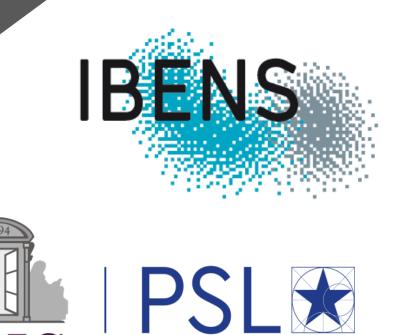
LncPlankton: A comprehensive collection of plankton long noncoding RNAs





<u>Ahmed Debit</u>^{1*}, Pierre Vincens¹, Chris Bowler¹, Helena Cruz de Carvalho^{1,2*}

¹ Institut de Biologie de l'ENS (IBENS), École normale supérieure, CNRS, INSERM, Université PSL, Paris, France ² Université Paris Est-Créteil (UPEC), Faculté des Sciences et Technologie, Créteil, France

* Correspondence: debit@bio.ens.psl.eu | cruz@bio.ens.psl.eu



Background

For a long time, long noncoding RNAs (IncRNAs) have been viewed as transcriptional noise. However, they're now being recognized as key regulatory molecules across the Eukarya domain, including plants, animals, and fungi. Despite this, we still don't have a clear understanding of how these molecules occur in the marine environment.

To address this knowledge gap, we created LncPlankton, the first comprehensive database of marine planktonic IncRNAs. By combining predictions from ten coding potential tools using majority voting, we identified over 2 million lncRNAs spread across 414 marine planktonic species from more than nine different phyla. We also developed a user-friendly, open-access interface to explore the database (https://www.lncplankton.bio.ens.psl.eu/).

Our goal is for LncPlankton to become a valuable resource for IncRNAs, supporting both small-scale and large-scale research in various marine planktonic species, and enabling comparative analysis that extends far beyond the marine environment.

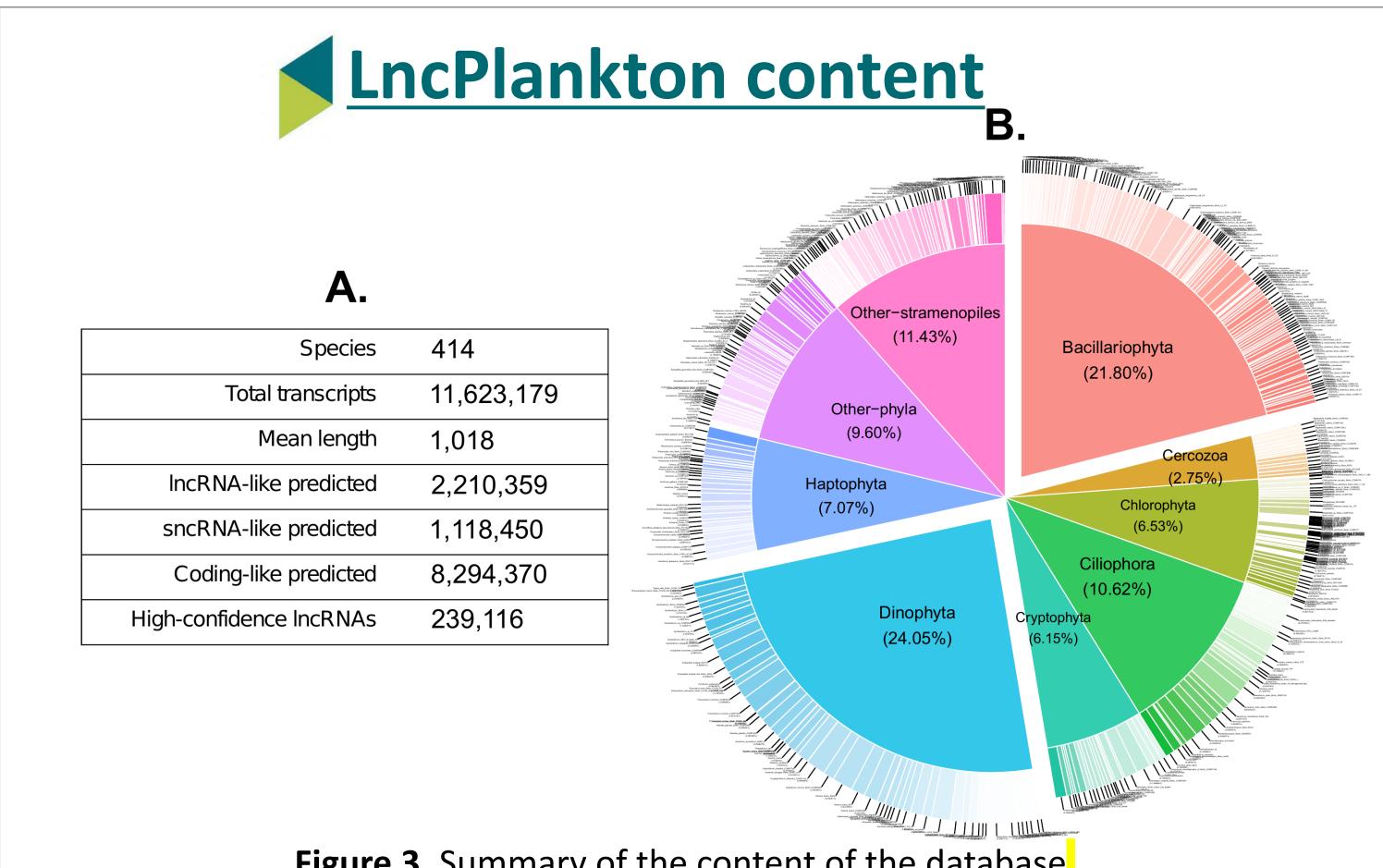


Figure 3. Summary of the content of the database

Materials & Methods



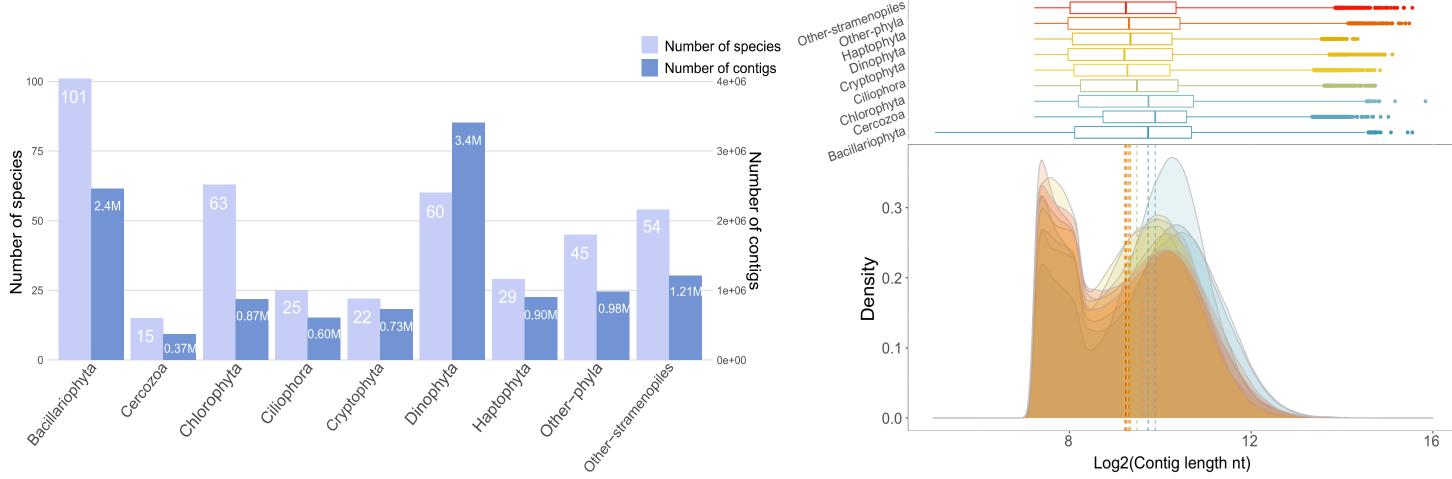


Figure 1. Data used in LncPlankton

- Transcriptomic data of 406 species MMETSP project (Keeling et al., 2014) + 6 diatom species not covered by the MMETSP project.
- Transcriptomes of *P. tricornutum* (Cruz de Carvalho et al., 2016) and Thalassiosira pseudonana (Goldman J.A. et al., 2019) used an inhouse assembly pipeline.
- Data analysis pipeline

Our method shows a high accuracy and low variability

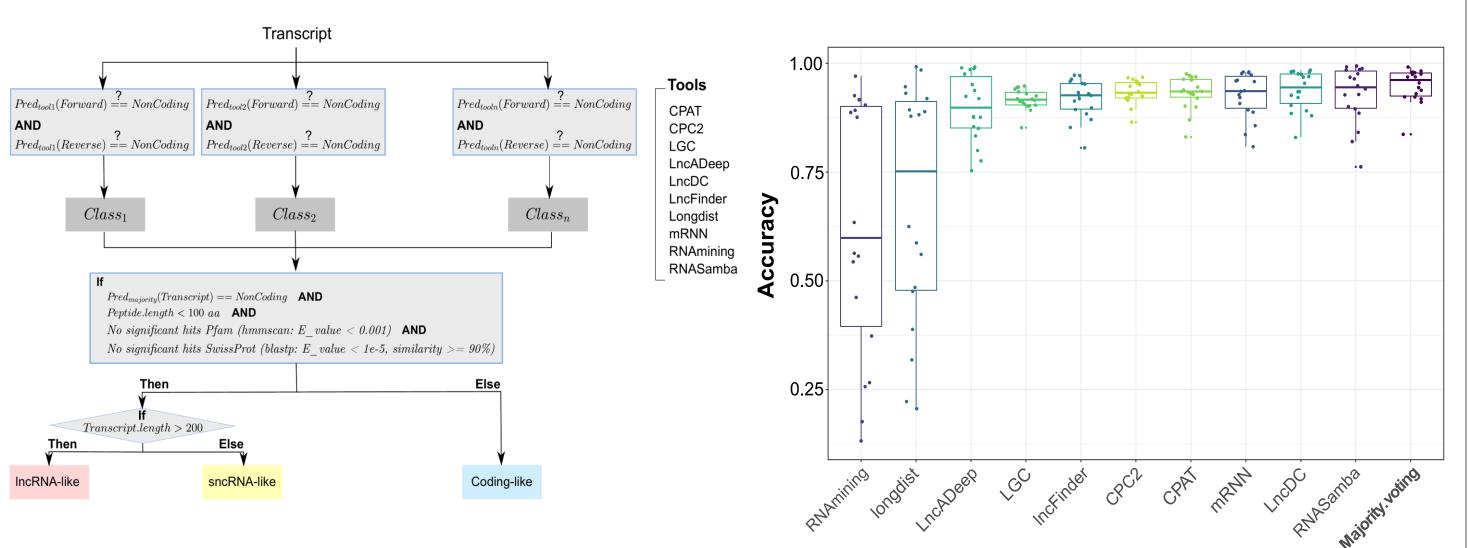


Figure 2. The pipeline used for the prediction of lncRNAs combines 10 coding potential tools in a majority voting setting

Benchmarking data: Different sets of coding and non-coding datasets related to 18 species of different chordate clades. The datasets were independent of the training sets used in the construction of the pretrained models related to each coding potential tool.

References

- Keeling et al., (2014); The Marine Microbial Eukaryote Transcriptome Sequencing Project (MMETSP): Illuminating the functional diversity of eukaryotic life in the oceans through transcriptome sequencing; PLoS Biology, 12.
- Cruz de Carvalho et al., (2016); Noncoding and coding transcriptome responses of a marine diatom to phosphate fluctuations; New Phytologist, 210, 497–510.
- Goldman J.A. et al., (2019); Fe limitation decreases transcriptional regulation over the diel cycle in the model diatom Thalassiosira pseudonana; PLOS One, 14.

LncPlankton Ul

The UI provides modules for browsing, searching, downloading IncRNA data per species and/or per phylum, interactive graphs, and an online BLAST service. A shiny application was also integrated, allowing the user to customize and visualize the classification.

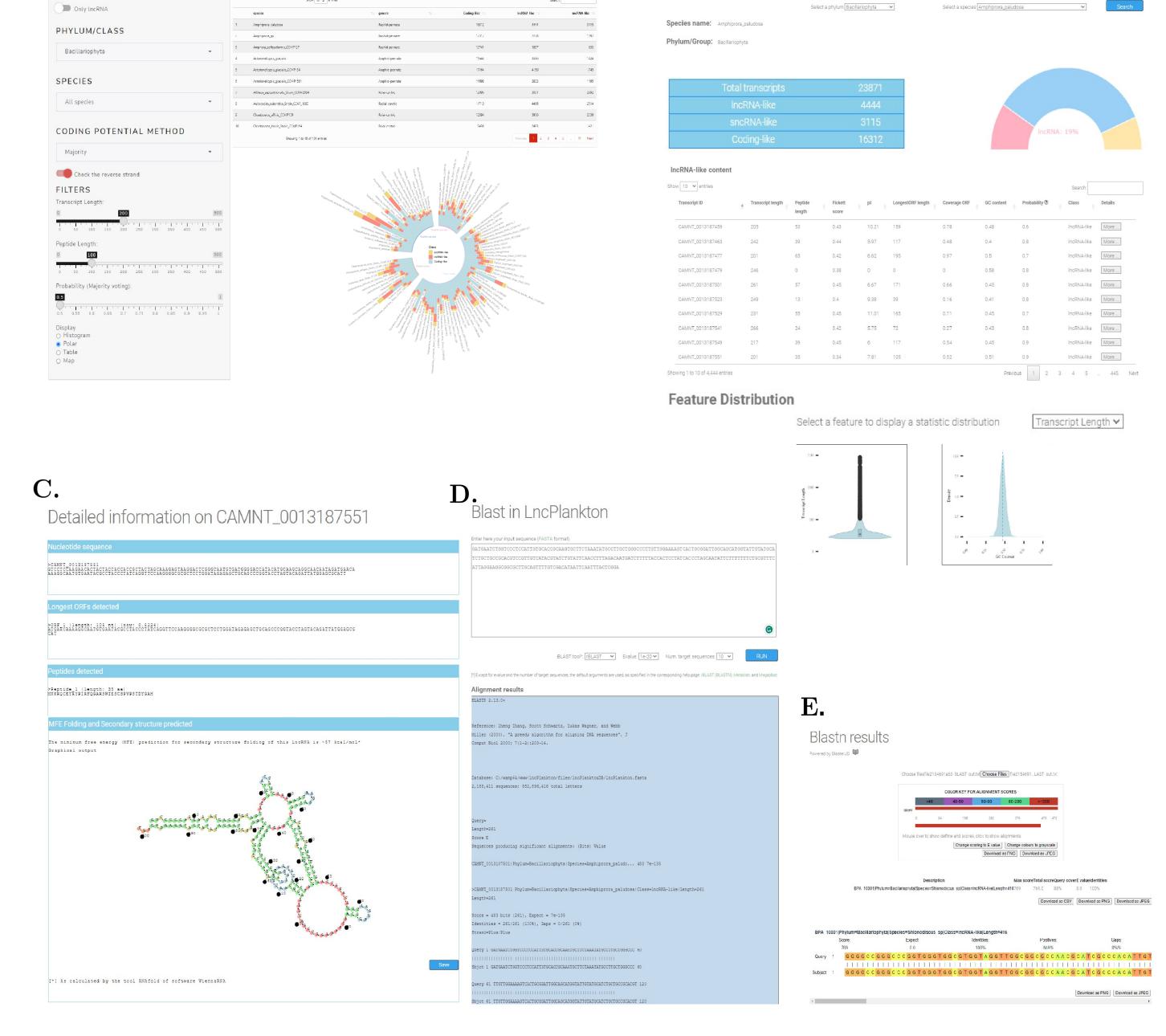


Figure 4. Screenshots of LncPlankton web interface and modules

Discussion & Conclusion

- We implemented a meta-learner using a majority voting-based ensemble learning technique for the prediction of IncRNAs in marine planktonic species (https://gitlab.com/a.debit/votinglnc)
- The meta-learner promoted the heterogeneity (multiple and diverse ML algorithms), and the diversity (integrated of different features describing the transcripts).
- Our method is robust and shows a low variability of the prediction across the different testing datasets.
- The majority voting tool offers us a **reliable** choice for IncRNAs identification, beyond what a single tool can offer at the moment.
- LncPlankton is the largest data repository on lncRNAs in marine species.
- We anticipate LncPlankton will contribute significantly to future efforts aimed at deciphering the biology of marine lncRNAs.