

algae not detectable with microscopy

Study 2: Expand and validate molecular sequences present in the **database**, with focus on the potentially toxic microalgal species

Study 3: Validate data obtained from satellite sensors, e.g., functional types, through the study area

3. Study area



**TEM & SEM** 

for species-level

identification

different DNA extraction, PCR (markers: database (e.g., silva, V4, V9, rbcL), sequencing



1.sampling: phytoplankton net

validate Phytoplankton

Functional Types with

observational data

2.microalgal strains isolation

identification

& counting:

inverted LM

3.culture growth & maintenance 4.molecular analysis: (LSU, V4, V9 & rbcL)

5.sequences compared or inserted in databases

silva

pr2)

Study 3:



test the quality of the satellite information (statistical approaches)





## WORK IN PROGRESS

- Until now 155 strains of microalgae have been isolated and cultured
- Sanger sequencing (LSU and 18S) was performed on 65 strains, identifying species such as:

Biecheleriopsis adriatica, Scripsiella trochoidea, Thalassiosira profunda, Chaetoceros socialis, Emiliania huxleyi, Skeletonema marinoi and Dicrateria inornata

Frame the QR code and watching the video of unidentified algal cultures



**next steps:** DNA extraction and TEM for identification

## • Next step will be to obtain the rbcL sequence for those species that lack the

sequences in the database

Università degli Studi di Palermo

NATIONAL BIODIVERSITY FUTURE CENTER