

Figure 1: Map of the studied sites

| Code | Taxa |
|------|---|
| AST | Asterionella+Asterionellopsis+Asteroplanus |
| CHA | Chaetoceros |
| CRY | Cryptophytes |
| DIT | Ditylum |
| EUG | Euglenophytes |
| GUI | Guinardia |
| GYM | Gymnodinium+Gyrodinium |
| LEP | Leptocylindrus |
| NIT | Nitzschia+Hantzschia |
| PLE | Pleurosigma+Gyrosigma |
| PRO | Prorocentrum |
| PRP | Protoperidinium+Archaeoperidinium+Peridinium |
| PSE | Pseudo-nitzschia |
| RHI | Rhizosolenia+Neocalyptrella |
| SCR | Scrippsiella+Enciculifera+Pentapharsodinium+Bysmatrum |
| SKE | Skeletonema |
| THL | Thalassionema+Lioloma |
| THP | Thalassiosira+Porosira |

Table 1: Name and composition of the phytoplanktonic groups used in the paper, based on¹

| Name of site | Location | Region | Number of points? | Mean rainfall | Mean temperature | Species we chose to keep |
|--------------|----------|--------|-------------------|---------------|------------------|--------------------------|
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Table 2: Attempt of summary for our locations (for now, only the main titles, we'll see if we have to complete)

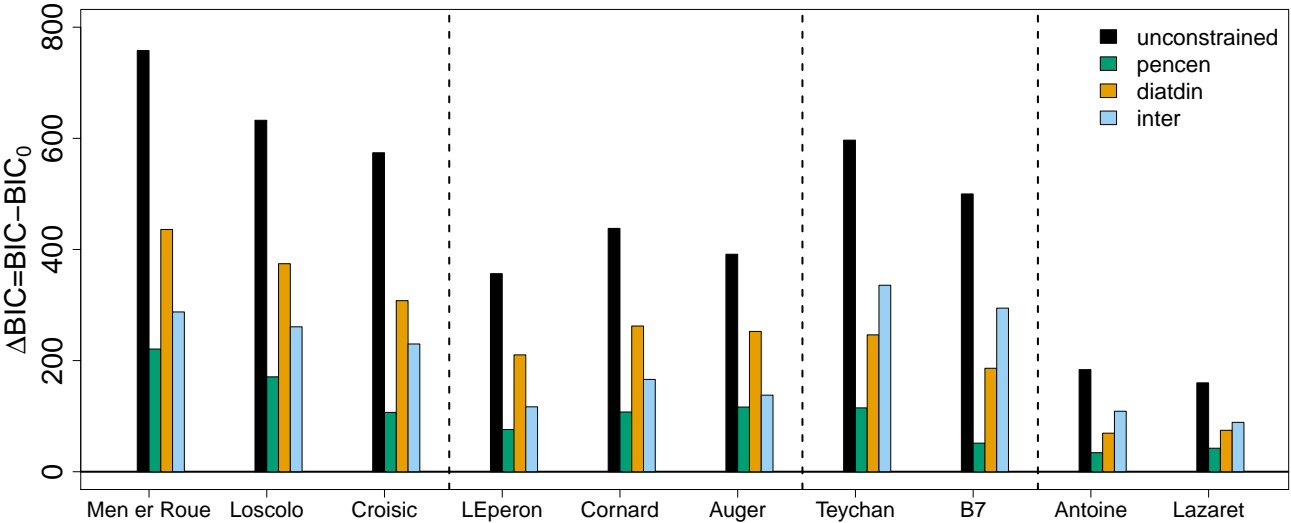


Figure 2: Comparison of BIC with different interaction matrices, compared to the null model (diagonal interaction matrix, allowing only intragroup interactions), for four different sites separated by dashed lines (Brittany, Oléron, Arcachon and Mediterranean Sea) and 10 different subsites. Different interaction matrices may allow all interactions between taxa (unconstrained), only interactions within pennate diatoms, centric diatoms, dinoflagellates, or other phytoplanktonic taxa (pecten), only interactions within diatoms, dinoflagellates or other taxa (diatdin), or only interactions between taxa belonging to these different groups. As model structures (length of the times series taken into account) are different between sites and subsites, groups of bars should not be compared.

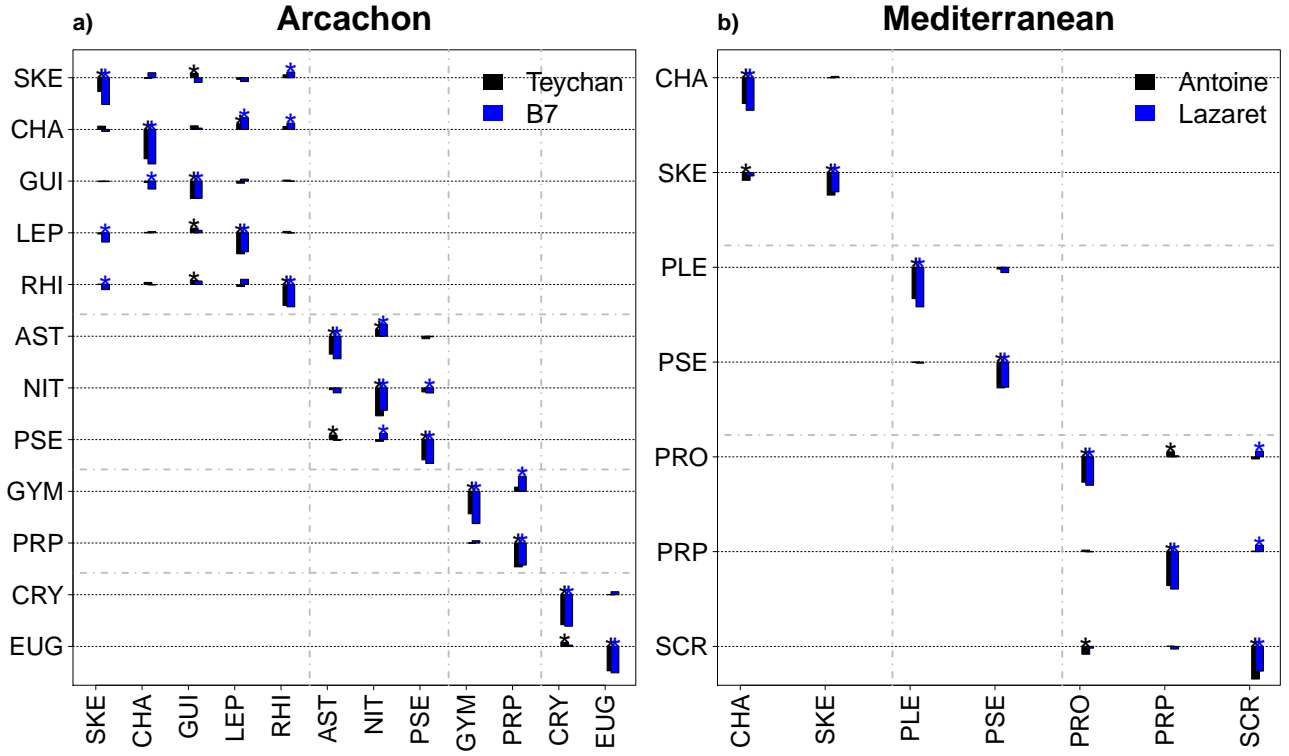


Figure 3: Interaction matrices estimated in Arcachon (a) and in the Mediterranean Sea (b). Only interactions between clades (pennate and centric diatoms, dinoflagellates, other planktonic taxa) are allowed. The figure should be read as taxon i having effect e_{ji} on taxon j . The scale for the coefficient values is given at the bottom left of panel a). 95% significance of coefficients was determined by bootstrapping and is marked by asterisks (*). The identity matrix was subtracted to the interaction matrix ($\mathbf{B}-\mathbf{I}$) in order to make effects on growth rates comparable. Composition of planktonic groups is given in Table 1.

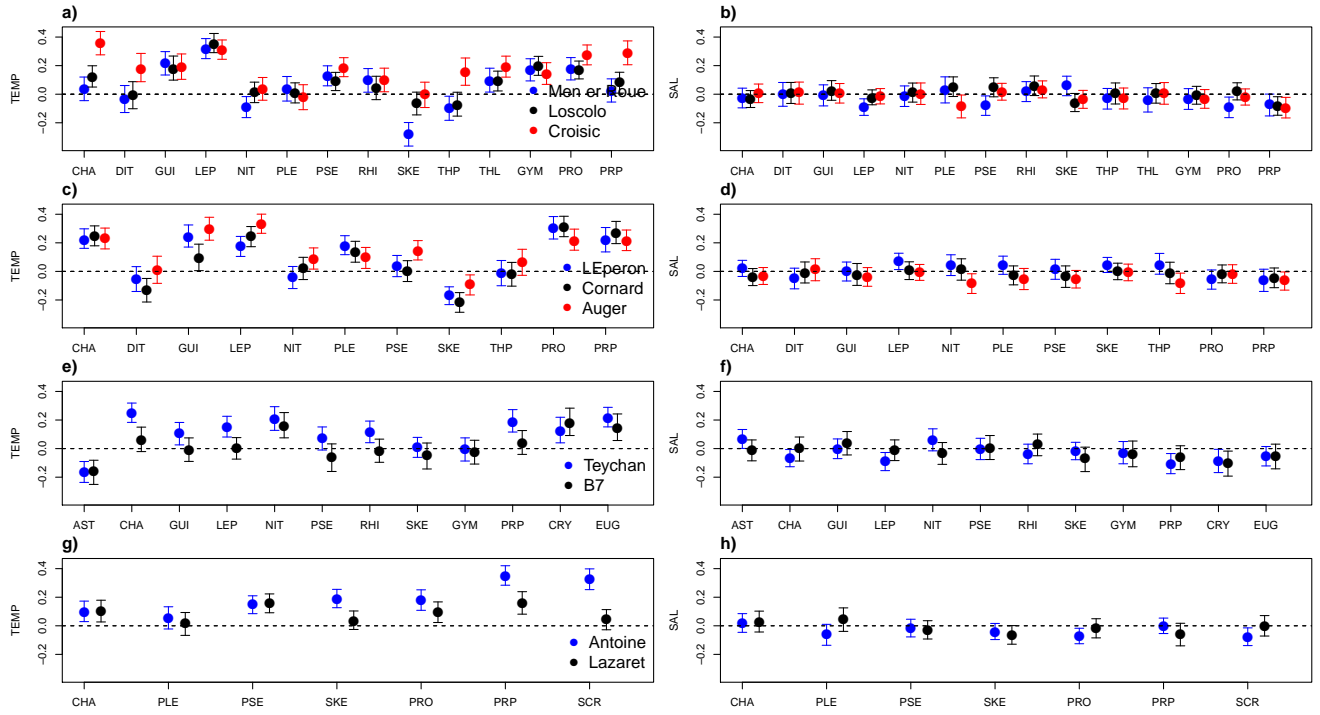


Figure 4: Effect of abiotic variables (temperature, TEMP or salinity, SAL) on phytoplankton group in Brittany (a, b), Oléron (c, d), Arcachon (e, f) and in the Mediterranean Sea (g, h). Each color corresponds to a different site. Error bar corresponds to the 95% confidence interval around the estimated coefficient. All variables were normalized before estimation.

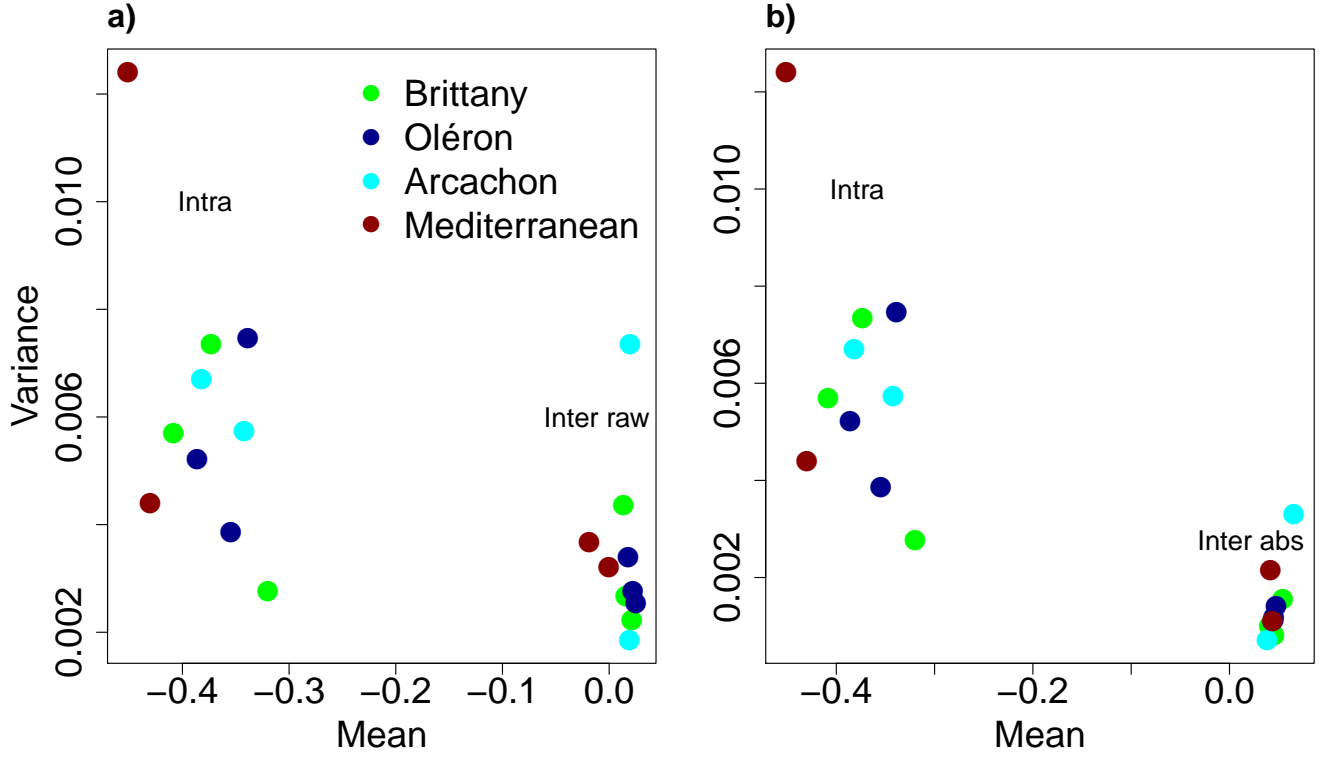


Figure 5: Variance of the coefficient in the interaction matrix ($\mathbf{B-I}$), as a function of their mean, for 10 sites in 4 regions, with a model allowing interactions only within clads (see above). The mean-variance relation was either computed with raw values of intergroup interactions (a) or absolute values of the intergroup coefficients (b). Intragroup coefficients were not modified.

References for the meta-analysis

| Code | Ref | Dimension | Type of organisms | System |
|----------------------|--|-----------|-------------------------------|----------------------------|
| Griffiths2015_Phyto1 | [2] | 7 | Phytoplankton | Coastal site |
| Griffiths2015_Phyto2 | [2] | 7 | Phytoplankton | Offshore site |
| Huber2006_Phyto | [3] | 10 | Phytoplankton | Lake |
| Huber2006_Ciliate | [3] | 10 | Ciliates | Lake |
| Ives1999_Zoo1 | [4], conditional least square estimate | 9 | Zooplankton | Lake |
| Ives1999_Zoo2 | [4], total least square estimate | 9 | Zooplankton | Lake |
| Ives2003_Plank1 | [5] | 4 | Plankton | Lake |
| Ives2003_Plank2 | [5] | 4 | Plankton | Lake with high planktivory |
| Ives2003_Plank3 | [5] | 4 | Plankton | Lake with low planktivory |
| Klug2000_Phyto | [6] | 2 | Phytoplankton | Lake |
| Klug2000_Zoo | [6] | 3 | Zooplankton | Lake |
| Klug2001_TaxoAlgae | [7] | 5 | Taxonomic groups of plankton | Lake |
| Klug2001_MorphoAlgae | [7] | 4 | Functional groups of plankton | Lake |
| Lindegren2009_Fish | [8] | 3 | Fish | Atlantic Ocean |
| Vik2008_LynxHare | [9] | 2 | Lynx/Hare | Terrestrial |
| Yamamura2006_Insects | [10] | 3 | Insects | Terrestrial |

Table 3: References used [TO COMPLETE]

References

- [1] T. Hernández Fariñas, C. Bacher, D. Soudant, C. Belin, and L. Barillé. Assessing phytoplankton realized niches using a French national phytoplankton monitoring network. *Estuarine, coastal and shelf science*, 159:15–27, 2015.
- [2] J.R. Griffiths, S. Hajdu, A.S. Downing, O. Hjerne, U. Larsson, and M. Winder. Phytoplankton community interactions and environmental sensitivity in coastal and offshore habitats. *Oikos*, 125(8):1134–1143, 2015.
- [3] V. Huber and U. Gaedke. The role of predation for seasonal variability patterns among phytoplankton and ciliates. *Oikos*, 114(2):265–276, 2006.
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- [6] J.L. Klug, J.M. Fischer, A.R. Ives, and B. Dennis. Compensatory dynamics in planktonic community responses to pH perturbations. *Ecology*, 81(2):387–398, 2000.
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- [8] M. Lindegren, C. Möllmann, A. Nielsen, and N.C. Stenseth. Preventing the collapse of the Baltic cod stock through an ecosystem-based management approach. *Proceedings of the national academy of sciences*, 106(34):14722–14727, 2009.
- [9] J.O. Vik, C.N. Brinch, S. Boutin, and N.C. Stenseth. Interlinking hare and lynx dynamics using a century’s worth of annual data. *Population ecology*, 50(3):267–274, 2008.
- [10] K. Yamamura, M. Yokozawa, M. Nishimori, Y. Ueda, and T. Yokosuka. How to analyze long-term insect population dynamics under climate change: 50-year data of three insect pests in paddy fields. *Population ecology*, 48(1):31–48, 2006.