

# 1 Understanding the resilient carbon cycle response to the 2014–2015 2 Blob event in the Gulf of Alaska using a regional ocean 3 biogeochemical model

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10 **Abstract.** Marine heatwaves (MHWs), characterized by anomalously high sea surface temperatures, are increasing in  
11 frequency and intensity and strongly impact ocean circulation, biogeochemistry, and marine ecosystems. During the 2014–  
12 2015 MHW (commonly called the Blob) in the NE subarctic Pacific, moored observations at Ocean Station Papa (OSP; 145°W,  
13 50°N) showed a moderate decrease in oceanic pCO<sub>2</sub>, contrary to the increase expected from warming-induced solubility  
14 reduction alone. Using a regional model that reproduces the observed pCO<sub>2</sub> variability and trend at OSP, we show that this  
15 decline resulted from a decrease in dissolved inorganic carbon (DIC) supply that outweighed the warming-driven increase in  
16 pCO<sub>2</sub>. The DIC reduction was primarily caused by weakened vertical transport associated with enhanced upper-ocean  
17 stratification and reduced Ekman pumping prior to the onset of the Blob, which suppressed the upwelling of DIC-rich  
18 subsurface waters. Horizontal transport also contributed locally, particularly at OSP. These results demonstrate that anomalous  
19 physical circulation, rather than biological processes, was the primary driver of the enhanced CO<sub>2</sub> uptake during the Blob and  
20 highlight the importance of resolving physical transport mechanisms when assessing carbon cycle responses to extreme  
21 warming events.

## 22 1 Introduction

23 Approximately 25% of historical anthropogenic carbon emission has been absorbed by the oceans (Friedlingstein et al., 2022),  
24 and the North Pacific is one of the major regions for ocean carbon uptake (Takahashi et al., 2009). The carbon exchange  
25 between the atmosphere and ocean is closely related to ocean temperature through its influences on the solubility of carbon  
26 dioxide (CO<sub>2</sub>) in seawater. In recent years, anomalously high ocean temperatures, known as marine heatwaves (MHWs;  
27 Hobday et al., 2016), have been frequently observed. These events are defined by prolonged periods of unusually high sea

28 surface temperature (SST) anomalies and their intensity. With the ongoing progression of global warming, the frequency and  
29 intensity of MHWs have been increasing (Frölicher et al., 2018; Oliver et al., 2018). In particular, the persistent MHW that  
30 occurred in the subarctic NE Pacific from the winter of 2013 to 2015, known as *the Blob* (Bond et al., 2015), had severe  
31 impacts on surrounding ecosystems (Cavole et al., 2016; Smale et al., 2019) and fisheries (Barbeaux et al., 2020). This  
32 anomalously high ocean temperature was attributed to weaker surface winds, which reduced ocean surface heat loss, and  
33 weakened horizontal and vertical mixing (Bond et al., 2015; Di Lorenzo and Mantua, 2016).

34

35 Biogeochemical variables are also impacted by MHWs. In particular, low-oxygen and acidification become more pronounced  
36 during MHWs (Gruber et al., 2021; Burger et al., 2022; Li et al., 2024b). These changes are driven by alterations in ocean  
37 circulation due to changes in wind and air–sea interactions associated with atmospheric anomalies that induce the MHWs,  
38 and also by the direct effects of high temperatures (e.g. enhanced stratification, reduced solubility, and changes in biological  
39 activity). During the Blob, both surface dissolved oxygen and inorganic carbon (DIC) were significantly decreased partly due  
40 to reduced solubility caused by increased water temperatures (Mogen et al., 2022; based on ocean model outputs from NorESM  
41 and MRI simulations). Franco et al. (2021) showed a decrease in surface DIC during the Blob using in situ time-series  
42 measurements.

43

44 For the air–sea CO<sub>2</sub> exchange, the uptake in the North Pacific decreased by 29±11% during persistent MHWs mainly due to  
45 the direct effect of increased water temperature (Mignot et al., 2022). However, this reduction in ocean carbon uptake is  
46 primarily driven by the substantial increase in oceanic CO<sub>2</sub> outgassing in the tropical Pacific. Focusing on the subarctic regions,  
47 the oceanic CO<sub>2</sub> uptake increased during the MHWs. For example, mooring data from Ocean Station Papa (OSP; 50°N, 145°W)  
48 in the eastern North Pacific, which is one of the longest-running observation sites, showed low surface DIC and oceanic partial  
49 pressure of CO<sub>2</sub> (pCO<sub>2</sub>) during a more recent MHW (Kohlman et al., 2024). In the Gulf of Alaska (GOA), air–sea CO<sub>2</sub> flux  
50 exhibited a negative anomaly (stronger ocean uptake) of approximately 45% relative to the climatological monthly means  
51 during the Blob based on a machine-learning based reconstruction (Duke et al., 2023). The enhanced ocean carbon uptake is  
52 caused by the decrease in oceanic pCO<sub>2</sub> in the subarctic North Pacific, which is driven by the effects of reduced surface ocean  
53 DIC. However, the processes responsible for this DIC reduction is not fully understood (Mignot et al., 2022; Li et al., 2024a).  
54 Furthermore, estimates of oceanic pCO<sub>2</sub> changes derived from observation-based products are subject to considerable  
55 uncertainty, as observational data in the subarctic North Pacific are temporally and spatially sparse. Consequently,  
56 discrepancies among data products restrict the robustness of their assessments (Li et al., 2024a).

57

58 To elucidate the mechanisms by which the MHWs lead to the observed changes in oceanic pCO<sub>2</sub> and air–sea CO<sub>2</sub> fluxes,  
59 numerical models can provide the carbon mass balance and exploring underlying physical and biogeochemical processes.  
60 Physical processes that control the regional ocean circulation, biogeochemical cycling, and air–sea gas exchanges often occur  
61 at scales smaller than several tens of kilometres, which remain unresolved in global earth system models (Gruber et al., 2021).

62 In the subarctic NE Pacific, reproducing the full variability of oceanic pCO<sub>2</sub> in models is complicated by boundary currents  
63 and eddies that transport macro- and micro-nutrients, alkalinity and DIC. A previous study using a coarse resolution model  
64 (Mckinley et al., 2006) has shown that although models captured the variability of individual components of oceanic pCO<sub>2</sub> on  
65 seasonal and interannual timescales, they have not fully reproduced the total variability of oceanic pCO<sub>2</sub>, and their ability to  
66 simulate the temporal patterns of air–sea CO<sub>2</sub> fluxes remains limited.

67  
68 There are several regional modeling studies exploring biogeochemical processes in the GOA. Coyle et al. (2012) developed  
69 regional, biogeochemical simulations for the northern GOA based on the ROMS circulation model with the horizontal  
70 resolution of 3 km from 2001 to 2004 focusing on the coastal regions. The biogeochemical model component is based on the  
71 GOANPZ model (Hinckley et al., 2009). The model reproduced the spring phytoplankton bloom from the continental shelves  
72 to the open ocean. This model subsequently was used to analyze ocean acidification with the inclusion of carbonate chemistry  
73 (Siedlecki et al., 2017). Hauri et al. (2020) developed a hindcast simulation for the similar northern GOA domain. They also  
74 used the ROMS physical model with 4.5 km horizontal resolution. The biogeochemical component is based on the COBALT  
75 biogeochemistry model (Stock et al., 2014) with specific modifications to capture coastal ecosystems. Following these previous  
76 studies, we constructed a regional biogeochemical and carbon cycle model for the GOA. The model has been validated against  
77 a suite of physical and biogeochemical observations (Ito et al., 2026) This study expands the model to include a validated  
78 carbon cycle component. The simulated temporal variability of oceanic pCO<sub>2</sub> is validated with the NOAA (Pacific Marine  
79 Environmental Laboratory’s Ocean Climate Stations and Carbon groups) mooring at OSP (Emerson et al., 2011; Cronin et al.,  
80 2015). The model outputs are used to understand the underlying mechanisms and to quantify the changes in oceanic pCO<sub>2</sub> in  
81 this critical carbon sink during the Blob. This paper includes the description of the model and the observational data used for  
82 model validation (section 2), and the results of oceanic pCO<sub>2</sub> variations during the Blob and its causes (section 3). Section 4  
83 discusses the associated changes in CO<sub>2</sub> flux, the potential impacts of high ocean temperature anomalies on the regional carbon  
84 cycle, their effects on the broader ocean basin, and possible future implications. Finally, Section 5 summarizes the main  
85 findings of this study.

## 86 **2 Data and Methods**

### 87 **2.1 Model description**

88 The regional ocean circulation and biogeochemistry model used in this study followed the configuration described in Ito et al.  
89 (2026), thus only a brief description was provided here, while full details can be found in their paper. The model was based on  
90 MITgcm (Marshall et al., 1997a, b) combined with an ocean biogeochemical model, Biogeochemistry with Light, Iron,  
91 Nutrients and Gases (BLING) version 2 (Dunne et al., 2020) including 10 tracers, DIC, alkalinity, O<sub>2</sub>, PO<sub>4</sub>, NO<sub>3</sub>, dissolved Fe,  
92 silica, dissolved organic P, and dissolved organic N. The model was driven by 3-hourly averaged atmospheric variables from  
93 the JRA-55-do reanalysis data (Tsujino et al., 2018), including the surface air temperature, humidity, 10 m wind stress,

94 precipitation (both rain and snow), river runoff, and downward shortwave and longwave radiation. The earlier simulations of  
95 Ito et al. (2026) included a positive bias in sea surface salinity (SSS) of approximately +0.2 psu. To maintain SSSs close to the  
96 observations, we applied Newtonian relaxation to SSS toward monthly climatology from World Ocean Atlas 2023 (Reagan et  
97 al., 2024) with a restoring timescale of 30 days in the surface grid box of 10 m thickness. The model had a horizontal resolution  
98 of 10 km with 42 vertical layers on a latitude-longitude z-level grid. The bathymetry was generated by interpolating the  
99 ETOPO2 global 2-minute resolution topography dataset. The model domain had a southern open boundary at 42°N and a  
100 western open boundary at 160°W. Vertical mixing was parameterized using the K-profile Parameterization (KPP) scheme  
101 (Large et al., 1994) as implemented in MITgcm. The default configuration, including the non-local convection term, was used  
102 for this study. KPP diffusivities were applied to momentum and all tracers. A sea ice model, implemented using the MITgcm  
103 sea ice package (Losch et al., 2010) and coupled to the ocean component, was used to represent sea ice dynamics and  
104 thermodynamics, as well as their influence on surface heat, freshwater, and momentum fluxes.

105

106 Both the initial and boundary conditions in our model also followed Ito et al. (2026). Open boundary conditions were set to  
107 the climatological values from GLODAPv2 (Lauvset et al., 2022) for most of the biogeochemical properties. However, for  
108 DIC, the open boundary conditions included time-dependent anthropogenic carbon rather than climatology. Temporal changes  
109 in DIC due to anthropogenic influences were imposed according to the rate of change in atmospheric CO<sub>2</sub> fractions measured  
110 at the Mauna Loa Observatory (Keeling et al., 2001) and its spatial structure was set to the anthropogenic carbon estimates  
111 from the GLODAPv2. The model integration was performed through 2017, and the physical open boundary conditions were  
112 taken from the oceanic reanalysis data of Simple Ocean Data Assimilation (SODA) version 3 (Carton and Giese, 2008; Carton  
113 et al., 2018).

114

115 Several biogeochemical parameters were adjusted to improve the model representation of the carbon cycle. These  
116 modifications primarily involved processes related to strength of photosynthesis and remineralization. The modeled default  
117 parameters for photosynthetic rates caused excessively high rates in coastal waters, resulting in unrealistically elevated primary  
118 production. To address this, the self-shading effect of phytoplankton was turned on, which regulated the light available for  
119 phytoplankton growth in the highly productive coastal waters. The bio-optical parameterization of Manizza et al. (2005) was  
120 used in our configuration but the background attenuation coefficient in this bio-optical model was reduced to one-quarter of  
121 its default value to re-calibrate the net primary production (NPP). Furthermore, the simulated oceanic pCO<sub>2</sub> was highly  
122 sensitive to the production of organic matter and the subsequent remineralization. A key factor was the export ratio which  
123 determines the fraction of NPP that sank as particulate organic matter from the surface layer and was subsequently recycled  
124 in the subsurface waters. Among the relevant parameters, the partitioning of particulate organic versus dissolved organic matter  
125 exerted a particularly strong influence on the representation of oceanic pCO<sub>2</sub>. To better capture observed oceanic pCO<sub>2</sub>, we  
126 adjusted this parameter so that 30% of NPP is converted to dissolved organic matter, compared with a default value of 10%,  
127 which meant that more organic matter was recycled in the surface layer than the default configuration.

## 128 2.2 Observations

129 To evaluate the model skills in reproducing the observed oceanic pCO<sub>2</sub>, SeaFlux data (Fay et al., 2021) were used for spatial  
130 comparisons, and NOAA CO<sub>2</sub> mooring data (Emerson et al., 2011; Cronin et al., 2015) were used for temporal comparisons  
131 at OSP. SeaFlux data consisted of six global observation-based gridded products that reconstruct spatiotemporally continuous  
132 estimated of air–sea CO<sub>2</sub> fluxes using different gap-fill methods on the surface oceanic pCO<sub>2</sub> observations. Because the spatial  
133 resolution of SeaFlux was one degree in latitude and longitude, our model output was remapped to the same resolution using  
134 nearest-neighbour interpolation for comparison. In June 2007, a surface CO<sub>2</sub> mooring time series was initiated at OSP by S.  
135 Emerson for the study of North Pacific Carbon Cycle, and the mooring observation has been continued by NOAA Office of  
136 Climate Observations. This data provided both physical and biogeochemical variables. For comparison with the model,  
137 temperature, salinity, and both oceanic and atmospheric pCO<sub>2</sub> data were obtained from the mooring dataset archived at  
138 [https://www.ncei.noaa.gov/access/ocean-carbon-acidification-data-system/oceans/Moorings/Papa\\_145W\\_50N.html](https://www.ncei.noaa.gov/access/ocean-carbon-acidification-data-system/oceans/Moorings/Papa_145W_50N.html) (last  
139 access: 1 Oct 2025). For comparison, OSP in the model was represented by averaging all grid points within a 24 km radius of  
140 50°N, 145°W.

141  
142 Model skill was further evaluated against the spatial patterns of monthly SST anomalies from the OISST dataset (0.25°  
143 resolution; Reynolds et al., 2007; Huang et al., 2021) and the HadISST dataset (1° resolution; Rayner et al., 2003).

## 144 2.3 Analysis methods

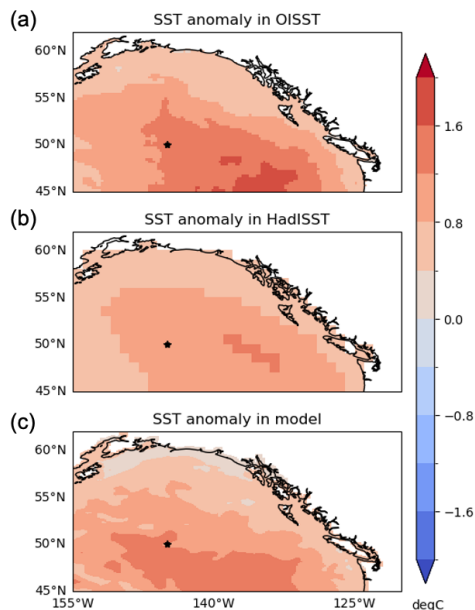
145 The oceanic pCO<sub>2</sub> anomalies were decomposed as:

$$146 \delta pCO_2 \approx \frac{\partial pCO_2}{\partial T} \delta T + \frac{\partial pCO_2}{\partial S} \delta S + \frac{\partial pCO_2}{\partial DIC^*} \delta DIC^* + \frac{\partial pCO_2}{\partial ALK^*} \delta ALK^* \quad (1)$$

148  
149 where oceanic pCO<sub>2</sub> was a function of SST (T), SSS (S), sea surface salinity-normalized DIC (DIC\*) and sea surface salinity-  
150 normalized alkalinity (ALK\*). The contributions of these four variables to the oceanic pCO<sub>2</sub> anomalies were calculated using  
151 the Python toolbox PyCO2SYS (Humphreys et al., 2022) version 1.8.3. For the input variables, DIC and alkalinity were  
152 normalized to a salinity of 35 to eliminate the influence from evaporation and dilution (Keeling et al., 2004). The partial  
153 derivatives were evaluated at each time step by perturbing each variable independently while holding the others constant. For  
154 this calculation, the constants were set to the median of the temporal variations, and the perturbations were applied with an  
155 amplitude equal to the standard deviation. The results were obtained for each grid, and the values at OSP were as follows:  
156  $\partial pCO_2 / \partial T = 15.27 \mu atm / ^\circ C$ ,  $\partial pCO_2 / \partial S = 25.26 \mu atm / psu$ ,  $\partial pCO_2 / \partial DIC^* = 2.15 \mu atm / \mu mol kg^{-1}$ ,  $\partial pCO_2 /$   
157  $\partial ALK^* = -1.76 \mu atm / \mu mol kg^{-1}$ . In the oceanic pCO<sub>2</sub> decomposition analysis, all input variables were detrended and  
158 deseasonalized using the STL function from the statsmodels.tsa.seasonal module in Python.

160 **3.1 Observational evidence of  $\Delta p\text{CO}_2$  during marine heatwave**

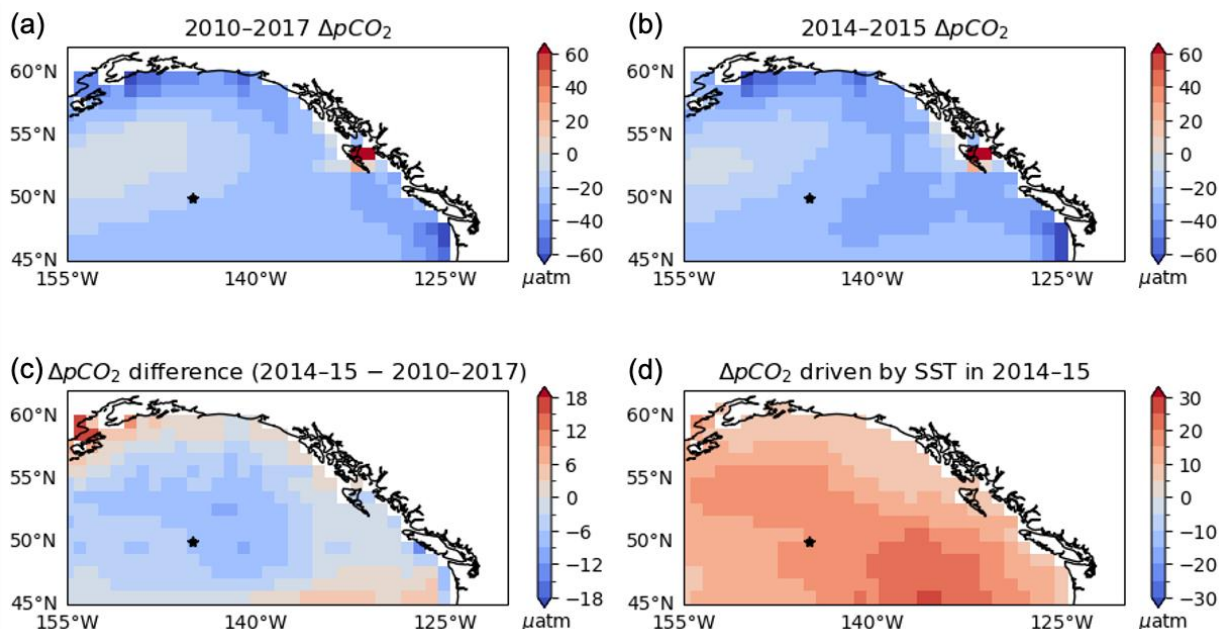
161 To assess the magnitude of the SST anomalies during the Blob and evaluate whether the model was consistent with the  
 162 observations, we compared the observed SST anomaly fields (Fig. 1). Both OISST and HadISST showed positive temperature  
 163 anomalies across the GOA. The SST anomalies in OISST were larger than those in HadISST, particularly in the southeastern  
 164 region. The modeled SST anomalies showed good agreement with both OISST and HadISST in terms of their spatial patterns  
 165 and overall amplitudes. Given this consistency, the model robustly represented the observed SST anomalies, suggesting that  
 166 model biases in SST were unlikely to influence the interpretation of the SST-driven component of oceanic  $p\text{CO}_2$ .



167  
 168 **Figure 1: Annual SST anomaly during Blob (2014–2015) relative to 2010–2017 from (a) OISST, (b) HadISST, and (c) the model.**  
 169 **The star indicates where OSP is located.**

170  
 171 Seaflex  $p\text{CO}_2$  data revealed the basin-scale surface oceanic  $p\text{CO}_2$  climatologies and anomalies during the Blob. Annual mean  
 172 climatology (2010–2017) showed negative  $\Delta p\text{CO}_2$  (oceanic  $p\text{CO}_2$  is smaller than atmospheric  $p\text{CO}_2$ ), indicating that the entire  
 173 domain of the model acted as a sink for atmospheric  $\text{CO}_2$  in the climatological mean field (Fig. 2a). Positive SST anomalies  
 174 were expected to reduce  $\text{CO}_2$  saturation in the ocean and consequently increase oceanic  $p\text{CO}_2$ . However, observation-based  
 175 data showed a decrease in oceanic  $p\text{CO}_2$ , accompanied by a larger amplitude of  $\Delta p\text{CO}_2$  than non-heatwave conditions during  
 176 the Blob (Fig. 2c). This indicated that the surface ocean absorbed more  $\text{CO}_2$  from the atmosphere during the Blob (2014–2015),  
 177 especially in the central GOA (blue areas in Fig. 2c). This trend contrasted with the rising SST, which increases oceanic  $p\text{CO}_2$   
 178 and reduces ocean carbon uptake (Fig. 2d). An estimated contribution from the positive SST anomalies on  $\Delta p\text{CO}_2$  should result

179 in an increase in  $\Delta p\text{CO}_2$  across the entire region (Fig. 2d). Therefore, elevated SST cannot account for the observed decrease  
180 in  $\Delta p\text{CO}_2$  during the Blob.

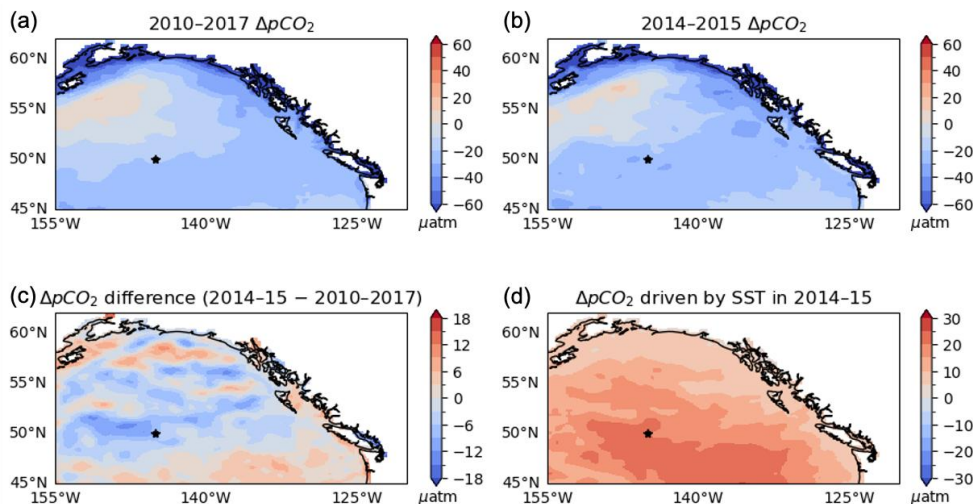


181

182 **Figure 2:  $\Delta p\text{CO}_2$  (oceanic  $p\text{CO}_2$  minus atmospheric  $p\text{CO}_2$ ):** (a) mean over 2010–2017, (b) mean during the Blob period from 2014–  
183 2015, (c) difference between (b) and (a), and (d) the component driven by SST changes during the Blob in SeaFlux. The SST-driven  
184 oceanic  $p\text{CO}_2$  in (d) is calculated as the difference between  $p\text{CO}_2$  computed under climatological mean conditions (2010–2017 mean  
185 SST and SSS from SODA version 3.4.2, and DIC and alkalinity from GLODAPv2) and  $p\text{CO}_2$  computed using SST from 2014–2015,  
186 while keeping the other variables fixed at their climatological values, with PyCO2SYS. Blue indicates  $\text{CO}_2$  uptake by the ocean, and  
187 red indicates  $\text{CO}_2$  outgassing to the atmosphere. The star in each panel marks the location of OSP.

### 188 3.2 Model validation

189 Before using the model to address the underlying mechanism behind the oceanic  $p\text{CO}_2$  decreases during the Blob, the ability  
190 of the model to reproduce existing observations must be evaluated. We first examined the model output for  $\Delta p\text{CO}_2$ . Consistent  
191 with the observations, the model indicated that increasing SST cannot account for the changes in oceanic  $p\text{CO}_2$  under the Blob,  
192 not only at OSP but across the entire GOA (Fig. 3). Compared to the observations (Fig. 2),  $\Delta p\text{CO}_2$  driven by SST under the  
193 Blob in the model was approximately 5  $\mu\text{atm}$  higher around OSP. Despite the moderate overestimation in thermal-driven  $p\text{CO}_2$   
194 anomaly, the modeled  $\Delta p\text{CO}_2$  showed a significant decrease across the central GOA, consistent with the SeaFlux dataset and  
195 the mooring observation at OSP. This consistency was underscoring the robustness of the modeled response. Throughout the  
196 region, the effect of warming-induced increase in the oceanic  $p\text{CO}_2$  was offset by the oceanic  $p\text{CO}_2$  reduction driven by  
197 decreased DIC, which was consistent with the previous work by Kohlman et al. (2024).

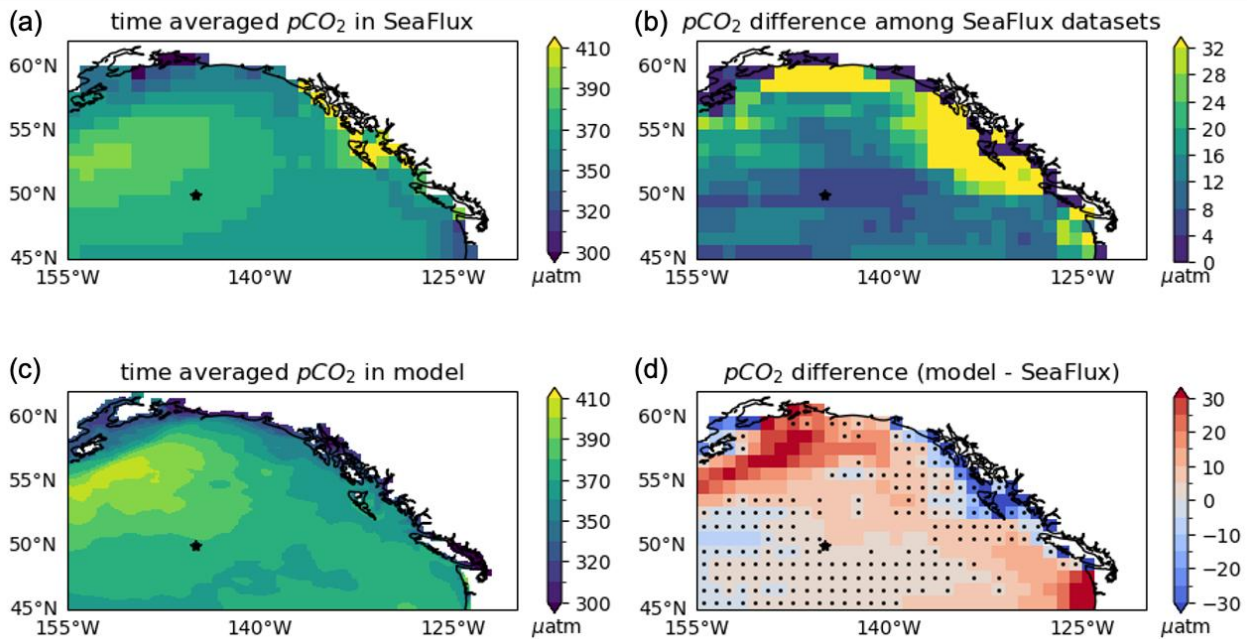


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199 **Figure 3: Same as Fig. 2 but for the model results.**

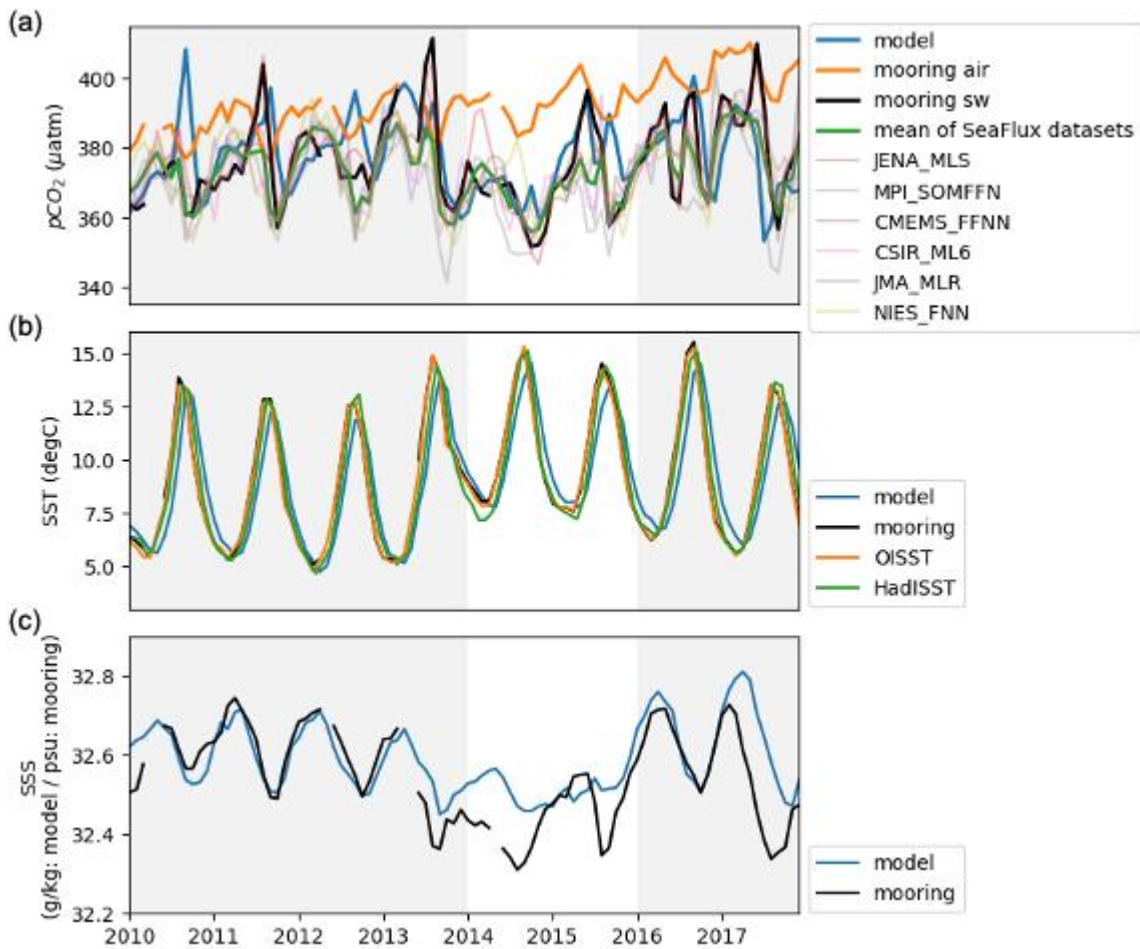
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201 Compared to the ensemble mean of the time-averaged oceanic pCO<sub>2</sub> from 2010 to 2017 in the SeaFlux dataset, the spatial  
 202 correlation between SeaFlux and the model was moderately positive ( $r=0.30$ ), and the model generally reproduced the broad  
 203 climatological spatial distribution of the observed SeaFlux oceanic pCO<sub>2</sub> (Fig. 4a, c). The model overestimated oceanic pCO<sub>2</sub>  
 204 in the northwestern area, which likely arised from model circulation biases in this region. Compared to satellite observations,  
 205 the model had slightly weaker horizontal geostrophic flow, and the core of the subpolar circulation was shifted northward (Ito  
 206 et al. 2026). The modeled vertical stratification was slightly weakened relative to the climatological observations, and this  
 207 weak stratification bias led to a higher oceanic pCO<sub>2</sub> in the model. To evaluate the uncertainties in the SeaFlux dataset,  
 208 variability among the different observational ensembles must be considered. In the open ocean, modeled oceanic pCO<sub>2</sub> was  
 209 within the range of SeaFlux ensemble variability (Fig. 4b, d). Therefore, in most areas, the model was within the uncertainty  
 210 bounds of the observation-based oceanic pCO<sub>2</sub>.



211  
 212 **Figure 4: Comparison of oceanic  $p\text{CO}_2$  distributions, averaged over 2010–2017, between the model and SeaFlux. The star in each**  
 213 **panel indicates the location of OSP. Dots in (d) indicate the grid points where the modeled climatology falls within the range of**  
 214 **differences among the SeaFlux ensemble datasets.**

215  
 216 The model also reproduced the temporal variation in oceanic  $p\text{CO}_2$  to a reasonable extent. Compared with the mooring time  
 217 series at OSP, the model captured the observed variability and magnitude in both biogeochemical and physical variables (Fig.  
 218 5). For oceanic  $p\text{CO}_2$ , the model reproduced a statistically significant fraction of the mooring  $p\text{CO}_2$  variability ( $r=0.44$ ,  
 219 normalized-RMSE=1.01), and fell within the spread of the SeaFlux ensemble data. During the Blob, both the mooring  
 220 observations and model showed pronounced decline in oceanic  $p\text{CO}_2$ . Correspondingly,  $\Delta p\text{CO}_2$ , which represented the  
 221 difference between the black or blue line and the orange line in Fig. 4a also decreased. The model slightly overestimated SSS  
 222 ( $\sim 0.1$  psu) compared to the mooring during the Blob, but this did not significantly compromise the stratification in the model,  
 223 because the density was primarily governed by sea water temperature in this region.



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**Figure 5: Comparison of time series between the model and mooring observations at OSP for (a) oceanic pCO<sub>2</sub>, (b) SST, and (c) SSS. In (a), atmospheric pCO<sub>2</sub> from the mooring observations is additionally shown in orange (mooring air) to illustrate variation in  $\Delta p\text{CO}_2$ . Oceanic pCO<sub>2</sub> from the SeaFlux ensemble mean is also shown in green, with individual ensembles indicated by thin lines with different colors. In (b) and (c), the model is blue and mooring (seawater, sw) is black. In (b), OISST is shown in orange and HadISST in green. The unshaded period corresponds to 2014–2015, during which the Blob occurred.**

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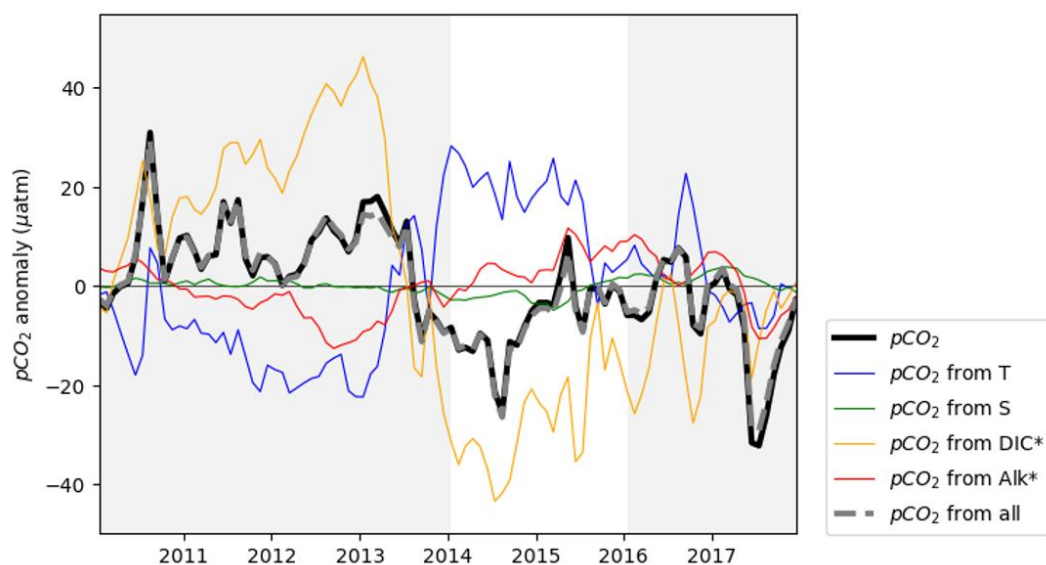
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Direct observational constraints for evaluating the modeled carbonate system were limited for this region. We compared the model DIC and TALK outputs to CMEMS-FFNN (Chau et al., 2024), one of the SeaFlux products. It should be noted that in CMEMS-FFNN, TALK was estimated using a locally interpolated alkalinity regression (Carter et al., 2016, 2018) based on SST, SSS, nitrate, and silicate, while DIC was reconstructed from oceanic pCO<sub>2</sub> and TALK. In the mean spatial distributions for 2010–2017 (Fig. S1), both DIC and TALK were generally higher in the model than in CMEMS-FFNN, particularly in the northwestern part of the model domain. However, these differences were relatively small in the central GOA near OSP, and this spatial feature was consistent with biases seen in other variables, such as nutrients and oceanic pCO<sub>2</sub>. At OSP, the model showed higher DIC concentrations than CMEMS-FFNN (+11.4  $\mu\text{mol kg}^{-1}$  during the Blob), particularly in autumn when CMEMS-FFNN had relatively low DIC values (Fig. S2). Nevertheless, the temporal variability agreed well between the two

239 products, although the model had a smaller seasonal amplitude. For TALK, the model generally showed higher concentrations  
 240 than CMEMS-FFNN ( $+7.55 \mu\text{mol kg}^{-1}$  during the Blob), except briefly in 2015. For both variables, the model reproduced the  
 241 decrease in concentrations during the Blob that was also seen in CMEMS-FFNN, indicating consistent temporal changes before,  
 242 during, and after the Blob between the two products. Compared with the in-situ observations (Franco et al., 2021), both the  
 243 model and CMEMS-FFNN showed statistically significant differences in TALK. For DIC, a statistically significant difference  
 244 was found in the model, whereas no significant difference was detected in CMEMS-FFNN. However, a more rigorous  
 245 assessment of model fidelity for DIC and TALK would require additional observational data.

### 246 3.3 Decomposition of $p\text{CO}_2$ variability

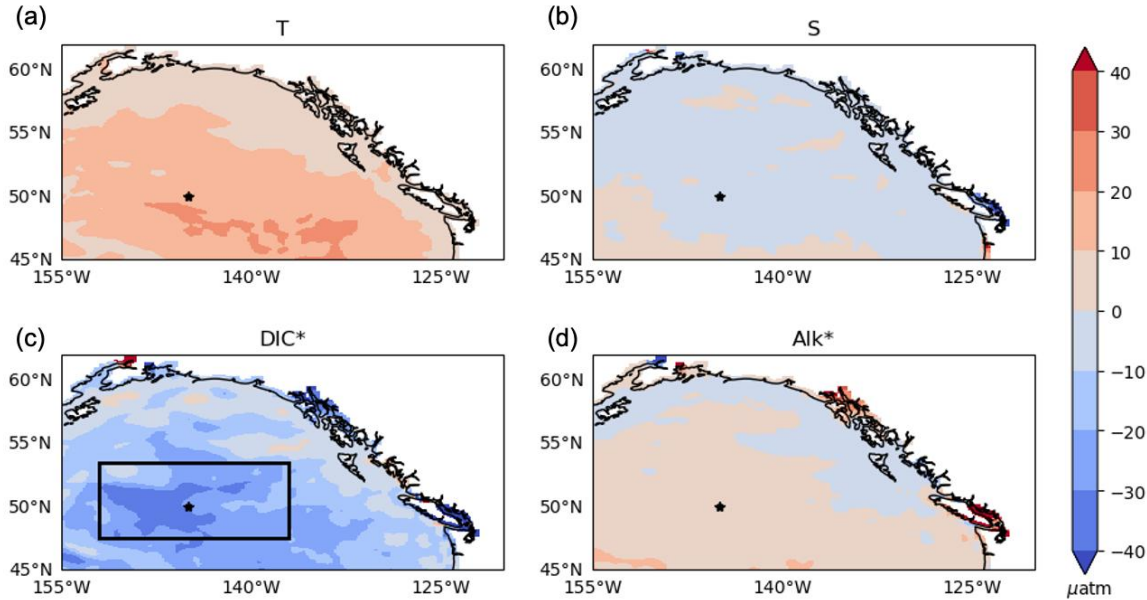
247 Oceanic  $p\text{CO}_2$  fluctuations can be explained by four components: temperature, salinity,  $\text{DIC}^*$  and  $\text{ALK}^*$  (Eq. 1). The  
 248 decomposition was applied to the variability of oceanic  $p\text{CO}_2$  at OSP, revealing that the warming-induced increased in oceanic  
 249  $p\text{CO}_2$  was fully compensated by the opposing changes in DIC during the Blob. First, the detrended and deseasonalized oceanic  
 250  $p\text{CO}_2$  time series was calculated and averaged within the 24 km radius of OSP. Similarly, the time series of SST, SSS,  $\text{DIC}^*$ ,  
 251 and  $\text{ALK}^*$  were calculated and Eq. (1) was applied to estimate the oceanic  $p\text{CO}_2$  anomalies (Fig. 6). The oceanic  $p\text{CO}_2$  changes  
 252 from SSS and  $\text{ALK}^*$  were small, while SST and  $\text{DIC}^*$  changes were the main drivers of the oceanic  $p\text{CO}_2$  changes. The  
 253 mooring observations showed that oceanic  $p\text{CO}_2$  increased due to SST changes during the Blob by about  $+20 \mu\text{atm}$ , but  
 254 decreasing oceanic  $p\text{CO}_2$  from  $\text{DIC}^*$  changes were even larger, around  $-30 \mu\text{atm}$ . Therefore, the net changes in oceanic  $p\text{CO}_2$   
 255 caused by SST and DIC are compensated, and the net oceanic  $p\text{CO}_2$  change of around  $-10 \mu\text{atm}$  was primarily driven by the  
 256 DIC, explaining the oceanic  $p\text{CO}_2$  decreases during the Blob.



257  
 258 **Figure 6:** Time series of the contributions to oceanic  $p\text{CO}_2$  anomaly referenced to 2010–2017 from SST (blue), SSS (green),  $\text{DIC}^*$   
 259 (yellow),  $\text{ALK}^*$  (red), and the combined effect of all variables (grey dashed) at OSP. The black line is the oceanic  $p\text{CO}_2$  directly  
 260 output from the model. All variables are detrended and deseasonalized. The unshaded period corresponds to 2014–2015, during

261  
262

which the Blob occurred. The grey dash line is the sum of individual components that closely matches the total pCO<sub>2</sub>, supporting the linearity of Eq (1).



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**Figure 7: Spatial patterns of the contributions to oceanic pCO<sub>2</sub> from (a) SST, (b) SSS, (c) DIC\*, and (d) Alk\* during 2014–2015. Blue shows CO<sub>2</sub> uptake, and red shows CO<sub>2</sub> outgassing. The star in each panel marks the location of OSP. The black box (47.5–53.5°N, 208–223°E) in (c) indicates the central GOA area used for the budget analysis shown in Figs. 8 and 9.**

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The decrease in DIC during the Blob happened not only at OSP but throughout the whole central GOA. Figure 7 showed the spatial distributions of the oceanic pCO<sub>2</sub> changes calculated from the model outputs of each variable, same as in Fig. 6 during the Blob. The characteristic described above, namely, the mutual compensation between the SST and DIC, also held in the entire region. These two factors counteracted each other, resulting in a relatively small decrease in oceanic pCO<sub>2</sub> due to the larger decrease in DIC. The magnitude of the oceanic pCO<sub>2</sub> declined peaks in the central GOA around the location of OSP.

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### 3.4 Simulated DIC mass balance

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To investigate the factors causing the significant decrease in DIC during the Blob, the surface ocean DIC mass balance was examined by the diagnosis of the DIC tendency terms. In the simplest form, the DIC mass balance was explained by three components: physical transport, biological activity, and air–sea gas exchange.

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$$\frac{dC}{dt} = (\text{transport}) + (\text{biological activity}) + (\text{air} - \text{sea gas transfer}) \quad (2)$$

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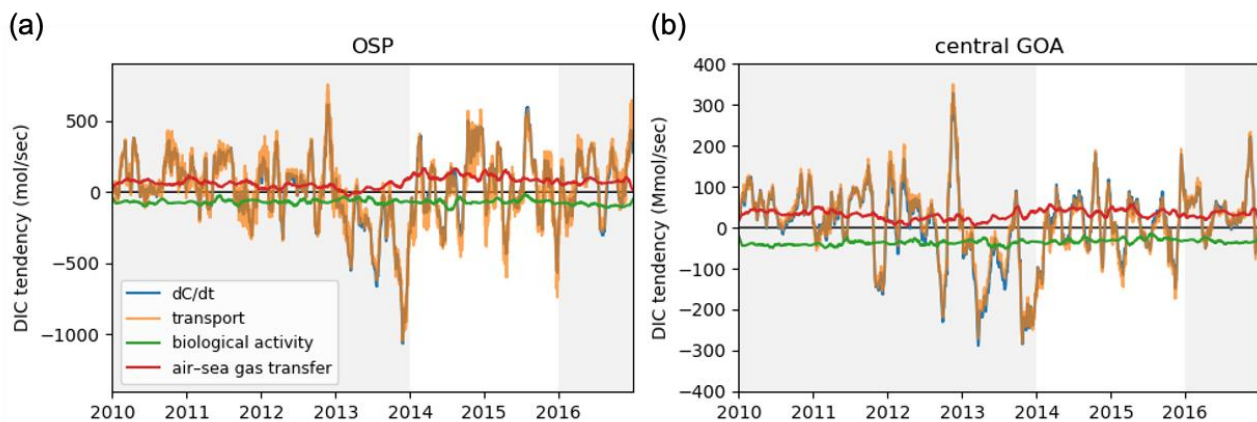
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On the right-hand side of Eq. 2, the transport term included resolved advective transport convergence and parameterized mixing terms. Zonal, meridional and vertical advection of DIC, parameterized mixing, as well as the total transport convergence were

281 calculated online and recorded as daily means. Biological activity included the net effects of photosynthetic carbon fixation,  
282 phytoplankton mortality, remineralization of dissolved and particulate organic matter, and the production and dissolution of  
283 calcium carbonates. These carbon tendency terms ( $\frac{dC}{dt}$ ) were calculated for each timesteps and were recorded as daily averages.  
284 The tendency terms calculated at OSP were integrated over the 24 km radius centred at OSP. To put OSP into context within  
285 the larger region (central GOA), rates were calculated within 47.5–53.5°N, 208–223°E, shown with a black box in Fig. 7c. In  
286 both cases, the tendencies were integrated from the surface to 177.5 m, with units of molC s<sup>-1</sup>. This depth range was greater  
287 than the maximum mixed layer depth diagnosed in our simulation and thus guarantees that vertical integration contained the  
288 entire mixed layer regardless of seasonal variability.

289  
290 Figure 8 showed the time series of each carbon tendency component at OSP (within 24 km radius) and in the central GOA  
291 (box in Fig 7c), after removing the linear trend and mean seasonal cycle and applying a 30-day moving window average. First,  
292 the sum of these three tendency components (Eq. 2) exactly matched the DIC tendency (i.e. left-hand side of Eq. 2). The  
293 variability of DIC was almost completely explained by the transport term throughout the entire period, while the effects of the  
294 other two components were relatively minor. At OSP, an extremely negative anomaly in DIC transport convergence rapidly  
295 developed in the winter of 2013, coinciding with the onset of the Blob. This anomaly was unprecedented compared to other  
296 periods. In the central GOA, a negative DIC transport anomaly appeared slightly earlier, in early 2013, and, as at OSP,  
297 intensified again in the winter of 2013. These anomalies led to a pronounced DIC decrease at the onset of the Blob, driven by  
298 changes in physical transport processes.



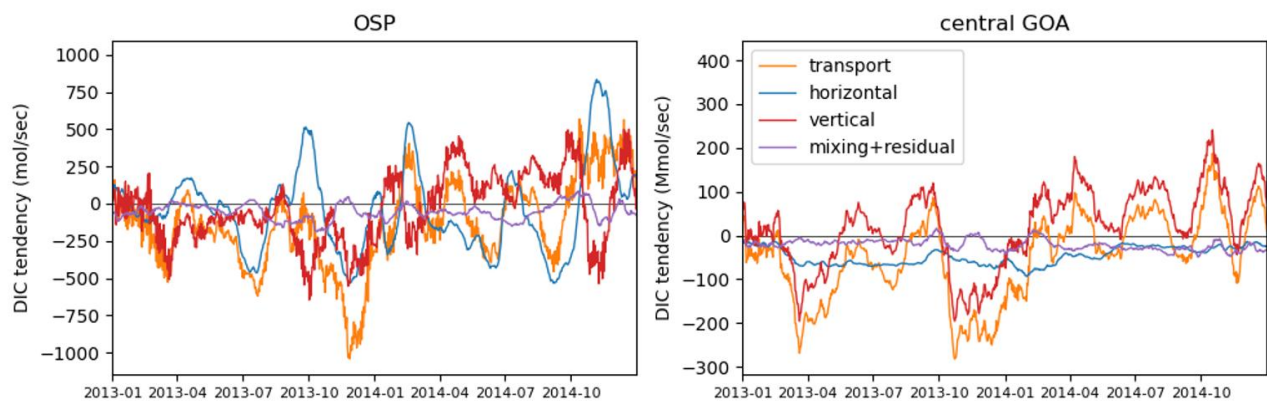
299  
300 **Figure 8: Time series of vertically integrated DIC tendencies from the surface to 177.5 m at (a) OSP and (b) the central GOA which**  
301 **is defined in Fig. 7c. The time derivative of DIC is shown in blue; changes in DIC due to transport are shown in orange, due to**  
302 **biological activity in green, and due to air–sea gas transfer in red. All variables are detrended, deseasonalized, and smoothed using**  
303 **a 30-day moving average. The unshaded period corresponds to 2014–2015, during which the Blob occurred.**

304

305 Among the physical processes responsible for the DIC reduction during the Blob, the vertical transport dominated over the  
306 large-scale domain, whereas locally the effects of the horizontal transport were comparable to those of the vertical transport.  
307 To better understand the transport-driven DIC changes, the transport term can be decomposed into individual components  
308 (horizontal and vertical advection) and the parameterized turbulent mixing terms. The background physical transport of DIC  
309 in the GOA was governed by both vertical and horizontal advection processes. Upwelling supplied DIC-enriched subsurface  
310 waters to the surface, while the prevailing westerlies drove a southward Ekman transport of DIC-rich surface waters from the  
311 northern part of the domain, where surface DIC concentrations were higher, to the southern part of the domain (Fig. S1a).

312  
313 Figure 9 showed the time series of each advective component of DIC tendency at OSP and in the central GOA. In the central  
314 GOA, the transport-driven DIC changes were primarily controlled by the vertical transport. This was because the small-scale  
315 horizontal transport within the computational domain was averaged out, and the transport across the domain boundaries can  
316 only play secondary roles in the regional DIC budget. Focusing on the DIC decrease in the winter of 2013, a pronounced DIC  
317 reduction associated with decreased vertical transport was evident, indicating suppressed upward transport of DIC-rich waters  
318 from the ocean interior to the surface layer due to enhanced stratification caused by elevated water temperatures.

319  
320 At OSP, the effect of the horizontal transport was more important locally relative to the larger domain in the central GOA.  
321 Changes in DIC reflected the combined contributions of the vertical and horizontal transport, and their relative contributions  
322 were quantified. In the winter of 2013, the net contribution of the horizontal transport accounted for approximately half of the  
323 total DIC decrease ( $-531.0 \text{ molC s}^{-1}$ ), while the remaining half was attributable to a reduction in the vertical transport ( $-533.7$   
324  $\text{molC s}^{-1}$ ). This DIC decrease associated with horizontal transport resulted from strengthened south-easterly currents in the  
325 winter of 2013. As shown in Fig. S3, anomalous south-easterly currents intensified from autumn to winter 2013, reducing the  
326 southward transport of DIC-rich surface waters that was typically supplied by Ekman transport from the north. However, as  
327 noted above, such south-easterly flow anomalies were not clearly evident over the entire model domain. Figure S3 also shows  
328 a clear shift toward negative DIC anomalies from 2013 to 2014. Consequently, the local DIC reduction at OSP during the Blob  
329 was driven by two mechanisms: anomalous south-to-north transport of low-DIC waters and suppressed upward transport of  
330 DIC-rich subsurface waters. These transport anomalies were consistent with the reduced Ekman transport associated with the  
331 weakened Aleutian Low that generated the anomalously high sea level pressure and SST (Bond et al., 2015; Hartmann et al.,  
332 2015).



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**Figure 9: Time series of vertically integrated DIC tendencies from the surface to 177.5 m at (a) OSP and (b) the central GOA, which is defined in Fig. 7c, for each transport component: horizontal advection (blue), vertical advection (red), and the mixing and residual term (purple), calculated as the remainder after subtracting the advective components from the total transport (orange). All variables are detrended, deseasonalized, and smoothed using a 30-day moving average. The unshaded period corresponds to 2014–2015, during which the Blob occurred.**

339

### **3.5 Drivers of the simulated weakening of physical transport**

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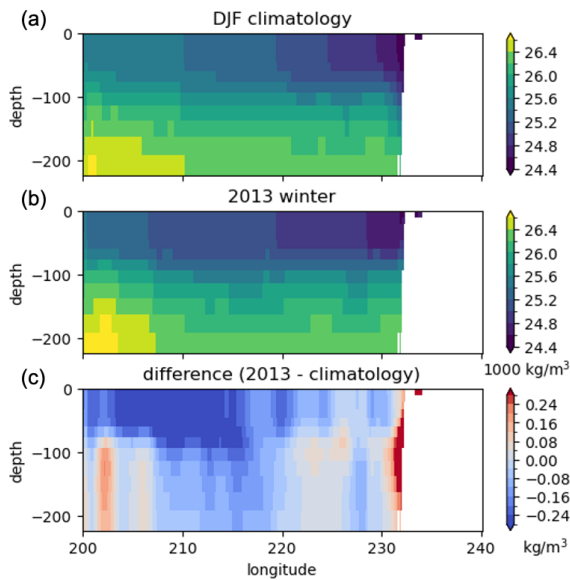
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The reduction in vertical DIC transport in the central GOA was driven by two key mechanisms: enhanced stratification associated with SST warming and weakened Ekman pumping caused by reduced wind-driven circulation. Compared with the DJF climatology, potential density in the winter 2013 exhibited substantially lower between 160°W and 140°W (Fig. 10), indicating strengthened stratification that suppressed the upward transport of DIC-rich subsurface waters and reduced the DIC supply to the surface.

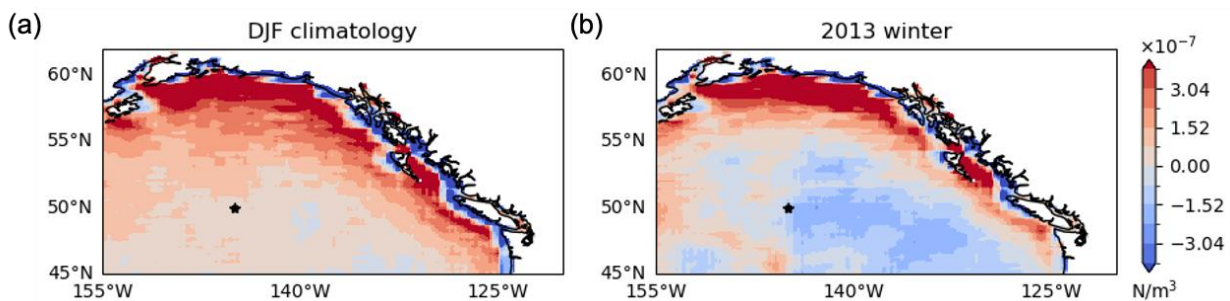


345

346 **Figure 10: Vertical cross-sections of potential density along 50.5°N for (a) the DJF-mean climatology over 2010–2017, (b) 2013, and**  
 347 **(c) their difference (2013 minus climatology).**

348

349 In addition, although the GOA was typically characterized by Ekman upwelling, the period immediately preceding the onset  
 350 of the Blob was marked by weakened wind stress curl and Ekman downwelling (Fig. 11), further suppressing the vertical DIC  
 351 supply. These potential density and wind stress curl changes were fully consistent with both the marine heatwave itself and  
 352 the weakening of the Aleutian Low that initiated the Blob. Together, these early-stage changes reduced surface DIC and exerted  
 353 a substantial influence on oceanic pCO<sub>2</sub> throughout the Blob period. This highlights the critical role of early physical forcing  
 354 in shaping the prolonged carbon cycle response during the Blob.



355

356 **Figure 11: Spatial patterns of wind stress curl ( $\text{curl}\tau_z$ ) for (a) the DJF-mean climatology over 2010–2017 and (b) 2013. Positive values**  
 357 **indicate cyclonic flow associated with Ekman upwelling. The star in each panel marks the location of OSP.**

#### 358 4 Discussion

359 Both the model and the SeaFlux product showed enhanced oceanic CO<sub>2</sub> uptake during the Blob. This increase was primarily  
360 driven by changes in surface ocean pCO<sub>2</sub>. Here, CO<sub>2</sub> flux was treated as a complementary diagnostic of the integrated air–sea  
361 CO<sub>2</sub> response to the Blob, given its nonlinear dependence on wind speed, temperature, and ΔpCO<sub>2</sub>. The spatial distribution of  
362 the CO<sub>2</sub> flux indicated net oceanic CO<sub>2</sub> uptake over most of the GOA domain, while enhanced outgassing (ocean to  
363 atmosphere), which was not evident in SeaFlux ensemble mean, was simulated in the northwestern region (Fig. S4), consistent  
364 with the elevated oceanic pCO<sub>2</sub> in this area (Figs. 2 and 3). This positive bias in the northwestern GOA was also reflected in  
365 other variables.

366  
367 In the area-integrated CO<sub>2</sub> flux time series for the central GOA (Fig. S5), the model exhibited weaker uptake than the SeaFlux  
368 ensemble mean during all periods, with values of −8.2 Tg yr<sup>−1</sup> versus −10.9 Tg yr<sup>−1</sup> before the Blob (2010–2013), −13.4 Tg  
369 yr<sup>−1</sup> versus −19.0 Tg yr<sup>−1</sup> during the Blob (2014–2015), and −12.9 Tg yr<sup>−1</sup> versus −14.8 Tg yr<sup>−1</sup> after the Blob (2017),  
370 respectively (the model versus the SeaFlux ensemble mean). Given that both products use JRA55 winds and that differences  
371 in CO<sub>2</sub> solubility are negligible, this discrepancy likely arised from differences in ΔpCO<sub>2</sub> and/or the gas transfer coefficient.  
372 Both the model and the SeaFlux ensemble showed overall good agreement with the mooring observations (Fig. 5), although  
373 the degree of agreement varies by year. Therefore, the weaker modeled CO<sub>2</sub> uptake did not necessarily imply reduced model  
374 fidelity.

375  
376 Our results showed that the reduction in DIC during the Blob was primarily driven by physical processes such as Ekman  
377 transport and vertical entrainment. Beyond the oceanic pCO<sub>2</sub> changes, numerous biological impacts of MHWs have also been  
378 reported. These include observed declines in annual net community production (Yang et al., 2018), reductions and taxonomic  
379 shifts in phytoplankton biomass, transitions in plankton communities from larger to smaller taxa (Du and Peterson, 2018; Peña  
380 et al., 2018; Batten et al., 2022). Whether ecosystem impacts can return to pre-MHW conditions remains uncertain (Suryan et  
381 al., 2021). The magnitude and nature of these biological responses vary greatly among regions, and a key factor is whether the  
382 area is primarily limited by nitrate or iron to the biological production (Peña et al., 2018; Hayashida et al., 2020; Wyatt et al.,  
383 2022). Because the GOA is an iron-limited region, it is likely less sensitive to MHW-induced macro-nutrient reductions than  
384 regions farther south in the subtropics and the transition zone (30–45°N, Wyatt et al., 2022). Nevertheless, model-based  
385 analyses will be essential for assessing long-term and basin-scale impacts. The biogeochemical model used in this study is  
386 relatively simple, and it does not explicitly represent shifts in plankton community composition. Revisiting this problem with  
387 a more sophisticated ecosystem model would be warranted to assess such ecological changes and determine their duration and  
388 magnitude.

389

390 Compared to the study conducted over broader spatial domains (Mignot et al., 2022), our results revealed a distinct response  
391 of oceanic pCO<sub>2</sub> to MHWs in the GOA. This highlights the importance of investigating the impacts of MHWs on oceanic  
392 pCO<sub>2</sub> in other regions as well. For instance, persistent positive SST anomalies have also been reported in the western North  
393 Pacific, particularly in the Oyashio region (40–43°N and 143–147°E, Miyama et al., 2021), which is one of the major carbon  
394 uptake areas in the basin. However, it remains unclear how oceanic pCO<sub>2</sub> in this region responded to such anomalies. It is  
395 important to expand the model domain to a broader area including the western North Pacific.

396

397 Abnormally high ocean temperatures like the Blob have been increasing in both frequency and duration (Frölicher et al., 2018;  
398 Oliver et al., 2018). The North Pacific experienced anomalously high SSTs in other years such as 2019 (Amaya et al., 2020)  
399 and 2023 (Dong et al., 2025). These MHWs exhibit even larger SST anomalies than the Blob and differ in several key  
400 characteristics, such as the timing of the peak warming (Amaya et al., 2020). Consequently, their impacts on oceanic pCO<sub>2</sub>,  
401 carbon cycling, and marine ecosystems may differ from those during the Blob. Indeed, the 2023 MHW showed a markedly  
402 different behaviour on a global scale, with an approximately 10% reduction in oceanic CO<sub>2</sub> uptake, substantially larger than  
403 in previous events (Müller et al., 2025). These contrasts call for event-specific investigations of individual MHWs. In this  
404 context, a similar analysis of the 2005 warm anomaly event in the GOA showed qualitatively comparable results to the Blob,  
405 in that the DIC decrease and associated reduction in oceanic pCO<sub>2</sub> were primarily controlled by changes in vertical transport.  
406 Although not shown, these results suggest that the proposed mechanism may operate under certain warm anomaly conditions  
407 in the GOA. However, its applicability is not necessarily universal and may depend on the characteristics and intensity of  
408 individual events, particularly for recent stronger MHWs.

409

410 The model employed in this study was integrated only through 2017 due to the availability of the open boundary conditions.  
411 Future work should extend the integration period, for instance by applying alternative boundary conditions, to assess the  
412 impacts of the 2019 and 2023 MHWs on oceanic pCO<sub>2</sub>.

## 413 **5 Conclusions**

414 From the winter of 2013 to 2015, the eastern North Pacific experienced an anomalously high in SST. Contrary to expectations  
415 based on the reduced CO<sub>2</sub> solubility under warming, oceanic pCO<sub>2</sub> did not rise. Instead, it decreased across the entire region,  
416 with a particularly large decrease in the open ocean (Figs. 2 and 3). To investigate why oceanic pCO<sub>2</sub> broadly decreased during  
417 the Blob, the simulation results from the regional ocean circulation and biogeochemistry model were analyzed. The model  
418 represents the oceanic pCO<sub>2</sub> remarkably well compared to the observations (Figs. 4 and 5). The decomposition of the oceanic  
419 pCO<sub>2</sub> anomalies into four components shows that the variability in the oceanic pCO<sub>2</sub> is primarily dominated by changes in  
420 SST and DIC. Furthermore, the effects of these two factors generally compensate for one another. During the Blob, the  
421 reduction in oceanic pCO<sub>2</sub> due to a decrease in DIC was stronger in magnitude than the warming-induced increase (Figs. 6 and

422 7). The pronounced reduction in DIC under the Blob is attributable not to biological processes, but rather to the anomalous  
423 physical transport (Fig. 8). In particular, reduced vertical DIC transport in the central GOA, driven by enhanced stratification  
424 and weakened Ekman pumping, was the primary mechanism (Figs. 9–11). Consequently, these changes in the vertical  
425 circulation decreased the surface DIC concentrations, leading to a subsequent decline in the oceanic pCO<sub>2</sub> during the Blob.  
426

#### 427 **Data availability**

428 The monthly and daily model output data used in this study are archived on Zenodo (doi: 10.5281/zenodo.18462325). The  
429 data will be made publicly available upon publication of the associated article.

#### 430 **Author contribution**

431 TI conceptualized and designed the model, and YA performed the simulations and analyses. AT contributed observational  
432 comparison and interpretation. YA wrote the original draft of the manuscript. TI, AT, CR, and JM reviewed and edited the  
433 manuscript and contributed to scientific discussion. All authors approved the final version of the manuscript.

#### 434 **Competing interests**

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

#### 436 **Acknowledgements**

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438 the public domain (Campin et al., 2025)

#### 439 **References**

- 440 Amaya, D. J., Miller, A. J., Xie, S.-P., and Kosaka, Y.: Physical drivers of the summer 2019 North Pacific marine heatwave,  
441 *Nat. Commun.*, 11, 1903, doi:10.1038/s41467-41020-15820-w, 2020.
- 442 Barbeaux, S. J., Holsman, K., and Zador, S.: Marine heatwave stress test of ecosystem-based fisheries management in the Gulf  
443 of Alaska Pacific cod fishery, *Front. Mar. Sci.*, 7, 703, doi:10.3389/fmars.2020.00703, 2020.
- 444 Batten, S. D., Ostle, C., Hélaouët, P., and Walne, A. W.: Responses of Gulf of Alaska Plankton Communities to a Marine Heat  
445 Wave, *Deep Sea Research Part II: Topical Studies in Oceanography*, 195, 105002, doi:10.1016/j.dsr2.2021.105002, 2022.

- 446 Bond, N. A., Cronin, M. F., Freeland, H., and Mantua, N.: Causes and impacts of the 2014 warm anomaly in the NE Pacific,  
447 *Geophys. Res. Lett.*, 42, 3414–3420, doi:10.1002/2015gl063306, 2015.
- 448 Burger, F. A., Terhaar, J., and Frölicher, T. L.: Compound marine heatwaves and ocean acidity extremes, *Nat. Commun.*, 13,  
449 4722, doi:10.1038/s41467-022-32120-7, 2022.
- 450 Campin, J.-M., Heimbach, P., Losch, M., Forget, G., edhill3, Adcroft, A., amolod, Menemenlis, D., dfer22, Jahn, O., Hill, C.,  
451 Scott, J., dngoldberg, stephdut, Mazloff, M., Fox-Kemper, B., antnguyen13, Doddridge, E., Fenty, I., Bates, M., Wang, O.,  
452 Smith, T., AndrewEichmann, N., mitllheisey, Lauderdale, J., Martin, T., Abernathey, R., samarkhatiwala, hongandyan, and  
453 Escobar, I.: MITgcm/MITgcm: checkpoint69e (Version checkpoint69e), Zenodo, doi:10.5281/zenodo.15320163, 2025.
- 454 Carter, B. R., Williams, N. L., Gray, A. R., and Feely, R. A.: Locally interpolated alkalinity regression for global alkalinity  
455 estimation, *Limnol. Oceanogr.-Meth.*, 14, 268–277, doi:1002/lom3.10087, 2016.
- 456 Carter, B. R., Feely, R. A., Williams, N. L., Dickson, A. G., Fong, M. B., and Takeshita, Y.: Updated methods for global  
457 locally interpolated estimation of alkalinity, pH, and nitrate, *Limnol. Oceanogr.-Meth.*, 16, 119–131, doi:10.1002/lom3.10232,  
458 2018.
- 459 Carton, J. A., and Giese, B. S.: A Reanalysis of Ocean Climate Using Simple Ocean Data Assimilation (SODA), *Mon. Weather*  
460 *Rev.*, 136, 2999–3017, doi:10.1175/2007MWR1978.1, 2008.
- 461 Carton, J. A., Chepurin, G. A., and Chen, L.: SODA3: A new ocean climate reanalysis, *J. Clim.*, 31, 6967–6983,  
462 doi:10.1175/JCLI-D-17-0149.1, 2018.
- 463 Cavole, L., Demko, A., Diner, R., Giddings, A., Koester, I., Pagniello, C., Paulsen, M.-L., Ramírez-Valdez, A., Schwenck, S.,  
464 Zill, M., and Franks, P.: Biological Impacts of the 2013–2015 Warm-Water Anomaly in the Northeast Pacific: Winners, Losers,  
465 and the Future, *Oceanography*, 29, 273–285, doi:10.5670/oceanog.2016.32, 2016.
- 466 Chau, T.-T.-T., Gehlen, M., Metzl, N., and Chevallier, F.: CMEMS-LSCE: a global, 0.25°, monthly reconstruction of the  
467 surface ocean carbonate system, *Earth Syst. Sci. Data*, 16, 121–160, doi:10.5194/essd-16-121-2024, 2024.
- 468 Coyle, K. O., Cheng, W., Hinckley, S. L., Lessard, E. J., Whitledge, T., Hermann, A. J., and Hedstrom, K.: Model and field  
469 observations of effects of circulation on the timing and magnitude of nitrate utilization and production on the northern Gulf of  
470 Alaska shelf, *Prog. Oceanogr.*, 103, 16–41, doi:10.1016/j.pocean.2012.03.002, 2012.
- 471 Cronin, M. F., Pelland, N. A., Emerson, S. R., and Crawford, W. R.: Estimating diffusivity from the mixed layer heat and salt  
472 balances in the North Pacific, *J. Geophys. Res.-Oceans*, 120, 7346–7362, doi:10.1002/2015jc011010, 2015.
- 473 Di Lorenzo, E., and Mantua, N.: Multi-year persistence of the 2014/15 North Pacific marine heatwave. *Nature Climate Change*,  
474 6 (11), 1042–1047, doi:10.1038/nclimate3082, 2016.
- 475 Dong, T., Zeng, Z., Pan, M., Wang, D., Chen, Y., Liang, L., Yang, S., Jin, Y., Luo, S., Liang, S., Huang, X., Zhao, D., Ziegler,  
476 A. D., Chen, D., Li, L. Z. X., Zhou, T., and Zhang, D.: Record-breaking 2023 marine heatwaves, *Science*, 389, 6758, pp. 369–  
477 374, doi: 10.1126/science.adr0910, 2025.
- 478 Du, X., and Peterson, W. T.: Phytoplankton community structure in 2011–2013 compared to the extratropical warming event  
479 of 2014–2015, *Geophys. Res. Lett.*, 45, 1534–1540, doi:10.1002/2017GL076199, 2018.

480 Duke, P. J., Hamme, R. C., Ianson, D., Landschützer, P., Ahmed, M. M. M., Swart, N. C., and Covert, P. A.: Estimating marine  
481 carbon uptake in the northeast Pacific using a neural network approach, *Biogeosciences*, 20, 3919–3941, doi:10.5194/bg-20-  
482 3919-2023, 2023.

483 Dunne, J. P., Bociu, I., Bronselaer, B., Guo, H., John, J. G., Krasting, J. P., Stock, C. A., Winton, M., and Zadeh, N.: Simple  
484 Global Ocean Biogeochemistry with Light, Iron, Nutrients and Gas version 2 (BLINGv2): Model description and simulation  
485 characteristics in GFDL's CM4.0, *J. Adv. Model. Earth Sy.*, 12, e2019MS002008, doi:10.1029/2019MS002008, 2020.

486 Emerson, S., Sabine, C., Cronin, M. F., Feely, R., Cullison Gray, S. E., and DeGrandpre, M.: Quantifying the flux of CaCO<sub>3</sub>  
487 and organic carbon from the surface ocean using in situ measurements of O<sub>2</sub>, N<sub>2</sub>, pCO<sub>2</sub>, and pH, *Global Biogeochem. Cy.*,  
488 25, GB3008, doi:10.1029/2010GB003924, 2011.

489 Fay, A. R., Gregor, L., Landschützer, P., McKinley, G. A., Gruber, N., Gehlen, M., Iida, Y., Laruelle, G. G., Rödenbeck, C.,  
490 Roobaert, A., and Zeng, J.: SeaFlux: harmonization of air–sea CO<sub>2</sub> fluxes from surface pCO<sub>2</sub> data products using a  
491 standardized approach, *Earth Syst. Sci. Data*, 13, 4693–4710, doi:10.5194/essd-13-4693-2021, 2021.

492 Franco, A. C., Ianson, D., Ross, T., Hamme, R. C., Monahan, A. H., Christian, J. R., Davelaar, M., Johnson, W. K., Miller, L.  
493 A., Robert, M., and Tortell, P. D.: Anthropogenic and climatic contributions to observed carbon system trends in the northeast  
494 Pacific, *Global Biogeochem. Cy.*, 35, doi:10.1029/2020GB006829, 2021.

495 Friedlingstein, P., O'Sullivan, M., Jones, M. W., Andrew, R. M., Gregor, L., Hauck, J., Le Quéré, C., Luijkx, I. T., Olsen, A.,  
496 Peters, G. P., Peters, W., Pongratz, J., Schwingshackl, C., Sitch, S., Canadell, J. G., Ciais, P., Jackson, R. B., Alin, S. R.,  
497 Alkama, R., Arneeth, A., Arora, V. K., Bates, N. R., Becker, M., Bellouin, N., Bittig, H. C., Bopp, L., Chevallier, F., Chini, L.  
498 P., Cronin, M., Evans, W., Falk, S., Feely, R. A., Gasser, T., Gehlen, M., Gkritzalis, T., Gloege, L., Grassi, G., Gruber, N.,  
499 Gürses, Ö., Harris, I., Hefner, M., Houghton, R. A., Hurtt, G. C., Iida, Y., Ilyina, T., Jain, A. K., Jersild, A., Kadono, K., Kato,  
500 E., Kennedy, D., Klein Goldewijk, K., Knauer, J., Korsbakken, J. I., Landschützer, P., Lefèvre, N., Lindsay, K., Liu, J., Liu,  
501 Z., Marland, G., Mayot, N., McGrath, M. J., Metzl, N., Monacci, N. M., Munro, D. R., Nakaoka, S.-I., Niwa, Y., O'Brien, K.,  
502 Ono, T., Palmer, P. I., Pan, N., Pierrot, D., Pockock, K., Poulter, B., Resplandy, L., Robertson, E., Rödenbeck, C., Rodriguez,  
503 C., Rosan, T. M., Schwinger, J., Séférian, R., Shutler, J. D., Skjelvan, I., Steinhoff, T., Sun, Q., Sutton, A. J., Sweeney, C.,  
504 Takao, S., Tanhua, T., Tans, P. P., Tian, X., Tian, H., Tilbrook, B., Tsujino, H., Tubiello, F., van der Werf, G. R., Walker, A.  
505 P., Wanninkhof, R., Whitehead, C., Willstrand Wranne, A., Wright, R., Yuan, W., Yue, C., Yue, X., Zaehle, S., Zeng, J., and  
506 Zheng, B.: Global Carbon Budget 2022, *Earth Syst. Sci. Data*, 14, 4811–4900, doi: 10.5194/essd-14-4811-2022, 2022.

507 Frölicher, T. L., Fischer, E. M., and Gruber, N.: Marine heatwaves under global warming, *Nature*, 560, 360–364,  
508 doi:10.1038/s41586-018-0383-9, 2018.

509 Gruber, N., Boyd, P. W., Frölicher, T. L., and Vogt, M.: Biogeochemical extremes and compound events in the ocean, *Nature*,  
510 600, 395–407, doi:10.1038/s41586-021-03981-7, 2021.

511 Hartmann, D. L.: Pacific sea surface temperature and the winter of 2014, *Geophys. Res. Lett.*, 42, 1894–1902,  
512 doi:10.1002/2015GL063083, 2015.

513 Hauri, C., Schultz, C., Hedstrom, K., Danielson, S., Irving, B., Doney, S. C., Dussin, R., Curchitser, E. N., Hill, D. F., and  
514 Stock, C. A.: A regional hindcast model simulating ecosystem dynamics, inorganic carbon chemistry, and ocean acidification  
515 in the Gulf of Alaska, *Biogeosciences*, 17, 3837–3857, doi:10.5194/bg-17-3837-2020, 2020.

516 Hayashida, H., Matear, R. J., and Strutton, P. G.: Background nutrient concentration determines phytoplankton bloom response  
517 to marine heatwaves, *Global Change Biol.*, 26, 4800–4811, doi:10.1111/gcb.15255, 2020.

518 Hinckley, S., Coyle, K. O., Gibson, G., Hermann, A. J., and Dobbins, E. L.: A biophysical NPZ model with iron for the Gulf  
519 of Alaska: reproducing the differences between an oceanic HNLC ecosystem and a classical northern temperate shelf  
520 ecosystem, *Deep-Sea Res. Pt. II*, 56, 2520–2536, doi:10.1016/j.dsr2.2009.03.003, 2009.

521 Hobday, A. J., Alexander, L. V., Perkins, S. E., Smale, D. A., Straub, S. C., Oliver, E. C., Benthuisen, J. A., Burrows, M. T.,  
522 Donat, M. G., Feng, M., Holbrook, N. J., Moore, P. J., Scannell, H. A., Sen Gupta, A., and Wernberg, T.: A hierarchical  
523 approach to defining marine heatwaves, *Prog. Oceanogr.*, 141, 227–238, doi:10.1016/j.pocean.2015.12.014, 2016.

524 Huang, B., Liu, C., Banzon, V., Freeman, E., Graham, G., Hankins, B., Smith, T., and Zhang, H.-M.: Improvements of the  
525 Daily Optimum Interpolation Sea Surface Temperature (DOISST) Version 2.1, *J. Climate*, 34, 2923–2939, doi:10.1175/JCLI-  
526 D-20-0166.1, 2021.

527 Humphreys, M. P., Lewis, E. R., Sharp, J. D., and Pierrot, D.: PyCO2SYS v1.8: marine carbonate system calculations in  
528 Python, *Geosci. Model Dev.*, 15, 15–43, doi:10.5194/gmd-15-15-2022, 2022.

529 Ito, T., Timmerman, A. H. V., Bjorklund, A., Stanley, S. I., Abe, Y., Reinhard, C. T., and Montoya, J.: Eddy-induced iron  
530 transport sustains the biological productivity in the Gulf of Alaska. *J. Geophys. Res.-Oceans*, 131, e2025JC022996, doi:  
531 10.1029/2025JC022996, 2026.

532 Keeling, C. D., Piper, S. C., Bacastow, R. B., Wahlen, M., Whorf, T. P., Heimann, M., and Meijer, H. A.: Exchanges of  
533 Atmospheric CO<sub>2</sub> and 13CO<sub>2</sub> with the Terrestrial Biosphere and Oceans from 1978 to 2000. I. Global Aspects, *SIO*  
534 *REFERENCE*, p. 29, <https://escholarship.org/uc/item/09v319r9> (last access: 06 Oct 2025), 2001.

535 Keeling, C. D., Brix, H., and Gruber, N.: Seasonal and long-term dynamics of the upper ocean carbon cycle at Station ALOHA  
536 near Hawaii, *Global Biogeochem. Cy.*, 18, GB4006, doi:10.1029/2004GB002227, 2004.

537 Kohlman, C., Cronin, M. F., Dziak, R., Mellinger, D. K., Sutton, A., Galbraith, M., Robert, M., Thomson, J., Zhang, D., and  
538 Thompson, L.: The 2019 marine heatwave at ocean station papa: A multi-disciplinary assessment of ocean conditions and  
539 impacts on marine ecosystems, *J. Geophys. Res.-Oceans*, 129(6), e2023JC020167, doi:10.1029/2023JC020167, 2024.

540 Large, W. G., McWilliams, J. C., and Doney, S. C.: Ocean vertical mixing: a review and a model with a nonlocal boundary  
541 layer parameterization, *Rev. Geophys.*, 32, 363–403, doi:10.1029/94RG01872, 1994.

542 Lauvset, S. K., Lange, N., Tanhua, T., Bittig, H. C., Olsen, A., Kozyr, A., Alin, S., Álvarez, M., Azetsu-Scott, K., Barbero, L.,  
543 Becker, S., Brown, P. J., Carter, B. R., da Cunha, L. C., Feely, R. A., Hoppema, M., Humphreys, M. P., Ishii, M., Jeansson,  
544 E., Jiang, L.-Q., Jones, S. D., Lo Monaco, C., Murata, A., Müller, J. D., Pérez, F. F., Pfeil, B., Schirnick, C., Steinfeldt, R.,  
545 Suzuki, T., Tilbrook, B., Ulfsbo, A., Velo, A., Woosley, R. J., and Key, R. M.: GLODAPv2.2022: the latest version of the  
546 global interior ocean biogeochemical data product, *Earth Syst. Sci. Data*, 14, 5543–5572, doi:10.5194/essd-14-5543-2022,  
547 2022.

- 548 Li, C., Burger, F. A., Raible, C. C., and Frölicher, T. L.: Observed Regional Impacts of Marine Heatwaves on Sea-Air CO<sub>2</sub>  
549 Exchange, *Geophys. Res. Lett.*, 51, e2024GL110379, doi:10.1029/2024GL110379, 2024a.
- 550 Li, C., Huang, J., Liu, X., Ding, L., He, Y., and Xie, Y.: The ocean losing its breath under the heatwaves, *Nat. Commun.*, 15,  
551 6840, doi:10.1038/s41467-024-51323-8, 2024b.
- 552 Losch, M., Menemenlis, D., Campin, J.-M., Heimbach, P., and Hill, C.: On the formulation of sea-ice models. Part 1: Effects  
553 of different solver implementations and parameterizations, *Ocean Model.*, 33, 129–144, doi:10.1016/j.ocemod.2009.12.008,  
554 2010.
- 555 Manizza, M., Le Quéré, C., Watson, A. J., and Buitenhuis, E. T.: Bio-optical feedbacks among phytoplankton, upper ocean  
556 physics and sea-ice in a global model, *Geophys. Res. Lett.*, 32, L05603, doi:10.1029/2004gl020778, 2005.
- 557 Marshall, J., Adcroft, A., Hill, C., Perelman, L., and Heisey, C.: A finite-volume, incompressible Navier Stokes model for  
558 studies of the ocean on parallel computers, *J. Geophys. Res.-Oceans*, 102, 5753–5766, doi:10.1029/96JC02775, 1997a.
- 559 Marshall, J., Hill, C., Perelman, L., and Adcroft, A.: Hydrostatic, quasi-hydrostatic, and nonhydrostatic ocean modeling, *J.*  
560 *Geophys. Res.-Oceans*, 102, 5733–5752, doi:10.1029/96JC02776, 1997b.
- 561 McKinley, G. A., Takahashi, T., Buitenhuis, E., Chai, F., Christian, J.R., Doney, S. C., Jiang, M. S., Lindsay, K., Moore, J.  
562 K., Le Quéré, C., Lima, I., Murtugudde, R., Shi, L., and Wetzel, P.: North Pacific carbon cycle response to climate variability  
563 on seasonal to decadal timescales, *J. Geophys. Res.*, 111, C07S06, doi:10.1029/2005JC003173, 2006.
- 564 Mignot, A., Schuckmann, K. V., Landschützer, P., Gasparin, F., Gennip, S. V., Perruche, C., Lamouroux, J., and Amm, T.:  
565 Decrease in air-sea CO<sub>2</sub> fluxes caused by persistent marine heatwaves, *Nat. Commun.*, 13, 1–9, doi:10.1038/s41467-022-  
566 31983-0, 2022.
- 567 Miyama, T., Minobe, S., and Goto, H.: Marine Heatwave of Sea Surface Temperature of the Oyashio Region in Summer in  
568 2010–2016, *Front. Mar. Sci.*, 7:576240, doi:10.3389/fmars.2020.576240, 2021.
- 569 Mogen, S. C., Lovenduski, N. S., Dallmann, A. R., Gregor, L., Sutton, A. J., Bograd, S. J., Quiros, N. C., Di Lorenzo, E.,  
570 Hazen, E. L., Jacox, M. G., Buil, M. P., and Yeager, S.: Ocean Biogeochemical Signatures of the North Pacific Blob, *Geophys.*  
571 *Res. Lett.*, 49, e2021GL096938, doi:10.1029/2021GL096938, 2022.
- 572 Müller, J. D., Gruber, N., Schneuwly, A., Bakker, D. C. E., Gehlen, M., Gregor, L., Hauck, J., Landschützer, P., and McKinley,  
573 G. A.: Unexpected decline in the ocean carbon sink under record-high sea surface temperatures in 2023. *Nat. Clim. Chang.* 15,  
574 978–985, doi:10.1038/s41558-025-02380-4, 2025.
- 575 Oliver, E. C. J., Donat, M. G., Burrows, M. T., Moore, P. J., Smale, D. A., Alexander, L. V., Benthuisen, J. A., Feng, M., Sen  
576 Gupta, A., Hobday, A. J., Holbrook, N. J., Perkins-Kirkpatrick, S. E., Scannell, H. A., Straub, S. C., and Wernberg, T.: Longer  
577 and more frequent marine heatwaves over the past century, *Nat. Commun.*, 9, 1324, doi:10.1038/s41467-018-03732-9, 2018.
- 578 Peña, M. A., Nemcek, N., and Robert, M.: Phytoplankton responses to the 2014–2016 warming anomaly in the northeast  
579 subarctic Pacific Ocean, *Limnol. Oceanogr.*, 64, pp. 515-525, doi:10.1002/lno.11056, 2018.
- 580 Rayner, N. A., Parker, D. E., Horton, E. B., Folland, C., Alexander, L., Rowell, D., Kent, E., and Kaplan, A.: Global analyses  
581 of sea surface temperature, sea ice, and night marine air temperature since the late nineteenth century, *J. Geophys. Res.*, 108,  
582 4407, doi:10.1029/2002JD002670, 2003.

583 Reagan, J. R., Seidov, D., Wang, Z., Dukhovskoy, D., Boyer, T. P., Locarnini, R. A., Baranova, O. K., Mishonov, A. V.,  
584 Garcia, H. E., Bouchard, C., Cross, S. L., and Paver, C. R.: World Ocean Atlas 2023, Volume 2: Salinity, A. Mishonov,  
585 Technical Editor, NOAA Atlas NESDIS 90, doi:10.25923/70qt-9574, 2024.

586 Reynolds, R. W., Smith, T. M., Liu, C., Chelton, D. B., Casey, K. S., and Schlax, M. G.: Daily high-resolution-blended analyses  
587 for sea surface temperature, *J. Climate*, 20, 5473–5496, doi:10.1175/2007JCLI1824.1, 2007.

588 Siedlecki, S. A., Pilcher, D. J., Hermann, A. J., Coyle, K., and Mathis, J.: The Importance of Freshwater to Spatial Variability  
589 of Aragonite Saturation State in the Gulf of Alaska, *J. Geophys. Res.-Oceans*, 122, 8482–8502, doi:10.1002/2017JC012791,  
590 2017.

591 Smale, D. A., Wernberg, T., Oliver, E. C. J., Thomsen, M., Harvey, B. P., Straub, S. C., Burrows, M. T., Alexander, L. V.,  
592 Benthuisen, J. A., Donat, M. G., Feng, M., Hobday, A. J., Holbrook, N. J., Perkins-Kirkpatrick, S. E., Scannell, H. A., Sen  
593 Gupta, A., Payne, B. L., and Moore, P. J.: Marine heatwaves threaten global biodiversity and the provision of ecosystem  
594 services, *Nat. Clim. Change*, 9, 306–312, doi:10.1038/s41558-019-0412-1, 2019.

595 Stock, C. A., Dunne, J. P., and John, J. G.: Global-scale carbon and energy flows through the marine planktonic food web: An  
596 analysis with a coupled physical–biological model, *Prog. Oceanogr.*, 120, 1–28, doi: 10.1016/j.poccean.2013.07.001, 2014.

597 Suryan, R. M., Arimitsu, M. L., Coletti, H. A., Hopcroft, R. R., Lindeberg, M. R., Barbeaux, S. J., Batten, S. D., Burt, W. J.,  
598 Bishop, M. A., Bodkin, J. L., Brenner, R., Campbell, R. W., Cushing, D. A., Danielson, S. L., Dorn, M. W., Drummond, B.,  
599 Esler, D., Gelatt, T., Hanselman, D. H., Hatch, S. A., Haught, S., Holderied, K., Iken, K., Irons, D. B., Kettle, A. B., Kimmel,  
600 D. G., Konar, B., Kuletz, K. J., Laurel, B. J., Maniscalco, J. M., Matkin, C., McKinstry, C. A. E., Monson, D. H., Moran, J.  
601 R., Olsen, D., Palsson, W. A., Pegau, W. S., Piatt, J. F., Rogers, L. A., Rojek, N. A., Schaefer, A., Spies, I. B., Straley, J. M.,  
602 Strom, S. L., Sweeney, K. L., Szymkowiak, M., Weitzman, B. P., Yasumiishi, E. M., and Zador, S. G.: Ecosystem response  
603 persists after a prolonged marine heatwave, *Sci. Rep.*, 11, 6235, doi:10.1038/s41598-021-83818-5, 2021.

604 Takahashi, T., Sutherland, S. C., Wanninkhof, R., Sweeney, C., Feely, R. A., Chipman, D. W., Hales, B., Friederich, G.,  
605 Chavez, F., Sabine, C., Watson, A., Bakker, D. C., Schuster, U., Metzl, N., Yoshikawa-Inoue, H., Ishii, M., Midorikawa, T.,  
606 Nojiri, Y., Körtzinger, A., Steinhoff, T., Hoppema, M., Olafsson, J., Arnarson, T. S., Tilbrook, B., Johannessen, T., Olsen, A.,  
607 Bellerby, R., Wong, C., Delille, B., Bates, N., and de Baar, H. J.: Climatological mean and decadal change in surface ocean  
608 pCO<sub>2</sub>, and net sea–air CO<sub>2</sub> flux over the global oceans, *Deep-Sea Res. Pt. 2*, 56, 554–577, doi:10.1016/j.dsr2.2008.12.009,  
609 2009.

610 Tsujino, H., Urakawa, S., Nakano, H., Small, R. J., Kim, W. M., Yeager, S. G., Danabasoglu, G., Suzuki, T., Bamber, J. L.,  
611 Bentsen, M., Böning, C. W., Bozec, A., Chassignet, E. P., Curchitser, E., Dias, F. B., Durack, P. J., Griffies, S. M., Harada,  
612 Y., Ilicak, M., Josey, S. A., Kobayashi, C., Kobayashi, S., Komuro, Y., Large, W. G., Sommer, J. L., Marsland, S. J., Masina,  
613 S., Scheinert, M., Tomita, H., Valdivieso, M., and Yamazaki, D.: JRA-55 based surface dataset for driving ocean-sea-ice  
614 models (JRA55-do), *Ocean Model.*, 130, 79–139, doi:10.1016/j.ocemod.2018.07.002, 2018.

615 Yang, B., Emerson, S. R., and Peña, M. A.: The effect of the 2013–2016 high temperature anomaly in the subarctic Northeast  
616 Pacific (the “Blob”) on net community production, *Biogeosciences*, 15, 6747–6759, doi: 10.5194/bg-15-6747-2018, 2018.

617 Wyatt, A. M., Resplandy, L., and Marchetti, A.: Ecosystem impacts of marine heat waves in the northeast Pacific,  
618 *Biogeosciences*, 19, 5689–5705, doi:10.5194/bg-19-5689-2022, 2022.

619