

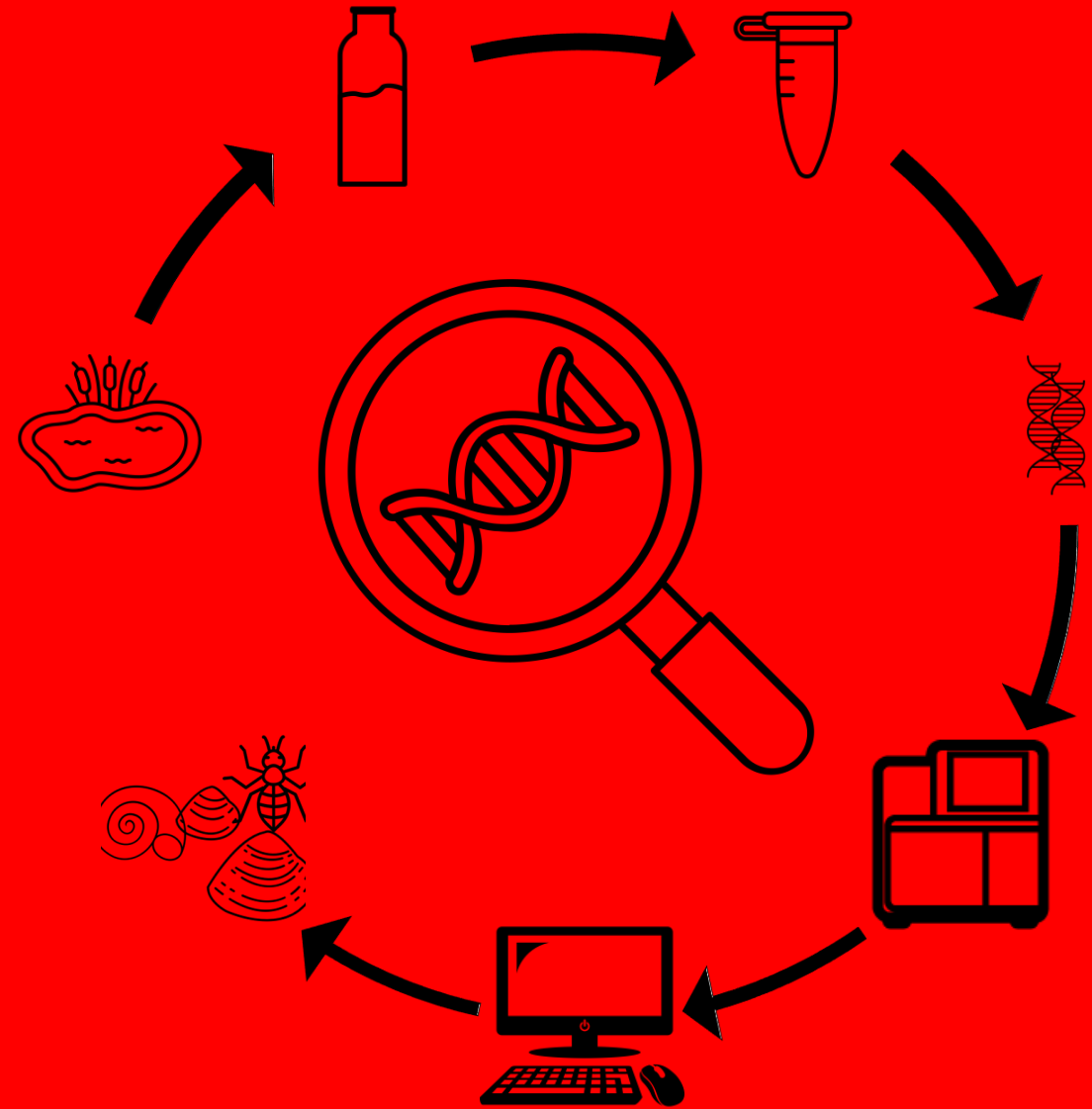
Environmental DNA

River University – 3 August 2023

Kari-Anne van der Zon

PhD Candidate in Ecology

Université de Strasbourg



Introduction

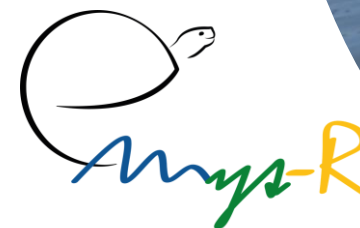
2012 – 2017: Bachelor Chemistry,
Utrecht University, Netherlands

2017 – 2020: Master Environmental Sciences,
Aquatic Ecology and Water Quality Management,
Wageningen University and Research, Netherlands

Since 2022 : PhD Research **“Ecology of man-made pond networks
for wetland biodiversity conservation”**,

University of Strasbourg, France

Supervised by Isabelle Combroux, Kathrin Theissingen
and Corinne Grac



www.emysr.cnrs.fr

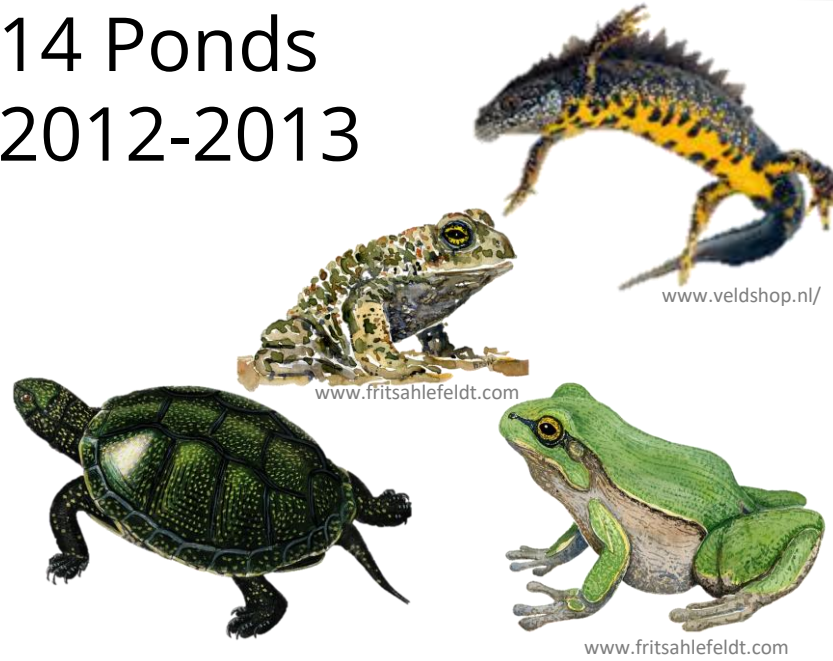
Woerr, France

6 Ponds, 2 lagunas
2011-2012 or 2015
+ 2 basins and gravel pit
rehabilitations



Neuburg, Germany

14 Ponds
2012-2013



Silene, Latvia

27 ponds
2013 or 2018



Actions

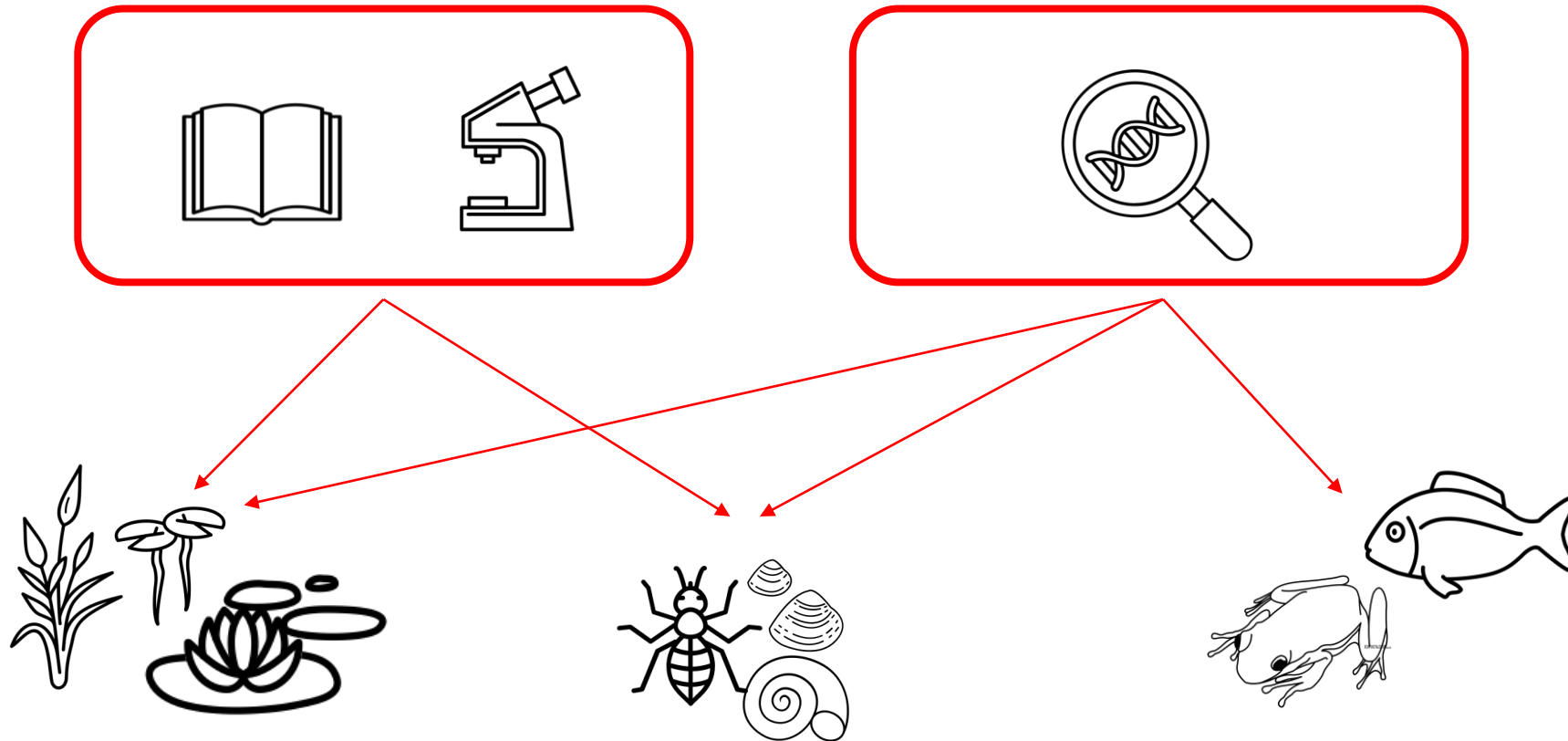
Target species

Example of pond

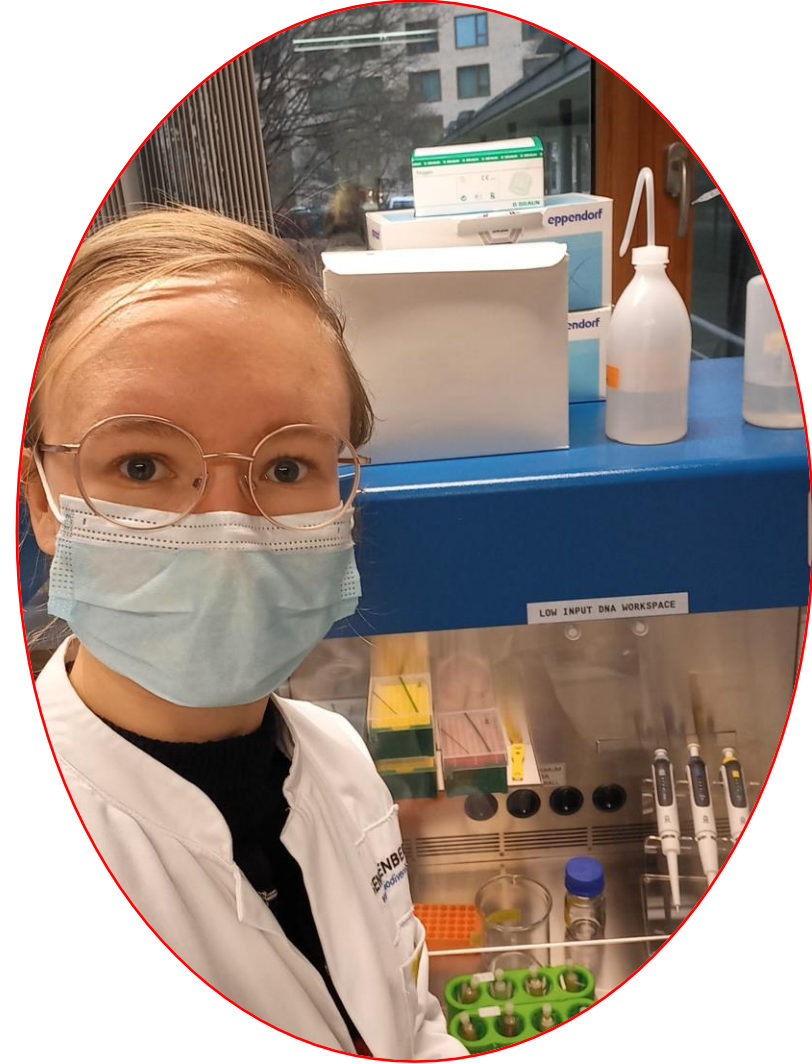


My project

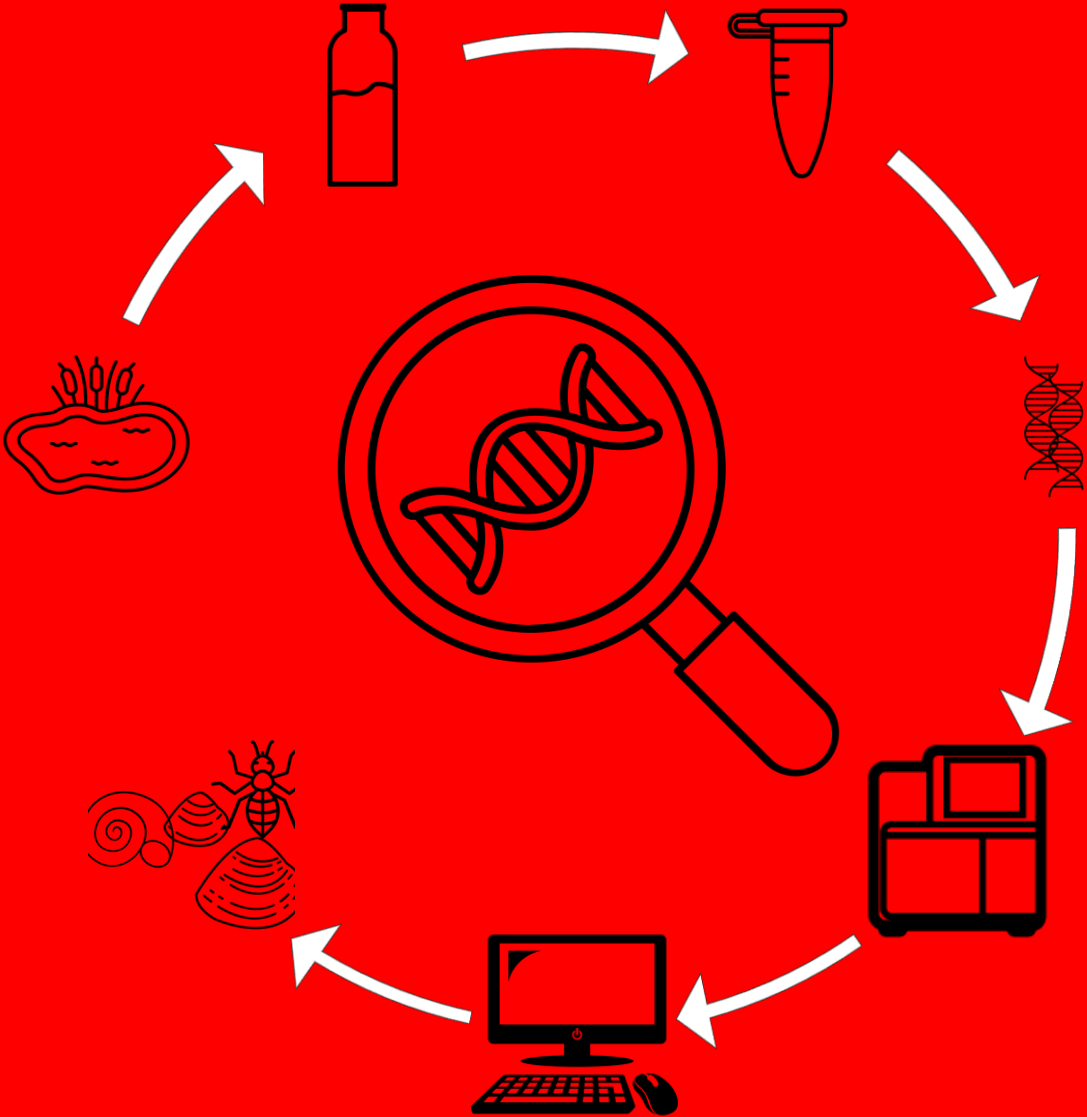
What variables determine the **biodiversity** and **habitat provisioning function** of **created pond networks**?



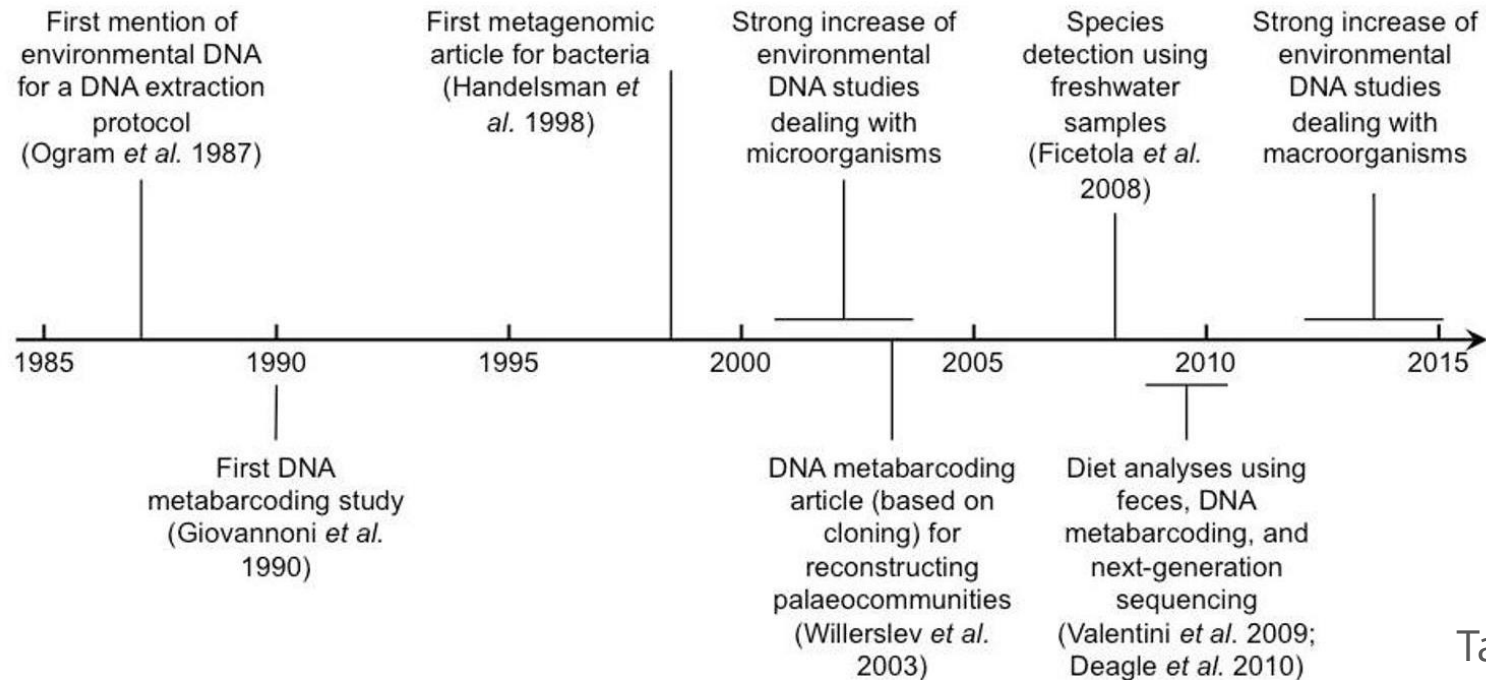
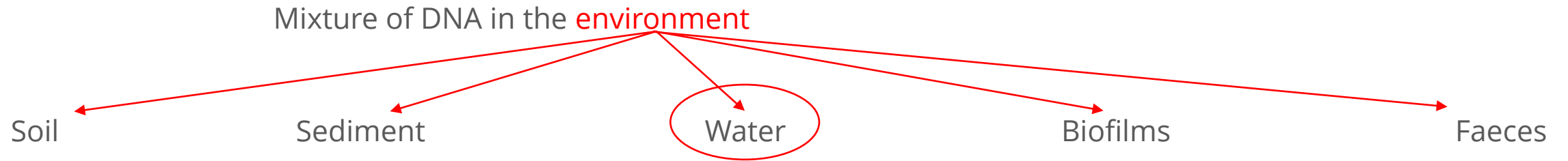
eDNA work in my project



Freshwater Environmental DNA



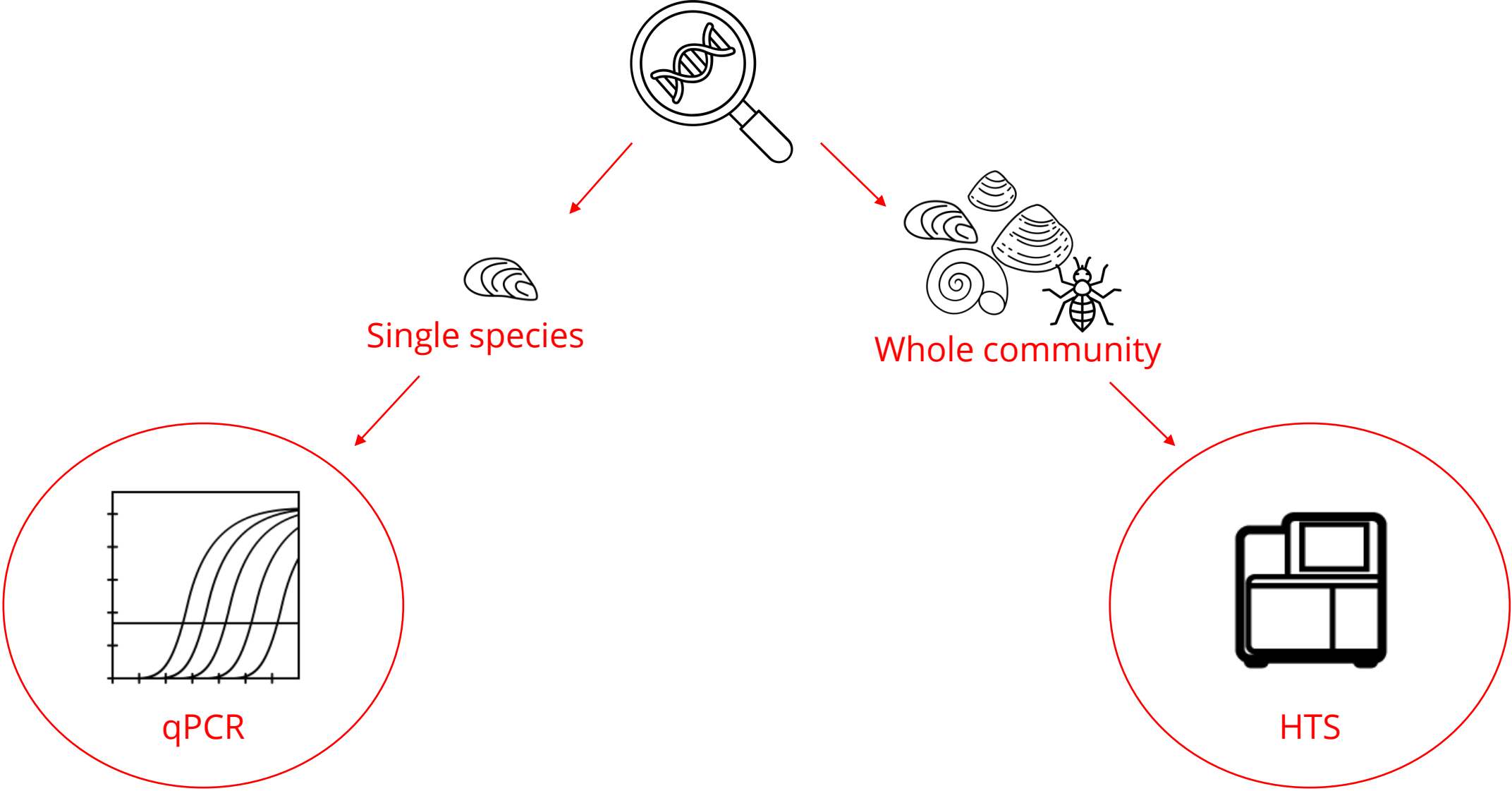
What is environmental DNA?



A young field

Taberlet *et al.* (2018)

Barcoding and metabarcoding



Example barcoding: Great Crested Newt survey

Traditional survey:

4 to 6 visits between mid-March and mid-May

eDNA qPCR:

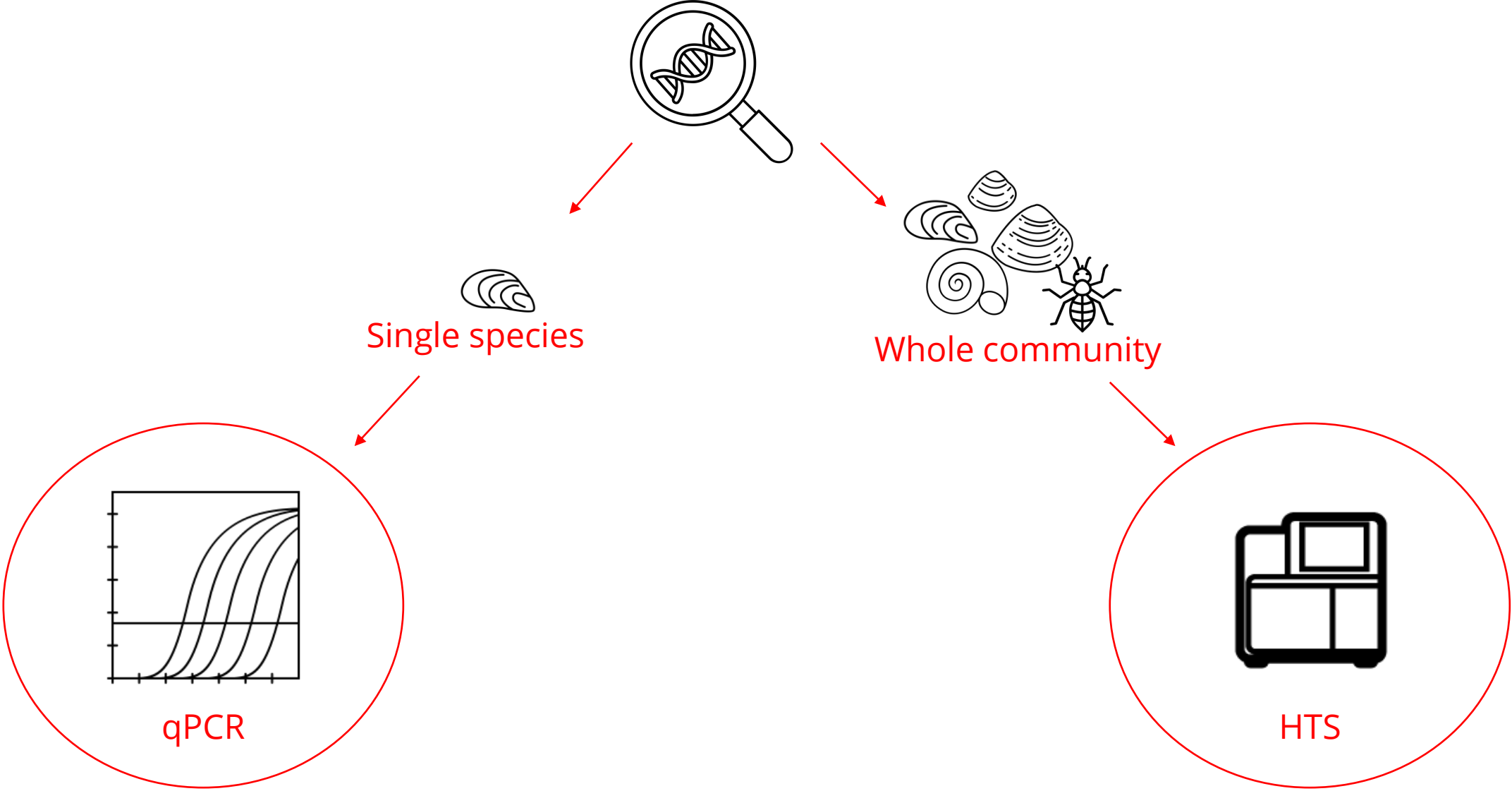
1 visit between March and August



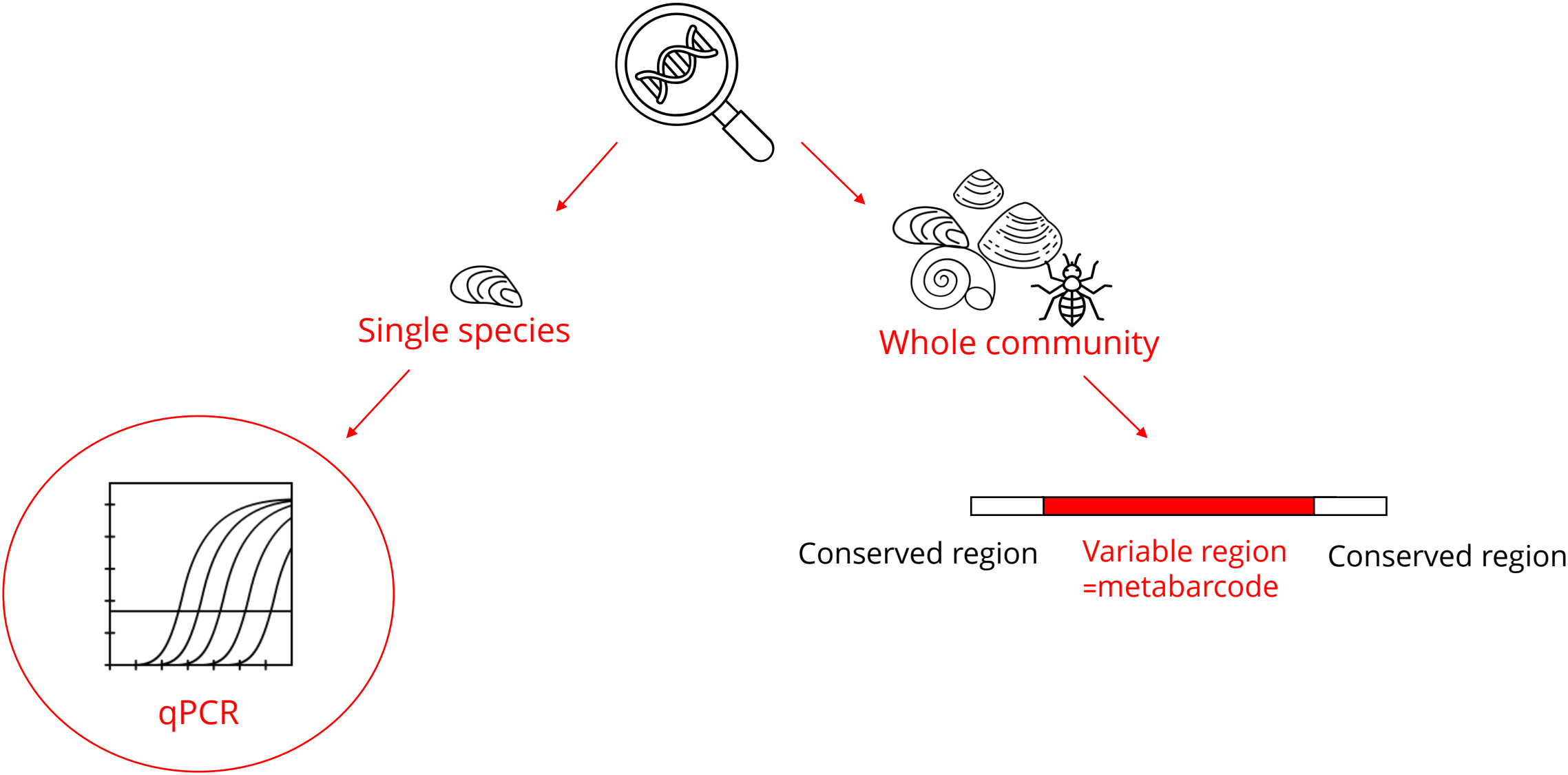
Rees et al. (2014), Rees et al. (2017)

Triturus cristatus

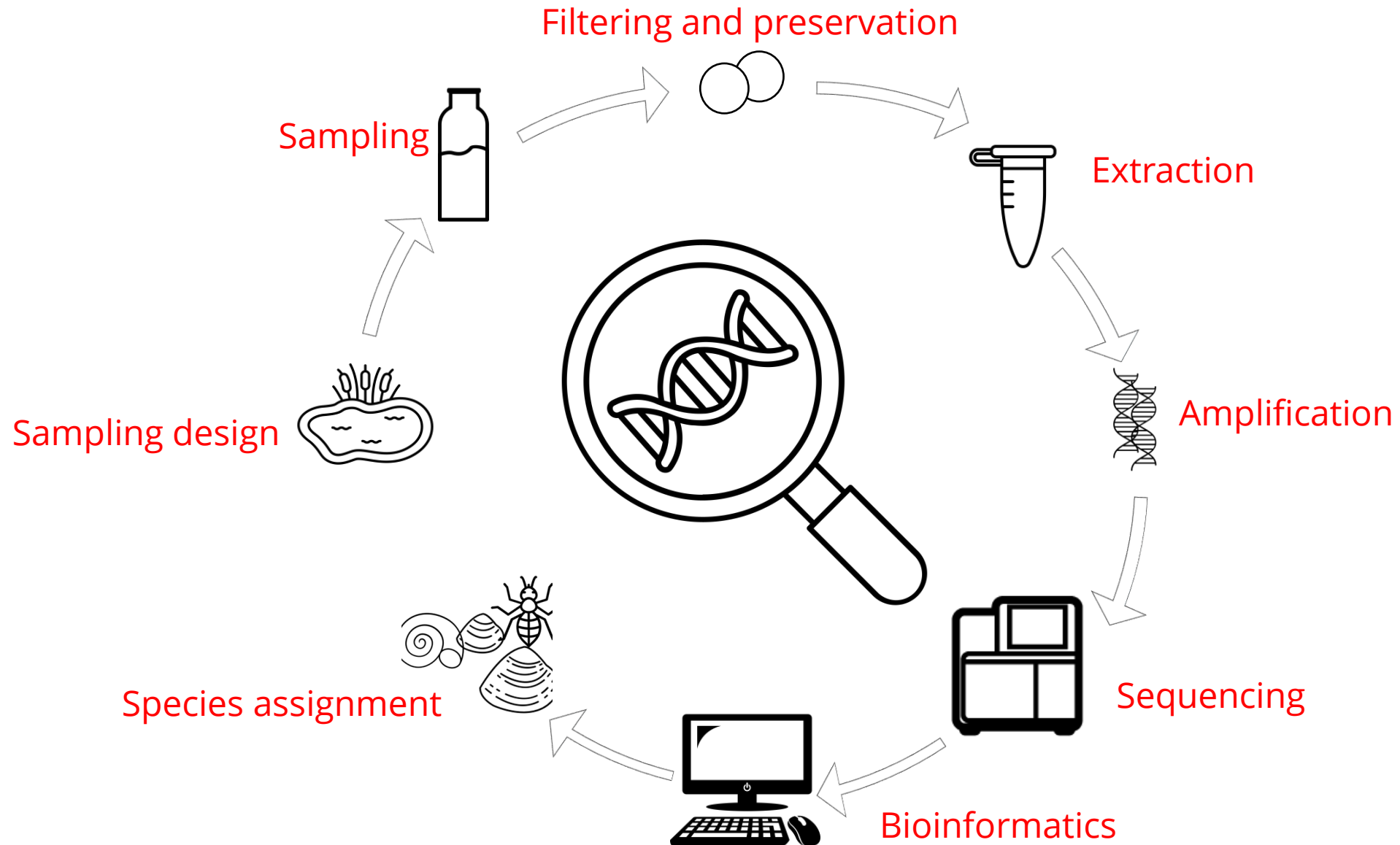
Barcoding and metabarcoding



Barcoding and metabarcoding



Aquatic eDNA metabarcoding workflow



Comparison with traditional methods

Keck et al. (2022)

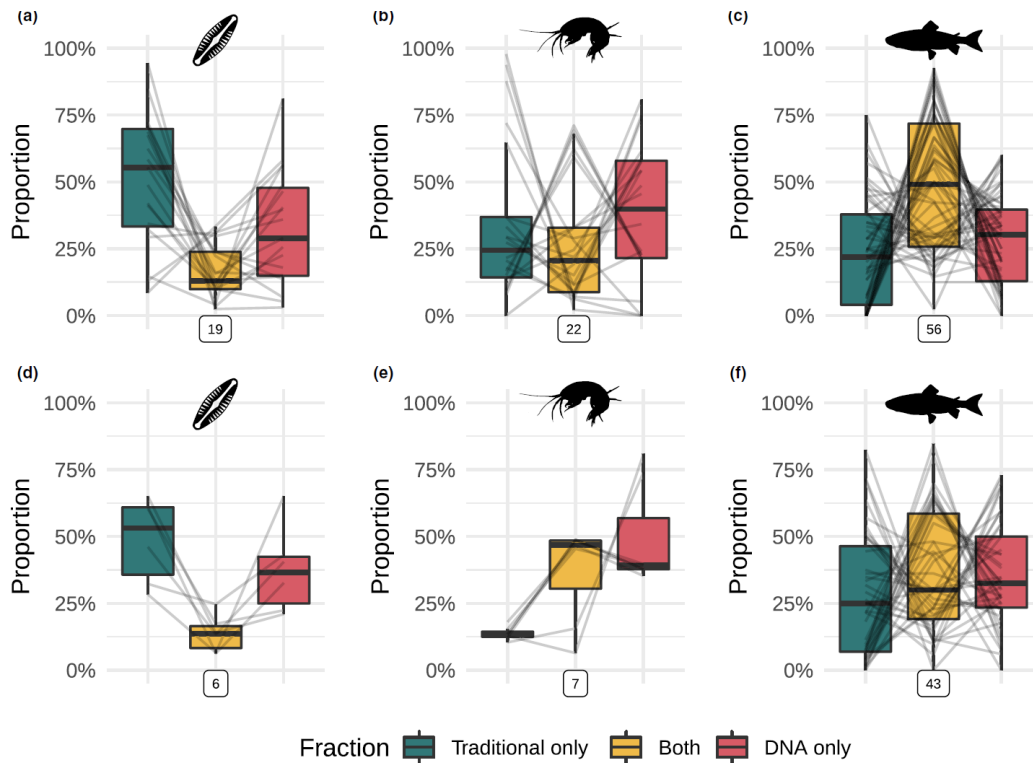


FIGURE 4 Relative fraction of diversity detected by the traditional method only, by DNA metabarcoding only and by both methods. Data are presented for different groups of organisms identified at species level only. Boxplots show medians, first and third quartiles, and full ranges (limited to $1.5 \times$ interquartile range). Grey lines connect values from the same comparison. Framed numbers below each panel indicate the number of comparisons represented. (a-c) Gamma diversity for plankton and microphytobenthos, macroinvertebrates and fish. (d-f) Alpha diversity for plankton and microphytobenthos, macroinvertebrates and fish

Fish: as good as or better than traditional methods

Results plankton, microphytobenthos and macroinvertebrate communities very different from traditional methods

Plant aquatic eDNA metabarcoding

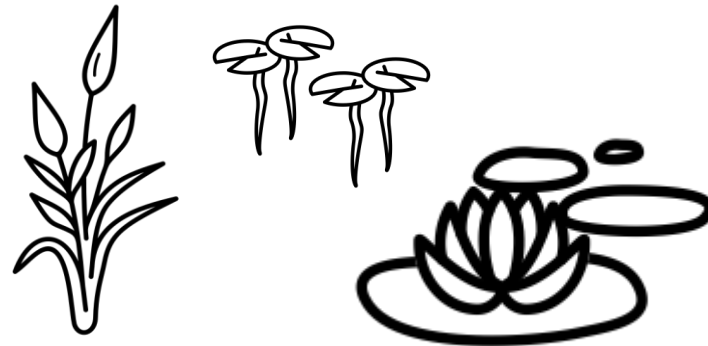
Traditional survey :

When plants flower

Overestimate floating leaved big plants

eDNA metabarcoding :

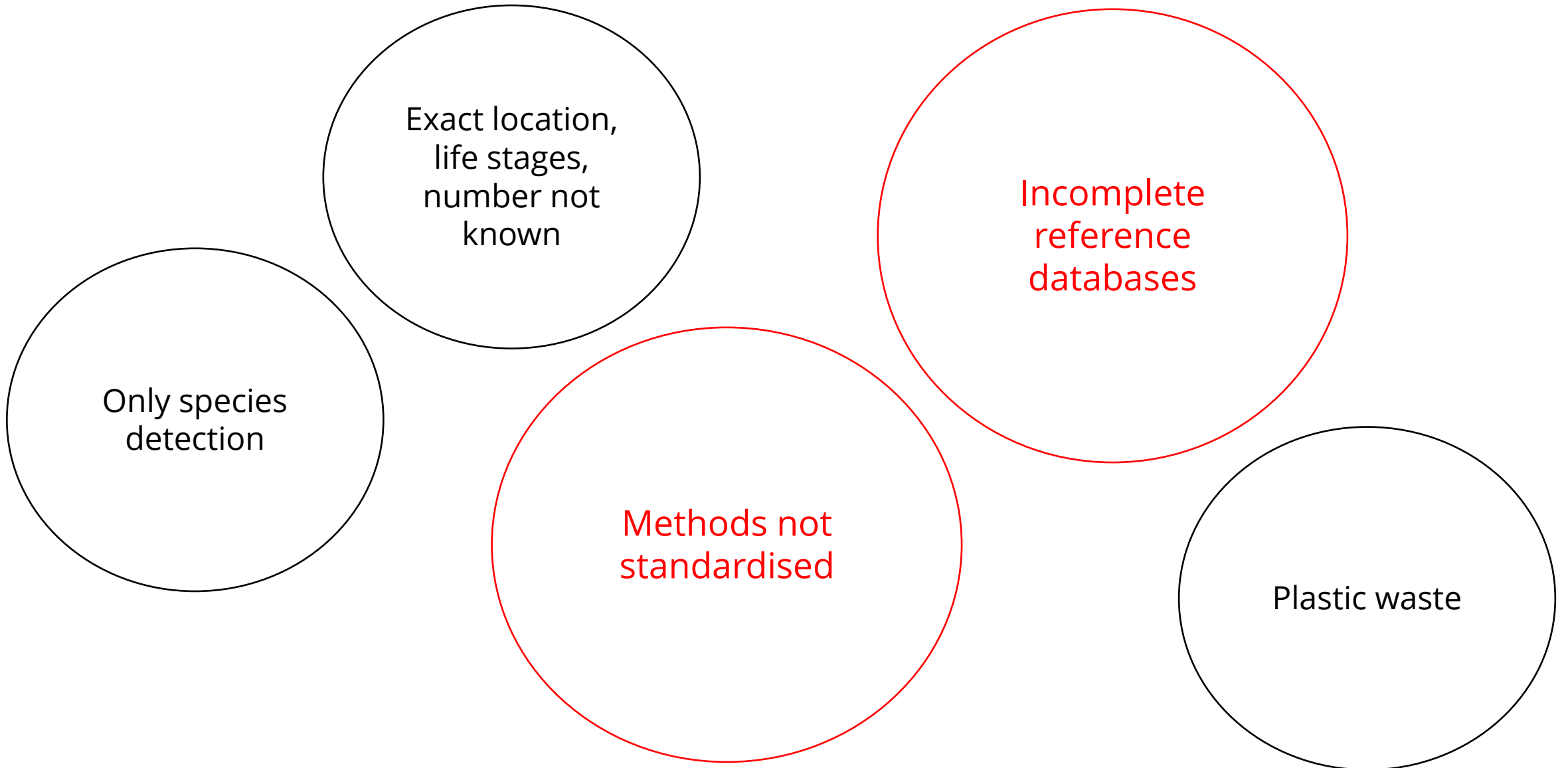
When plants are present



Need multiple
metabarcodes

Espinosa Prieto et al. (2023)

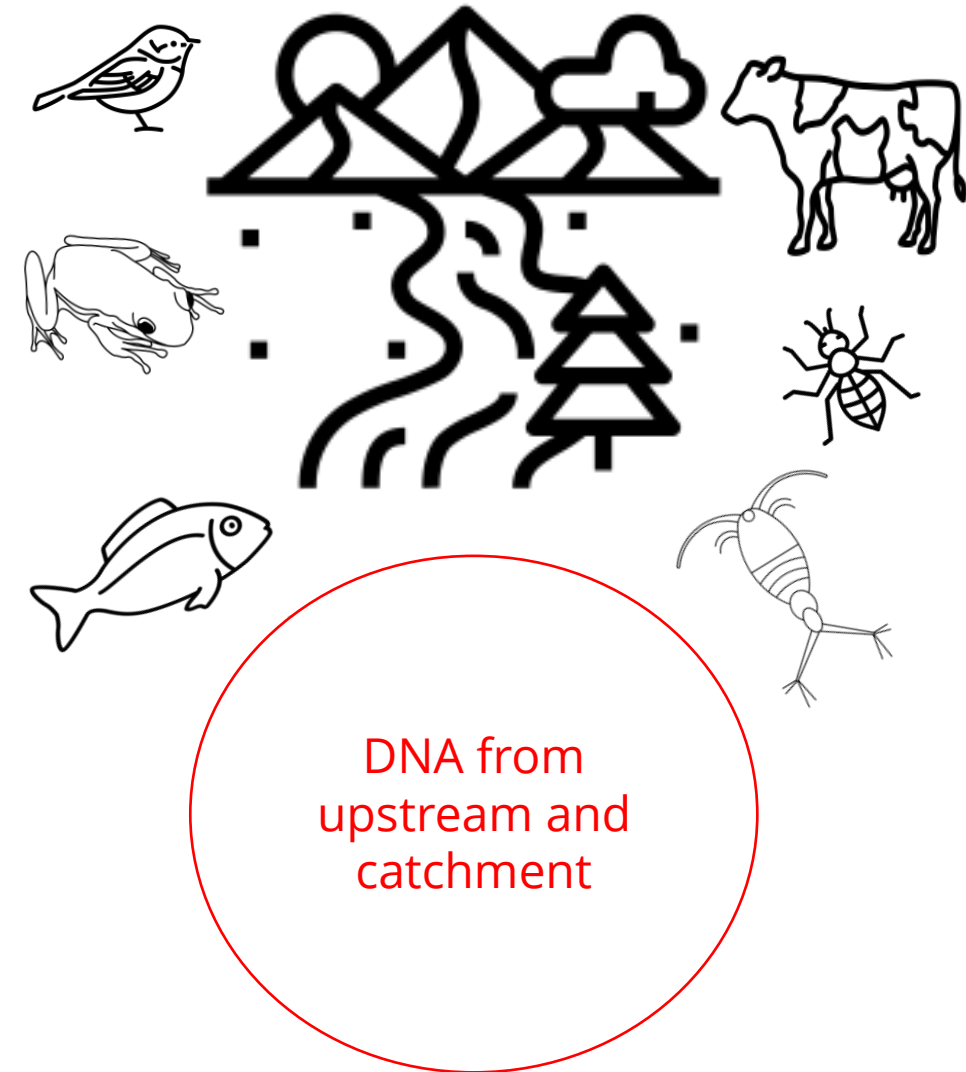
Constraints



Example: whole river biodiversity assessment

Cannon et al. (2016)

Group	Primer pair	No. species detected
Mammals	16Smam	17
Insects+Arachnids	COI_ZBJ_Art	15
Birds	Aves12S	8
Fish	FishCB	8
Copepods	Cop28S	4
Amphibians and reptiles	AmpCB	2
Vascular plants	trnL	-
Algae	23SrDNA	-
Fungi	FungusITS	-
Bryophytes	BryoTrnL	-
Diatoms	Diatom18S	-
Archaea	Archaea16S	-



Example: bivalve biodiversity

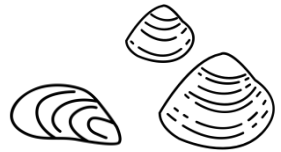
Traditional surveys:

Difficult to find

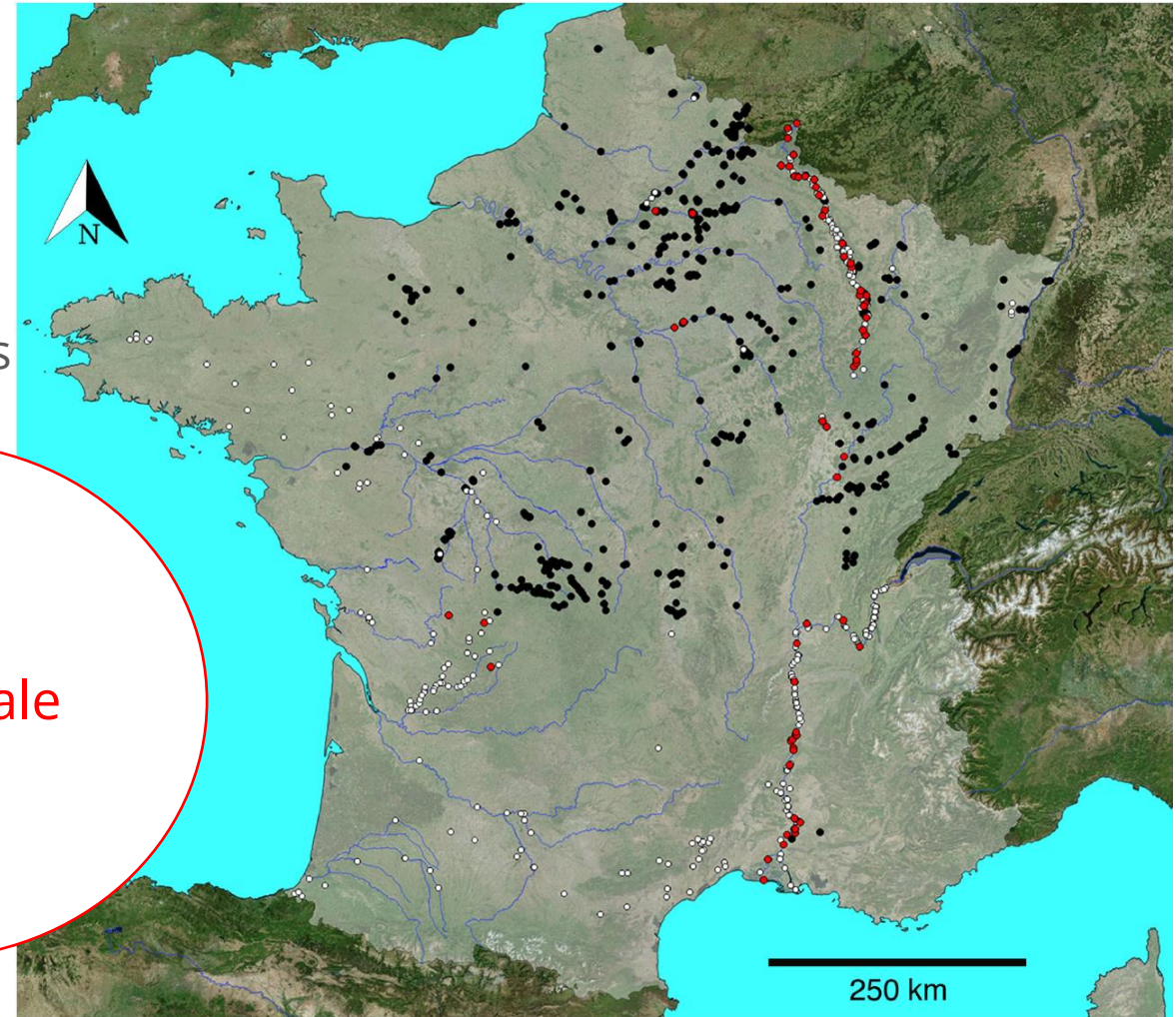
Difficult to identify

Different methods (scuba divers) different species

Often need to kill



Large scale



Prié et al. (2020)

Fig. 6 *Unio crassus* in France. White dots: eDNA sampling sites with absence; red dots: eDNA sampling sites with presence; black dots: all available data prior to eDNA

deployment, with no limit of time. The data acquired down stream the Rhône River and in the south-west is completely new

Conclusions

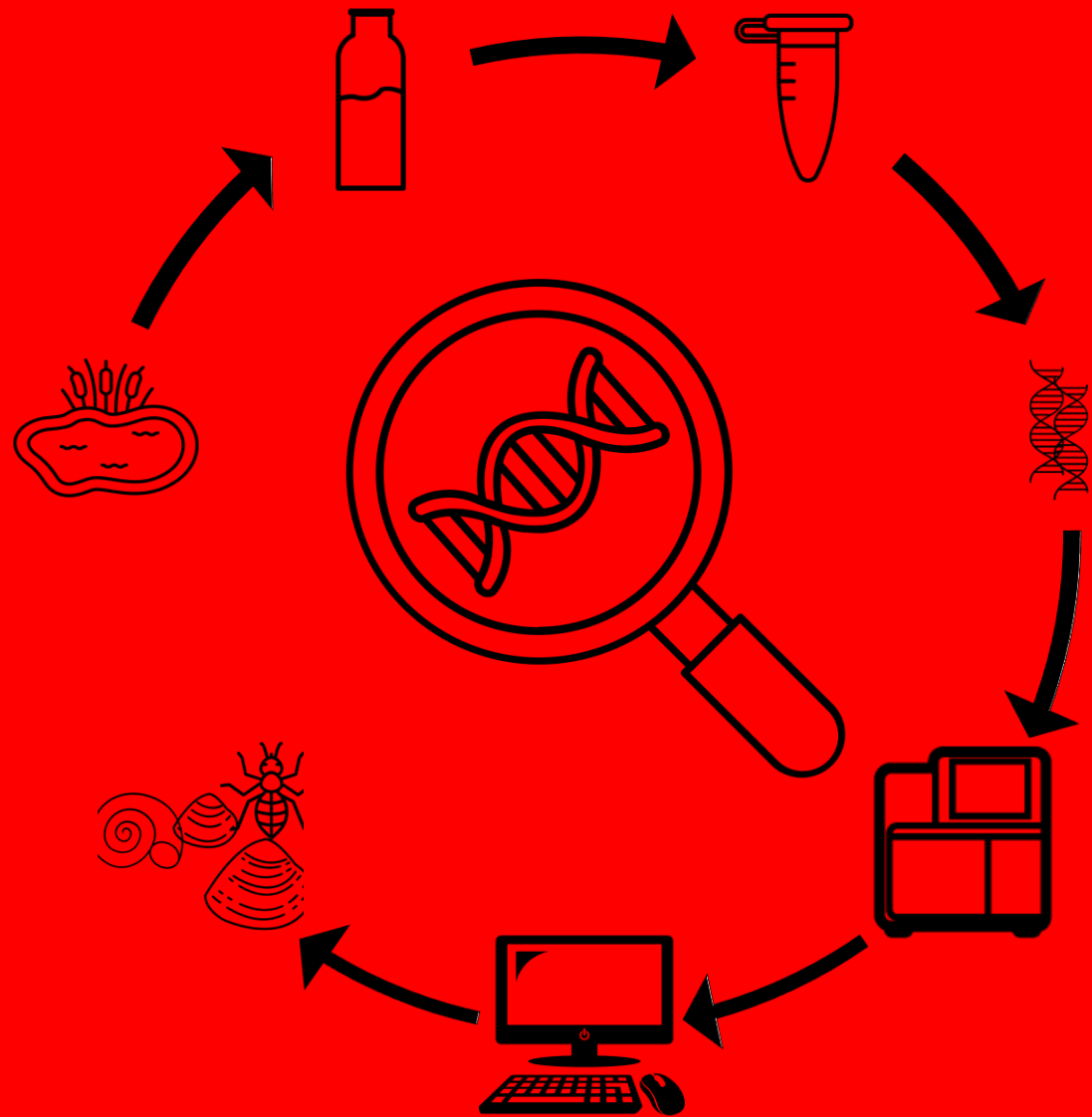
Applied for
detection of
invasive and
rare species

Works well for
fish, amphibian
and bivalve
communities

Results plankton,
microphytobenthos
and
macroinvertebrate
communities very
different from
traditional methods

“Another tool in
the toolbox”

Questions?



References

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