

1 **Physical mechanisms for biological carbon uptake during the onset of the spring**  
2 **phytoplankton bloom in the northwestern Mediterranean Sea (BOUSSOLE site)**

3 Liliane Merlivat<sup>1</sup>, Michael Hemming<sup>2</sup>, Jacqueline Boutin<sup>1</sup>, David Antoine<sup>3, 4</sup>, Vincenzo  
4 Vellucci<sup>5</sup>, Melek Golbol<sup>5</sup>, Gareth A. Lee<sup>6</sup>, Laurence Beaumont<sup>7</sup>

5 <sup>1</sup> Sorbonne Université, CNRS/IRD/MNHN, LOCEAN, IPSL, Paris, France

6 <sup>2</sup> Coastal and Regional Oceanography Lab, Centre for Marine Science and Innovation, UNSW  
7 Sydney, Sydney, NSW 2052 Australia

8 <sup>3</sup> Remote Sensing and Satellite Research Group, School of Earth and Planetary Sciences,  
9 Curtin University, Perth, WA 6845, Australia

10 <sup>4</sup> Sorbonne Université, CNRS, Laboratoire d'Océanographie de Villefranche, Villefranche sur  
11 Mer 06230, France

12 <sup>5</sup> Sorbonne Université, CNRS, Institut de la Mer de Villefranche, Villefranche sur Mer 06230,  
13 France

14 <sup>6</sup> Centre for Ocean and Atmospheric Sciences, School of Environmental Sciences, University  
15 of East Anglia, Norwich Research Park, Norwich NR4 7TJ, UK


16 <sup>7</sup> Division Technique INSU-CNRS, 92195 Meudon CEDEX, France

17

18 Correspondence to: [liliane.merlivat@gmail.com](mailto:liliane.merlivat@gmail.com)

19 Keywords: Bloom onset ; Carbon uptake ; Air-sea interaction; Mediterranean sea

20 Abstract

21 Several trigger mechanisms have been proposed for the onset of the phytoplankton spring  
22 bloom. Among these, that phytoplankton cells begin to bloom when they experience higher  
23 average light levels in shallower mixed layers, a result of the surface net heat fluxes becoming  
24 positive and wind strength decreasing. We study the impact of these two forcings in the   
25 northwestern Mediterranean Sea. We take advantage of hourly measurements of oceanic and  
26 atmospheric parameters collected at two neighboring moorings during the months of March  
27 and April in the years 2016 to 2019, combined with glider data in 2016. The onset of the

28 surface phytoplankton growth is concomitant with the start of significant biological activity  
29 detected by a sudden decrease in dissolved inorganic concentrations derived from  
30 measurements in the upper 10 m of the water column. A rapid reduction in wind stress  
31 following high-wind events is observed at the same time. A resulting shallow mixing layer  
32 favors carbon uptake by phytoplankton lasting a few days. Simultaneously, the air-sea net  
33 heat flux switches from negative to positive linked to changes of the latent air-sea heat flux,  
34 which is proportional to the wind speed. This results in an increased thermal stratification of  
35 the ocean's surface layers. In 2016, glider data show that the mixing layer is significantly  
36 shallower than the mixed layer at the onset of the surface phytoplankton bloom. We conclude  
37 that decreases in the mixing and mixed layer depths lead to the onset of the phytoplankton  
38 bloom due to the relaxation of wind speed following storms. We estimate net daily  
39 community production in the mixing layer over periods of 3 days between 2016 and 2019 as  
40 between  $38 \text{ mmol C m}^{-2}$  and  $191 \text{ mmol C m}^{-2}$ . These results have important implications as  
41 biological processes play a major role in the seasonal evolution of surface  $\text{pCO}_2$  and thereby  
42 the rate of reduction of atmospheric  $\text{CO}_2$  by exchange at the air-sea interface.

43

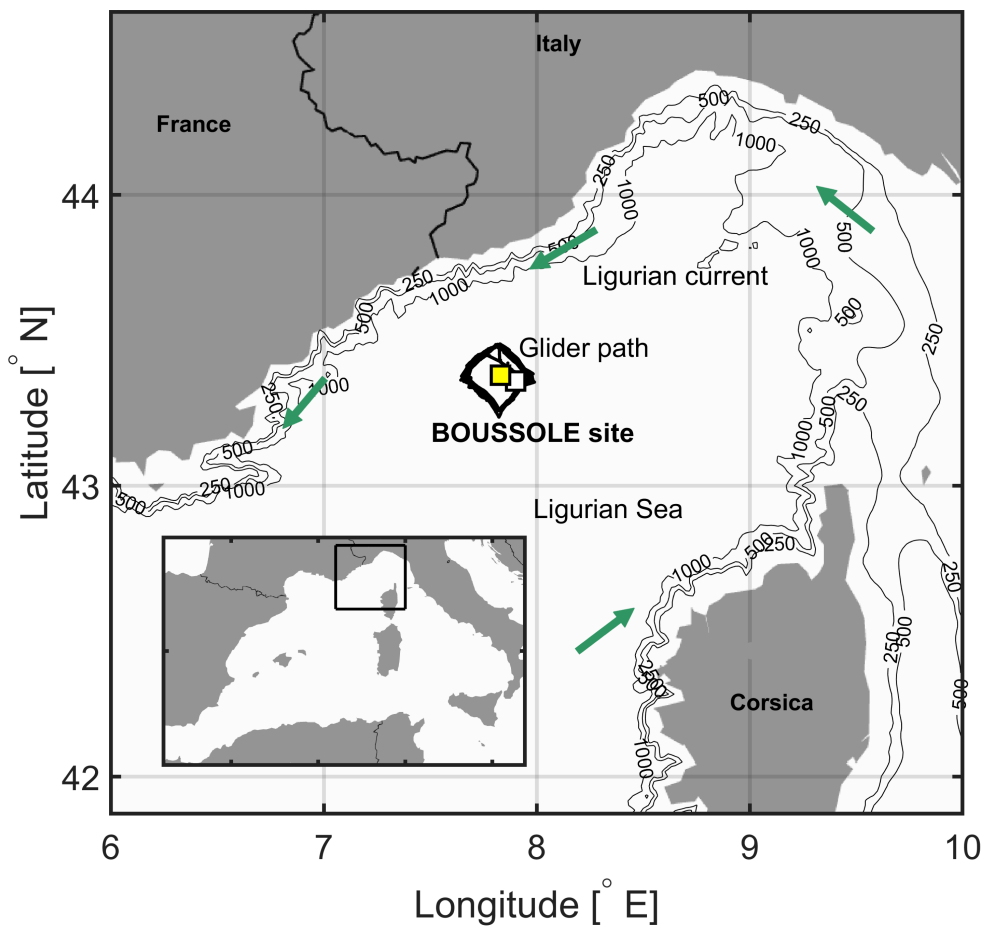
44

45

## 46 Introduction

47 Investigating the physical and biological mechanisms contributing to changes in air-sea  
48 carbon dioxide ( $\text{CO}_2$ ) fluxes is essential. It is a critical component of the global carbon cycle  
49 and the climate system with the ocean removing about a quarter of the  $\text{CO}_2$  emitted into the  
50 atmosphere by human activities over the last decade. It depends on the gradient of  $\text{CO}_2$  partial  
51 pressure,  $\text{pCO}_2$ , between the atmosphere and the surface ocean [Takahashi et al., 2009;  
52 Rödenbeck et al., 2013; Merlivat et al., 2015].  $\text{pCO}_2$  in the ocean mixed surface layer  
53 depends on the relative contribution of dynamic transport, thermodynamics and  
54 biogeochemistry. As the biological carbon pump lowers  $\text{pCO}_2$  and Dissolved Inorganic  
55 Carbon (DIC) at the sea surface, the production of plankton biomass during a bloom can be  
56 sufficient to reverse the annual cycle of surface-ocean  $\text{CO}_2$  uptake. A decrease in surface DIC,  
57 after physical processes have been eliminated, indicates the beginning of phytoplankton  
58 spring growth. Conceptual frameworks have been proposed to explain when depth-integrated

59 phytoplankton biomass starts accumulating during blooms [Behrenfeld, 2010; Sverdrup,  
60 1953]. The timing of the initiation of the surface spring phytoplankton bloom depends in  
61 particular on atmospheric forcing. The physical processes of wind stress, heat flux and  
62 vertical mixing control the depth of the mixed/mixing layer and thus the availability of light  
63 [Siegel et al, 2002, Chiswell, 2011; Taylor and Ferrari, 2011; Brody and Lozier, 2015;  
64 Enriquez and Taylor, 2015, Rumyantseva et al, 2019]. Thus, to better understand which  
65 fundamental physical processes control the timing of the onset of the surface bloom, in situ  
66 measurements are needed at the appropriate hourly to daily timescale. It is important to draw  
67 attention to the atmospheric forcing timescale. In biogeochemical ocean modeling studies,  
68 typically built on atmospheric models, reanalysis or satellite data, time resolution of at best a  
69 few days are required [Mahadevan et al, 2012].



70

71 Figure 1. The area of the northwestern Mediterranean Sea showing the French and Italian  
72 Rivas, the island of Corsica, the main current branches (green arrows) in the Ligurian Sea  
73 and the location of the BOUSSOLE buoy (white square) and the AZUR meteorological buoy  
74 (yellow square). The black line indicates the glider's path in March-April 2016.

75 In the northwestern Mediterranean Sea, the BOUSSOLE (Bouée pour l'acquisition de Séries  
76 Optiques à Long Terme; Antoine et al., 2006, 2008a,b) buoy is located in the central area of  
77 the Liguro-Provençal basin (Fig.1), where the dominant ocean currents are generally weak  
78 ( $<20 \text{ cm s}^{-1}$ ) [Millot, 1999; Antoine et al., 2008b; Niewiadomska et al., 2008]. The main  
79 processes that govern the  $\text{pCO}_2$  variations at the sea surface on an annual scale, after removal  
80 of the temperature effect, are vertical convective mixing and biological production [Begovic  
81 and Copin-Montégut, 2002; Copin-Montégut et al., 2004; Hood and Merlivat, 2001; Merlivat et  
82 al., 2018]. Intense convection resulting from repeated high wind events in winter or early  
83 spring when atmospheric temperatures are typically low brings nutrients to the surface layer  
84 [Andersen and Prieur, 2000; Antoine et al., 2008b; Marty et al., 2002; Pasqueron de  
85 Fommervault et al., 2015].

86 The objective of this study is to examine the impact of atmospheric forcing on surface  
87 phytoplankton bloom initiation in the northwestern Mediterranean Sea. To this aim we use  
88 concurrent hourly data of seawater carbon chemistry measured at depths less or equal to 10 m  
89 at the BOUSSOLE mooring [Merlivat et al., 2018] and atmospheric flux measurements at the  
90 Azur meteorological buoy 3.1 nm (5.8 km) away from BOUSSOLE in March and April 2016  
91 to 2019. Additional measurements of phytoplankton chlorophyll-a (Chla), temperature and  
92 density were provided by an underwater glider deployed in March-April 2016. The initiation  
93 of biological carbon uptake is revealed by the sudden decrease in DIC measured at the  
94 BOUSSOLE buoy.

95 In the Material and Methods section, we provide the description of data processing,  
96 theoretical framework, and calculations used to evaluate mixing regimes in the water column  
97 and net community production. The results section presents measured atmospheric and  
98 biogeochemical parameters observed in March-April, and NCP estimates from 2016 to 2019.  
99 The Discussion section relates our results to previous findings and presents our conclusions  
100 regarding the drivers of phytoplankton bloom initiation.

101


## 102 2 Material and methods

### 103 2.1- Data from the BOUSSOLE mooring

104 CARIOCA sensors [Merlivat and Brault, 1995] installed on the BOUSSOLE buoy ( $43^{\circ}22'N$ ,  
105  $7^{\circ}54'E$ ; Fig.1) monitored hourly partial pressure,  $\text{pCO}_2$ , at both 3 and 10 m alongside salinity

106 and temperature (SEABIRD Scientific SBE 37 MicroCAT CTDs). Protocols for calibration of  
107 in situ measurements are described in [Merlivat et al, 2018]. CARIOCA measurements of  
108 pCO<sub>2</sub> and temperature are combined with total alkalinity (Alk) derived from salinity  
109 [Merlivat et al., 2018] to calculate DIC. The dissociation constants of Mehrbach et al., (1973)  
110 refitted by Dickson and Millero (1987) are used, as recommended by Alvarez et al. (2014) for  
111 the Mediterranean Sea. For the March-April months mean salinity is 38.3, corresponding with  
112 mean Alk equal to 2571.4 μmol kg<sup>-1</sup> [Copin et al, 2004]. The expected precision of the  
113 CARIOCA pCO<sub>2</sub>, and DIC is 5 μatm, and 3 μmol kg<sup>-1</sup>, respectively [Merlivat et al, 2018].

## 114 2.2- Glider observations

115 An iRobot Seaglider model 1KA (SG537) with an ogive fairing operated by the University of  
116 East Anglia (UEA) [see Hemming et al., 2017] was deployed during the BOUSSOLE  
117 monthly cruise (#169) [Golbol et al 2000] and recovered 1 month later with a ship of  
118 opportunity. Diving to 1000 m, 147 dives were completed between March 7 and April 5 2016  
119 repeatedly-sampling a square-shaped path surrounding the BOUSSOLE buoy (Fig. 1) with  
120 corners pointing to cardinal directions. Out of 147 dives, a number of them contain data that  
121 are within 10 km of the BOUSSOLE mooring, i.e. when it was sampling in the eastern corner  
122 of the diamond path. The glider was equipped with a non-pumped SEABIRD Scientific SBE  
123 9 CTD instrument package providing temperature and salinity, and a WET Labs Eco Puck  
124 sensor providing relative phytoplankton Chla fluorescence, factor  calibrated into Chla  
125 concentration units (mg m<sup>-3</sup>).

## 126 2.3- Azur meteorological buoy

127 The Azur meteorological buoy has been deployed since 1999 by the French weather forecast  
128 Agency (Meteo-France) at 43°23'N, 7°50'E (Fig. 1). It provides hourly measurements of  
129 surface wind speed extrapolated to 10 m height, U<sub>10</sub> (m s<sup>-1</sup>), net incoming and emitted long-  
130 wave radiation, incoming short-wave radiation, and the sensible and latent heat fluxes. These  
131 are needed to estimate the net air-sea heat budget at the sea surface (<http://mistrals.sedoo.fr>).  
132 U<sub>10</sub> is used to calculate wind stress, τ, as follows:  $\tau = \rho_a C_d U_{10}^2$ , where ρ<sub>a</sub> is air density (1.2  
133 kg m<sup>-3</sup>), and C<sub>d</sub> is the drag coefficient (1.4 x 10<sup>-3</sup>). Photosynthetically Active Radiation (PAR,  
134 mol photons m<sup>-2</sup> d<sup>-1</sup>) is related to solar irradiance (i.e. short-wave flux), SW, following PAR =  
135 0.473 SW [Papaioannou et al., 1993].

## 136 2.4- Estimation of mixed and mixing layer depths

137 The mixing layer depth,  $Z_{mx}$ , is the upper part of a mixed layer of uniform density where  
 138 active turbulence occurs [Brainerd and Gregg 1995]. Several recent studies have highlighted  
 139 the significance of this layer to understand the phytoplankton phenology of phytoplankton  
 140 blooms [Brody and Lozier, 2015; Taylor and Ferrari, 2011]. Autonomous profiling floats  
 141 equipped with bio-optical sensors have also highlighted the importance of mixing rather than  
 142 mixed layers to characterize the onset and the development of a phytoplankton bloom. Wind-  
 143 driven and buoyancy-driven regimes control the characteristics and depth of active mixing  
 144 [Brody and Lozier, 2014, 2015, Ramuyantseva et al, 2015, Lacour et al, 2019, Pellichero et al,  
 145 2020]. Wind-driven mixing dominates in the presence of weak buoyancy forcing when heat  
 146 fluxes are small. In that case, the dominant mixing length scale is equal to the Ekman length  
 147 layer depth scale.

148 The daily mean mixed layer depth,  $Z_{ml}$  is derived for 2016 from glider vertical profiles of  
 149 temperature and salinity using the algorithm provided by Holte and Talley, (2009) estimated  
 150 using the potential density algorithm.

151 To estimate  $Z_{mx}$ ,  $\tau$  is used to calculate the Ekman layer depth,  $Z_{ek}$ ,  $Z_{ek}=\gamma u_*/f$ , where  $\gamma$  is an  
 152 empirical constant (0.5),  $f$  is the Coriolis parameter ( $10^{-4} s^{-1}$ ) over the BOUSSOLE area, and  
 153  $u_*$  is the turbulent friction velocity,  $u_* = \sqrt{\frac{\tau}{\rho_w}}$  ( $m s^{-1}$ ), with  $\rho_w$  the density of the surface sea  
 154 water [Lacour et al, 2019]. When heat fluxes are large and positive, Enrikez and Taylor  
 155 (2015) proposed to express the mixing depth  $Z_{mx}$  in terms of the surface forcing, i.e. the  
 156 surface stress and a stabilizing surface buoyancy flux  $B_0$ , as follows:

$$157 \quad \frac{1}{Z_{mx}^2} = \frac{f^2}{(C_3 u_*)^2} + \frac{f B_0}{(C_4 u_*^2)^2} \quad (1)$$

158 With  $C_3=1$ ,  $C_4=0.57$ ,  $B_0 = Q \lambda g / c_p \rho_0$ , where  $Q$  is the net surface heat flux,  $\lambda$  the thermal  
 159 expansion coefficient ( $1.65 \cdot 10^{-4} \text{ } ^\circ\text{C}^{-1}$ ),  $g$  the gravitational acceleration ( $9.81 \text{ m s}^{-2}$ ),  $c_p$  the  
 160 specific heat of water ( $4 \cdot 10^3 \text{ J kg}^{-1} \text{ } ^\circ\text{C}^{-1}$ ) and  $\rho_0$  the pure water density ( $1000 \text{ kg m}^{-3}$ ).

## 161 2.5-Estimation of euphotic zone depth

162 The euphotic depth,  $Z_{eu}$ , is generally derived as the depth where PAR is 1% of its surface  
 163 value.  $Z_{eu}$  can be estimated from Chla at the surface using the method described by Lee et al.  
 164 (2007) based on the equation of Morel and Berthon (1989):

$$165 \quad Z_{eu} = 34 (\text{Chla})^{-0.39} \quad (2)$$

166 For the period between 2017 and 2019, GlobColour merged Chla products were used, which  
167 are based on satellite observations with a resolution of 25 km, and a binning period of 8 days  
168 (<http://www.oceancolour.org>). For 2016, glider surface Chla was used to derive Zeu instead  
169 of satellite measurements. We excluded glider Chla between 05:00 and 20:00 UTC affected  
170 by sunlight-induced fluorescence quenching, and we applied a depth-constant offset to glider  
171 measurements using Chla water samples available on March 7.

## 172 2.6-Estimation of average mixing layer irradiance

173 We calculate the average mixing layer irradiance,  $I$ , function of the incident surface  
174 irradiance, PAR, mixing layer depth,  $h$ , and the diffuse attenuation coefficient,  $K_d$ , estimated  
175 from surface chlorophyll-a concentrations (Venables and Moore, 2010 ).

$$K_d = 0.05 + 0.057 C_{\text{Chla}}^{0.58}$$

$$176 \quad I = \frac{\text{PAR}}{K_d h} (1 - e^{-K_d h}) \quad (3)$$

177

## 178 2.7 – Estimation of net community production

179 Biological Net Community Production rates, NCP ( $\text{mmol m}^{-2} \text{day}^{-1}$ ), are calculated for  
180 separate 3-day periods in March-April, 2016 to 2019 from DIC concentrations derived from  
181 CARIOCA measurements, assuming that measurements in the top 10 m of the water column  
182 are representative of an homogeneous mixing layer [Boutin and Merlivat, 2009; Merlivat et al,  
183 2015; Pelichero et al, 2020]. In the study area, current velocities are expected to be small  
184 [Millot, 1999; Antoine et al., 2008; Niewiadomska et al., 2008], hence to estimate NCP, we  
185 assume that the contribution of horizontal advection was negligible. Vertical mixing events  
186 are identified by an increase in DIC resulting from upward movement of high DIC Levantine  
187 Intermediate Water (LIW) at approximately 200 m depth [Copin-Montégut and Bégovic,  
188 2002]. We isolated time periods when local physical processes were largely one-dimensional  
189 in order to study changes in biological and chemical parameters that occurred during rapid  
190 transitions from deep mixing to intermittent stratification. In 2016, over a four-day period,  
191 March 18-21, the diurnal cycle of DIC values characterized by a maximum in the morning  
192 followed by a minimum at the end of the day indicates the onset of organic matter formation.  
193 The eddy diffusion term is negligible as the DIC gradient at the base of the mixing layer is  
194 very weak, entrainment is not expected as changes in DIC are observed during periods of

195 stratification. For the identified periods, biological production and air-sea exchange are the  
196 dominant processes responsible for daily changes in DIC.

197 NCP is thus estimated from temporal variation of the DIC integrated over  $Z_{\text{mx}}$  ( $\text{DIC}_{\text{int}}$ ), and  
198 corrected for the air-sea  $\text{CO}_2$  flux contribution [Pelichero et al, 2020]:

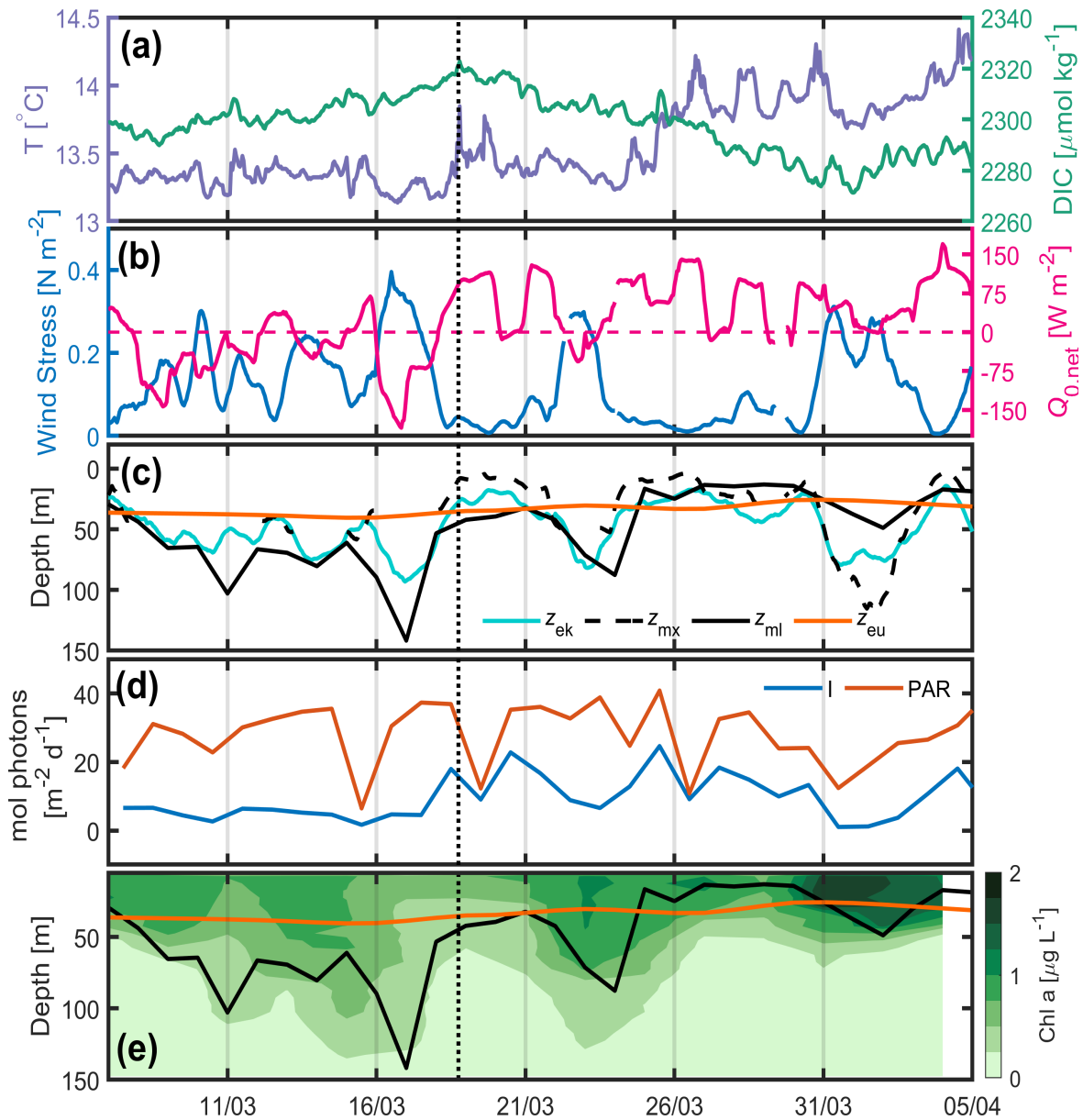
199 
$$\text{NCP} = \left( \frac{\Delta \text{DIC}_{\text{int}}}{\Delta t} \right)_{\text{bio}} = \left( \frac{\Delta \text{DIC}_{\text{int}}}{\Delta t} \right)_{\text{meas}} - k \times s \times (\text{pCO}_{2\text{atm}} - \text{pCO}_{2\text{sw}}) \quad (4)$$

200 The first term in equation 4,  $\left( \frac{\Delta \text{DIC}_{\text{int}}}{\Delta t} \right)_{\text{meas}}$ , is the daily variation of  $\text{DIC}_{\text{int}}$  calculated over 3  
201 days from the date identified as the start of phytoplankton bloom ( $t_0$ ), whilst the remaining  
202 part corresponds to DIC variability due to air-sea  $\text{CO}_2$  exchange, where  $k$  is the gas transfer  
203 velocity [Wanninkhof, 2014],  $s$  is the solubility of  $\text{CO}_2$  in seawater [Weiss, 1974], and  
204  $\text{pCO}_{2\text{sw}}$  and  $\text{pCO}_{2\text{atm}}$  are respectively the partial pressures of  $\text{CO}_2$ , in  $\mu\text{atm}$ , in sea water and in  
205 air [Merlivat et al, 2018].

## 206 3 Results

### 207 3.1- Description of the 2016 time-series





208

209 Figure 2. 2016 (a) Hourly DIC (green) and SST(purple) at the BOUSSOLE site, (b) hourly  
 210 wind stress (blue) and net heat flux (pink) at the Azur buoy site. The pink dotted line indicates  
 211 the change of sign of the net heat flux from negative to positive values, (c) daily depths of the  
 212 mixed layer (black), mixing layer (dotted black), Ekman layer (cyan) and euphotic layer  
 213 (orange) at the BOUSSOLE site. The black dotted line on panels (a), (b), (c) (d) and (e)  
 214 indicates the onset time of the bloom (March 18). (d) PAR (orange) and I, average mixing  
 215 layer irradiance (blue), (e) Glider-derived depths of the mixed layer (black curve, same as in  
 216 (c)), euphotic layer (orange), and nighttime 2-day binned chlorophyll concentration (green  
 217 coloring) retrieved between March 7 and April 5.

218 The temporal evolution of physical and biogeochemical parameters between March 7 and  
219 April 5 2016 is displayed in Fig. 2. It shows an initial increase in DIC until March 18 (Fig.  
220 2a), resulting from the vertical mixing of surface waters with the DIC-enriched underlying  
221 LIW [Copin-Montégut and Bégovic, 2002], followed by a decrease in DIC that corresponds  
222 to an increase of surface Chla (Fig 2e). Temperature increases intermittently over the same  
223 time period (Fig. 2a), with a sharp increase on March 18 and 25. Before March 18 frequent  
224 atmospheric fronts associated with wind forcing between 0.2 and 0.5 N m<sup>-2</sup> were observed  
225 (Fig. 2b).  $Q_{0,net}$  was negative most of the time leading up to March 18, but mostly positive  
226 afterwards (Fig. 2b). A clear anticorrelation was observed between  $\tau$  and  $Q_{0,net}$  (Fig. 2b). The  
227 mixed-layer depth derived from the glider measurements (< 20 km away from the Boussole  
228 site) deepened to > 70 m at times between March 8 and March 18, consistent with the passage  
229 of frequent atmospheric fronts. From March 18 to April 5,  $Z_{ml}$  shoaled from ~ 45 m to ~10 m,  
230 excluding two short periods around March 24 and April 2 when atmospheric fronts were  
231 passing (Fig. 2c). From March 18 to 31,  $Z_{mx}$  was for a considerable time either shallower than,  
232 or at similar depth to  $Z_{ml}$  and  $Z_{eu}$  (Fig. 2c). The start of the increase in PAR from March 15  
233 precedes the increase in irradiance by 3 days as a result of the decrease of the mixing layer  
234 depth observed only after March 18 (Fig. 2d).

235 From March 18 to April 5, an increase in glider Chla was observed in the upper layer (Fig.  
236 2e) as the result of the biomass accumulation within the water column following the initial  
237 growth of surface phytoplankton [Behrenfeld, M. and E. Boss, 2014]. The same was observed  
238 in the Southern Ocean [Pelichero et al, 2019].

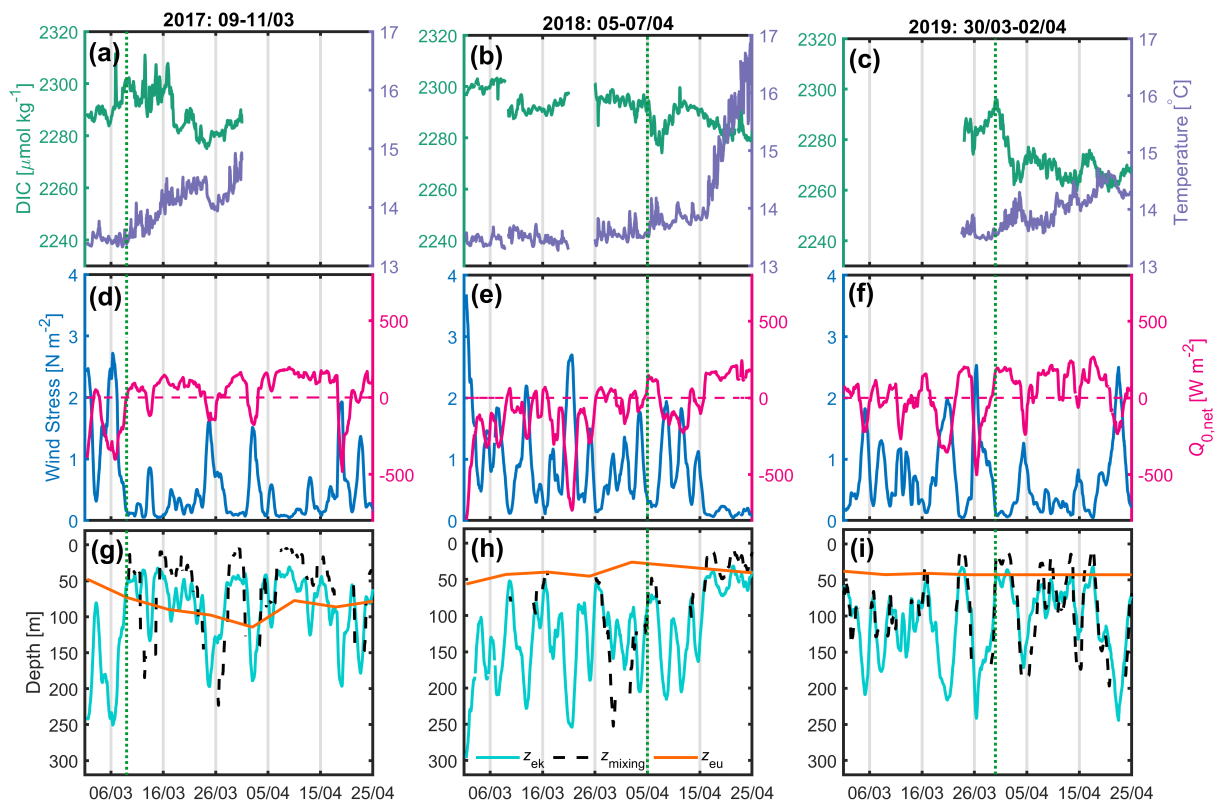
### 239 3.2- Description of the 2017-2018-2019 time-series

240 Only measurements at the BOUSSOLE and Azur buoys were available between 2017 and  
241 2019. Here we focus on the periods between March 1 and April 25. In 2017, from March 9,  
242 after the passage of a very active atmospheric front ( $\tau$  up to 2.7 N m<sup>-2</sup>), a decrease in DIC  
243 accompanied by an increase in temperature was observed for a period of 14 days despite a  
244 few short bursts of vertical mixing, typically lasting less than 1 day (Fig.3a, d, g). At the same  
245 time, the net surface heat flux switched from negative to positive most of the time. Over this  
246 14-day period,  $Z_{mx}$  was predominantly shallower than the  $Z_{eu}$  at times when DIC was  
247 decreasing.

248 In 2018 (Fig. 3b, e, h) and 2019 (Fig. 3c, f, i), the weather records throughout the month of  
249 March are very similar, showing a succession of very strong wind events at the start of the

250 investigated time periods, with  $\tau$  values of up to  $3.7 \text{ N m}^{-2}$  and  $Z_{ek}$  values as deep as 300 m.  
 251 On April 5, 2018, following a period of high wind stress resulting in the formation of a deep  
 252 mixing layer ( $\sim 250 \text{ m}$ ), a decrease in DIC and an increase in SST were observed over 3 days.  
 253 At the same time, the net heat flux oscillated around zero. In 2019, CARIOCA data were only  
 254 available after March 24. On March 30, 3 days after high winds, a decrease in DIC and an  
 255 increase in temperature lasted 3 days, while simultaneously the net heat flux reversed from  
 256 negative to positive.

257



258

259 Figure 3. (a, b, c) 2017, 2018 and 2019 DIC (green) and SST (purple) at the BOUSSOLE site,  
 260 (d, e, f) wind stress (blue) and net heat flux (red) at the Azur buoy site. (d, e, f) The red dotted  
 261 line indicates the change of sign of the net heat flux from negative to positive values, (g, h, i)  
 262 depths of the mixing layer (dotted black), Ekman layer (cyan) and euphotic layer (orange) at  
 263 the BOUSSOLE site. The vertical dotted green line indicates the onset of the decrease of DIC.

264 For the period between 2017 and 2019, only satellite chlorophyll-a concentrations with a  
 265 binning period of 8 days were available, which is too large to calculate the average mixing  
 266 irradiance as it is highly variable on a daily scale (Fig 2.d).

267 3.3- NCP

268 NCP was estimated (Eq. 3) for 3-day periods over the four time-series between 2016 and  
 269 2019 (Table 1). Because we compute NCP only during periods when  $Z_{ml}$  and  $Z_{mx}$  shoal and  
 270 the air-sea  $CO_2$  flux is very small, the measured total daily changes of DIC,  $dDIC/dt$   
 271 (correlation coefficient,  $r^{*2} > 0.70$ ), is expected to represent the biological consumption of  
 272 DIC. In 2016, 2017, and 2019, daily biological carbon uptake fluxes are between 38 and 191  
 273  $mmol\ m^{-2}\ d^{-1}$ . PAR and I vary respectively from 30 to 53 and 13 to 25  $mol\ photons\ m^{-2}\ d^{-1}$  for  
 274 the four considered periods (Table 2).

	$T_0$	$dDIC/dt^a$	$r^{*2\ b}$	$\Delta DIC / \Delta t^c$	$r^{*2\ d}$	air-sea flux $CO_2^e$	NCP <sup>f</sup>
		$\mu mol\ kg^{-1}d^{-1}$		$mmol\ m^{-2}d^{-1}$		$mmol\ m^{-2}d^{-1}$	$mmol\ m^{-2}\ d^{-1}$
<b>2016</b>	March 18	-3.0	0.80	-37	0.68	1	38
10m							
<b>2017</b>	March 9	-4.5	0.84	-188	0.86	3	191
3m							
<b>2018</b>	April 5	-6.5	0.70			17	
3m							
<b>2019</b>	March 30	-8.3	0.93	-163	0.63	4	167
10m							

275

276 Table 1. Net community production computed in the mixing layer over 3 days (NCP) after the  
 277 bloom onset ( $T_0$ ). (a, b) variation of DIC at the measurement depth and linked correlation  
 278 coefficient; (c, d) change of integrated DIC over  $Z_{mx}$  and linked correlation coefficient ;(e)  
 279  $CO_2$  flux from the atmosphere to the ocean; (f), absolute value of calculated NCP is  
 280 considered in order to provide positive values throughout the presentation and discussion in  
 281 the manuscript.

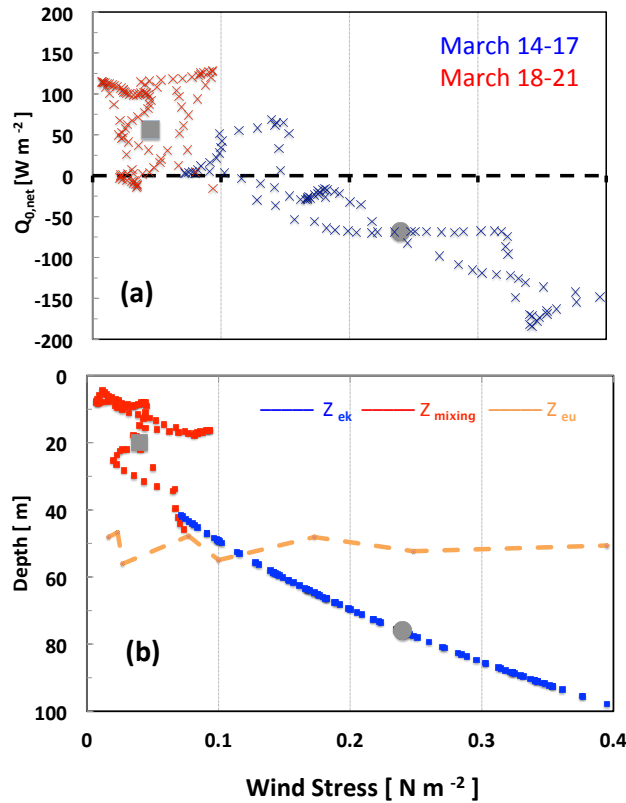
282

283 4 Discussion

284 4.1 Analysis of the physical drivers of the bloom initiation over the 4 years, 2016-2019

285 4.1.1 In 2016, the consumption of carbon indicated by the decrease in DIC from March 18  
 286 attests the initiation of the formation of phytoplankton biomass (Fig. 2.). Between 2

287 successive days, March 17 and 18, a rapid decrease in daily wind stress from 0.24 to 0.04 N  
 288  $\text{m}^{-2}$  results in a change of sign of the net heat flux from -68 to +56  $\text{W m}^{-2}$  (Fig.4a) and a  
 289 shoaling  $Z_{\text{mx}}$ , which is for the most part shallower than  $Z_{\text{eu}}$  (Fig.4b) after March 18.



290

1

291 Figure 4. Changes of physical parameters (hourly values) at the onset of the 2016 bloom  
 292 during 2 consecutive periods of 4 days, March 14-17 (blue) and March 18-21 (red) as a  
 293 function of wind stress (a) net surface heat flux (b) depths of the Ekman, mixing and euphotic  
 294 layer. Grey circles and grey squares indicate mean values respectively on March 17 and  
 295 March 18. For comparison, the orange line shows the euphotic layer depth (March 14-21).

296 This illustrates the very fast change in heat flux and mixing layer thickness resulting from the  
 297 decrease in wind stress. The main drivers to explain the initiation of near surface spring  
 298 phytoplankton blooms proposed by [Taylor and Ferrari, 2011, Brody-Lozier, 2014, 2015] are  
 299 a decrease in wind strength, a positive surface heat flux, and a decrease in the dominant  
 300 mixing length scale. Our results indicate that a decrease in  $\tau$  is a common component for these  
 301 last two mechanisms (Fig.4). The variability of  $Q_{0,\text{net}}$  is strongly controlled by the variability  
 302 of the latent heat flux proportional to the wind speed. From March 17 to 18, the latent heat

303 flux changed by  $84 \text{ W m}^{-2}$  representing nearly 2 thirds of the total change of  $Q_0$ , with the  
304 remainder originating from an increased short-wave flux. Hence,  $Q_{0,\text{net}}$  increased as  $\tau$   
305 decreased. In the same way  $Z_{\text{mx}}$ , which is related to  $\tau$  and  $Q_{0,\text{net}}$ , decreased. This highlights the  
306 major contribution of wind stress relaxation in controlling the net surface heat flux, and  
307 subsequent stratification leading to shallower  $Z_{\text{mx}}$

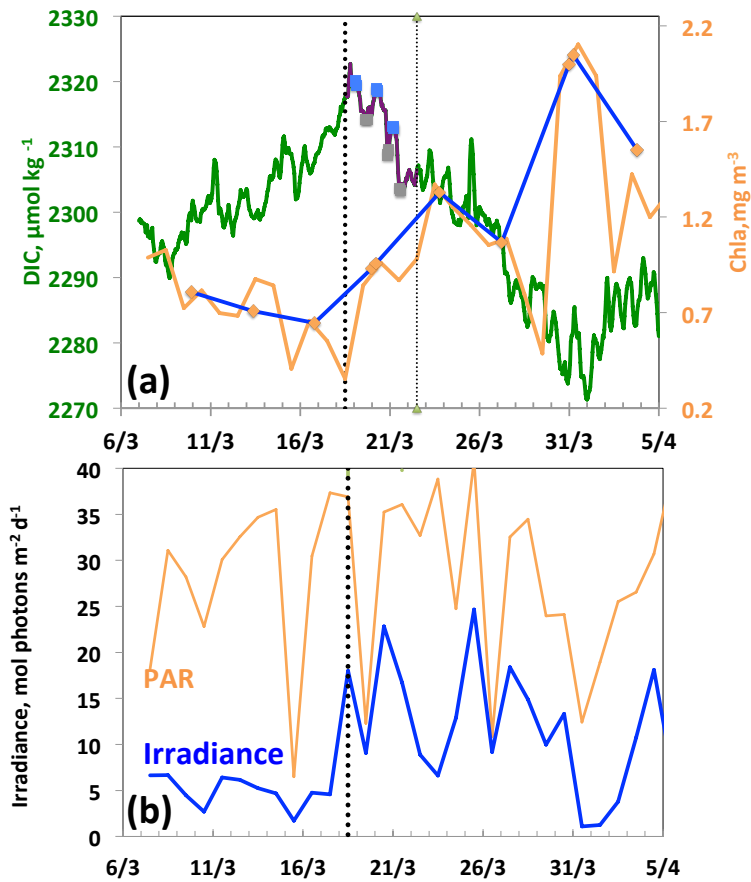
308 4.1.2 Over the 4 years, 2016-2019, during the months of March and April, at the very  
309 beginning of spring, the conditions required for initiating the spring bloom are very similar  
310 although the range of wind stress values is smaller in 2016 (Fig. 2, Fig. 3). The wind stress  
311 values are low following storm events. For all years except 2018,  $Z_{\text{mx}}$  is smaller than  $Z_{\text{eu}}$ . The  
312 average atmospheric and oceanographic conditions during the onset of the bloom are shown  
313 in Table 2. The decrease of wind-driven mixing is then a dominant physical mechanism,  
314 namely mixed layer shoaling which best predicts the timing of the spring bloom [Brody  
315 Lozier, 2014, 2015]. Likewise, in the Southern Ocean, Pelichero et al (2020) showed that a  
316 bloom is initiated only when the wind-driven mixed layer decreases, even if the net heat flux  
317 has been positive for some months.

#### 318 4.2 DIC, Chla, Irradiance at the ocean surface in 2016

319 Between 18 and 21 March, a diurnal cycle in DIC is observed (Figure 5a) characterized by a  
320 maximum in the morning followed by a minimum at the end of the day, indicating the onset  
321 of organic matter formation. The decrease in DIC is accompanied by an increase of glider  
322 surface Chla on 18 March and a simultaneous increase in the average mixing layer irradiance  
323 (Fig. 5 b). It is worth noting that the Chla maximum does not occur until 31 March, 13 days  
324 after the decrease in DIC. This maximum is the one identified in most cases by the Chla  
325 satellite measurements with a binning period of 8 days. It follows that satellite data cannot  
326 accurately characterize the conditions that prevail for the onset of a bloom, as these are  
327 dictated by a great temporal variability of atmospheric forcings. Due to the seasonal  
328 increase of surface irradiance in March-April, as well as the availability of nutrients following  
329 vertical mixing with intermediate nutrient-enriched waters [Begovic and Copin-Montégut,  
330 2002], the conditions for the onset of a bloom are met.

331

332



333

334 Figure 5. From March 7 to April 5, (a) DIC and Chla. The purple line and the squares  
 335 (blue, morning; grey, evening) indicate the 3 days biological diurnal DIC changes during the  
 336 period considered to compute NCP. The blue and orange lines indicate the surface Chla when  
 337 the glider was at a distance of less than 5 km (blue) and less than 20 km (orange) respectively  
 338 from the Boussole buoy. (b) PAR and I average mixing layer irradiance. The vertical dotted  
 339 black line indicates the onset of the bloom on March 18.

#### 340 4.3- Biological carbon uptake

341 In 2016, 2017, and 2019, at the onset of the bloom  $Z_{\text{mx}}$  is shallower than  $Z_{\text{eu}}$  (Table 2). Two  
 342 factors account for the observed large range of NCP variability at the time of triggering the  
 343 bloom:  $d\text{DIC}/dt$  and  $Z_{\text{ml}}$ . In 2016 and 2017, DIC decreased over a period of 14 days, whereas  
 344 it lasted for 3 days in 2018 and 2019, as a result of short-lived high wind events ( $\tau > 1 \text{ N m}^{-2}$ ).  
 345 On April 2, 2018, a high surface phytoplankton Chla,  $1.9 \text{ mg m}^{-3}$ , was measured by satellite  
 346 (Globcolour data) suggesting a strong biological signal. However, as  $Z_{\text{eu}}$  is significantly  
 347 shallower than  $Z_{\text{mx}}$  (26 m vs. 65m; Fig. 3h), it would be incorrect to estimate NCP integrated  
 348 over  $Z_{\text{mx}}$  using the DIC gradient measured at 3 m depth and likewise to compute the average  
 349 mixing layer irradiance.

350 At the nearby DyFAMed site (Dynamique des Flux Atmospheriques en Mediterranee), using  
 351 a 20-year O<sub>2</sub> time series, Coppola et al. (2018) estimated an annual net community carbon  
 352 production equal to 7.1 mol m<sup>-2</sup> yr<sup>-1</sup>, or 19.4 mmol m<sup>-2</sup> d<sup>-1</sup>. Additionally, using  
 353 determinations of primary production from <sup>14</sup>C in-situ incubations and of carbon export fluxes  
 354 from sediment traps at DyFAMed, Marty and Chiaverini (2002) calculated average daily  
 355 primary productions between 19.6 and 53 mmol m<sup>-2</sup> d<sup>-1</sup>. These estimates may have  
 356 underestimated the annual NCP values as they were calculated from monthly observations  
 357 while phytoplankton production varies on a daily timescale. Our DIC measurements highlight  
 358 the large short-term variability of carbon consumption over a few days in March-April when  
 359 bloom-triggering atmospheric conditions are met, such as a seasonal light increase and the  
 360 availability of nutrients.

	T <sub>0</sub>	Wind stress N m <sup>-2</sup>	Net heat flux W m <sup>-2</sup>	PAR mol photons m <sup>-2</sup> d <sup>-1</sup>	Z <sub>mixing</sub> m	Z <sub>Euphotic</sub> m	Irradiance mol photons m <sup>-2</sup> d <sup>-1</sup>
<b>2016</b>	March 18	0.04 +/-0.02	56+/-36	37+/-3	20+/-9	47	18
<b>2017</b>	March 9	0.12+/-0.02	45+/-21	30+/-3	28+/-12	71	13
<b>2018</b>	April 5	0.30+/-0.24	127+/-12	42+/-6	62+/-9	29	
<b>2019</b>	March 30	0.10+/-0.02	183+/-9	53+/-2	20+/-5	43	25

361

362 Table 2. 2016-2019. Atmospheric drivers of the onset of the bloom: average values and  
 363 standard deviation on the day of the start of the bloom, T<sub>0</sub>.

364

### 365 5- Concluding remarks

366 Around March and April in the northwestern Mediterranean Sea, the seasonal change in  
 367 lighting conditions and atmospheric forcing are the essential parameters that control the  
 368 triggering of near-surface phytoplankton blooms. These mechanisms have been highlighted  
 369 through the coupling of high-frequency physical and biogeochemical in situ observations. For  
 370 years 2016 to 2019, a bloom started in March or early April identified by a decrease in surface  
 371 DIC, following a period of intense mixing. The rapid decrease in surface wind speed entailed  
 372 a switch from negative to positive air-sea net heat fluxes, essentially due to changes in the



373 latent air-sea heat flux proportional to wind speed. These analysis support the hypothesis that  
374 decreases in the depth of active mixing, a result of the transition from buoyancy-driven to  
375 wind-driven mixing, control the timing of the surface spring bloom.

376 We have shown that the onset of surface phytoplankton growth, identified in 2016 by a  
377 simultaneous initial decrease in DIC and an increase in surface Chla, precedes by a few days  
378 the surface chlorophyll maximum as detected by satellite measurements with a binning period  
379 of 8 days. This time span does not allow to identify precisely the contribution of atmospheric  
380 drivers to trigger the onset of the growth of phytoplankton as it occurs on a daily basis.

381

382

383 Data availability

384 BOUSSOLE data (2016-2019) are available in the SEANOE data base (Merlivat Liliane,  
385 Boutin Jacqueline (2020). Mediterranean Sea surface CO<sub>2</sub> partial pressure and temperature  
386 data. SEANOE. <https://doi.org/10.17882/56709>

387 Azur Meteorological buoy: <http://mistrals.sedoo.fr>

388 SChl, 8-day, 4 km x4 km resolution, level 3 mapped ocean color product distributed by the  
389 European Space Agency, available at <http://www.ocean colour/org>

390 All glider data is archived at the British Oceanographic Data Centre  
391 (BODC, [https://www.bodc.ac.uk/data/bodc\\_database/gliders/](https://www.bodc.ac.uk/data/bodc_database/gliders/) ).

392 Author contributions: VV and MG were respectively in charge of the BOUSSOLE mooring  
393 buoy maintenance and monthly water sampling. LB was responsible for the laboratory  
394 preparation and calibration of the CARIOCA sensors. GAL led the UEA glider mission in  
395 March-April 2016. DA and VV provided data and funding through the BOUSSOLE project  
396 and contributed to writing and editing of the manuscript.

397 The authors declare that they have no conflict of interest.

398 Acknowledgments: Michael Hemming was a PhD student at the University of East Anglia  
399 and Sorbonne University under the supervision of Prof. Jan Kaiser, Prof. Karen J. Heywood,  
400 Dr Dorothee Bakker and Dr Jacqueline Boutin, funded by the Defence, Science and

401 Technology Laboratory (Contract no. DSTLX1000092277) with cooperation with Direction  
402 Générale de l'Armement (DGA)-. The assistance of the UEA Glider Group in keeping the  
403 glider flying is also gratefully acknowledged. The BOUSSOLE time series project is funded  
404 by the Centre National d'Etudes Spatiales (CNES) and the European Space Agency  
405 (ESA/ESRIN contract 4000119096/17/I-BG). The authors acknowledge Météo-France for  
406 supplying the data from the Azur buoy and the HyMeX database teams (ESPRI/IPSL and  
407 SEDOO/Observatoire Midi-Pyrénées) for their help in accessing their data. Crew and  
408 Captains of R/V Tethys II are warmly thanked for assistance with operations at sea.

409

#### 410 References

411 Álvarez, M., Sanleón-Bartolomé, H., Tanhua, T., Mintrop, L., Luchetta, A., Cantoni, C.,  
412 Schroeder, K., and Civitarese, G.: The CO<sub>2</sub> system in the Mediterranean Sea: a basin wide  
413 perspective, *Ocean Sci.*, 10, 69–92, <https://doi.org/10.5194/os-10-69-2014>, 2014.

414 Andersen, V. and Prieur, L.: One-month study in the open NW Mediterranean Sea  
415 (DYNAPROC experiment, May 1995): Overview of hydrobiogeochemical structures and  
416 effects of wind events, *Deep-Sea Res. I*, 47, 397–422, 2000.

417 Antoine, D. M. Chami, H. Claustre, F. D'Ortenzio, A. Morel, G. Bécu, B. Gentili, F. Louis, J.  
418 Ras, E. Roussier, A.J. Scott, D. Tailliez, S. B. Hooker, P. Guevel, J.-F. Desté, C. Dempsey  
419 and D. Adams. 2006, BOUSSOLE : a joint CNRS-INSU, ESA, CNES and NASA Ocean  
420 Color Calibration and Validation Activity. NASA Technical memorandum, N° TM-2006-  
421 214147, NASA/GSFC, Greenbelt, USA.

422 Antoine, D., P. Guevel, J.-F. Desté, G. Bécu, F. Louis, A.J. Scott and P. Bardey: The  
423 «BOUSSOLE» buoy; a new transparent-to-swell taut mooring dedicated to marine optics:  
424 design, tests and performance at sea, *Journal of Atmospheric and Oceanic Technology*, 25,  
425 968-989, 2008a.

426 Antoine, D., F. d'Ortenzio, S. B. Hooker, G. Bécu, B. Gentili, D. Tailliez, and A. J. Scott:  
427 Assessment of uncertainty in the ocean reflectance determined by three satellite ocean color  
428 sensors (MERIS, SeaWiFS and MODIS-A) at an offshore site in the Mediterranean Sea  
429 (BOUSSOLE project), *Journal of Geophysical Research*, 113(C7), 2008b.

430 Begovic , M., and C. Copin-Montegut, Processes controlling annual variations in the partial  
431 pressure of fCO<sub>2</sub> in surface waters of the central northwestern Mediterranean sea (Dyfamed  
432 site), *Deep-Sea Research II*, 49, 2031-2047, 2002

433 Behrenfeld, M. J, Abandoning Sverdrup's critical depth hypothesis on phytoplankton blooms.  
434 *Ecology*, 91(4), 977-989, 2010

435 Behrenfeld, M. and E. Boss, Resurrecting the ecological underpinnings of ocean plankton  
436 blooms. *Annual Review of Marine Science*, 6, 167-194, DOI: 10.1146/annurev-marine-  
437 052913-021325, 2014

438 Brainerd, K. E., and Gregg, M. C, Surface mixed and mixing layer depths. *Deep Sea Research*  
439 *I: Oceanographic Research Papers*, 42:1521–1543, 1995

440 Brody, S., and Lozier, M. Changes in dominant mixing length scale as a driver of  
441 phytoplankton bloom initiation in the North Atlantic, *Geophysical Research Letters* 41, 3197–  
442 3206, 2014.

443 Brody, S. R., & Lozier, M. S. , Characterizing upper-ocean mixing and its effect on the spring  
444 phytoplankton bloom with in situ data. *ICES Journal of Marine Science*, 72(6), 1961-1970,  
445 2015.

446 Chiswell, S. M.: Annual cycles and spring blooms in phytoplankton: don't abandon Sverdrup  
447 completely. *Marine Ecology Progress Series*, 443, 39-50, 2011.

448 Copin-Montégut, C., Begovic, M.: Distributions of carbonate properties and oxygen along the  
449 water column (0– 2000 m) in the central part of the NW Mediterranean Sea (Dyfamed site).  
450 Influence of winter vertical mixing on air– sea CO<sub>2</sub> and O<sub>2</sub> exchanges. *Deep-Sea Res., Part 2*,  
451 *Top. Stud. Oceanogr.* 49, 2049– 2066, 2002.

452 Copin-Montégut, C., M. Bégovic, and L. Merlivat.: Variability of the partial pressure of CO<sub>2</sub>  
453 on diel to annual time scales in the Northwestern Mediterranean Sea, *Mar Chem*, 85(3-4),  
454 169-189, 2004.

455 Coppola, L., Legendre, L., Lefevre, D., Prieur, L., Taillandier, V., & Riquier, E. D. :  
456 Seasonal and inter–annual variations of dissolved oxygen in the northwestern Mediterranean  
457 Sea (DYFAMED site). *Progress in Oceanography*, 2018.

458 Dickson, A. G., and F. J. Millero : A comparison of the equilibrium constants for the  
459 dissociation of carbonic acid in seawater media, *Deep Sea Research Part A. Oceanographic*  
460 *Research Papers*, 34(10), 1733-1743 1987.

461 Enriquez, R.M., Taylor, J.R: Numerical simulations of the competition between wind-driven  
462 mixing and surface heating in triggering spring phytoplankton blooms. *ICES J. Mar. Sci. J. du*  
463 *Cons. fsv071*, 2015.

464 Golbol, M., Vellucci, V., Antoine, D : BOUSSOLE, <https://doi.org/10.18142/1>, 2000.

465 Hemming, M. P., Kaiser, J., Heywood, K. J., Bakker, D. C., Boutin, J., Shitashima, K., Onken,  
466 R. : Measuring pH variability using an experimental sensor on an underwater glider. *Ocean*  
467 *Science*, 13 , 427–442,2017.

468 Holte, J., & Talley, L. : A new algorithm for finding mixed layer depths with applications to  
469 Argo data and Subantarctic Mode Water formation. *Journal of Atmospheric and Oceanic*  
470 *Technology*, 26(9), 1920-1939, 2009.

471 Hood, E. M., and L. Merlivat : Annual and interannual variations of fCO<sub>2</sub> in the  
472 northwestern Mediterranean Sea: Results from hourly measurements made by CARIOCA  
473 buoys, 1995-1997, *J Mar Res*, 59, 113-131, 2001.

474 Lacour, L., Briggs, N., Claustre, H., Ardyna, M., & Dall'Olmo, G.: The intraseasonal  
475 dynamics of the mixed layer pump in the subpolar North Atlantic Ocean: A  
476 Biogeochemical - Argo float approach. *Global Biogeochemical Cycles*, 33(3), 266–281,2019.

477 Lee, Z., Weidemann, A., Kindle, J., Arnone, R., Carder, K. L., & Davis, C. : Euphotic  
478 zone depth: Its derivation and implication to ocean-color remote sensing. *Journal of*  
479 *Geophysical Research: Oceans* , 112 (C3),2007.

480 Mahadevan, A., D'asaro, E., Lee, C., & Perry, M. J. : Eddy-driven stratification initiates  
481 North Atlantic spring phytoplankton blooms. *Science*, 337(6090), 54-58, 2012.

482 Marty, J.-C., Chiavérini, J. : Seasonal and interannual variations in phytoplankton  
483 production at DYFAMED time-series station, northwestern Mediterranean Sea. *Deep*  
484 *Sea Res. Part II* 49, 2017–2030, 2002.

485 Mehrbach, C., C. H. Culberson, J. E. Hawley, and R. M. Pytkowicz : Measurement of the  
486 apparent dissociation constants of carbonic acid in seawater at atmospheric pressure, *Limnol*  
487 *Oceanogr*, 18(6), 897-907, 1973.

488 Merlivat, L., and Brault, P.: CARIOCA BUOY, Carbon Dioxide Monitor, *Sea Technol.*, 23–  
489 30, 1995.

490 Merlivat, L., Boutin, J., & Antoine, D. : Roles of biological and physical processes in driving  
491 seasonal air–sea CO<sub>2</sub> flux in the Southern Ocean: New insights from CARIOCA pCO<sub>2</sub>.  
492 *Journal of Marine Systems*, 147, 9-20, 2015.

493 Merlivat, L., Boutin, J., Antoine, D., Beaumont, L., Golbol, M., & Vellucci, V. : Increase of  
494 dissolved inorganic carbon and decrease of pH in near surface waters of the Mediterranean  
495 Sea during the past two decades. *Biogeosciences*, 15(18), 5653-5662, 2018.

496 Millot: Circulation in the Western Mediterranean Sea, *Journal of Marine Systems*, 20, 423–  
497 442, 1999.

498 Morel, A., and J. F. Berthon . : Surface pigments, algal biomass profiles, and potential  
499 production of the euphotic layer: relationships reinvestigated in review of remote-sensing  
500 applications, *Limnol. Oceanogr.*, 34, 1545–1562, 1989.

501 Niewiadomska, Katarzyna, Claustre, Hervé, Prieur, Louis, d'Ortenzio, Fabrizio, (2008),  
502 Submesoscale physical - biogeochemical coupling across the Ligurian current (northwestern  
503 Mediterranean) using a bio - optical glider, *Limnology and Oceanography*, 53, doi:  
504 10.4319/lo.2008.53.5\_part\_2.2210.

505 Papaioannou, G., Papanikolaou, N., and Retalis, D. : Relationships of photosynthetically  
506 active radiation and shortwave irradiance. *Theoretical and Applied Climatology*, 48: 23–27,  
507 1993.

508 Pasqueron de Fommervault, O., Migon, C., D'Ortenzio, F., Ribera d'Alcalà, M., and Coppola,  
509 L.: Temporal variability of nutrient concentrations in the northwestern Mediterranean sea  
510 (DYFAMED time-series station), *Deep-Sea Res. Pt. I*, 100, 1–12, 2015.

511 Pellichero, V., Boutin, J., Claustre, H., Merlivat, L., Sallée, J.-B., & Blain, S. : Relaxation of  
512 wind stress drives the abrupt onset of biological carbon uptake in the Kerguelen bloom: a  
513 multisensor approach. *Geophysical Research Letters*, 47, e2019GL085992

514 <https://doi.org/10.1029/2019GL085992>, 2020.

515 Rödenbeck, C., Keeling, R. F., Bakker, D. C. E., Metzl, N., Olsen, A., Sabine, C., and  
516 Heimann, M.: Global surface-ocean pCO<sub>2</sub> and sea–air CO<sub>2</sub> flux variability from an  
517 observation- driven ocean mixed-layer scheme, *Ocean Sci.*, 9, 193–216, doi:10.5194/os-9-  
518 193-2013, 2013.

519 Rumyantseva, A, Henson, S, Martin, A , Thompson, A. F, Damerell G.M, Kaiser J, Heywood,  
520 K.J.: Phytoplankton spring bloom initiation: The impact of atmospheric forcing and light in  
521 the temperate North Atlantic Ocean, *Progress in Oceanography*, 2019.

522 Siegel, D. A., S. C. Doney, and J. A. Yoder (2002), The North Atlantic spring phytoplankton  
523 bloom and Sverdrup’s critical depth hypothesis, *Science*, 296, 730– 733.

524 Sverdrup, H. U.: On vernal blooming of phytoplankton. *Conseil Exp. Mer*, 18, 287-295,1953.

525 Takahashi, T., Sutherland, S. C., Wanninkhof, R., Sweeney, C., Feely, R. A., Chipman, D. W.,  
526 Hales, B., Friederich, G., Chavez, F., Sabine, C., Watson, A., Bakker, D. C. E., Schuster, U.,  
527 Metzl, N., Yoshikawa-Inoue, H., Ishii, M., Midorikawa, T., Nojiri, Y., Kortzinger, A.,  
528 Steinhoff, T., Hoppema, M., Olafsson, J., Arnarson, T. S., Tillbrook, B., Johannessen, T. and  
529 Olsen, A., Bellerby, R., Wong, C. S., Delille, B., Bates, N. R., and de Baar, H. J. W.:  
530 Climatological mean and decadal change in surface ocean pCO<sub>2</sub> and net sea-air CO<sub>2</sub> flux  
531 over the global oceans, *Deep-Sea Res. II*, 56, 554–577, 2009.

532 Taylor, J. R., & Ferrari, R.: Shutdown of turbulent convection as a new criterion for the onset  
533 of spring phytoplankton blooms. *Limnology and Oceanography*, 56(6), 2293-2307, 2011.

534 Venables, H. and Moore, C.M.: Phytoplankton and light limitation in the Southern Ocean:  
535 Learning from high nutrient, high chlorophyll areas. *Journal of Geophysical Research:*  
536 *Oceans*, 115(C2), 2010.

537 Wanninkhof, R.: Relationship between wind speed and gas exchange over the ocean revisited.  
538 *Limnology and Oceanography: Methods*, 12(6), 351–362, 2014.

539 Weiss, R.: Carbon dioxide in water and seawater: the solubility of a non-ideal gas, *Marine*  
540 *chemistry* , 2 (3), 203–215, 1974

541

542

543 Figures and tables

544 Figure 1. The area of the northwestern Mediterranean Sea showing the French and Italian  
545 Rivièras, the island of Corsica, the main current branches (green arrows) in the Ligurian Sea  
546 and the location of the BOUSSOLE buoy (white square) and the AZUR meteorological buoy  
547 (yellow square). The black line indicates the glider's path in March-April 2016.

548 Figure 2. 2016 (a) Hourly DIC (green) and SST (purple) at the BOUSSOLE site, (b) hourly  
549 wind stress (blue) and net heat flux (pink) at the Azur buoy site. The pink dotted line indicates  
550 the change of sign of the net heat flux from negative to positive values, (c) daily depths of the  
551 mixed layer (black), mixing layer (dotted black), Ekman layer (cyan) and euphotic layer  
552 (orange) at the BOUSSOLE site. The black dotted line on panels (a), (b), (c) (d) and (e)  
553 indicates the onset time of the bloom (18 March). (d) PAR (orange) and average mixing layer  
554 irradiance (blue), (e) Glider-derived depths of the mixed layer (black curve, same as in (c)),  
555 euphotic layer (orange), and nighttime 2-day binned chlorophyll concentration (green  
556 coloring) retrieved between March 7 and April 5.

557 Figure 3. (a, b, c) 2017, 2018 and 2019 DIC (green) and SST (purple) at the BOUSSOLE site,  
558 (d, e, f) wind stress (blue) and net heat flux (red) at the Azur buoy site. The red dotted line  
559 indicates the change of sign of the net heat flux from negative to positive values, (g, h, i)  
560 depths of the mixing layer (dotted black), Ekman layer (cyan) and euphotic layer (orange) at  
561 the BOUSSOLE site. The vertical dotted green line indicates the onset of the bloom.

562 Figure 4. Changes of physical parameters (hourly values) at the onset of the 2016 bloom  
563 during 2 consecutive periods of 4 days, March 14-17 (blue) and March 18-21 (red) as a  
564 function of wind stress (a) net surface heat flux (b) depths of the Ekman, mixing and euphotic  
565 layer. Grey circles and grey squares indicate mean values respectively on March 17 and  
566 March 18. For comparison, the orange line shows the euphotic layer depth (March 14-21).

567 Figure 5. From March 7 to April 5, (a) DIC and Chla. The purple line and the squares  
568 (blue, morning; grey, evening) indicate the 3 days biological diurnal DIC changes during the  
569 period considered to compute NCP. The blue and orange lines indicate the surface Chla when  
570 the glider was at a distance of less than 5 km (blue) and less than 20 km (orange) respectively  
571 from the BOUSSOLE buoy. (b) PAR and I average mixing layer irradiance. The vertical  
572 dotted black line indicates the onset of the bloom on March 18.

573 Table 1. Net community production computed in the mixing layer over 3 days (NCP) after the  
574 bloom onset ( $T_0$ ). (a, b) variation of DIC at the measurement depth and linked correlation  
575 coefficient; (c, d) change of integrated DIC over  $Z_{mx}$  and linked correlation coefficient ;(e)  
576  $CO_2$  flux from the atmosphere to the ocean; (f), absolute value of calculated NCP is  
577 considered in order to provide positive values throughout the presentation and discussion in  
578 the manuscript.

579 Table 2. 2016-2019. Atmospheric drivers of the onset of the bloom: average values and  
580 standard deviation on the day of the start of the bloom,  $T_0$ .

581