



HAL
open science

Spatio-temporal variability of a chlorophyll-a based biomass index and influence of coastal sources of enrichment in the Algerian Basin

Romaissa Harid, Hervé Demarcq, Mehdiya-Asma Keraghel, Malik Ait-Kaci, Mohamed Zerrouki, Nour-El-Islam Bachari, Fouzia Houma

► To cite this version:

Romaissa Harid, Hervé Demarcq, Mehdiya-Asma Keraghel, Malik Ait-Kaci, Mohamed Zerrouki, et al.. Spatio-temporal variability of a chlorophyll-a based biomass index and influence of coastal sources of enrichment in the Algerian Basin. CONTINENTAL SHELF RESEARCH, 2022, 232, pp.104629. 10.1016/j.csr.2021.104629 . hal-03708536

HAL Id: hal-03708536

<https://hal.umontpellier.fr/hal-03708536>

Submitted on 8 Jan 2024

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution - NonCommercial 4.0 International License

1 **Spatio-Temporal variability of a Chlorophyll-*a* based biomass index and**
2 **influence of coastal sources of enrichment in the Algerian Basin**

3
4 Romaïssa Harid ^{a,b,*}, Hervé Demarcq ^b, Mehdiya-Asma Keraghel ^a, Malik Ait-Kaci ^a, Mohamed
5 Zerrouki ^a, Nour-El-Islam Bachari ^c, Fouzia Houma ^a

6
7 ^a ECOSYSMarL: Laboratoire des Écosystèmes Marins et Littoraux , École Nationale Supérieure
8 des Sciences de la Mer et de l’Aménagement du Littoral (ENSSMAL), Campus Universitaire de
9 Dely Ibrahim Bois des Cars, B.P. 19, 16320, Alger, Algeria

10 ^b MARBEC, IRD, Ifremer, CNRS, Univ Montpellier, Sète, Avenue Jean Monnet, CS 30171,
11 34203 Sète cedex, France

12 ^c USTHB : Université des Sciences et Technologie Houari Boumedién, Département Écologie et
13 Environnement, BP 32 Bab Ezzouar, 16111 Alger, Algeria

14
15 **The e-mail address and ORCID iD of each authors:**

16 (r.harid@enssmal.dz, <https://orcid.org/0000-0003-1418-7091>), herve.demarcq@ird.fr,
17 <https://orcid.org/0000-0003-1995-1183>), (ma.keraghel@enssmal.dz, [https://orcid.org/0000-0002-](https://orcid.org/0000-0002-5145-4913)
18 [5145-4913](https://orcid.org/0000-0002-5145-4913)), (m.ait-kaci@enssmal.dz), (m.zerrouki@enssmal.dz, [https://orcid.org/0000-0002-](https://orcid.org/0000-0002-8842-9535)
19 [8842-9535](https://orcid.org/0000-0002-8842-9535)), (n.bechari@usthb.dz, <https://orcid.org/0000-0001-6686-2429>),
20 (f.houmabachari@enssmal.dz)

21 * **Corresponding author:** Romaïssa Harid /r.harid@enssmal.dz/ romaïssa.harid@hotmail.fr
22 /+213 556 222 824

23 **Highlights**

- 24 • A specific fortnightly climatology of 1-km resolution Chlorophyll-*a* has been generated
25 over the Algerian basin from MODIS data to explore the coastal domain.
- 26 • The Algerian Basin is characterised by two extreme seasons of high and low biomass,
27 separated by sharp transitions, while the coastal area reveals a separated dynamic.
- 28 • Discontinuous and intense coastal cross-shore gradients reveal specific coastal sources of
29 enrichments.
- 30 • A Chl-*a* based integrated index was defined to determine the importance of coastal
31 enrichments.
- 32 • The Chl-*a* biomass in Algerian coastal waters is mainly associated with enrichments from
33 wadis and cities with a pronounced seasonal effect.

34 **Abstract.** This study investigates the spatial distribution and temporal variability of chlorophyll-
35 *a* (Chl-*a*) biomass in the Algerian Basin (AB) along its meridional and cross-shore dimensions,
36 focusing on coastal enrichments. After correcting most atmospheric disturbances in the daily
37 MODIS Level-2 data series between 2003 and 2018, a fortnightly climatology of 1-km
38 resolution Chl-*a* has been generated to account for specific coastal features previously poorly
39 evidenced from 4-km level-3 data. The AB is characterised by two extreme seasons of high and
40 low biomass, separated by sharp transitions, that characterise the offshore domain. The coastal
41 area (<10 km) reveals an intense and distinct dynamic associated with highly productive local
42 hotspots rather than seasonal variability. A biomass index is proposed as the horizontally
43 integrated Chl-*a* concentration from the coastline to the most offshore extension of the 0.5 mg m⁻³
44 Chl-*a* isopleth. This index separately quantifies the cumulative biomass of both offshore and
45 coastal domains with large alongshore variability. Low values (<5 g m⁻²) were observed in the
46 offshore area during summer and high values during the spring blooms (up to 40 g m⁻²), while

47 maximum values ($>50 \text{ g m}^{-2}$) were locally observed in the coastal domain. The narrow coastal
48 area alone represents 44% of the total biomass, with coastal hotspots where the enrichment is up
49 to 5 times higher than offshore. Multivariate modelling of the potential factors favouring coastal
50 enrichments shows that the phytoplanktonic biomass in coastal waters is mainly associated with
51 enrichments from wadis and seasonally from city sewage as well as by the presence of a bay. A
52 separate source of enrichment is undoubtedly associated with the presence of aquaculture cages.

53 **Keywords:** Ocean colour; alongshore variability; wadi; anthropic enrichment; MODIS.

54 1. Introduction

55 Chlorophyll-*a* (Chl-*a*) biomass is associated with the net primary production in marine
56 ecosystems, and marine physical and biochemical processes strongly influence its variability. A
57 precise description of the spatio-temporal variability of Chl-*a* biomass is necessary to
58 understand coastal marine systems functioning. The Chl-*a* concentration in the southwestern
59 Mediterranean Sea (Med) is closely related to winter mixing and summer stratification. The Med
60 ecosystem is increasingly threatened by human activities in coastal areas, as well as by a
61 continuing warming trend (Vargas-Yáñez et al., 2010). A recent study conducted by Keraghel et
62 al. (2020) highlights that the southwestern Med is a net sink of carbon dioxide, even compared to
63 the Med as a whole, which is a significant contributor at the global level (Khaliwala et al., 2013).
64 Scientists are still trying to understand the evolution of the Med ecosystem to better assess
65 current and future changes and consider solutions to mitigate some impacts of global warming.

66 The Algerian Basin (AB) is classified as mesotrophic (D'Ortenzio and Ribera d'Alcalà, 2009;
67 Harid et al., 2018; O'Reilly and Werdell, 2019). Water exchange across Gibraltar has a
68 significant influence on its general circulation (Béranger et al., 2005; Millot, 1989; Peliz et al.,
69 2009) and controls its nutrient content (Bethoux et al., 2002; Crispi and Pacciaroni, 2009; Elbaz-
70 Poulichet et al., 2001; Huertas et al., 2009), with a direct influence on the Chl-*a* based

71 phytoplanktonic biomass. In winter, the presence of distinct water bodies indicates the eastward
72 movements of anticyclonic eddies (Olita et al., 2011), generated by instabilities in the Algerian
73 current (Millot et al., 1990), which enrich the surface water and increase its primary production.
74 The Atlantic water flow is characterised by a transit time of two to four months between
75 Gibraltar and the Algerian coasts (Millot, 1999), strongly influencing the seasonal Chl-*a* signal
76 (Salgado-Hernanz et al., 2019). In summer, the stability of water masses limits the nutrient input
77 (Moutin and Prieur, 2012), leading to a decrease in phytoplankton production.

78 The continental shelf of the AB is very narrow (15 km on average) and is neglected in most
79 studies. Nonetheless, the shelf is the richest domain of the AB and shelter more complex
80 interactions than in the offshore domain. Ocean colour remote sensing has provided high-quality
81 observations in this respect for over twenty years on the abundance and distribution of Chl-*a*
82 concentration, which is considered a proxy for phytoplankton biomass (Cullen, 1982; Strickland,
83 1965). Turbid waters in the Med are rare compared to other seas (Morel et Prieur, 1977).
84 According to Antoine et al. (1995), the coastal case-2 waters (where other constituents as
85 mineral particles are also present) in the whole Med are estimated to be 5%. Currently, ocean
86 colour analysis coupled with in-situ data could be used to characterise and monitor
87 phytoplankton blooms (Barale et al., 2008; Cerino et al., 2019; Groom et al., 2019; Lavigne et
88 al., 2015; Mayot et al., 2016; Palmiéri et al., 2018). In previous studies, a significant limitation
89 was related to the poor representation of coastal patterns due to Level-3 data at 4-km resolution
90 (as in Lavigne et al., (2015), Marañón et al., (2020), Mayot et al., (2016), Volpe, (2012), Volpe et
91 al., (2018)).

92 To this end, our study proposes a practical approach to improve the quality of the standard
93 MODIS-Aqua Level-2 1-km resolution (swatch) Chl-*a* product, specifically in coastal
94 environments, where the higher data resolution provides more detailed information. A
95 comparison between simultaneous in-situ and satellite Chl-*a* was performed to assess the

96 accuracy of MODIS Chl-*a* data in AB. In addition, a cumulative Chl-*a* biomass index (I_B) was
97 developed to synthesise the spatial patterns and variability of Chl-*a*. This paper describes the
98 seasonal climatology of I_B in AB from 16 years of data (2003-2018), focusing on offshore and
99 coastal areas separately. Finally, a discussion on the influence of different sources of coastal
100 enrichment on the Chl-*a* biomass in the AB is presented.

101 **2. Methods**

102 **2.1 Study area**

103 The Algerian Basin (Fig. 1) is a major energetic area for mesoscale activity throughout the Med
104 (Amores et al., 2013; Pessini et al., 2018). Millot and Taupier-Letage (2005) described the East
105 flowing Algerian current, which carries Atlantic surface water, is 50-100 km wide and 100-200
106 m thick with a speed of some 10s cm s^{-1} (El-Geziry and Bryden, 2010). It generally follows the
107 continental slope and generates small eddies of 10-100 km diameter, lasting a few weeks or
108 months. Periodically, this current forms a growing meander of 50-100 km; it can detach to form
109 an anticyclonic eddy of 100-200 km diameter that encompasses the entire thickness of the Med
110 water (El-Geziry and Bryden, 2010; Fani et al., 2014; Millot, 1989). Some oceanic eddies persist
111 for up to three years, circulating in the AB in a cyclonic circuit (Millot and Taupier-Letage,
112 2005). Thus, the AB acts as a reservoir where Atlantic waters accumulate before flowing either
113 eastwards (surface waters) or northwards (deep waters) of the Med (Millot, 1999). Indeed, this
114 buffer zone decouples the inflow and outflow of Med surface waters.

115 **2.2 Satellite data sources**

116 We used daily Level-2 Chl-*a* concentration data from the MODIS-Aqua sensor from 2003 to
117 2018 at 1-km nominal resolution. The data set consists of 5844 daily observations from 15020
118 individual orbits acquired from NASA's Ocean Color website (NASA's Ocean Color Web, 2019).
119 Each daily data field was remapped over the AB, between 35°N-40°N and 6°W-10°E (Fig. 1), at

120 a spatial resolution of 96 pixels per degree of latitude and longitude. The equivalent daily
121 MODIS Level-3 mapped data set at 4-km spatial resolution was obtained from the NASA Ocean
122 Data Processing System. This data set was compared to the MODIS Level-2 data set to
123 demonstrate permanent coastal Chl-*a* patterns. The climatological period from 1 to 15 January
124 2003-2018 (Fig. 2) illustrates the differences between both spatial resolutions of 1-km and 4-km.
125 In this work, a corrected version of the MODIS Level-2 data at 1-km was used to adequately
126 describe the AB Chl-*a* variability in the coastal and offshore areas.

127 **2.3 Cloud masking improvement of MODIS Level-2 Chl-*a* data**

128 This section describes the specific processing steps applied for the first time to the standard
129 cloud-corrected Level-2 Chl-*a* fields to detect and remove spurious patterns that affect data
130 quality, even in fortnightly averages (Fig. 2b). Specifically, we noticed the presence of (i)
131 partially cloudy pixels at the edge of the cloud mask, resulting in spurious high Chl-*a* values and
132 (ii) noisy pixels. Three criteria were used to discriminate these contaminated pixels: 1) a
133 maximum allowable value associated with a realistic Local Chl-*a* Gradient (LG), 2) a maximum
134 value of daily Chl-*a* change during 3-day periods (Temporal Variation, “TV”), and 3) their
135 position as Isolated Pixels (IP) inside the cloud mask. The pixel values corresponding to any of
136 these criteria cited here are selected and replaced by the missing value.

137 *2.3.1 Local Gradient (LG) criteria*

138 First, we applied a 3x3 Gaussian filter (Eq. 1) followed by a Sobel Gradient filter (Sobel, 1990)
139 as follows:

$$140 \quad k(x, y) = \frac{1}{2\pi\sigma^2} \exp\left(-\frac{x^2+y^2}{2\sigma^2}\right) \quad (1)$$

141 where, $k(x,y)$ is the matrix of the kernel used to convolute the original image; x and y are the
142 pixel’s position along the abscissa and ordinate axes respectively. σ^2 is the variance of the 3x3
143 pixel matrix.

144 The horizontal and vertical components of the Sobel gradient (Eq. 2 and Eq. 3, respectively)
 145 were separately computed and combined into the final gradient (Eq. 4):

$$146 \quad G_h[x][y] = k(x, y) * \begin{Bmatrix} -1 & -2 & -1 \\ 0 & 0 & 0 \\ 1 & 2 & 1 \end{Bmatrix} \quad (2)$$

$$147 \quad G_v[x][y] = k(x, y) * \begin{Bmatrix} -1 & 0 & 1 \\ -2 & 0 & 2 \\ -1 & 0 & 1 \end{Bmatrix} \quad (3)$$

$$148 \quad Sobel_G [x][y] = \sqrt{G_h^2[x][y] + G_v^2[x][y]} \quad (4)$$

149 As spurious gradients are mainly associated with atmospheric perturbations in the vicinity of
 150 clouds, a maximum threshold value of the local Sobel gradient of Chl-*a* was considered to detect
 151 outlier pixels (Fig. 3a), at a maximum distance of 5 km from cloud borders (black areas in Fig.
 152 3b). After several tests, two thresholds were defined: 0.4 mg m⁻³ km⁻¹ and 1 mg m⁻³ km⁻¹
 153 respectively for the coastal zone (distance from the coast <5 km) (Fig. 3c) and the offshore area.

154 2.3.2 Temporal Variability (TV) criteria

155 Each daily individual Chl-*a* pixel value of a given day (Day₀) was compared to the average value
 156 of the previous day (Day₋₁) and the following day (Day₊₁) when one or both values are available
 157 to detect anomalous Chl-*a* variations through time, as given by Eq. 5:

$$158 \quad TV = Day_0 - \left(\frac{Day_{-1} + Day_{+1}}{2} \right) \quad (5)$$

159 This difference was then compared with the maximum threshold of temporal variability set at 2
 160 mg m⁻³. Furthermore, this test was applied in the offshore domain only (distance from coast >20
 161 km) to consider the higher spatio-temporal variability of the coastal environment (Fig. 3e).

162 2.3.3 Isolated Pixel (IP) removal

163 This criterion is used to eliminate pixels closely associated with a cloud structure. We considered
 164 that Chl-*a* pixels bordered by more than five cloudy pixels (including those isolated in a cloud)

165 belong to the same atmospheric structure and should be eliminated (Fig. 3f).

166 2.3.4 *Combination of criteria*

167 The three criteria mentioned above were cumulated, and the pixels marked by at least one
168 criterion were removed. The resulting daily data are significantly less noisy, as shown in Fig. 3h.

169 A fortnightly climatology of Chl-*a* at 1-km resolution is then computed for the whole AB.

170 **2.4 Comparison between in-situ and satellite Chl-*a* data**

171 We compared the resulting 1-km satellite Chl-*a* data with an initial set of 70 high-performance
172 liquid chromatography (HPLC) measurements of surface Chl-*a* concentration obtained during
173 the SOMBA (*Système d'Observation à la Mer dans le Bassin Algérien*) cruise in the AB between
174 August 13 and September 10, 2014 (Mortier et al., 2014). Further details about the cruise are
175 available at <https://doi.org/10.17600/14007500>. Calibration precision was estimated to be 0.4%
176 from the HPLC 1200 instrument used for the Chl-*a* measurements. A total of 34 measurements
177 were retained according to their correspondence with satellite passes within ± 6 hours. The
178 nearest pixel at 1-km resolution was considered. A representativity error (RE) was also
179 considered, based on the proximity of a high Chl-*a* gradient, which we defined at a value of 0.01
180 $\text{mg m}^{-3} \text{ km}^{-1}$, compared with the effective Chl-*a* gradient measured in a 5x5 pixel matrix (Error
181 bars in Fig. 4). In contrast, this spatial variability is generally low for pixels far from eddies (Fig.
182 4b and 4e). We also checked the proximity of the HPLC measurements to the clouds (less than 5
183 km); only two measurements deviate from this criterion (Fig. 4c and 4d).

184 **2.5 Biomass index**

185 A modified version of the coastal Chl-*a*-based index developed by Demarcq et al., (2007) was
186 applied. Based on the Chl-*a* concentrations, a threshold (β) of Chl-*a* concentration was set to
187 compute the proposed integrated Chl-*a* biomass index (I_B) in AB. This threshold was chosen as a
188 value observed during all seasons (Fig. 7b). A value of 0.5 mg m^{-3} was selected according to this

189 criterion for the present study. The integrated Chl-*a* biomass index was calculated at each coastal
190 point with the following formula:

$$191 \quad I_B = \left(\sum_{dist=Dist_{min}}^{Dist(\beta, max)} \overline{Chla} \right) \times Dist_{(\beta, max)} \quad (6)$$

192 where, the average Chl-*a* is calculated by the formula:

$$193 \quad \overline{Chla} = \left(\sum_{i=1}^{n(\beta, max)} Chla \right) / n_{(\beta, max)} \quad (7)$$

194 where, $n_{(\beta, max)}$ is the position of the most distant pixel of the cross-shore transect.

195 The cross-shore distance associated with the index $Dist_{(\beta, max)}$ is the most distant position where
196 $Chl-a \geq \beta$ with the constraint $Dist_{(\beta, max)} \leq max$, is the maximum allowable distance. When this
197 distance is determined, all pixel values from $Dist_{min}$ to $Dist_{(\beta, max)}$ are considered, regardless of
198 their value; sometimes slightly $< \beta$. If no data $\geq \beta$ was found within a profile, the computation of
199 I_B was performed only for the first valid pixel at the only $Dist_{min}$ position, regardless of its value.
200 The maximum distance for $Dist_{(\beta, max)}$ in km was chosen at 10 km for the coastal area and 80 km
201 for the next offshore area, according to the average structure of the cross-profiles (Fig. 6b). It is
202 important to note that for continuity reasons, the first (inner) pixel of the offshore area was
203 considered the pixel immediately offshore of the last of the coastal areas. The integrated coastal
204 and offshore biomass indices were then calculated for the entire Algerian coastline or part of it
205 (Table 1).

206 To estimate the relationships between Chl-*a* biomass (Chl-*a* or I_B) classes and factor variables
207 representative of the different sources of coastal enrichment, we applied the Generalized Linear
208 Model (GLM) approach. All data analysis was done using the “stats” package version 3.4.4 of
209 the R software.

210 **2.6 Physical oceanographic data**

211 *2.6.1 Altimetry data*

212 Geostrophic ocean currents and total kinetic energy (TKE) were extracted from the CMEMS

213 (Copernicus Marine Environmental Service) database of the *SEA-LEVEL GLO PHY L4 REP*
214 *OBSERVATIONS 008 047* altimetry data product (<http://marine.copernicus.eu>, last accessed
215 February 27, 2019), for the same period, and remapped in the AB at a spatial resolution of 0.25°
216 (Fig. 1 for geostrophic currents and Fig. S2b for TKE).

217 2.6.2 *Mixed Layer Depth*

218 The mixed layer depth (MLD) has been defined in previous studies (as Lavigne et al., (2015) and
219 Volpe, (2012)) using in-situ data in the Med (AB included). In this study, the monthly
220 climatology (1969 to 2013) of the MLD was used as defined by Houpert et al. (2015) (data from
221 <https://www.seanoe.org/data/00354/46532/>).

222 2.6.3 *Wadis outflows*

223 Outflow data measured by the Algerian National Agency for Hydraulic Resources (*Agence*
224 *Nationale des Ressources Hydrauliques*, ANRH, <http://www.anrh.dz/>) were used to evaluate the
225 possible influence of wadis (temporary rivers) on the Chl-*a* variability along the Algerian shelf.

226 3. Results

227 We show that a high-resolution fortnightly climatology very significantly improves the
228 description of the spatio-temporal variability of Chl-*a* (including abrupt seasonal changes) and a
229 spatio-temporal view of the enrichment sources. We explore here the coastal and offshore
230 domains along meridional and cross-shore transects, focusing on the coastal domain, to
231 understand the main seasonal dynamics of these enrichments.

232 3.1 Impact of data resolution on the description of seasonal patterns

233 Firstly, we compared the standard (original) MODIS data (1-km) to the corrected (this work)
234 MODIS data (1-km). The improvement is particularly high during winter (characterised by a
235 large cloud cover) with better detection of atmospherically contaminated pixels (Fig. 5). An
236 example of the impact of Chl-*a* outlier values in the spatial distribution of the time series
237 averages is represented in Fig. 5a and Fig. 5b for a fortnightly, and in Fig. 5c and Fig. 5d for a
238 monthly climatological average. This correction produces a moderate decrease in Chl-*a* mainly

239 during the productive season, reinforcing the describing cross-shore profile. The resulting
240 fortnightly climatology of MODIS Level-2 Chl-*a* data (at 1-km resolution) in the AB can be
241 found online at <https://doi.org/10.5281/zenodo.5390383>.

242 The new fortnightly climatology at 1-km resolution was compared to the 4-km resolution used in
243 all previous studies. Seasonal variability of Chl-*a* from the corrected MODIS 1-km Level 2 data
244 (Fig. 6b) was explored along average cross-shore transects and compared to the MODIS 4-km
245 Level-3 data (Fig. 6a). A closer look at the shorter distances (0-10 km) (Fig. 6c and 6d) shows
246 that, as expected, the improvement is very significant and highlights much stronger cross-shore
247 patterns (Fig. 6e and 6f for the most contrasted months of March and August), both in terms of
248 Chl-*a* concentration average and seasonal patterns. The coastal Chl-*a* (0-10 km) from the 1-km
249 data is 37% higher than that from 4-km data (49% and 46% respectively at distances of 2 and 4-
250 km from the coast). The 4-km product cannot detect a significant part of the coastal enrichment,
251 representing 44% of the production of the AB from 1-km data, while only 25% are detected from
252 4-km data. Consequently, the spatial resolution impacts the scale of the description and more
253 importantly the high contribution of the coastal area in the regional marine productivity.

254 **3.2 MODIS Level-2 data validation in the AB**

255 The in-situ Chl-*a* data range between 0.062 and 0.307 mg m⁻³ (Fig. 4). These values are typical
256 for the AB offshore area during the oligotrophic season in the surface layer. Both sources of Chl-
257 *a* data span nearly the same magnitude. The HPLC data have a slightly lower mean and median
258 (respectively 0.100 mg m⁻³ and 0.094 mg m⁻³) than the satellite data (0.105 mg m⁻³ and 0.104 mg
259 m⁻³). It should be noted that the satellite has a vertically integrating effect (exponentially
260 decreasing) on the estimated Chl-*a* value. In contrast, the in-situ measurements represent
261 exclusively punctual surface values at 1 m depth and the remaining ones at 2 m depth. The final
262 error associated with the Chl-*a* satellite data was estimated at 0.025 (8% of the average) by the
263 RMSD (Root Mean Square Deviation) between Chl-*a* HPLC and MODIS Level-2 data.

264 No in-situ measurements were available in the coastal areas of the AB. Nevertheless, Pieri et al.,
265 (2015) have found that the OC3M standard algorithm (used in our work) gives valid results in
266 the Western Mediterranean Sea when the Chl-*a* concentration does not exceed 1 mg m⁻³. In our
267 case, the Chl-*a* exceeds 1 mg m⁻³ generally in the three first kilometers from coast (i.e. the 3 first
268 pixels) and only during the high production season (December to March, as shown in Fig. 6 and
269 7). To estimate the importance of the likely overestimation of the values >1 mg m⁻³, we apply an
270 empirical correction model with two levels of intensity, by reducing the values >1 mg m⁻³ by a
271 factor of two and by a factor of three. The results show a relatively modest overestimation of
272 respectively 6% and 9% of the Chl-*a* in these two extreme cases. This allows us to assume that a
273 likely overestimation of the Chl-*a* values in the very coastal area does not significantly impact
274 our conclusions.

275 **3.3 Cross-shore and seasonal variability of Chl-*a***

276 Chl-*a* cross-shore sections (Fig. 6b) indicate that the lowest Chl-*a* concentrations are observed
277 from May to October at all locations, both in coastal and offshore areas. The highest Chl-*a*
278 concentrations, representing the productive season, are observed from November to April. In the
279 coastal area, the maximum Chl-*a* can exceed 2 mg m⁻³ in winter (Fig. 7a) and 0.5 mg m⁻³ in
280 summer (Fig. 7a). However, in the offshore area, beyond 10 km from the coast, the average
281 minimum reach 0.5 mg m⁻³ in winter and 0.2 mg m⁻³ in summer (Fig. 6b and Fig. 7b). Beyond 10
282 km, the Chl-*a* concentration becomes stable during all seasons (Fig. 6b and Fig. 7a). Indeed, we
283 chose the distance of 10 km from the coast as the shortest distance at which Chl-*a* seasonality
284 becomes weak and stops increasing towards the offshore (Fig. 6b and 7b).

285 The cross-shore gradient is well pronounced throughout the year and increases exponentially
286 towards the coast (Fig. 6b). The intensity of this gradient is maximum during the productive
287 season and is always maximal at the coast and regularly decreases with increasing distance from
288 the coast (Fig. 6b), ranging from 0.2 to 2 mg m⁻³ onshore (Fig. 6b) and 0.1 to 0.5 mg m⁻³ offshore

289 (Fig. 6b).

290 **3.4 Meridian variability and seasonality of Chl-*a* biomass index**

291 The integrated cross-shore biomass index (Section 2.6) was computed from each coastal point
292 northwards to integrate the Chl-*a* concentration up to a variable distance (Fig. 8b) where Chl-*a*
293 drops below the predefined value of 0.5 mg m^{-3} (see methods). The aim was to explore and
294 quantify the integrated coastal Chl-*a* biomass along the Algerian coastline (Fig. 8d). The value
295 was carefully chosen as the best threshold $\beta = 0.5 \text{ mg m}^{-3}$ that intersects the different average
296 cross-shore climatological profiles over the year (Fig. 7b). The red line in Fig. 8a represents the
297 maximum distance effectively reached during the productive season (we considered Chl-*a* <0.5
298 mg m^{-3} as oligotrophic and does not represent an enrichment). The resulting distance (Fig. 8b)
299 shows that the productive area varies over time from a few kilometres in summer (cyan area in
300 Fig. 8b) with a minimum of 1 km (when only one sea pixel is considered) to a maximum of 80
301 km in winter, mainly reached near the Alboran Sea in the west. The longitudinal gradient along
302 the coastline shows an apparent decrease of this distance eastward, which is well in line with the
303 decreasing influence of the enrichment of Atlantic waters entering the Alboran Sea and moving
304 eastwards.

305 The average Chl-*a* along the same transect (Fig. 8c) shows extreme variability between regions,
306 from values $<0.5 \text{ mg m}^{-3}$ in summer (this is possible when a minimum of one sea pixel is
307 considered) to values $>2 \text{ mg m}^{-3}$ between April and November, i.e., during low-biomass season.
308 The resulting biomass index I_B (Fig. 8d) is defined as the product of the previous distance by the
309 average Chl-*a* concentration along the same transect (Eq. 6). This index represents the spatial
310 integration of the most elevated Chl-*a* values along the cross-shore transect, while the vertical
311 dimension is partially considered by the attenuation depth of the satellite measurement.
312 Nevertheless, this depth represents a variable fraction of the euphotic layer, according to the
313 shape of the vertical Chl-*a* profile.

314 The Chl-*a* (Fig. 8c) represents a proxy of the average phytoplankton biomass over the cross-
315 shore profile, while the spatially integrated index (I_B , Fig. 8d) is predominantly determined by
316 the length of the profile (Fig 8b). The I_B index is expressed in g m^{-2} and varies between 20 and
317 50 g m^{-2} during the productive season (Fig. 8d) with a regular eastward decrease. Several peaks
318 in phytoplanktonic biomass are observed (I_B is $>45 \text{ g m}^{-2}$; between 2.2°W and 0.5°E from
319 January to March, between 5.1°E to 5.5°E in January and February, and at 7.7°E from January to
320 March-April). These peaks correspond to pronounced extensions of $\text{Dist}_{0.5\text{mg}}$ (Fig. 8b) as near the
321 Alboran region or mostly to higher Chl-*a* values in the central and eastern parts of the AB (Fig.
322 8c). I_B is $<10 \text{ g m}^{-2}$ everywhere during the low-biomass season, except in the Algiers and Annaba
323 bays (Fig. 8d). I_B is, in fact, practically zero from June to September in many locations where the
324 Chl-*a* concentration barely exceeds 0.5 mg m^{-3} (Fig. 8c).

325 **4. Discussion**

326 The construction of a data set at 1-km spatial resolution allows investigating and extracting the
327 specific enrichments of coastal origin, distinguishable from the large-scale seasonal cycle.
328 Previous descriptions of the climatological cycle in the AB were based on monthly averages
329 (instead of fortnightly in this study) and at a much higher spatial resolution. Therefore, our
330 description of the cross-shore gradient of Chl-*a* and its variability along a longitudinal gradient
331 becomes much more realistic and highlights the high importance of the coastal domain ($<10 \text{ km}$)
332 in the whole basin.

333 **4.1 Processes governing Chl-*a* variability in AB**

334 *4.1.1 Seasonal variability*

335 AB is characterised by two contrasting seasons (Fig. 7a): an early 3.5-month high-biomass
336 season (mid-December to March) and a 4.5-month low-biomass season (June to mid-October),
337 characterised by intense stratification. The two seasons are separated by quasi symmetrical and
338 sharp 2-month transition periods in spring and autumn (Fig. 7a). The extreme precocity of the

339 productive period (between October and November), i.e. during a low sun elevation, clearly
340 shows that light is not the main limiting factor in the occurrence of winter blooms. The
341 seasonality of Chl-*a* is closely related to the dynamics of the Mixed Layer Depth (MLD, Fig. 7a,
342 brown line and Fig. S1 in supplementary material), which is maximum (40 m to 60 m) in winter
343 between December and February and very low (15 m) in summer from June to September.

344 It is well known that winter and spring blooms in the region are almost exclusively driven by the
345 nutrient input following autumn and winter vertical mixing (Fani et al., 2014; Huertas et al.,
346 2012; Lazzari et al., 2012; Millot et al., 1990; Pasqueron de Fommervault et al., 2015), as
347 reflected by our biomass index (Fig. 8d). Moreover, the results show that high Chl-*a* values
348 dominate several well-defined coastal areas outside the productive season, from April to
349 November (Fig. 8c). On a large scale, the variability of Chl-*a* concentration in the AB is known
350 to be driven by the inflow of nutrient-rich Atlantic waters that enter the Alboran Sea through the
351 Gibraltar strait (Taupier-Letage and Millot, 1988) and progress eastwards along the AB. The
352 same conclusions have also been drawn more recently by several authors (Fani et al., 2014;
353 Huertas et al., 2012; Lazzari et al., 2012; Pasqueron de Fommervault et al., 2015). Consequently,
354 the eastward propagating eddies modulate the circulation of water masses beyond the continental
355 shelf (Pessini et al., 2020), generating intense vertical mixing (Millot et al., 1990). Its positive
356 influence on productivity is perceptible up to the eastern part of the country and is reinforced by
357 nutrient enrichments from the bottom (Millot et al., 1990). The offshore vertical mixing is
358 considered to be the main factor influencing winter-spring enrichments before the summer-
359 autumn stratification period.

360 In addition to these two well-known potential sources of enrichment, we identified a third coastal
361 source: the presence of nutrients of coastal origin, generally associated with bays or gulfs, which
362 enhance local phytoplankton growth. The integrated Chl-*a* biomass index (I_B , Fig. 8d) is used in
363 this study as a proxy for the primary production dynamic in the AB. Better than local Chl-*a*

364 averages, it adequately describes the longitudinal variability of spring blooms due to its cross-
365 shore integrative capability. In other words, the Chl-*a* averages (Fig. 8c) gives a clear view of the
366 origin of the enrichment effects without considering their spatial importance. In contrast, the I_B
367 (Fig. 8d) integrates both components.

368 However, this index in Fig. 8d (and the associated average Chl-*a*) does not distinguish between
369 coastal and offshore sources of enrichment. We, therefore, divided it into an inshore and an
370 offshore component, as detailed in Section 2.6. The coastal area is defined as the distance
371 between the coast and the 0.5 mg m^{-3} isopleth position, with a maximum distance of 10 km (Fig.
372 6b). The offshore component is then defined as the area beyond this variable spatial limit up to a
373 maximum distance of 80 km. The maximum 10 km limit was chosen to best separate the coastal
374 and offshore signals, from the Chl-*a* signature (Fig. 9a-b) and the corresponding integrated
375 biomass index (Fig. 9c-d).

376 4.1.2 Coastal enrichment

377 Beyond the spatially averaged seasonal signal computed in both domains (Fig. 7a), the results
378 give precise insights about their regional alongshore variability, which is well distinguished by
379 the biomass index (Fig. 9c-d). The coastal biomass index (I_B , Fig. 9c) highlights the increase in
380 the duration of the productive season varies, that from 4 to 6 months from East to West and from
381 4 to 8 months (and more) in the coastal areas in the form of spatially distinct peaks of values >20
382 g m^{-2} . Some locations, such as the Algiers Bay (3.2°E) and the Annaba Bay (7.9°E), show high
383 index values almost yearly. In contrast, no Chl-*a* peaks are visible in the offshore area (Fig. 9d),
384 even in the locations where the highest coastal peaks are observed (Fig. 9c). The offshore area
385 exhibits a much more homogeneous spatial pattern with high cumulated biomass during the
386 central part of the productive season, with a maximum between January and February, except
387 near the Alboran Sea, where the maximum occurs one month later. The duration of the
388 productive season considering the I_B at 10 g m^{-2} limit, varies from 5 months (December to April)

389 west of 1°E with maximum values constantly between 30 g m⁻² and 40 g m⁻², then is 4 months in
390 most of the area up to 7°E, to a minimum duration <2 months (mid-January to February) with
391 maximum values <25 g m⁻². Between 5°E and 6°E longitude, a distinct offshore maximum in the
392 biomass index (Fig. 9d and Fig. 11, blue curve) is observed during the productive season. This
393 feature probably corresponds to the relative permanence of the mesoscale cyclonic EAG (Eastern
394 Algerian Gyre, Fig. 1) as described by Testor et al., (2005) and where high Chl-*a* concentrations
395 are regularly found (Taupier-Letage, 2003).

396 The same processing was applied to compare these results with the equivalent information from
397 the 4-km Level 3 data and presented in Supplementary Fig. S2. As previously shown, the coastal
398 fraction of the enrichment is here only 25% (44% from 1-km data), and only two areas with Chl-
399 *a* peaks would be partially detected, regardless of the season. This clearly shows that 1-km data
400 are a minimum requirement to adequately explore the relative importance of the different sources
401 of enrichment in the coastal regions.

402 Coastal and offshore averages of the biomass index were calculated for all areas where high
403 biomass index values were observed, hereafter referred to as **High-Biomass Coastal Zones** (or
404 **HBC**), and outside these zones referred to as **Low-Biomass Coastal Zones** (or **LBC**). The aim is
405 to separate the specific effects of local (coastal) enrichments from the large-scale enrichments
406 (Table 1, left-hand side). Compared to LBC, HBC logically dominate the total biomass by a 4-
407 fold ratio (+300%) in summer, and almost double by +80% in winter (Table 1, and Fig. S3a). The
408 importance of HBC is still high in the offshore domain in summer (+97%), while almost no
409 difference is observed (+7%) in winter during the productive season (see Fig. S3b for more
410 details). The annual cumulative biomass index associated with the HBC represents 88% of the
411 coastal domain, despite a corresponding coastline fraction of 42%. Because of the relative
412 importance of the winter period (December to March) in the productive season, we summarised
413 the overall significance of the coastal sector by its ratio to the offshore sector (Table 1, right-

414 hand side). This ratio shows that the coastal domain slightly dominates from January to February
415 in both LBC and HBC (respective values of 0.64 and 0.73). Nevertheless, this ratio strongly
416 increases in the adjacent months (December and March), with values of 2.57 and 4.14 for LBC
417 and HBC respectively, highlighting the importance of HBC in coastal areas.

418 *4.1.3 Large scale longitudinal variability*

419 We specifically examined three profiles of the average biomass index and the MLD and TKE
420 (Fig. 10), both averaged from the coast to 80 km offshore, to explore the longitudinal variability
421 of the phytoplankton biomass over the year. Three cross-shore transects from the coastline to 80
422 km offshore were examined at three longitudes (1°W, 4.5°E and 7.5°E, blue dashed line in Fig.
423 8a). They were chosen because of their independence from the observed enrichments in the
424 coastal domain (Fig. 8c).

425 Firstly, the results show a substantial eastward decrease in the intensity of the productive season,
426 as shown by the biomass index (Fig. 10, orange bars), with annual cumulative values of 285, 152
427 and 86 g m⁻² from West to East. This quantifies well the observations of a progressive decrease
428 in the influence of the rich Atlantic waters eastward, following its progressive nutrient depletion.
429 The shape of the productive season is stable at 1°W and 4.5°E, with a variable maximum centred
430 at the first fortnight of February. A precise computation of this chronology along the entire
431 Algerian coast (data not shown) shows that this central position is relatively stable from 1°W to
432 8.7°E (with minor variability <1 fortnight). In contrast, there is a positive shift of almost a
433 fortnight (first fortnight of March) between 3°W and 1°W near the Alboran Sea, where a
434 maximum shift of two fortnights is observed (data not shown). The productive season is more
435 extended, due to the higher nutrient content of Atlantic waters.

436 We know that winter mixing due to wind and currents is the main cause of nutrient availability in
437 the euphotic layer of the region (Conan et al., 2018; Fernandez et al., 2006; Kessouri et al., 2018;
438 Millot, 1989). A close relationship was effectively observed between the seasonality of the MLD

439 and, to a lesser extent, the TKE (see Fig. S1 for spatial mapping of these two parameters) and the
440 dynamics of the productive season captured by the biomass index. The MLD deepens eastwards
441 with a simultaneous winter maximum (45 m to 60 m, see Fig. S1a) from January to February
442 while the Modified Atlantic Waters (MAW) are drifted eastwards (Font et al., 1998). The
443 productive season follows the winter mixing (average MLD >20 m, blue line) by about one
444 month in the western and central part of the region (Fig. 10a-b). At the same time, a relative
445 synchrony is progressively reached further East (Fig. 10c). The TKE (computed from
446 geostrophic currents, i.e., including eddy energy and permanent currents) is almost permanently
447 high at 1°W near the Alboran Sea (Fig. 10a) and not in phase with the biomass index.

448 It is very likely that, along with the significant enrichment effect of the MAW, water mixing is an
449 important factor in the initiation of surface productivity but is probably not a limiting factor near
450 the Alboran Sea with high-energy levels related to both Alboran gyres. Lower energy levels are
451 observed further East at 4.5°E (Fig. 10b) with a slight relationship to the biomass index, while a
452 higher association is observed at 7.5°E (Fig. 10c). This well-defined seasonal pattern explains
453 that the higher MLD (50 to 60 m) observed in January-February, induces a rapid mixing of
454 surface waters and a rapid increase in planktonic biomass during the second half of February.
455 Average wind speed (data not shown) do not show any relationship with the surface water
456 mixing, with low winter wind values from October to February. The MAW trajectory determines
457 the large-scale variability of planktonic biomass in the AB. It is strongly constrained by nutrient
458 availability, while the different sources of coastal enrichment represent a significant contribution
459 throughout the year and unexpected relative importance of about two-thirds during the central
460 part of the productive season and progressively higher during the rest of the year.

461 **4.2 Sources of coastal enrichment**

462 Previous studies using Chl-*a* variability as a proxy of the phytoplankton biomass variability
463 along the Algerian basin have focused on large spatial scales, mainly over the continental shelf

464 (Mayot et al., 2016; Pieri et al., 2015; Salgado-Hernanz et al., 2019). They have generally
465 ignored small scales and coastal waters, except locally for sanitary purposes or risk assessment.
466 The annual average of the I_B averaged spatially from all seasons was calculated in the coastal
467 domain (0-10 km, green curve in Fig. 11) and in the offshore domain (beyond 10 km, blue curve
468 in Fig.11). The ratio (I_B Coastal/ I_B Offshore) is considered a relative indicator of the local coastal
469 enrichment (red curve in Fig. 11). The I_B coastal peaks are variable in space and time (Fig. 11)
470 and indicate many distinct anomalies. In the Arzew, Bou-Ismaïl, Algiers and Annaba bays, the I_B
471 average is $>20 \text{ g m}^{-2}$ throughout the seasons (Fig. 11). These anomalies correspond mainly to
472 sandy coasts (orange bars in Fig. 11). It is important to note that many microphytobenthos
473 species (some diatoms, cyanobacteria, chlorophyceae and/or flagellates) prefer shallow sandy
474 coastal environments for their development (Cook and Røy, 2006; Hassan et al., 2006).
475 Nevertheless, the origin of these high production areas remains unclear.

476 Several bays are associated with wadis (temporary rivers) in many locations along the coast (in
477 blue in Fig. 11). These wadis are characterised by shallow flows (Fig. 12), generally $<15 \text{ m}^3 \text{ s}^{-1}$
478 during all year seasons. For example, in the Bou-Ismaïl Bay (Fig. 12a), the Mazafran flow varies
479 from $<4 \text{ m}^3 \text{ s}^{-1}$ in summer (June to October) and from 4 to $13 \text{ m}^3 \text{ s}^{-1}$ in winter. In Algiers Bay
480 (Fig. 12b), the El-Harrach flow presents a very similar pattern. The Algiers city is affected by
481 intensive urbanisation (~70% of the coastline up to 800 m inland is urbanised), which has caused
482 significant environmental degradation of the coastal area and impacted the coastal morphology
483 (Rabehi et al., 2019). In Annaba Bay, the Seybouse wadi (Fig. 11, wadi 1) input is highly
484 concentrated in PO_4 and NH_4 compared to Mediterranean rivers (Ounissi et al., 2014), and
485 presents a potential risk of eutrophication (Ziouch et al., 2020). The observed peaks of *Chl-a*
486 (and I_B) are associated with different contributions (Table 2): the type of coast (sandy and rocky),
487 the presence of wadis and large cities, and the presence of bays.

488 In many cases, coastal enrichments are not the result of a single factor: for example, a biomass

489 peak is observed at 7.3°E associated with a wadi (wadi 2). The peak disappears shortly at 7.5°E,
490 despite the presence of sandy coast. In Oran Bay, a coastal peak is observed despite the absence
491 of a sandy coast and wadis (Fig. 11). However, Oran is a large city (>100,000 inhabitants) (Fig.
492 11), that induces a significant marine pollution due to wastewater discharged into the sea,
493 increasing nutrients in coastal waters. Another enrichment is observed at approximately 1.3°W,
494 where floating aquaculture cages are installed (Fig. 11). These aquaculture facilities are
495 considered a significant source of local enrichment (Cao et al., 2007). The low flows of the
496 wadis on the Algerian coast suggest that suspended matter (SM) from terrestrial inputs is weak in
497 coastal waters. In contrast, a significant source of SM may originate from local sediment
498 resuspension, especially on sandy coasts, where sediment resuspension generates a considerable
499 flux of nutrients (Robinson and Hill, 2005).

500 It is also known that the inner part of gulfs and bays trap nutrients from territorial inputs by
501 modifying local hydrodynamics that limit nutrient dispersal, thus maintaining high coastal
502 production with little influence on offshore production. It is important to note that, all previous
503 studies (Colella et al., 2016; Okubo, 1973; Pingree and Maddock, 1979; Signell and Geyer,
504 1991; Taillandier et al., 2020; Wolanski and Hamner, 1988) never considered potential coastal
505 influences. In winter and early spring, the richer coastal waters are often mixed with the offshore
506 waters and therefore contribute to the production beyond the continental shelf up to 10 km from
507 the coast, as shown by our biomass index.

508 **4.3 Modelling approach**

509 As previously mentioned, the Algerian coastal waters were divided into two classes: HBC (High-
510 Biomass Coastal Zones) and LBC (Low-Biomass Coastal Zones) (Fig. 11). The HBCs
511 consequently refer to the highest values of Chl-*a* and I_B, and the LBCs to the lowest values.
512 Their separation into two classes was visually optimised by defining specific thresholds for Chl-
513 *a* and I_B variables, summarised in Table 3 (response variables). We defined three qualitative

514 variables: the presence of wadis, the type of coast (sandy/rocky), and the existence of a Bay. The
515 City is defined as a quantitative variable with four levels: 0 (no-city), 25 k, 75 k, and 200 k
516 inhabitants.

517 Table 2 summarises the respective characteristics of LBC (numbered 1-16) and HBC (numbered
518 1-15) as manually selected in Fig. 11. The presence of Cities, Wadis and Bays are positively
519 related to the detection of High-biomass coastal areas. At the same time, the type of coast
520 appears to be irrelevant, mainly compared to the presence of a Bay.

521 Linear qualitative models (General Linear Models) were performed to evaluate the interactions
522 between either the coastal Chl-*a* concentration or the biomass index (I_B) and the four explanatory
523 variables, as shown in Table 3. The two specific areas HBC-1 and HBC-8 were excluded from
524 the modelling because these two areas are specifically influenced by aquaculture floating cages
525 that are not associated with the explanatory variables. A general model (m1 in Tab. 3) is first
526 tested by combining all seasons to test the separation between HBC and LBC, as presented in
527 Fig. 11. This model explains 67% of the variability, with a unique City effect. The Wadi effect is
528 absent, probably because of its association with City. On the opposite, in winter, the Chl-*a*
529 response variable (m2 model, 79%) is primarily associated with the presence of a Wadi, then to
530 City. In contrast, the I_B response variable (m3 model) is mainly related to a Bay and City
531 presence. The winter I_B model (m3) is very similar in explaining the biomass variability (77%),
532 with a dominance of City presence (as previously with Chl-*a*) as well as to a significant Bay
533 effect. In these winter models, the Wadi effect is only evidenced by the Chl-*a* variable that most
534 reflects the influence of local enrichments rather than their spatial extension, associated with the
535 biomass index (I_B), highlighting the Bay effect.

536 During summer (low-biomass season), the Chl-*a* based model (m4 in Tab. 3) shows only 57% of
537 explanation, with a unique Wadi effect despite the generally low flow of wadis in winter (Fig.
538 12). The equivalent model for the biomass index (m5) explains 78% of the variability, with a

539 dominance of Bay presence, while the Wadi effect is still present. The lower biomass variability
540 in summer (not shown) is better explained by the spatially integrated biomass index (I_B). The
541 later highlights a Bay effect, even if the main variability of the coastal enrichments is probably
542 dominated by wadis and underneath by the influence of cities sewage.

543 Therefore, we can argue that anthropic effects (presence of a City and a Wadi) dominate the
544 biomass variability in the coastal areas along the Algerian coast, much more than “natural”
545 effects such as the coast type and the presence of a bay. However, the presence of Bay is also of
546 primary importance for trapping enriched water within the coastal domain. Another significant
547 point is undoubtedly the strong positive effect of aquaculture cages in two specific country
548 locations (Fig. 11). It is noteworthy that marine aquaculture has developed considerably over the
549 last decade, with a national initiative plan whose objective was to produce 100,000 tonnes of fish
550 and shellfish by 2020 horizon (FAO, 2019).

551 **5. Conclusion**

552 Satellite-based *Chl-a* is an important proxy of phytoplanktonic biomass that allows us to
553 disentangle very different dynamics between the coastal and offshore domains of the Algerian
554 Basin (AB), characterised by a very narrow continental shelf. We show that a specific fortnightly
555 climatology of 1-km resolution *Chl-a* generated from MODIS data makes possible this
556 identification. The AB is characterised by two extreme high and low biomass seasons, separated
557 by short 2-month transition periods. The offshore variability is closely related to large-scale
558 processes governed by the influence of Atlantic waters and a progressive eastward decrease in
559 biomass. The coastal domain reveals a very distinct dynamic associated with highly productive
560 hotspots rather than a well-defined seasonality. The irregular morphology and nature of the
561 Algerian coast (bays, gulfs, rocky or sandy coasts) is shaped by numerous terrestrial and
562 temporary inputs that affect its local productivity. A *Chl-a* based spatially integrated index allows

563 us to quantify the importance of these coastal enrichments. At the same time, a modelling
564 approach shows that seasonal wadis and city sewages, along with the presence of a bay, explain
565 up to 79% of the presence of these productive hotspots. A separate source of enrichment is
566 undoubtedly associated with the recent presence of aquaculture cages. Finally, considering
567 phytoplanktonic communities and the in-situ determination of water quality would be beneficial
568 to understand the biological consequences of these enrichments.

569 **Acknowledgements**

570 We applied the SDC (Sequence Determines Credit) approach for the sequence of authors. We
571 would like to thank the space agency NASA for providing the MODIS satellite images used in
572 this paper (<https://oceancolor.gsfc.nasa.gov/>). The authors are grateful to the whole team of the
573 SOMBA-2014 cruise for providing the in-situ data used in this work. We thank the three
574 anonymous reviewers for their helpful suggestions that greatly improved this manuscript. The
575 combination of three funding sources supported this research: a PhD scholarship from the
576 MESRS (Algerian government), a PhD scholarship from the Algerian-French program PROFAS
577 B+ 2018-2019 (MERS-Algeria) and a partnership project (IRD-French) Fellowship.

References

- Amores, A., Monserrat, S., Marcos, M., 2013. Vertical structure and temporal evolution of an anticyclonic eddy in the Balearic Sea (western Mediterranean): DESCRIPTION OF AN EDDY (BALEARIC SEA). *J. Geophys. Res. Oceans* 118, 2097–2106. <https://doi.org/10.1002/jgrc.20150>
- Antoine, D., Morel, A., André, J.-M., 1995. Algal pigment distribution and primary production in the eastern Mediterranean as derived from coastal zone color scanner observations. *J. Geophys. Res.* 100, 16193. <https://doi.org/10.1029/95JC00466>
- Barale, V., Garcia-Gorriz, E., Hoepffner, N., Stips, A., 2008. Near-Coastal Features of the NW Mediterranean Sea - Space and Time Heterogeneity of Atmospheric Forcing, Vertical Mixing and Algal Blooming, from Satellite Observations and Model Simulations (1997-2007). <https://doi.org/10.13140/RG.2.1.2425.9285>
- Béranger, K., Mortier, L., Crépon, M., 2005. Seasonal variability of water transport through the Straits of Gibraltar, Sicily and Corsica, derived from a high-resolution model of the Mediterranean circulation. *Progress in Oceanography, Mediterranean physical oceanography and biogeochemical cycles: Mediterranean general circulation and climate variability* 66, 341–364. <https://doi.org/10.1016/j.pocean.2004.07.013>

- Bethoux, J.P., Durieu de Madron, X., Nyffeler, F., Tailliez, D., 2002. Deep water in the western Mediterranean: peculiar 1999 and 2000 characteristics, shelf formation hypothesis, variability since 1970 and geochemical inferences. *Journal of Marine Systems* 33–34, 117–131. [https://doi.org/10.1016/S0924-7963\(02\)00055-6](https://doi.org/10.1016/S0924-7963(02)00055-6)
- Cao, L., Wang, W., Yang, Y., Yang, C., Yuan, Z., Xiong, S., Diana, J., 2007. Environmental Impact of Aquaculture and Countermeasures to Aquaculture Pollution in China. *Env Sci Pollut Res* 11. <http://dx.doi.org/10.1065/espr2007.05.426>
- Cerino, F., Fornasaro, D., Kralj, M., Giani, M., Cabrini, M., 2019. Phytoplankton temporal dynamics in the coastal waters of the north-eastern Adriatic Sea (Mediterranean Sea) from 2010 to 2017. *NC* 34, 343–372. <https://doi.org/10.3897/natureconservation.34.30720>
- Colella, S., Falcini, F., Rinaldi, E., Sammartino, M., Santoleri, R., 2016. Mediterranean Ocean Colour Chlorophyll Trends. *PLoS One* 11. <https://doi.org/10.1371/journal.pone.0155756>
- Conan, P., Testor, P., Estournel, C., D’Ortenzio, F., Durieu de Madron, X., 2018. Observing Winter Mixing and Spring Bloom in the Mediterranean. *Eos* 99. <https://doi.org/10.1029/2018EO105887>
- Cook, P.L.M., Røy, H., 2006. Advective relief of CO₂ limitation in microphytobenthos in highly productive sandy sediments. *Limnology and Oceanography* 51, 1594–1601. <https://doi.org/10.4319/lo.2006.51.4.1594>
- Crispi, G., Pacciaroni, M., 2009. Long-term numerical evolution of the nitrogen bulk content in the Mediterranean Sea. *Estuarine, Coastal and Shelf Science* 83, 148–158. <https://doi.org/10.1016/j.ecss.2007.12.015>
- Cullen, J.J., 1982. The Deep Chlorophyll Maximum: Comparing Vertical Profiles of Chlorophyll *a*. *Can. J. Fish. Aquat. Sci.* 39, 791–803. <https://doi.org/10.1139/f82-108>
- Demarcq, H., Barlow, R., Hutchings, L., 2007. Application of a chlorophyll index derived from satellite data to investigate the variability of phytoplankton in the Benguela ecosystem. *African Journal of Marine Science* 29, 271–282. <https://doi.org/10.2989/AJMS.2007.29.2.11.194>
- D’Ortenzio, F., Ribera d’Alcalà, M., 2009. On the trophic regimes of the Mediterranean Sea: a satellite analysis. *Biogeosciences* 139–148. <https://doi.org/10.5194/bg-6-139-2009>
- Elbaz-Poulichet, F., Guieu, C., Morley, N.H., 2001. A Reassessment of Trace Metal Budgets in the Western Mediterranean Sea. *Marine Pollution Bulletin* 42, 623–627. [https://doi.org/10.1016/S0025-326X\(01\)00065-0](https://doi.org/10.1016/S0025-326X(01)00065-0)
- El-Geziry, T.M., Bryden, I.G., 2010. The circulation pattern in the Mediterranean Sea: issues for modeller consideration. *Journal of Operational Oceanography* 3, 39–46. <https://doi.org/10.1080/1755876X.2010.11020116>
- Fani, F., Nuccio, C., Lazzara, L., Massi, L., Battocchi, C., Penna, A., 2014. *Fibrocapsa japonica* (Raphidophyceae) occurrence and ecological features within the phytoplankton assemblage of a cyclonic eddy, offshore the Eastern Alboran Sea. *Medit. Mar. Sci.* 15, 250. <https://doi.org/10.12681/mms.398>
- FAO, 2019. FAO Fisheries & Aquaculture - Vue générale du secteur aquacole national - Algérie. National Aquaculture Sector Overview Fact Sheets. Texte par Oussaid, M. Dans: Division des pêches de la FAO [en ligne]. http://www.fao.org/fishery/countrysector/naso_algeria/fr#tcN700B1 (accessed 8.24.21).
- Fernandez, V., Umlauf, L., Dobricic, S., Burchard, H., Nadia, P., 2006. Validation and intercomparison of two vertical-mixing schemes in the Mediterranean Sea. *Ocean Science Discussions* 3. <https://doi.org/10.5194/osd-3-1945-2006>

- Fondriest Environmental, 2014. Sediment Transport and Deposition. Environmental Measurement Systems. <https://www.fondriest.com/environmental-measurements/parameters/hydrology/sediment-transport-deposition/> (accessed 2.27.21).
- Font, J., Millot, C., Salas, J., Julià, A., Chic, O., 1998. The drift of Modified Atlantic Water from the Alboran Sea to the eastern Mediterranean. *SCIENTIA MARINA* 62, 6. <https://doi.org/10.3989/scimar.1998.62n3211>
- Google Maps, 2021. Algeria - Google Maps. <https://www.google.com/maps/place/Alger> (accessed 1.14.21).
- Groom, S., Sathyendranath, S., Ban, Y., Bernard, S., Brewin, R., Brotas, V., Brockmann, C., Chauhan, P., Choi, J., Chuprin, A., Ciavatta, S., Cipollini, P., Donlon, C., Franz, B., He, X., Hirata, T., Jackson, T., Kampel, M., Krasemann, H., Lavender, S., Pardo-Martinez, S., Mélin, F., Platt, T., Santoleri, R., Skakala, J., Schaeffer, B., Smith, M., Steinmetz, F., Valente, A., Wang, M., 2019. Satellite Ocean Colour: Current Status and Future Perspective. *Front. Mar. Sci.* 6, 485. <https://doi.org/10.3389/fmars.2019.00485>
- Harid, R., Ait Kaci, M., Keraghel, M.A., Zerrouki, M., Houma-Bachari, F., 2018. Seasonal and Interannual Variability of Primary Production and Chlorophyll Concentrations in the Algerian Basin: Application of Ocean Color, in: Kallel, A., Ksibi, M., Ben Dhia, H., Khélifi, N. (Eds.), *Recent Advances in Environmental Science from the Euro-Mediterranean and Surrounding Regions*, Advances in Science, Technology & Innovation. Springer International Publishing, Cham, pp. 1641–1643. https://doi.org/10.1007/978-3-319-70548-4_475
- Hassan, G.S., Espinosa, M.A., Isla, F.I., 2006. Modern Diatom Assemblages in Surface Sediments from Estuarine Systems in the Southeastern Buenos Aires Province, Argentina. *J Paleolimnol* 35, 39. <https://doi.org/10.1007/s10933-005-6444-8>
- Houpert, L., Testor, P., Durrieu de Madron, X., Somot, S., D’Ortenzio, F., Estournel, C., Lavigne, H., 2015. Seasonal cycle of the mixed layer, the seasonal thermocline and the upper-ocean heat storage rate in the Mediterranean Sea derived from observations. *Progress in Oceanography* 132, 333–352. <https://doi.org/10.1016/j.pocean.2014.11.004>
- Huertas, I.E., Ríos, A.F., García-Lafuente, J., Navarro, G., Makaoui, A., Sánchez-Román, A., Rodríguez-Galvez, S., Orbi, A., Ruiz, J., Pérez, F.F., 2012. Atlantic forcing of the Mediterranean oligotrophy: ATLANTIC FORCING OF MEDITERRANEAN OLIGOTROPHY. *Global Biogeochem. Cycles* 26, n/a-n/a. <https://doi.org/10.1029/2011GB004167>
- Huertas, I.E., Ríos, A.F., García-Lafuente, J., Makaoui, A., Rodríguez-Galvez, S., Sánchez-Román, A., Orbi, A., Ruiz, J., Pérez, F.F., 2009. Anthropogenic and natural CO₂ exchange through the Strait of Gibraltar. *Biogeosciences* 6, 647–662. <https://doi.org/10.5194/bg-6-647-2009>
- Keraghel, M.A., Louanchi, F., Zerrouki, M., Aït Kaci, M., Aït-Ameur, N., Labaste, M., Legoff, H., Taillandier, V., Harid, R., Mortier, L., 2020. Carbonate system properties and anthropogenic carbon inventory in the Algerian Basin during SOMBA cruise (2014): Acidification estimate. *Marine Chemistry* 221, 103783. <https://doi.org/10.1016/j.marchem.2020.103783>
- Kessouri, F., Ulses, C., Estournel, C., Marsaleix, P., D’Ortenzio, F., Severin, T., Taillandier, V., Conan, P., 2018. Vertical Mixing Effects on Phytoplankton Dynamics and Organic Carbon Export in the Western Mediterranean Sea. *J. Geophys. Res. Oceans* 123, 1647–1669. <https://doi.org/10.1002/2016JC012669>

- Khatiwala, S., Tanhua, T., Mikaloff Fletcher, S., Gerber, M., Doney, S.C., Graven, H.D., Gruber, N., McKinley, G.A., Murata, A., Ríos, A.F., Sabine, C.L., 2013. Global ocean storage of anthropogenic carbon. *Biogeosciences* 10, 2169–2191. <https://doi.org/10.5194/bg-10-2169-2013>
- Lavigne, H., D’Ortenzio, F., Ribera D’Alcalà, M., Claustre, H., Sauzède, R., Gacic, M., 2015. On the vertical distribution of the chlorophyll & concentration in the Mediterranean Sea: a basin scale and seasonal approach. *Biogeosciences Discuss.* 12, 4139–4181. <https://doi.org/10.5194/bgd-12-4139-2015>
- Lazzari, P., Solidoro, C., Ibello, V., Salon, S., Teruzzi, A., Béranger, K., Colella, S., Crise, A., 2012. Seasonal and inter-annual variability of plankton chlorophyll and primary production in the Mediterranean Sea: a modelling approach. *Biogeosciences* 9, 217–233. <https://doi.org/10.5194/bg-9-217-2012>
- Marañón, E., Van Wambeke, F., Uitz, J., Boss, E.S., Pérez-Lorenzo, M., Dinasquet, J., Haëntjens, N., Dimier, C., Taillandier, V., 2020. Deep maxima of phytoplankton biomass, primary production and bacterial production in the Mediterranean Sea during late spring (preprint). *Biodiversity and Ecosystem Function: Microbial Ecology & Geomicrobiology*. <https://doi.org/10.5194/bg-2020-261>
- Mayot, N., D’Ortenzio, F., Ribera d’Alcalà, M., Lavigne, H., Claustre, H., 2016. Interannual variability of the Mediterranean trophic regimes from ocean color satellites. *Biogeosciences* 13, 1901–1917. <https://doi.org/10.5194/bg-13-1901-2016>
- Millot, C., 1999. Circulation in the Western Mediterranean Sea. *Journal of Marine Systems* 20, 423–442. [https://doi.org/10.1016/S0924-7963\(98\)00078-5](https://doi.org/10.1016/S0924-7963(98)00078-5)
- Millot, C., 1989. La circulation générale en Méditerranée occidentale: Aperçu de nos connaissances et projets d’études. *geo* 98, 497–515. <https://doi.org/10.3406/geo.1989.20925>
- Millot, C., Taupier-Letage, I., 2005. Circulation in the Mediterranean Sea. *Hdb Env Chem* 5, 38. <https://doi.org/DOI 10.1007/b107143>
- Millot, C., Taupier-Letage, I., Benzohra, M., 1990. The Algerian eddies. *Earth-Science Reviews* 27, 203–219. [https://doi.org/10.1016/0012-8252\(90\)90003-E](https://doi.org/10.1016/0012-8252(90)90003-E)
- Morel, A., Prieur, L., 1977. Analysis of variations in ocean color1: Ocean color analysis. *Limnol. Oceanogr.* 22, 709–722. <https://doi.org/10.4319/lo.1977.22.4.0709>
- Mortier, L., Alt Aneur, N., Taillandier, V., 2014. SOMBA-GE-2014 cruise, Téthys II R/V. <https://doi.org/10.17600/14007500>
- Moutin, T., Prieur, L., 2012. Influence of anticyclonic eddies on the Biogeochemistry from the Oligotrophic to the Ultraoligotrophic Mediterranean (BOUM cruise). *Biogeosciences* 9, 3827–3855. <https://doi.org/10.5194/bg-9-3827-2012>
- NASA’s Ocean Color Web, 2019. Available online: <http://oceancolor.gsfc.nasa.gov/> (accessed on 29 October 2019).
- Okubo, A., 1973. Effect of shoreline irregularities on streamwise dispersion in estuaries and other embayments. *Netherlands Journal of Sea Research* 6, 213–224. [https://doi.org/10.1016/0077-7579\(73\)90014-8](https://doi.org/10.1016/0077-7579(73)90014-8)
- Olita, A., Sorgente, R., Ribotti, A., Fazioli, L., Perilli, A., 2011. Pelagic primary production in the Algero-Provençal Basin by means of multisensor satellite data: focus on interannual variability and its drivers. *Ocean Dynamics* 61, 1005–1016. <https://doi.org/10.1007/s10236-011-0405-8>

- O'Reilly, J.E., Werdell, P.J., 2019. Chlorophyll algorithms for ocean color sensors - OC4, OC5 & OC6. *Remote Sensing of Environment* 229, 32–47. <https://doi.org/10.1016/j.rse.2019.04.021>
- Ounissi, M., Ziouch, O.-R., Aounallah, O., 2014. Variability of the dissolved nutrient (N, P, Si) concentrations in the Bay of Annaba in relation to the inputs of the Seybouse and Mafragh estuaries. *Marine Pollution Bulletin* 80, 234–244. <https://doi.org/10.1016/j.marpolbul.2013.12.030>
- Palmiéri, J., Dutay, J.-C., D'Ortenzio, F., Houpert, L., Mayot, N., Bopp, L., 2018. The Mediterranean subsurface phytoplankton dynamics and their impact on Mediterranean bioregions. *Biogeosciences Discussions* 1–38. <https://doi.org/10.5194/bg-2018-423>
- Pasqueron de Fommervault, O., D'Ortenzio, F., Mangin, A., Serra, R., Migon, C., Claustre, H., Lavigne, H., Ribera d'Alcalà, M., Prieur, L., Taillandier, V., Schmechtig, C., Poteau, A., Leymarie, E., Dufour, A., Besson, F., Obolensky, G., 2015. Seasonal variability of nutrient concentrations in the Mediterranean Sea: Contribution of Bio-Argo floats. *J. Geophys. Res. Oceans* 120, 8528–8550. <https://doi.org/10.1002/2015JC011103>
- Peliz, Á., Teles-Machado, A., Marchesiello, P., Dubert, J., Lafuente, J.G., 2009. Filament generation off the Strait of Gibraltar in response to Gap winds. *Dynamics of Atmospheres and Oceans* 46, 36–45. <https://doi.org/10.1016/j.dynatmoce.2008.08.002>
- Pessini, F., Cotroneo, Y., Olita, A., Sorgente, R., Ribotti, A., Jendersie, S., Perilli, A., 2020. Life history of an anticyclonic eddy in the Algerian basin from altimetry data, tracking algorithm and in situ observations. *Journal of Marine Systems* 207, 103346. <https://doi.org/10.1016/j.jmarsys.2020.103346>
- Pessini, F., Olita, A., Cotroneo, Y., Perilli, A., 2018. Mesoscale eddies in the Algerian Basin: do they differ as a function of their formation site? *Ocean Sci.* 14, 669–688. <https://doi.org/10.5194/os-14-669-2018>
- Pieri, M., Massi, L., Lazzara, L., Nuccio, C., Lapucci, C., Maselli, F., 2015. Assessment of three algorithms for the operational estimation of [CHL] from MODIS data in the Western Mediterranean Sea. *European Journal of Remote Sensing* 48, 383–401. <https://doi.org/10.5721/EuJRS20154822>
- Pingree, R.D., Maddock, L., 1979. The tidal physics of headland flows and offshore tidal bank formation. *Marine Geology* 32, 269–289. [https://doi.org/10.1016/0025-3227\(79\)90068-9](https://doi.org/10.1016/0025-3227(79)90068-9)
- Rabehi, W., Guerfi, M., Mahi, H., 2019. La baie d'Alger, un espace côtier prisé, entre pressions d'urbanisation et gouvernance territoriale. *Geo-Eco-Marina* 25, 113–130. <https://doi.org/10.5281/ZENODO.3609744>
- Robinson, A.R., Hill, M.N. (Eds.), 2005. *Biological-physical interactions in the sea*, 1. Harvard Univ. Press publication 2005. ed, *The sea The global coastal ocean*. Harvard Univ. Press, Cambridge, Mass.
- Salgado-Hernanz, P.M., Racault, M.-F., Font-Muñoz, J.S., Basterretxea, G., 2019. Trends in phytoplankton phenology in the Mediterranean Sea based on ocean-colour remote sensing. *Remote Sensing of Environment* 221, 50–64. <https://doi.org/10.1016/j.rse.2018.10.036>
- Signell, R.P., Geyer, W.R., 1991. Transient eddy formation around headlands. *Journal of Geophysical Research: Oceans* 96, 2561–2575. <https://doi.org/10.1029/90JC02029>
- Sobel, I., 1990. An Isotropic 3x3 Gradient Operator, *Machine Vision for Three Dimensional Scenes*, Freeman, H. Academic Press 376–379.

- Strickland, J., 1965. Production of organic matter in primary stages of the marine food chain, in: Chemical Oceanography, Academic Press, London, edited by: J. P. Riley and G. Skirrow. Chemical Oceanography, Academic Press 477–610.
- Taillandier, V., Prieur, L., D’Ortenzio, F., Ribera d’Alcalà, M., Pulido-Villena, E., 2020. Profiling float observation of thermohaline staircases in the western Mediterranean Sea and impact on nutrient fluxes (preprint). Biogeochemistry: Open Ocean. <https://doi.org/10.5194/bg-2019-504>
- Taupier-Letage, I., 2003. Biological response to mesoscale eddies in the Algerian Basin. J. Geophys. Res. 108, 3245. <https://doi.org/10.1029/1999JC000117>
- Taupier-Letage, Millot, C., 1988. Surface circulation in the Algerian basin during 1984. Oceanologia Acta 9, 79–85. <https://archimer.ifremer.fr/doc/00267/37811/>
- Testor, P., Send, U., Gascard, J.-C., Millot, C., Taupier-Letage, I., Béranger, K., 2005. The mean circulation of the southwestern Mediterranean Sea: Algerian Gyres. Journal of Geophysical Research: Oceans 110. <https://doi.org/10.1029/2004JC002861>
- Vargas-Yáñez, M., Zunino, P., Benali, A., Delpy, M., Pastre, F., Moya, F., García-Martínez, M. del C., Tel, E., 2010. How much is the western Mediterranean really warming and salting? Journal of Geophysical Research: Oceans 115. <https://doi.org/10.1029/2009JC005816>
- Volpe, G., 2012. Seasonal to interannual phytoplankton response to physical processes in the Mediterranean Sea from satellite observations. Remote Sensing of Environment 13. <https://doi.org/10.1016/j.rse.2011.09.020>
- Volpe, G., Colella, S., Brando, V., Forneris, V., La Padula, F., Di Cicco, A., Sammartino, M., Bracaglia, M., Artuso, F., Santoleri, R., 2018. The Mediterranean Ocean Colour Level 3 Operational Multi-Sensor Processing (preprint). Remote Sensing/Biological Processes/Surface/Mediterranean Sea. <https://doi.org/10.5194/os-2018-73>
- Wolanski, E., Hamner, W.M., 1988. Topographically Controlled Fronts in the Ocean and Their Biological Influence. Science 241, 177–181. <https://doi.org/10.1126/science.241.4862.177>
- Ziouch, O.R., Laskri, H., Chenaker, H., Ledjedel, N.E., Daifallah, T., Ounissi, M., 2020. Transport of nutrients from the Seybouse River to Annaba Bay (Algeria, SW Mediterranean). Marine Pollution Bulletin 156, 111231. <https://doi.org/10.1016/j.marpolbul.2020.111231>

578 **Figure captions**

579 **Figure 1.** March climatological average of the Chl-*a* concentration (mg m^{-3}) between 2003 and
 580 2018 in the Algerian Basin (Mediterranean Sea). The average current velocity (m s^{-1}) for the
 581 same period (black arrows), the 800 m isobath (red line), and the 0.5 mg m^{-3} isopleth (dark green
 582 line) is superimposed. The stations of the SOMBA-2014 cruise are also superimposed (white
 583 dots). The large eddies in black (Alg. WG and Alg. EG) are deduced from the average sea level
 584 anomaly between 2003 and 2018. The average position of the Alboran eddies (Alb. WG and Alb.
 585 EG) are added.

586 **Figure 2.** Comparison of Chl-*a* concentration for the first fortnight of January (2003-2018
587 climatology) from (a) MODIS Level-3 data at 4-km resolution and (b) MODIS Level-2
588 (uncorrected data) at 1-km resolution.

589 **Figure 3.** Definition of the outlier pixels criteria for improving the cloud masking (example of
590 07 January 2014). (a) Local (3x3 matrix) Sobel gradient ($\text{mg m}^{-3} \text{ km}^{-1}$). (b) Distance (D1) from
591 the cloud borders, in km (clouds are in white). (c) Distance (D2) from the shoreline, in km. (d to
592 f): Pixels identified as cloudy from, (d) the gradient criteria, (e) the temporal variation (TV)
593 criteria, (f) from isolated pixels within cloud (IP) criteria. The result of the combination of all
594 three criteria is shown in (h) versus the original MODIS Level-2 data (g).

595 **Figure 4.** Linear relationship (red line) (a) between MODIS satellite corrected data (Y axis) and
596 in-situ Chl-*a* data (X axis) during the SOMBA cruise (14 August to 10 September 2014) in the
597 Algerian Basin. (b), (c), (d), and (e): spatial variability of the daily Chl-*a* field associated with
598 four selected stations. The stations represented with a grey cross in (c) and (d) were removed
599 from the comparison on the basis of a gradient threshold criteria or because their proximity to
600 clouds.

601 **Figure 5.** Results of the elimination of outlier pixels from the original uncorrected MODIS
602 Level-2 data (left column), respectively for a fortnightly average (example of 1-15/01/2014) (a
603 and b), and for a monthly climatological average (example of January 2003-2018) (c and d).

604 **Figure 6.** Hovmöller diagrams of the cross-shore seasonality (from the coast line to 120 km
605 offshore) of the Chl-*a* concentration in the Algerian Basin averaged between 2.2°W and 8.7°E,
606 from fortnightly climatologies (2003-2018) computed from (a) MODIS Level-3 data at 4-km
607 resolution, (b) MODIS Level-2 corrected data at 1-km resolution. The black line separates the 10
608 km distance from the coast. (c) and (d) highlight the data for this coastal area. The average cross-
609 shore profiles in March (green line) and August (blue line) are shown in (e) and (f).

610 **Figure 7.** Average seasonality of the Chl-*a* concentration from 2003 to 2018 in the Algerian
611 Basin from fortnightly Chl-*a* averages (a) in the 0-10 km coastal sector (green plain line) and
612 offshore (blue plain line). The corresponding 4-km resolution Chl-*a* MODIS data (dotted lines)
613 as well as the depth of the mixed Layer (MLD, orange dasher line) are superimposed. (b)
614 Zonally averaged cross-shore Chl-*a* transect from the coast to 50 km for the High-Biomass
615 (green), the transition (yellow), and the Low-Biomass (blue) seasons. The horizontal line shows
616 the 0.5 mg m⁻³ limit used to compute the integrated index, detailed in Fig. 8 and Fig. 9.

617 **Figure 8.** Time-space diagrams of the longitudinal variability (2.2°W to 8.7°E) of three Chl-*a*
618 associated indices in the Algerian Basin from August to July. The four distances considered (a)
619 were: the distance of 10 km from the coast (green line), the average maximum distance of the 0.5
620 mg m⁻³ isopleth, the $Dist_{0.5mg}^{max}$ (red line), the 80 km maximum distance allowed for I_B (gray solid
621 line), and the middle distance from the northern coastlines (gray dashed line). (b) the distance
622 from the coast of the Chl-*a* concentration ≥ 0.5 mg m⁻³. (c) the Chl-*a* averaged over the same
623 area. (d) the integrated biomass index (IB) from whole basin.

624 **Figure 9.** Time-space diagrams of the alongshore variability (2.2°W 8.7°E) of the Chl-*a* and
625 biomass index I_B: (a) the Chl-*a* averaged in the coastal area (0-10 km) and (b) in the offshore
626 area (10- $Dist_{0.5mg}^{max}$ km), the biomass index (I_B) integrated from (c) the coastline to a maximum
627 distance of 10 km, and (d) from 10 km to the maximum distance $Dist_{0.5mg}^{max}$.

628 **Figure 10.** Average seasonality of the integrated biomass index (orange bars) at three locations :
629 1°W (a), 4.5°E (b), and 7.5° (c) situated outside coastal influences (see Fig. 8a for precise
630 locations). The climatological values of the Mixed Layer Depth (MLD, blue line), and the Total
631 Kinetic Energy (TKE, black line) at the same locations are superimposed.

632 **Figure 11.** Longitudinal variability of the offshore (blue line) and coastal (green line) yearly
633 averages of the integrated biomass index, with the “I_B coastal / offshore” ratio superimposed (red

634 line). Areas with high coastal biomass index are highlighted with green rectangles (1-15) by
635 comparison with Low-Biomass coastal Chl-*a* areas (1-16) left blank. The wadis discharges, the
636 coast type (sandy coast in yellow and rocky coast in blue), and the size of nearby cities (red dots)
637 are superimposed. The coast type is deduced from the “Google Map” images (Google Maps,
638 2021). The wadis names are respectively : (1) Seybouse Wadi, (2) El-Kebir Wadi, (3) Z'Hor
639 Wadi, (4) El-Kebir Wadi, (5) Soummam Wadi, (6) Bou-Douaou Wadi, (7) El-Hamiz Wadi, (8)
640 El-Harrach Wadi, (9) Mazafran Wadi, (10) Cheliff Wadi, (11) El-Hammam Wadi.

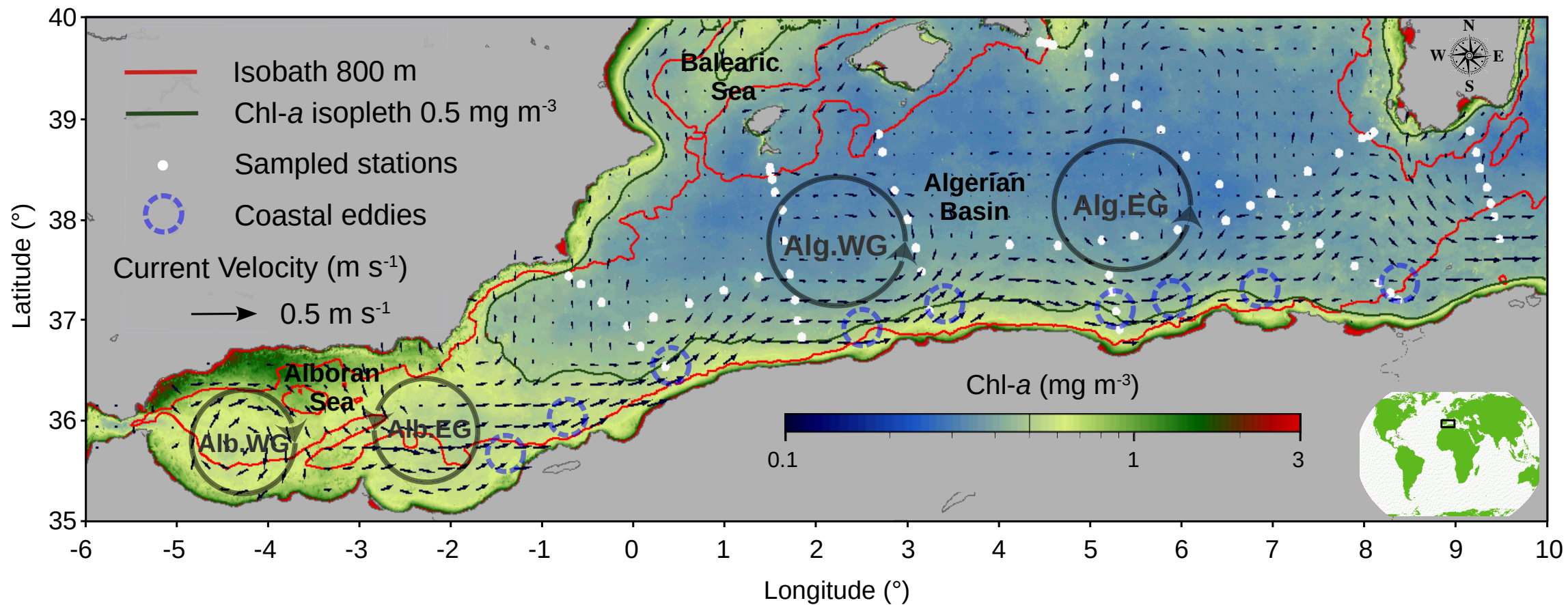
641 **Figure 12.** Seasonal variability of the Mazafran and the El-harrach wadis flows (in $\text{m}^3 \text{s}^{-1}$, orange
642 line) and the corresponding Chl-*a* concentration (green line) averaged at the isobath <50 m from
643 2003 to 2012 : (a) in the Bou-Ismaïl bay and (b) in the Algiers bay. The maps (a' and b') show
644 the Chl-*a* yearly average at each location.

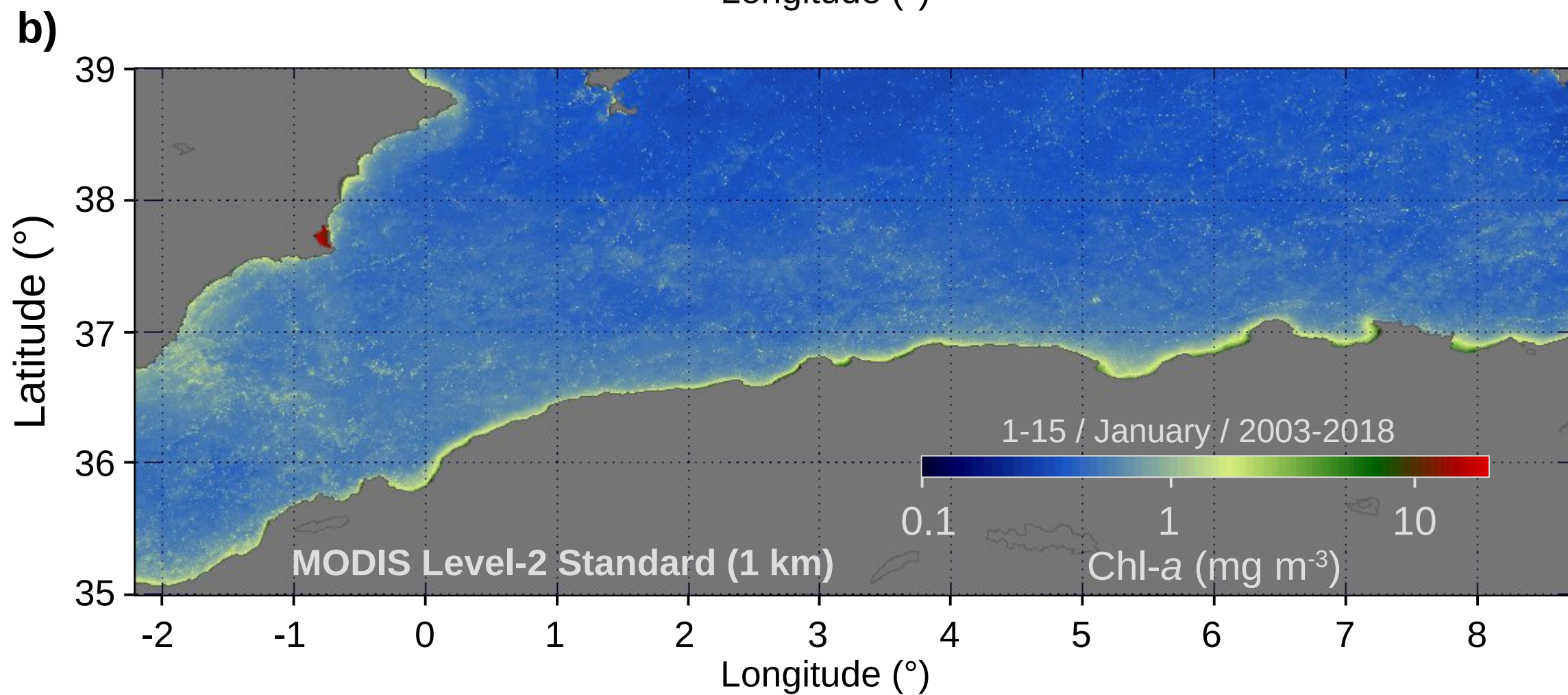
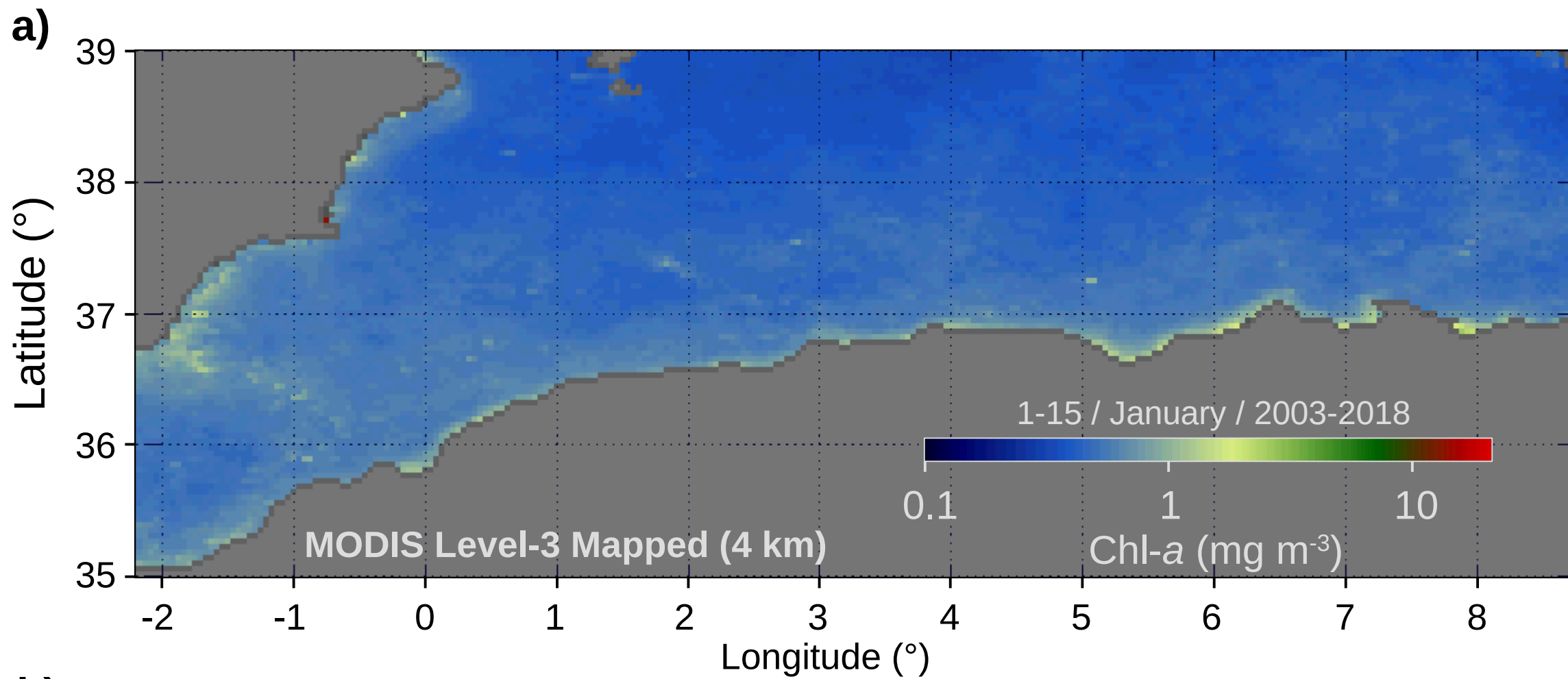
645 **Table 1.** Spatially integrated biomass index (I_B , in g m^{-2}) seasonally cumulated between 2003
646 and 2018 for regions of high and low biomass (as displayed in Fig. 11) for the coastal and
647 offshore domains. The last line shows the relative importance of the High vs Low I_B index and
648 the right part of the table shows the ratio between the coastal and offshore domains for both
649 types of regions.

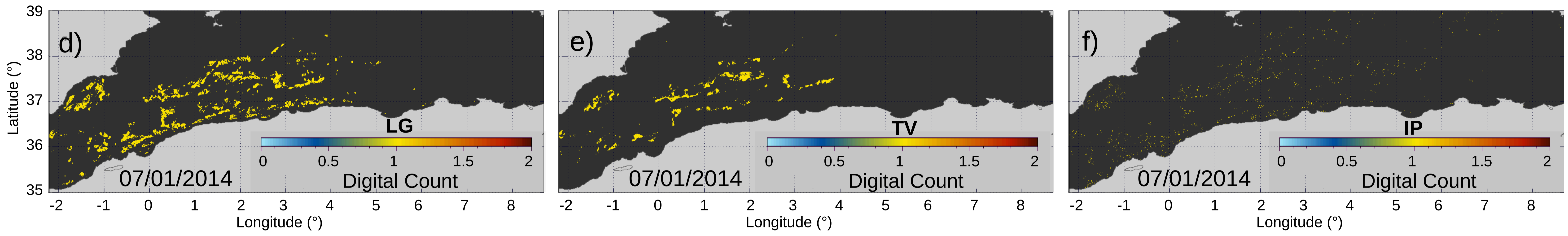
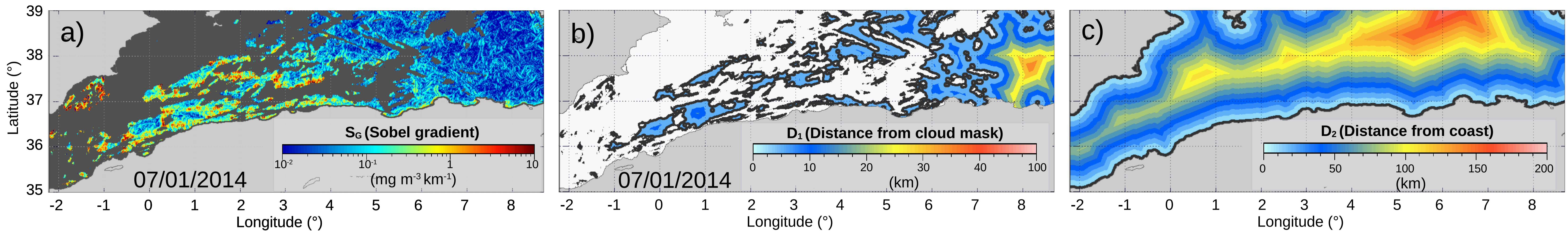
650 **Table 2.** Summary of the characteristics of the LBC (numbered 1-16) and HBC (numbered 1-
651 15) regions as manually selected in Fig. 11. Both Chl-*a* and I_B variables were averaged for each
652 LBC (white rectangles, 1-16) and HBC (green rectangles, 1-15) of Fig. 11. The City size [0-3] is
653 defined by respectively: 0 (no city), 1: [0 50k] inhabitants (small red dot), 2 [50k- 100k]
654 (intermediate red dot), 3: >100k (large red dot). The Coast-type is either Sandy(S) or Rocky(R).
655 The small black dots represent aquaculture cages.

656 **Table 3.** Parameters of the linear models calculated to evaluate the interactions between the
657 coastal Chl-*a* biomass or the I_B index and the four aforementioned variables of different of

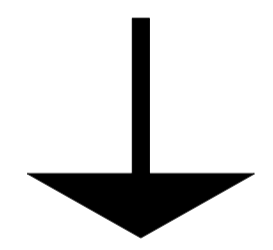
658 coastal enrichment sources. The averages of I_B and Chl-*a* in winter and summer are calculated
659 according to the High and the Low-Biomass months shown in Fig. 7a.



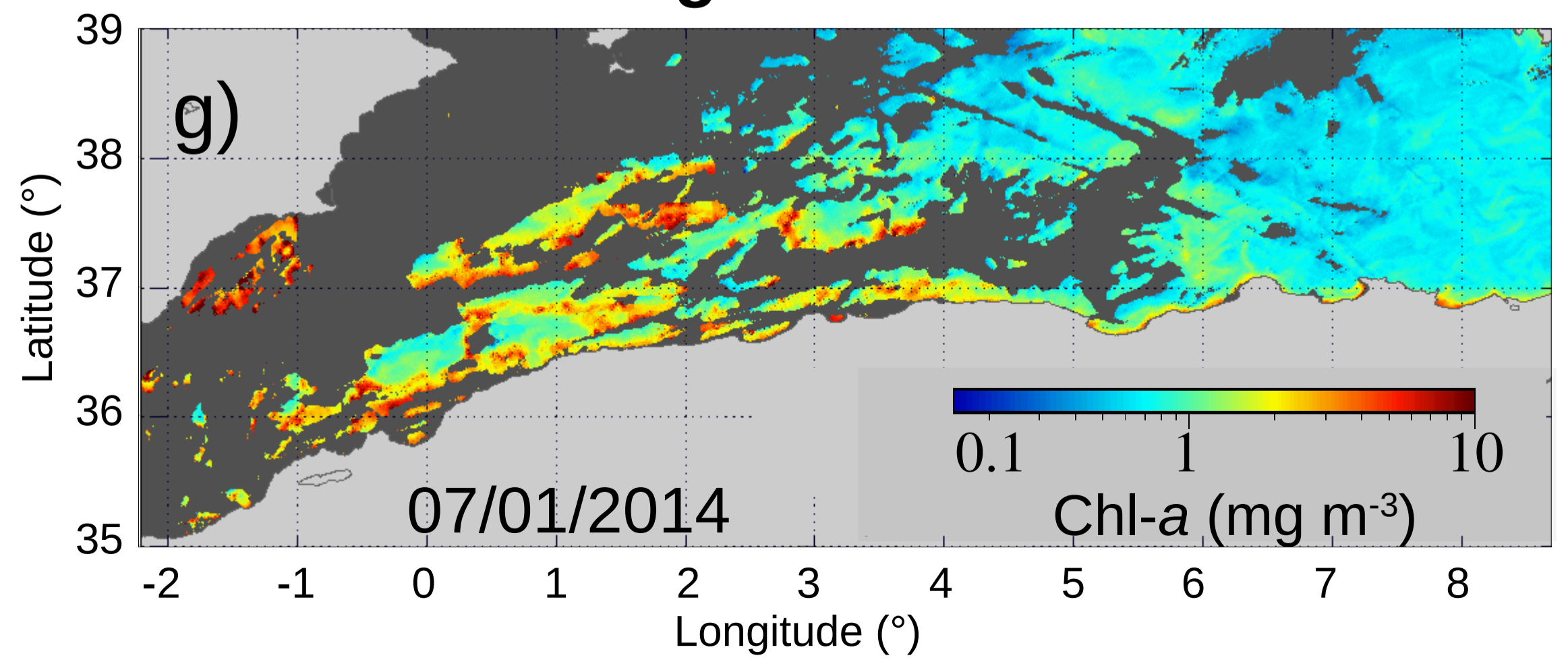




Combination

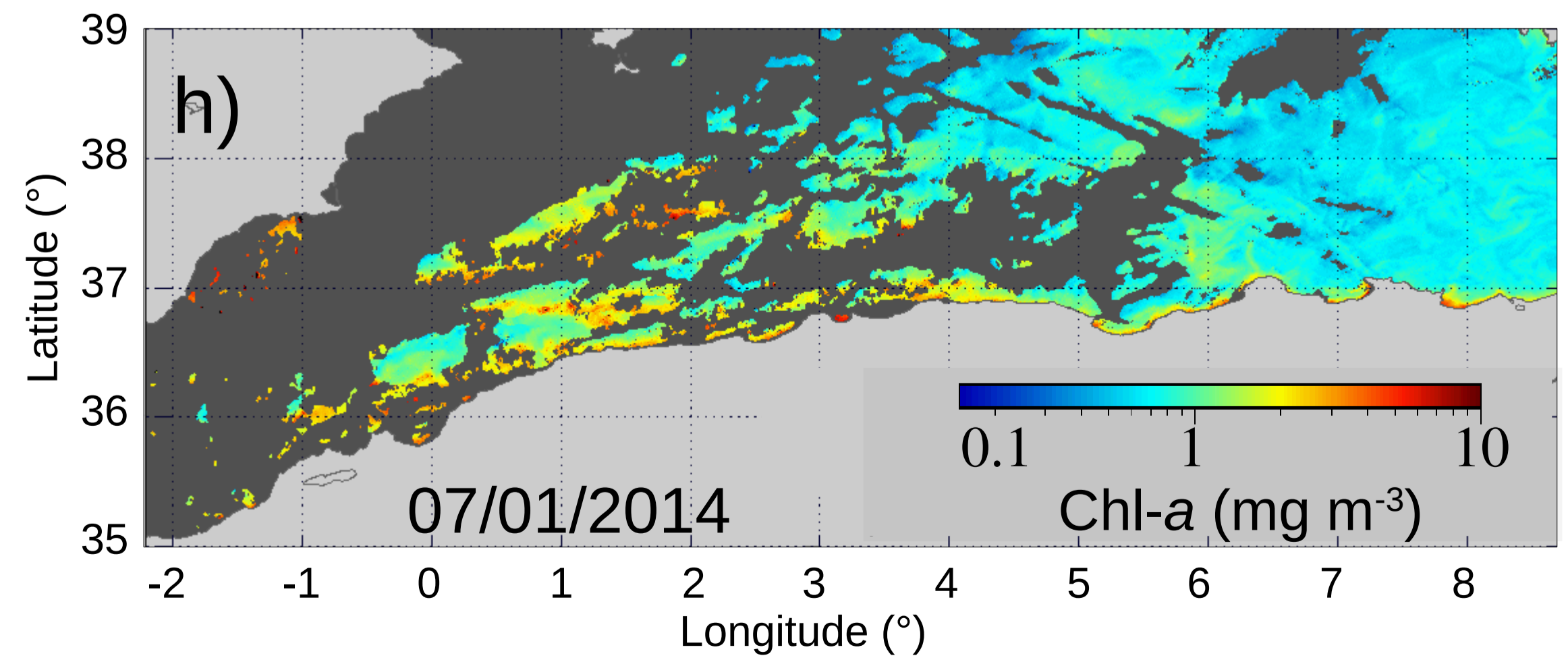


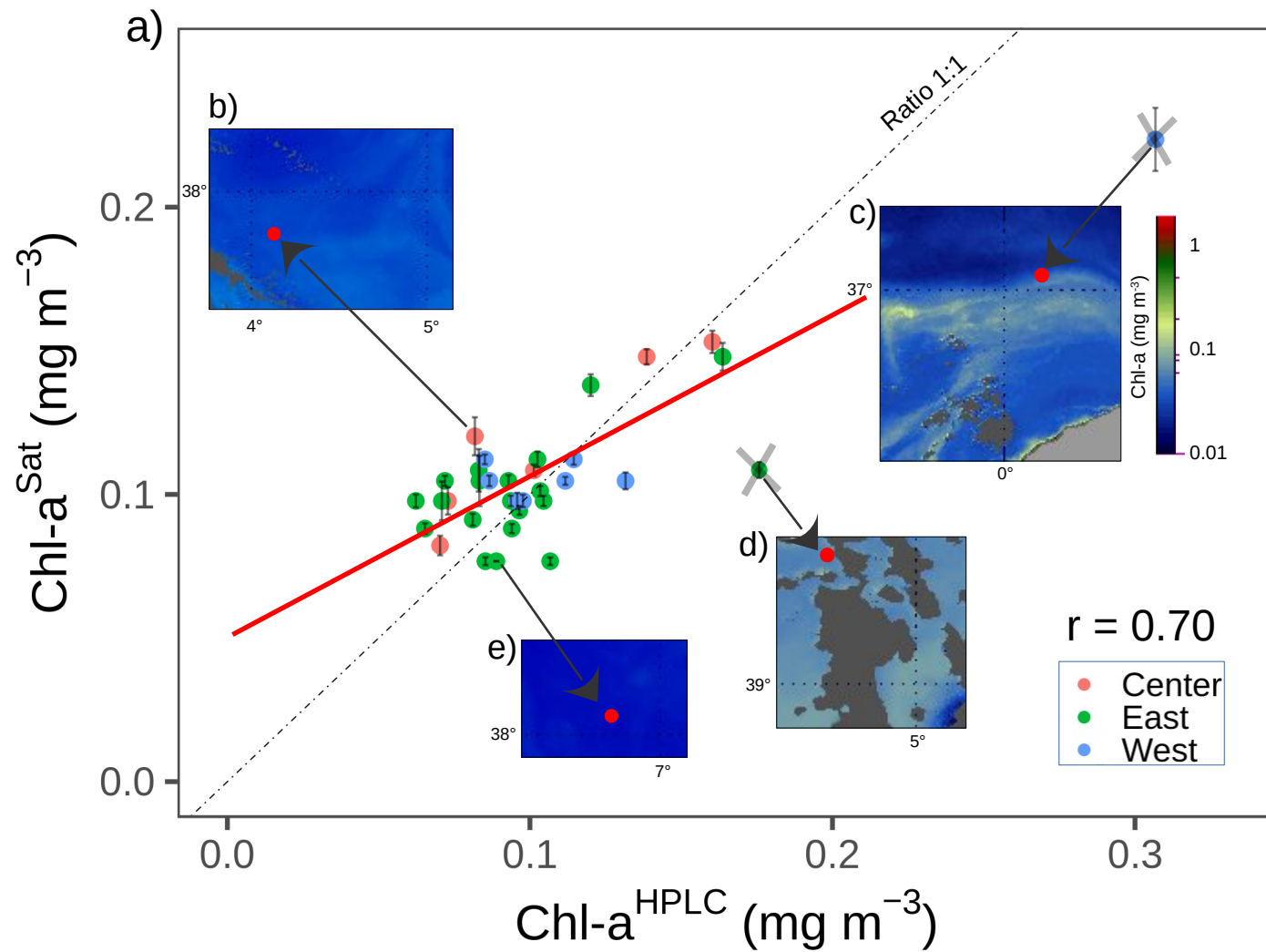
Original data



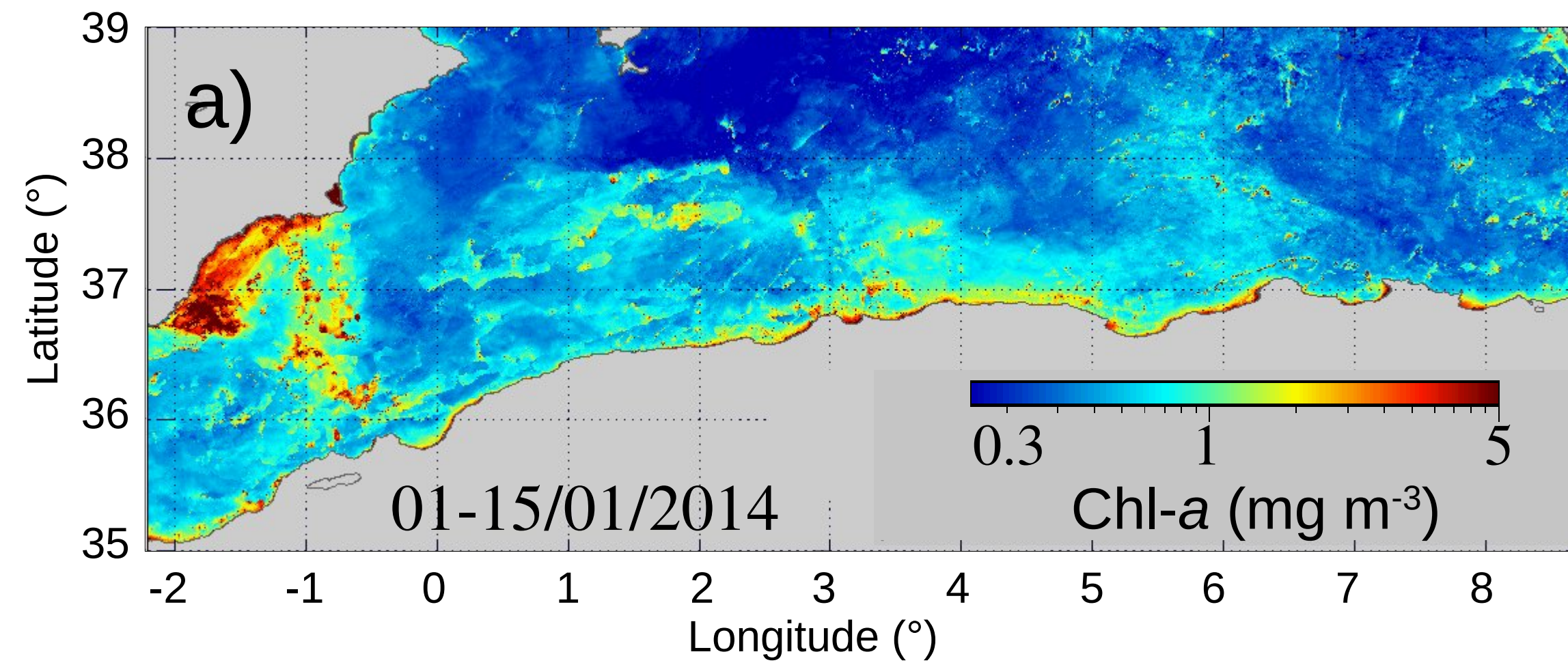
1 day

Corrected data

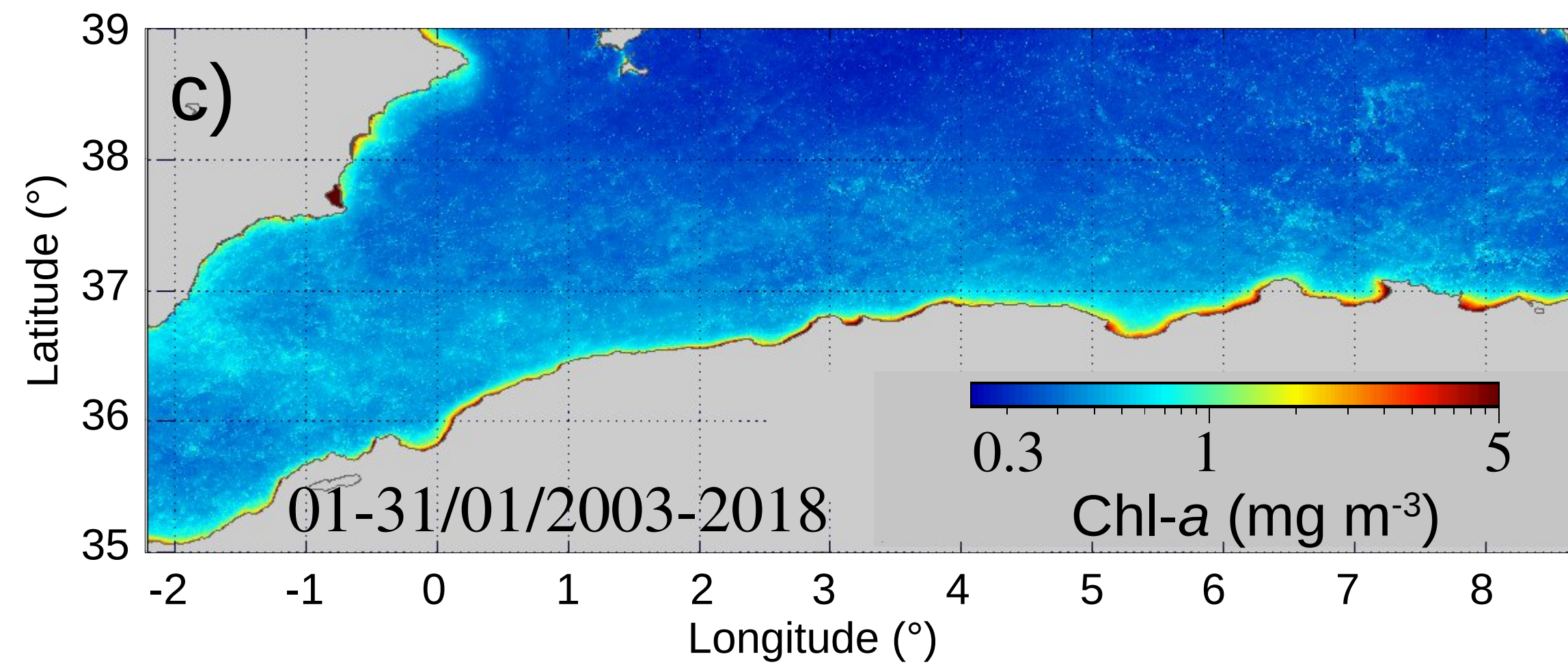
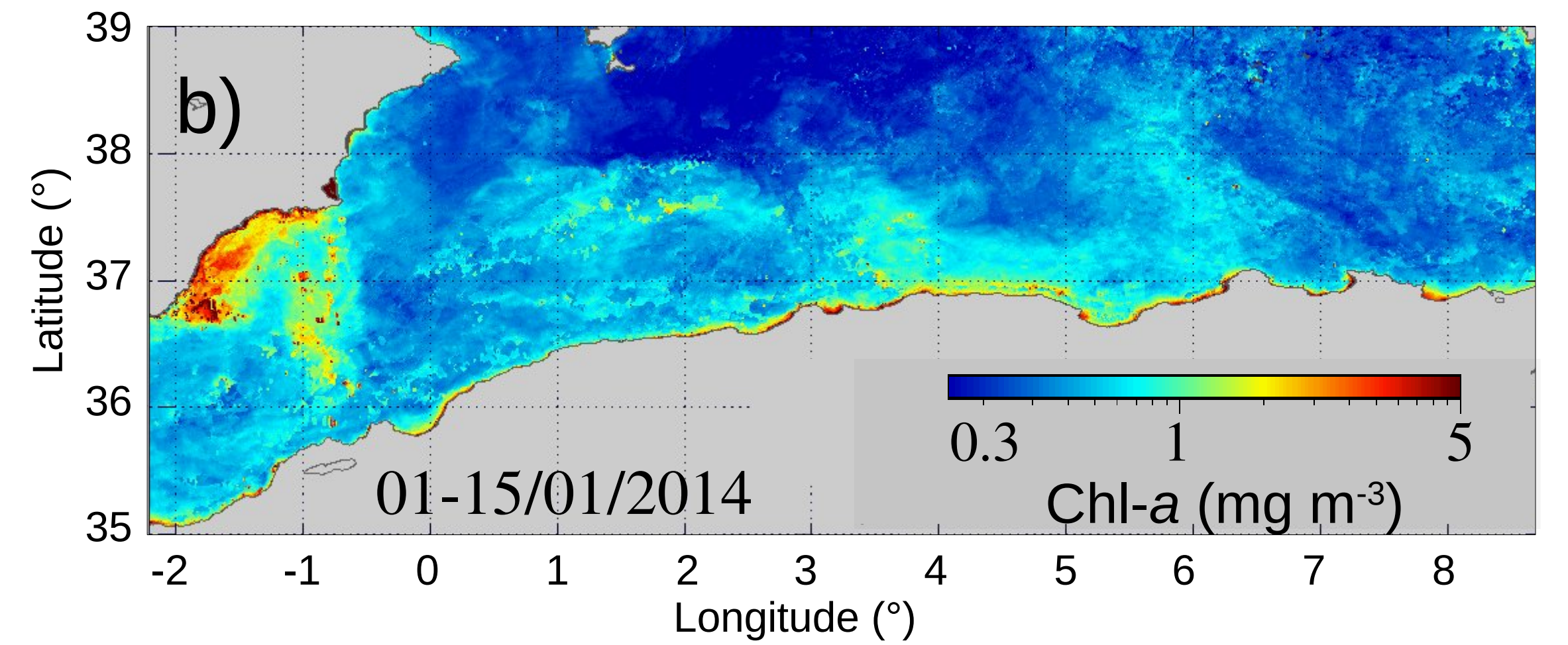




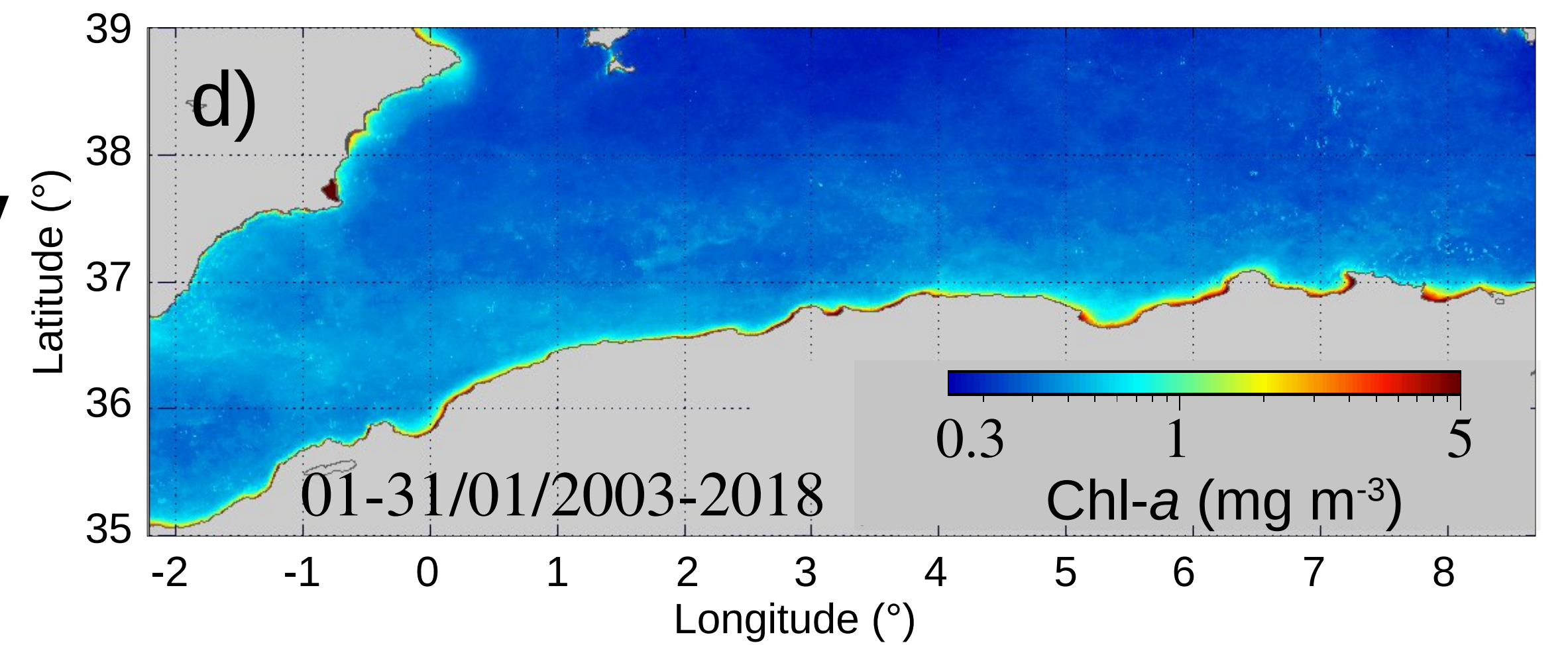
Original data



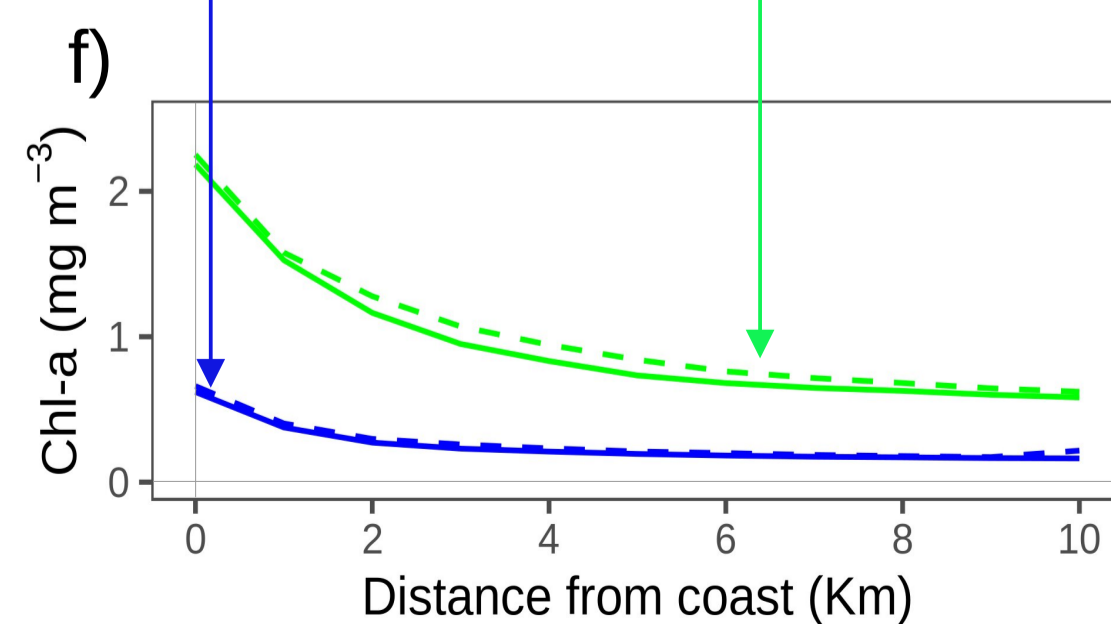
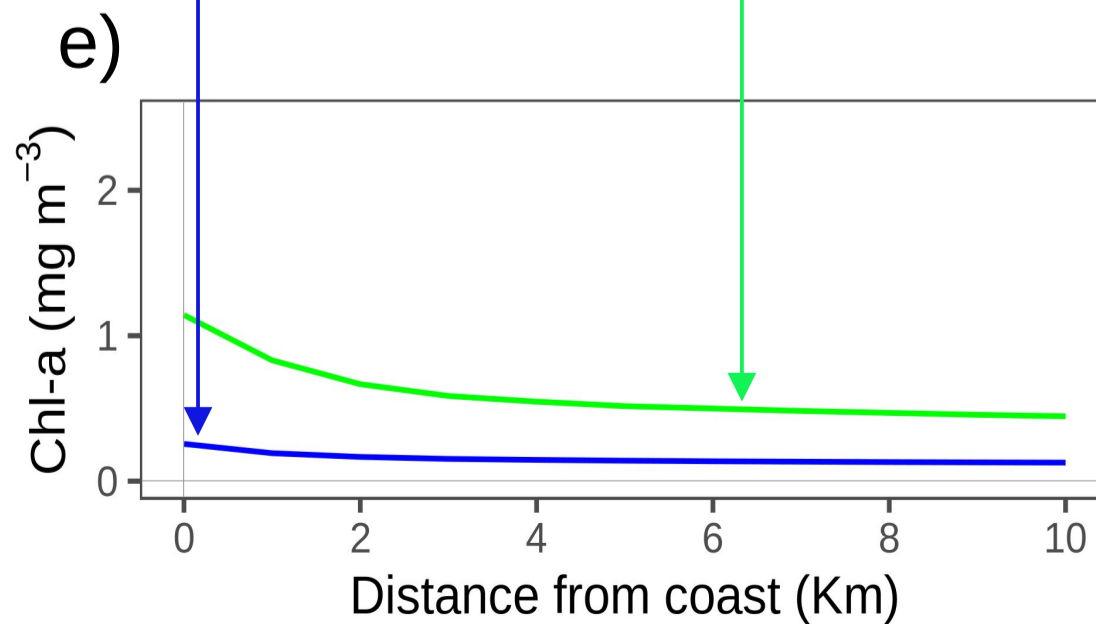
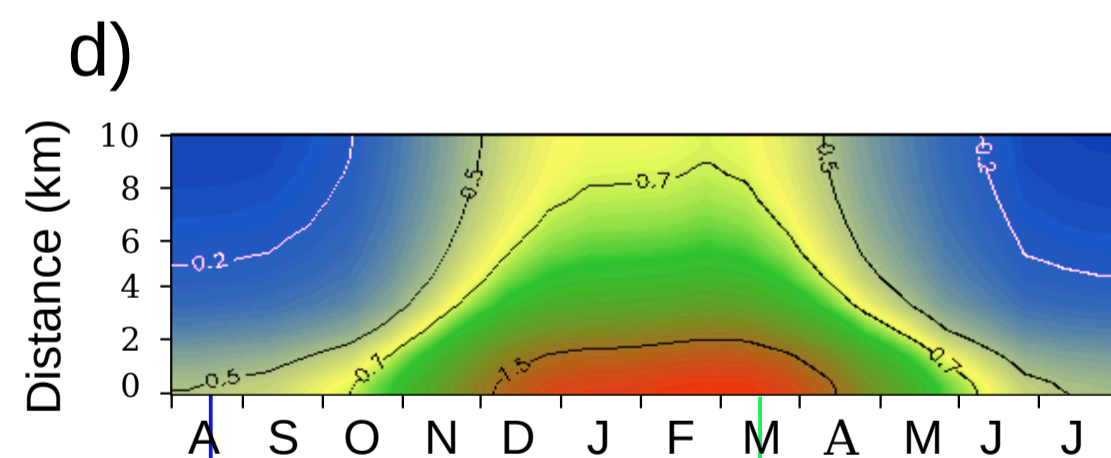
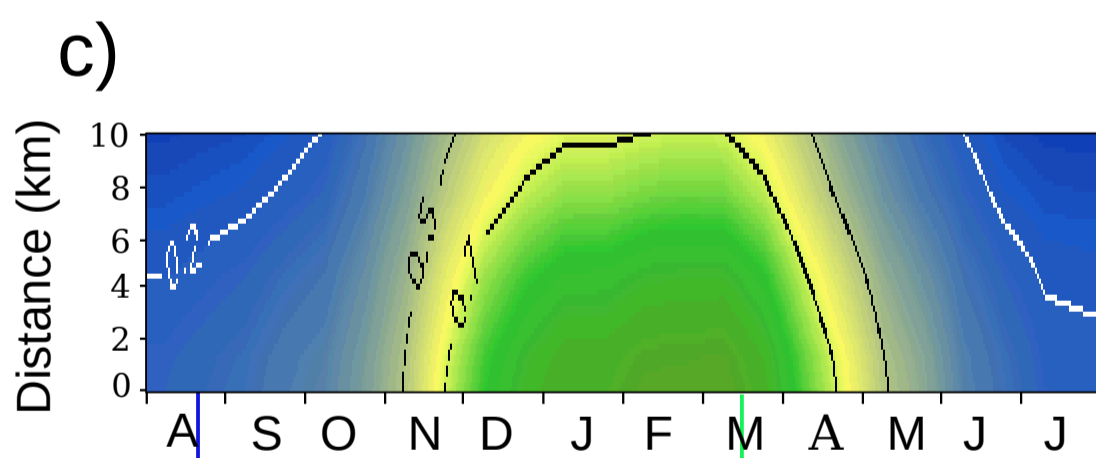
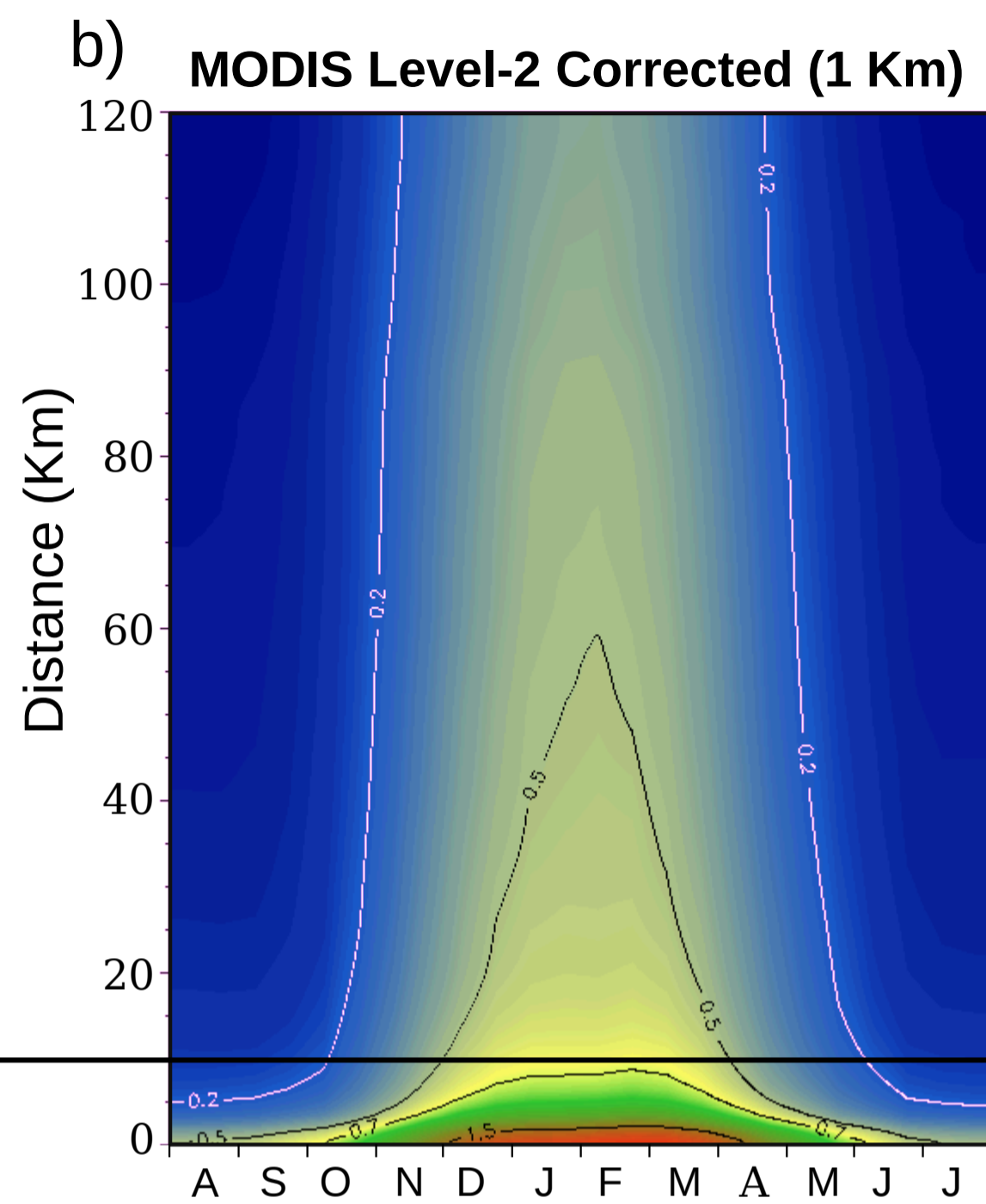
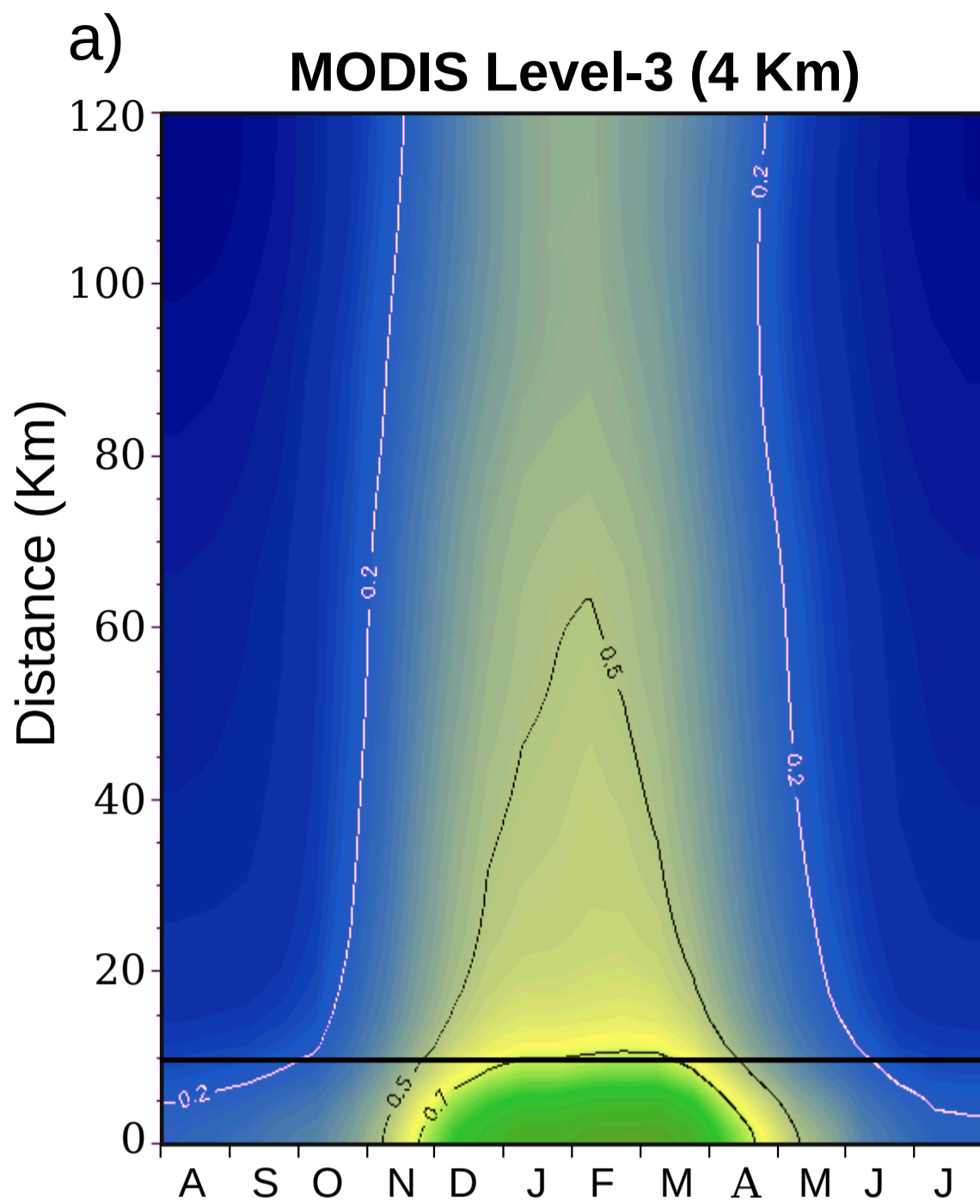
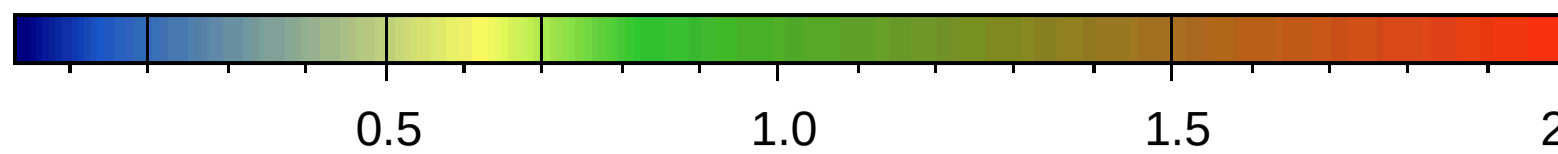
Fortnightly average

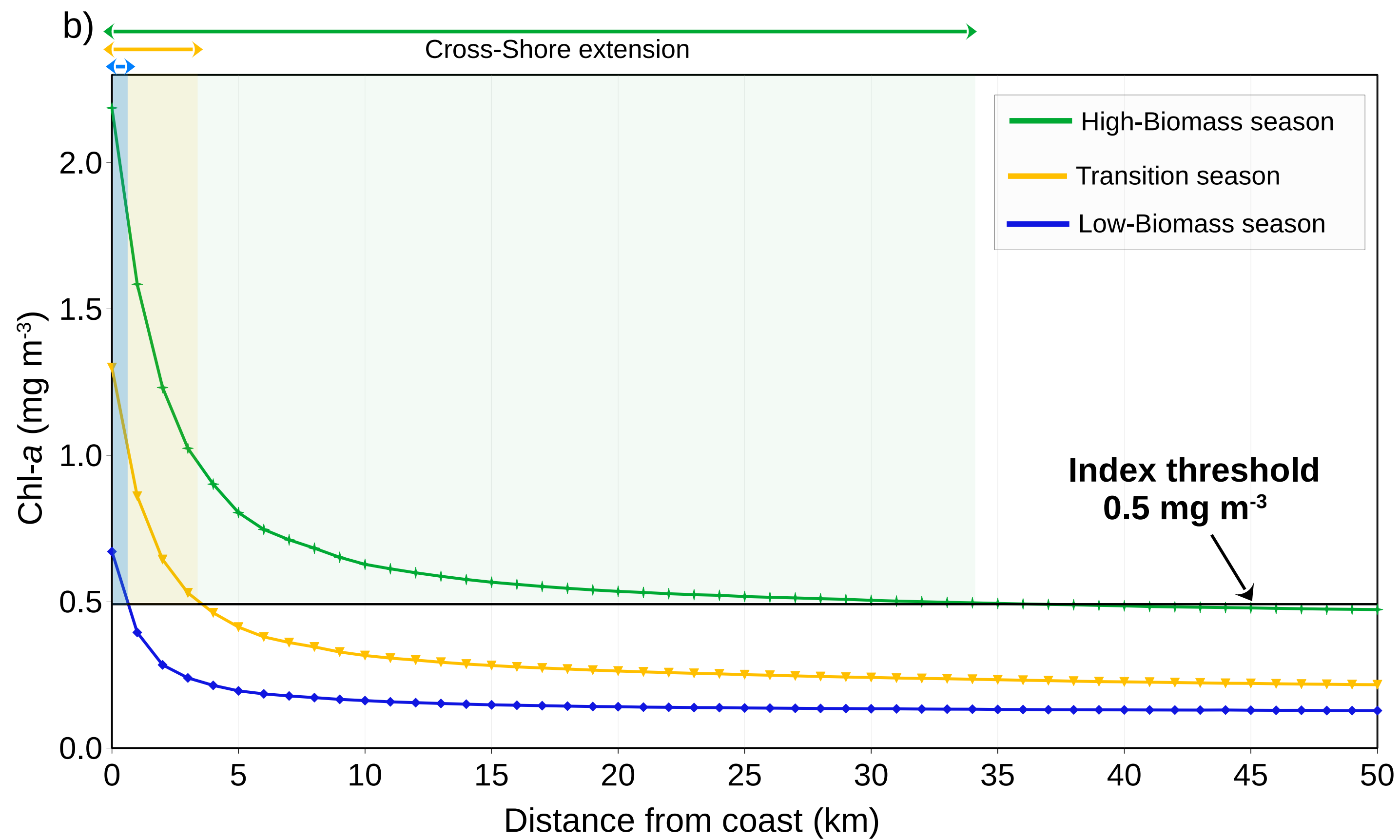
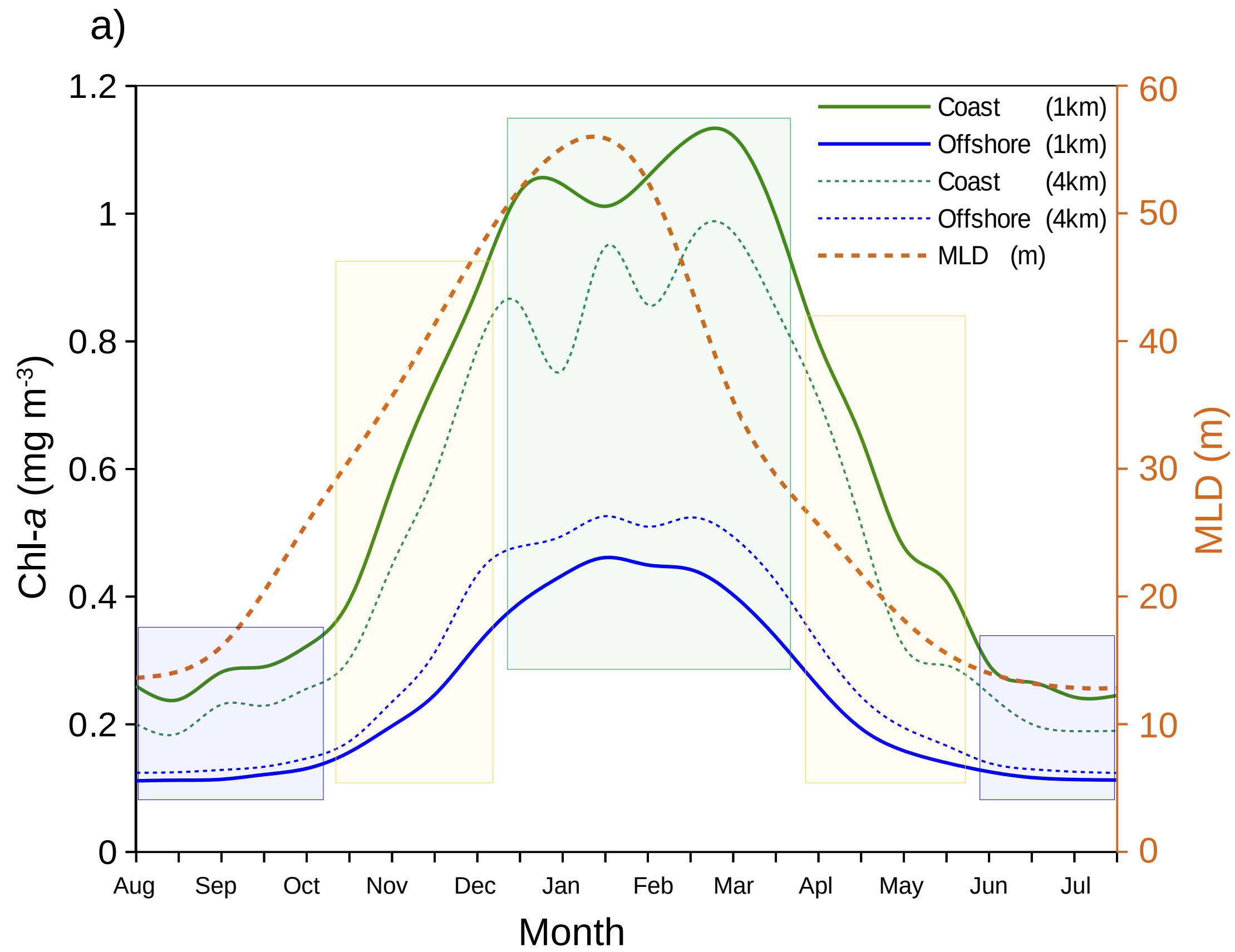


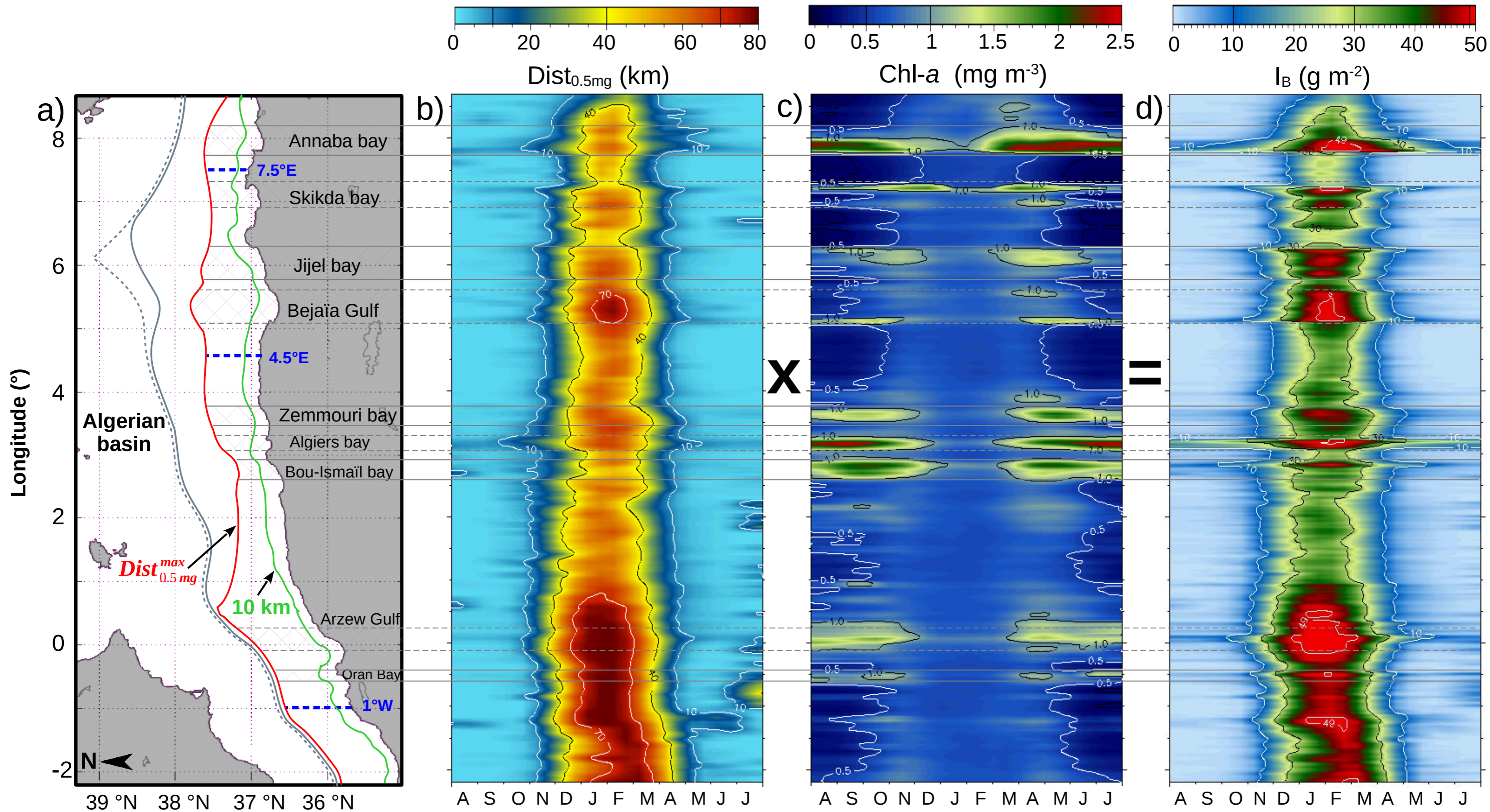
Monthly climatology

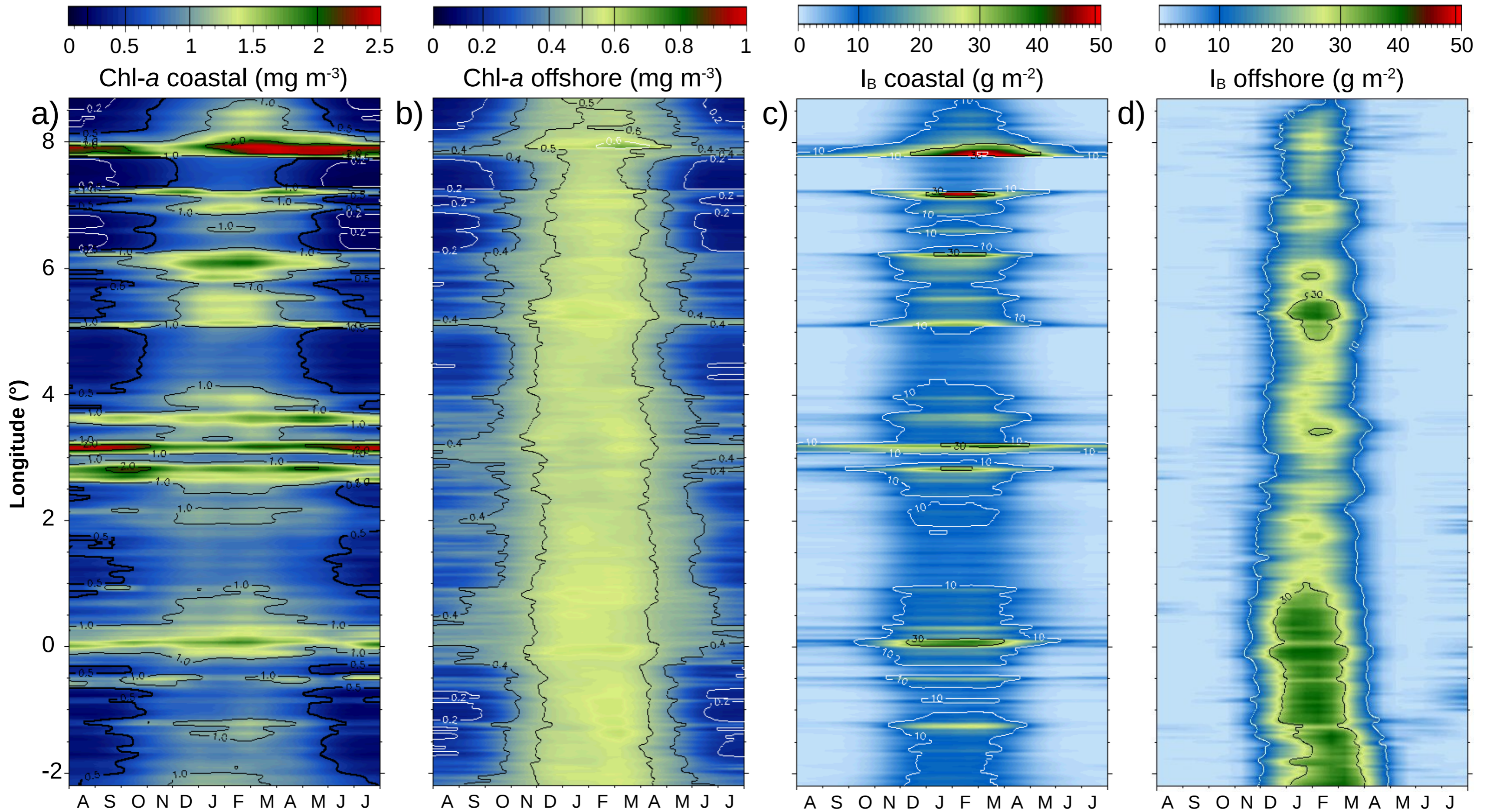


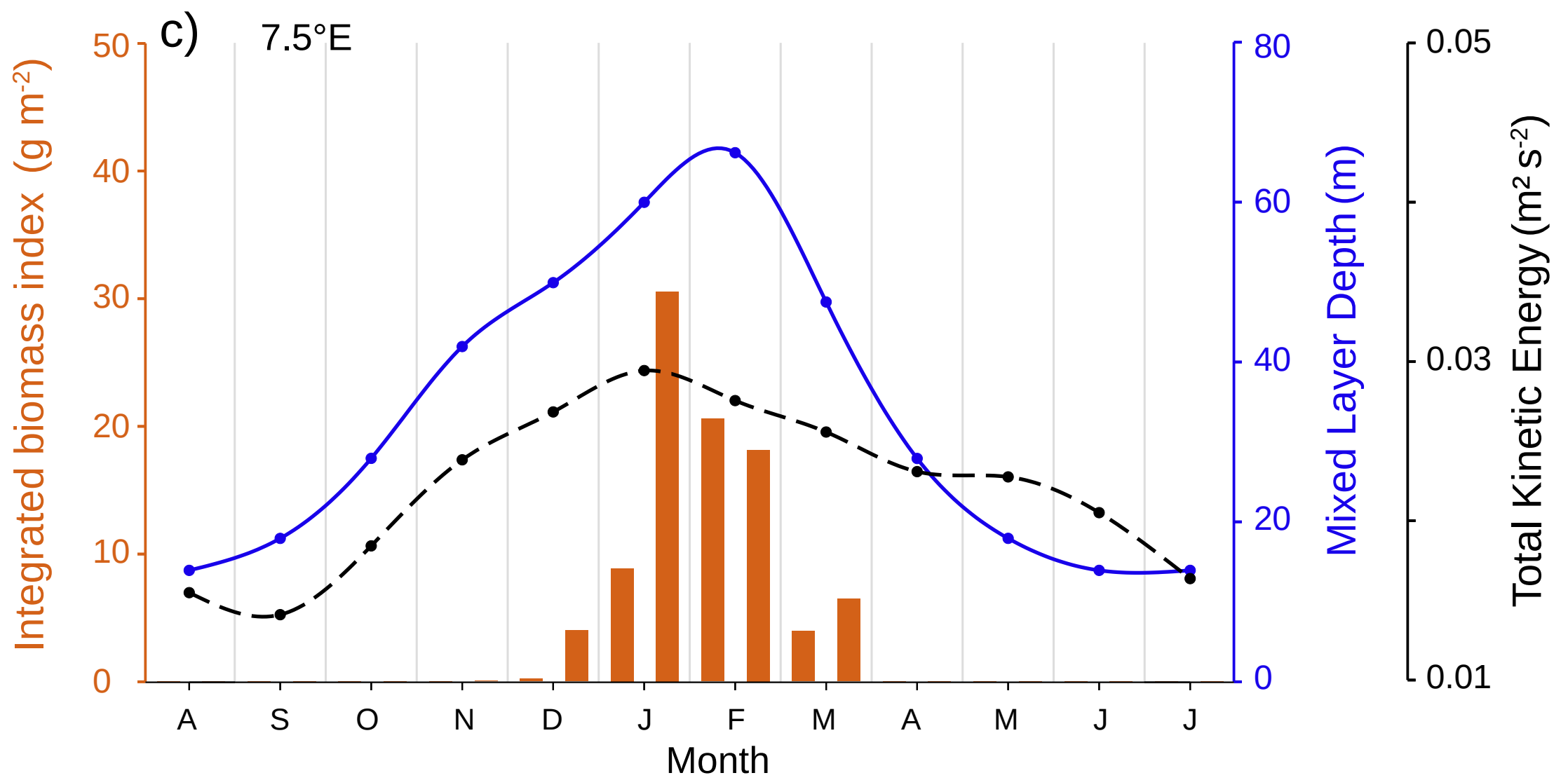
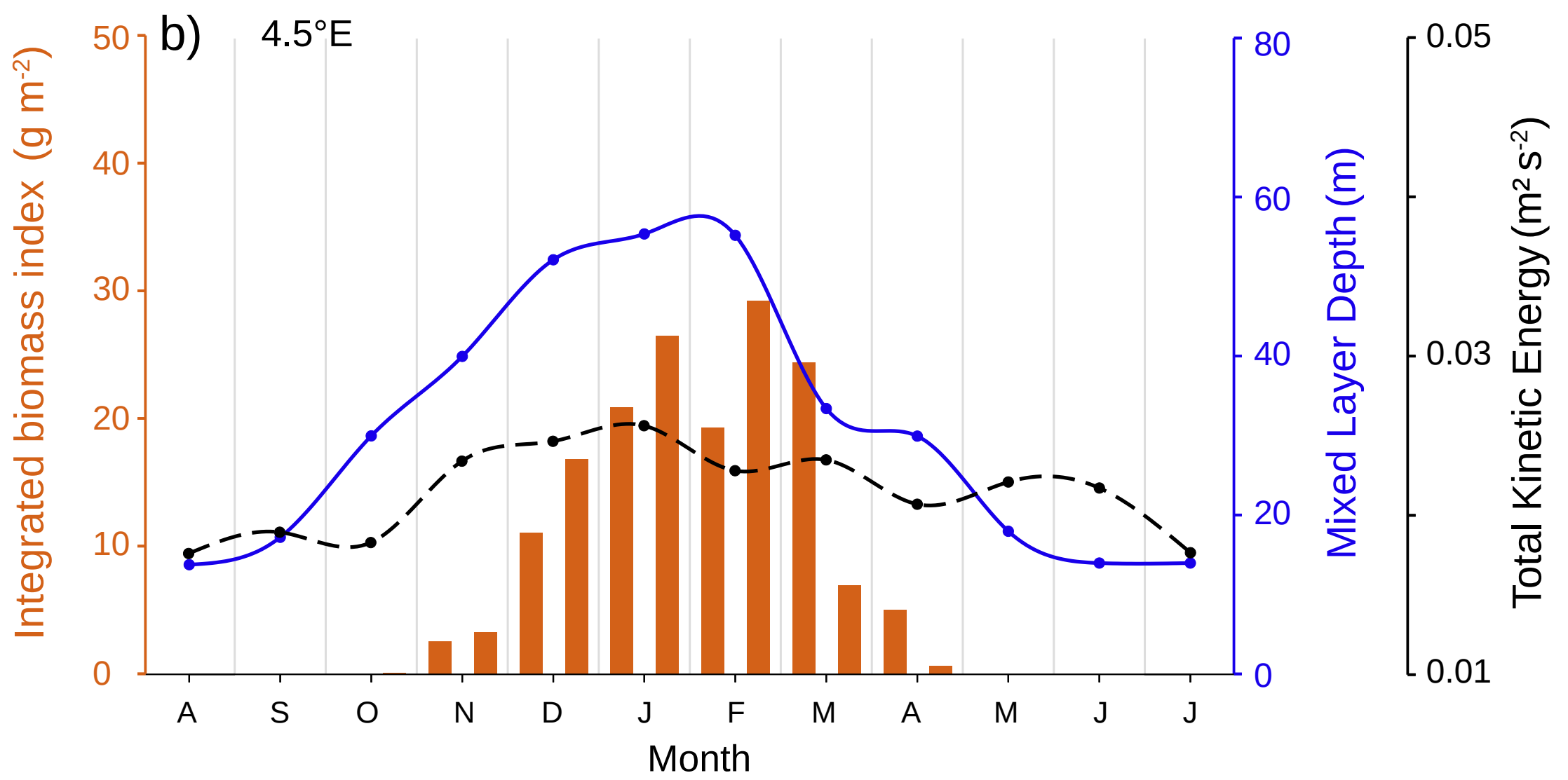
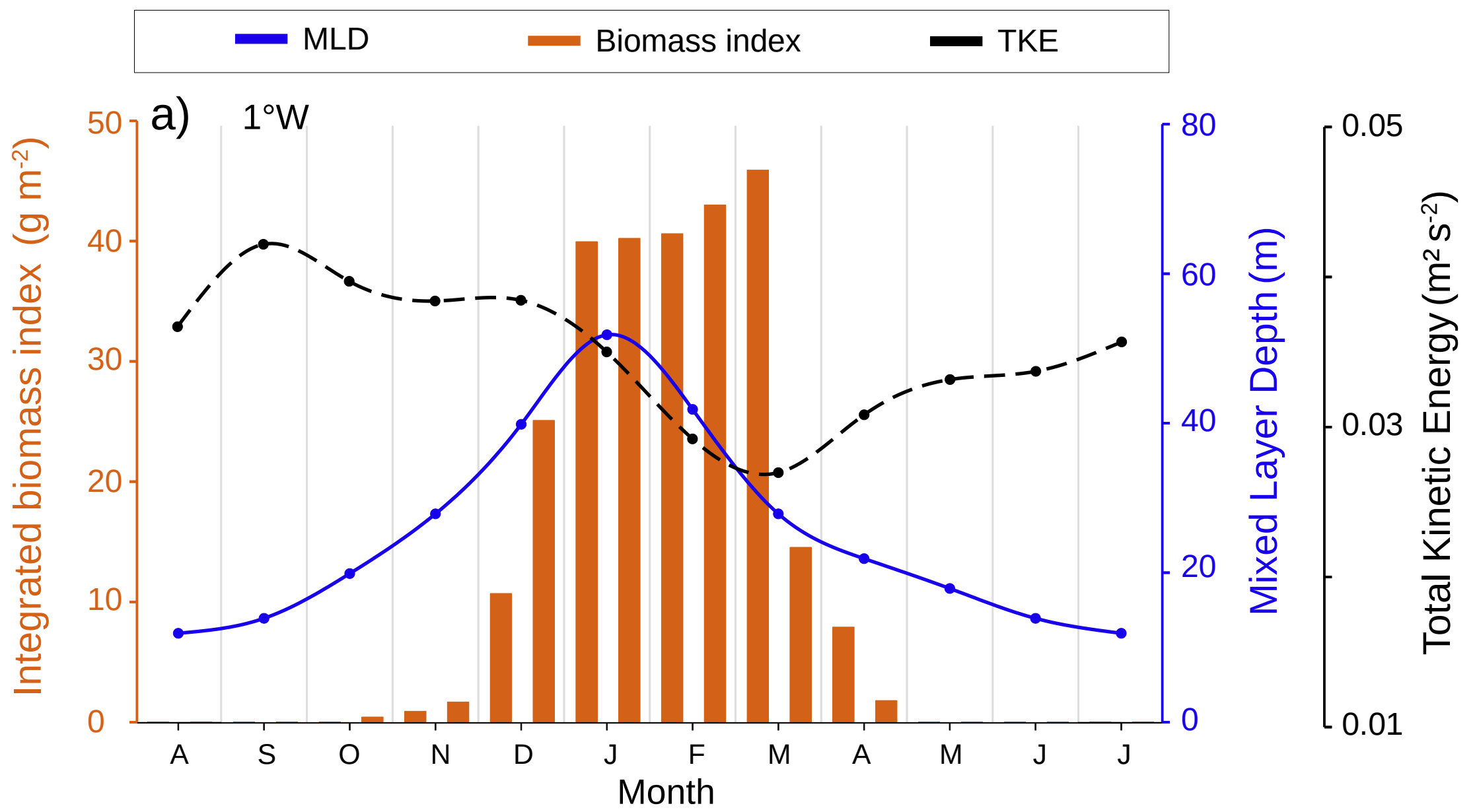
Chl-a (mg m^{-3})

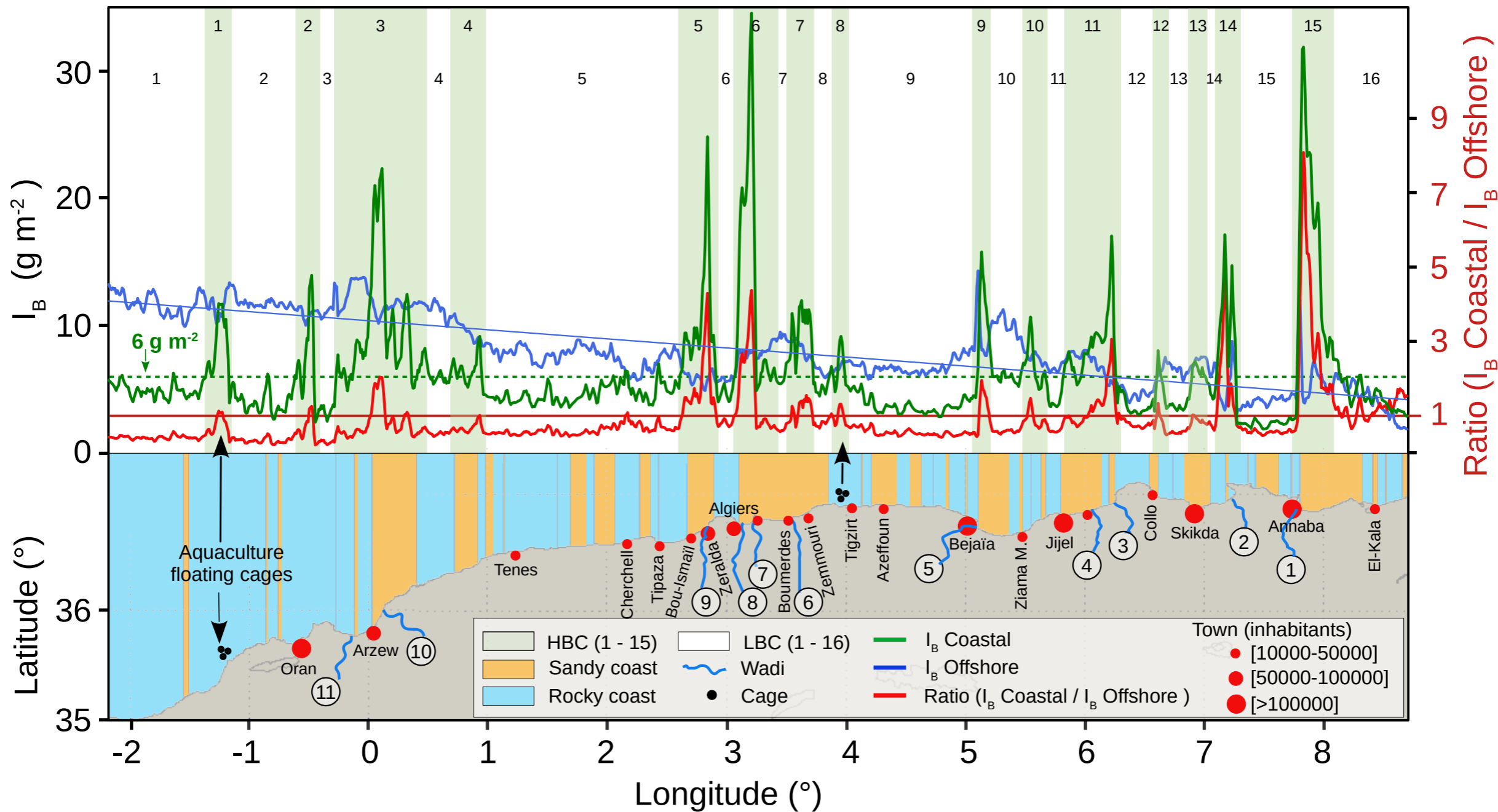


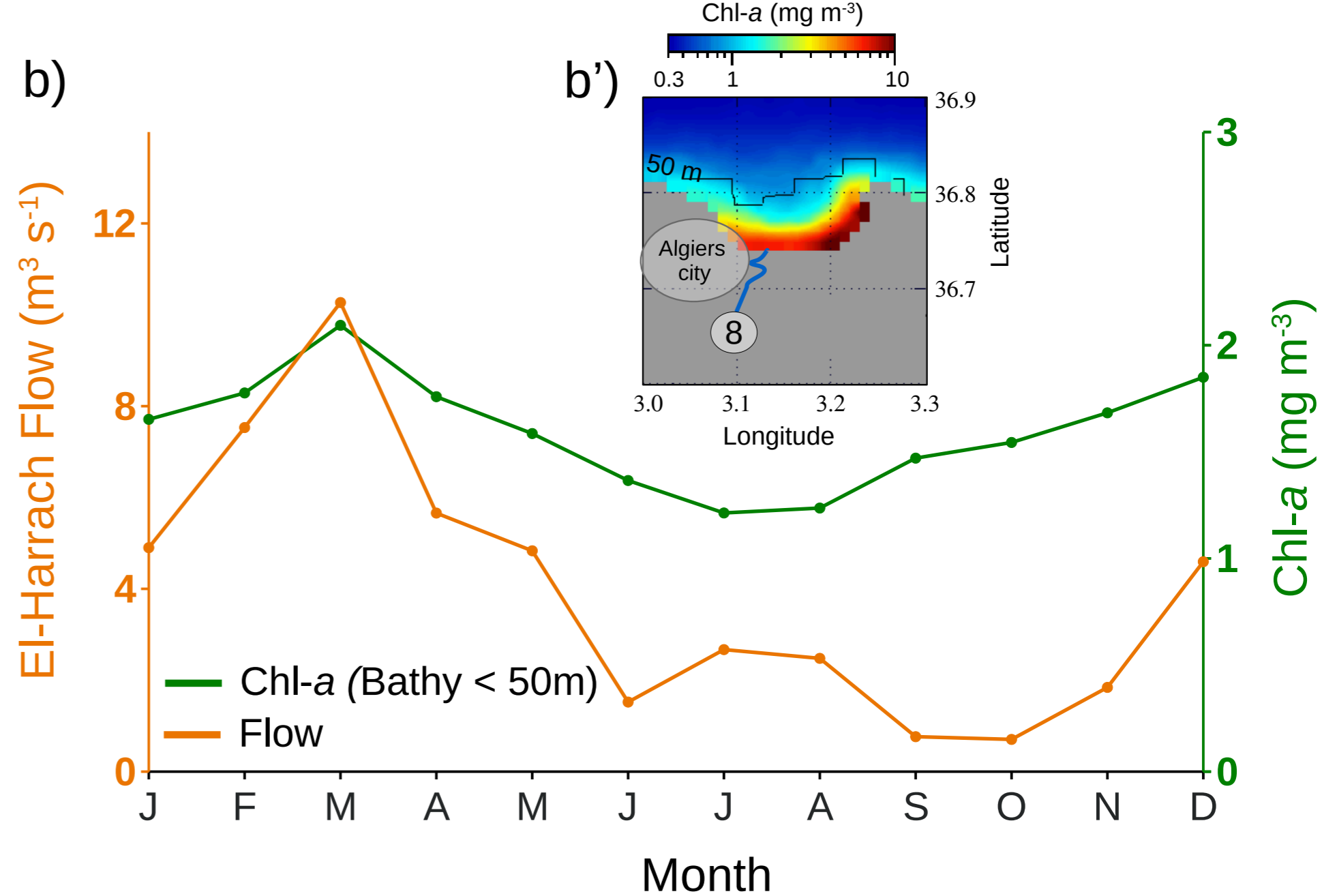
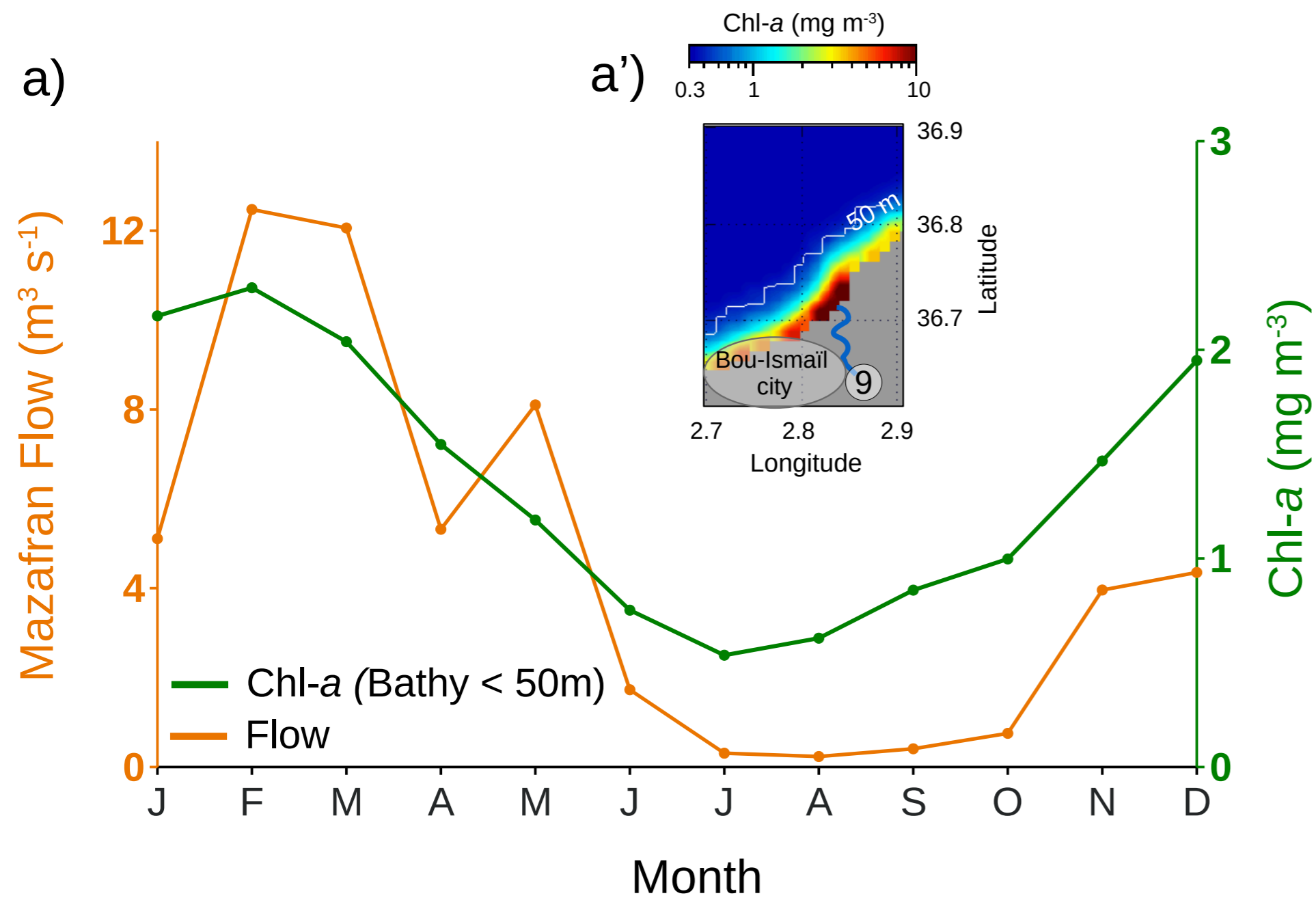












Season	I_B Coastal ($g\ m^{-2}$)		I_B Offshore ($g\ m^{-2}$)		I_B Coastal / I_B Offshore	
	<i>summer</i>	<i>winter</i>	<i>summer</i>	<i>winter</i>	<i>Jan & Feb</i>	<i>Dec & Mar</i>
LBC	0.7	10.1	0.317	23.2	0.64	2.57
HBC	2.9	18.2	0.626	24.8	0.73	4.14
HBC / LBC (%)	+305%	+80%	+97%	+7%		

LBC n°	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16
Chl- <i>a</i> (mg m ⁻³)	0.67	0.50	0.46	0.76	0.63	0.72	0.69	0.77	0.55	0.76	0.65	0.54	0.54	0.60	0.37	0.61
I _B (g m ⁻²)	4.9	4.0	3.2	5.7	4.6	4.7	5.6	5.3	3.8	5.9	4.8	3.7	3.8	4.5	2.4	4.2
City size [0-3]					●				●							●
Coast-type	R	R	R	R	R	R	S	S	R	S	R	R	R	R	S	S
Wadi presence																
Bay (or gulf)																B

HBC n°	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15
Chl- <i>a</i> (mg m ⁻³)	0.71	0.75	1.00	0.81	1.11	1.47	1.14	1.11	1.12	0.84	1.04	0.63	0.85	1.22	1.59
I _B (g m ⁻²)	8.6	8.0	9.7	6.5	10.0	13.2	9.4	8.7	11.6	7.9	8.6	7.2	6.5	11.5	14.4
City size [0-3]		●	●		●	●	●		●	●	●	●	●		●
Coast-type	R, ●	R	S	S	S	S	S	R, ●	S	R	S	S	S	R	S
Wadi presence			~~~~~		~~~~~	~~~~~	~~~~~		~~~~~					~~~~~	~~~~~
Bay (or gulf)		B	B		B	B	B		B	B	B		B	B	B

Model	Season	Response variables	Explanatory variables	p-value	Model %
m1	All seasons	[HBC ; LBC]	Coast type	-	67%
			Bay	-	
			Wadi	-	
			City	**	
m2	Winter	Chl-a \geq 1.5	Coast type	-	79%
			Bay	-	
			Wadi	***	
			City	*	
m3		I _B \geq 13	Coast type	-	77%
			Bay	**	
			Wadi	-	
			City	**	
m4	Summer	Chl-a \geq 0.5	Coast type	-	57%
			Bay	-	
			Wadi	*	
			City	-	
m5		I _B \geq 2.4	Coast type	-	78%
			Bay	*	
			Wadi	*	
			City	-	

Statistical signification of p-value (correlation is significant with p-value < 0.05 (5%)):
*** < 0.1 % ; ** < 1% ; * < 5% ; 5% < . < 10% ; - > 10%