

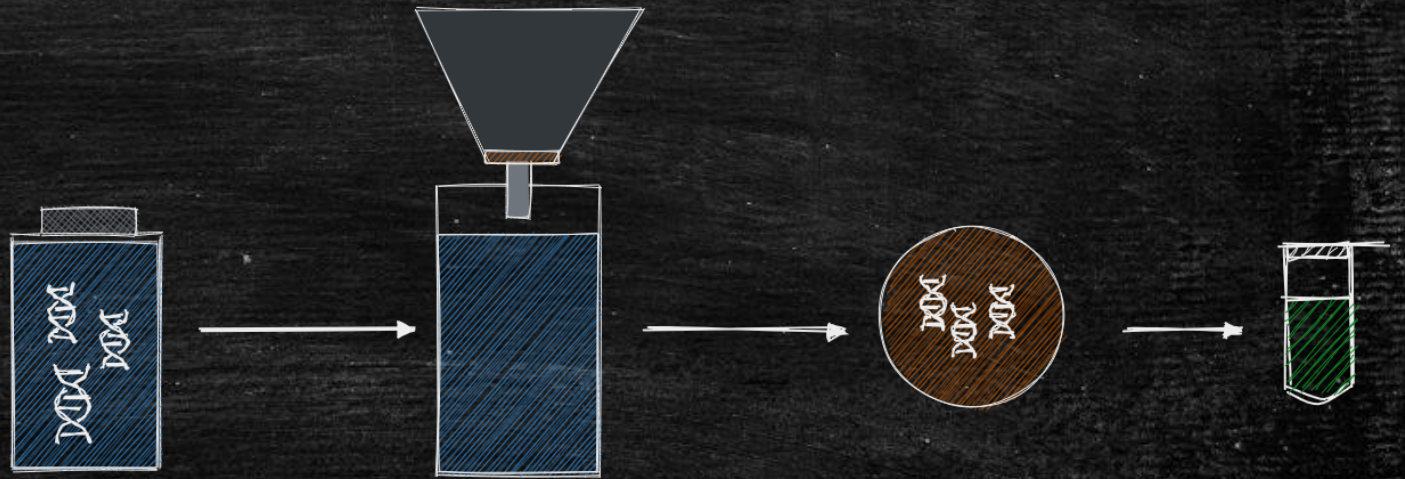
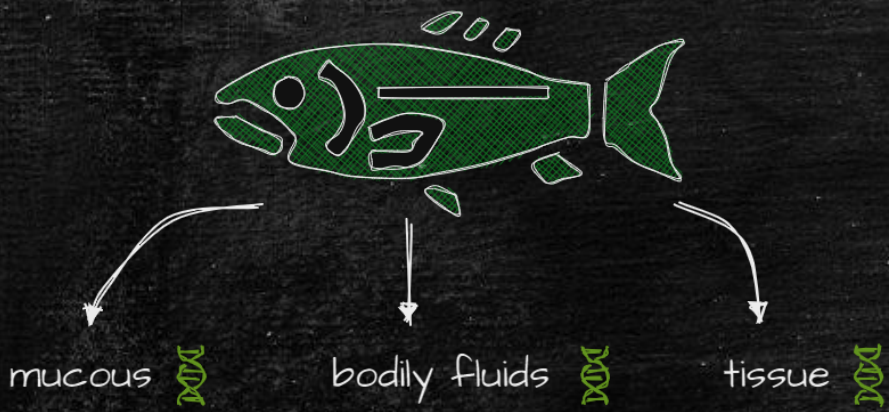
Environmental DNA effectively characterizes pre-construction fish communities at offshore wind sites

Shannon J. O'Leary, Sam Chin, Jason E. Adolf, Keith J. Dunton

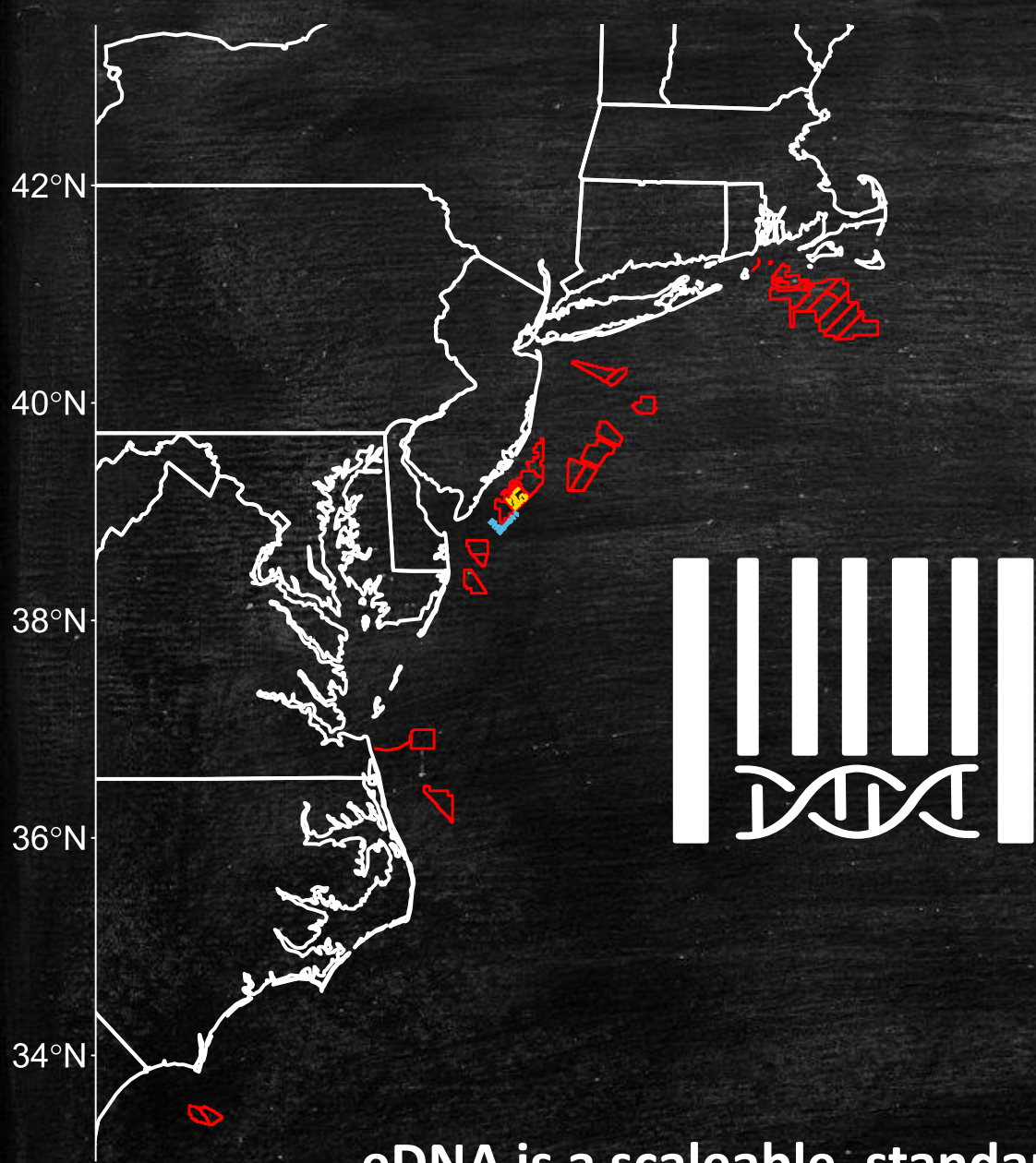
DNA is shed as cellular or extracellular material into the surrounding water

collect & filter water from aquatic systems

extract DNA from filters

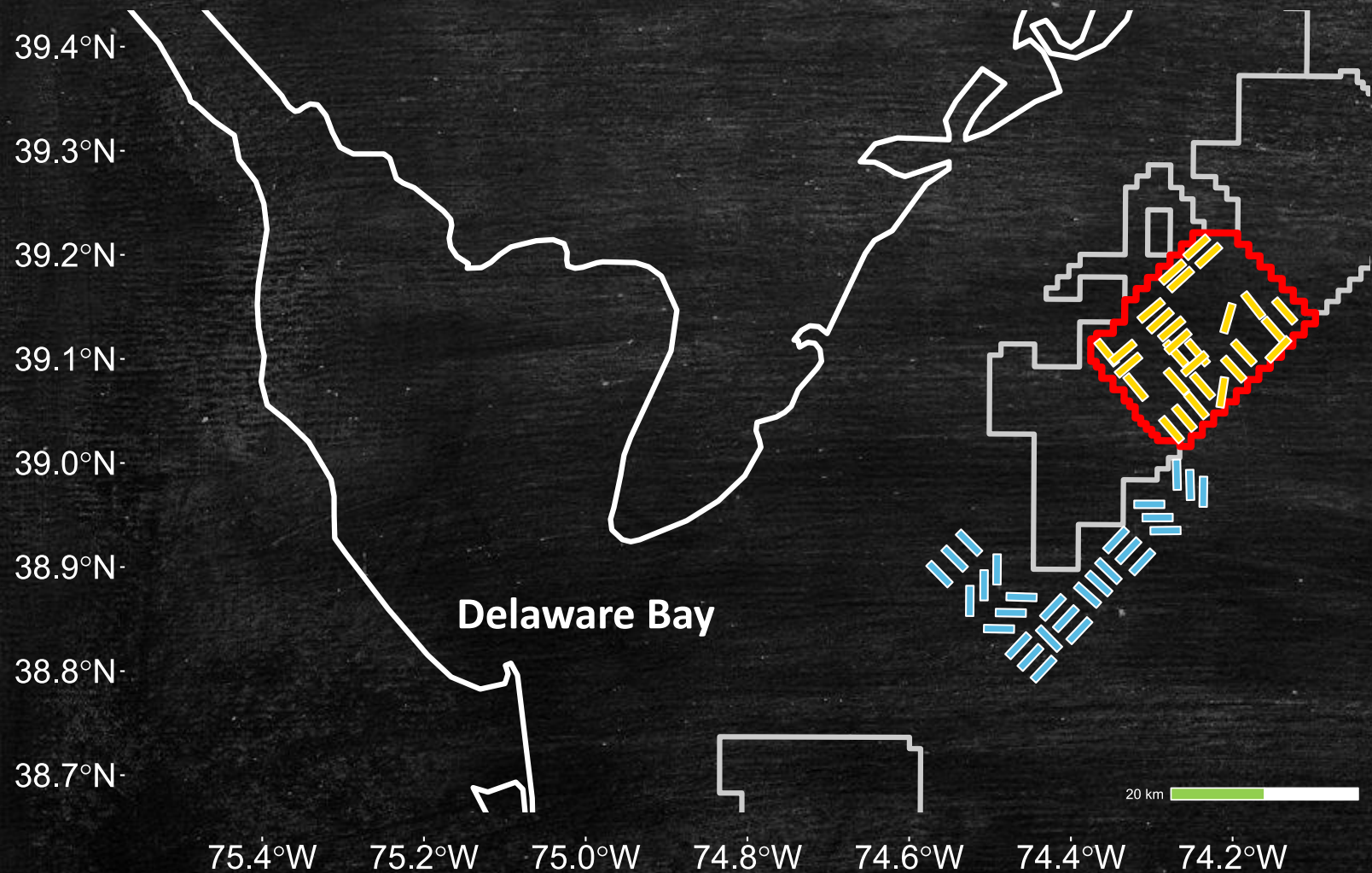


Environmental DNA is DNA captured from an environmental sample without the need for pre-isolating specific targets



- Low cost methods (scaleable)
- Simple sampling protocol (standardize across habitats)
- Easy access & few permitting issues
- minimal damage to habitat, target/bycatch species

eDNA is a scaleable, standardized, low-impact, low-equipment tool monitoring of the impact of large structures such as windfarms on ecosystems

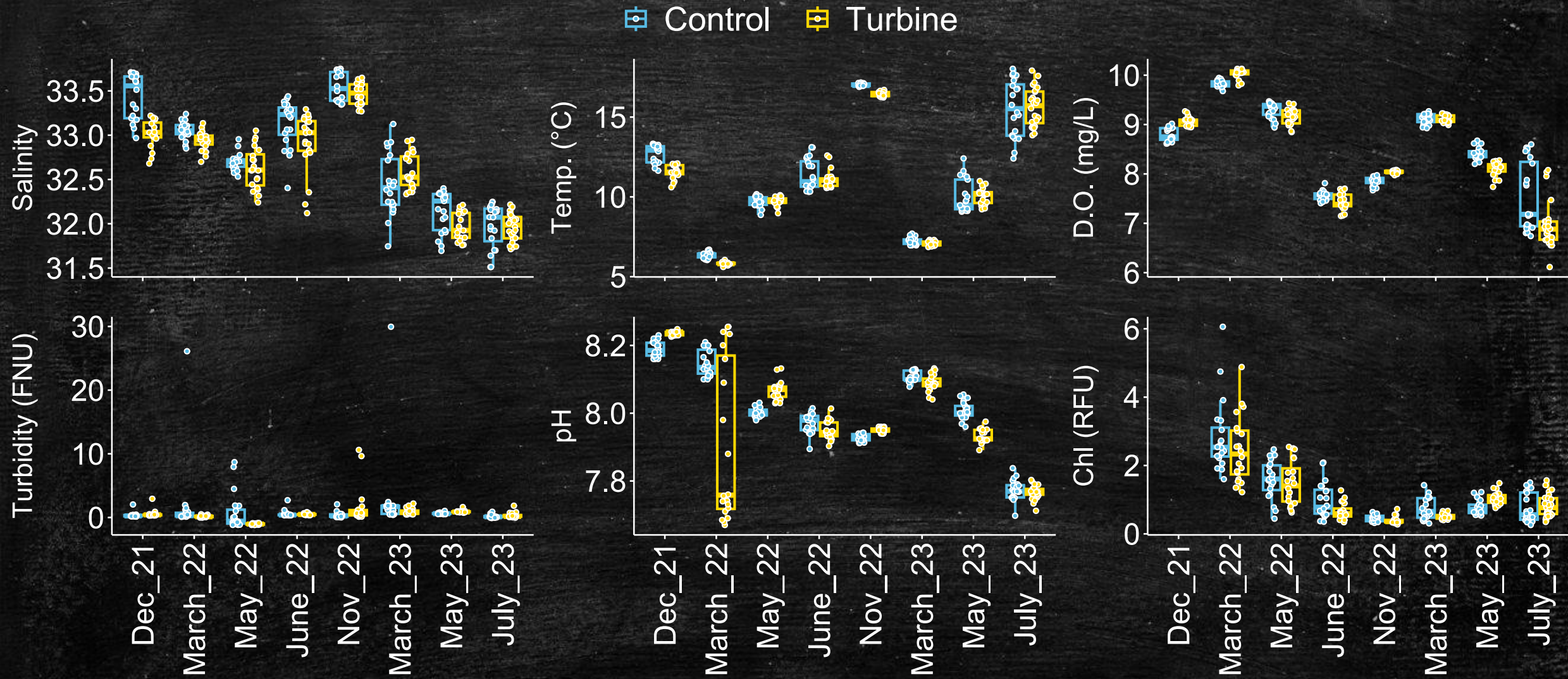


Offshore wind project with planned capacity of 1,100 MW to be operational in 2024

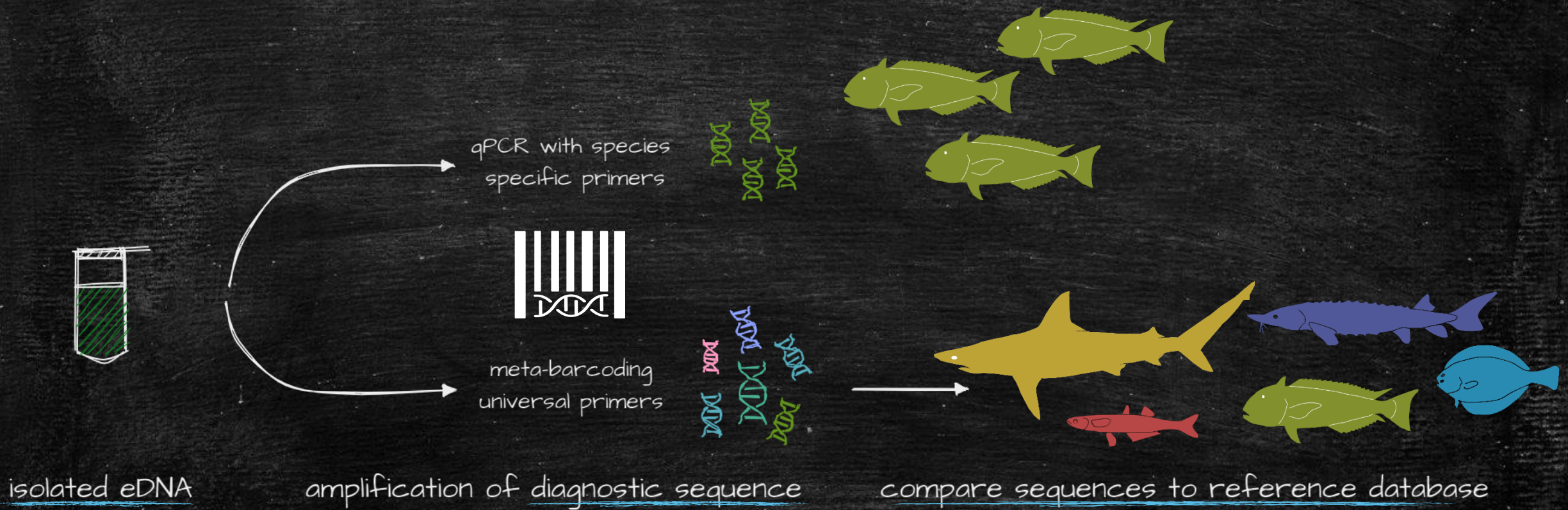
Turbine and **Control** sites consist of 28 stations

Quarterly **sampling bouts** for continuous monitoring of fish communities

Biomonitoring using environmental DNA to occur before, during, and after construction



seasonal variability of environmental conditions is larger than between-site variation



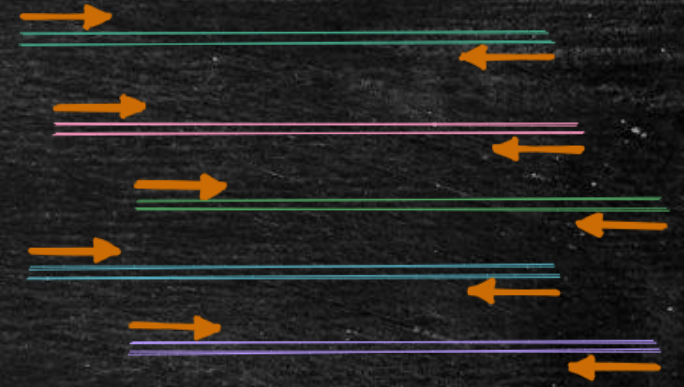
eDNA is useful for species-specific assessments and to characterize entire communities

even "universal" primers have bias that needs to be accounted for

e.g. Modified 16S primers (Riaz) amplified 2-3x more elasmobranchs compared to the standard set



isolated eDNA



amplify diagnostic locus

next-generation sequencing
(100 - 200k reads per sample)

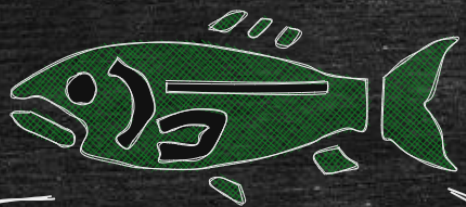
Identify taxonomic groups present in sample



identify unique amplicon variants (ASVs)
match ASVs to reference (presence)
determine no. reads per ASV (rel. abund)

Metabarcoding can be used to characterize benthic fish communities

DNA is shed as cellular or extracellular material into the surrounding water



mucous  bodily fluids  tissue 

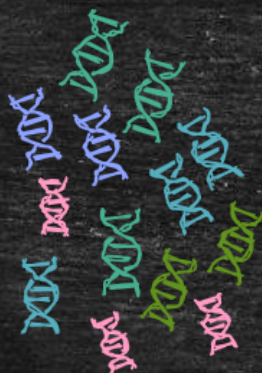
Temperature

UVB radiation

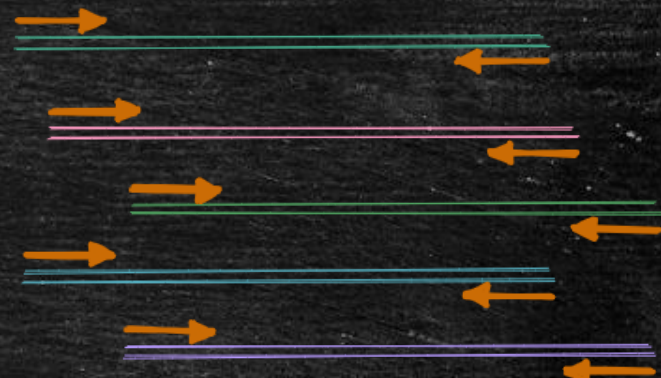
Acidity

Currents

Enzymes



isolated eDNA



amplify diagnostic locus

next-generation sequencing
(100 - 200k reads per sample)

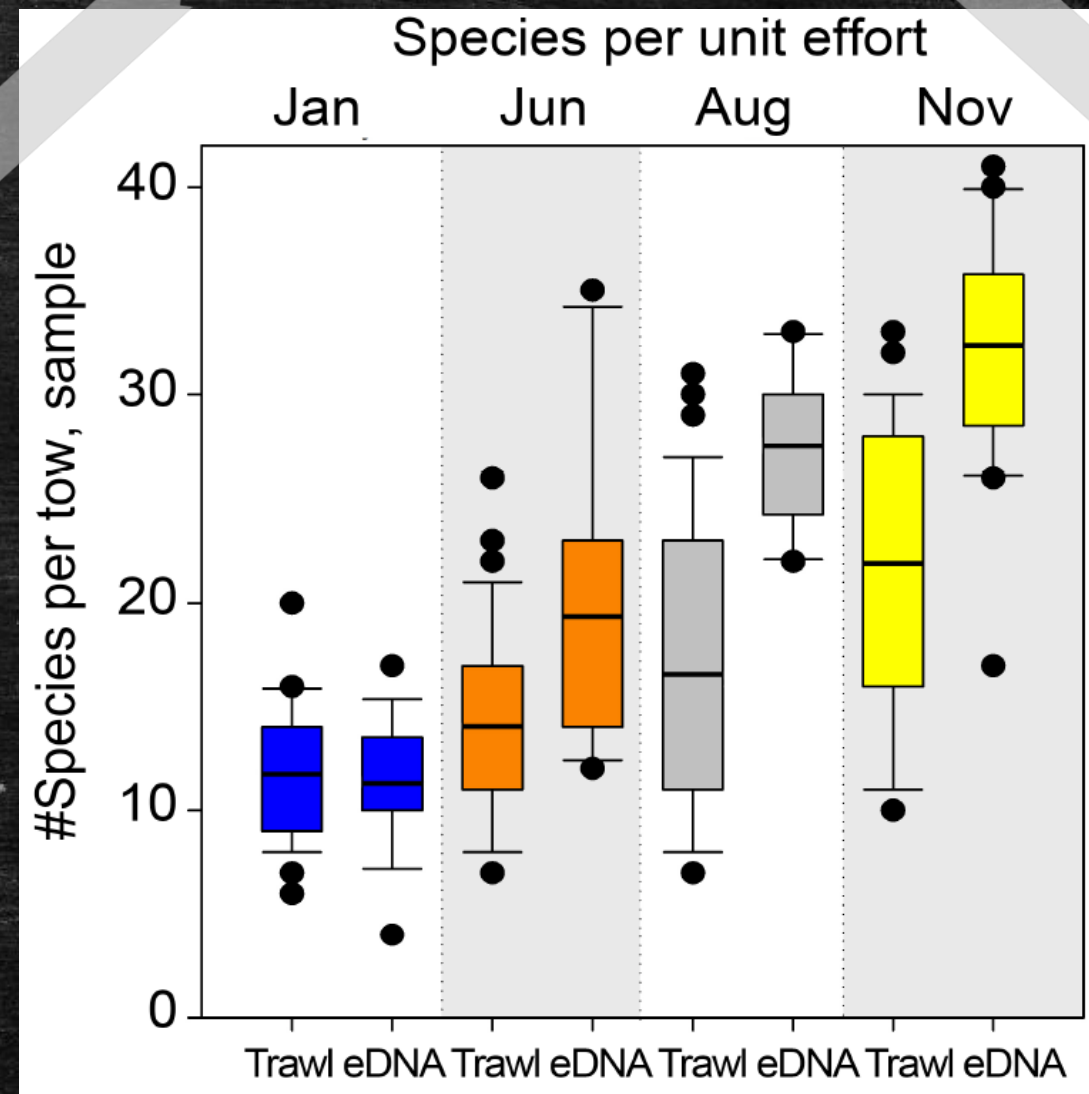
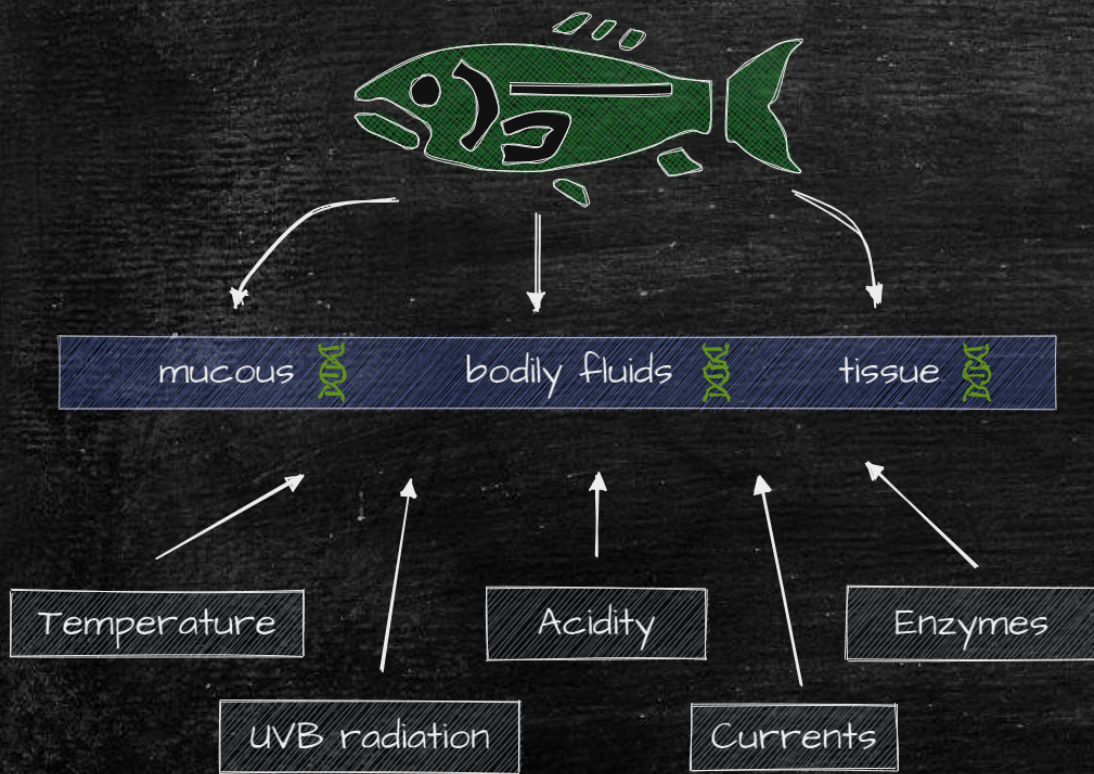
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Metabarcoding is semi-quantitative: characterize presence & relative abundance

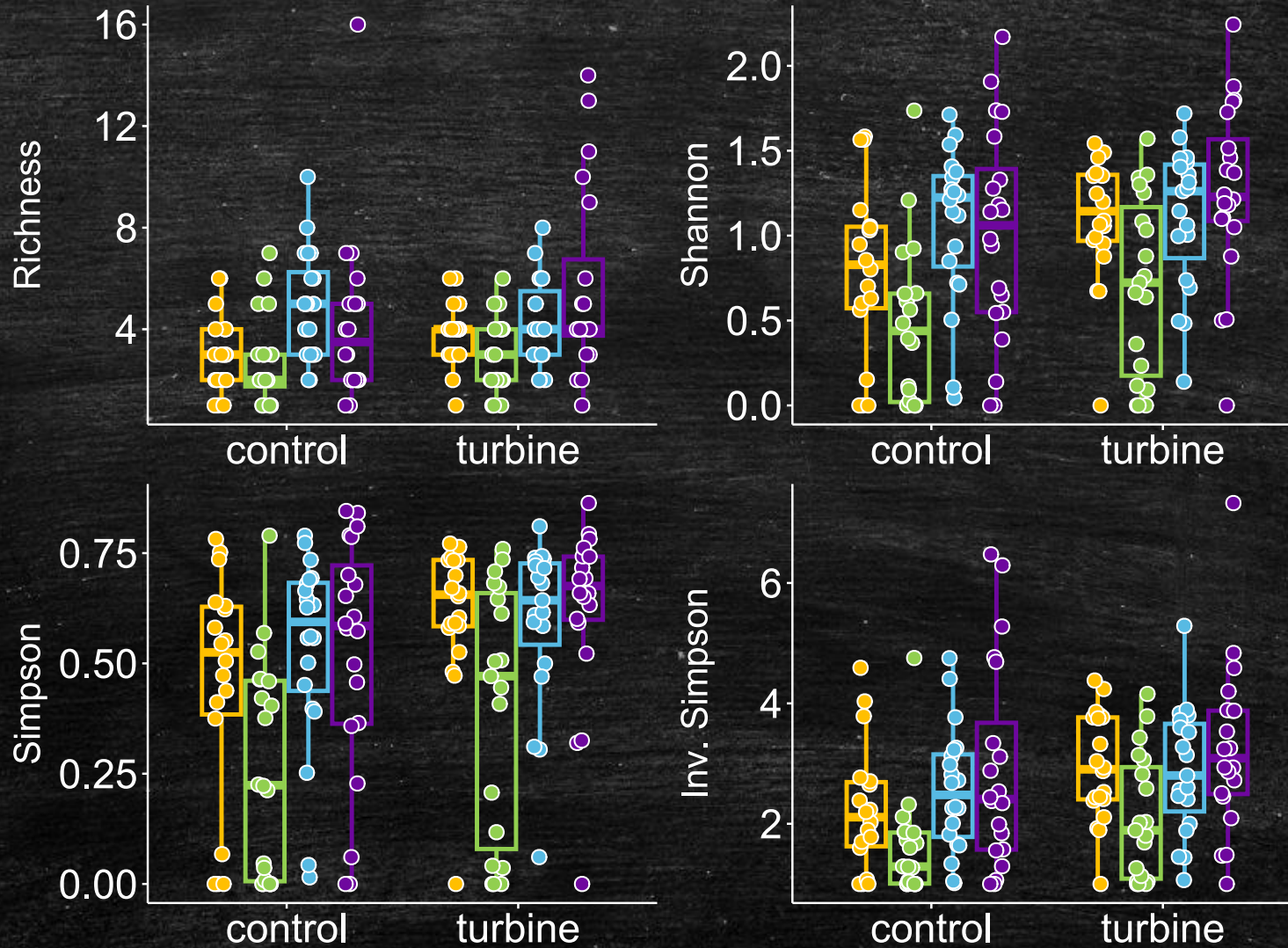
DNA is shed as cellular or extracellular material into the surrounding water



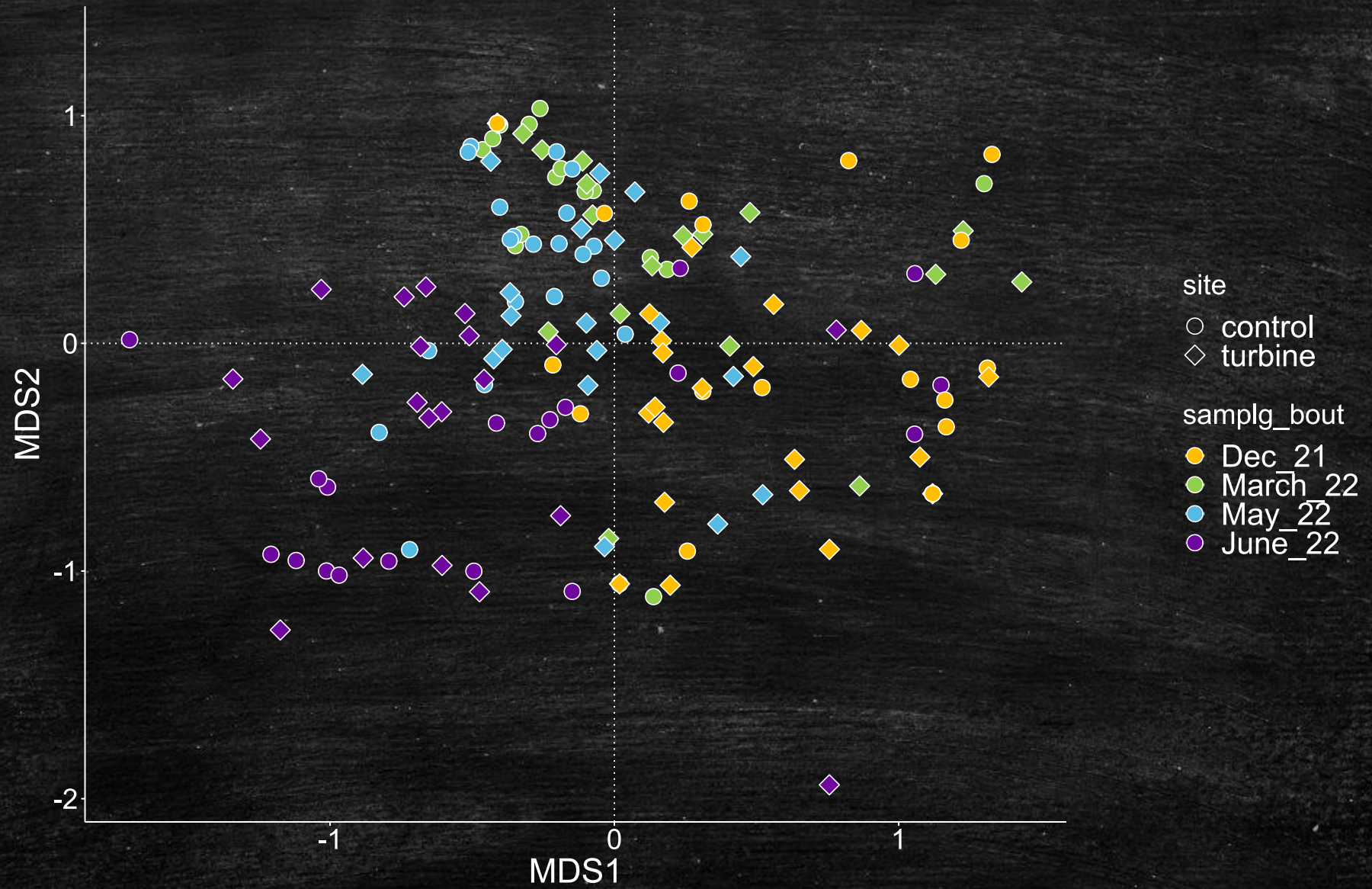
Stoeckle et al. 2020

Metabarcoding is semi-quantitative: characterize presence & relative abundance

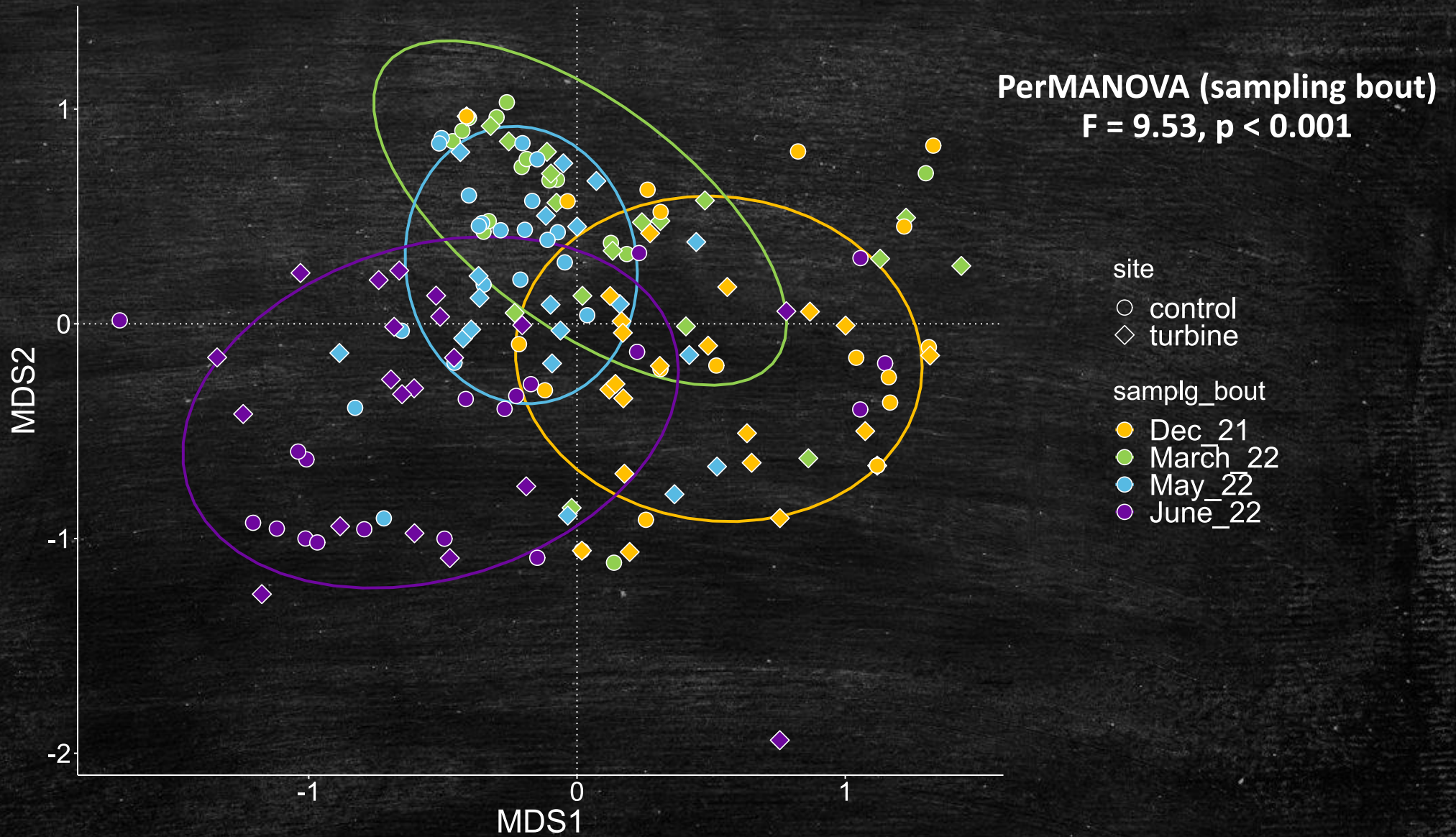
samplg_bout ⊠ Dec_21 ⊠ March_22 ⊠ May_22 ⊠ June_22



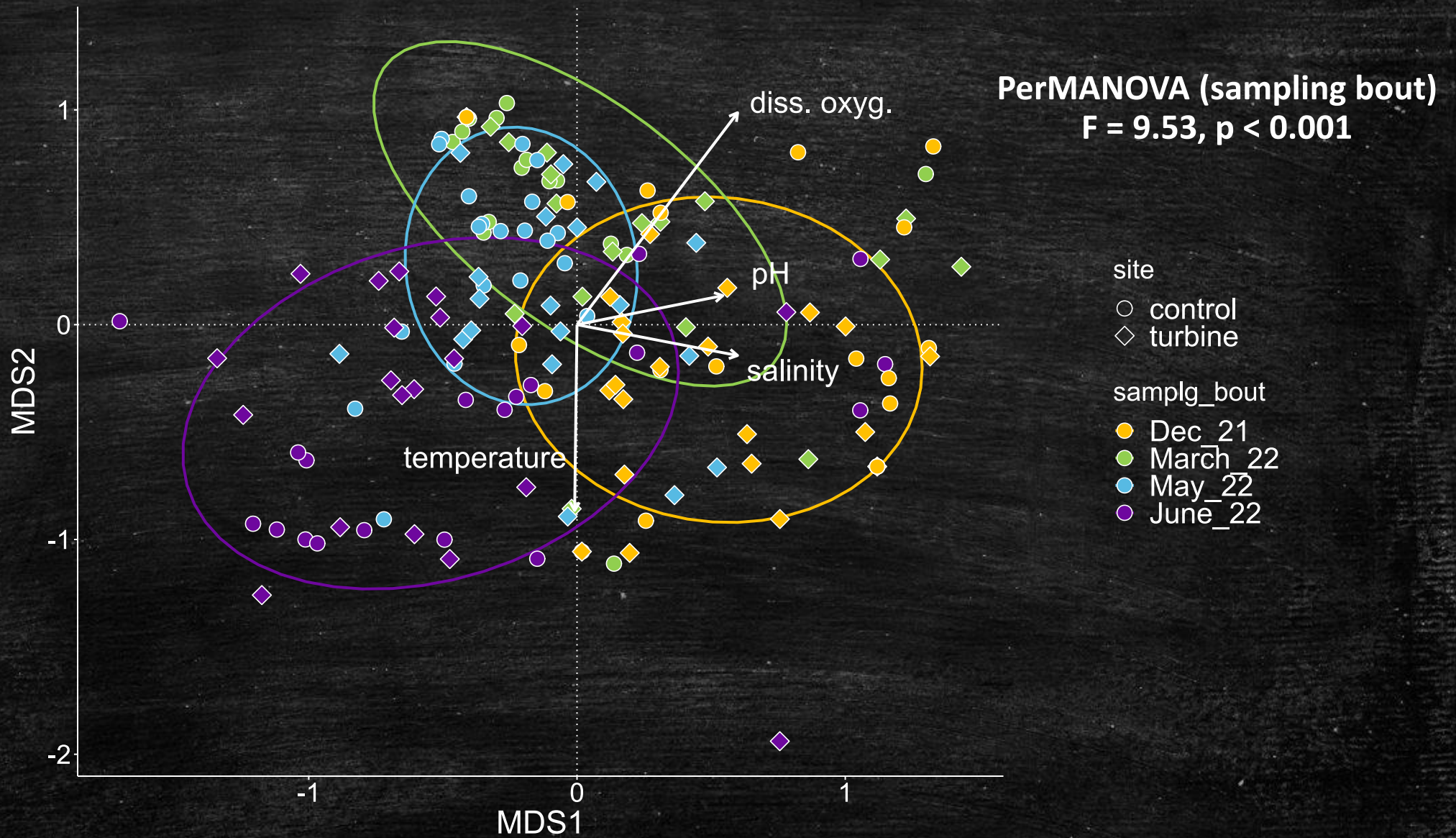
Differences in species abundance and diversity are driven by seasonal differences



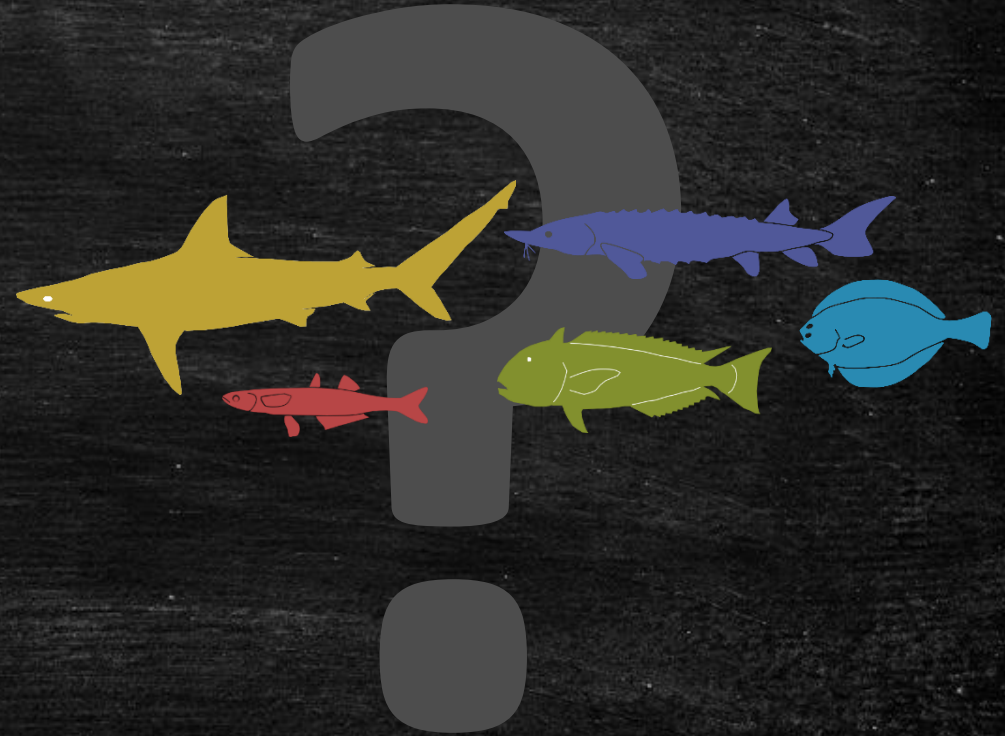
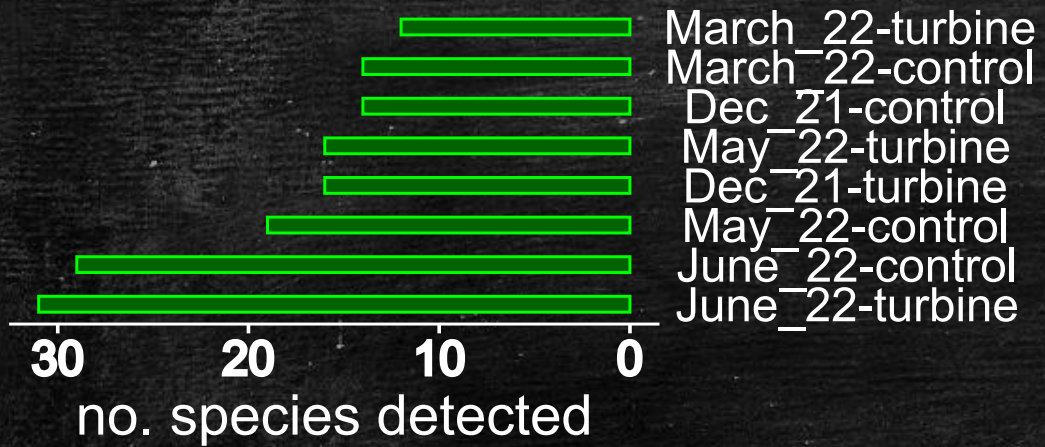
Structural diversity is driven by environmental variables and sampling season



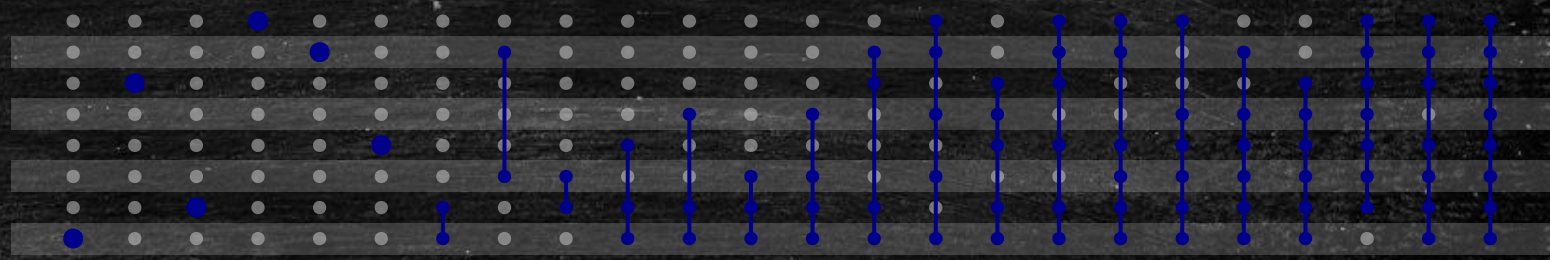
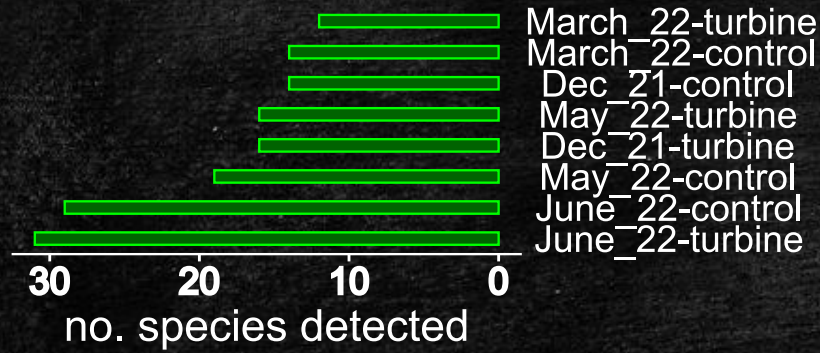
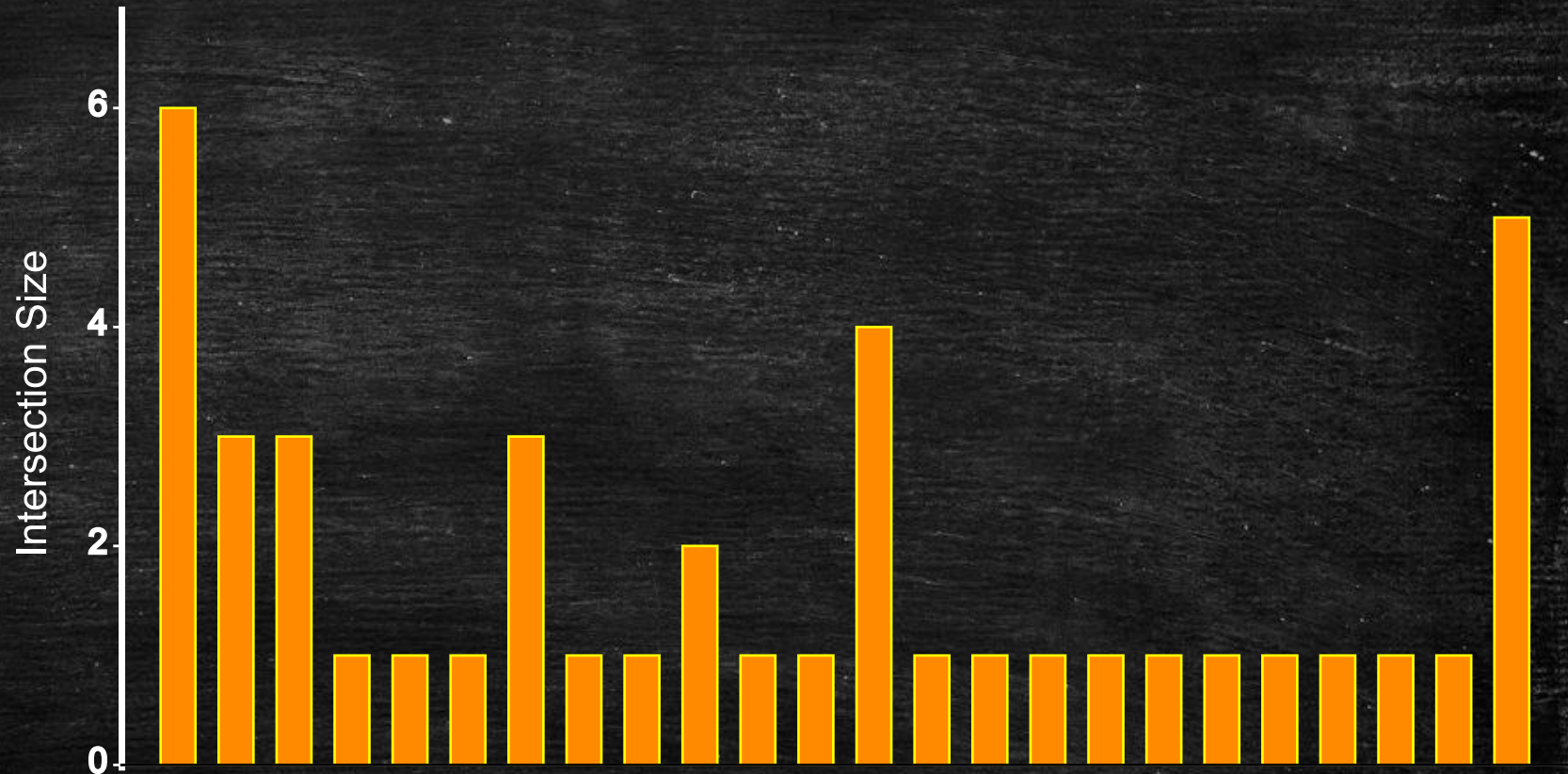
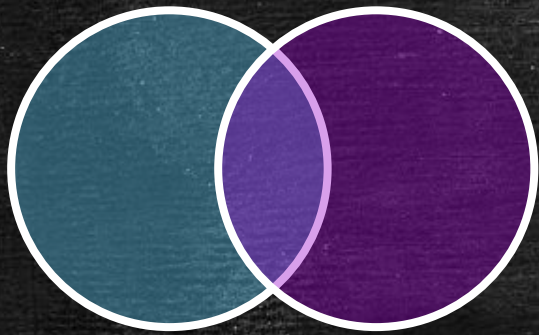
Structural diversity is driven by environmental variables and sampling season



Structural diversity is driven by environmental variables and sampling season



**A total of 43 species were detected across all stations and sampling bouts
(mostly the usual suspects)**

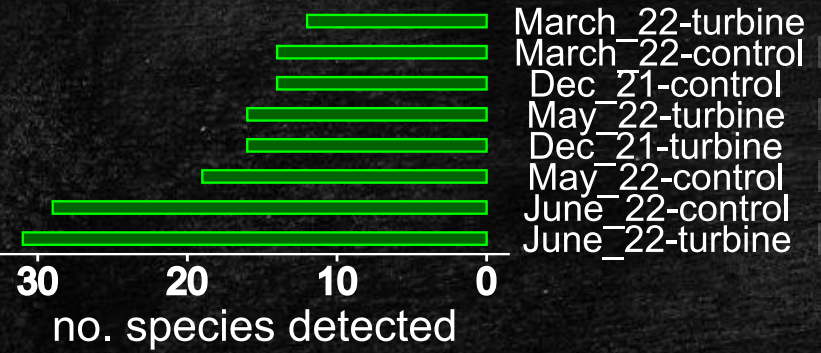
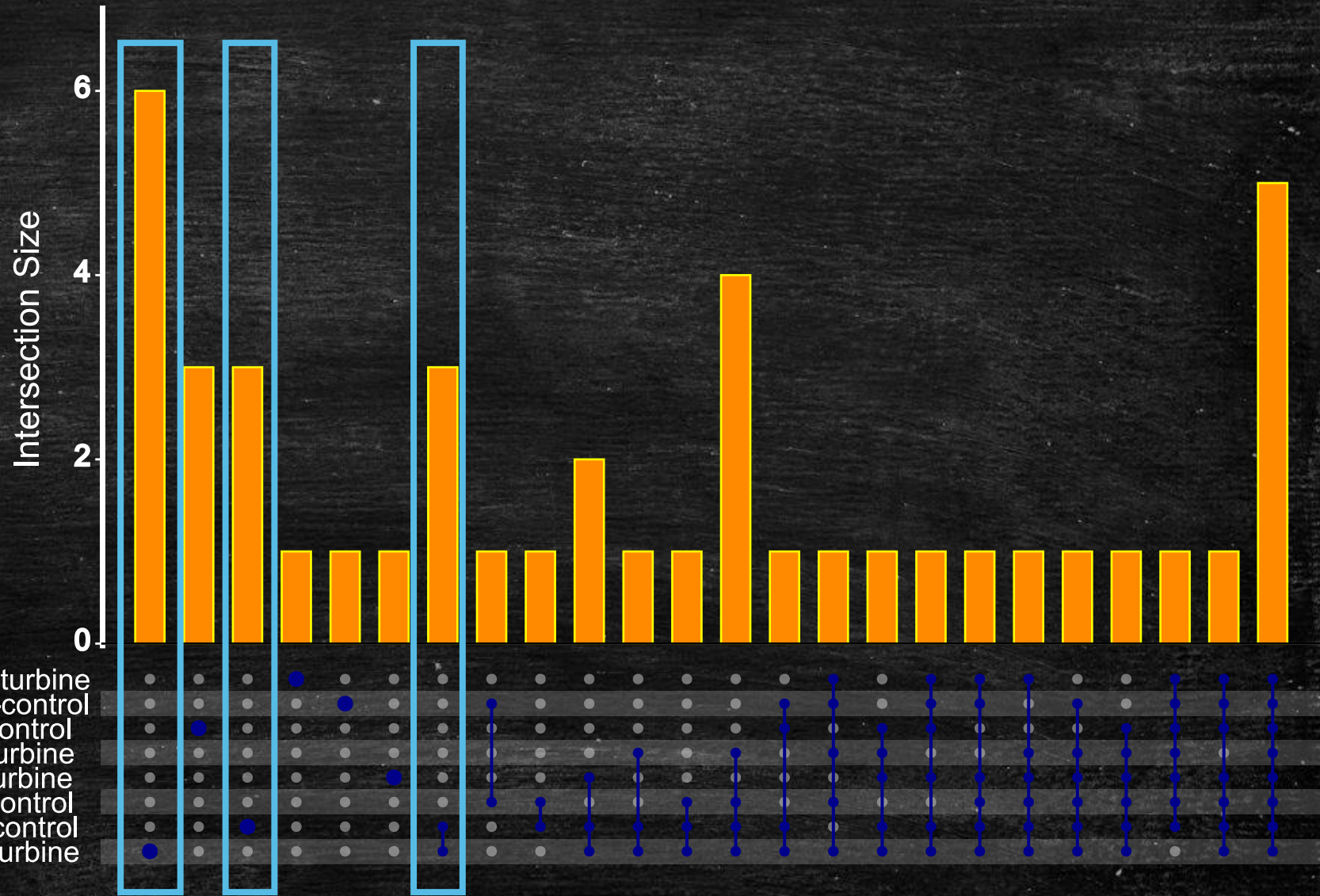


Five species were detected across all sampling bouts and seasons
June and December exhibit most unique species

White catfish
 Tautog
 Red drum
 Rough tail stingray
 Clearnose skate
 Thresher shark

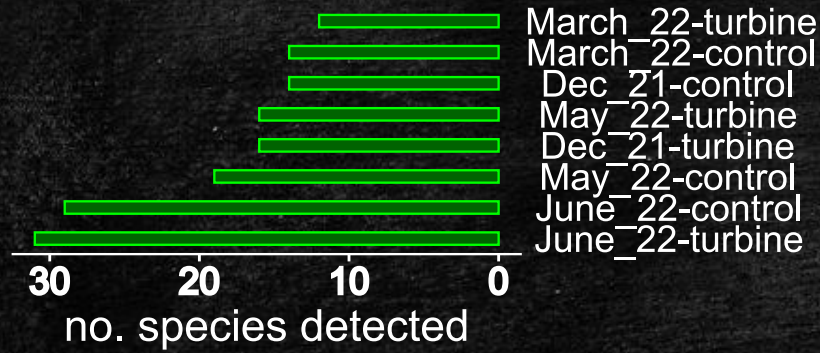
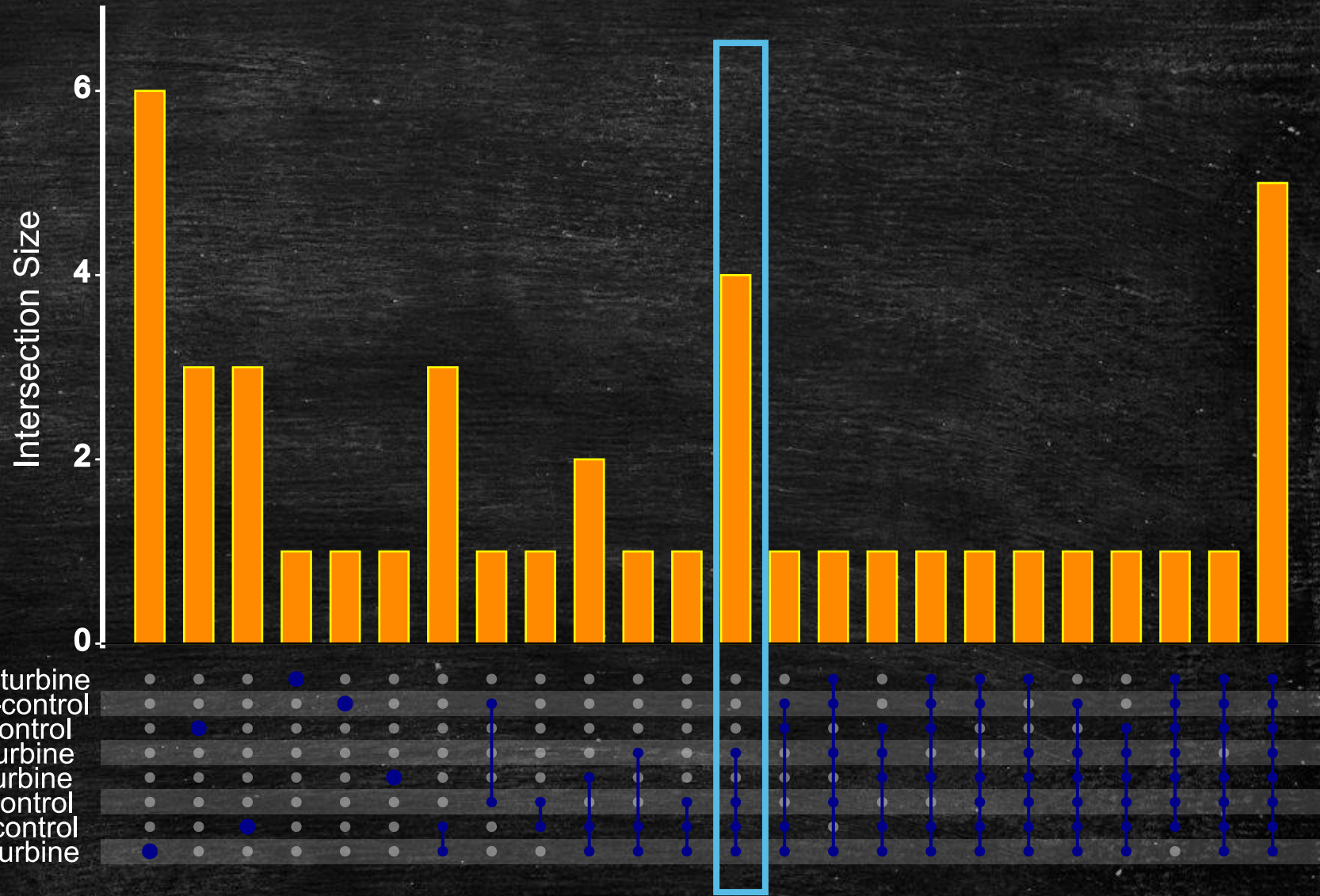
Cobia
 American anglerfish
 (Atl. Herring)

Thread herring
 Cunner
 Fourspot flounder



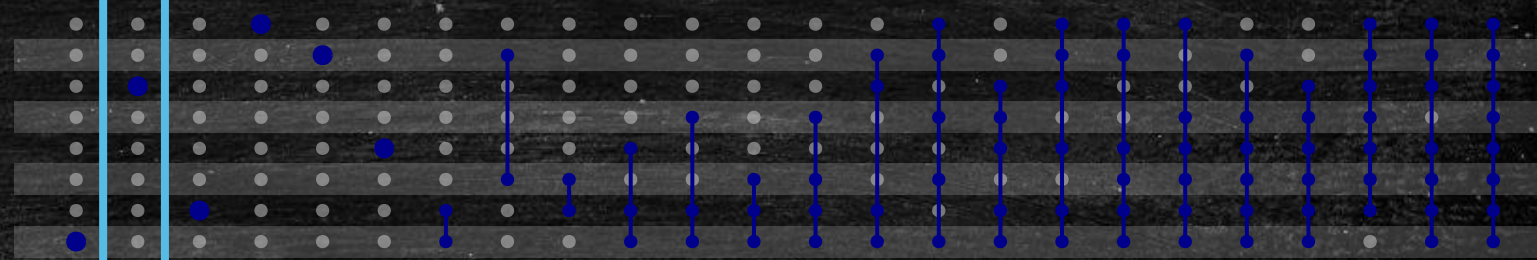
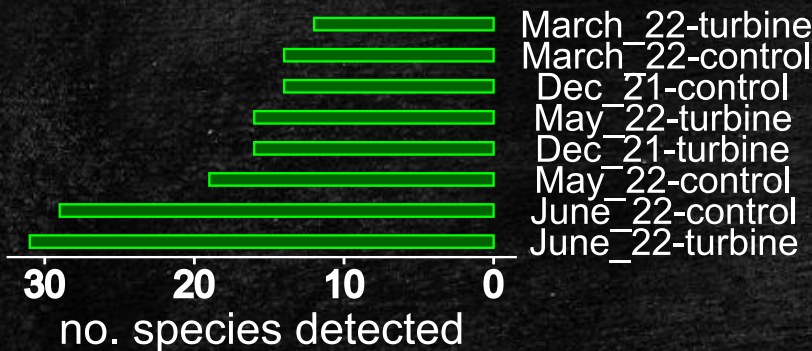
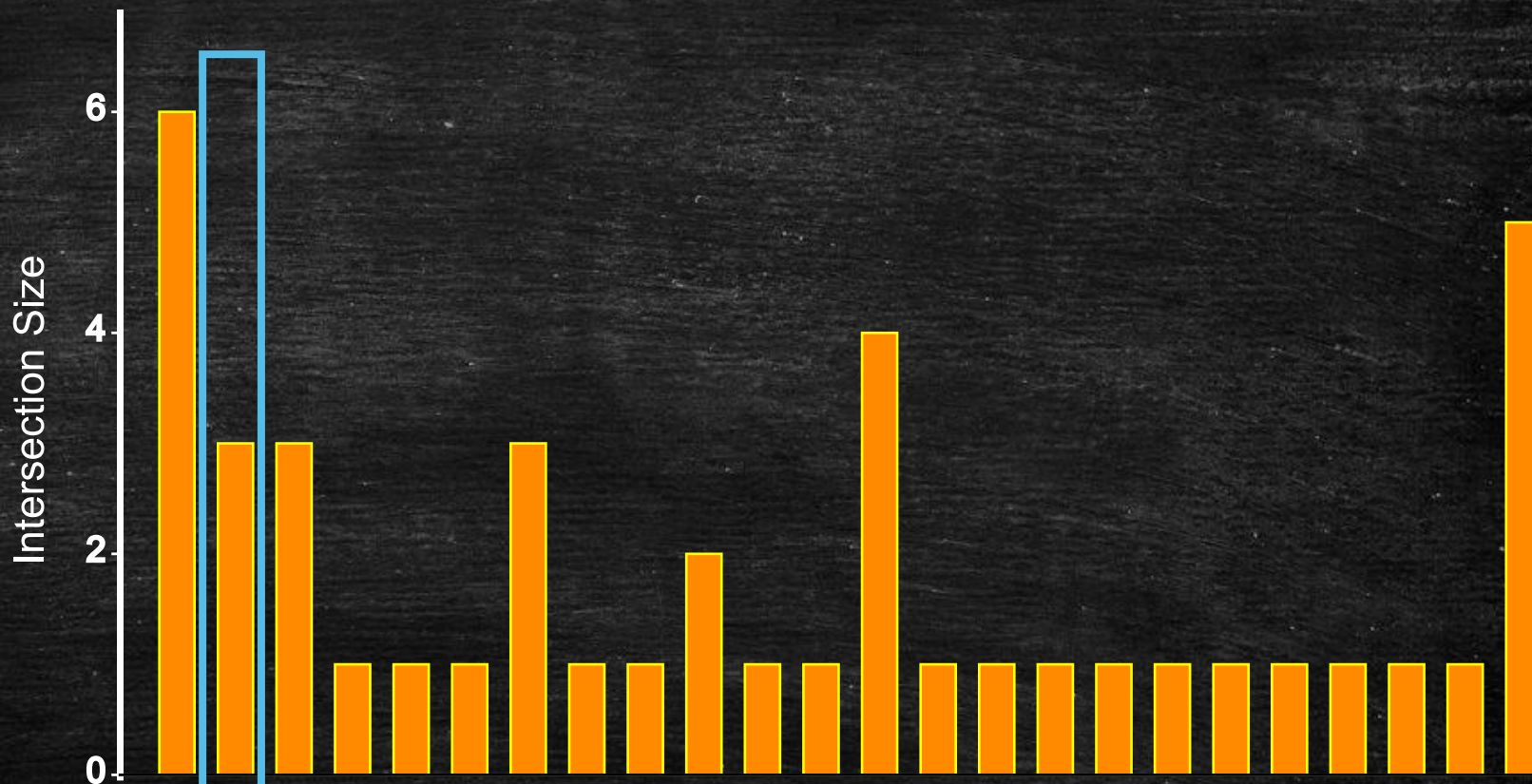
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White perch
 Silver hake
 Atlantic mackerel
 Atl/North Sand Lance



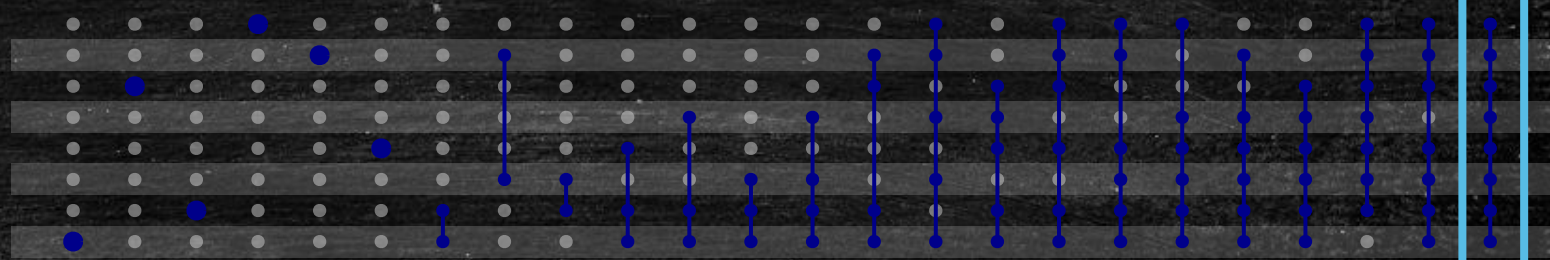
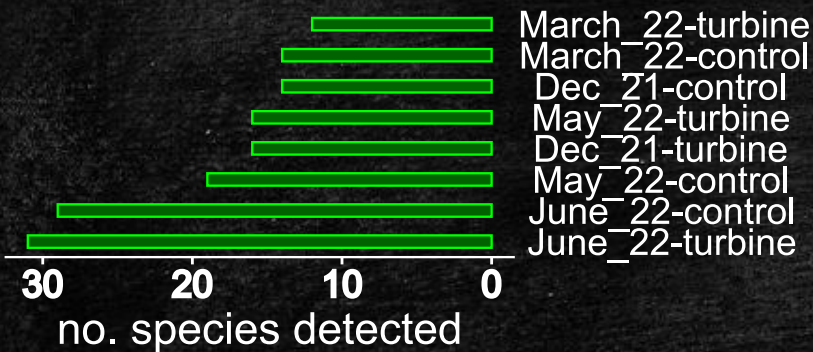
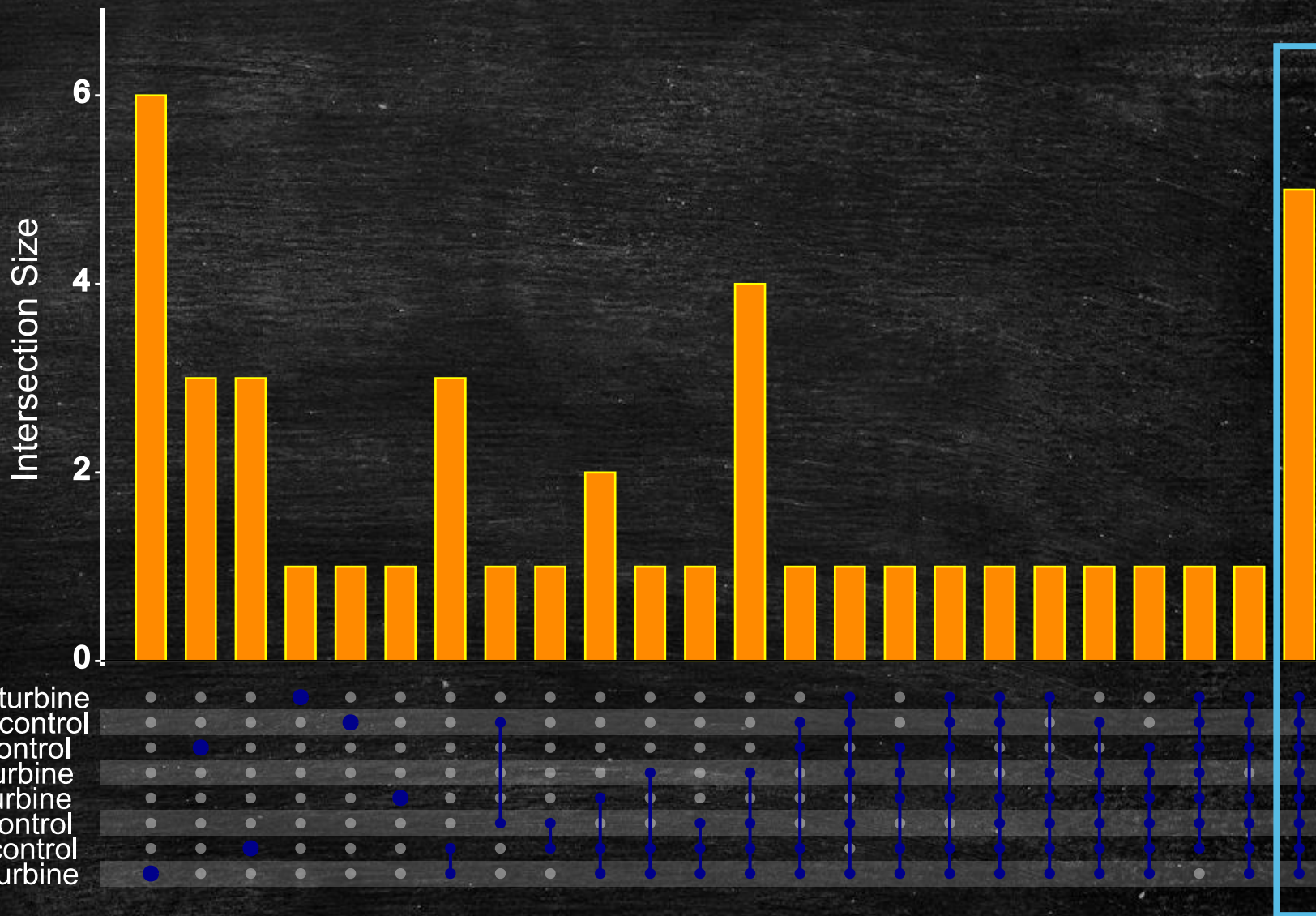
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Atlantic cod
 Inshore lizardfish
 Tuna sp



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Spiny dogfish
 White/spotted hake
 Black seabass
 American butterfish
 Menhaden/Herring



Five species were detected across all sampling bouts and seasons
June and December exhibit most unique species



Detections include species consistently caught at 35 – 65% of stations and those detected at only one or two during certain sample bouts

DNA captured from environmental samples without pre-isolating specific targets

- Low cost, simple, flexible sampling methods (scaleable)
- Standardize methods across wide range
- minimal damage to habitat, target/bycatch species
- few permitting issues
- Need more calibration studies to understand “gear bias” (DNA degradation, relative abundance estimates)



Specimen are extracted from their ecosystems with potential impact on habitat from fishing

- Biological data of captured specimen
- Existing long-term data sets
- Gear-bias well-understood
- Established methods for analysis & application to stock assessments

Potential and Pitfalls of using eDNA for biomonitoring compared to traditional sampling



Questions?



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Library Prep & Sequencing: Sabeena Nazar (BASLab)