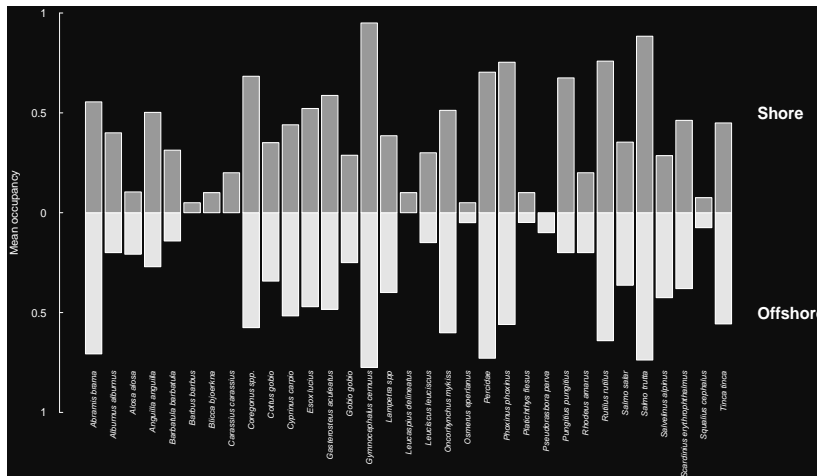
Type 2: High alkaline shallow coastal lakes

Type 3: High alkaline very shallow inland lakes

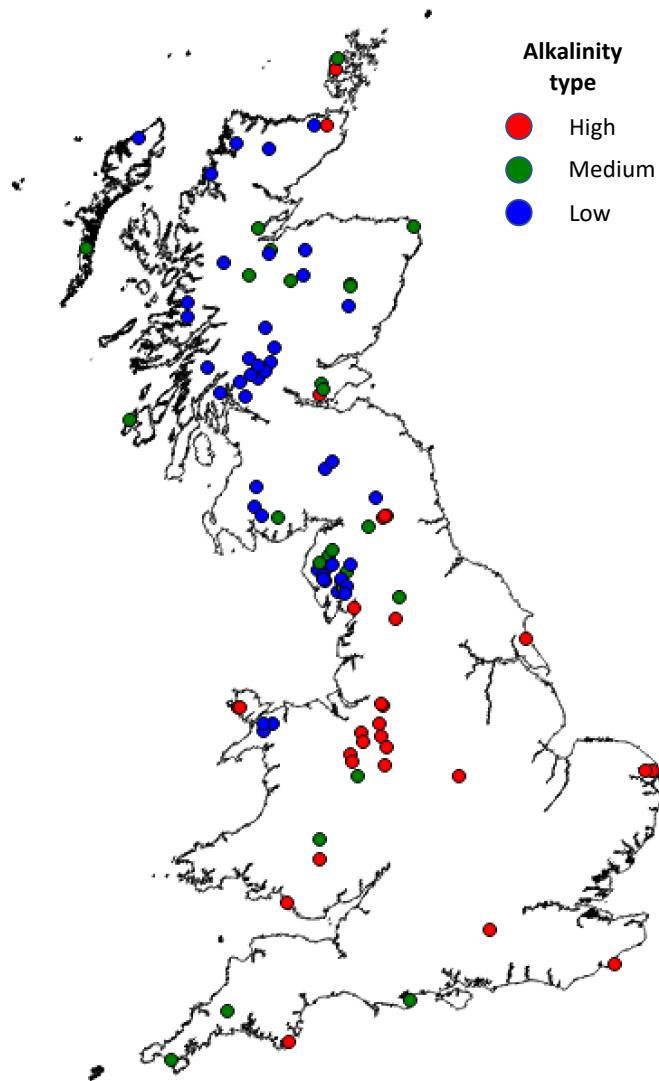


# Recommended sampling strategy for UK lakes (aim = biodiversity monitoring)

- Samples collected from the shore
- Winter sampling (November – February)
- 20 samples per lake (2 litres each)



# Building a data set of representative UK lakes



## 101 water bodies

- Wide geographical spread and coverage of lake types and status
- 28 lakes of “reference conditions”
- 40 fish taxa detected of which 16 restricted to <5% of sites
- Median richness 7 taxa per site (2-18).

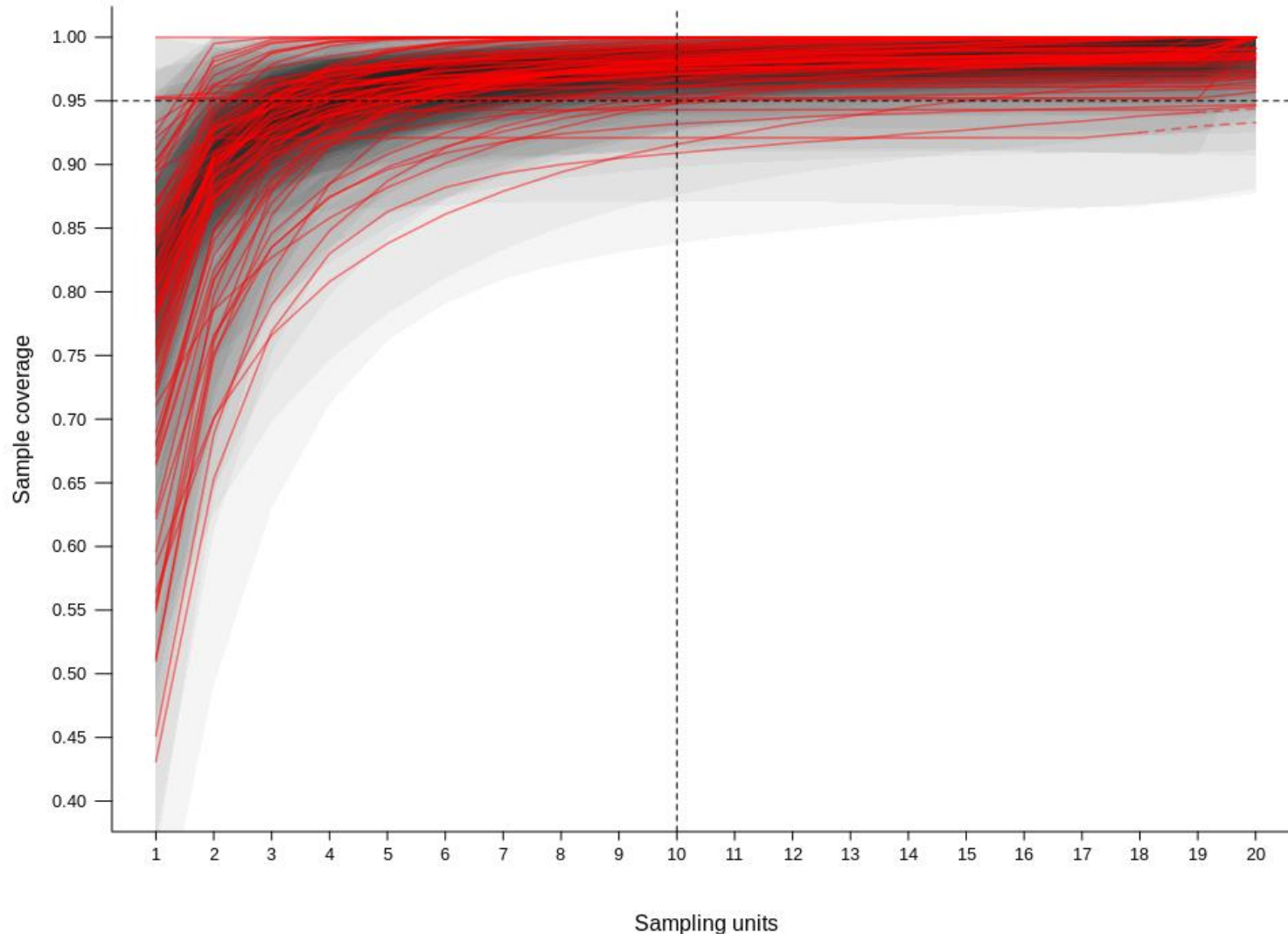
### Morpho-edaphic index:

Integrated measure of natural productivity

High = high alkalinity, shallow

Low = Low alkalinity, deep

## Sample coverage, based on Chao & Jost (2013)



How many samples are needed to reach 95% sample coverage (sampling threshold)?

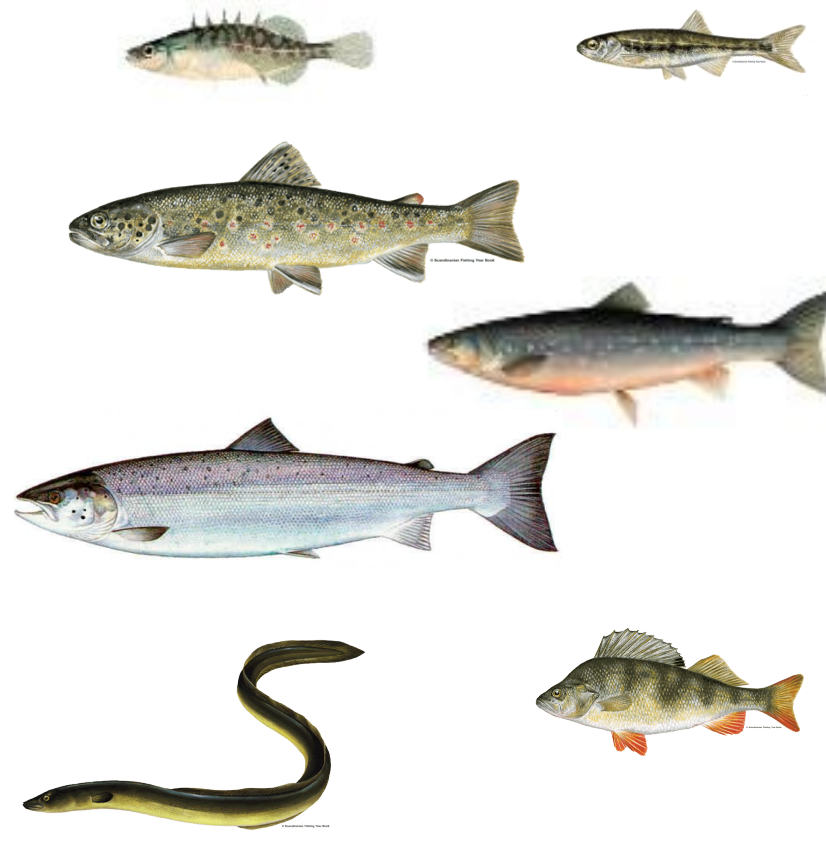
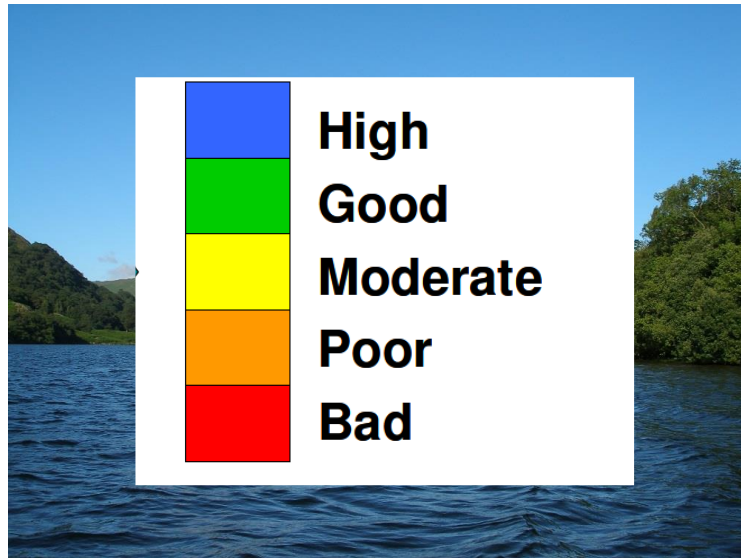
- Range: 1-25
- Mean = 5.7 (SD 4.7)
  
- 10 samples: 89% of 101 lakes
- 20 samples: 96% of 101 lakes

Sampling threshold vs **species richness**:  
 $r_s = 0.42, p < 0.05$

Sampling threshold vs **mean occupancy**:  
 $r_s = -0.83, p < 0.05$

Sampling threshold vs **lake size**:  
 $r_s = -0.09, p > 0.3$

# Can we detect anthropogenic pressure using species relative abundance alone where there are few species?

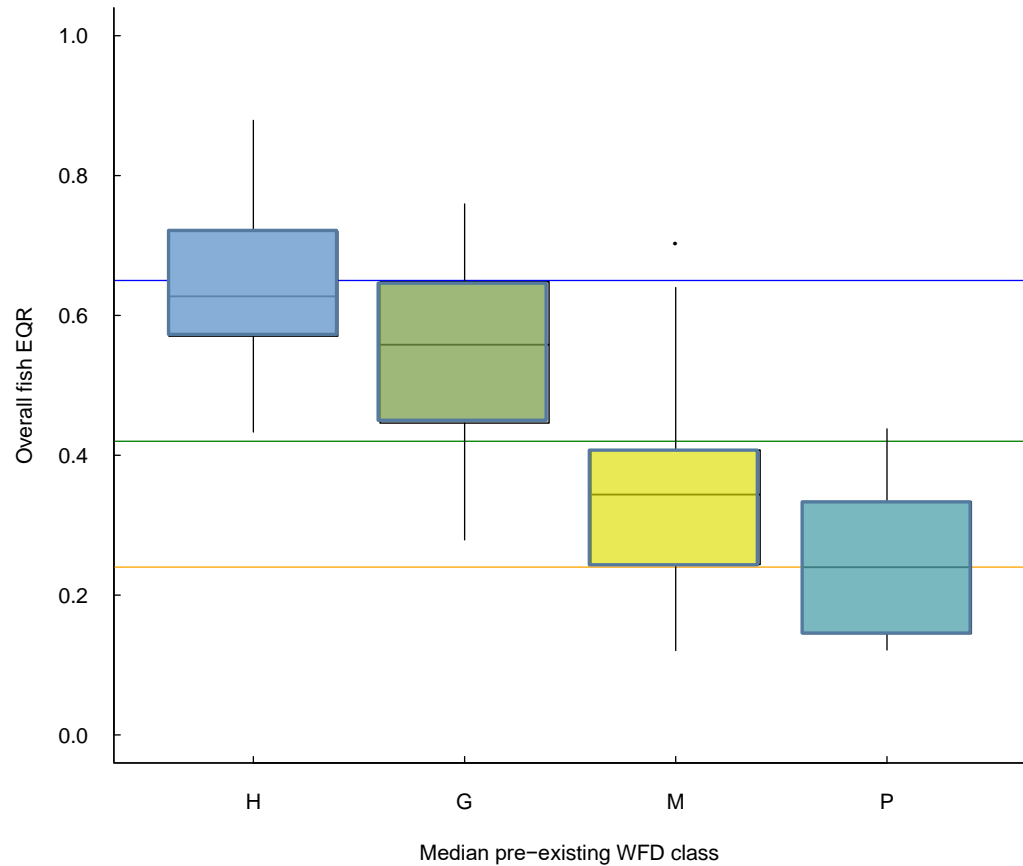


# Fish metrics considered for WFD tool

---

- Composition using **taxonomic data** – occupancy or read shares for 24 taxa found in >5% of sites
- Composition using **functional groupings/guilds** e.g. piscivores v insectivores, pelagic v littoral, phytophilous vs lithophilous. Limited utility
- **Taxon richness** per site
- Richness or read shares of **non-native taxa**
- Community index based on ranked position of each species on **eutrophication pressure axis** (ASPT for fish)

# Deriving class boundaries on fish EQR scale



Class boundaries of fish tool positioned to maximise class agreement and minimize classification bias, based on 'typical' pre-existing WFD class (WoE, not 1oAo) for eutrophication sensitive BQEs

|                  |     | fish class |    |    |     | sum |
|------------------|-----|------------|----|----|-----|-----|
|                  |     | H          | G  | M  | P/B |     |
| WFD median class | H   | 18         | 19 | 0  | 0   | 37  |
|                  | G   | 7          | 18 | 7  | 0   | 32  |
|                  | M   | 1          | 4  | 14 | 6   | 25  |
|                  | P/B | 0          | 1  | 2  | 4   | 7   |
| sum              |     | 26         | 42 | 23 | 10  | 101 |

54% same class, 99% +/- 1 class

Bias: -0.15 classes

# People contributing to this research



## UoH team:

Lori Lawson-Handley  
Christoph Hahn  
Helen Kimbell  
Paul Nichols  
Harriet Johnson  
Hayley Watson  
Rose Wilcox  
Rob Donnelly  
Joe Li  
Rosie Blackman  
Marco Benucci  
Lynsey Harper  
Cristina Di-Muri  
Graham Sellers

## Fera team:

Neil Boonham  
Eleanor Jones  
Ian Adams

## CEH team:

Ian Winfield  
Dan Read  
Anna Oliver  
Ben James  
Janice Fletcher

## Agency team:

Kerry Walsh  
Graeme Peirson  
Willie Duncan  
Alistair Duguid  
Sean Morrison  
Tristan Hatton-Ellis

## University of Stirling

Nigel Willby  
Colin Bull  
Alan Law

## Funding bodies

