

1   **Title:** Regulation of a surface chlorophyll hotspot by wind-driven upwelling and eddy circulation  
2   in the Santa Barbara Channel, Southern California

3

4   **Authors:** Rachel D. Simons<sup>1\*</sup> and Dylan Catlett<sup>2</sup>

5

6   <sup>1</sup> Earth Research Institute, University of California, Santa Barbara, CA, USA

7   <sup>2</sup> Woods Hole Oceanographic Institution, Wood Hole, MA, USA

8   \* Corresponding Author. Address: Email address: [simons@eri.ucsb.edu](mailto:simons@eri.ucsb.edu)

9

10   **Keywords:** upwelling, eddies, chlorophyll, phytoplankton, ROMS, nitrogen

11

12     **Abstract**  
13

14     While most of the U.S. Pacific coast is dominated by strong wind-driven upwelling in the spring  
15     and regular seasonal cycles of phytoplankton primary production, the Southern California Bight  
16     has weak intermittent wind-driven upwelling and low phytoplankton concentrations due to its  
17     irregular sheltered coastline. However, the Santa Barbara Channel (SBC), located in the  
18     northern Southern California Bight, contains an anomalous hotspot of phytoplankton biomass.  
19     We use 3D ocean circulation and particle tracking models, an empirical temperature-nitrate  
20     relationship, and satellite observations of surface chlorophyll from 1998 to 2007 to determine  
21     how wind-driven upwelling and cyclonic eddy circulation govern phytoplankton dynamics in the  
22     SBC. Our findings show that elevated surface chlorophyll in the spring is driven by the coupling  
23     of wind-driven upwelling and cyclonic eddy circulation and requires the presence of both high  
24     nitrate from wind-driven upwelling and prolonged residence times from eddy circulation. Long  
25     residence times, created by persistent cyclonic eddy circulation, allow nitrate transported into  
26     the surface layer by wind-driven upwelling to be retained in the SBC long enough to meet the  
27     required timescales for phytoplankton nutrient uptake and accumulation, overcoming the  
28     typically weak wind-driven upwelling of the Southern California Bight. Typical spring upwelling  
29     periods (SUPs) have high levels of surface chlorophyll, which are produced by the balanced  
30     coupling of upwelling strength and cyclonic eddy circulation. For two years of our study, 1998  
31     and 1999, the SUP did not correlate with elevated surface chlorophyll. Corresponding to a  
32     strong El Niño event, the 1998 SUP had exceptionally weak wind-driven upwelling, which did  
33     not produce sufficient levels of nitrate to stimulate elevated surface chlorophyll. For the 1999  
34     SUP, a strong La Niña event produced unusually strong wind-driven upwelling, which resulted in  
35     very high levels of nitrate, but suppressed eddy circulation and residence times to the point  
36     where phytoplankton biomass could not accumulate in the SBC. These anomalous SUPs  
37     illustrate that when wind-driven upwelling is too strong or weak, the balance between wind-  
38     driven upwelling and cyclonic eddy circulation is disrupted, resulting in a dramatic reduction of

39 surface chlorophyll, which may become more frequent with climate-driven upwelling changes in  
40 the future.

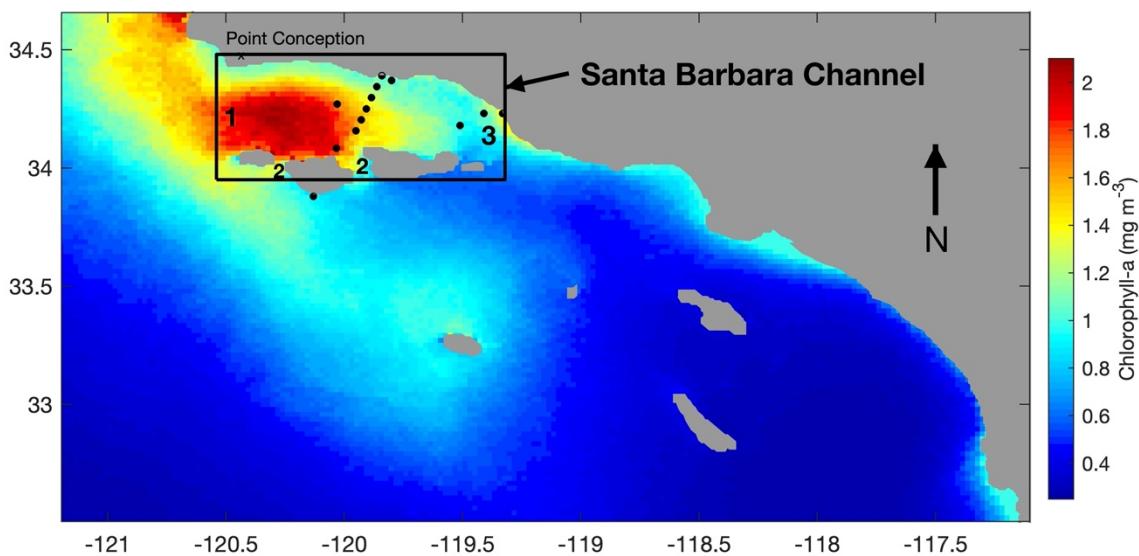
41     **1.0 Introduction**

42         While continental margins contain only a fraction of the world's oceans, they have been  
43         estimated to produce 10-15% of the global primary production (Thunell et al. 2007; Fennel and  
44         Wilkin 2009; Levin et al. 2015). Eastern boundary upwelling systems (EBUSs), which include  
45         the California EBUS, are among the most productive continental margin ecosystems in the  
46         world and are of particular concern as they are being increasingly impacted from climate  
47         change, the harvesting of marine resources, and coastal development (Levin et al. 2015).  
48         Encompassing the west coast of the United States, the California EBUS is one of the largest  
49         EBUSs in the world and is defined by a seasonal cycle of primary production. Wind-driven  
50         upwelling in the California EBUS is strongest in the spring and early summer when alongshore  
51         winds drive offshore Ekman transport and nutrients are vertically transported into the surface  
52         layer stimulating primary production (Legaard and Thomas 2006; Henson and Thomas 2007;  
53         García-Reyes and Largier 2012). Messie and Chavez (2015) found that in addition to nutrient  
54         supply, physical export of nutrients is an important regulator of primary production in the  
55         California EBUS. Physical export can limit primary production if nutrients introduced to the  
56         surface layer are transported off the shelf before uptake by phytoplankton can occur. Thus, to  
57         understand the variability of primary production in EBUSs, the physical processes that retain  
58         nutrients in the surface layer on the shelf need to be understood (Lachkar and Gruber 2011;  
59         Messié and Chavez 2015).

60         Located in the southern region of the California EBUS is the Southern California Bight  
61         (SC Bight, Fig. 1), which encompasses the southern California coast and the eight Channel  
62         Islands. While most of the California EBUS is dominated by strong spring upwelling and regular  
63         seasonal cycles of primary production, the SC Bight has weak, intermittent upwelling due to its  
64         irregular, sheltered coastline, which results in low levels of primary production year round  
65         (Winant and Dorman 1997; Dorman and Winant 2000). However, the Santa Barbara Channel  
66         (SBC), located within the northern SC Bight, regularly contains anomalously high phytoplankton

67 biomass and primary productivity compared to the rest of the SC Bight (Fig. 1, Mantyla et al.  
68 1995; Henderikx Freitas et al. 2017; Santora et al. 2017). Due in part to this unusually  
69 productive environment, the SBC supports a uniquely biodiverse ecosystem of extensive kelp  
70 forests, intertidal habitats, and sandy beaches along with large populations of seabirds, marine  
71 mammals, fishes, and plankton (Beers et al. 1986; Fiedler et al. 1998; Miller et al. 2011; Dugan  
72 and Hubbard 2016).

73



74  
75 **Figure 1:** Mean surface chlorophyll-a concentrations ( $\text{mg m}^{-3}$ ) in the Southern California Bight  
76 from satellite data (Kahru et al. 2012, 2015, <http://spg-satdata.ucsd.edu>). Location of the Santa  
77 Barbara Channel (SBC) is identified by the black box. SBC boundaries are identified as 1-West,  
78 2-South, and 3-East. The solid black circles show the sample locations of the CalCOFI and  
79 UCSB Plumes and Blooms programs.

80  
81 The SBC is a 50 km wide, 100 km long, and 500 m deep basin, which is bounded to the  
82 north by the Southern California coast and the south by the Northern Channel Islands (Fig. 1).  
83 To the north of the SBC, the oceanographic regime is characterized by the cold, nutrient-rich  
84 waters of the southward flowing California Current (CC), which can extend up to 1,000 km  
85 offshore (Hickey 1993; Bray et al. 1999). To the south of the SBC, the oceanographic regime is  
86 dominated by the warm, nutrient-poor waters of the northward flowing Southern California

87 Countercurrent (SCC, Hickey 1993; Bray et al. 1999). The SBC is the transition zone where  
88 these two regimes meet, creating a dynamic physical and biological environment.

89 The two dominant circulation patterns associated with phytoplankton accumulation in the  
90 SBC are wind-driven upwelling and cyclonic eddy circulation (Anderson et al. 2006; Brzezinski  
91 and Washburn 2011; Matson et al. 2019). For this paper, wind-driven upwelling will be referred  
92 to as “upwelling”, and cyclonic eddy circulation as “eddy circulation”. Driven by persistent,  
93 steady winds from the west, upwelling occurs along the mainland coast of the SBC and is  
94 strongest in the spring and early summer (Winant and Dorman 1997; Harms and Winant 1998;  
95 Melton et al. 2009). Upwelling also occurs in the SBC in the late summer and early fall, but is  
96 frequently interrupted by wind reversals, which transport warm nutrient poor waters back  
97 towards the coastline (Washburn and McPhee-Shaw 2013). McPhee-Shaw et al. (2007)  
98 estimated that spring upwelling accounted for  $70\% \pm 15\%$  of the total annual nutrient delivery to  
99 the inner-shelf. During spring upwelling, large phytoplankton blooms, covering more than 50%  
100 of the SBC, are observed, but smaller blooms can occur at any time during the year (Otero and  
101 Siegel 2004; Santoro et al. 2010). Similar to the California EBUS, upwelling intensity in the SBC  
102 is affected by the El Niño Southern Oscillation (ENSO) with strong El Niño events suppressing  
103 upwelling and strong La Niña events enhancing it (Schwing et al. 2000; Jacox et al. 2018).

104 Eddy circulation is present periodically throughout the year in the western SBC and  
105 ranges in diameter from 5 to 40 km, at times penetrating down to 300 m. Eddy circulation  
106 occurs when a poleward pressure gradient balances equatorward wind stress (Harms and  
107 Winant 1998; Dever et al. 1998; Oey et al. 2004) and persists for an average of two weeks,  
108 ranging from a few days to months depending on ocean conditions (Nishimoto and Washburn  
109 2002; Beckenbach and Washburn 2004). Eddy circulation is also affected by the ENSO. When  
110 El Niño conditions cause atypically weak upwelling and wind relaxation, the equatorward wind  
111 stress balances the poleward pressure gradient for unusually long periods of time allowing eddy  
112 circulation to persist for months. In contrast, La Niña conditions cause unusually strong

113 equatorward wind stress, which dominates the poleward pressure gradient and does not allow  
114 stable eddy circulation to form (Nishimoto and Washburn 2002; Simons et al. 2015).

115 Based on a field study of water column measurements from 16 seasonal channel-wide  
116 cruises, Brzezinski and Washburn (2011) found that 80% of the variability in phytoplankton  
117 primary production was explained by the presence of upwelling and eddy circulation. Through  
118 their work along with Anderson et al. (2006), Krause et al. (2013), and Simons et al. (2015), the  
119 following hypotheses were proposed connecting upwelling and eddy circulation to the  
120 anomalously high primary production in the SBC. Nutrient-rich water is introduced into the  
121 surface layer of the SBC by three processes; (1) upwelling along the mainland coast, (2)  
122 advection of upwelled water from the west into the southern SBC by eddy circulation, and (3)  
123 eddy circulation through the uplift of isopycnals, also known as eddy pumping (McGillicuddy  
124 2016). After these processes occur, eddy circulation retains nutrient-rich water in the SBC by  
125 creating an environment of long residence times, allowing phytoplankton to accumulate.

126 In this study, we integrate a high-resolution 3D Regional Ocean Modeling System  
127 (ROMS) with satellite and field observations to determine the interactive roles of upwelling, eddy  
128 circulation, and nutrient transport in regulating mesoscale phytoplankton blooms in the SBC.  
129 The ROMS is used to identify the presence and strength of upwelling and eddy circulation in the  
130 SBC over a 10-year period from 1998-2007. Driven by the ROMS, a 3D particle tracking model  
131 is used to estimate residence time and the source location of water in the SBC. Surface  
132 chlorophyll-a concentrations from satellite data are used as a proxy for phytoplankton biomass.  
133 We compare mean decadal, annual, and interannual patterns of modeled upwelling, eddy  
134 circulation, nutrient availability, source water location, and residence times to surface  
135 chlorophyll-a concentrations.

136

## 137 **2.0 Methods**

### 138 ***2.1 Ocean circulation modeling and eddy detection***

139        The 3D ocean circulation model used for this study was a high-resolution Regional  
140    Ocean Modeling System (ROMS) applied to the SC Bight (Shchepetkin and McWilliams 2005;  
141    Dong and McWilliams 2007; Dong et al. 2009). Our analysis was conducted on 6-hr averaged  
142    offline solutions of velocity and temperature from the ROMS for 1998-2007. The model domain,  
143    shown in Fig. 1, contains the SC Bight, which includes the SBC. The model grid spans a  
144    latitude of 32°17' N to 34°45' N and a longitude of 117°06' W to 121°12' W with 1 km horizontal  
145    resolution and 40 vertical levels. The model has been validated for seasonal, annual, and  
146    interannual mesoscale circulation (Dong et al. 2009; Ohlmann and Mitarai 2010) and has  
147    successfully reproduced upwelling events in the SC Bight (Dong et al. 2011) and the interannual  
148    variability of eddy circulation in the SBC (Simons et al. 2015). As shown in Fig. 1, our study  
149    area is the SBC, which spans a latitude of 33°57' N to 34°29' N and a longitude of 119°20' W to  
150    120°34' W.

151        Since our goal is to determine how upwelling and eddy circulation regulate surface  
152    chlorophyll-a concentrations in the SBC, we focus on the SBC surface layer defined as the top  
153    30 m of the water column. This definition is based on depth measurements of the SBC euphotic  
154    zone of  $24\text{ m} \pm 9.5\text{ m}$  from Brzezinski and Washburn (2011) and observations by Krause et al.  
155    (2013), which showed that 30 m is generally near or below the base of the surface layer.  
156    Venrick (1998) also observed that chlorophyll in the SBC was consistently concentrated in the  
157    upper 25 m of the water column.

158        Using the ROMS, eddy circulation in the SBC was quantified using an eddy detection  
159    algorithm developed by Nencioli et al. (2010). This eddy detection algorithm was developed  
160    using vector geometry and identifies the center, shape, lifespan, and trajectory of all eddies in  
161    horizontal flow fields. In previous work, this algorithm has been successfully applied to the  
162    ROMS solutions used in the study (Nencioli et al. 2010; Dong et al. 2012; Simons et al. 2015)  
163    and to field observations of eddies in the South China Sea (Chen et al. 2011; Wang et al. 2021)

164 and the North Pacific Ocean (Liu et al. 2012). For this analysis, the algorithm was applied to  
165 horizontal velocity fields from the ROMS solutions that were depth-averaged over the surface  
166 layer, the top 30 m of the water column, and provided a dataset of eddy presence, location,  
167 size, and lifespan every six-hours for the 10-year modeling period of 1998-2007.

168 **2.2 Estimating residence time and source water zone with particle tracking**

169 To estimate the residence time and source location of water in the SBC, a 3D particle  
170 tracking model (PTM) was used. Driven by the stored flow fields from the ROMS, the PTM used  
171 tri-linear interpolation to identify the velocity of the particles within a grid cell and then moved the  
172 particles in time using a fourth-order accurate Adams-Bashforth-Moulton predictor-corrector  
173 method (Carr et al. 2008; Mitarai et al. 2009). Particles were released daily throughout the SBC  
174 surface layer horizontally on a 1 km<sup>2</sup> grid and vertically every 5 m from 5 m to 30 m below the  
175 surface. Particles were tracked passively forward and backwards in time in separate simulations  
176 for 60 days, resulting in 54,744 particles released daily and a total of 199.8 million particles  
177 tracked for the 10-year modeling period.

178 The forward particle tracking simulations were used to calculate residence time in the  
179 SBC. First, the horizontal location and tracking time for all particles released over the previous  
180 60 days were identified for each day of the 10-year modeling period. Then, the mean tracking  
181 time of the particles located within each grid cell (1 km<sup>2</sup>) was calculated for all grid cells in the  
182 SBC. This method created a 2D distribution of mean tracking times or residence time (RT) for  
183 each day of the modeling period where RT represented the mean length of time that particles  
184 remained within a grid cell. The residence time within the eddy (RT<sub>eddy</sub>) was determined using  
185 the eddy detection scheme to identify the horizontal area of the eddy and then averaging the RT  
186 over the eddy's area.

187 The reverse particle tracking simulations were used to calculate the percentage of water  
188 in the SBC surface layer coming from four different source water zones: West, East, South, and  
189 SBC. The source water zone of a particle was defined by the boundary, West, East, or South

190 (Fig. 1), where the particle first entered the SBC. The particles that entered the SBC through  
191 the West boundary were assumed to originate from the CC and through the East boundary were  
192 assumed to originate from the SCC. The particles that remained in the SBC during the 60-day  
193 reverse tracking time were identified as originating from the SBC. Particles that entered the  
194 SBC through the South boundary could have come from either the CC or the SCC. These  
195 calculations provided the daily percentage of water in the SBC originating from each of the four  
196 source water zones.

## 197 ***2.2 Satellite observations of surface chlorophyll***

198 To quantify the spatial and temporal patterns of chlorophyll concentrations in the SBC,  
199 satellite-derived five-day composites of surface chlorophyll-a concentrations (Chl) with 1 km<sup>2</sup>  
200 resolution from 1998-2007 were used. The Chl data set was derived from three ocean color  
201 sensors, SeaWiFS, MODIS-Aqua, MODIS-Terra, and MERIS, assembled by the Scripps  
202 Photobiology Group at U.C. San Diego (Kahru et al. 2012, 2015, <http://spg-satdata.ucsd.edu>)  
203 and has been used in many studies of the California Current System (e.g. Kahru et al. 2012;  
204 McClatchie et al. 2016; Jacox et al. 2016b). Chl was used as a proxy for phytoplankton biomass  
205 in this study. While known biases can arise in the relationships between Chl and phytoplankton  
206 carbon biomass due to variability in phytoplankton physiological status, it is generally thought  
207 that phytoplankton biomass is the first-order determinant of Chl variability in productive coastal  
208 regions like the SBC (Behrenfeld et al. 2005; Siegel et al. 2013).

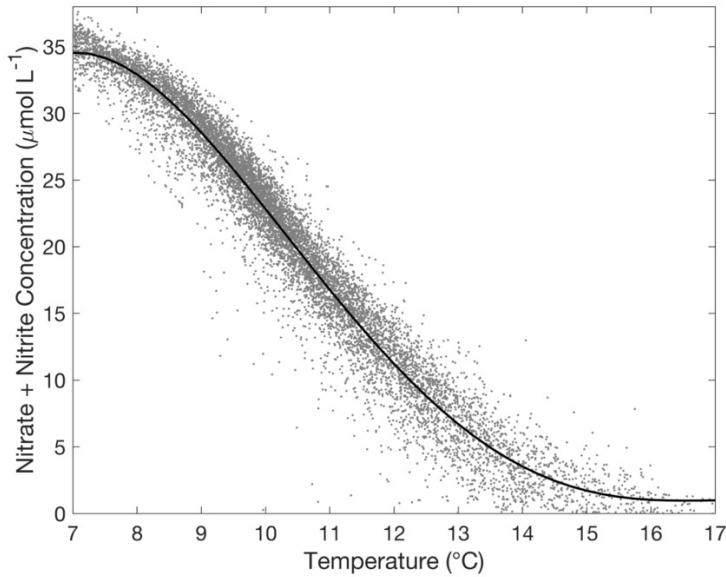
## 209 ***2.3 Estimating nutrient concentrations***

210 The concentrations of macronutrients (nitrate + nitrite, phosphate, and silicate) tend to  
211 be highly correlated in the California EBUS, and nitrogen is typically the limiting macronutrient of  
212 phytoplankton production in this region (Messié and Chavez 2015; Deutsch et al. 2021). Many  
213 previous studies have shown a stable relationship between temperature and inorganic nitrogen  
214 in the SBC and SC Bight (Hayward and Venrick 1998; McPhee-Shaw et al. 2007; Omand et al.  
215 2012; Snyder et al. 2020). In the California EBUS, prior work has shown that temperature is an

216 adequate predictor of nitrate in areas, like the SBC, that are located within the same latitude and  
217 50 km of the shoreline (Palacios et al. 2013; Jacox et al. 2018). Thus, the maximum available  
218 nitrate + nitrite concentrations ( $N_{max}$ ) in the SBC surface layer were calculated using an  
219 empirical temperature to nitrate + nitrite relationship derived from water samples collected in the  
220 SBC and the 3D temperature solutions from the ROMS. Water sample data came from the  
221 CalCOFI (calcofi.org) and the UCSB Plumes and Blooms program (Fig. 1, Catlett et al. 2021).  
222 To avoid artifacts in the relationship due to nutrient uptake by phytoplankton, only water  
223 samples collected from depths below the surface layer, deeper than 30 m, were used. The  
224 empirical temperature to nitrate + nitrite relationship or  $N_{max}$  model was derived using 10,311  
225 water samples collected at 14 stations in the SBC from depths of 35 m to 517 m over 1980-  
226 2018 (Fig. 1). To calculate  $N_{max}$  from ROMS temperature, the following fourth-order polynomial  
227 using a least squares fit was derived from the water sample data (Fig. 2):

$$228 N_{max} = -0.0108T^4 + 0.5899T^3 - 11.3447T^2 + 87.2050T - 196.3239$$

229 where  $N_{max}$  is the maximum available nitrate + nitrite concentration ( $\mu\text{mol L}^{-1}$ ) and  $T$  ( $^{\circ}\text{C}$ ) is  
230 temperature in the surface layer, the top 30 m of the water column. To create a 3D  $N_{max}$  data set  
231 for the SBC surface layer,  $N_{max}$  was calculated from ROMS temperature horizontally at each  
232 grid cell and vertically every meter from the surface to 30 m depth every 6-hr for the 10-year  
233 modeling period, 1998-2007.  $N_{max}$  in the eddy was determined using the eddy detection  
234 scheme to identify the horizontal eddy area and then averaging  $N_{max}$  horizontally and vertically  
235 over the eddy area.



236  
 237 **Figure 2:** Nitrate + nitrite concentrations ( $\mu\text{mol L}^{-1}$ ) vs. temperature ( $^{\circ}\text{C}$ ) from field samples of  
 238 UCSB Plumes and Bloom and CalCOFI programs. Black line shows polynomial fit used to  
 239 model  $N_{\text{max}}$ .

240  
 241

242 **2.4 Identifying  $N_{\text{max}}$  transport mechanisms**

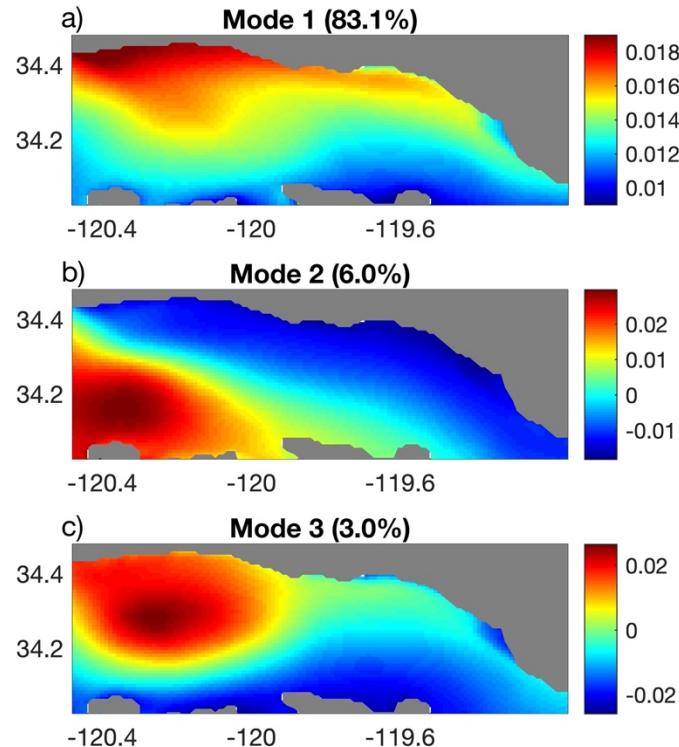
243 Upwelling and eddy circulation are the primary processes associated with increased  
 244 nutrients and Chl in the SBC (Anderson et al. 2006; McPhee-Shaw et al. 2007; Brzezinski and  
 245 Washburn 2011). The strength of upwelling is usually quantified in EBUSs using surface wind  
 246 stress or upwelling indices, which are a function of surface wind stress. However, the wind data  
 247 used to force the ROMS was not available with the offline solutions of the SC Bight. Thus, to  
 248 determine the presence and strength of the mechanisms transporting  $N_{\text{max}}$  into the SBC surface  
 249 layer, Empirical Orthogonal Function (EOF) analysis was used on 2D depth-averaged  $N_{\text{max}}$  over  
 250 the 10-year modeling period. The EOF method is a linear statistical method that identifies the  
 251 spatiotemporal variability within a dataset by transforming the dataset into a set of empirical  
 252 orthogonal functions or modes. The patterns of these modes can then be correlated with  
 253 dynamic mechanisms in the ocean (Thomson and Emery 2014). The advantage of this method  
 254 was that only mechanisms that were strong enough to transport elevated  $N_{\text{max}}$  into the surface  
 255 layer were identified. This is particularly useful for the SBC where field studies have shown that

256 while upwelling favorable winds tend to be most persistent in the spring, they occur  
257 intermittently all year round in the SBC (Dorman and Winant 2000; Oey et al. 2001). The first  
258 EOF mode explained 83.1% of the  $N_{max}$  variability. Representing a pattern of wind-driven  
259 upwelling, the mode 1 map (Fig. 3(a)) displayed a strong north-south gradient with the highest  
260 positive values along the mainland coast and lowest values along the north coast of the islands.  
261 Along the mainland coast, a west-east gradient is shown with higher positive values that extend  
262 farther from shore in the western than the eastern SBC, indicating that upwelling may be  
263 stronger in the western SBC. The second and third modes are associated with eddy circulation.  
264 The second EOF mode explained 6.0% of the  $N_{max}$  variability. Representing advection of  
265 elevated  $N_{max}$  water into the south-western SBC by eddy circulation, the mode 2 map (Fig. 3(b))  
266 displayed the highest positive values in the south-western SBC with negative values along the  
267 mainland coast and eastern SBC. The third EOF mode explained 3.0% of the  $N_{max}$  variability.  
268 Representing retention of elevated  $N_{max}$  by eddy circulation in the western SBC, positive values  
269 in the mode 3 map (Fig. 3(c)) show a circular shape in the western SBC with negative values in  
270 the eastern SBC. By analyzing horizontal distributions of phytoplankton primary production  
271 (PPP) during seasonal channel-wide cruises from 2001-2006, Brzezinski and Washburn (2011)  
272 found that wind-driven upwelling was the primary process driving high PPP in the SBC and eddy  
273 circulation was the secondary process, supporting our results.

## 274 **2.5 Analysis**

275 Regulation of Chl in the SBC by upwelling and eddy circulation was assessed by  
276 comparing satellite observations of Chl to modeled parameters of  $N_{max}$ , RT,  $RT_{eddy}$ , source water  
277 zone, eddy size and lifespan, and  $N_{max}$  EOF modes. Physical controls on the spatiotemporal  
278 distribution of Chl in the SBC were investigated across the decadal, annual, interannual, and  
279 synoptic timescales. Where time-series of channel-wide mean values are presented, Chl and  
280 modeled parameters were horizontally averaged over the SBC and, when applicable, depth-  
281 averaged over the surface layer, the top 30 m of the water column. Horizontal distributions of

282 modeled parameters were computed by depth-averaging over the surface layer. Where direct  
283 comparisons of Chl and modeled parameters are presented, Chl was interpolated to match the  
284 horizontal resolution of modeled parameters or modeled parameters were averaged over the 5-  
285 day Chl composites.



286

287 **Figure 3:**  $N_{\max}$  Empirical Orthogonal Function maps for (a) mode 1, (b) mode 2, and (c) mode 3

288

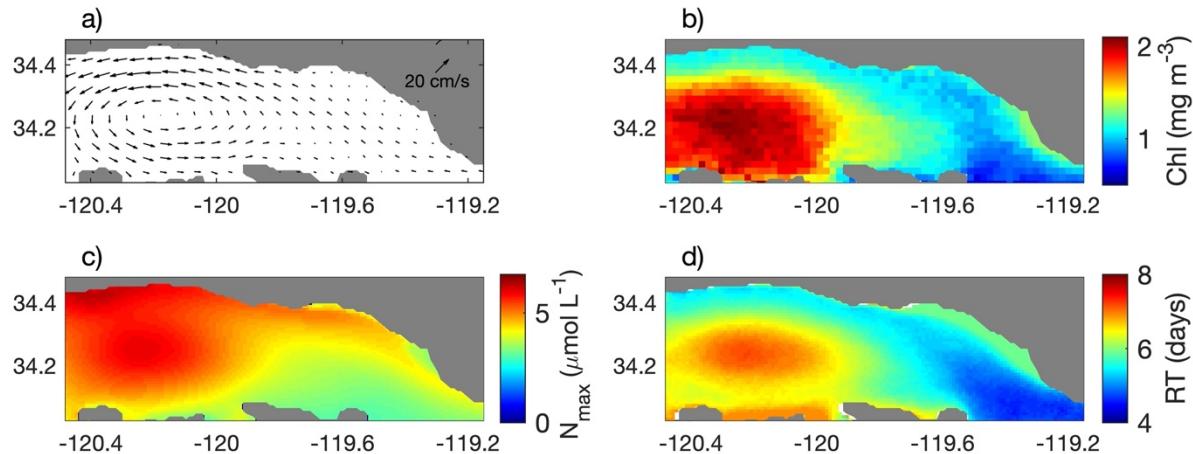
## 289 **3.0 Results**

### 290 **3.1 Mean decadal horizontal distributions**

291 Fig. 4 shows the 10-year mean horizontal distributions of currents, Chl,  $N_{\max}$ , and RT in  
292 the SBC. The mean current pattern in the surface layer (Fig. 4(a)) show that strong eddy  
293 circulation, centered mid-channel, dominates the mean circulation in the western SBC with  
294 weaker, primarily westward currents observed in the eastern SBC. The magnitude of the mean  
295 and maximum currents in Fig. 4(a) are  $4 \text{ cm s}^{-1}$  and  $13 \text{ cm s}^{-1}$  respectively. The highest mean  
296 Chl are observed in the western SBC where eddy circulation dominates and decreases to the

297 north and east with the lowest values at the southeastern boundary (Fig. 4(b)). This spatial  
 298 distribution of Chl has been observed in previous studies (Otero and Siegel 2004; Brzezinski  
 299 and Washburn 2011; Henderikx Freitas et al. 2017). The mean Chl varies from  $0.1\text{-}3.4\text{ mg m}^{-3}$   
 300 across  $1\text{ km}^2$  pixels. The spatial distribution of mean RT reflects the eddy circulation with the  
 301 highest values of  $\sim 7$  days in the central western SBC and lowest values of  $\sim 4$  days in the  
 302 southeastern SBC (Fig. 4(d)). The spatial distribution of mean  $N_{\max}$  shows a distinct east-west  
 303 gradient with the highest values of  $\sim 6\text{ }\mu\text{mol L}^{-1}$  observed in the western SBC and the lowest  
 304 values of  $\sim 3\text{ }\mu\text{mol L}^{-1}$  observed in the eastern SBC. Unlike mean Chl and RT, the mean  $N_{\max}$   
 305 distribution shows two distinct regions of high concentrations, a circular shaped region in the  
 306 central western SBC and a region along the mainland coast in the north-western SBC. The  
 307 circular region is located near the center of the eddy circulation (Fig. 4(a)) and coincides with  
 308 the area of the highest mean Chl and RT (Fig. 4(b,d)).

309

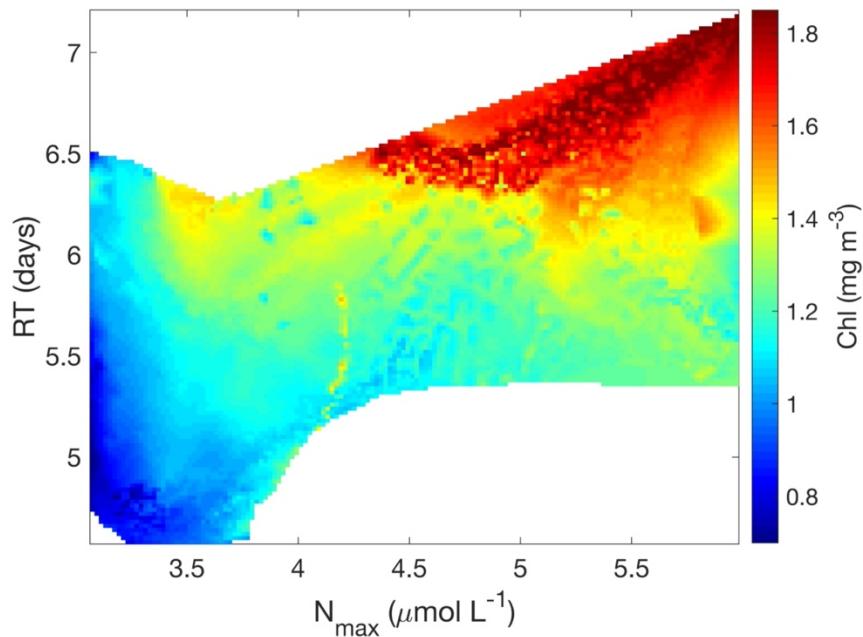


310  
 311 **Figure 4:** 10-year mean: (a) ROMS currents, (b) Chl ( $\text{mg m}^{-3}$ ), (c)  $N_{\max}$  ( $\mu\text{mol L}^{-1}$ ), and (d) RT  
 312 (days).  $N_{\max}$ , RT, and currents are depth-averaged over the surface layer.  
 313

314 To show the relationship between mean  $N_{\max}$ , RT, and Chl, mean Chl is shown as a  
 315 function of mean  $N_{\max}$  and RT in Fig. 5. Elevated mean Chl, defined by Chl greater than  $1.4\text{ mg}$   
 316  $\text{m}^{-3}$ , is found where mean  $N_{\max}$  and RT are simultaneously greater than  $\sim 4\text{ }\mu\text{mol L}^{-1}$  and  $\sim 5.5$   
 317 days, respectively. These conditions coincide with eddy circulation in the western SBC. In

318 contrast, although mean  $N_{\max}$  is elevated along the north-western mainland coast (Fig. 4(c)), the  
319 mean RT in this area is relatively short, less than 5.5 days, suggesting that  $N_{\max}$  is advected  
320 away from the coast and into the open channel before elevated Chl can be produced. Thus, it  
321 appears that for long-term horizontal means, the elevated Chl in the SBC is being produced by  
322 a combination of long RT and high  $N_{\max}$ .

323



324  
325  
326 **Figure 5:** Chl ( $\text{mg m}^{-3}$ ) as a function of  $N_{\max}$  ( $\mu\text{mol L}^{-1}$ ) and RT (days). White represents where  
327 Chl data are not available for  $N_{\max}$ -RT combinations.  
328  
329

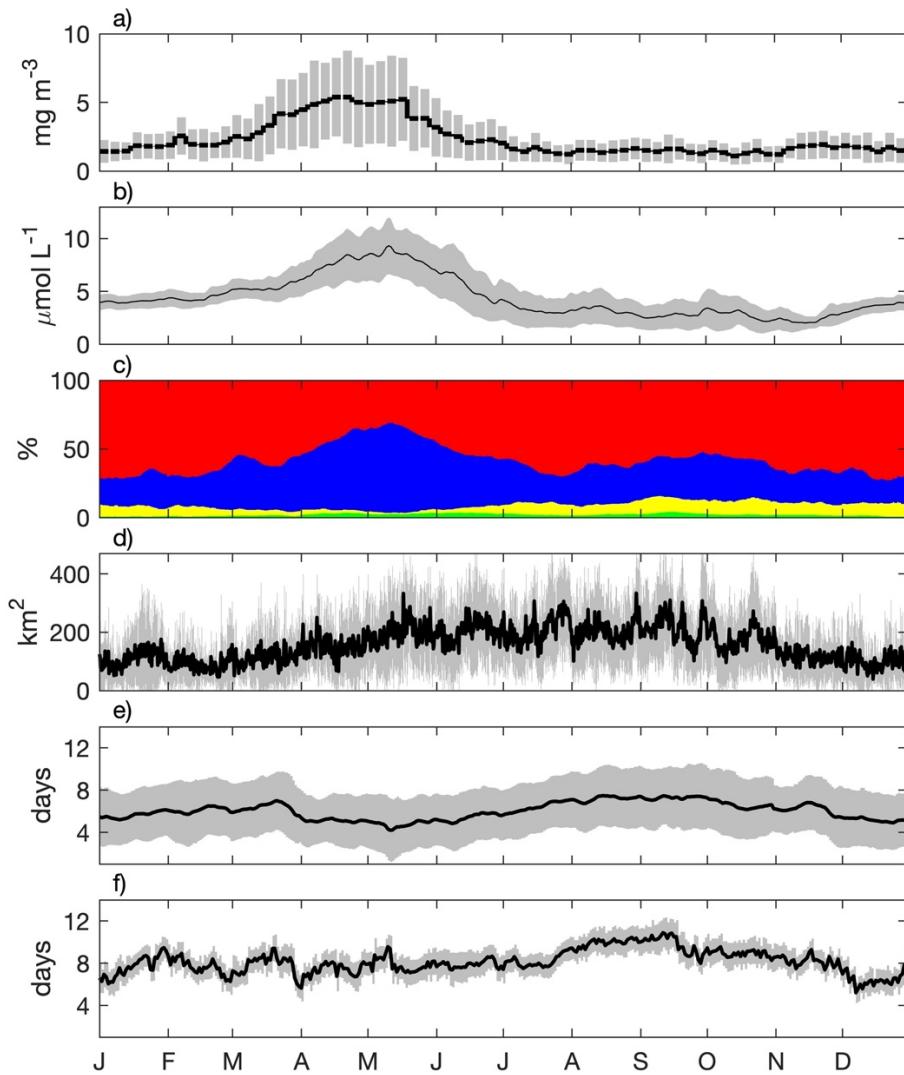
### 330 **3.2 Mean annual cycle**

331 Mean annual cycles of Chl,  $N_{\max}$ , source water zone, eddy size, RT, and  $RT_{\text{eddy}}$  are  
332 displayed in Fig. 6. Chl shows a strong seasonal cycle with elevated values from March through  
333 June and maximum values of 3.8-5.4  $\text{mg m}^{-3}$  in April and May (Fig. 6(a)). Low mean Chl is  
334 observed throughout the rest of the year, averaging 1.6  $\text{mg m}^{-3}$ . This seasonal cycle of mean  
335 Chl in the SBC has been observed in prior studies (Shipe and Brzezinski 2001; Otero and  
336 Siegel 2004; Brzezinski and Washburn 2011; Henderikx Freitas et al. 2017). Mean  $N_{\max}$  follows  
337 a seasonal cycle similar to mean Chl with elevated values in March through June and maximum

338 values in April and May of 6.1-9.5  $\mu\text{mol L}^{-1}$  (Fig. 6(b)). Low mean  $N_{\max}$  is observed throughout  
339 the rest of the year with an average of 3.3  $\mu\text{mol L}^{-1}$ . By averaging mean  $N_{\max}$  in Fig. 6(b) over  
340 the 5-day composite window used for Chl (Fig. 6(a)), the annual cycle of mean Chl and  $N_{\max}$  is  
341 shown to be highly correlated with a Pearson's squared correlation coefficient ( $r^2$ ) of 0.83.  
342 Brzezinski and Washburn (2011) observed a similar relationship in the seasonal cycles of  
343 chlorophyll and nitrate in the SBC.

344 The mean annual cycles of physical properties, including source water zone, eddy size,  
345 RT, and  $RT_{\text{eddy}}$ , are shown in Fig. 6(c-f). Of these four physical properties, the annual cycle of  
346 source water from the West zone, representing the CC, is the most similar to the annual cycles  
347 of  $N_{\max}$  and Chl ( $r^2$  of 0.70-0.71) with maximum values of 40-65% from March to June (Fig. 6(c)).  
348 When the mean percent of water from the West zone is low in July-February, water in the SBC  
349 originates primarily from the East zone or the SCC, accounting for 45-61% of the SBC source  
350 water. The mean percent of water from inside the SBC ranges from 1-4% and from the South  
351 zone ranges from 5-8% throughout the year without a defined seasonal cycle.

352 Eddy circulation is detected throughout the year as indicated by the eddy size in Fig.  
353 6(d)). Eddy size is directly proportional to the strength of eddy circulation and ranges from a few  
354  $\text{km}^2$  up to 450  $\text{km}^2$ . The mean annual cycle of eddy size is much less pronounced than the  
355 mean annual cycles of Chl,  $N_{\max}$ , and West zone source water and displays its largest values in  
356 May through October. These results are supported by field studies that observed the presence  
357 of eddy circulation most consistently in summer and fall (Harms and Winant 1998; Dever et al.  
358 1998; Winant et al. 2003). Mean RT lacks a defined seasonal cycle and ranges from 4.2 to 7.5  
359 days (Fig. 6(e)). The daily standard deviation of RT is fairly large, ranging from 2.0 to 3.2 days  
360 and indicating that for any single day the RT can vary widely. Mean  $RT_{\text{eddy}}$  (Fig. 6(f)) is  
361 consistently higher than the mean RT, ranging from 5.2 to 10.9 days over the year with a small  
362 standard deviation from 0.7-1.6 days and no clear seasonal cycle.

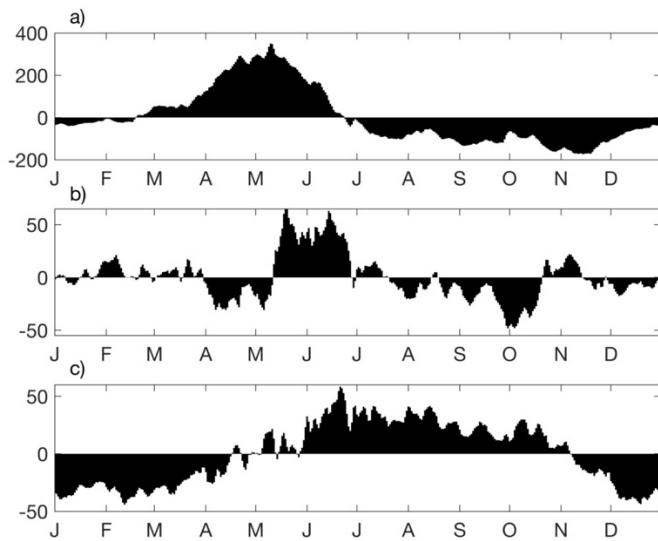


365 **Figure 6:** Mean annual cycle of (a) Chl ( $\text{mg m}^{-3}$ ), (b)  $\text{N}_{\text{max}}$  ( $\mu\text{mol L}^{-1}$ ), (c) source water zone (%)  
 366 with West zone (blue), East zone (red), South zone (yellow), and SBC (green), (d) eddy size  
 367 ( $\text{km}^2$ ), (e) RT (days), and (f)  $\text{RT}_{\text{eddy}}$  (days). Gray areas show  $\pm$  one standard deviation.  
 368

369 The mean annual cycle of the principal components for  $\text{N}_{\text{max}}$  EOF modes 1, 2 and 3

370 (PC1, PC2, and PC3) is shown in Fig. 7. From the EOF analysis in Section 2.4, the  $\text{N}_{\text{max}}$   
 371 transport mechanisms represented by each mode are (1) wind-driven upwelling, (2) advection of  
 372 high  $\text{N}_{\text{max}}$  water into the south-western SBC by eddy circulation, and (3) retention of  $\text{N}_{\text{max}}$  within  
 373 the SBC by eddy circulation. Mean PC1 shows a similar annual cycle as mean Chl,  $\text{N}_{\text{max}}$ , and  
 374 West zone source water (Fig. 6(a-c)) with positive values in March through June and maximum

375 values in April and May, reinforcing that upwelling is the primary transport process of  $N_{max}$  to the  
 376 surface layer in spring. Although much smaller in variance than PC1, mean PC2 and PC3,  
 377 representing the transport of  $N_{max}$  by eddy circulation, have peak positive values in late May and  
 378 June, when PC1 decreases and then becomes negative (Fig. 7). This pattern suggests that as  
 379 upwelling weakens, eddy circulation plays a larger role in transporting  $N_{max}$  in the SBC surface  
 380 layer. Mean PC3 is consistently positive in June through October (Fig. 7(c)), which is when  
 381 mean Chl and  $N_{max}$  are at their lowest and mean PC1 is negative.

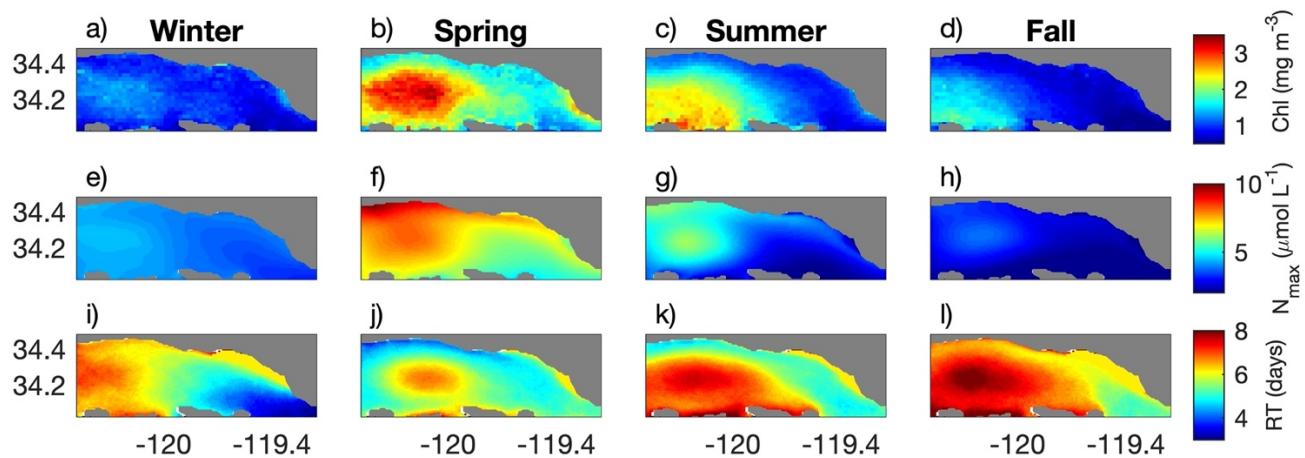


382  
 383 **Figure 7:** Mean annual cycle of principal components for  $N_{max}$  EOF: (a) mode 1, (b) mode 2,  
 384 and (c) mode 3  
 385

386 To evaluate the spatial variability in the seasonal cycle of mean Chl,  $N_{max}$ , and RT, these  
 387 parameters are averaged over the four seasons and shown in Fig. 8. The horizontal  
 388 distributions of Chl and  $N_{max}$  in spring show the highest concentrations and are most similar to  
 389 their decadal horizontal means (Fig. 4(b,c)), implying that spring activity is driving the decadal  
 390 mean distributions. Both Chl and  $N_{max}$  in spring (Fig. 8(b,f)) display distinct along channel  
 391 gradients with high concentrations in the western SBC, low concentrations in the eastern SBC,  
 392 and maximum concentrations in the central western SBC where decadal mean eddy circulation

393 is observed (Fig. 4(a)). Compared to spring, Chl and  $N_{\max}$  display decreasing concentrations  
 394 throughout the SBC in summer and fall, although both parameters continue to show along-  
 395 channel gradients with the highest concentrations in the western SBC. In contrast, RT displays  
 396 high values in the western SBC of similar magnitude throughout the year with the spatial  
 397 distribution of RT varying in shape with the season (Fig. 8(i-l)). Although eddy circulation is  
 398 present throughout the year, the eddy is not stationary and moves primarily along-channel in the  
 399 western SBC with a tendency to propagate westward (Harms and Winant 1998; Beckenbach  
 400 and Washburn 2004; Simons et al. 2015). Shown in Fig. 8(i-l), the oval shape of the longest RT,  
 401 greater than approximately 6 days, reflects the along-channel movement of the eddy, which is  
 402 smallest in spring, largest in summer and fall, and at its least constrained in winter. Overall, Fig.  
 403 8 emphasizes the need for both high  $N_{\max}$  and long RT to be present to produce elevated Chl.

404

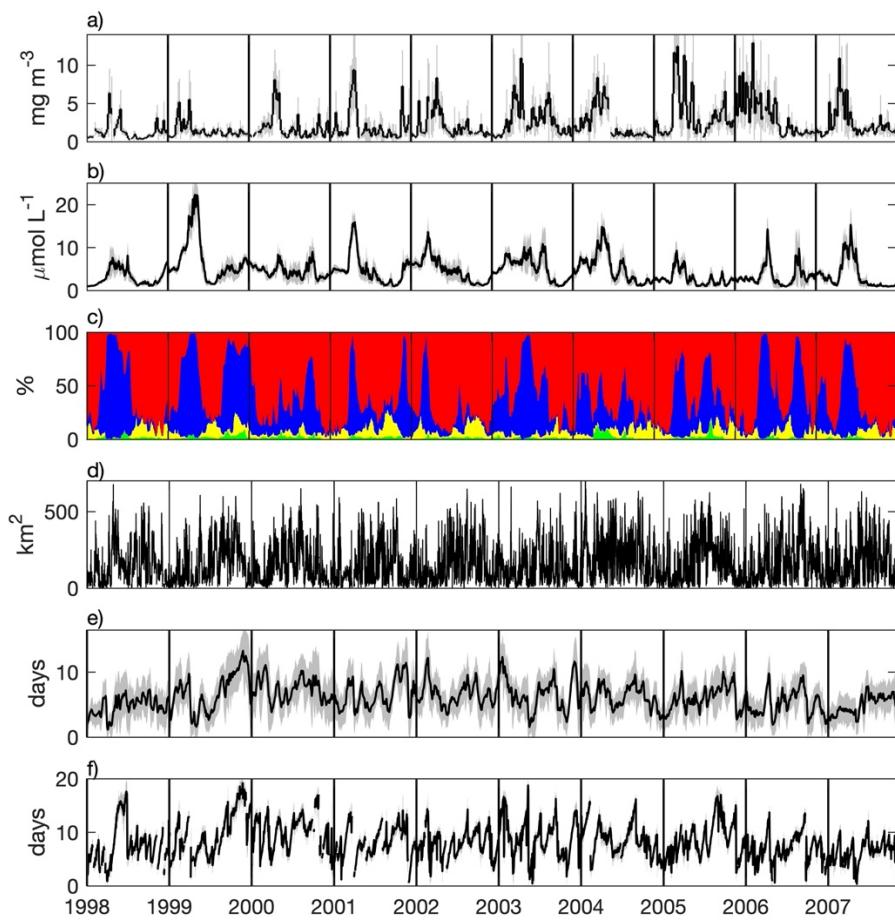


405  
 406 **Figure 8:** Mean seasonal cycle of Chl ( $\text{mg m}^{-3}$ ) for (a) winter (Dec-Feb), (b) spring (Mar-May),  
 407 (c) summer (Jun-Aug), and (d) fall (Sep-Nov),  $N_{\max}$  ( $\mu\text{mol L}^{-1}$ ) for (e) winter, (f) spring, (g)  
 408 summer, and (h) fall, and RT (days) for (i) winter, (j) spring, (k) summer, and (l) fall  
 409

410 **3.3 Interannual variations in the spring upwelling period**

411 Time-series of Chl and  $N_{\max}$  over the 10-year modeling period are shown in Fig. 9(a,b).  
 412 Although varying in magnitude between years, the maximum Chl and  $N_{\max}$  are consistently  
 413 observed in the spring. Similarly, PC1 or upwelling strength shows the largest positive values in

414 the spring (Fig. 10(a)). Based on these results as well as the mean annual cycles, our  
 415 interannual analysis focuses on the spring upwelling period (SUP), the time of year with the  
 416 strongest upwelling and highest Chl and  $N_{\max}$ . The SUP is defined by contiguous positive  
 417 values of PC1 that occur in February through July. From 1998 to 2007, the SUP ranges in  
 418 length from 42 to 127 days with an annual average of 76 days (Table 1, Fig. 10(a)). Mean  
 419 values of Chl,  $N_{\max}$ , PC1, RT and  $RT_{\text{eddy}}$  for yearly SUPs are shown in Table 1. For every year  
 420 except 1998 and 1999 when a strong ENSO cycle occurred, the SUP is associated with  
 421 elevated mean Chl ranging from 3.7 to 8.3  $\text{mg m}^{-3}$  and elevated mean  $N_{\max}$  ranging from 6.0 to  
 422 10.7  $\mu\text{mol L}^{-1}$  (Table 1, Fig. 9(a,b)). Mean Chl and  $N_{\max}$  from times of the year other than the  
 423 SUP are 1.6  $\text{mg m}^{-3}$  and 3.3  $\mu\text{mol L}^{-1}$  respectively.



424  
 425 **Figure 9:** SBC time-series of (a) Chl ( $\text{mg m}^{-3}$ ), (b)  $N_{\max}$  ( $\mu\text{mol L}^{-1}$ ), (c) source water zone (%)  
 426 with West zone (blue), East zone (red), South zone (yellow), and SBC (green), (d) eddy size  
 427 ( $\text{km}^2$ ), (e) RT (days), and (f)  $RT_{\text{eddy}}$  (days). Gray areas show  $\pm$  one standard deviation.

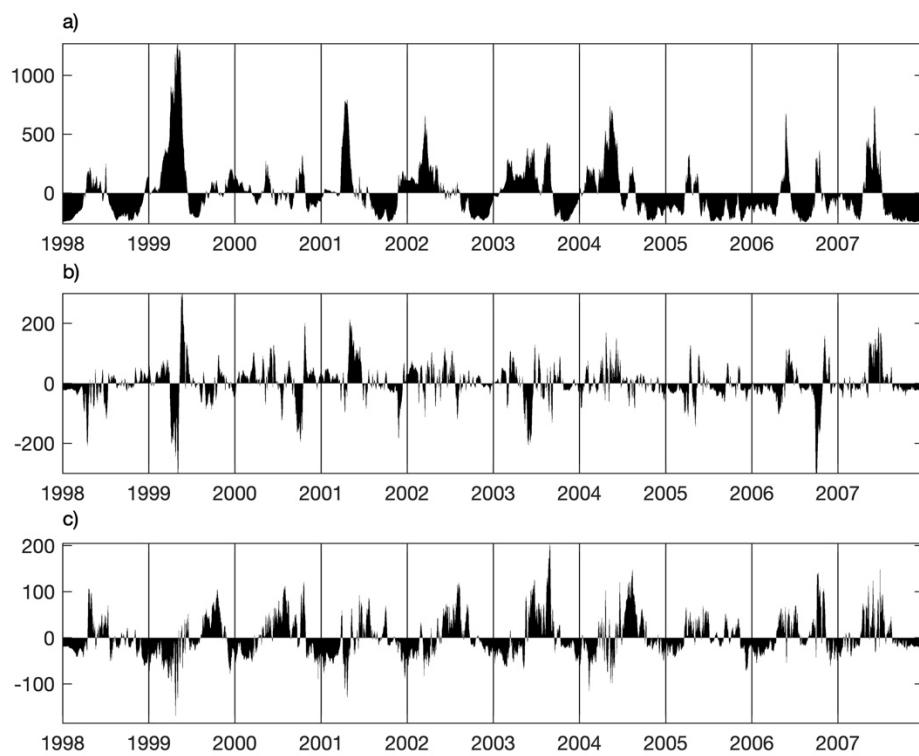
428

Parameter	Units	Year										Yearly Mean
		1998	1999	2000	2001	2002	2003	2004	2005	2006	2007	
Start date		Apr 19	Feb 9	Apr 24	Mar 23	Feb 19	May 13	Mar 21	Mar 27	May 3	Apr 18	Mar 28
End date		Jul 7	Jun 15	Jun 4	May 20	May 20	Jul 19	Jun 23	May 23	Jun 16	Jul 19	Jun 15
Length	days	92	127	42	59	91	68	95	58	45	83	76
Chl	mg m <sup>-3</sup>	2.2	2.2	4.4	5.3	3.7	4.0	4.9	8.3	4.0	4.2	4.3
N <sub>max</sub>	μmol L <sup>-1</sup>	5.6	12.5	6.0	10.7	8.2	7.4	9.4	6.0	7.6	8.7	8.2
Upwelling strength (PC1)		88	561	103	432	263	218	363	119	218	291	266
RT	days	4.8	5.5	6.4	5.9	6.5	4.9	6.2	5.3	4.7	5.0	5.5
RT <sub>eddy</sub>	days	11.1	8.1	10.3	6.6	7.0	7.1	8.4	8.5	6.0	6.5	8.0
MEI		1.57	-0.88	-0.10	0.04	0.32	0.07	0.31	0.67	0.57	-0.13	0.3
Maximum eddy lifespan	days	56	30	27	11	22	22	11	6	20	8	21
Eddy size upwelling	km <sup>2</sup>	263	116	156	130	62	80	201	137	131	159	144
Eddy size relaxation	km <sup>2</sup>	131	131	237	209	131	242	285	208	282	224	208
Maximum RT <sub>eddy</sub>	days	17.6	13.8	12.9	9.2	14.0	13.5	13.2	11.5	8.4	12.1	12.6
N <sub>max</sub> in eddy - N <sub>max</sub> <sup>*</sup>	μmol L <sup>-1</sup>	1.7	8.5	6.4	7.3	5.0	4.4	6.9	4.1	8.4	5.4	5.8
Maximum N <sub>max</sub> Flux	μmol L <sup>-1</sup> day <sup>-1</sup>	3.8	13.8	1.7	4.1	2.1	4.4	3.7	2.5	3.7	4.7	4.4

429

430  
431

**Table 1:** Yearly data for the spring upwelling period (SUP). Mean values are displayed unless otherwise indicated. \*At the end of the SUP.

434 **Figure 10: Principal components for  $N_{\max}$  EOF: (a) mode 1, (b) mode 2, and (c) mode 3.**436 **Section 3.3.1 *Typical spring upwelling period***

437 The strength and timing of  $N_{\max}$  transport processes during the SUP are critical to  
 438 understanding how upwelling and eddy circulation regulate Chl in the SBC. The SUP consists of  
 439 three main phases; (1) the upwelling phase, when upwelling starts and then increases in  
 440 strength, (2) the transition phase, when upwelling transitions from increasing to decreasing  
 441 strength, and (3) the relaxation phase, when upwelling strength declines until the end of the  
 442 SUP. To examine these three phases, time-series of Chl,  $N_{\max}$ , PC1, eddy size, RT, and  $RT_{\text{eddy}}$   
 443 during a typical SUP are examined (Figs. 11(a-f) and 12). The 2001 SUP is presented as a  
 444 typical SUP for several reasons. First, PC1 displays a clear three phase cycle. Second, the  
 445 2001 SUP has the best agreement between modeled and observed temperature in the SBC  
 446 (see SI). Third, the 2001 SUP has a neutral multivariate ENSO index (MEI), which tracks El  
 447 Niño and La Niña events in the Pacific Ocean (Table 1, Fig. S1). Although the following analysis

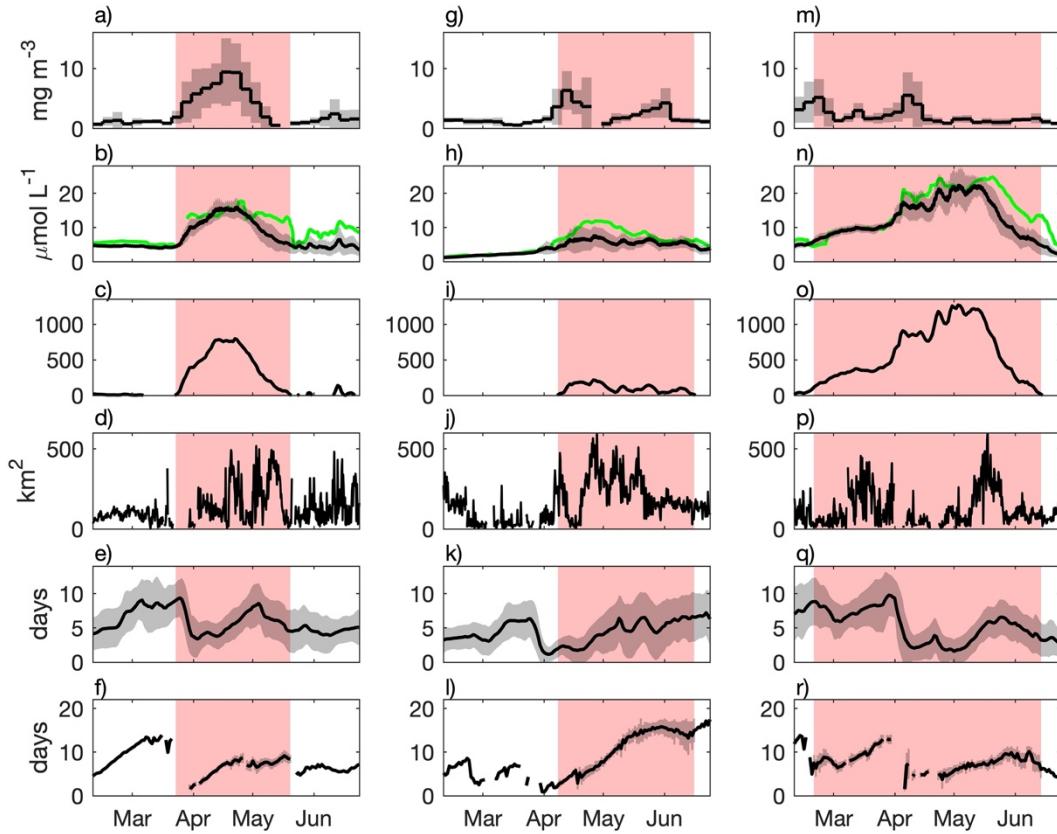
448 focuses on the 2001 SUP, the 2000 and 2002-2007 SUPs display similar results, which are  
449 shown for comparison in Table 1 and Figs. 9 and 10.

450 In the upwelling phase of the 2001 SUP, upwelling begins and increases in strength as  
451 displayed by the increasing values of PC1 from March 23 to April 12 in Fig. 11(c). Chl and  $N_{max}$   
452 also steadily increase during this phase (Fig. 11(a,b)). The horizontal advection of upwelled  
453 water from the CC into the SBC results in a rapid drop in RT (Fig. 11(e)) and little to no eddy  
454 circulation as represented by the small eddy size (Fig. 11(d)). Shown for April 2<sup>nd</sup> in Fig.  
455 12(a,d), the spatial relationship of  $N_{max}$  and RT during the upwelling phase displays the intrusion  
456 of high  $N_{max}$  water from the west, which corresponds to a shortened RT and the absence of  
457 eddy circulation. Next in the transition phase, upwelling strength stops increasing as shown by  
458 the plateauing values of PC1 from April 13 to 22 in Fig. 11(c). This coincides with maximum Chl  
459 and  $N_{max}$  and increasing RT and eddy size (Fig. 11 (a-b,d-e)). During the transition phase on  
460 April 14<sup>th</sup>, the horizontal distribution of  $N_{max}$  shows values greater than 10  $\mu\text{mol L}^{-1}$  across the  
461 entire SBC with the highest concentrations along the mainland coast (Fig. 12(b)). The  
462 horizontal distribution of RT on April 14<sup>th</sup> shows increased RT and eddy circulation compared to  
463 the upwelling phase as advection from the west weakens (Fig. 12(e)).

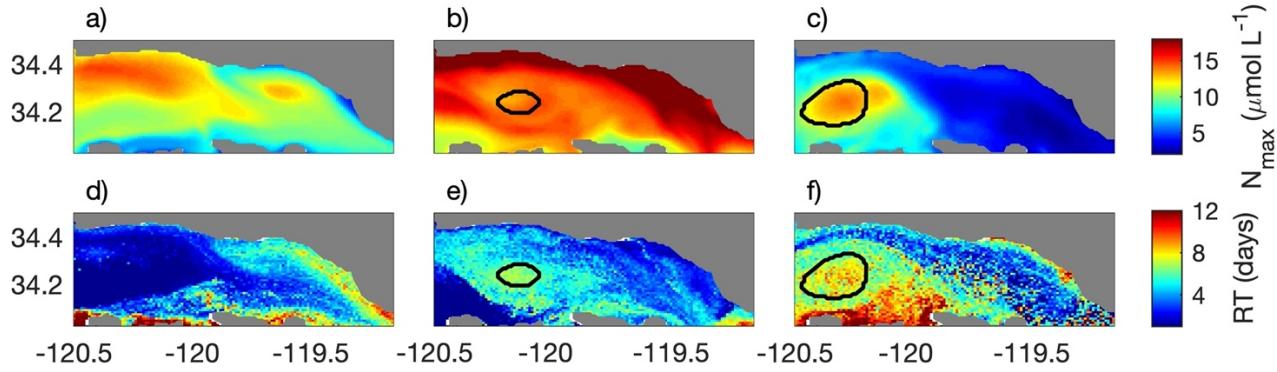
464 The relaxation phase, the last phase of the SUP, displays the greatest deviation from  
465 typical upwelling-relaxation cycles in EBUSs due to the presence of eddy circulation. In this  
466 phase, upwelling strength decreases, shown by the steady decline in PC1 from April 23 to May  
467 20 (Fig. 11(c)). Chl and  $N_{max}$  similarly decline during this phase (Fig. 11(a,b)). Horizontal  
468 distributions of  $N_{max}$  and RT on May 12<sup>th</sup> (Fig. 12(c,f)) show that the influx of high  $N_{max}$  water  
469 from the west has stopped and eddy circulation has strengthened. Eddy circulation has trapped  
470 high  $N_{max}$  water in the western SBC as shown by the high  $N_{max}$  and long RT located within the  
471 eddy area (Fig. 12(c,f)).  $N_{max}$  within the eddy, shown in Fig. 11(b), is similar in magnitude to  
472  $N_{max}$  in the SBC during the upwelling and transition phases, but is greater than  $N_{max}$  in the SBC  
473 by 3-7  $\mu\text{mol L}^{-1}$  throughout the relaxation phase as eddy circulation retains high  $N_{max}$  water. The

474 largest difference between  $N_{\max}$  within the eddy and  $N_{\max}$  in the SBC of  $7.3 \mu\text{mol L}^{-1}$  is observed  
 475 at the end of the SUP. Similar to the 2001 SUP, the other typical years, 2000 and 2002-2007,  
 476 all show an increase in eddy size from the upwelling to the relaxation phase and  $N_{\max}$  in the  
 477 eddy higher than  $N_{\max}$  in the SBC at the end of SUP (Table 1).

478



479  
 480 **Figure 11:** Chl ( $\text{mg m}^{-3}$ ) for (a) 2001, (g) 1998, and (m) 1999;  $N_{\max}$  ( $\mu\text{mol L}^{-1}$ , black line) and  
 481  $N_{\max}$  in eddy ( $\mu\text{mol L}^{-1}$ , green line) for (b) 2001, (h) 1998, and (n) 1999; Positive values of PC1  
 482 for (c) 2001, (i) 1998, and (o) 1999; eddy size ( $\text{km}^2$ ) for (d) 2001, (j) 1998, and (p) 1999; RT  
 483 (days) for (e) 2001, (k) 1998, and (q) 1999;  $RT_{\text{eddy}}$  (days) for (f) 2001, (l) 1998, and (r) 1999.  
 484 Pink areas identify the SUP. Gray areas show  $\pm$  one standard deviation.  
 485  
 486



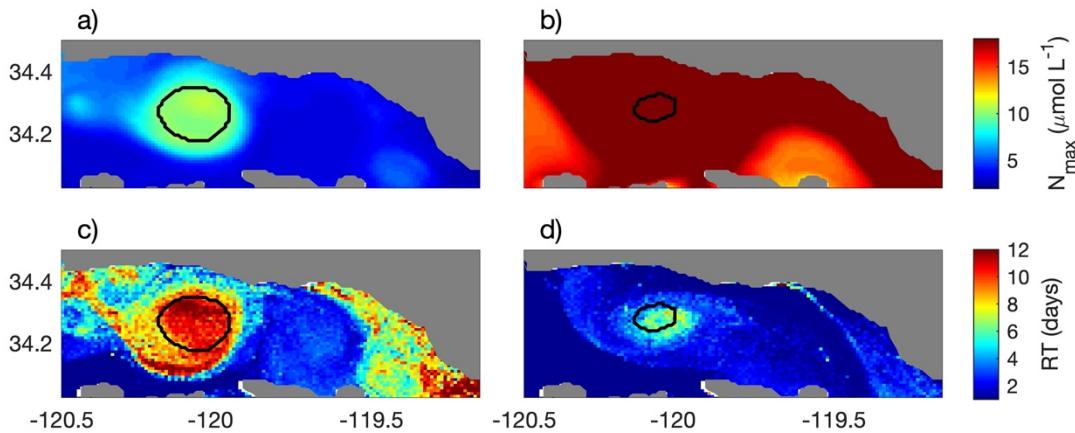
487  
488 **Figure 12:**  $N_{\max}$  ( $\mu\text{mol L}^{-1}$ ) for (a) upwelling phase on April 2, 2001, (b) transition phase on April  
489 14, 2001, and (c) relaxation phase on May 12, 2001. RT (days) for (d) upwelling phase on April  
490 2, 2001, (e) transition phase on April 14, 2001, and (f) relaxation phase on May 12, 2001. Black  
491 line identifies the eddy area.  $N_{\max}$  and RT have been depth-averaged over the surface layer.  
492  
493

### 494 3.3.2 Spring upwelling period during extreme *El Niño* and *La Niña* conditions

495 Patterns of upwelling and eddy circulation during 1998 and 1999 SUPs deviate the most  
496 from the typical years and coincide with an exceptionally strong ENSO cycle, an extreme El  
497 Niño event in 1998 followed by an extreme La Niña event in 1999. The SUP in both 1998 and  
498 1999 have anomalously low mean Chl of  $2.2 \text{ mg m}^{-3}$ , which is 40-74% lower than the other  
499 years of 2000-2007 in the study period (Table 1 and Fig. 11(g,m)). While strong El Niño events  
500 historically disrupt primary productivity in the California EBUS, La Niña events are often  
501 associated with enhanced upwelling and phytoplankton accumulation (Bograd et al. 2000;  
502 Schwing et al. 2000; Bograd and Lynn 2001).

503 The 1998 SUP coincided with an extreme El Niño, which resulted in a mean PC1,  
504 upwelling strength, and  $N_{\max}$  of 88 and  $5.6 \mu\text{mol L}^{-1}$  respectively, the lowest of all yearly SUPs  
505 (Table 1, Fig. 11(h,i)). Taking place from April 19 to July 7, the 1998 SUP does not display  
506 distinct phases of upwelling, transition, and relaxation, but instead shows very weak upwelling of  
507 similar magnitude throughout the SUP. In addition, the 1998 SUP exhibits an anomalous  
508 pattern of eddy circulation compared to the typical SUPs of 2000-2007. Unlike a typical SUP  
509 where eddy circulation increases in strength from the upwelling phase to the relaxation phase,

510 eddy circulation is strong and continuous for most the 1998 SUP as shown by the eddy size in  
 511 Fig. 11(j). During the 1998 SUP, eddy circulation displays a lifespan, defined as a continuously  
 512 tracked eddy, of 56 days from May 3 to June 28. Present for 60% of the 1998 SUP, this eddy  
 513 lifespan is twice as long as the longest eddy lifespan for the other SUPs (Table 1) and almost  
 514 five times longer than the mean SUP eddy lifespan of 12 days.  $RT_{eddy}$  is exceptionally long with  
 515 an average of 14.3 days over the 56-day eddy lifespan, which is 79% greater than the SUP  
 516 mean eddy lifespan of 8 days (Table 1). Horizontal distributions of  $N_{max}$  and  $RT$  on May 7<sup>th</sup>, the  
 517 midpoint of the 1998 SUP, show that  $N_{max}$  within the eddy is only slightly greater than  $N_{max}$   
 518 outside of the eddy and that  $RT$  is much higher within the eddy than outside (Fig. 13(a,c)).  
 519 These horizontal distributions are distinctly different from all phases of the 2001 SUP (Fig. 12).  
 520 These results imply that in the presence of very weak upwelling, prolonged periods of eddy  
 521 circulation produce long  $RT$ , but not high  $N_{max}$  or elevated Chl.



522  
 523 **Figure 13:**  $N_{max}$  ( $\mu\text{mol L}^{-1}$ ) on (a) May 7, 1998 and (b) May 1, 1999.  $RT$  (days) on (c) May 7,  
 524 1998 and (d) May 1, 1999. Black line identifies the eddy area.  $N_{max}$  and  $RT$  have been depth-  
 525 averaged over the surface layer.  
 526

527 In contrast, 1999 is an extreme La Niña year with record high upwelling for the entire  
 528 California EBUS (Schwing et al. 2000). This strong upwelling is reflected in the model  
 529 predictions for the 1999 SUP, which has the highest mean PC1, upwelling strength, of 561 and  
 530  $N_{max}$  of  $12.5 \mu\text{mol L}^{-1}$  of all the SUPs (Table 1). The 1999 SUP is the longest SUP, lasting 127

531 days from February 9 to June 15. Although the 1999 SUP has unusually high  $N_{max}$ , it does not  
532 show elevated Chl with a mean Chl of  $2.2 \text{ mg m}^{-3}$ , the same as the 1998 SUP, which has the  
533 weakest upwelling (Table 1). Similar to the 2001 SUP, the 1999 SUP shows three distinct  
534 phases of upwelling, transition, and relaxation shown by the patterns of  $N_{max}$  and PC1 (Fig.  
535 11(n,o)). However, eddy circulation deviates from the typical SUPs by remaining small  
536 throughout the transition phase, which is 38 days from April 7 to May 15 (Fig. 11(p)). During the  
537 1999 transition phase, upwelling strength (PC1) and  $N_{max}$  are atypically high, averaging 1,044  
538 and  $18.9 \mu\text{mol L}^{-1}$  respectively (Fig. 11(n,o)), but RT is unusually low, averaging only 2.6 days  
539 (Fig. 11(p)). This very low RT indicates that water is being advected through the SBC very  
540 quickly. To approximate the  $N_{max}$  flux through the SBC,  $N_{max}$  is divided by RT for all of the  
541 SUPs. All SUPs have a daily  $N_{max}$  flux of  $4.7 \mu\text{mol L}^{-1} \text{ day}^{-1}$  or less except for the 1999 SUP  
542 transition phase where  $N_{max}$  flux ranges from 4.8 to  $13.8 \mu\text{mol L}^{-1} \text{ day}^{-1}$  (Table 1). The horizontal  
543 distributions of  $N_{max}$  and RT during the transition phase on May 1, 1999 (Fig. 13(b,d)) are very  
544 different than those of the 2001 SUP transition phase (Fig. 12(b,e)) with unusually high  $N_{max}$  and  
545 low RT throughout the SBC. The eddy is also small on May 1<sup>st</sup>, containing only slightly elevated  
546 RT (Fig. 13 (b,d)). These results suggest that the exceptionally strong upwelling in 1999  
547 advected nutrients through the SBC too quickly for Chl accumulation.

548

#### 549 **4.0 Discussion and Conclusions**

550 Our results suggest that spring Chl in the SBC is regulated by the coupling of upwelling  
551 and eddy circulation and that elevated spring Chl requires the presence of both high  $N_{max}$  from  
552 upwelling and long RTs from eddy circulation at mean decadal (Figs. 4-5), mean annual (Figs.  
553 6-8), interannual (Figs. 9-11), and synoptic timescales (Figs. 12-13). Unlike most of the coast  
554 in the SC Bight that is dominated by the warm poleward flowing SCC, the SBC is a transition  
555 zone where the cold equatorward flowing CC and the warm poleward flowing SCC meet, which,

556 along with the bathymetry, creates a pattern of cyclonic eddy circulation (Hickey 1993; Harms  
557 and Winant 1998; Oey et al. 2004). Our results show that eddy circulation is present throughout  
558 the year in the SBC (Figs. 6(d) and 9(d)) and a critical process driving the Chl hotspot in SBC.

559 **4.1 Upwelling and eddy circulation stimulate a productivity hotspot in the SBC**

560 In the California EBUS, the time required after an upwelling event for phytoplankton to  
561 deplete the surface ocean of new nutrients is 3-10 days (Dugdale et al. 1997; Botsford et al.  
562 2006; Wilkerson et al. 2006; Krause et al. 2013). Our study predicts a 10-year mean RT of 6.0  
563  $\pm$  2.7 days (mean  $\pm$  1 standard deviation) in the SBC, which meets the above criteria for nutrient  
564 uptake. In addition, eddy circulation creates a horizontal oval pattern of long mean RTs in the  
565 western SBC, which persists in the spring, summer, and fall and correlates with similar patterns  
566 of high mean  $N_{max}$  and Chl in the spring (Fig. 8). When comparing the 10-year mean horizontal  
567 distributions of RT and  $N_{max}$  to Chl, both elevated RT and  $N_{max}$  are required to produce high Chl,  
568 greater than  $\sim$ 1.4 mg m<sup>-3</sup>, with minimum threshold values of  $\sim$ 4.0  $\mu$ mol L<sup>-1</sup> for  $N_{max}$  and  $\sim$ 5.5  
569 days for RT (Fig. 5).

570 On a mean annual scale, the modeling predicts the strongest upwelling in the spring  
571 (Fig. 7(a)), which coincides with elevated Chl, high  $N_{max}$ , and a large influx of water from the CC  
572 (Fig. 6(a-c)). These results reflect previous studies of the SBC that observed high Chl,  
573 phytoplankton primary production, and nutrients in the spring followed by low values the rest of  
574 the year (Otero and Siegel 2004; Brzezinski and Washburn 2011; Henderikx Freitas et al.  
575 2017). The mean annual SUP ranges from March to June as determined by PC1 (Fig. 7(a)) and  
576 coincides with the transport of  $N_{max}$  by eddy circulation from late-May to June, represented by  
577 PC2 and PC3 (Fig. 7(b-c)). While mean RT and  $RT_{eddy}$  remain above the nutrient uptake and  
578 depletion timescales throughout the year (Fig. 6(e,f)), mean elevated Chl is only observed  
579 during the SUP when mean  $N_{max}$  is high. While positive PC2 and PC3 are present during the  
580 SUP (Fig. 7(b,c)), they are also positive at other times of the year when elevated Chl is not

581 observed. This indicates that eddy circulation without upwelling is unlikely to transport new  
582 nutrients into the surface layer in sufficient quantities to stimulate elevated Chl on the spatial  
583 scale of this study.

584 On an interannual scale, the SUP is observed for every year of the 10-year modeling  
585 period, beginning as early as February and ending as late as July. The SUP ranges in length  
586 from 1-4 months and occurs most frequently in April and May. A typical SUP consists of three  
587 phases; upwelling, transition, and relaxation. When upwelling increases in strength during the  
588 upwelling phase, eddy circulation is weak and RT is short due to the strong westward advection  
589 of high  $N_{max}$  water from the CC. During the transition and relaxation phases when upwelling  
590 strength plateaus and then weakens, eddy circulation strengthens, and RT increases.  
591 Consequently, eddy circulation traps and retains nutrients introduced by upwelling in the SBC.  
592 This process is clearly demonstrated by  $N_{max}$  in the eddy remaining elevated throughout the  
593 relaxation phase while  $N_{max}$  outside of the eddy decreases (Figs. 11(b) and 12(c), Table 1).  
594 Overall, eddy retention of high  $N_{max}$  water during the relaxation phase appears to be a critical  
595 factor in regulating the Chl hotspot in the SBC.

596 Due to its sheltered coastline, upwelling in the SC Bight is weak and intermittent  
597 compared to the Northern and Central California EBUS (Winant and Dorman 1997; Dorman and  
598 Winant 2000; Strub and James 2000). However, the SBC contains Chl in the spring that is  
599 higher than the rest of the SC Bight and similar in magnitude the coast north of Point  
600 Conception (Fig. 1, Mantyla et al. 1995; Henderikx Freitas et al. 2017; Kilpatrick et al. 2018).  
601 We hypothesize that eddy circulation when coupled with the relatively weak upwelling amplifies  
602 the seasonal Chl cycle in the SBC.

603 ***4.2 ENSO events point toward implications for understanding climate-driven changes in  
604 upwelling***

605 There has been much speculation about the impact of climate change on the intensity of  
606 upwelling in EBUSs and how this will affect phytoplankton primary production (Bograd et al.

607 2023). ENSO is one of the major drivers of interannual variability in the SC Bight and has been  
608 observed to disrupt typical patterns in spring primary productivity (Bograd and Lynn 2001; Shipe  
609 and Brzezinski 2001; Shipe et al. 2002). In 1998, an extreme El Niño event occurred in the  
610 California EBUS, producing unusually weak upwelling, unseasonably warm water temperatures,  
611 and record low Chl (Lynn et al. 1998; Kudela and Chavez 2002). For the 1998 SUP in the SBC,  
612 the mean Chl was less than 50% of the yearly average and only slightly greater than  
613 background levels (Table 1). The model predicts the weakest upwelling and the lowest  $N_{max}$  for  
614 the 1998 SUP of the 10-year modeling period. In addition to weak upwelling, the pattern of eddy  
615 circulation deviates significantly from a typical SUP with strong continuous eddy circulation  
616 lasting for two months. This is the longest period of uninterrupted eddy circulation for all SUPs  
617 and is very unusual as the average eddy lifespan for the SUPs is only 12 days. Although the  
618 prolonged eddy circulation produces record long  $RT_{eddy}$  during the 1998 SUP, neither the strong  
619 eddy circulation or the weak upwelling transports sufficient  $N_{max}$  into the surface layer to  
620 produce elevated Chl.

621 Following the extreme El Niño in 1998, an extreme La Niña event occurred in 1999 with  
622 record high upwelling throughout the California EBUS (Bograd et al. 2000; Schwing et al. 2000).  
623 The 1999 SUP displays the strongest upwelling of the 10-year modeling period with a mean  
624 PC1 and  $N_{max}$  that are 100% and 50% greater respectively than the yearly average (Table 1).  
625 However, the strong upwelling did not produce elevated Chl as the 1999 SUP has the lowest  
626 mean Chl of the 10-year modeling period, matching the mean Chl of the 1998 SUP (Table 1).  
627 During the 1999 SUP, maximum upwelling strength and  $N_{max}$  occurs in April and May. In  
628 contrast, eddy circulation and RT are unusually small during these months due to the fast  
629 eastward flow of CC water entering the SBC. Our results show that the flux of  $N_{max}$  through the  
630 SBC in April and May of 1999 is greater than the maximum flux of  $N_{max}$  for all other SUPs. Thus,  
631 we hypothesize that the unusually strong upwelling resulted in anomalously low Chl by

632 suppressing eddy circulation and RT to the point where phytoplankton biomass was not able to  
633 accumulate in the SBC.

634           Conceptual models of shelf chlorophyll for the Northern and Central California EBUS  
635 suggest that maximum shelf chlorophyll occurs when the strength of upwelling favorable winds  
636 fall into an optimal mid-range between weak and strong (Botsford et al. 2003; Stone et al. 2020).  
637 This optimal upwelling injects sufficient nutrients into the surface layer to stimulate  
638 phytoplankton growth, while allowing adequate retention of upwelled waters for phytoplankton  
639 nutrient uptake and accumulation. Work by Jacox et al. (2016a) also supports this conceptual  
640 model with ROMS simulations of the Northern and Central California EBUS for 13 years  
641 including the 1998 El Niño and 1999 La Niña. By comparing the SUPs from the typical years to  
642 the two anomalous years of 1998 and 1999, we propose the following conceptual model for the  
643 SBC SUP on how Chl is regulated by the interaction of upwelling and eddy circulation. High Chl  
644 production in typical years is associated with optimal conditions of intermediate upwelling  
645 strength and  $N_{max}$  flux, which allows sufficient injection of  $N_{max}$  into the surface layer and eddy  
646 circulation to retain  $N_{max}$  in the SBC. However, when upwelling strength and  $N_{max}$  flux are  
647 greater than optimal conditions, eddy circulation is too weak to retain  $N_{max}$  in the SBC, and  $N_{max}$   
648 is advected out of the SBC too quickly to produce high Chl. In contrast, when upwelling strength  
649 and  $N_{max}$  flux are below optimal conditions,  $N_{max}$  is too low to produce high Chl, and strong eddy  
650 circulation does not contribute enough  $N_{max}$  to the surface layer to produce high Chl.

651           Decadal climate oscillations may also impact our observations, but our ability to resolve  
652 these impacts is limited by the relatively short modeling period. In particular, the North Pacific  
653 Gyre Oscillation (NPGO) is thought to drive decadal variations in the southern CC ecosystem,  
654 including the SBC, but remained in a cold phase for most of the study period except for a short  
655 warm phase in 2005 and 2006 (Di Lorenzo et al. 2008; Catlett et al. 2021). The 2005 and 2006  
656 SUPs are associated with  $N_{max}$  slightly below, but close to, average (Table 1), suggesting that  
657 the NPGO warm phase did not exert a significant influence on SBC upwelling strength. Although

658 Chl is above average for the 2005 SUP and below average for the 2006 SUP, highly anomalous  
659 Chl are not observed in either year. Previous observations showed that anomalous advection of  
660 SCC water into the SBC in 2005 and 2006 was associated with unusual dinoflagellate bloom  
661 events (Catlett et al. 2021). However, since the multi-decadal climate variability and the  
662 variability in phytoplankton composition on Chl cannot be resolved in the present study, future  
663 work should be conducted on the influence of decadal climate oscillations on the SBC.

664 Climate-driven changes in upwelling patterns in EBUSs are a major ecological concern  
665 (Bograd et al. 2023). The weak upwelling and warm temperatures of the 1998 El Niño represent  
666 conditions of long marine heatwaves, extended periods of unusually warm water temperatures,  
667 which have detrimental effects on the ecology and have become more frequent in the SBC and  
668 EBUSs globally due to climate change (Cavole et al. 2016; Benthuysen et al. 2020; Michaud et  
669 al. 2022). In contrast, Bakun (1990) hypothesized that under climate change, upwelling rates in  
670 EBUSs would intensify due to the differential heating response between land and water. More  
671 recent studies have shown that upwelling has intensified across some EBUSs, although  
672 observed changes are spatially variable (Sydeman et al. 2014; Wang et al. 2015). For the  
673 California EBUS, a trend in increased upwelling strength has been observed for the Northern  
674 and Central California EBUS, but is not for the Southern California EBUS including the SC Bight  
675 (Jacox et al. 2015; Quilfen et al. 2021). The La Niña conditions in 1999 represent the potential  
676 consequences for the SBC and other coastal zones when strong upwelling reduces RT to an  
677 extent that does not allow nutrient uptake by phytoplankton. As the balance between the  
678 upwelling and eddy circulation regulates the Chl hotspot in the SBC, disruptions to this balance,  
679 such as those shown during the 1998 and 1999 SUPs, provide a framework for predicting the  
680 ecological consequences of climate-driven changes in upwelling for the future.

681  
682

683 **Acknowledgements**

684 This research was supported by the National Science Foundation Biological Oceanography

685 Program (OCE-2023693).

686

687

688 **References**

- 689 Anderson C, Brzezinski M, Washburn L, Kudela R (2006) Circulation and environmental  
690 conditions during a toxicogenic *Pseudo-nitzschia australis* bloom in the Santa Barbara  
691 Channel, California. *Mar Ecol Prog Ser* 327:119–133.  
692 <https://doi.org/10.3354/meps327119>
- 693 Bakun A (1990) Global climate change and intensification of coastal ocean upwelling. *Science*  
694 247:198–201. <https://doi.org/10.1126/science.247.4939.198>
- 695 Beckenbach E, Washburn L (2004) Low-frequency waves in the Santa Barbara Channel  
696 observed by high-frequency radar. *J Geophys Res* 109:C02010.  
697 <https://doi.org/10.1029/2003JC001999>
- 698 Beers JR, Trent JD, Reid FMH, Shanks AL (1986) Macroaggregates and their phytoplanktonic  
699 components in the Southern California Bight. *J Plankton Res* 8:475–487.  
700 <https://doi.org/10.1093/plankt/8.3.475>
- 701 Behrenfeld MJ, Boss E, Siegel DA, Shea DM (2005) Carbon-based ocean productivity and  
702 phytoplankton physiology from space. *Global Biogeochem Cycles* 19:GB1006.  
703 <https://doi.org/10.1029/2004GB002299>
- 704 BenthuySEN JA, Oliver ECJ, Chen K, Wernberg T (2020) Editorial: Advances in understanding  
705 marine heatwaves and their impacts. *Front Mar Sci* 7:147.  
706 <https://doi.org/10.3389/fmars.2020.00147>
- 707 Bograd S, D'Giacomo PM, Durazo R, et al (2000) The State of the California Current, 1999–  
708 2000: Forward to a new regime? *Calif Coop Fish Invest Rep* 41:26–52
- 709 Bograd SJ, Jacox MG, Hazen EL, et al (2023) Climate change impacts on Eastern Boundary  
710 Upwelling Systems. *Annu Rev Mar Sci* 15:303–328. <https://doi.org/10.1146/annurev-marine-032122-021945>
- 712 Bograd SJ, Lynn RJ (2001) Physical-biological coupling in the California Current during the  
713 1997–99 El Niño-La Niña Cycle. *Geophys Res Lett* 28:275–278.  
714 <https://doi.org/10.1029/2000GL012047>
- 715 Botsford LW, Lawrence CA, Dever EP, et al (2006) Effects of variable winds on biological  
716 productivity on continental shelves in coastal upwelling systems. *Deep Sea Res Part II:*  
717 *Top Stud Oceanogr* 53:3116–3140. <https://doi.org/10.1016/j.dsr2.2006.07.011>
- 718 Botsford LW, Lawrence CA, Dever EP, et al (2003) Wind strength and biological productivity in  
719 upwelling systems: an idealized study. *Fish Oceanogr* 12:245–259.  
720 <https://doi.org/10.1046/j.1365-2419.2003.00265.x>
- 721 Bray NA, Keyes A, Morawitz WML (1999) The California Current system in the Southern  
722 California Bight and the Santa Barbara Channel. *J Geophys Res* 104:7695–7714.  
723 <https://doi.org/10.1029/1998JC900038>

- 724 Brzezinski MA, Washburn L (2011) Phytoplankton primary productivity in the Santa Barbara  
725 Channel: Effects of wind-driven upwelling and mesoscale eddies. *J Geophys Res*  
726 116:C12013. <https://doi.org/10.1029/2011JC007397>
- 727 Carr SD, Capet XJ, McWilliams JC, et al (2008) The influence of diel vertical migration on  
728 zooplankton transport and recruitment in an upwelling region: estimates from a coupled  
729 behavioral-physical model. *Fish Oceanogr* 17:1–15. <https://doi.org/10.1111/j.1365-2419.2007.00447.x>
- 731 Catlett D, Siegel DA, Simons RD, et al (2021) Diagnosing seasonal to multi-decadal  
732 phytoplankton group dynamics in a highly productive coastal ecosystem. *Prog Oceanogr*  
733 197:102637. <https://doi.org/10.1016/j.pocean.2021.102637>
- 734 Cavole L, Demko A, Diner R, et al (2016) Biological impacts of the 2013–2015 warm-water  
735 anomaly in the Northeast Pacific: Winners, losers, and the future. *Oceanog* 29:273–285.  
736 <https://doi.org/10.5670/oceanog.2016.32>
- 737 Chen G, Hou Y, Chu X (2011) Mesoscale eddies in the South China Sea: Mean properties,  
738 spatiotemporal variability, and impact on thermohaline structure. *J Geophys Res*  
739 116:C06018. <https://doi.org/10.1029/2010JC006716>
- 740 Deutsch C, Frenzel H, McWilliams JC, et al (2021) Biogeochemical variability in the California  
741 Current System. *Prog Oceanogr* 196:102565.  
742 <https://doi.org/10.1016/j.pocean.2021.102565>
- 743 Dever EP, Hendershott MC, Winant CD (1998) Statistical aspects of surface drifter observations  
744 of circulation in the Santa Barbara Channel. *J Geophys Res* 103:24781–24797.  
745 <https://doi.org/10.1029/98JC02403>
- 746 Di Lorenzo E, Schneider N, Cobb KM, et al (2008) North Pacific Gyre Oscillation links ocean  
747 climate and ecosystem change. *Geophys Res Lett* 35:L08607.  
748 <https://doi.org/10.1029/2007GL032838>
- 749 Dong CM, Idica EY, McWilliams JC (2009) Circulation and multiple-scale variability in the  
750 Southern California Bight. *Prog Oceanogr* 82:168–190.  
751 <https://doi.org/10.1016/j.pocean.2009.07.005>
- 752 Dong CM, Lin XY, Liu Y, et al (2012) Three-dimensional oceanic eddy analysis in the Southern  
753 California Bight from a numerical product. *J Geophys Res-Oceans* 117:.  
754 <https://doi.org/10.1029/2011jc007354>
- 755 Dong CM, McWilliams JC (2007) A numerical study of island wakes in the Southern California  
756 Bight. *Cont Shelf Res* 27:1233–1248. <https://doi.org/10.1016/j.csr.2007.01.016>
- 757 Dong CM, McWilliams JC, Hall A, Hughes M (2011) Numerical simulation of a synoptic event in  
758 the Southern California Bight. *J Geophys Res-Oceans* 116:C05018.  
759 <https://doi.org/10.1029/2010jc006578>
- 760 Dorman CE, Winant CD (2000) The Structure and Variability of the Marine Atmosphere around  
761 the Santa Barbara Channel. *Monthly Weather Rev* 128:261.  
762 [https://doi.org/10.1175/1520-0493\(2000\)128<0261:TSAVOT>2.0.CO;2](https://doi.org/10.1175/1520-0493(2000)128<0261:TSAVOT>2.0.CO;2)

- 763 Dugan JE, Hubbard DM (2016) Sandy Beaches. In: Mooney HA, Zavaleta E (eds) Ecosystems  
764 of California. University of California Press, pp 389–409
- 765 Dugdale RC, Davis CO, Wilkerson FP (1997) Assessment of new production at the upwelling  
766 center at Point Conception, California, using nitrate estimated from remotely sensed sea  
767 surface temperature. *J Geophys Res* 102:8573–8585.  
768 <https://doi.org/10.1029/96JC02136>
- 769 Fennel K, Wilkin J (2009) Quantifying biological carbon export for the northwest North Atlantic  
770 continental shelves. *Geophys Res Lett* 36:L18605.  
771 <https://doi.org/10.1029/2009GL039818>
- 772 Fiedler PC, Reilly SB, Hewitt RP, et al (1998) Blue whale habitat and prey in the California  
773 Channel Islands. *Deep Sea Res Part II: Top Stud Oceanogr* 45:1781–1801.  
774 [https://doi.org/10.1016/S0967-0645\(98\)80017-9](https://doi.org/10.1016/S0967-0645(98)80017-9)
- 775 García-Reyes M, Largier JL (2012) Seasonality of coastal upwelling off central and northern  
776 California: New insights, including temporal and spatial variability. *J Geophys Res*  
777 117:C03028. <https://doi.org/10.1029/2011JC007629>
- 778 Harms S, Winant CD (1998) Characteristic patterns of the circulation in the Santa Barbara  
779 Channel. *J Geophys Res-Oceans* 103:3041–3065. <https://doi.org/10.1029/97jc02393>
- 780 Hayward TL, Venrick EL (1998) Nearsurface pattern in the California Current: coupling between  
781 physical and biological structure. *Deep Sea Res Part II: Top Stud Oceanogr* 45:1617–  
782 1638. [https://doi.org/10.1016/S0967-0645\(98\)80010-6](https://doi.org/10.1016/S0967-0645(98)80010-6)
- 783 Henderikx Freitas F, Siegel DA, Maritorena S, Fields E (2017) Satellite assessment of  
784 particulate matter and phytoplankton variations in the Santa Barbara Channel and its  
785 surrounding waters: Role of surface waves. *J Geophys Res Oceans* 122:355–371.  
786 <https://doi.org/10.1002/2016JC012152>
- 787 Henson SA, Thomas AC (2007) Phytoplankton scales of variability in the California Current  
788 System: 1. Interannual and cross-shelf variability. *J Geophys Res* 112:C07017.  
789 <https://doi.org/10.1029/2006JC004039>
- 790 Hickey BM (1993) Chapter 2. Physical Oceanography. In: Dailey MD, Reish DJ, Anderson JW  
791 (eds) *Ecology of the Southern California Bight*. University of California Press, pp 19–70
- 792 Jacox MG, Bograd SJ, Hazen EL, Fiechter J (2015) Sensitivity of the California Current nutrient  
793 supply to wind, heat, and remote ocean forcing. *Geophys Res Lett* 42:5950–5957.  
794 <https://doi.org/10.1002/2015GL065147>
- 795 Jacox MG, Edwards CA, Hazen EL, Bograd SJ (2018) Coastal Upwelling Revisited: Ekman,  
796 Bakun, and Improved Upwelling Indices for the U.S. West Coast. *J Geophys Res*  
797 *Oceans* 123:7332–7350. <https://doi.org/10.1029/2018JC014187>
- 798 Jacox MG, Hazen EL, Bograd SJ (2016a) Optimal environmental conditions and anomalous  
799 ecosystem responses: constraining bottom-up controls of phytoplankton biomass in the  
800 California Current System. *Sci Rep* 6:27612. <https://doi.org/10.1038/srep27612>

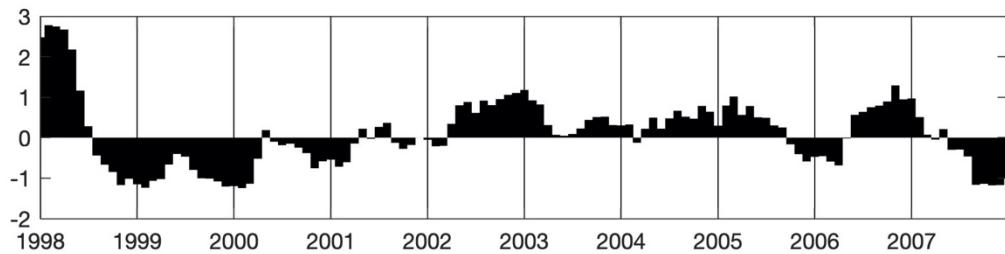
- 801 Jacox MG, Hazen EL, Zaba KD, et al (2016b) Impacts of the 2015–2016 El Niño on the  
802 California Current System: Early assessment and comparison to past events. *Geophys*  
803 *Res Lett* 43:7072–7080. <https://doi.org/10.1002/2016GL069716>
- 804 Kahru M, Kudela RM, Anderson CR, Mitchell BG (2015) Optimized merger of ocean chlorophyll  
805 algorithms of MODIS-Aqua and VIIRS. *IEEE Geosci Remote Sensing Lett* 12:2282–  
806 2285. <https://doi.org/10.1109/LGRS.2015.2470250>
- 807 Kahru M, Kudela RM, Manzano-Sarabia M, Greg Mitchell B (2012) Trends in the surface  
808 chlorophyll of the California Current: Merging data from multiple ocean color satellites.  
809 *Deep Sea Res Part II: Top Stud Oceanogr* 77–80:89–98.  
810 <https://doi.org/10.1016/j.dsr2.2012.04.007>
- 811 Kilpatrick T, Xie S, Miller AJ, Schneider N (2018) Satellite observations of enhanced chlorophyll  
812 variability in the Southern California Bight. *J Geophys Res: Oceans* 123:7550–7563.  
813 <https://doi.org/10.1029/2018JC014248>
- 814 Krause JW, Brzezinski MA, Siegel DA, Thunell RC (2013) Biogenic silica standing stock and  
815 export in the Santa Barbara Channel ecosystem. *J Geophys Res Oceans* 118:736–749.  
816 <https://doi.org/10.1029/2012JC008070>
- 817 Kudela RM, Chavez FP (2002) Multi-platform remote sensing of new production in central  
818 California during the 1997–1998 El Niño. *Progr Oceanogr* 54:233–249.  
819 [https://doi.org/10.1016/S0079-6611\(02\)00051-4](https://doi.org/10.1016/S0079-6611(02)00051-4)
- 820 Lachkar Z, Gruber N (2011) What controls biological production in coastal upwelling systems?  
821 Insights from a comparative modeling study. *Biogeosci* 8:2961–2976.  
822 <https://doi.org/10.5194/bg-8-2961-2011>
- 823 Legaard KR, Thomas AC (2006) Spatial patterns in seasonal and interannual variability of  
824 chlorophyll and sea surface temperature in the California Current. *J Geophys Res*  
825 111:C06032. <https://doi.org/10.1029/2005JC003282>
- 826 Levin LA, Liu K-K, Emeis K-C, et al (2015) Comparative biogeochemistry–ecosystem–human  
827 interactions on dynamic continental margins. *J Mar Syst* 141:3–17.  
828 <https://doi.org/10.1016/j.jmarsys.2014.04.016>
- 829 Liu Y, Dong C, Guan Y, et al (2012) Eddy analysis in the subtropical zonal band of the North  
830 Pacific Ocean. *Deep Sea Res Part I: Oceanogr Res Papers* 68:54–67.  
831 <https://doi.org/10.1016/j.dsr.2012.06.001>
- 832 Lynn KJ, Baumgaktnek T, Garcia J, et al (1998) The state of the California Current, 1997–1998:  
833 Transition to El Niño Conditions. *Calif Coop Fish Invest Rep* 39:25–49
- 834 Mantyla AW, Venrick EL, Hayward TL (1995) Primary production and chlorophyll relationships,  
835 derived from ten years of CalCOFI measurements. *Calif Coop Fish Invest Rep* 36:159–  
836 166
- 837 Matson PG, Washburn L, Fields EA, et al (2019) Formation, development, and propagation of a  
838 rare coastal Coccolithophore bloom. *J Geophys Res: Oceans* 124:3298–3316.  
839 <https://doi.org/10.1029/2019JC015072>

- 840 McClatchie S, Jacox MG, Ohman MD, et al (2016) State of the California Current 2015–16:  
841 Comparisons with the 1997–98 El Nino. *Calif Coop Fish Invest Rep* 57:5–61
- 842 McGillicuddy DJ (2016) Mechanisms of physical-biological-biogeochemical interaction at the  
843 oceanic mesoscale. *Annu Rev Mar Sci* 8:125–159. <https://doi.org/10.1146/annurev-marine-010814-015606>
- 845 McPhee-Shaw EE, Siegel DA, Washburn L, et al (2007) Mechanisms for nutrient delivery to the  
846 inner shelf: Observations from the Santa Barbara Channel. *Limnol Oceanogr* 52:1748–  
847 1766. <https://doi.org/10.4319/lo.2007.52.5.1748>
- 848 Melton C, Washburn L, Gotschalk C (2009) Wind relaxations and poleward flow events in a  
849 coastal upwelling system on the central California coast. *J Geophys Res* 114:C11016.  
850 <https://doi.org/10.1029/2009JC005397>
- 851 Messié M, Chavez FP (2015) Seasonal regulation of primary production in eastern boundary  
852 upwelling systems. *Progr Oceanogr* 134:1–18.  
853 <https://doi.org/10.1016/j.pocean.2014.10.011>
- 854 Michaud KM, Reed DC, Miller RJ (2022) The Blob marine heatwave transforms California kelp  
855 forest ecosystems. *Commun Biol* 5:1143. <https://doi.org/10.1038/s42003-022-04107-z>
- 856 Miller RJ, Reed DC, Brzezinski MA (2011) Partitioning of primary production among giant kelp  
857 (*Macrocystis pyrifera*), understory macroalgae, and phytoplankton on a temperate reef.  
858 *Limnol Oceanogr* 56:119–132. <https://doi.org/10.4319/lo.2011.56.1.0119>
- 859 Mitarai S, Siegel DA, Watson JR, et al (2009) Quantifying connectivity in the coastal ocean with  
860 application to the Southern California Bight. *J Geophys Res-Oceans* 114:C10026.  
861 <https://doi.org/10.1029/2008JC005166>
- 862 Nencioli F, Dong C, Dickey T, et al (2010) A vector geometry-based eddy detection algorithm  
863 and its application to a high-resolution numerical model product and high-frequency  
864 radar surface velocities in the Southern California Bight. *J Atmos Ocean Technol*  
865 27:564–579. <https://doi.org/10.1175/2009JTECHO725.1>
- 866 Nishimoto M, Washburn L (2002) Patterns of coastal eddy circulation and abundance of pelagic  
867 juvenile fish in the Santa Barbara Channel, California, USA. *Mar Ecol Prog Ser* 241:183–  
868 199. <https://doi.org/10.3354/meps241183>
- 869 Oey L-Y, Wang D-P, Hayward T, et al (2001) “Upwelling” and “cyclonic” regimes of the near-  
870 surface circulation in the Santa Barbara Channel. *J Geophys Res* 106:9213–9222.  
871 <https://doi.org/10.1029/1999JC000129>
- 872 Oey L-Y, Winant C, Dever E, et al (2004) A model of the near-surface circulation of the Santa  
873 Barbara Channel: Comparison with observations and dynamical interpretations. *J Phys  
874 Oceanogr* 34:23–43. [https://doi.org/10.1175/1520-0485\(2004\)034<0023:AMOTNC>2.0.CO;2](https://doi.org/10.1175/1520-0485(2004)034<0023:AMOTNC>2.0.CO;2)
- 876 Ohlmann JC, Mitarai S (2010) Lagrangian assessment of simulated surface current dispersion  
877 in the coastal ocean. *Geophys Res Lett* 37:L17602.  
878 <https://doi.org/10.1029/2010GL044436>

- 879 Omand MM, Feddersen F, Guza RT, Franks PJS (2012) Episodic vertical nutrient fluxes and  
880 nearshore phytoplankton blooms in Southern California. *Limnol Oceanogr* 57:1673–  
881 1688. <https://doi.org/10.4319/lo.2012.57.6.1673>
- 882 Otero M, Siegel D (2004) Spatial and temporal characteristics of sediment plumes and  
883 phytoplankton blooms in the Santa Barbara Channel. *Deep Sea Res Part II: Top Stud*  
884 *Oceanogr* 51:1129–1149. [https://doi.org/10.1016/S0967-0645\(04\)00104-3](https://doi.org/10.1016/S0967-0645(04)00104-3)
- 885 Palacios DM, Hazen EL, Schroeder ID, Bograd SJ (2013) Modeling the temperature-nitrate  
886 relationship in the coastal upwelling domain of the California Current: Modeling the T-N  
887 Relationship. *J Geophys Res Oceans* 118:3223–3239.  
888 <https://doi.org/10.1002/jgrc.20216>
- 889 Quilfen Y, Shutler J, Piolle J-F, Autret E (2021) Recent trends in the wind-driven California  
890 current upwelling system. *Remote Sens Environ* 261:112486.  
891 <https://doi.org/10.1016/j.rse.2021.112486>
- 892 Santora JA, Sydeman WJ, Schroeder ID, et al (2017) Persistence of trophic hotspots and  
893 relation to human impacts within an upwelling marine ecosystem. *Ecol Appl* 27:560–574.  
894 <https://doi.org/10.1002/eap.1466>
- 895 Santoro AE, Nidzieko NJ, Dijken GL van, et al (2010) Contrasting spring and summer  
896 phytoplankton dynamics in the nearshore Southern California Bight. *Limnol Oceanogr*  
897 55:264–278. <https://doi.org/10.4319/lo.2010.55.1.0264>
- 898 Schwing FB, Moore CS, Ralston S, Sakuma KM (2000) Record coastal upwelling in the  
899 California Current. *Calif Coop Fish Invest Rep* 41:148–160
- 900 Shchepetkin AF, McWilliams JC (2005) The regional oceanic modeling system (ROMS): a split-  
901 explicit, free-surface, topography-following-coordinate oceanic model. *Ocean Model*  
902 9:347–404. <https://doi.org/10.1016/j.ocemod.2004.08.002>
- 903 Shipe RF, Brzezinski MA (2001) A time series study of silica production and flux in an eastern  
904 boundary region: Santa Barbara Basin, California. *Global Biogeochem Cycles* 15:517–  
905 531. <https://doi.org/10.1029/2000GB001297>
- 906 Shipe RF, Passow U, Brzezinski MA, et al (2002) Effects of the 1997–98 El Niño on seasonal  
907 variations in suspended and sinking particles in the Santa Barbara basin. *Progr*  
908 *Oceanogr* 54:105–127. [https://doi.org/10.1016/S0079-6611\(02\)00045-9](https://doi.org/10.1016/S0079-6611(02)00045-9)
- 909 Siegel DA, Behrenfeld MJ, Maritorena S, et al (2013) Regional to global assessments of  
910 phytoplankton dynamics from the SeaWiFS mission. *Remote Sensing of Environment*  
911 135:77–91. <https://doi.org/10.1016/j.rse.2013.03.025>
- 912 Simons RD, Nishimoto MM, Washburn L, et al (2015) Linking kinematic characteristics and high  
913 concentrations of small pelagic fish in a coastal mesoscale eddy. *Deep Sea Res Part I:*  
914 *Oceanogr Res Papers* 100:34–47. <https://doi.org/10.1016/j.dsr.2015.02.002>
- 915 Snyder JN, Bell TW, Siegel DA, et al (2020) Sea surface temperature imagery elucidates  
916 spatiotemporal nutrient patterns for offshore kelp aquaculture siting in the Southern  
917 California Bight. *Front Mar Sci* 7:22. <https://doi.org/10.3389/fmars.2020.00022>

- 918 Stone HB, Banas NS, MacCready P, et al (2020) Linking chlorophyll concentration and wind  
919 patterns using satellite data in the Central and Northern California Current System. *Front*  
920 *Mar Sci* 7:551562. <https://doi.org/10.3389/fmars.2020.551562>
- 921 Strub PT, James C (2000) Altimeter-derived variability of surface velocities in the California  
922 Current System: 2. Seasonal circulation and eddy statistics. *Deep Sea Res Part II: Top*  
923 *Stud Oceanogr* 47:831–870. [https://doi.org/10.1016/S0967-0645\(99\)00129-0](https://doi.org/10.1016/S0967-0645(99)00129-0)
- 924 Sydeman WJ, García-Reyes M, Schoeman DS, et al (2014) Climate change and wind  
925 intensification in coastal upwelling ecosystems. *Science* 345:77–80.  
926 <https://doi.org/10.1126/science.1251635>
- 927 Thomson RE, Emery WJ (2014) The Spatial Analyses of Data Fields. In: *Data Analysis Methods*  
928 in *Physical Oceanography*. Elsevier, pp 313–424
- 929 Thunell R, Benitez-Nelson C, Varela R, et al (2007) Particulate organic carbon fluxes along  
930 upwelling-dominated continental margins: Rates and mechanisms. *Global Biogeochem*  
931 *Cycles* 21:GB1022. <https://doi.org/10.1029/2006GB002793>
- 932 Venrick EL (1998) The phytoplankton of the Santa Barbara Basin: Patterns of chlorophyll and  
933 species structure and their relationships with those of surrounding stations. *Calif Coop*  
934 *Fish Invest Rep* 39:124–132
- 935 Wang D, Gouhier TC, Menge BA, Ganguly AR (2015) Intensification and spatial homogenization  
936 of coastal upwelling under climate change. *Nature* 518:390–394.  
937 <https://doi.org/10.1038/nature14235>
- 938 Wang X, Du Y, Zhang Y, et al (2021) Influence of two eddy pairs on high-salinity water intrusion  
939 in the Northern South China Sea during fall-winter 2015/2016. *J Geophys Res: Oceans*  
940 126:e2020JC016733. <https://doi.org/10.1029/2020JC016733>
- 941 Washburn L, McPhee-Shaw E (2013) Coastal Transport Processes Affecting Inner-Shelf  
942 Ecosystems in the California Current System. *oceanog* 26:34–43.  
943 <https://doi.org/10.5670/oceanog.2013.43>
- 944 Wilkerson FP, Lassiter AM, Dugdale RC, et al (2006) The phytoplankton bloom response to  
945 wind events and upwelled nutrients during the CoOP WEST study. *Deep Sea Res Part*  
946 *II: Top Stud Oceanogr* 53:3023–3048. <https://doi.org/10.1016/j.dsr2.2006.07.007>
- 947 Winant CD, Dever EP, Hendershott MC (2003) Characteristic patterns of shelf circulation at the  
948 boundary between central and southern California. *J Geophys Res* 108:3021.  
949 <https://doi.org/10.1029/2001JC001302>
- 950 Winant CD, Dorman CE (1997) Seasonal patterns of surface wind stress and heat flux over the  
951 Southern California Bight. *J Geophys Res* 102:5641–5653.  
952 <https://doi.org/10.1029/96JC02801>
- 953
- 954

955      **Supplementary Information**  
956



957  
958      Figure S1: MEI (pls.noaa.gov/enso/mei, Wolter and Timlin 1993, 1998)  
959  
960

961      Comparison of temperature from ROMS to field observations during the SUP

962           The accuracy of the temperature predictions from ROMS in the SBC surface layer is  
963           important for this study as  $N_{max}$ , representing nutrient availability, is estimated from temperature  
964           (see Section 2.4). To determine the accuracy of the temperature during the SUP, field  
965           observations of temperature at 8 stations in the SBC (Figure S2) are compared to temperature  
966           predictions from ROMS. The stations are located along the SBC coastline; four (ALE, NAP,  
967           MKO, and CAR) on the mainland, three (PEL, SCP, and TRL) on the north coast of Santa Cruz  
968           Island, and one on the coast of Anacapa Island (ANS). At these stations, continuously logged  
969           temperature data at 14 m depth starting from 2000-2004 through 2007 is available from the  
970           Santa Barbara Coastal Long-Term Ecological Research site (sbcler.msi.ucsb.edu, Washburn et  
971           al. 2022).

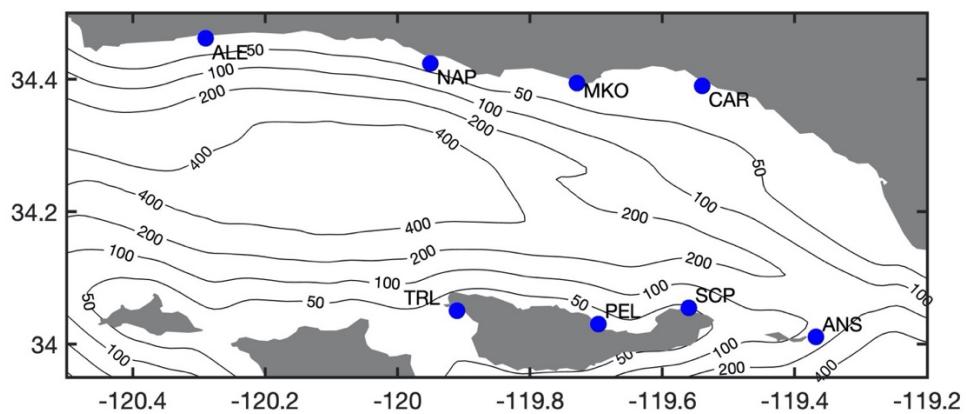
972           For each year from 2000-2007, the root-mean-square error (RMSE) between the  
973           temperature from ROMS and the field station during the SUP is calculated and presented in  
974           Tables S1. The model and field temperature show good agreement with RMSE ranging from  
975           0.6 to 1.6°C per year when averaged over all stations. The year that shows the best  
976           agreement is the 2001 SUP with a RMSE ranging from 0.6 to 0.8°C per station. ROMS and  
977           field temperature data for the 2001 SUP is shown in Figure 3S for six stations.

978

Station	2000	2001	2002	2003	2004	2005	2006	2007
ALE	1.2	0.7	1.1	1.6	0.9	1.4	1.2	1.0
NAP	NA	0.6	1.1	1.3	1.4	1.5	1.2	1.4
MKO	NA	NA	NA	2.8	2.3	1.1	2.1	1.9
CAR	NA	0.8	1.0	2.3	2.3	1.2	1.9	1.8
PEL	1.4	0.5	1.1	1.0	1.1	1.3	1.6	1.7
SCP	1.3	0.5	1.3	1.0	1.0	1.4	1.5	1.6
TRL	1.2	0.6	0.8	1.0	0.8	1.8	1.2	1.5
ANS	NA	NA	NA	1.6	1.6	1.5	1.3	1.6
Mean	1.3	0.6	1.1	1.6	1.4	1.4	1.5	1.6

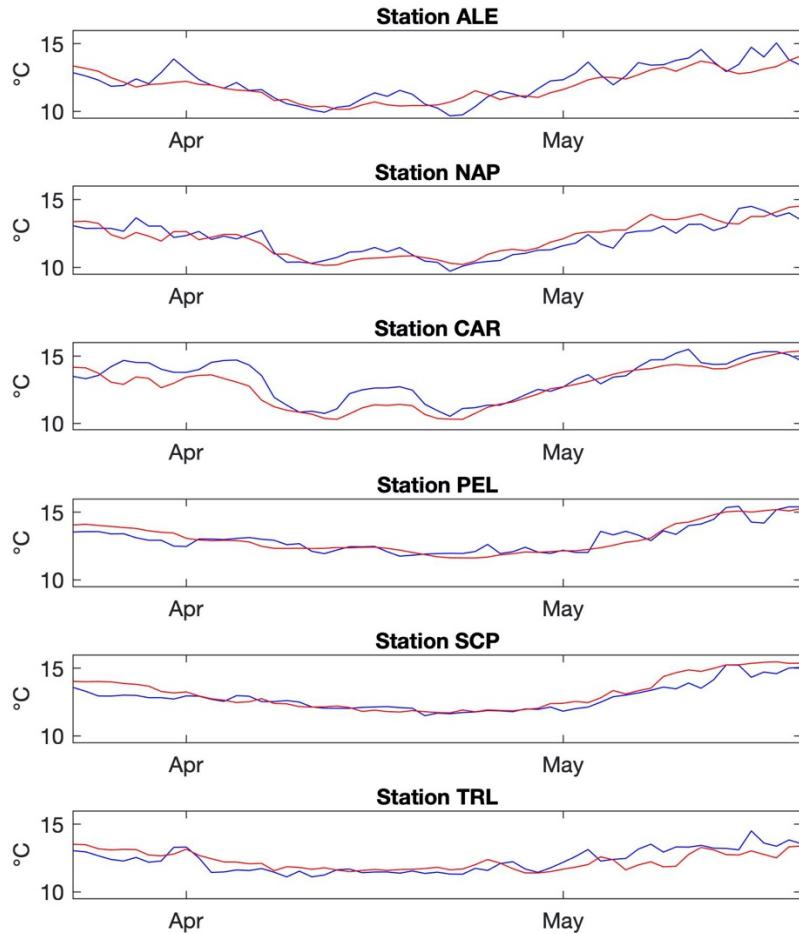
979 Table S1: Temperature RMSE ( $^{\circ}\text{C}$ ) between ROMS and field station data

980



981  
982  
983

Figure S2: Santa Barbara Channel with bathymetry (meters) and temperature stations (blue circles).



984

985 Figure S3: Temperature data for the 2001 SUP. The blue line is the field data, and the red line  
 986 is the ROMS data.

987

988 References

989 Washburn L, Gotschalk C, Salazar D (2022) SBC LTER: Ocean: Currents and Biogeochemistry:  
 990 Moored CTD and ADCP data from Alegria Reef Mooring (ALE), ongoing since 1999

991

992 Wolter K, Timlin MS (1993) Monitoring ENSO in COADS with a Seasonally Adjusted Principal  
 993 Component Index. Proceedings of the 17th Climate Diagnostics Workshop, Norman, 18-23  
 994 October 1992 52-57

995

996 Wolter K, Timlin MS (1998) Measuring the strength of ENSO events: How does 1997/98 rank?  
 997 Weather 53:315-324. <https://doi.org/10.1002/j.1477-8696.1998.tb06408.x>

998

999