

Supporting Information

Biogenic Carbon Pool Production Maintains the Southern Ocean Carbon Sink

Yibin Huang^{1,2}, Andrea J. Fassbender^{2,1*}, Seth M. Bushinsky³

¹Department of Ocean Sciences, University of California, Santa Cruz, CA, USA, 95064

²NOAA/OAR Pacific Marine Environmental Laboratory, Seattle, WA, USA, 98115

³Department of Oceanography, University of Hawaii at Mānoa, Honolulu, HA, USA, 96822

*Correspondence to: A. J. Fassbender, andrea.j.fassbender@noaa.gov

This PDF file includes:

Supplementary text 1 to 1.8;

Figure S1 to S24;

Table S1 to S3;

SI References

44 Supplementary Text

45 1.1 Constraining the C:N Ratio of Dissolved Organic Matter (C:N_{DOM}) Production

46

47 C:N_{DOM} is constrained using a recently published compilation of global shipboard
48 observations (Hansell et al., 2021; **Fig. S18a**) that includes concurrent measurements of dissolved
49 organic carbon (DOC), total dissolved nitrogen (TDN), and inorganic nitrogen (IN: including
50 nitrate and nitrite). Dissolved organic nitrogen (DON) is computed by subtracting IN from TDN.
51 In seawater, a considerable fraction of TDN is attributed to IN, particularly in high-latitude regions.
52 Thus, the DON values derived by differencing two large values may be subject to a significant
53 error. Unfortunately, accuracies for TDN and IN are not reported in the compiled dataset. We use
54 an analytical error of 4% for TND and 2% for IN measurements to approximate the propagated
55 error associated with our DON estimate (σ_{DON} , **Eq. S7**):

56

$$57 \sigma_{\text{DON}} = \sqrt{(\text{TND} \times 0.04)^2 + (\text{IN} \times 0.02)^2} \quad \text{Eq. S7}$$

58

59 We discard all DON values below σ_{DON} .

60

61 Since a large fraction of DOM is refractory (1), the C:N ratio of the bulk DOM pool does not
62 reflect the labile DOM production ratio. Thus, we estimate the C:N_{DOM} for labile DOM production
63 using the seasonal DOC and DON inventory changes. Since the DOM seasonal cycle is rarely
64 resolved (i.e., there are few wintertime observations), we use two approaches to infer the
65 wintertime DOM value (DOM_{winter}) using summertime DOM (DOM_{summer}) observations. The
66 first method is to set DOM_{winter} to the mean of DOM_{summer} values above the annual maximum
67 MLD (MLD_{max}) following Baetge, *et al.* (2) (**Eq. S8**):

68

$$69 \text{DOM}_{\text{winter}} = \frac{\int_0^{\text{MLD}_{\text{max}}} \text{DOM}_{\text{summer}} dz}{\text{MLD}_{\text{max}}} \quad \text{Eq. S8}$$

70

71 where MLD_{max} is determined using an existing monthly Argo climatology product (3). A visual
72 example of this method is shown **Fig. S21**. The second method is to assume that DOM_{winter} is
73 equivalent to the DOM_{summer} profile concentration at the base of annual MLD_{max} (4). After
74 estimating DOM_{winter}, we can calculate the seasonal change in DOC and DON within z_{eu} to
75 quantify C: N_{DOM} (**Eq. S9**):

76

$$77 \text{C: N}_{\text{DOM}} = \frac{\frac{\Delta \text{DOC}}{\int_0^{z_{\text{eu}}} \text{DOC}_{\text{summer}} dz - z_{\text{eu}} \times \text{DOC}_{\text{winter}}}}{\frac{\Delta \text{DON}}{\int_0^{z_{\text{eu}}} \text{DON}_{\text{summer}} dz - z_{\text{eu}} \times \text{DON}_{\text{winter}}}} \quad \text{Eq. S9}$$

78

79 Before averaging the C: N_{DOM} values into 5° meridional bins, we remove implausible values
80 (e.g., <5 or >25) that we attribute to unresolved horizontal advection or EP effects. The meridional
81 patterns of C: N_{DOM} derived from the two DOM_{winter} reconstruction approaches generally agree
82 within their uncertainties (**Fig. S18b**). In this study, we use the C: N_{DOM} values derived from the
83 Baetge, *et al.* (2) approach as the subsequent carbon pool partitioning results exhibit better
84 agreement with ship-board observations (**Fig. 2d in the main text**).

86 1.2 Compilation of Shipboard Measurements of Sinking Particle Flux

87 We compile published shipboard observations of POC and PIC sinking flux (F_{POC} and F_{PIC})
 88 determined by ^{234}Th - ^{238}U disequilibrium in the SO during the productive season (**Fig. S20**; Henson,
 89 *et al.* (5), Le Moigne, *et al.* (6), Rosengard, *et al.* (7)). In these compiled dataset, F_{PIC} is estimated
 90 from F_{POC} multiplied by the PIC:POC ratio determined from contemporaneous observations of
 91 suspended particles (7). To maintain a consistent integration depth, we scale the F_{POC} and F_{PIC}
 92 values measured at different depth horizons to the base of z_{eu} using the Martin Curve and applying
 93 a global average attenuation coefficient (b) of 0.84 (8). The z_{eu} normalized F_{POC} and F_{PIC} values
 94 are then averaged into 5° meridional bins. Finally, the seasonal flux is inferred as F_{POC} multiplied
 95 by the average number of days in the productive season at each latitude band (**Fig. S1b**), assuming
 96 a steady flux over the course of the productive season.
 97

98 1.3 Residual Potential Nitrate Growth Rate and Dissolved Iron

99 We analyze the meridional pattern of Residual Potential Nitrate Growth (RNPG, (9)) and
 100 dissolved iron to gain insight into the environmental conditions that may favor coccolithophore
 101 blooms. RNPG is an index describing the residual potential growth rate for nitrate-dependent algae
 102 ($\mu_{\text{NO}_3^-}$) relative to silicate-dependent algae (μ_{Si}), under the observed nutrient conditions (9, 10)
 103 (**Eq. S15**):
 104

$$105 \text{RNPG} = \mu_{\text{NO}_3^-} - \mu_{\text{Si}} \quad \text{Eq. S15}$$

106
 107 The sign of RNPG indicates which type of algal group is expected to outcompete the other,
 108 assuming trace metal limitation is relieved. $\mu_{\text{NO}_3^-}$ can be modeled as the Michaelis-Menten
 109 formulation (**Eq. S16**):
 110

$$111 \mu_{\text{NO}_3^-} = \mu_{\text{NO}_3^-_{\text{max}}} \times \frac{\text{NO}_3^-}{K_{\text{NO}_3^-_{\text{max}}} + \text{NO}_3^-} \quad \text{Eq. S16}$$

112
 113 where $\mu_{\text{NO}_3^-_{\text{max}}}$ represents the maximum nitrate-dependent growth rate in the absence of substrate
 114 limitation, $K_{\text{NO}_3^-_{\text{max}}}$ is the half-saturation coefficient for the growth, and NO_3^- is the seawater
 115 concentration from float measurements. Unlike with NO_3^- , it is thought that phytoplankton are
 116 unable to fully exhaust seawater silicate (Si). Thereby, a residual Si term (Si_0) is added in the μ_{Si}
 117 parameterization as follows (**Eq. S17**):
 118

$$119 \mu_{\text{Si}} = \mu_{\text{Si}_{\text{max}}} \times \frac{\text{Si} - \text{Si}_0}{K_{\text{Si}_{\text{max}}} + \text{Si}_0} \quad \text{Eq. S17}$$

120
 121 where Si_0 is set to 0.7 umol L^{-1} (9). We estimate seawater Si along the float trajectory using the
 122 CANYON-B algorithm (11). The maximum growth and half-saturation coefficient are dependent
 123 on the nutrient condition, which can be modeled as a function of substrate concentration (S,
 124 corresponding to seawater NO_3^- and Si concentration in our study, **Eq. S18, and Eq. S19**):
 125

126
127
128
129
130
131
132
133
134
135
136
137
138
139
140
141
142
143
144
145
146
147
148
149
150
151
152
153
154
155
156
157
158
159
160
161
162
163
164
165
166
167
168

$$\mu_{S_max} = \frac{S}{S+2} \times 3 \quad \text{Eq. S18}$$

$$K_S = \frac{S}{S+3} \times 5.5 \quad \text{Eq. S19}$$

Dissolved iron concentrations are extracted from a gridded, global, monthly climatology product that was created using machine learning to gap fill shipboard observations (12). We interpolate the dissolved iron product in 3-dimensions (location, time, and sampling depth) to align with each float profile before averaging the result into 5° meridional bins for comparison with our PIC production estimates (Fig. S7).

1.4 Annual CO₂ Flux Estimate in the Southern Ocean and Comparison with Prior Studies

Our float-based estimate of the annual Southern Ocean (south of 35°S) carbon sink (F_{CO_2} ; -0.43 ± 0.14 Pg C yr⁻¹; where negative values indicate ocean carbon uptake) is larger in magnitude than prior float-based estimates (-0.08 ± 0.55 Pg C yr⁻¹, Gray, *et al.* (13), and -0.35 ± 0.19 Pg C yr⁻¹, Bushinsky, *et al.* (14)), indicating a stronger CO₂ sink. This is somewhat surprising because, unlike prior studies that applied a pH-dependent pH adjustment to quality-controlled pH values that yields lower pCO_2 estimates, we expected to find a weaker annual CO₂ sink in the Southern Ocean. However, it is worth noting that prior studies relied on float observations from May 2014 to April 2017, reflecting significantly fewer observations than our study. To test the importance of data density, we recalculate F_{CO_2} using only the float observations from May 2014 to April 2017 and replicating all flux calculation procedures from prior studies (13) except for the pH-dependent pH adjustment. This includes the use of ERA-interim wind speed and the division of the Southern Ocean into five frontal regions, with the PAZ further partitioned into the PFZ and ASZ. We find the SO to be a source of CO₂ to the atmosphere ($+0.28 \pm 0.076$ Pg C yr⁻¹), matching expectations. Recent finding by Stammerjohn, *et al.* (15) similarly found that greater data coverage from the current, larger float array leads to an elevated SO CO₂ sink estimate. Therefore, the difference in data coverage and/or inter-annual variability could be contributing to the larger SO CO₂ uptake found in our work using float observations from 2014 to 2021.

Notably, our float-based SO F_{CO_2} estimate still suggests a weaker annual sink than estimates derived from shipboard observations upscaled to the entire SO using machine learning techniques (~ -1.1 Pg C yr⁻¹ in the south of 35°S) (14, 16, 17). These gap-filled data products have known seasonal sampling biases in their training datasets due to a lack of wintertime observations in high-latitude regions that are anticipated to exhibit strong seasonal CO₂ outgassing (13, 18). A recent study by Long, *et al.* (19), using atmospheric CO₂ gradient observations from aircraft, reported nearly double the CO₂ influx (-0.53 ± 0.23 Pg C yr⁻¹; average over 2009-2018) of our float-based estimate in the region south of 45° S (-0.26 ± 0.07 Pg C yr⁻¹). Further study is needed to reduce the uncertainty in Southern Ocean F_{CO_2} estimates by considering potential discrepancies in methodology, inter-annual variability, and sampling coverage.

1.5 Potential Biases in Float-based POC and PIC Estimates and the Implications in Carbon Pool Partitioning

169 Float-based POC concentrations ($\text{POC}_{\text{b}_{\text{bp}}}$) in our study are derived from float b_{bp} observations
 170 using a global empirical relationship (20). This might not be suitable in regions with
 171 coccolithophore blooms (e.g., 47-57 °S in our study, **Fig. S7**) where highly-refractive calcite
 172 particles (coccoliths) could lead to elevated b_{bp} signals (9). Such a bias in our $\text{POC}_{\text{b}_{\text{bp}}}$ estimate
 173 could distort the magnitude and meridional pattern of net biological production solved from the
 174 $\text{POC}_{\text{b}_{\text{bp}}}$ budget $\left(\frac{\partial \text{POC}_{\text{b}_{\text{bp}}}}{\partial t}\right)_{\text{Bio}}$. Further, since the net biological production of suspended PIC
 175 $\left(\frac{\partial \text{PIC}_{\text{b}_{\text{bp}}}}{\partial t}\right)_{\text{Bio}}$ is calculated from $\frac{\partial \text{POC}_{\text{b}_{\text{bp}}}}{\partial t}\bigg|_{\text{Bio}}$ multiplied by the satellite derived surface PIC:POC
 176 ratio (**Eq. 7** in the main text), $\frac{\partial \text{PIC}_{\text{b}_{\text{bp}}}}{\partial t}\bigg|_{\text{Bio}}$ will be influenced by the assumption of a constant
 177 PIC:POC ratio throughout the euphotic zone column.

178 Our work focuses on variability in the export potential of distinct biogenic carbon pools
 179 $\left(\frac{\partial \text{DIC}}{\partial t}\bigg|_{\text{Bio_POC}}, \frac{\partial \text{DIC}}{\partial t}\bigg|_{\text{Bio_DOC}}, \frac{\partial \text{DIC}}{\partial t}\bigg|_{\text{Bio_PIC}}\right)$ and the associated particle sinking fluxes (F_{POC} and F_{PIC}).
 180 The former relies on linking multiple chemical tracer budgets (**Eq. 4-5 in the main text**), which
 181 eliminates the relevance of a potential $\text{POC}_{\text{b}_{\text{bp}}}$ and $\text{PIC}_{\text{b}_{\text{bp}}}$ bias. However, such a bias would
 182 propagate into the subsequent computations of F_{POC} and F_{PIC} , which are determined by subtracting
 183 $\frac{\partial \text{POC}_{\text{b}_{\text{bp}}}}{\partial t}\bigg|_{\text{Bio}}$ (or $\frac{\partial \text{PIC}_{\text{b}_{\text{bp}}}}{\partial t}\bigg|_{\text{Bio}}$) from $\frac{\partial \text{DIC}}{\partial t}\bigg|_{\text{Bio_POC}}$ (or $\frac{\partial \text{DIC}}{\partial t}\bigg|_{\text{Bio_PIC}}$). It is worth pointing out that the
 184 magnitudes of biological terms solved from suspended particles are one-tenth of the corresponding
 185 chemically derived export potential values (**Fig. S22**). Therefore, F_{POC} and F_{PIC} are less sensitive
 186 to potential errors in the $\text{POC}_{\text{b}_{\text{bp_bio}}}$ and $\text{PIC}_{\text{b}_{\text{bp_bio}}}$ estimates.

187

188 1.6 Tracer Budgets

189 We use a 1-dimensional mass balance model approach to account for processes affecting the
 190 time rate of concentration change ($\text{mmol m}^{-3} \text{d}^{-1}$) of biologically relevant tracers (DIC, TA, NO_3^- ,
 191 and $\text{POC}_{\text{b}_{\text{bp}}}$) that are observed by profiling floats within the euphotic zone (z_{eu}) (**Eq. S1**):

192

$$\begin{aligned}
 & \frac{dT_{(\text{DIC,TA,NO}_3^-, \text{POC}_{\text{b}_{\text{bp}}})}}{dt}\bigg|_{\text{Obs}_{z_{\text{eu}}}} = \\
 & \frac{\partial T_{(\text{DIC})}}{\partial t}\bigg|_{\text{Gas}_{z_{\text{eu}}}} + \frac{\partial T_{(\text{DIC,TA,NO}_3^-, \text{POC}_{\text{b}_{\text{bp}}})}}{\partial t}\bigg|_{\text{Phys}_{z_{\text{eu}}}} + \frac{\partial T_{(\text{DIC,TA,NO}_3^-, \text{POC}_{\text{b}_{\text{bp}}})}}{\partial t}\bigg|_{\text{EP}_{z_{\text{eu}}}} + \\
 & \frac{\partial T_{(\text{DIC,TA,NO}_3^-, \text{POC}_{\text{b}_{\text{bp}}})}}{\partial t}\bigg|_{\text{Bio}_{z_{\text{eu}}}} \quad \text{Eq. S1}
 \end{aligned}$$

196

197 where subscripts on the right-hand side represent air-sea gas exchange (Gas), physical transport
 198 and mixing (Phys), evaporation and precipitation (EP), and biological activity (Bio), respectively.
 199 The computation of air-sea gas exchange for the dissolved inorganic carbon (DIC) budget is
 200 described in the methods section.

201 When z_{eu} is deeper than the mixed layer depth (MLD; defined by a temperature increase of
 202 0.2 °C relative to the 10 m temperature, following de Boyer Montégut, *et al.* (21), physical
 203 transport and mixing at the base of z_{eu} consists of diapycnal diffusion and wind-induced Ekman
 204 pumping (shown here for DIC, **Eq. S2**):

205

206

$$\frac{\partial \text{DIC}}{\partial t} |_{\text{Phys}_{z_{\text{eu}}}} = \underbrace{\left(K_{z_{\text{zeu}}} \times \frac{\partial \text{DIC}}{\partial z} \Big|_{z_{\text{eu}}} \right)}_{\text{Diapycnal diffusion}} + \underbrace{\left(w_{z_{\text{eu}}} \times \int_0^{z_{\text{eu}}} \frac{\partial \text{DIC}}{dz} \right)}_{\text{Ekman pumping}} / z_{\text{eu}}; \text{ for } z_{\text{eu}} > \text{MLD} \quad \text{Eq. S2}$$

207

208

209

210

211

212

213

214

215

216

217

218

219

220

221

222

223

224

225

226

227

228

229

230

231

232

233

234

235

236

237

238

239

240

where $K_{z_{\text{zeu}}}$ is the diapycnal diffusivity coefficient, $\frac{\partial \text{DIC}}{\partial z} \Big|_{z_{\text{eu}}}$ is the vertical DIC gradient across z_{eu} , $w_{z_{\text{eu}}}$ is the Ekman pumping velocity computed from satellite-derived wind stress following Signorini, *et al.* (22), and $\int_0^{z_{\text{eu}}} \frac{\partial \text{DIC}}{dz}$ is the depth integrated DIC vertical gradient. In this study, we set K_Z to 10^{-4} m s^{-2} at the base of the mixed layer (23) with an exponential decay to the background value of 10^{-5} m s^{-2} over 20 m following 1/e scaling (24). We recognize the present choice of K_Z is somewhat arbitrary. Nevertheless, our analysis focuses on the spring/summer season when the upper layer water column is stratified and stable. Therefore, we believe the diapycnal diffusion during this period is minimal and the magnitude of K_Z has a limited effect on the biological term estimate.

During periods when the MLD exceeds z_{eu} , $\frac{\partial \text{DIC}}{\partial t} |_{\text{Phys}_{z_{\text{eu}}}}$ is scaled from the physical transport and mixing at the base MLD $\left(\frac{\partial \text{DIC}}{\partial t} |_{\text{Phys}_{\text{MLD}}} \right)$ by assuming the impacts of physical transport and mixing occurring at the base of MLD will be equally distributed throughout the entire mixed water column (**Eq. S3**):

$$\frac{\partial \text{DIC}}{\partial t} |_{\text{Phys}_{z_{\text{eu}}}} = \frac{\partial \text{DIC}}{\partial t} |_{\text{Phys}_{\text{MLD}}} \times \frac{z_{\text{eu}}}{\text{MLD}}; \text{ for } z_{\text{eu}} < \text{MLD} \quad \text{Eq. S3}$$

Compared with $\frac{\partial \text{DIC}}{\partial t} |_{\text{Phys}_{z_{\text{eu}}}}$ shown in **Eq. S2**, $\frac{\partial \text{DIC}}{\partial t} |_{\text{Phys}_{\text{MLD}}}$ encompasses an additional term, entrainment, to account for tracer changes induced by MLD changes (**Eq. S4**):

$$\frac{\partial \text{DIC}}{\partial t} |_{\text{Phys}_{\text{MLD}}} = \underbrace{\left(K_{z_{\text{MLD}}} \times \frac{\partial \text{DIC}}{\partial z} \Big|_{\text{MLD}} \right)}_{\text{Diapycnal diffusion}} + \underbrace{\left(w_{\text{MLD}} \times \int_0^{\text{MLD}} \frac{\partial \text{DIC}}{dz} \right)}_{\text{Ekman pumping}} + \underbrace{\left(\text{DIC}_{\text{base}} - \overline{\text{DIC}_{\text{MLD}}} \right) \times \frac{\partial \text{MLD}}{\partial t}}_{\text{Entrainment}} / \text{MLD}$$

Eq. S4

Entrainment can be modeled as a product of the MLD time rate of change $\left(\frac{\partial \text{MLD}}{\partial t} \right)$ and the difference between the average DIC concentration within MLD $\left(\overline{\text{DIC}_{\text{MLD}}} \right)$ and DIC concentration at the base of MLD $\left(\text{DIC}_{\text{base}} \right)$. The entrainment term is set to zero when the MLD shoals $\left(\frac{\partial \text{MLD}}{\partial t} < 0 \right)$.

The evaporation and precipitation term is quantified using the ratio of tracer to salinity at initial time t_1 $\left(\frac{T_{(\text{DIC,TA,NO}_3^-, \text{POC}_{\text{bBP}})}}{\text{Sal}} \Big|_{t_1} \right)$ multiplied by the salinity time rate of change due to EP $\left(\frac{\partial \text{Sal}}{\partial t} \Big|_{\text{EP}_{z_{\text{eu}}}} \right)$ (**Eq. S5**):

$$\frac{\partial T_{(\text{DIC,TA,NO}_3^-, \text{POC}_{\text{bBP}})}}{\partial t} \Big|_{\text{EP}_{z_{\text{eu}}}} = \frac{\partial \text{Sal}}{\partial t} \Big|_{\text{EP}_{z_{\text{eu}}}} \times \frac{T_{(\text{DIC,TA,NO}_3^-, \text{POC}_{\text{bBP}})}}{\text{Sal}} \Big|_{t_1} \quad \text{Eq. S5}$$

241 $\frac{\partial \text{Sal}}{\partial t} |_{\text{EP}_{\text{zeu}}}$ is computed from the difference between observed salinity changes and estimated
 242 physically driven salinity changes (**Eq. S6**):

$$243 \quad \frac{\partial \text{Sal}}{\partial t} |_{\text{EP}_{\text{zeu}}} = \frac{\partial \text{Sal}}{\partial t} |_{\text{obs}_{\text{zeu}}} - \frac{\partial \text{Sal}}{\partial t} |_{\text{Phys}_{\text{zeu}}} \quad \text{Eq. S6}$$

244
 245
 246 After accounting for abiotic terms, the biological term can be calculated as a residual.

247 248 249 **1.7 Reconstruction of DIC and TA for Different Productivity Scenarios**

250 To assess the role of biology in maintaining the Southern Ocean carbon sink, we reconstruct
 251 DIC and TA time series for different scenarios of modified productivity: only organic matter
 252 production and zero biological production (abiotic). Float-derived DIC values ($\text{DIC}_{\text{Obs}_n}$) can be
 253 expressed as a sum of the DIC concentration at the start of the productive season ($\text{DIC}_{\text{Obs}_1}$) and
 254 the time integral of DIC changes from biology ($\sum_{\text{day}_1}^{\text{day}_{n-1}} \frac{\partial \text{DIC}}{\partial t} |_{\text{Bio}}$), air-sea gas exchange
 255 ($\sum_{\text{day}_1}^{\text{day}_{n-1}} \frac{\partial \text{DIC}}{\partial t} |_{\text{Gas}}$), and other abiotic processes ($\sum_{\text{day}_1}^{\text{day}_{n-1}} \frac{\partial \text{DIC}}{\partial t} |_{\text{Other}}$) including
 256 evaporation/precipitation, and physical transport and mixing (**Eq. S10**):

$$257 \quad \text{DIC}_{\text{Obs}_n} = \text{DIC}_{\text{Obs}_1} + \sum_{\text{day}_1}^{\text{day}_{n-1}} \frac{\partial \text{DIC}}{\partial t} |_{\text{Bio}} + \sum_{\text{day}_1}^{\text{day}_{n-1}} \frac{\partial \text{DIC}}{\partial t} |_{\text{Gas}} + \sum_{\text{day}_1}^{\text{day}_{n-1}} \frac{\partial \text{DIC}}{\partial t} |_{\text{Other}} \quad \text{Eq. S10}$$

258
 259 Likewise, the time-series of DIC in the reconstructed, abiotic ocean ($\text{DIC}_{\text{abio}_n}$) can be written as
 260 follows (**Eq. S11**):

$$261 \quad \text{DIC}_{\text{abiotic}_n} = \text{DIC}_{\text{Obs}_1} + \sum_{\text{day}_1}^{\text{day}_{n-1}} \frac{\partial \text{DIC}}{\partial t} |_{\text{Gas}_{\text{abiotic}}} + \sum_{\text{day}_1}^{\text{day}_{n-1}} \frac{\partial \text{DIC}}{\partial t} |_{\text{Other}_{\text{abiotic}}} \quad \text{Eq. S11}$$

262
 263 For simplicity, we assume the “other” terms are equivalent between the observed and reconstructed
 264 scenarios ($\sum_{\text{day}_1}^{\text{day}_{n-1}} \frac{\partial \text{DIC}}{\partial t} |_{\text{Other}} \approx \sum_{\text{day}_1}^{\text{day}_{n-1}} \frac{\partial \text{DIC}}{\partial t} |_{\text{Other}_{\text{abiotic}}}$), and rearrange $\text{DIC}_{\text{abio}_n}$ by linking **Eq.**
 265 **S10** with **Eq. S11** (**Eq. S12**, same as **Eq. 12** in the main text):

$$266 \quad \text{DIC}_{\text{abiotic}_n} = \text{DIC}_{\text{Obs}_n} - \sum_{\text{day}_1}^{\text{day}_{n-1}} \frac{\partial \text{DIC}}{\partial t} |_{\text{Bio}} - \left(\sum_{\text{day}_1}^{\text{day}_{n-1}} \frac{\partial \text{DIC}}{\partial t} |_{\text{Gas}} - \sum_{\text{day}_1}^{\text{day}_{n-1}} \frac{\partial \text{DIC}}{\partial t} |_{\text{Gas}_{\text{abiotic}}} \right) \quad \text{Eq. S12}$$

267
 268 As shown in **Eq. S12**, the abiotic DIC time series can be inferred from the float-derived DIC at the
 269 corresponding time step, with an adjustment for the time integral of biological production and the
 270 gas exchange difference between the two scenarios. A similar approach can be used to derive the
 271 abiotic TA time series ($\text{TA}_{\text{abiotic}_n}$), which excludes the influence of gas exchange (**Eq. S13**, same
 272 as **Eq. 11** in the main text):

$$273 \quad \text{TA}_{\text{abiotic}_n} = \text{TA}_{\text{Obs}_n} - \sum_{\text{day}_1}^{\text{day}_{n-1}} \frac{\partial \text{TA}}{\partial t} |_{\text{Bio}} \quad \text{Eq. S13}$$

274
275
276
277
278
279

280 Since TA_{Obs_n} and $\frac{\partial TA}{\partial t}|_{Bio_n}$ are previously quantified terms, we can directly reconstruct $TA_{abiotic}$
 281 following **Eq. S13**.

282
 283 The $\frac{\partial DIC}{\partial t}|_{Gas_abiotic}$ term required to compute $DIC_{abiotic}$ in **Eq. S12** is determined iteratively.
 284 For the initial time step ($n=1$), we assume the DIC and TA concentrations and gas exchange terms
 285 are identical between the observed (biotic) and reconstructed (abiotic) ocean ($DIC_{abiotic_1} =$
 286 DIC_{Obs_1} ; $TA_{abiotic_1} = TA_{Obs_1}$; $\frac{\partial DIC}{\partial t}|_{Gas_abiotic_1} = \frac{\partial DIC}{\partial t}|_{Gas_1}$). It follows that there is also no
 287 difference between the biotic and abiotic gas exchange terms for the subsequent time step ($n = 2$,
 288 **Eq. S14**).

$$289 \quad DIC_{abiotic_2} = DIC_{Obs_2} - \sum_{day_1}^{day_1} \frac{\partial DIC}{\partial t}|_{Bio} - \underbrace{\left(\sum_{day_1}^{day_1} \frac{\partial DIC}{\partial t}|_{Gas} - \sum_{day_1}^{day_1} \frac{\partial DIC}{\partial t}|_{Gas_abiotic} \right)}_{set\ to\ 0} \quad \text{Eq. S14}$$

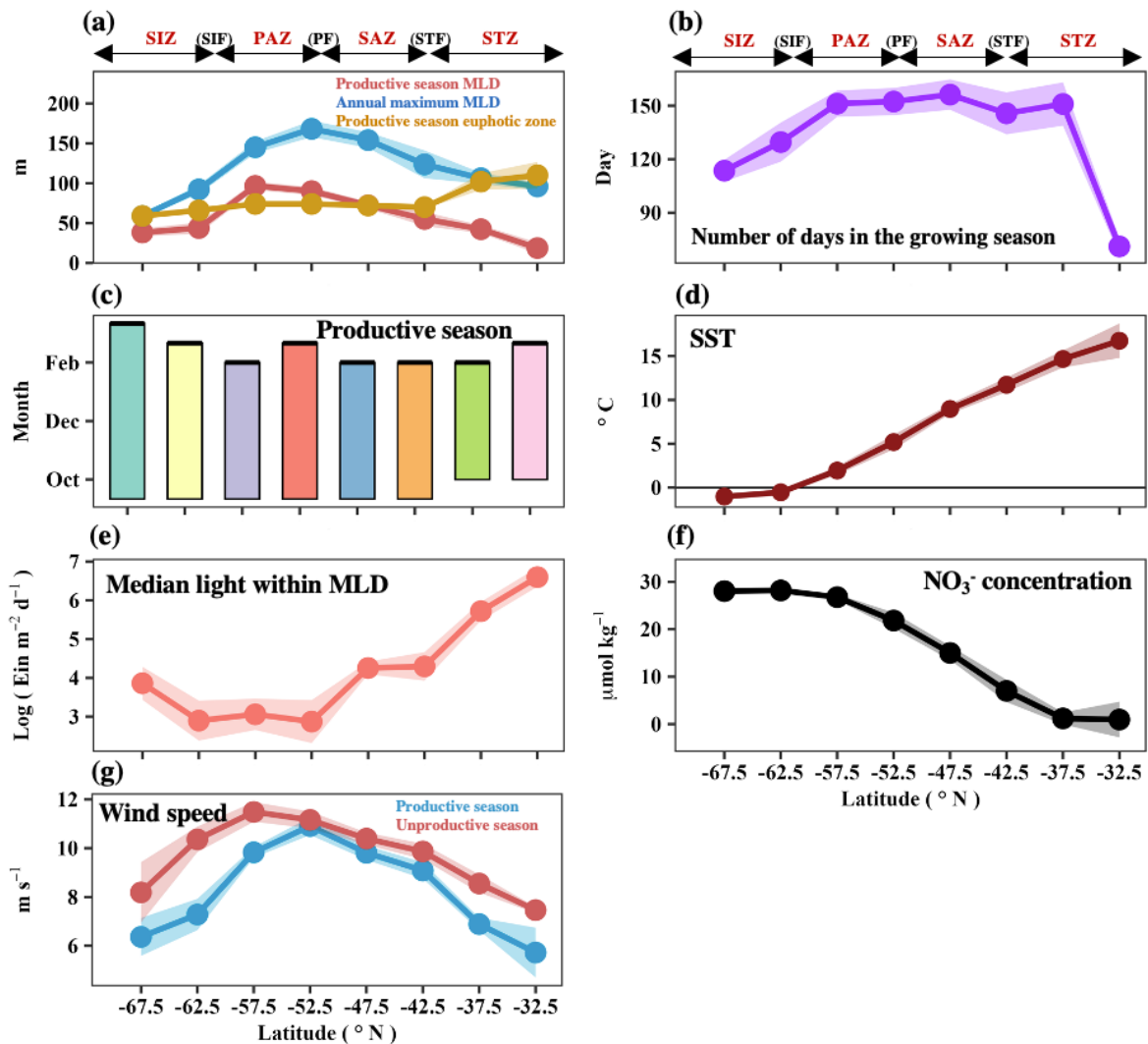
290
 291 With $DIC_{abiotic_2}$ and $TA_{abiotic_2}$ we can calculate the abiotic seawater partial pressure of CO_2
 292 ($pCO_{2_sea_abiotic_2}$) and the associated air-sea gas exchange ($\frac{\partial DIC}{\partial t}|_{Gas_abiotic_2}$). $DIC_{abiotic_3}$, and all
 293 subsequent time steps, can then be computed following **Eq. S12**. Through this procedure, we
 294 iteratively reconstruct the time-series of $DIC_{abiotic}$, $pCO_{2_sea_abiotic}$, and $\frac{\partial DIC}{\partial t}|_{Gas_abiotic}$. A visual
 295 example of our reconstruction for different productivity scenarios is provided in **Fig. S23**.

296
 297
 298 We could infer $DIC_{abiotic_n}$ from $DIC_{abiotic_n-1} + \frac{\partial DIC}{\partial t}|_{Gas_abiotic_n-1}$; however, any biases in the
 299 gas exchange estimates would accumulate iteratively, which could pull our abiotic terms far from
 300 the starting DIC value. Considering that the gas exchange velocity (k) used to parameterize air-sea
 301 CO_2 flux has a ~30% uncertainty (25, 26), we opt for a method that starts with the float-based DIC
 302 estimate at each time step to ensure we are not far from reality. This approach also ensures that
 303 pCO_2 changes caused by other physical processes, such as horizontal advection, are not
 304 unintentionally omitted.

305 306 **1.8 Exclusion of Additional pH-dependent pH Adjustment to Quality-controlled SOCCOM** 307 **Float pH for pCO_2 Computation**

308
 309 The calculation of pH from DIC and TA does not always agree well with directly measured
 310 pH values, and this bias can exhibit a pH dependency. Such a pH-dependent pH bias has been
 311 identified in numerous high-quality shipboard datasets in which at least three carbonate chemistry
 312 parameters (including pH) were measured (27-29). In effort to align float-based pCO_2 estimates
 313 with pCO_2 values that would be computed from high-quality DIC and TA observations, a pH-
 314 dependent pH adjustment (28) has been routinely applied to quality-controlled SOCCOM float pH
 315 observations before computing the pCO_2 values provided in SOCCOM data snapshots
 316 (<https://socom.princeton.edu/content/data-access>). However, implementation of this adjustment
 317 is based on the pH-dependent pH bias at 1500 m, which is then applied to the entire profile as an
 318 offset. Thus, the pH adjustment is not applied in a pH-dependent manner and does not correct for
 319 the issue presented in Williams, *et al.* (28) and Carter, *et al.* (27), Carter, *et al.* (29). Since 2019, a
 320 collaborative international effort entitled the Ocean Carbonate System Intercomparison Forum
 321 (OCSIF), supported by the U.S. Ocean Carbon and Biogeochemistry Program (

322 ocb.org/ocean-carbonate-system-intercomparison-forum/), has focused on identifying the origin
323 of the pH-dependent pH bias in high-quality shipboard observations. While efforts remain ongoing,
324 the current OCSIF recommendation is to use the quality-controlled float pH data directly, without
325 any type of additional pH adjustment, for computations of $p\text{CO}_2$.
326



328

329

330

331

332

333

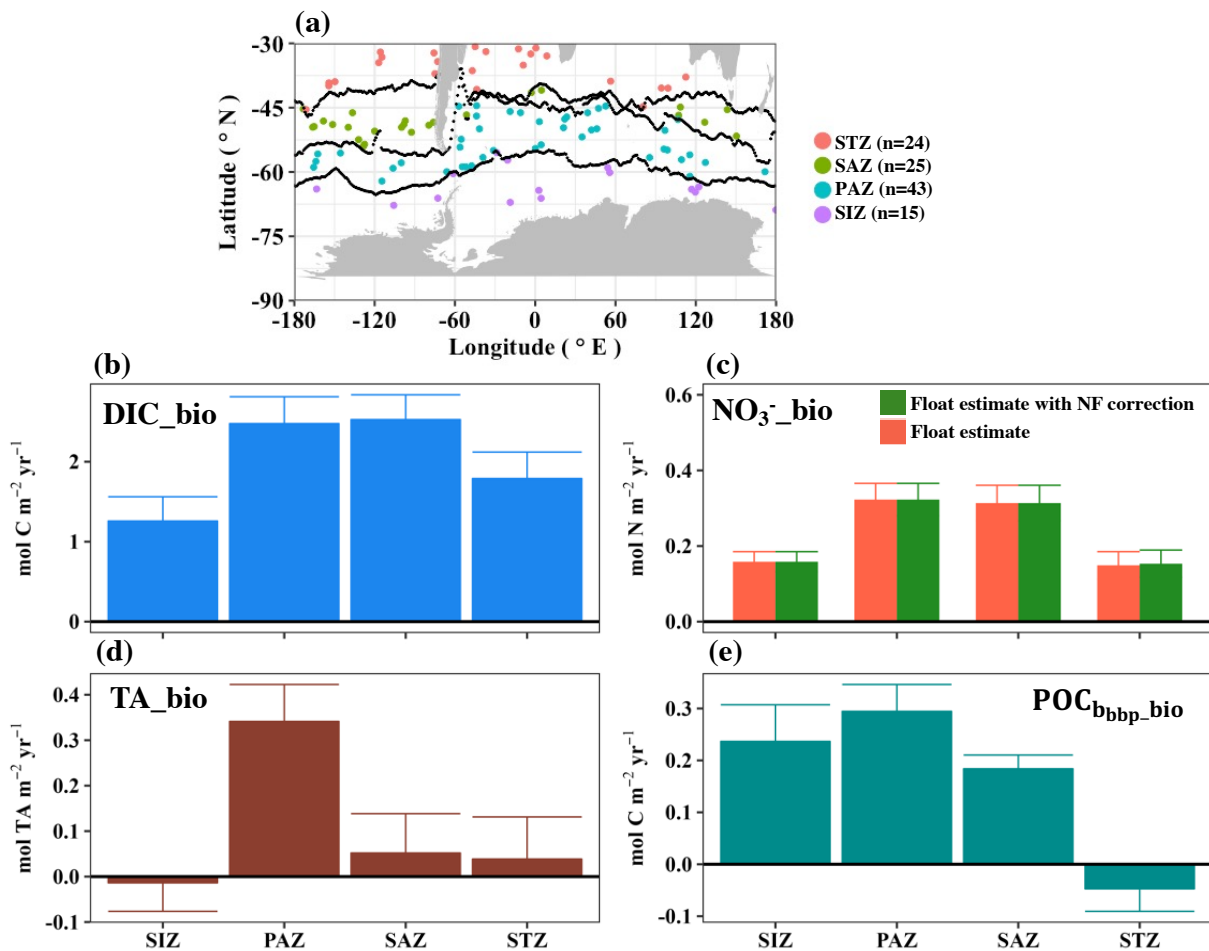
334

335

336

337

Fig. S1 The meridional patterns of environmental parameters during the Southern Ocean productive season, where shading indicates the propagated error. Median light within mixed layer depth (MLD) is computed based on the remotely sensed surface light field and its attenuation as inferred from float Chl-*a* profiles (30). Note that the frontal labels depicted on the top of the panels indicate rough locations of frontal regions, with a more precise representation appearing in **Fig. S2a**. STF: subtropical front; STZ: subtropical zone; SAZ: subantarctic zone; PF: polar front; PAZ: polar Antarctic zone; SIF: seasonal ice front; SIZ: sea ice zone; SST: sea surface temperature; NO_3^- : nitrate; MLD: mixed layer depth.



339
 340 **Fig. S2** Net biological production during the Southern Ocean productive season as averaged across
 341 the four main frontal zones. (a) Mean location of each float for each seasonal production estimate,
 342 with frontal zones labeled. Net biological term (bio) results for each frontal region determined
 343 from the (b) dissolved inorganic carbon (DIC), (c) nitrate (NO₃⁻), (d) total alkalinity (TA), and (e)
 344 particulate organic carbon (derived from particle backscattering coefficient; POC_{bbbp}_{bio}) tracer
 345 budgets. Error bars represent the propagated errors. Numbers in panel a indicate how many
 346 independent, float seasonal cycles were evaluated in each frontal zone. Zones as described in **Fig.**
 347 **S1**. NF: nitrogen fixation.

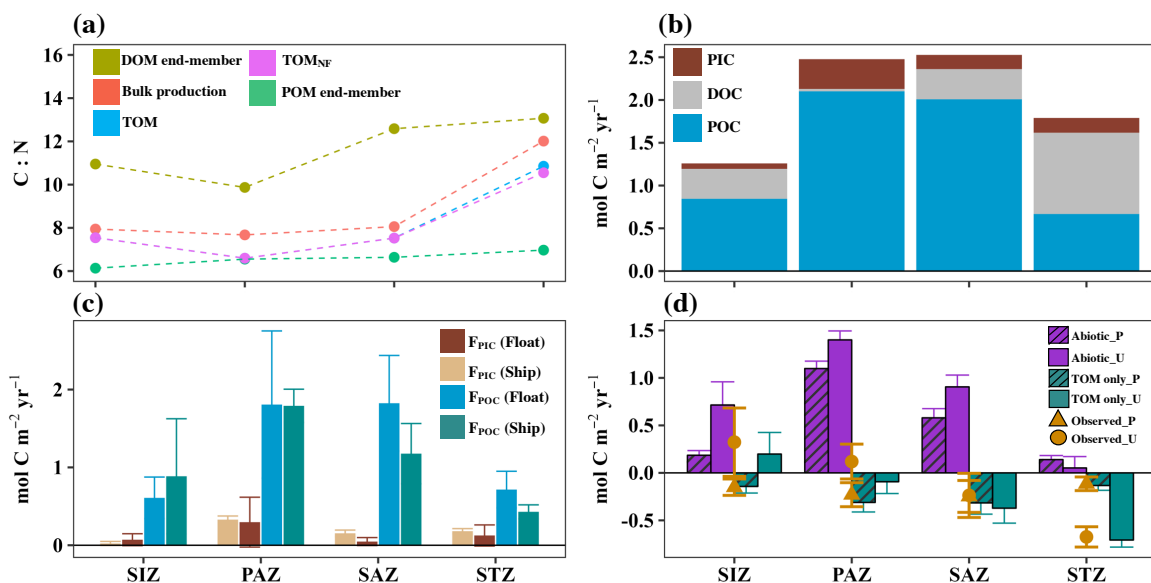
348

349

350

351

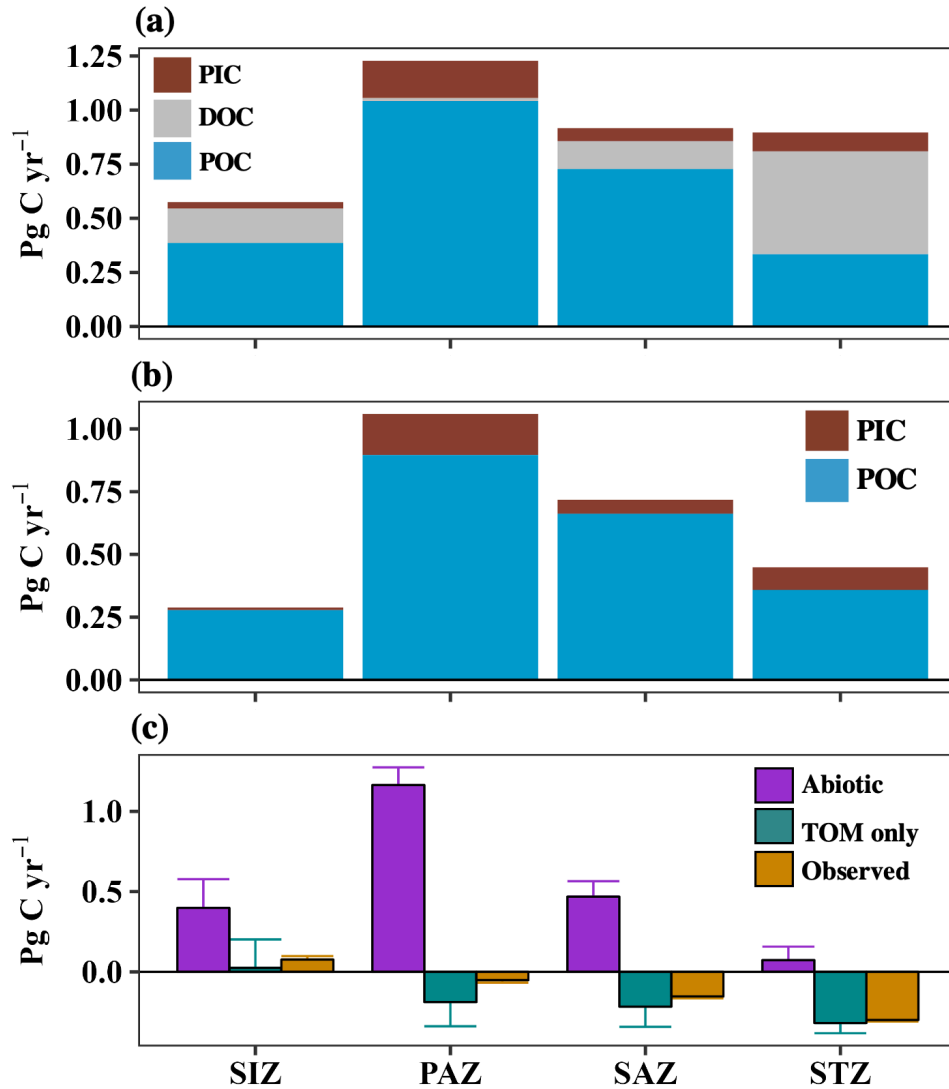
352



354
 355 **Fig. S3** (a) C:N ratio for biological production of different carbon pools, (b) magnitude and
 356 fraction of distinct carbon pool export potential, (c) sinking flux of particulate organic carbon (F_{POC})
 357 and particulate inorganic carbon (F_{PIC}) during the Southern Ocean productive season (P), and (d)
 358 influence of biology on air-sea CO₂ exchange for different productivity scenarios as averaged over
 359 the four main frontal zones. The dissolved organic matter (DOM) and particulate organic matter
 360 (POM) end-member C:N ratios are derived from ship-based observations. The bulk biological
 361 production, total organic matter (TOM) production, and TOM production with the correction for
 362 nitrogen fixation (TOM_{NF}) are derived from float observations and a biogeochemical inverse
 363 model. Error bars represent the propagated errors. Positive values in panel d indicate a source of
 364 carbon to the atmosphere. Zones as described in Fig. S1. DOM: dissolved organic matter; TOM:
 365 total organic matter; PIC: particulate inorganic carbon; POM: particulate organic matter; DOC:
 366 dissolved organic carbon; POC: particulate organic carbon. Abiotic: reconstructed, abiotic ocean;
 367 TOM only: reconstructed ocean with only total organic carbon production; Observed: observed,
 368 biotic ocean including both TOM and particulate inorganic carbon production. U: unproductive
 369 season.

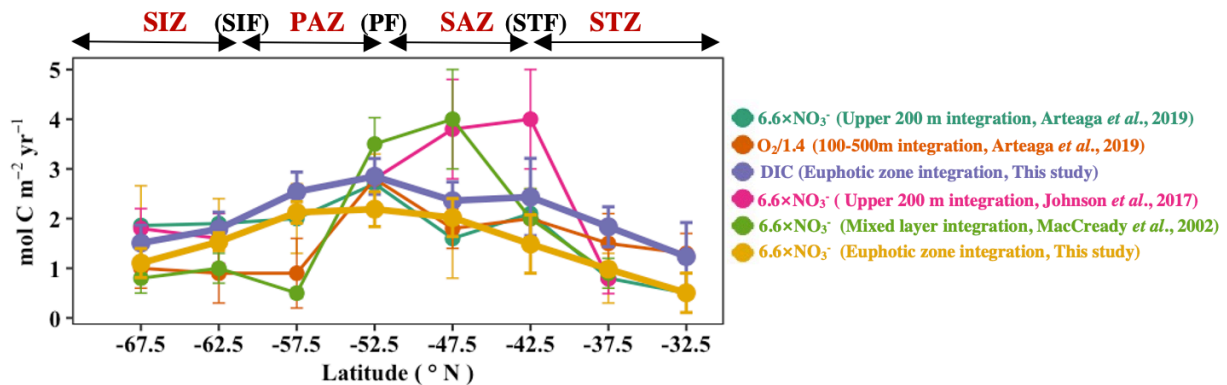
370

371



372
 373 **Fig. S4** Area-weighted cumulative annual (a) magnitude and (b) fraction of distinct carbon pool
 374 export potential, and (c) influence of biology on annual air-sea CO_2 exchange for different
 375 productivity scenarios as averaged over the four main frontal zones. Positive values in panel c
 376 indicate a source of carbon to the atmosphere. Error bars represent the propagated errors. Zones as
 377 described in **Fig. S1**. TOM: total organic matter; DOC: dissolved organic carbon. Note the annual
 378 biological production estimate is scaled from the float-estimated biological production during the
 379 productive season using data-constrained model estimates of the fraction of annual biological
 380 production that occurs during the productive period at each float location (*Method and Material*
 381 **in the main text**).

382
 383



385

386

387

388

389

390

391

392

393

394

395

396

397

398

399

400

401

402

403

404

405

406

407

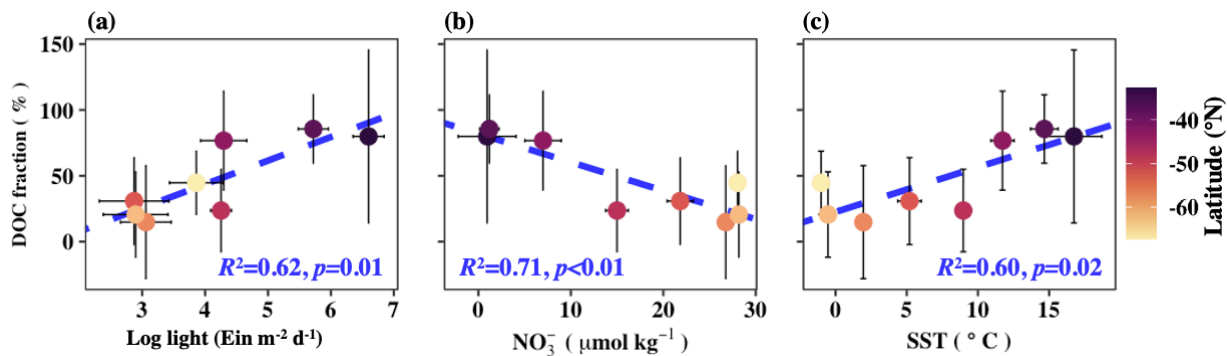
408

409

410

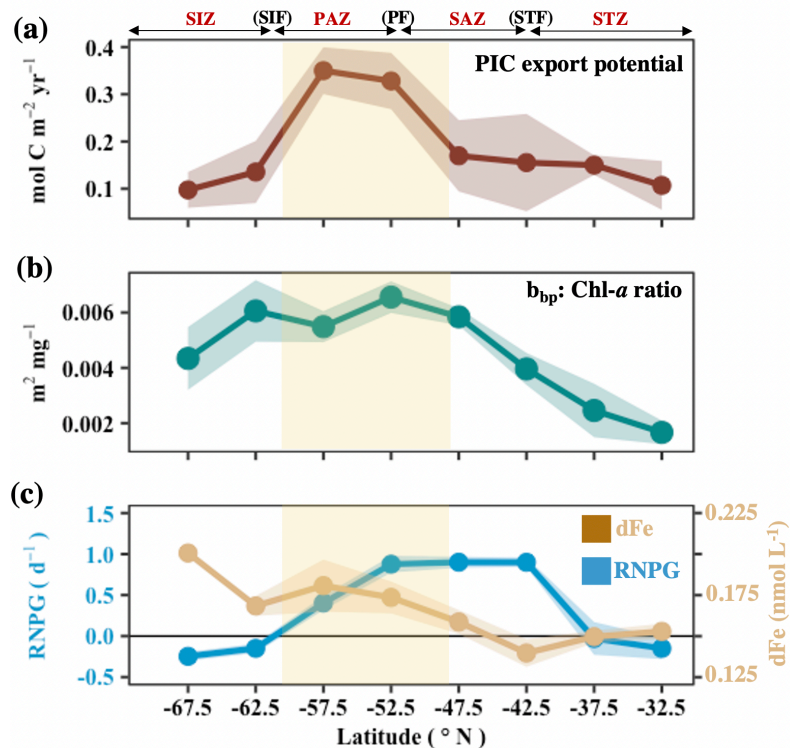
411

Fig. S5 Comparison of seasonal export potential during the Southern Ocean productive season solved from multiple studies. Export potential values derived from nitrate and oxygen are converted to units of carbon using the Redfield ratio to facilitate comparison. Error bars represent the propagated errors. Zones as described in **Fig. S1**. NO₃⁻: nitrate; DIC: dissolved inorganic carbon; O₂: oxygen.



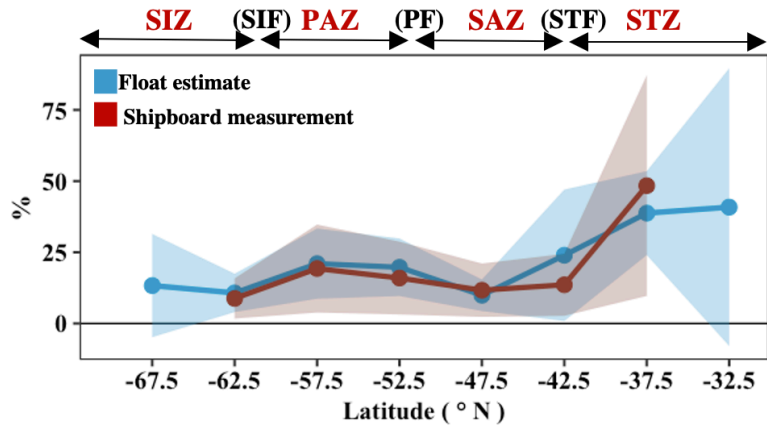
412
 413 **Fig. S6** Relationships between the fraction of total carbon export potential attributed to dissolved
 414 organic carbon (DOC) and various environmental parameters during the Southern Ocean
 415 productive season. Light availability in panel a refers to the median light field within the mixed
 416 layer (**Fig. S1e**). Error bars represent the propagated errors. NO_3^- : nitrate; SST: sea surface
 417 temperature.

418
 419
 420
 421
 422
 423
 424
 425
 426
 427
 428
 429
 430
 431
 432



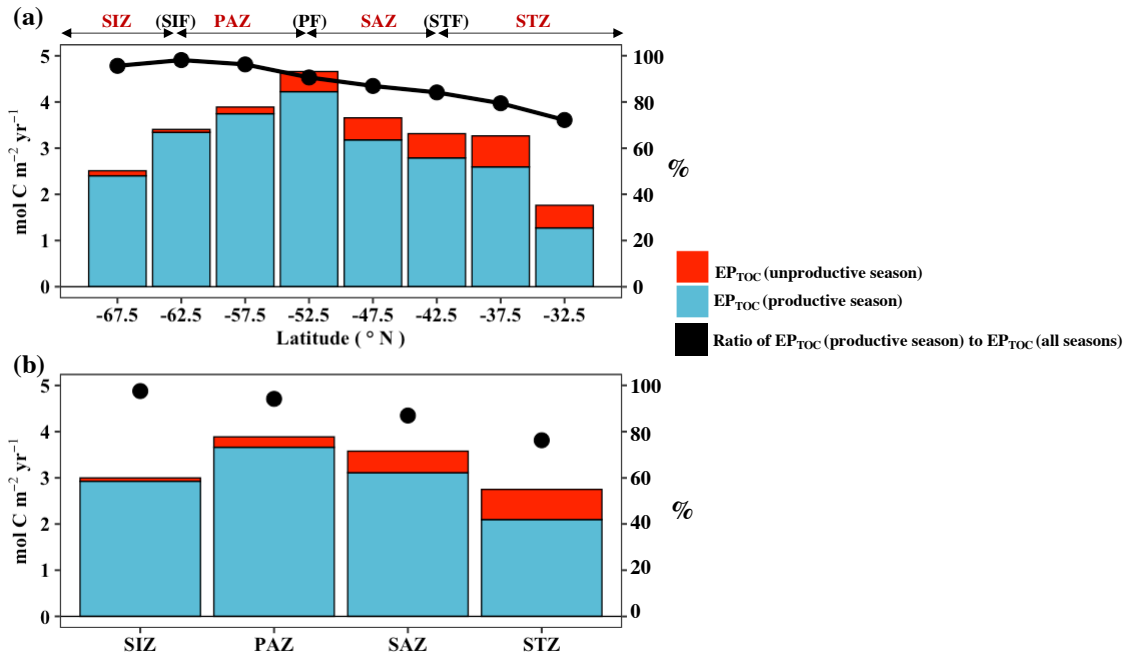
433
 434 **Fig. S7** The meridional pattern of (a) seasonal particulate inorganic carbon (PIC) export potential,
 435 (b) backscattering to Chlorophyll-*a* ratio (b_{bp} :Chl-*a*), and (c) dissolved iron (dFe) and residual
 436 nitrate potential growth rate (RNPG; detailed description in **Text S1.3**) during the Southern Ocean
 437 productive season. Yellow shading highlights the region with enhanced PIC production. Shading
 438 on each line reflects the propagated error. Zones as described in **Fig. S1**.

439
 440
 441
 442
 443
 444
 445
 446
 447
 448
 449



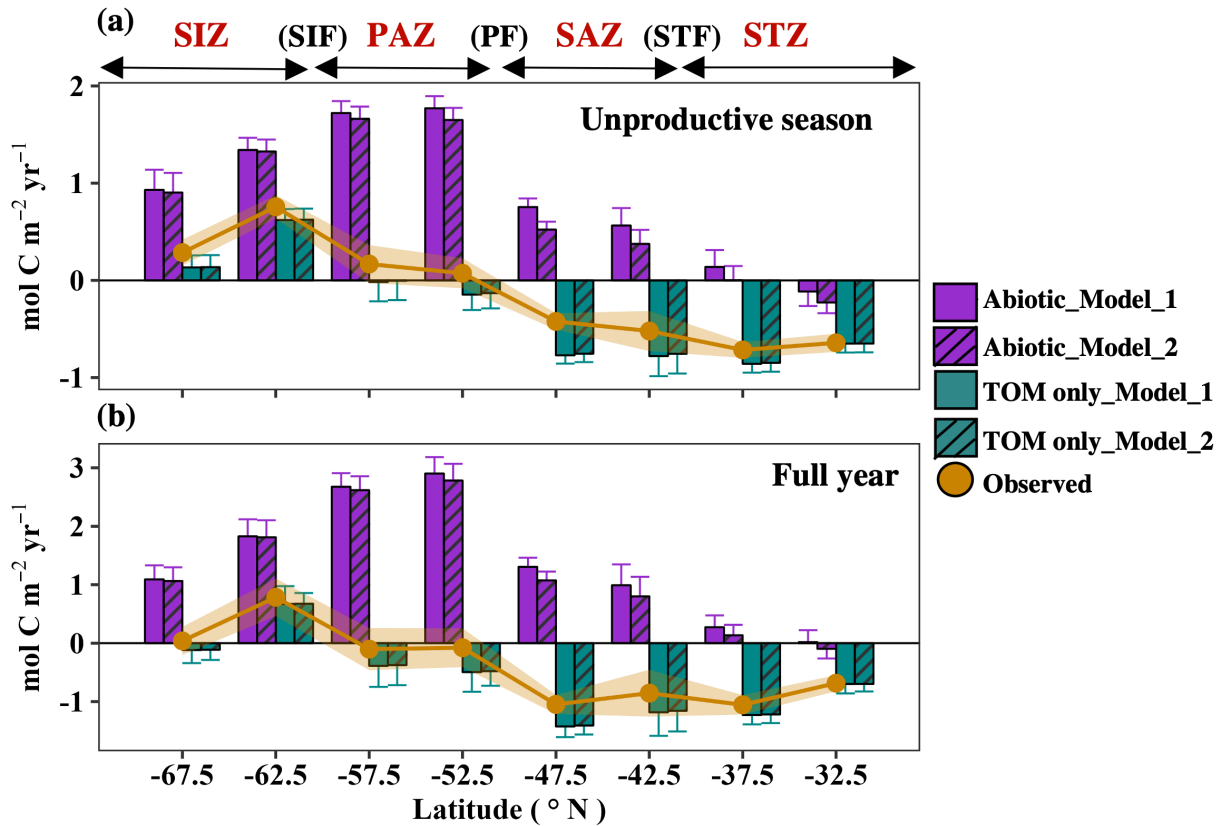
450
 451
 452
 453
 454
 455
 456
 457
 458
 459
 460
 461
 462
 463
 464
 465
 466
 467

Fig. S8 Meridional pattern (5° bin) of euphotic zone sinking particle rain ratio ($F_{PIC}/F_{POC} \times 100\%$) during the Southern Ocean productive season. Shading reflects the propagated error. Zones as described in **Fig. S1**.

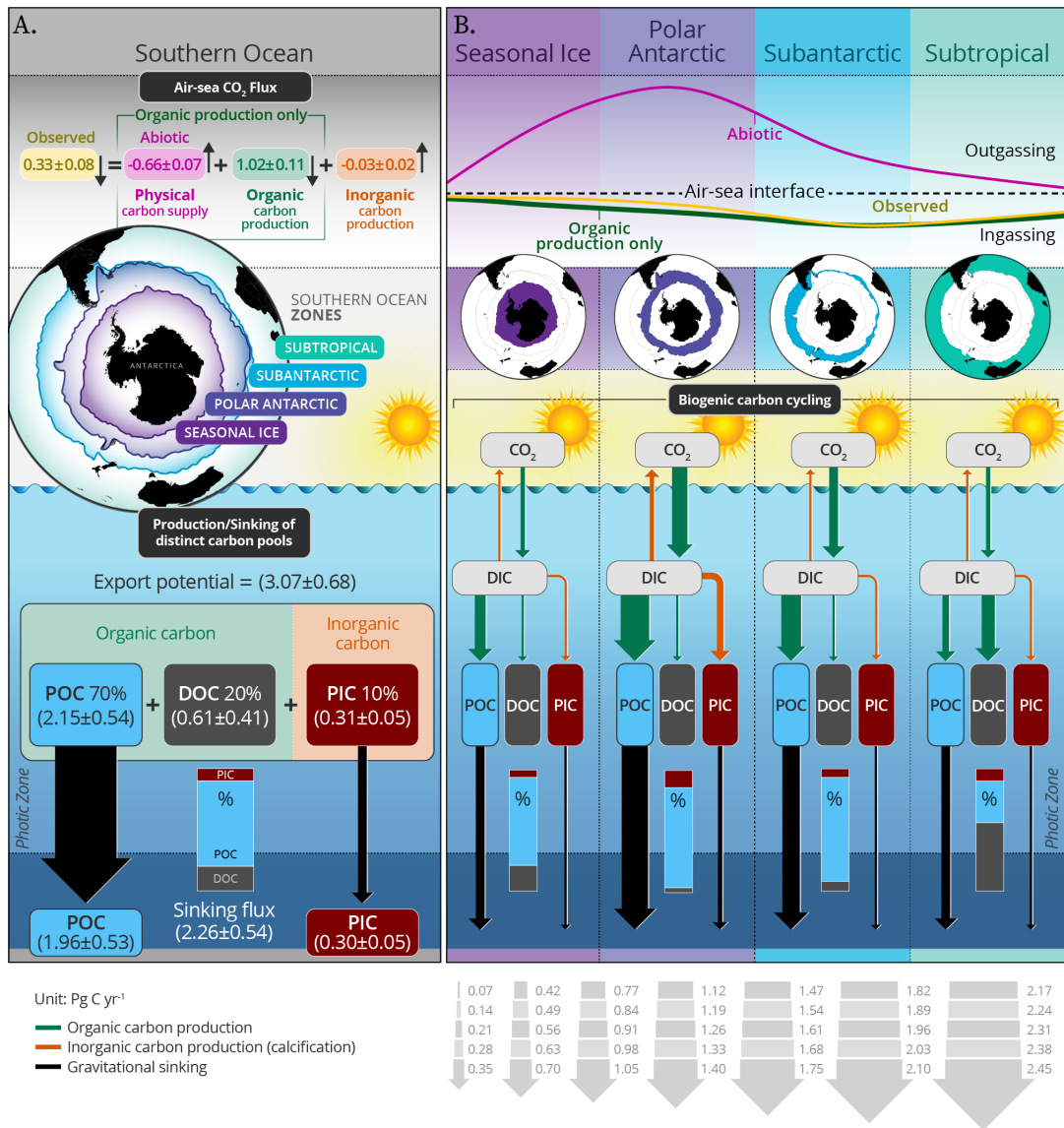


468
 469 **Fig. S9** Year 2008 to 2018 average euphotic zone integrated export potential of total organic
 470 carbon (EP_{TOC}) in the Southern Ocean for the productive and unproductive seasons derived from
 471 data-constrained biogeochemical Southern Ocean State Estimate (B-SOSE) model output (31).
 472 Results are shown for (a) 5° latitude bins and (b) the four main frontal regions. Frontal regions as
 473 described in **Fig. S1**.

474
 475
 476

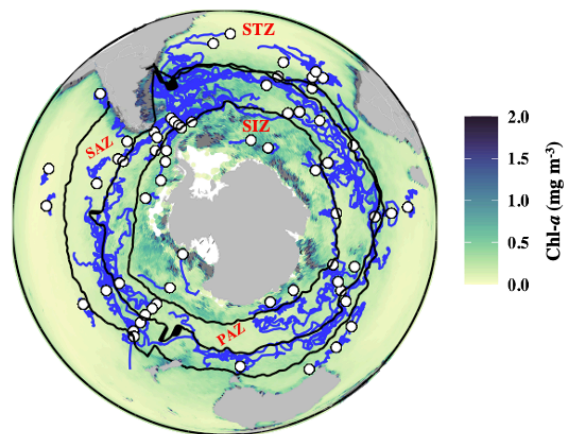


477
 478 **Fig. S10** Time cumulative air-sea CO₂ gas flux for different productivity scenarios during the (a)
 479 unproductive period and (b) the full year based on the different model settings. Gold lines show
 480 the observed CO₂ fluxes. Frontal zones as described in **Fig. S1**. Positive values indicate a source
 481 of carbon to the atmosphere. Error bars and shading represent the propagated uncertainty in each
 482 scenario. Model 1 (adopted in the main text): reconstruction results the abiotic and TOM
 483 production only scenarios in which the unproductive season contributes to the total annual
 484 production (*Materials and Methods* in main text); Model 2: reconstruction results the abiotic and
 485 TOM production only scenarios in which the unproductive season does not contribute to the total
 486 annual production.
 487
 488



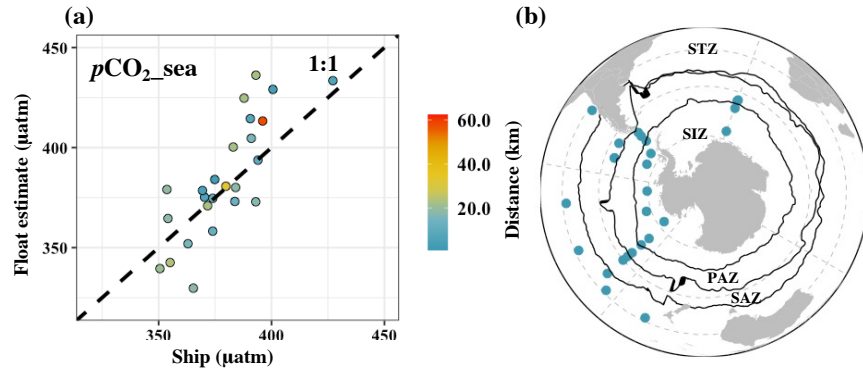
489
490
491
492
493
494
495
496
497
498
499
500
501
502

Fig. S11 Schematic of Southern Ocean carbon cycling during the productive season. Contributions of distinct biogenic carbon pools to the total biological carbon production and the associated impacts on air-sea CO₂ flux. Arrow sizes are proportional to the area-weighted, cumulative carbon flux magnitude within each frontal zone. Inset bar plots show the percentage contributions of each biogenic carbon pool to the export potential.



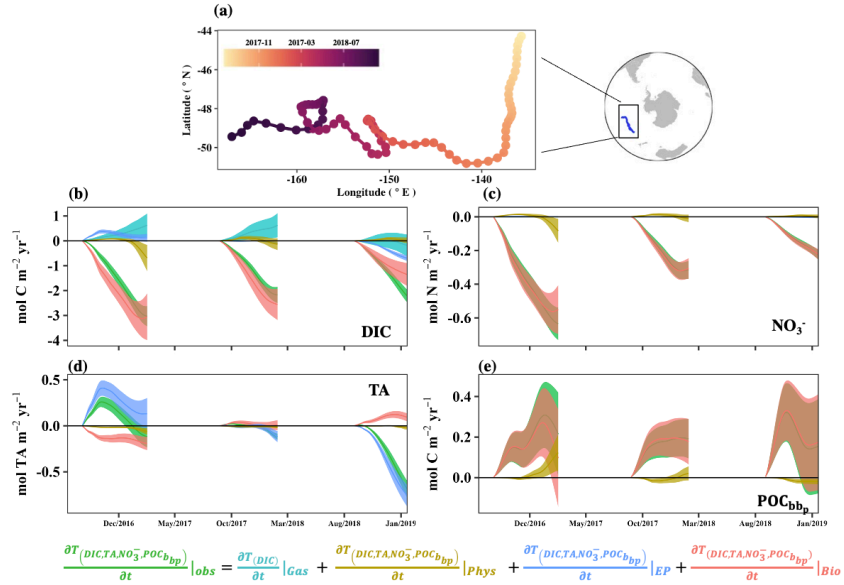
503 **Fig. S12** Trajectories of the floats used in this study. White dots show the deployment locations.
 504 The background color shows a climatology of remotely sensed surface chlorophyll-*a* during austral
 505 summer. Black lines show the climatological boundaries (mean of 2004–2014) of the Southern
 506 Ocean fronts determined using an Argo-based climatology of temperature and salinity (32). Zones
 507 as described in **Fig. S1**.
 508

509
 510
 511
 512
 513
 514
 515
 516
 517
 518
 519
 520
 521
 522
 523
 524
 525
 526



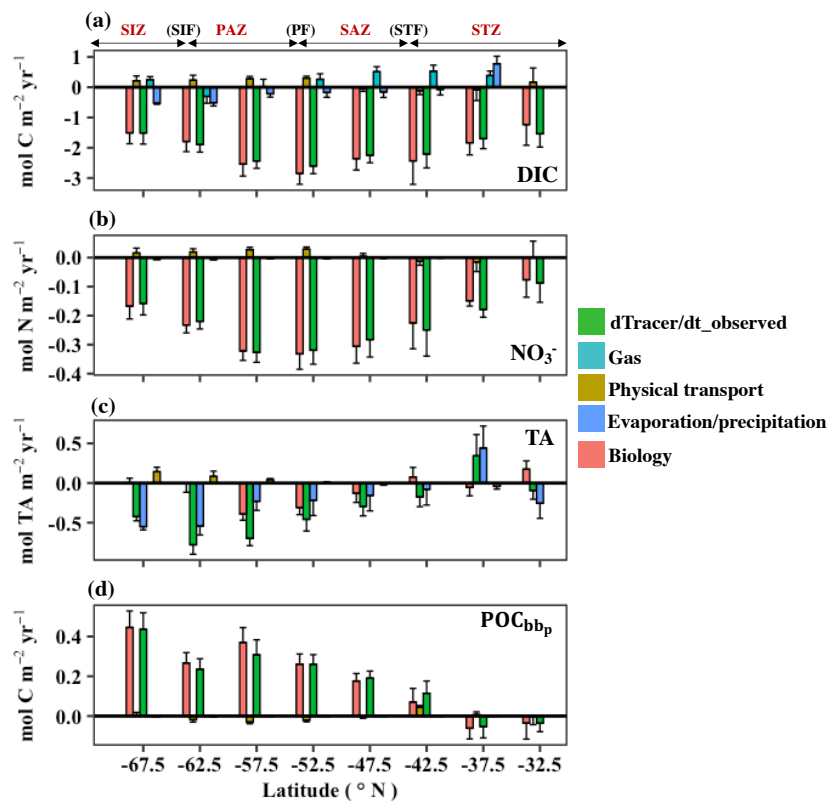
527
 528 **Fig. S13 (a)** Comparison of the sea surface partial pressure of carbon dioxide ($p\text{CO}_{2_sea}$) from float
 529 estimates and underway shipboard observations compiled in SOCAT (version 2022 (33)). **(b)** The
 530 float and ship samples were paired based on location ($\pm 0.2^\circ$) and sampling date (± 2 days). The
 531 color bar in **a** reflects the distance between ship and float samples. Frontal zones in panel **b** as
 532 described in **Fig. S1**.

533
 534
 535
 536
 537
 538
 539
 540
 541
 542
 543
 544
 545
 546
 547
 548

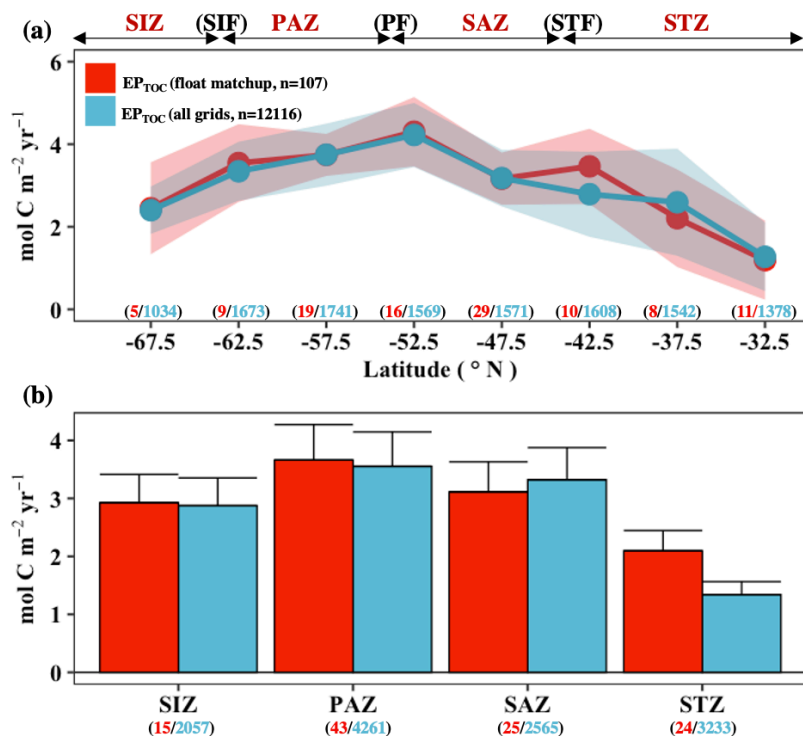


549
 550
 551
 552
 553
 554
 555
 556
 557
 558

Fig. S14 Time cumulative fluxes in each productive season within the euphotic zone (~77m) for a representative biogeochemical float (WMO ID: 5904761). Shading represents the propagated error. Positive values indicate the process causes an increase in the tracer inventory. DIC: dissolved inorganic carbon; NO₃⁻: nitrate; TA: total alkalinity; POC_{bbp}: particulate organic carbon derived from the backscattering coefficient.

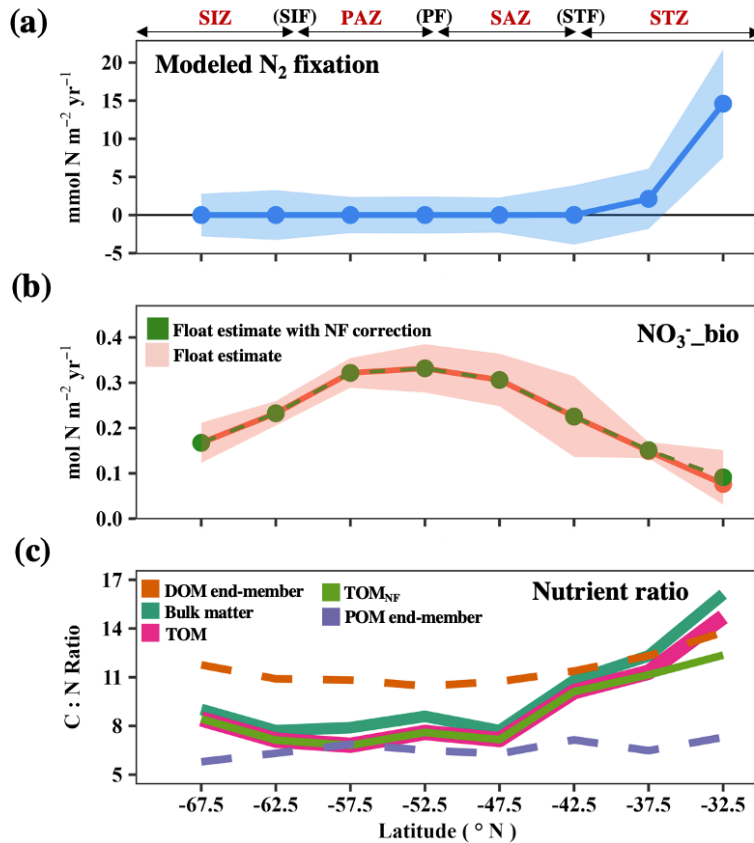


559
 560 **Fig. S15** Meridional pattern (5° bin) of tracer budget integrated over the euphotic zone during the
 561 Southern Ocean productive season. Error bars represent the propagated error. The positive value
 562 indicates the process leads to an increase in the tracer inventory. DIC: dissolved inorganic carbon;
 563 NO₃⁻: nitrate; TA: total alkalinity; POC_{bbp} : particulate organic carbon derived from the
 564 backscattering coefficient. Zones as described in **Fig. S1**.
 565



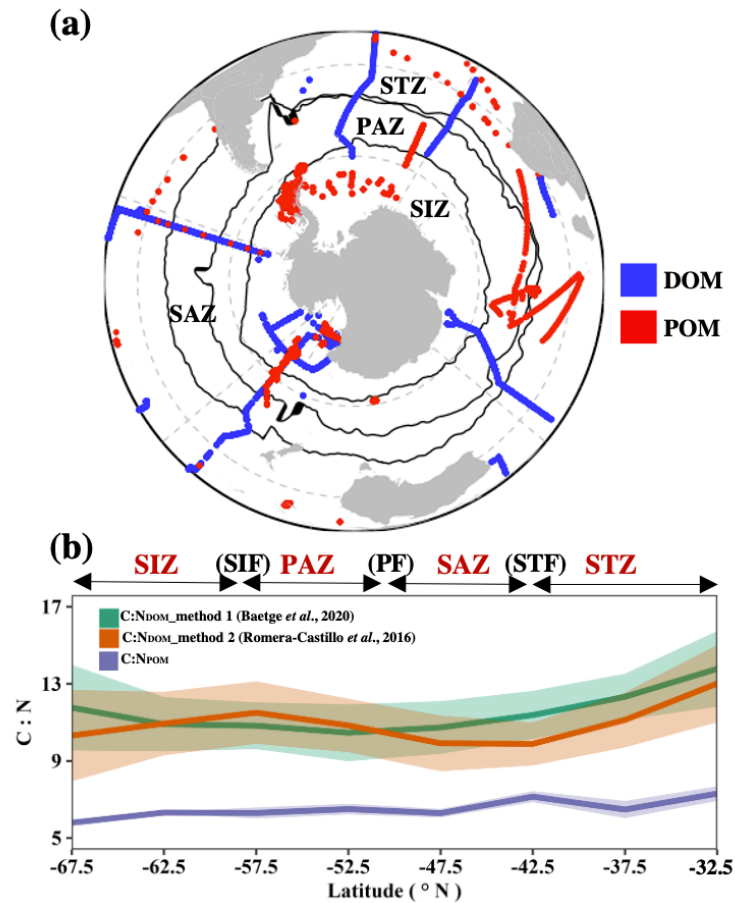
566
 567
 568
 569
 570
 571
 572
 573
 574
 575
 576
 577

Fig. S16 Evaluation, using data constrained biogeochemical Southern Ocean State Estimate (B-SOSE) model output (31), of how representative the float-based productivity estimates are of the entire Southern Ocean. Comparison of subsampled (along float trajectories) versus fully resolved (1°×1°) model fields of euphotic zone integrated export potential of total organic carbon (E_{TOC}) during the productive season. Results are shown as an average (a) across meridional bands and (b) the four main frontal regions. Shading and error bars reflect spatial variability in EP_{TOC} within each sub-region. Red and blue numbers near the bottom of each panel show the number of grid points used in each sub-region for computing the average value. The frontal regions are as described in **Fig. S1**.



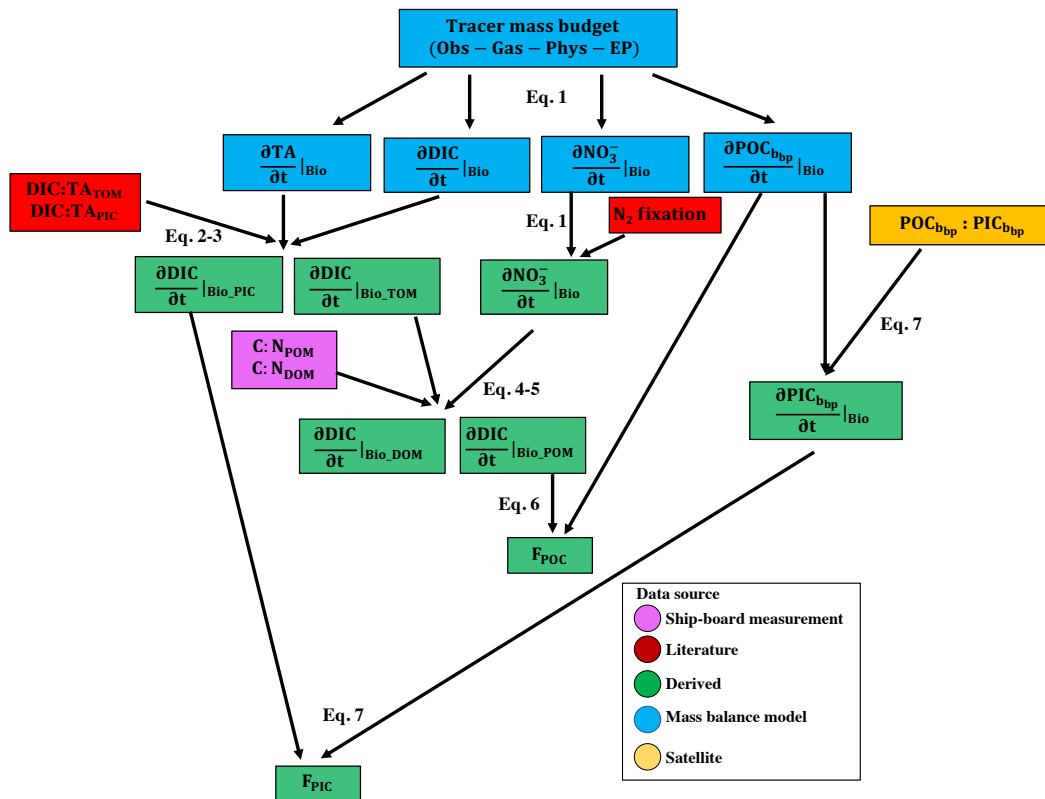
578
 579 **Fig. S17** The meridional pattern of (a) climatology of nitrogen fixation (NF) derived from a data-
 580 driven inverse model (34), and (b) net biological term solved from nitrate tracer budget ($\text{NO}_3^-_{\text{bio}}$),
 581 (c) carbon-to-nitrogen ratios (C:N) for biological production of different carbon pools during the
 582 Southern Ocean productive season. The dissolved organic matter (DOM) and particulate organic
 583 matter (POM) end-member C:N ratios are derived from the compilation of existing ship-based
 584 observations. The bulk biological production, total organic matter (TOM) production, and TOM
 585 production with the correction for nitrogen fixation (TOM_{NF}) are derived from float observations
 586 and a biogeochemical inverse model. Shading on each line reflects the propagated error. Zones as
 587 described in Fig. S1.

588
 589
 590
 591
 592



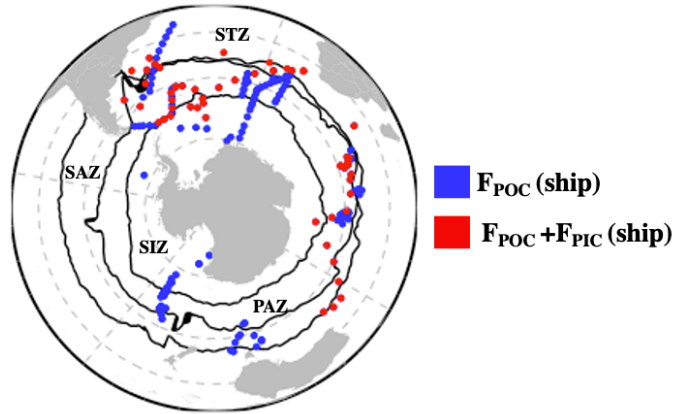
593
 594
 595
 596
 597
 598
 599
 600

Fig. S18 (a) Distribution of discrete particulate organic matter (POM) and dissolved organic matter (DOM) observations and the (b) meridional patterns of end-member C:N ratios for POM ($C:N_{POM}$) and DOM ($C:N_{DOM}$) produced during the Southern Ocean productive season. Shading represents the propagated error. Zones as described in Fig. S1. $C:N_{DOM_method\ 1}$ and $C:N_{DOM_method\ 2}$ represent the results from two different approaches described in Text S1.1.



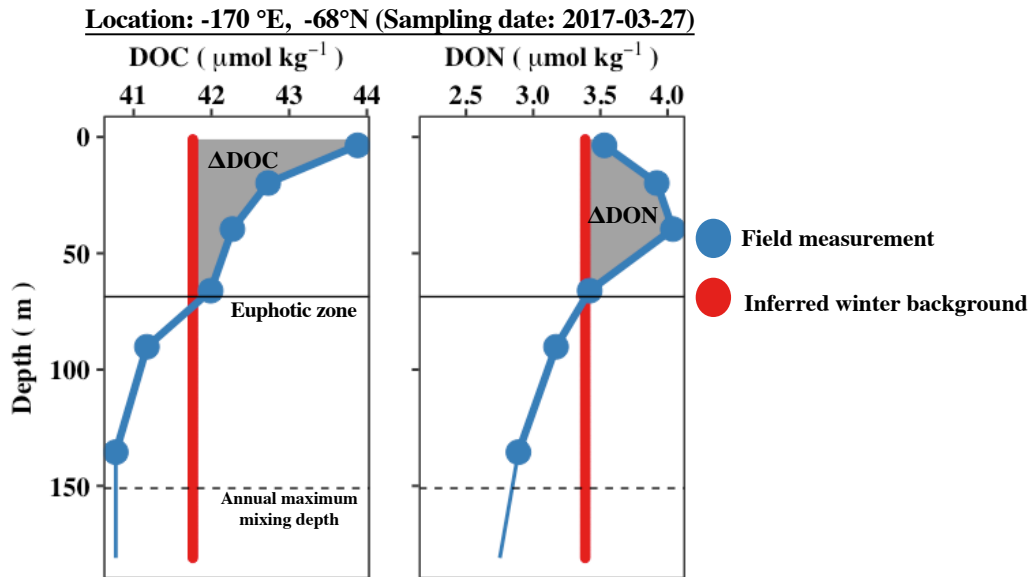
601
 602
 603
 604
 605
 606
 607
 608
 609
 610
 611
 612
 613
 614
 615

Fig. S19 A schematic to depict the workflow and data sources for multiple terms applied in distinct carbon pools partitioning. See detailed descriptions for each term in the method section.



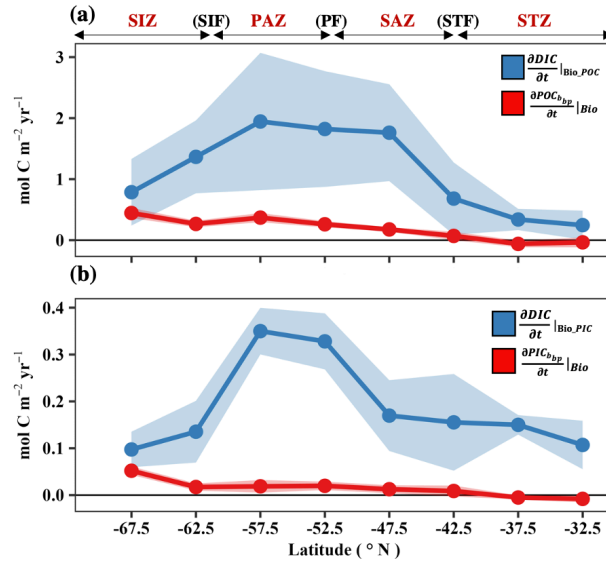
616
 617 **Fig. S20** Distribution of discrete particulate organic carbon (F_{POC}) and particulate inorganic carbon
 618 (F_{PIC}) sinking fluxes inferred from measurements of ^{234}Th - ^{238}U disequilibrium during the Southern
 619 Ocean productive seasons. See **Text S1.2** for a detailed description of the flux calculation and
 620 integration adjustment. Zones as described in **Fig. S1**.

621
 622
 623
 624
 625
 626
 627
 628
 629
 630
 631
 632
 633
 634
 635



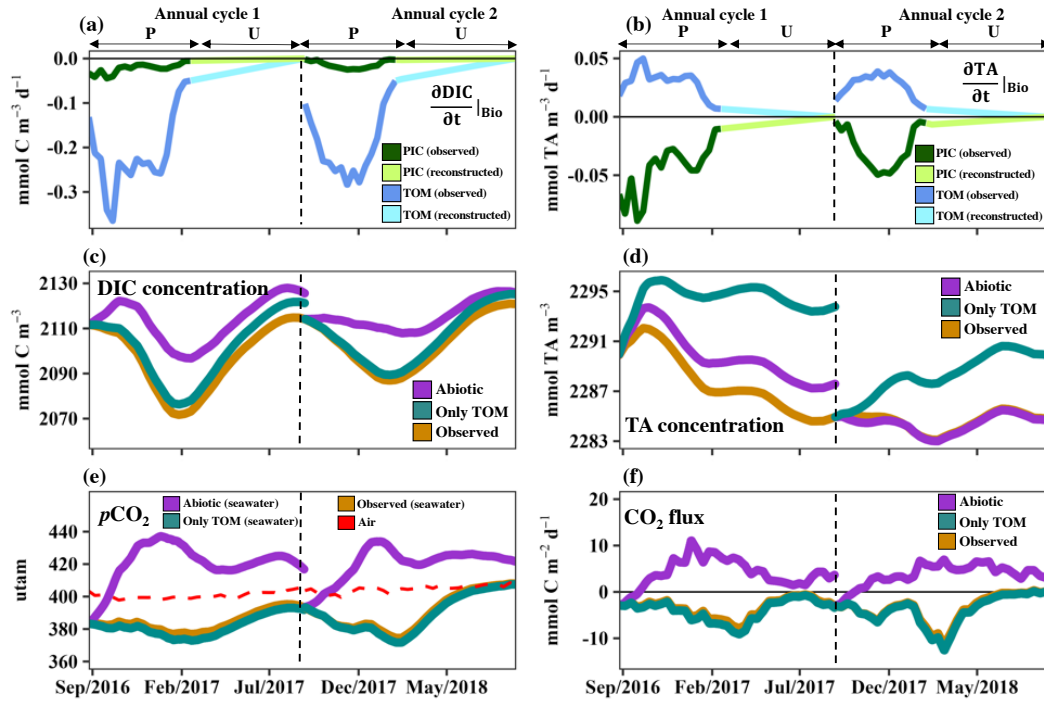
636
 637 **Fig. S21** An example illustrating the reconstruction of seasonal dissolved organic matter (ΔDOM)
 638 production ratios based on profiles of dissolved organic carbon (DOC) and dissolved organic
 639 nitrogen (DON) collected during the stratified summer. Method following Baetge, *et al.* (2).
 640

641
 642
 643
 644
 645
 646
 647
 648
 649
 650
 651



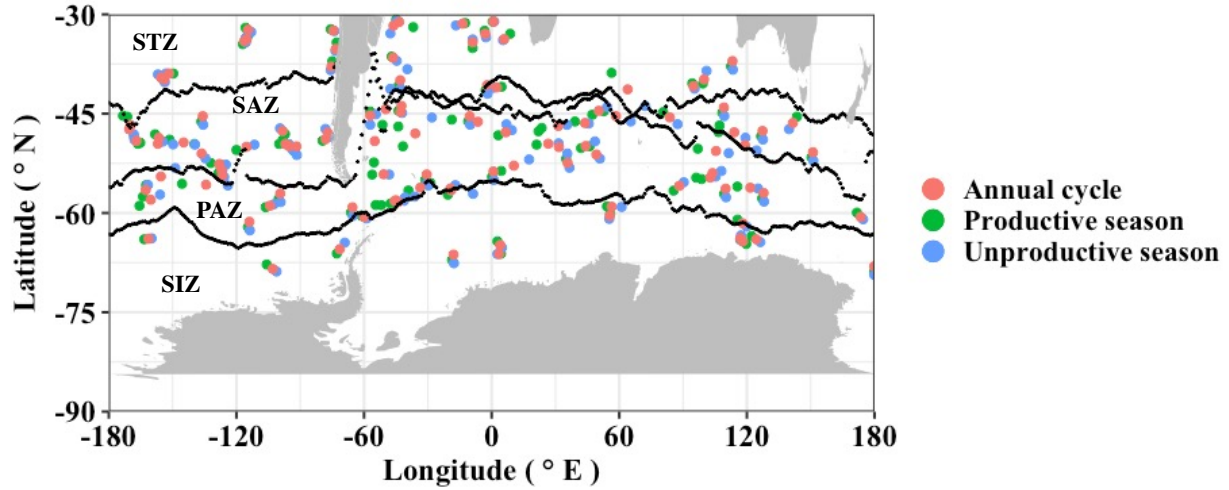
652
 653
 654
 655
 656
 657
 658
 659
 660
 661
 662
 663
 664

Fig. S22 Meridional pattern (5° bin) of euphotic zone net biological terms solved from suspended particles ($\frac{\partial \text{POC}_{\text{bp}}}{\partial t} |_{\text{Bio}}$ and $\frac{\partial \text{PIC}_{\text{bp}}}{\partial t} |_{\text{Bio}}$) and the DIC tracer budget ($\frac{\partial \text{DIC}}{\partial t} |_{\text{Bio_POC}}$ and $\frac{\partial \text{DIC}}{\partial t} |_{\text{Bio_PIC}}$) during the Southern Ocean productive season. Shading reflects the propagated error. Zones as described in **Fig. S1**. POC: particulate organic carbon; PIC: particulate inorganic carbon.



665
 666 **Fig. S23** Time-series of (a) biologically-induced dissolved inorganic carbon (DIC) change, (b)
 667 biologically-induced total alkalinity (TA) change, (c) DIC concentration, (d) TA concentration, (e)
 668 partial pressure of carbon dioxide ($p\text{CO}_2$), and (f) air-sea CO_2 gas flux over two annual cycles for
 669 a representative biogeochemical float (WMO ID: 5904761) under three different productivity
 670 scenarios: observed, abiotic, and total organic matter production only. Reconstructions of
 671 productivity during the unproductive (U) period are based on the fraction of total production
 672 occurring during the productive (P) period in B-SOSE model output (*Materials and Methods* in
 673 main text).

674
 675
 676
 677
 678
 679
 680
 681
 682
 683



684
 685
 686
 687
 688
 689
 690
 691
 692
 693
 694
 695
 696
 697
 698
 699
 700
 701
 702
 703
 704
 705
 706

Fig. S24 Mean location of each float during each annual cycle, productive season, or unproductive season. Black lines show the climatological locations (mean of 2004–2014) of Southern Ocean fronts determined using an Argo-based climatology of temperature and salinity (32). Zones as described in **Fig. S1**.

707 **Table S1.** List of World Meteorological Organization identification numbers (WMO IDs) for the
708 biogeochemical floats used in this study.
709

No.	WMO ID	No.	WMO ID	No.	WMO ID	No.	WMO ID
1	5904188	21	5904686	41	5905135	61	5906244
2	5904395	22	5904693	42	5905981	62	5906245
3	5904396	23	5904695	43	5905982	63	5906250
4	5904468	24	5904761	44	5905995		
5	5904469	25	5904765	45	5905996		
6	5904470	26	5904766	46	5905997		
7	5904473	27	5904768	47	5905998		
8	5904657	28	5904842	48	5906003		
9	5904658	29	5904844	49	5906007		
10	5904659	30	5904854	50	5906031		
11	5904660	31	5904856	51	5906032		
12	5904661	32	5904857	52	5906204		
13	5904662	33	5904980	53	5906208		
14	5904663	34	5904982	54	5906209		
15	5904673	35	5905075	55	5906211		
16	5904675	36	5905105	56	5906216		
17	5904679	37	5905109	57	5906218		
18	5904682	38	5905131	58	5906222		
19	5904683	39	5905132	59	5906224		
20	5904684	40	5905134	60	5906227		

710
711
712
713
714
715
716
717
718
719
720
721
722
723
724
725
726
727

728 **Table S2** Assignment of uncertainties used in the Monte Carlo error analysis. NO₃⁻: nitrate; TA:
 729 total alkalinity; POC_{bbp}: particulate organic carbon based on backscattering coefficient; Chl-*a*:
 730 Chlorophyll-*a* concentration based on Chlorophyll-fluorescence; CO₂: carbon dioxide; MLD:
 731 mixed layer depth; POM: particulate organic matter; DOM: dissolved organic matter.
 732

Variable	Uncertainty	Reference
pH measurement	±0.005	Johnson, <i>et al.</i> (35)
NO ₃ ⁻ measurement	±0.5 μmol kg ⁻¹	Johnson, <i>et al.</i> (20)
TA estimates	± 4.5 μmol kg ⁻¹	Bittig, <i>et al.</i> (11)
POC _{bbp} estimates	±30%	Graff, <i>et al.</i> (36)
Chl- <i>a</i> estimates	20%	Johnson, <i>et al.</i> (20)
CO ₂ gas model	±30%	Bender, <i>et al.</i> (25), Wanninkhof (26)
Background eddy diffusivity coefficient	±35%	Yang, <i>et al.</i> (23)
Ekman pumping velocity	±50%	Haskell, <i>et al.</i> (37)
Thickness of MLD	±0.5 m	Vertical resolution/4
Thickness of euphotic zone	5-15 m	Spatial variability at each latitude band (Fig. S1a)
C:N ratio for POM/DOM production	0.5-2.5	Spatial variability at each latitude band (Fig. S18a)
Model-simulated N ₂ fixation	10-30%	Spatial variability at each latitude band (Fig. S17a)

733

Table S3 Summary of ancillary parameters used in this study.

Parameters	Description	Reference /Link	Usage
Parameters used to partition carbon pools and in tracer budget closure			
Ice coverage	Monthly surface ice coverage; 25 km resolution	Global Sea Ice Concentration Climate Data (https://cds.climate.copernicus.eu/cdsapp#!/dataset/satellite-sea-ice-concentration?tab=overview)	Scale the air-sea CO ₂ exchange by ice fraction
Euphotic zone depth (Zeu)	Remote-sensed daily surface property. MODIS; 0.83°×0.083°; Level 3	Ocean color (https://oceandata.sci.gsfc.nasa.gov/directdataaccess/Level-3%20Mapped/Aqua-MODIS)	Determine the Zeu
Wind stress	Daily surface wind stress; 0.25° × 0.25°	Advanced Scatterometer product (https://manati.star.nesdis.noaa.gov/datasets/ASCATData.php)	Calculate the Ekman pumping velocity
Wind speed	Daily values; 0.16°×0.16°	NCEP-DOE reanalysis: II (https://psl.noaa.gov/data/gridded/data.ncep.reanalysis2.html)	Calculate the atmospheric pCO ₂ and air-sea CO ₂ fluxes
Air pressure			
Relative humidity			
N ₂ fixation	Monthly climatology simulated from inverse biogeochemical and prognostic ocean models; 1°×1°; 24 depth interval	Wang, <i>et al.</i> (34)	Correct NO ₃ ⁻ potential export
Particulate inorganic carbon (PIC)	Daily surface properties; 0.83°×0.083°; Level 3	Ocean color (https://oceandata.sci.gsfc.nasa.gov/directdataaccess/Level-3%20Mapped/Aqua-MODIS)	Derive the PIC:POC ratio to estimate seawater PIC inventory
Particulate organic carbon (POC)			
Dissolved organic matter (DOM)	Global compilation of dissolved organic carbon (DOC) and dissolved organic nitrogen (DON=total dissolved nitrogen – total inorganic nitrogen) observations	Hansell <i>et al.</i> , (2021) (https://www.ncei.noaa.gov/access/metadata/landing-page/bin/iso?id=gov.noaa.nodc:0227166)	Constrain the end-member C:N ratio of DOM and POM produced
Particulate organic matter (POM)	Global compilation of suspended particulate organic carbon (POC) and particulate organic nitrogen (PON) observations	Martiny, <i>et al.</i> (38) (https://www.bco-dmo.org/dataset/526747)	
Parameters used to validate or compare with float analysis			
Particulate organic/inorganic carbon sinking flux (F _{POC} /F _{PIC})	Compilation of POC and PIC sinking fluxes inferred from ²²³⁴ Th- ²³⁸ U	Henson, <i>et al.</i> (5), Le Moigne, <i>et al.</i> (6), Rosengard, <i>et al.</i> (7)	Compare with the float estimate
Export potential of organic carbon	Monthly climatology (2013-2021); 0.16°×0.16°	Biogeochemical Southern Ocean State Estimate (B-SOSE) (http://sose.ucsd.edu/) (31)	Evaluate the representativeness of the spatial variability of carbon export derived from the float locations
Parameters used to analyze light and nutrient availability			

Dissolved iron (dFe) concentration	Monthly climatology simulated from a data-driven model; 1°×1°; 33 depth interval	Huang, <i>et al.</i> (12) (https://zenodo.org/record/6994318#.Y0m2fuzMLjk)	Analyze the meridional pattern of dFe
SiO ₄ ⁻ concentration	Estimated from float temperature, salinity, and dissolved oxygen observations	Canyon-B algorithm (11)	Analyze the meridional pattern of SiO ₄ ⁻ dependent growth rate
Photosynthetically available radiation (PAR)	Remote-sensed daily surface properties; MODIS; 083°×0.083°; Level 3	Ocean color (https://oceandata.sci.gsfc.nasa.gov/directdataaccess/Level-3%20Mapped/Aqua-MODIS)	Analyze the meridional pattern of light availability

736
737
738
739
740
741
742
743
744
745
746
747
748
749
750
751
752
753
754
755
756
757
758
759
760
761
762
763
764
765
766
767
768
769
770
771
772

773 **References**

774

- 775 1. D. A. Hansell, M. V. Orellana, Dissolved organic matter in the global ocean: A primer.
776 *Gels* **7** (2021).
- 777 2. N. Baetge, J. R. Graff, M. J. Behrenfeld, C. A. Carlson, Net community production,
778 dissolved organic carbon accumulation, and vertical export in the western North Atlantic.
779 *Frontiers in Marine Science* **7** (2020).
- 780 3. J. Holte, L. D. Talley, J. Gilson, D. Roemmich, An Argo mixed layer climatology and
781 database. *Geophysical Research Letters* **44**, 5618-5626 (2017).
- 782 4. C. Romera-Castillo, R. T. Letscher, D. A. Hansell, New nutrients exert fundamental control
783 on dissolved organic carbon accumulation in the surface Atlantic Ocean. *Proceedings of*
784 *the National Academy of Sciences* **113**, 10497-10502 (2016).
- 785 5. S. Henson, F. Le Moigne, S. Giering, Drivers of carbon export efficiency in the global
786 ocean. *Global Biogeochem Cycles* **33**, 891-903 (2019).
- 787 6. F. A. Le Moigne, K. Pabortsava, C. L. Marcinko, P. Martin, R. J. Sanders, Where is mineral
788 ballast important for surface export of particulate organic carbon in the ocean? *Geophys*
789 *Res Lett* **41**, 8460-8468 (2014).
- 790 7. S. Z. Rosengard *et al.*, Carbon export and transfer to depth across the Southern Ocean Great
791 Calcite Belt. *Biogeosciences* **12**, 3953-3971 (2015).
- 792 8. J. H. Martin, G. A. Knauer, D. M. Karl, W. W. Broenkow, VERTEX: carbon cycling in
793 the northeast Pacific. *Deep-Sea Res* **34**, 267-285 (1987).
- 794 9. W. M. Balch *et al.*, Factors regulating the Great Calcite Belt in the Southern Ocean and its
795 biogeochemical significance. *Global Biogeochemical Cycles* **30**, 1124-1144 (2016).
- 796 10. I. Salter *et al.*, Carbonate counter pump stimulated by natural iron fertilization in the Polar
797 Frontal Zone. *Nature Geoscience* **7**, 885-889 (2014).
- 798 11. H. C. Bittig *et al.*, An alternative to static climatologies: robust estimation of open ocean
799 CO₂ variables and nutrient concentrations from T, S, and O₂ data using Bayesian Neural
800 Networks. *Frontiers in Marine Science* **5** (2018).
- 801 12. Y. Huang, A. Tagliabue, N. Cassar, Data-driven modeling of dissolved iron in the global
802 ocean. *Frontiers in Marine Science* **9** (2022).
- 803 13. A. R. Gray *et al.*, Autonomous biogeochemical floats detect significant carbon dioxide
804 outgassing in the high-latitude Southern Ocean. *Geophysical Research Letters* **45**, 9049-
805 9057 (2018).
- 806 14. S. M. Bushinsky *et al.*, Reassessing Southern Ocean air-Sea CO₂ flux estimates with the
807 addition of biogeochemical float observations. *Global Biogeochem Cycles* **33**, 1370-1388
808 (2019).
- 809 15. S. Stammerjohn *et al.*, Antarctica and the Southern Ocean. *Bulletin of the American*
810 *Meteorological Society* **102**, S317-S356 (2021).
- 811 16. P. Landschützer, N. Gruber, D. C. E. Bakker, Decadal variations and trends of the global
812 ocean carbon sink. *Global Biogeochemical Cycles* **30**, 1396-1417 (2016).
- 813 17. P. Landschützer, N. Gruber, D. C. E. Bakker, U. Schuster, Recent variability of the global
814 ocean carbon sink. *Global Biogeochemical Cycles* **28**, 927-949 (2014).
- 815 18. C. J. Prend *et al.*, Indo-Pacific Sector Dominates Southern Ocean Carbon Outgassing.
816 *Global Biogeochemical Cycles* **36** (2022).
- 817 19. M. C. Long *et al.*, Strong Southern Ocean carbon uptake evident in airborne observations.
818 *Science* **374**, 1275-1280 (2021).

- 819 20. K. S. Johnson *et al.*, Biogeochemical sensor performance in the SOCCOM profiling float
820 array. *Journal of Geophysical Research: Oceans* **122**, 6416-6436 (2017).
- 821 21. C. de Boyer Montégut, G. Madec, A. S. Fischer, A. Lazar, D. Iudicone, Mixed layer depth
822 over the global ocean: an examination of profile data and a profile-based climatology.
823 *Journal of Geophysical Research-Oceans* **109**, C12003 (2004).
- 824 22. S. R. Signorini, C. R. McClain, J. R. Christian, C. S. Wong, Seasonal and interannual
825 variability of phytoplankton, nutrients, TCO₂, pCO₂, and O₂ in the eastern subarctic Pacific
826 (ocean weather station Papa). *Journal of Geophysical Research: Oceans* **106**, 31197-31215
827 (2001).
- 828 23. B. Yang, S. R. Emerson, S. M. Bushinsky, Annual net community production in the
829 subtropical Pacific Ocean from *in-situ* oxygen measurements on profiling floats. *Global*
830 *Biogeochemical Cycles* **31**, 728–744 (2017).
- 831 24. O. M. Sun, S. R. Jayne, K. L. Polzin, B. A. Rahter, L. C. St. Laurent, Scaling turbulent
832 dissipation in the transition layer. *Journal of Physical Oceanography* **43**, 2475-2489 (2013).
- 833 25. M. L. Bender, S. Kinter, N. Cassar, R. Wanninkhof, Evaluating gas transfer velocity
834 parameterizations using upper ocean radon distributions. *Journal of Geophysical Research:*
835 *Oceans* **116** (2011).
- 836 26. R. Wanninkhof, Relationship between wind speed and gas exchange over the ocean
837 revisited. *Limnology and Oceanography: Methods* **12**, 351-362 (2014).
- 838 27. B. R. Carter, J. A. Radich, H. L. Doyle, A. G. Dickson, An automated system for
839 spectrophotometric seawater pH measurements. *Limnology and Oceanography: Methods*
840 **11**, 16-27 (2013).
- 841 28. N. L. Williams *et al.*, Calculating surface ocean pCO₂ from biogeochemical Argo floats
842 equipped with pH: An uncertainty analysis. *Global Biogeochemical Cycles* **31**, 591-604
843 (2017).
- 844 29. B. R. Carter *et al.*, Updated methods for global locally interpolated estimation of alkalinity,
845 pH, and nitrate. *Limnology and Oceanography: Methods* **16**, 119-131 (2018).
- 846 30. A. Morel *et al.*, Examining the consistency of products derived from various ocean color
847 sensors in open ocean (Case 1) waters in the perspective of a multi-sensor approach.
848 *Remote Sensing of Environment* **111**, 69-88 (2007).
- 849 31. A. Verdy, M. R. Mazloff, A data assimilating model for estimating Southern Ocean
850 biogeochemistry. *Journal of Geophysical Research: Oceans* **122**, 6968-6988 (2017).
- 851 32. D. Roemmich, J. Gilson, The 2004–2008 mean and annual cycle of temperature, salinity,
852 and steric height in the global ocean from the Argo Program. *Progress in Oceanography*
853 **82**, 81-100 (2009).
- 854 33. D. C. E. Bakker *et al.*, A multi-decade record of high-quality fCO₂ data in version 3 of the
855 Surface Ocean CO₂ Atlas (SOCAT). *Earth System Science Data* **8**, 383-413 (2016).
- 856 34. W. L. Wang, J. K. Moore, A. C. Martiny, F. W. Primeau, Convergent estimates of marine
857 nitrogen fixation. *Nature* **566**, 205-211 (2019).
- 858 35. K. S. Johnson *et al.*, Deep-Sea DuraFET: A pressure tolerant pH sensor designed for global
859 sensor networks. *Analytical chemistry* **88**, 3249-3256 (2016).
- 860 36. J. R. Graff *et al.*, Analytical phytoplankton carbon measurements spanning diverse
861 ecosystems. *Deep Sea Research Part I: Oceanographic Research Papers* **102**, 16-25
862 (2015).

- 863 37. W. Z. Haskell, A. J. Fassbender, J. S. Long, J. N. Plant, Annual net community production
864 of particulate and dissolved organic carbon from a decade of biogeochemical profiling float
865 observations in the Northeast Pacific. *Global Biogeochemical Cycles* **34** (2020).
- 866 38. A. C. Martiny, J. A. Vrugt, M. W. Lomas, Concentrations and ratios of particulate organic
867 carbon, nitrogen, and phosphorus in the global ocean. *Sci Data* **1**, 140048 (2014).
868