Bridging the gaps between particulate backscattering measurements and modeled particulate organic carbon in the ocean

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Abstract. Oceanic particulate organic carbon (POC) is a small but dynamic component of the global carbon cycle. Biogeochemical models historically focused on reproducing the sinking flux of POC driven by large fast-sinking particles (LPOC). However, suspended and slow-sinking particles (SPOC, here <100 µm) dominate the total POC (TPOC) stock, support a large fraction of microbial respiration, and can make sizable contributions to vertical fluxes.

Recent developments in the parameterization of POC reactivity in the PISCES model (PISCESv2_RC) have improved its ability to capture POC dynamics. Here we evaluated this model by matching a global 3D simulation and 1D simulations at 50 different locations with observations made from biogeochemical (BGC-) Argo floats and satellites. Our evaluation covers globally representative biomes between 0 and 1000 m depth and relies on: (1) a refined scheme for converting particulate backscattering at 700 nm (b_{bp700}) to POC, based on biome-dependent POC/b_{bp700} ratios in the surface layer that decrease to an asymptotic value at depth; (2) a novel approach for matching annual time series of BGC-Argo vertical profiles to PISCES 1D simulations forced by pre-computed vertical mixing fields; and (3) a critical evaluation of the correspondence between in-situ measurements of POC fractions, PISCES model tracers, and SPOC and LPOC estimated from high vertical resolution b_{bp700} profiles through separation of the baseline and spike signals.

We show that PISCES captures the major features of SPOC and LPOC across a range of spatiotemporal scales, from highly resolved profile time series to biome-aggregated climatological profiles. Model–observations agreement is usually better in the epipelagic (0–200 m) than in the mesopelagic (200–1000 m), with SPOC showing overall higher spatiotemporal correlation and smaller deviation (typically within a factor of 1.5). Still, annual mean LPOC stocks estimated from PISCES and BGC-Argo are highly correlated across biomes, especially in the epipelagic (r = 0.78; n = 50). Estimates of the SPOC/TPOC fraction converge around a median 85% (range 66–92%) globally. Distinct patterns of model-observations misfits are found in subpolar and subtropical gyres, pointing to the need to better resolve the interplay between sinking, remineralization and SPOC–LPOC interconversion in PISCES. Our analysis also indicates that a widely used satellite algorithm overestimates POC by several-fold at high latitudes during the winter. The approaches proposed here can help constrain the stocks, and ultimately budgets, of oceanic POC.

1 Introduction

The biological carbon pump (BCP) is the ensemble of processes that transfer the organic matter produced by plankton in the sunlit ocean surface to deeper layers (Volk and Hoffert, 1985). This vertical flux plays a central role in the Earth's climate, as it influences the oceans' capacity to absorb and ultimately store atmospheric CO_2 over centennial or millennial time scales (Kwon et al., 2009; Passow and Carlson, 2012). The BCP is also central to biogeochemical functioning of the ocean, as it determines the quality and quantity of organic matter available to the ocean interior (Arístegui et al., 2009; Hernández-León et al., 2020) and the seafloor ecosystems. The spatiotemporal patterns of organic matter supply and respiration influence the distribution of dissolved oxygen, inorganic carbon, remineralized nutrients and trace metals in the ocean interior (Bianchi et al., 2018; Duteil et al., 2012; Hayes et al., 2015, Oschlies et al., 2018; Weber et al., 2016), and their return pathways to the surface. In consequence, the BCP is intimately linked to, and feeds back on, upper ocean productivity.

Over the last decades, BCP research has placed emphasis on understanding the ecological and physical factors that control the gravitational export of particulate organic carbon (POC; see Table 1 for acronym definitions). This process is often represented as the product of export production (the fraction of net primary production exported below the euphotic layer) and transfer efficiency (Teff; the fraction of export production that reaches a given depth below the euphotic layer). Both variables vary widely across ocean biomes and along the seasonal cycle (Buesseler and Boyd, 2009; Passow and Carlson, 2012; Buesseler et al., 2020), and our understanding of the underlying mechanisms is still limited by the relatively small amount of in-situ measurements (Mouw et al. 2016). Biogeochemical models have been built, and their parameters tuned, to be able to reproduce these sparse observations of export production and vertical flux attenuation, sometimes reaching apparently contradictory results (Marsay et al., 2015). In comparison, the models' ability to represent marine particle concentrations has received less attention (Lam et al., 2011).

Marine particles are mainly composed of living microbial plankton cells, living metazoans and detritus ("marine snow" aggregates, fecal pellets, zooplankton feeding structures, biominerals), whose size ranges from <1 µm to several mm (Bishop et al., 1980; Mullin et al., 1965; Stemmann and Boss, 2012). Indeed, these particles feature wide variations in their physicochemical properties and degree of biological processing (Kharbush et al., 2020; Lam et al., 2015; Passow and Carlson, 2012; Stemmann and Boss, 2012), as well as microbial colonization (Baumas et al., 2021; Duret et al., 2019; Mestre et al., 2018), all of which change during the particles' lifetime. Gravitational sinking speed generally increases with particle size, although observations show a wide scatter around canonical Stokes' law predictions (Cael et al., 2021; Laurenceau-Cornec et al., 2019). Owing to this general relationship, particle populations are often partitioned into a few functional size classes: large particles, typically defined as larger than 50 or 100 µm, which usually sink at several tens or hundreds of m d^{-1}; and small

particles, which usually sink slowly (<10 m d^{-1}) or are suspended in the water column. In this study, POC is divided into small POC (SPOC) and large POC (LPOC), with a nominal cutoff at 100 µm (Table 2; section 2.2.2).

The traditional BCP paradigm posits that gravitational sinking of LPOC controls the vertical carbon flux (Sarmiento and Gruber, 2006). However, it has been known for decades that POC export is shaped by additional processes, such as the physical transport of particles by convection and subduction and the "active" particle flux mediated by the vertical migration of metazoans. Recently, these processes were collectively termed "particle injection pumps" (Boyd et al., 2019). In parallel, the role of dissolved organic carbon in vertical carbon export has been widely recognized (Jiao et al., 2010; Passow and Carlson, 2012; Legendre et al., 2015). Therefore, the BCP is increasingly seen as a diverse array of interconnected mechanisms.

One aspect that has recently received considerable attention is the role of suspended and slow-sinking particles (Alonso-González et al., 2010; Baker et al., 2017). Owing to their longer residence time, small particles usually dominate the POC stock (Aumont et al., 2017; Baker et al., 2017) and may support a proportional fraction of the respiration (Baltar et al., 2010a, 2010b; Belcher et al., 2016; García-Martín et al., 2021). Convective mixing (Bishop et al., 1986; Dall'Olmo and Mork, 2014; Lacour et al., 2019) and subduction (Llort et al., 2018; Omand et al., 2015; Resplandy et al., 2019) can transport SPOC into the mesopelagic layer, adding to other export mechanisms and potentially making large contributions to total POC export (Alonso-Gonzalez et al., 2010; Henson et al., 2015). Production of SPOC in the mesopelagic and below also results from the fragmentation of LPOC, caused by physical disaggregation (Takeuchi et al., 2019), bacterial solubilization and zooplankton activity (Briggs et al., 2020; Goldthwait et al., 2004; Mayor et al., 2020; Stemmann et al., 2004b). Moreover, SPOC is also produced through bacterial chemosynthesis in the dark ocean (Arístegui et al., 2009; Herndl and Reinthaler, 2014). Altogether, these findings illustrate how our limited knowledge of POC characteristics and cycling hampers mechanistic understanding of the BCP and mesopelagic carbon budgets (Giering et al., 2014).

Biogeochemical models designed to capture only gravitational POC sinking fail to represent POC stocks in the ocean interior. Aumont et al. (2017) recently showed that the Pelagic Interactions Scheme for Carbon and Ecosystem Studies model (PISCESv2; Aumont et al., 2015) underestimated POC by one order-of-magnitude or more below the epipelagic layer. This pitfall is likely common to any state-of-the-art model with similar structure (Laufkötter et al., 2016; Séférian et al., 2020). Aumont's work also showed that the model's fit to observed deep-ocean POC concentrations could be dramatically improved by treating detrital POC, both small and large, as a mixture of particles with different reactivity (or lability) towards bacterial degradation. In this scheme, termed the reactivity continuum (RC) parameterization, detrital POC degradation is computing after dividing it into many reactivity classes that approximately follow a continuous gamma distribution –hence its name. The most labile fractions are rapidly consumed below the upper mixed layer, such that vertically exported POC becomes progressively more refractory. This results in enhanced preservation of SPOC in the model and a much more realistic fraction

of SPOC with respect to total POC (TPOC) in the ocean interior. In addition, the RC scheme does not appreciably degrade model estimates of the gravitational POC flux.

100 Despite this breakthrough in the representation of POC fractions in PISCESv2_RC (hereafter "PISCES"), the new parameterization was evaluated using only sparse measurements (Druffel et al., 1992; Lam et al., 2011; 2015) based on large-volume filtration with in-situ pumps. This approach enables accurate determination of the mass and composition of the particulate fraction but cannot afford high-frequency sampling over extended spatiotemporal scales (Bishop, 1999; Boss et al., 2015; Gardner et al., 2006). During the last decade, the launching of the biogeochemical Argo (BGC-Argo) program of robotic

105 observations has ceased the historical undersampling of particles in the ocean interior (Claustre et al., 2020). BGC-Argo floats provide vertical profiles of temperature, salinity, bio-optical and chemical variables between 0–1000 m every 1 to 10 days in near-real time and are thus well-suited to study particles (~0.5 µm to ~2 mm in size; Table 2) in the mesopelagic layer, where the strongest POC gradient occurs. The rapidly growing fleet of BGC-Argo floats equipped with bio-optical sensors enables comparison between models and observations at global scales with enhanced spatiotemporal resolution. Unfortunately, BGC-

110 Argo floats measure only a bio-optical proxy of POC, the particulate backscattering coefficient (usually at 700 nm, b_{bp700}) and empirical conversion factors are needed to estimate POC (Bishop and Wood, 2008; Cetinić et al., 2012; Stramski et al., 2008). These conversion factors vary in response to several concurrent processes that alter particle abundance, size distribution, shape, composition, and ultimately optical properties (Boss et al., 2015; Giering et al., 2020).

115 In this study we compare SPOC and LPOC concentrations estimated from BGC-Argo floats to their PISCES-simulated counterparts, as well as satellite-retrieved surface POC concentration. The comparison is enabled by a novel empirical algorithm to convert b_{bp700} to POC. Observations and simulations are matched in 3D (biome-wide climatological scale) and 1D (at defined locations over an annual cycle). These complementary strategies allow us to evaluate the skill of PISCES at simulating POC stocks and fractions in globally representative biomes. We conclude with a list of recommendations to fully

120 exploit the potential of robotic particle observations combined with biogeochemical modelling.

2 Methods

2.1 Definition of vertical and horizontal domains

Studies of the BCP usually decompose the ocean into vertical domains: a surface layer where autotrophic activities dominate, and one or several ocean interior layers where heterotrophic processes dominate. Functional definitions based on light

125 penetration, peak export production, vertical mixing or long-term carbon sequestration are usually the most appropriate ones for process studies (Buesseler and Boyd, 2009; Buesseler et al., 2020; Guidi et al., 2015; Palevsky and Doney, 2018). Because this paper is mainly descriptive and combines observations and simulations, we will refer to layers defined by fixed depths: epipelagic (0–200 m), mesopelagic (200–1000 m) and bathypelagic (1000–4000 m).

130   Over the horizontal dimensions, our comparisons between observations and model results rely on the ocean biomes defined by Fay and McKinley (2014). These authors subdivided each ocean basin (Atlantic, Pacific, Indian, and Southern Ocean) into different biomes based on observed variables, namely: sea surface temperature, spring/summer chlorophyll *a* concentration (Chla), ice fraction, and maximum mixed layer depth (MLD), all on a $1° \times 1°$ grid. This division resulted in 17 regions ascribed to one of the following five biomes: the ice biome, the subpolar seasonally stratified biome, the subtropical seasonally stratified

135   biome, the subtropical permanently stratified biome, and the equatorial biome. The analyses reported herein focus on the following four biomes (Fig. 1): the seasonally stratified North Atlantic subpolar gyre (NASPG); the permanently stratified Atlantic and Pacific subtropical gyres, which were grouped together (STG); the seasonally stratified Southern Ocean (Subantarctic); and the Mediterranean Sea, which was added here owing to the abundance of BGC-Argo data, and represents a seasonally stratified subtropical biome. Fay and McKinley's definition allows biome boundaries to change from one year to

140   another. Here we analysed only data from the core of each biome, defined as the grid cells that never changed classification during the 1998–2010 satellite observation periods.

2.2 BGC-Argo observations

The global dataset acquired by the array of BGC-Argo floats was downloaded on 14 January 2020 from the Global Data Assembly Center hosted by Ifremer (ftp://ftp.ifremer.fr/ifremer/argo/dac/) (Argo, 2000). The selected floats were equipped

145   with a Seabird-Wetlabs ECO-Triplet sensor package including a Chla fluorometer (excitation at 470 nm; emission at 695 nm) and a backscattering sensor at 700 nm (b_{bp700}), in addition to the conductivity-temperature-depth (CTD) probe. The downloaded measurements had undergone the standard processing, which includes the application of calibration equations to raw sensor output and the performance of near-real-time quality control to both CTD (Wong et al., 2021) and Chla measurements (Schmechtig et al., 2018). Since no specific quality control procedure has been established yet for b_{bp700} profiles, the latter

150   were only quality-controlled according to the general criteria (Schmechtig et al., 2016). Thus, we used all b_{bp700} measurements with quality control flag ≤ 3 (equivalent results were obtained with flag ≤ 2). Two different processing pipelines were applied to different subsets of the BGC-Argo data, as described below.

2.2.1 Global gridded climatologies (3D approach)

The global dataset acquired between 2010 and 2019 was used to produce global gridded monthly and seasonal climatologies

155   for b_{bp700} and Chla. The measurements were binned onto the ORCA2_L31 grid used for NEMO-PISCES simulations (see 2.4.1), which has a horizontal resolution of about $2°$ that increases to $0.5°$ in the meridional direction in the equatorial domain, and 30 oceanic vertical levels between the surface and the ocean bottom. The thickness of the vertical bins increases progressively from 10 m at the surface to 339 m in the 22nd bin (870–1209 m), the deepest one containing BGC-Argo data. In each grid element, the average, median, range and data counts were computed. Profiles from the CSIRO and INCOIS data

assembly centers were not used because, at the time of download, they had not taken into consideration the new calibration files provided by the manufacturer. A total of 72460 profiles were used to calculate the global gridded climatologies.

2.2.2 Profile time series for individual floats sampling at higher resolution (1D approach)

A subset of the floats, deployed mostly by the projects NAOS, remOcean and Bio-Argo France (model NKE PROVOR CTS-4), were programmed to sample at higher temporal and vertical resolution than the Argo defaults (10 days and 10 m). These floats made vertical profiles between 1,000 m and the surface every 2, 5 or 10 days with a vertical resolution of 10 m between 1,000 and 250 (or 350) m, 1 m between 250 (or 350) and 10 m, and 0.2 m between 10 m and the sea surface. We processed this dataset with a dedicated pipeline to extract additional information on POC size fractions and their dynamics. Along each vertical profile we computed depth, conservative temperature, absolute salinity, σ_θ and spiciness (Flament, 2002) from the calibrated pressure, temperature and salinity using the R package *oce* (Kelley, 2011). The MLD was calculated as the shallowest depth where σ_θ exceeded the surface reference value by 0.03 kg m^{-3} (Bishop and Wood, 2009; Sallée et al., 2021). The surface reference corresponded to the σ_θ at 5 m after applying a 5-point running mean to the top 10 m of the profile. The 0.03 kg m^{-3} criterion provided sensible results across biomes and was consistent with the NEMO-simulated turbocline depth (see 2.5.2). Eleven additional MLD criteria were also calculated to assess the robustness of the approach (Fig. S1).

Following Briggs et al. (2011, 2020), each b_{bp700} vertical profile was smoothed with sequential 11-point running-minimum and running-maximum filters to separate the baseline from the spikes. The baseline signal corresponds to the bulk population of small particles, whose diameter is smaller than 100 µm and mostly between 0.5 and 30 µm (Dall'Olmo et al., 2009; Organelli et al., 2018). Each spike reflects the passage of a particle larger than about 100 µm in front of the sensor window. Previous studies inferred that backscattering spikes are caused mostly by phytodetrital aggregates, but also by large zooplankton and phytoplankton (Bishop and Wood, 2008; Briggs et al., 2011; Gardner et al., 2000). Assuming that backscattering sensors sample a volume of 10 mL (Briggs et al., 2020), we estimated that backscattering spikes concentration was typically between a few and <100 L^{-1}, consistent with previous independent estimates (McDonnell and Buesseler, 2010; Stemmann et al., 2008; Stemmann and Boss, 2012). Backscattering spikes were on average 4–10 times more abundant than chlorophyll fluorescence spikes. The b_{bp700} spikes larger than 0.008 m^{-1}, associated with particles larger than ~2 mm, were removed, with a negligible impact on the total spike signal (Briggs et al., 2020). Unlike Briggs et al. (2020), we did not subtract from the baseline profile the 850–900 m signal, which in that study was attributed to a background of small refractory particles with constant concentration. The baseline and spike signals were converted to SPOC and LPOC, respectively, as described in the next section.

All measurements were subsequently averaged into 18 vertical bins of progressively increasing thickness, such that the deepest bins contained at least 10 measurements. Finally, each profile was interpolated onto the L75 vertical grid commonly used in

NEMO simulations. This grid has 46 bins between the surface (0–1 m) and the deepest layer considered here (901–996 m). The profile time series was temporally binned into 5-day periods.

195   For the comparison to PISCES 1D simulations, BGC-Argo time series were cut into one-year periods (shifted by 6 months in the Southern hemisphere), which we will call coherent annual time series (CATS) hereafter. The CATS fulfilled the following conditions: (1) sampling dates spanned at least between days of year 25 and 340; (2) the float remained in the same region and did not cross major oceanic fronts according to the vertical-temporal evolution of temperature, salinity, σ_θ and spiciness; (3) bottom depth was > 1000 m for all profiles (bathymetry obtained from the 15 arc-second GEBCO 2019 product;
200   https://www.gebco.net/data_and_products/gridded_bathymetry_data/gebco_2019/gebco_2019_info.html); and (iv) the Chla and b_{bp700} sensors were stable according to both the vertical profiles and the continuous measurements acquired during drift at 1000 m between profiles. A total of 50 CATS from 28 different floats were selected, with 10–16 CATS in each biome, 32 (18) in the Northern (Southern) hemisphere (Table S1).

2.3 Conversion of b_{bp700} to POC

205   To convert the profiles of the backscattering coefficient at 700 nm (b_{bp700}) to POC we developed an empirical algorithm building on previous studies (Bol et al., 2018; Evers-King et al., 2017). The behaviour of this algorithm is summarized in Fig. 2 and discussed in section 4.2. Further details are provided in the Appendix. The algorithm estimates the POC/b_{bp700} ratio along the vertical profile between 0 and 1000 m, and proceeds in two steps. First, the POC/b_{bp700} ratio is calculated by prescribing a POC/b_{bp700} ratio in the surface layer ($z_{surf,biome}$) and an exponential decrease with depth in the underlying water column, which
210   converges asymptotically towards a constant deep value (c):

$$\frac{POC}{b_{bp700}}(z) = c + a_{biome} \cdot e^{-0.001 \cdot b \cdot (z - z_{surf,biome})}, \quad z > z_{surf,biome} \tag{1}$$

The a_{biome} coefficient is biome-specific, whereas the asymptote at depth is fixed at c = 1000 mmol C m^{-3} m. The $z_{surf,biome}$
215   corresponds to the 5% quantile of the climatological MLD in summer in a given biome (here ranging between 14 m in the Mediterranean and 41 m in the Subantarctic). The POC/$b_{bp,700}$ ratios at $z_{surf,biome}$, corresponding to a_{biome} + c in Eq. 1, are taken from the literature and range between 2600 and 4900 mmol C m^{-3} m (Fig. 2; Table A1). Second, the POC/$b_{bp,700}$ profile derived from Eq. 1 is modified by extrapolating a constant POC/$b_{bp,700}$ value, taken from a reference depth, z_{ref}, to the sea surface. In each vertical profile, z_{ref} is defined as the deepest of $z_{surf,biome}$ and the MLD:
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$$\frac{POC}{b_{bp700}}(z) = \frac{POC}{b_{bp700}}(z_{ref}), \; z \leq z_{ref}; \quad z_{ref} = \max(z_{surf,biome}, MLD); \tag{2}$$

The exponential decrease prescribed by Eq. 1 is similar to that proposed by Bol et al. 2018, except for the inclusion of the constant term c that prevents the ratio from becoming 0 at depth. The slope of the exponential decrease (b = -6.57) is constant in all biomes and based on our fit to the Cetinić et al. (2012) dataset, using the same depth bins as Bol et al. (2018), but additionally forcing the curve towards c at 1000 m.

The uncertainty of regional b_{bp700}-POC conversion factors in the epipelagic is typically <10% according to the standard error of the POC vs. b_{bp700} linear regression slopes (Table A1). The few available measurements in the mesopelagic suggest a POC/b_{bp700} uncertainty lower than a factor of two. Through this study, we will assume that model/observation ratios larger/smaller than 2/0.5 can safely be regarded as model over/underestimates, which possibly is a conservative criterion for the epipelagic layer.

The conversion of b_{bp700} to POC was done using different MLD data for the global climatologies and the CATS. For the global climatologies we used the Monthly Isopycnal & Mixed-layer Ocean Climatology (MIMOC) of Schmidtko et al. (2013), downloaded from https://www.pmel.noaa.gov/mimoc/, which was reprojected onto the ORCA2 horizontal grid. Although MIMOC is based on an algorithm that evaluates several MLD criteria, it has been shown to be globally consistent with the MLD based on a 0.03 kg m^{-3} σ_θ threshold (Holte and Talley, 2009; Sallée et al., 2021). For the float time series, we used the MLD defined by a 0.03 kg m^{-3} threshold computed for each individual profile.

2.4 Ocean color satellite data

Satellite observations for the 1997–2019 period were downloaded from GlobColour (https://www.globcolour.info), a merged multisensor dataset, on 2 March 2020. Monthly sea-surface POC fields based on the algorithm of Stramski et al. (2008) were used to compute monthly climatologies that were subsequently reprojected onto the ORCA2 grid.

2.5 PISCES simulations and matching with observations

Simulations were run using the ocean biogeochemistry model PISCESv2 (Aumont et al., 2015) with the RC parameterization for detrital POC (Aumont et al., 2017). The configuration of PISCES used here has 24 tracers: two classes of phytoplankton ("nanophytoplankton" and diatoms), detrital particles (small and big), and zooplankton (micro- and mesozooplankton), plus 18 additional tracers that comprise dissolved inorganic macronutrients and iron, inorganic carbon chemistry variables, dissolved organic carbon (DOC), and different particulate stocks of iron and silica. Phytoplankton growth depends on light, inorganic nitrogen, phosphorus, and iron, with an additional silicate requirement for diatoms. Microzooplankton and mesozooplankton consume the two classes of phytoplankton and small detrital particles with different preferences, and mesozooplankton also predate on microzooplankton. Detrital particles are produced through the mortality of phytoplankton and zooplankton (which are routed to small and large particles in different proportions), zooplankton sloppy feeding, and DOC coagulation. Production of large detritus also results from enhanced diatom mortality upon bloom collapse, aggregation of

255 small detritus, and zooplankton mortality and fecal pellet production (the latter two derived from a closure term that accounts for unresolved higher trophic levels). Small and large detrital particles are nominally smaller/larger than 100 μm and sink, respectively, at 2 and 50 m d^{-1}. Both small and large detritus are remineralized by implicit bacterial activity, and consumed by flux-feeding mesozooplankton. Remineralization follows first-order kinetics with an initial specific rate "k" of 0.035 d^{-1} (at 0°C) for freshly produced detritus in the upper mixed layer. This "k" decreases with depth as an emergent result of the RC

260 parameterization. To account for bacterial solubilization of aggregate-binding polymers, 10% of the degraded large detritus are routed to small detritus (this fraction is hard-coded based on previous calculations). The flux feeding rate depends on the particles' sinking flux, and thus attenuates the flux of large particles more strongly than that of small particles. Additionally, a variable fraction of the large detritus intercepted by flux feeders is fragmented into small detritus, attenuating up to around 50% of the large detritus sinking flux through the top 500 m during intense export events. Phytoplankton growth rates and

265 remineralization rates increase with temperature with a Q$_{10}$ of 1.9, whereas zooplankton growth rates have a Q$_{10}$ of 2.14.

To evaluate PISCES simulations against in situ POC measurements or their proxies, the correspondence between PISCES tracers (*in italics*) and observed POC fractions must be established. In this study we assumed that SPOC corresponds to the sum of PISCES-simulated nanophytoplankton (*PHY*), diatoms (*PHY2*), small detrital particles (*POC*) and microzooplankton

270 (*ZOO*), whereas LPOC corresponds to the sum of PISCES-simulated large detrital particles (*GOC*) and mesozooplankton (*ZOO2*) (Table 2). Total POC (TPOC) corresponds to the sum of those six PISCES tracers or, what is the same, SPOC + LPOC. Heterotrophic prokaryotes (*BACT*) are not a prognostic tracer in PISCES and are not explicitly included in our analysis. The correspondence between observed and simulated POC fractions, explicit and implicit, is discussed in section 4.3.

2.5.1 PISCES 3D simulations vs. biome-aggregated observations

275 For the global-scale comparison between PISCES outputs and observations from BGC-Argo and satellites, we used the simulation presented by Aumont et al. (2017), which was forced by pre-computed dynamical fields from a pre-industrial run of the ocean circulation model NEMO. Global monthly climatological fields of the PISCES tracers were used to compute seasonal climatologies of modeled SPOC, LPOC and TPOC. To enable direct comparison to the BGC-Argo observations, model output was resampled at locations where BGC-Argo profiles were available over the 2010–2019 period. Prior to

280 comparison with modeled fields, BGC-Argo observations were further screened to remove "outliers" in each biome and season. Outliers were defined as grid cells where the mean b$_{bp700}$ in the upper 50 m was above the 95% percentile or greater than 0.008 m^{-1} (Briggs et al., 2020). The same spatial resampling was applied to satellite retrieved POC.

2.5.2 PISCES 1D simulations vs. BGC-Argo coherent annual time series (CATS)

A more detailed comparison was undertaken by matching each of the CATS from individual BGC-Argo floats with a PISCES

285 water-column ("PISCES 1D") simulation. The match was based on the coherence between the seasonal cycle of MLD observed by the float and the turbulent layer simulated by NEMO. The pre-computed dynamical fields used to evaluate the match-ups

were obtained from an ocean-only historical simulation (NEMO v3.6) at 1° resolution with 75 vertical levels (ORCA1_L75 grid) forced with the JRA-55 atmospheric reanalysis that covered the 1958–2018 period (Tsujino et al., 2020) following the OMIP2 protocol (Griffies et al., 2016). The float-observed MLD and the NEMO-simulated turbocline depth (defined by

290 turbulent vertical diffusivity > $5 \cdot 10^{-4}$ m^2 s^{-1}) were compared over the entire annual cycle in all the model grid cells that had been visited by the float on a given year. The best model grid cell was selected based on model-observations correlation, root-mean-square error, and time lag in the onset date of summer stratification. An example of the metrics used to match NEMO dynamical fields and BGC-Argo MLD is provided in Fig. S1-S4.

295 PISCES 1D offline simulations were forced using the dynamical fields from the selected model grid cells. The same annual forcing, corresponding to the year of the BGC-Argo observations, was repeated over 5 simulation years. After 4 years of spin-up, the output from year 5 at 5-days resolution was used for the comparison to the BGC-Argo CATS. Initial conditions (climatological fields of inorganic nutrients and carbon chemistry variables) and boundary conditions (atmospheric deposition) were the same used for PISCES 3D (Aumont et al., 2015, 2017). Nutrient fields were restored towards the mean annual profile

300 below 300 m. This procedure avoided drift in nutrient stocks by replenishing the upper ocean with the same amount of nutrients each year, resulting in regular seasonal cycles after one year and identical cycles from year 4 onwards.

3 Results

3.1 Climatological POC fields

This section describes the comparison among TPOC fields estimated from BGC-Argo and satellite observations and PISCES

305 simulations across four ocean biomes. Fig. 3 compares monthly climatologies at the sea surface (0–20 m), Fig. 4 compares seasonal climatologies between 0–1000 m, and Fig. 5 displays skill metrics (Pearson's correlation, model/observations ratio and bias) for the vertical profiles shown in Fig. 4, as well as for the 1D simulations matched to BGC-Argo CATS.

3.1.1 Seasonally stratified subpolar biomes

In the subpolar biomes, near-surface TPOC ranged between ~1 mmol m^{-3} in the winter months and around 5 (Subantarctic) or

310 10 (NASPG) mmol m^{-3} in early summer. In these biomes, PISCES-simulated TPOC was within the 2.5–97.5% bounds of BGC-Argo observations for most months (Fig. 3). During the apex of the bloom (months 5–7), however, median PISCES estimates exceeded those obtained from BGC-Argo (by ~80%) and satellites (by ~15%). In the Subantarctic, this pattern extended through the fall. Satellite TPOC was in poor agreement with both PISCES and BGC-Argo TPOC outside the apex of the bloom, exceeding BGC-Argo by up to seven-fold (fourfold) in the NASPG (Subantarctic), as discussed in section 4.1.

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PISCES reproduced the vertical decrease of TPOC concentration down to 1000 m (Fig. 4) with generally good skill (Fig. 5), but some misfits were observed. In the North Atlantic subpolar gyre (NASPG), PISCES underestimated TPOC through the

epipelagic and the upper mesopelagic during the winter by ~40% (Fig. 4) because of too vigorous convection in the NEMO dynamical fields that kept phytoplankton under insufficient light exposure. In the Subantarctic biome, simulated TPOC exceeded BGC-Argo estimates in the upper portion of the mesopelagic layer in spring, and the overestimation pattern propagated downwards through the summer and fall. A similar but smaller overestimation pattern was observed in the NASPG in summer and fall.

3.1.2 Permanently and seasonally stratified subtropical biomes

In the oligotrophic biomes, monthly median surface TPOC displayed low seasonal amplitude. Total POC concentrations estimated from BGC-Argo data typically oscillated around 2 mmol m^{-3}, with a maximum/minimum ratio of around 1.6 in the Mediterranean and 1.3 in the Atlantic and Pacific subtropical gyres (STG). In the STG, satellite and BGC-Argo TPOC were in good agreement, which could be expected because subtropical b_{bp700}-POC conversion factors and satellite POC are based on the same study (Stramski et al., 2008). On the contrary, PISCES TPOC exceeded BGC-Argo TPOC by around twofold in the STG. In the Mediterranean, satellite TPOC exceeded BGC-Argo TPOC by ~80%, pointing to the differences in the respective POC estimation algorithms. PISCES TPOC was nearly fourfold higher than BGC-Argo TPOC at the surface in the Mediterranean, an overestimation that results from unrealistic physics in that basin caused by the too-coarse (2°) model grid (Tonani et al., 2008; Lebeaupin Brossier et al., 2011) (see 3.3 and 4.3). Vertical TPOC profiles evidenced the shortcomings of PISCES simulations in the oligotrophic gyres. Compared to BGC-Argo profiles, PISCES simulations produced too-sharp deep POC maxima and underestimated TPOC in the waters above and below (Fig. 4). These mismatch patterns prompted us to examine in greater detail the seasonal cycles of POC in different biomes.

3.2 Coherent annual time series of SPOC and LPOC: case studies

In this section we describe two CATS from BGC-Argo floats and their PISCES 1D counterparts (section 3.2.1). The floats, identified by their World Meteorological Organization (WMO) number, are the 6901486 in the NASPG (year 2015) and the 6901660 in the South Pacific STG (year 2017). The float 6901486 represents the most productive conditions of our dataset, with annual median (maximum) Chla of 0.60 (10.7) mg m^{-3} and TPOC of 5.5 (16.7) mmol m^{-3} in the near-surface layer (0–20 m). By contrast, float 6901660 represents the most oligotrophic waters, with annual median (maximum) Chla of 0.011 (0.036) mg m^{-3} and TPOC of 1.8 (2.5) mmol m^{-3} in the near-surface layer. In Fig. S5–S10 we provide additional examples of the CATS-PISCES 1D match-ups in the four biomes and in two subregions within the NASPG.

3.2.1 Labrador Sea (North Atlantic subpolar gyre)

Float 6901486 was deployed in June 2013 close to the Reykjanes Ridge in the Irminger Sea, NW Atlantic subpolar gyre. After drifting SW carried by the East Greenland Current, the float was trapped in the Labrador Sea cyclonic circulation between 2014 and July 2017, when it stopped communication after completing 344 profiles (cycles). During its multi-year sampling in the Labrador Sea (56-60°N latitude and 48–54°W longitude), over a bottom depth of around 3500 m, the float showed stable

physical and bio-optical records at 1000 m, broken only by winter convection events, and recurring annual patterns of spring-summer phytoplankton blooming and vertical carbon export as depicted by Chla and POC profiles. Here we describe the year 2015 (Fig. 6), characterized by deep convection during February and March, when the MLD generally exceeded 1000 m (Fig. S1). Epipelagic TPOC increased rapidly upon water-column re-stratification in mid-April and peaked in mid-May. A secondary bloom peaked in late June after a transient MLD deepening caused by stormy weather. Epipelagic TPOC decreased progressively thereafter until a small bloom was observed in October linked to pycnocline erosion. This bloom terminated rapidly and epipelagic TPOC reached the baseline level in late December. The SPOC fraction dominated epipelagic TPOC all year round, and highest LPOC fractions of nearly 20% were recorded at the apex of the spring-summer blooms. Distinct vertical particle export events were observed in May and June, matching the surface phytoplankton blooms, and August, when nutrient limitation likely triggered bloom collapse. These export pulses produced synchronous increases in SPOC and LPOC through the mesopelagic layer, though with different magnitudes. After reaching relative minima in October, mesopelagic SPOC and TPOC increased again in November, but they showed different vertical patterns.

The matching PISCES 1D simulation captured with good skill the patterns of SPOC and LPOC in the epipelagic and, to a lesser extent, the mesopelagic layer (Fig. 6). Excellent correlation (r = 0.92) and bias (-0.5%) between BGC-Argo and PISCES were found for vertically integrated epipelagic TPOC (Fig. 6i). A delayed start of the bloom was observed in the simulation, which can be partly attributed to a delay of around one week in the onset of permanent stratification in the model. It is also plausible that modelled phytoplankton reacted too weakly to the cessation of deep convection, which was captured by alternative MLD metrics in BGC-Argo profiles (Fig. 6a and b). Despite the general good agreement between observed and simulated SPOC, the model produced a conspicuous plume of SPOC that sank from the surface spring bloom into the mesopelagic layer, at the prescribed constant rate of 2 m d^{-1}, which was not found in the observations. In the core of this plume, PISCES POC exceeded BGC-Argo SPOC by more than twofold. A similar model-observations mismatch was observed in all the northern and southern subpolar CATS as well as in some CATS in the Mediterranean (Figures S5-S8 and S10). On average, simulated LPOC exceeded BGC-Argo LPOC by 36% and 96% in the epi- and mesopelagic layers, respectively. The largest overestimation was observed during the midsummer export event. On the other hand, LPOC underestimation was found during the May bloom between 0–400 m.

PISCES qualitatively reproduced the late summer peak of mesopelagic LPOC, which was observed in all the subpolar North Atlantic CATS. On the contrary, it failed to reproduce both the decrease of SPOC and LPOC in fall between 600-800 m and the LPOC increase below 800 m. The latter occurred in 6 out of 11 CATS in the NASPG, all located in the Labrador Sea. The apparent decoupling of deep mesopelagic LPOC from the overlying water column may be related to the insufficient temporal resolution of BGC-Argo profiling during that period compared to LPOC sinking speed, or reflect LPOC export events from surface waters not located vertically over the float (Siegel and Deuser, 1997).

3.2.2 South Pacific subtropical gyre

Float 6901660 was deployed in March 2015 in the western South Pacific STG, and drifted westwards until it deflected SW while approaching Tahiti. As of March 2021, the float was still active and had completed 244 cycles with stable continuous records at the 1000 m drift depth. Between July 2017 and June 2018 (18-21°S latitude and 148-157°E longitude), the period selected for the CATS analysis, the BGC-Argo profiles portrayed a stably stratified water column typical of the core of the subtropical gyres (Fig. 7). Vertical mixing events that reached a depth of around 100 m were observed in July, August and October. However, their effect on surface SPOC was hardly noticeable, indicating that turbulent entrainment of nutrients was too weak to stimulate new production significantly. A deep Chla maximum was present all year round between 150 and 200 m as identified by the maximum Chla gradient. This Chla maximum did not translate into a deep POC maximum. The fraction LPOC/TPOC was consistently around 6% in the epipelagic and 12% in the mesopelagic according to the vertically integrated stocks. The vertical-temporal distribution of LPOC was patchy in the lower mesopelagic (500–1000 m), perhaps reflecting the difficulty to detect rare aggregates from b_{bp700} spikes.

The epipelagic TPOC stock (driven by SPOC) simulated by PISCES matched well the observations, with a high temporal correlation coefficient (r = 0.69; Fig. 7i) despite the low seasonal variability in this tropical setting. PISCES overestimated SPOC between the base of the mixed layer and the deep Chla maximum, and underestimated SPOC below it. Thus, the low mean bias of epipelagic SPOC (9%) resulted from these mutually compensating biases. In the mesopelagic layer, vertically integrated PISCES SPOC was on average 45% lower than BGC-Argo SPOC and showed low negative temporal correlation to the observations. Regarding LPOC, the simulated stock was nearly twice as large as BGC-Argo estimates in the epipelagic, with the largest overestimation seen in the deep Chla maximum. In the mesopelagic, PISCES LPOC exceeded BGC-Argo estimates by 25% on average. Despite these deviations, PISCES simulations supported the increase in the LPOC/TPOC fraction between the epipelagic and the mesopelagic layers deduced from BGC-Argo profiles.

3.3 Coherent annual time series of SPOC and LPOC: generalized approach

Although each of the 50 CATS included in this study has unique features, some of the misfit patterns between PISCES and BGC-Argo data described in the previous section are common to most CATS in a given biome or, more broadly, in subpolar vs. subtropical biomes. In this section we generalize the quantitative comparison between the 50 BGC-Argo CATS and their PISCES 1D counterparts across the four biomes. We consider separately the epipelagic and mesopelagic domains, focusing on mean annual SPOC and LPOC standing stocks (Fig. 8), the SPOC/TPOC fraction (Fig. 9), and the mesopelagic Teff of SPOC and LPOC (Fig. 10).

Epipelagic SPOC stocks ranged between 193–425 mmol C m^{-2} (1.0–2.1 mmol C m^{-3} in concentration units) according to BGC-Argo observations. The SPOC range in the corresponding PISCES 1D simulations was 282–537 mmol C m^{-2} (1.4–2.7 mmol

C m^{-3}). Although the ranges of the different biomes overlapped, smaller stocks were usually found in the more oligotrophic

415 (STG and Mediterranean) biomes. The agreement between simulated and observed epipelagic SPOC was better in the STG and NASPG biomes, whereas simulated epipelagic SPOC exceeded observations by around 50% in the Mediterranean and Subantarctic biomes. Simulated and observed stocks were positively correlated (r = 0.45, p = 9·10^{-4}).

In the mesopelagic domain, observed and simulated SPOC stocks ranged between 145–283 and 105–308 mmol C m^{-2},

420 respectively (corresponding concentration ranges of 0.18–0.35 and 0.13–0.39 mmol C m^{-3}). The correlation between simulated and observed mesopelagic SPOC stocks was not significant (r = -0.08, p = 0.55). PISCES SPOC was similar to or slightly higher than BGC-Argo SPOC in the subpolar biomes, but up to two-fold lower in the STG and Mediterranean biomes, as previously shown in Fig. 4 and 7.

425 Epipelagic LPOC stocks were smaller and showed wider inter-biome variability than SPOC stocks, with around twofold higher LPOC in subpolar biomes. The positive correlation between simulated and observed LPOC was highly significant (r = 0.78, p = 2·10^{-11}). Yet, PISCES LPOC (range 19–83 mmol C m^{-2}, 0.10–0.42 mmol C m^{-3}) typically exceeded BGC-Argo LPOC (range 27–119 mmol C m^{-2}, 0.14–0.60 mmol C m^{-3}) by around 50%, and up to threefold for some CATS. A lower but still highly significant correlation was found in the mesopelagic (r = 0.64, p = 5·10^{-7}), where PISCES LPOC was usually within a factor

430 of 1.5 of observations, except for the NASPG biome where it exceeded BGC-Argo LPOC by around twofold.

The SPOC/TPOC fraction showed low variability in the epipelagic layer (Fig. 9), with a median (range) of 89% (84–94%) and 85% (75–91%) for BGC-Argo and PISCES, respectively. The simulated SPOC/TPOC fraction was within ±10% of BGC-Argo estimates for all biomes except the NASPG, where PISCES tended to underestimate the observed SPOC/TPOC fraction.

435 A significant positive correlation (r = 0.56, p = 2·10^{-5}) was found between simulations and observations in the epipelagic layer. The SPOC/TPOC fraction was lower in the mesopelagic layer according to both BGC-Argo (75–90%) and PISCES (66–85%) estimates. In this case, CATS from the Subantarctic biome showed excellent model-observations agreement, whereas PISCES was 20% lower than BGC-Argo estimates in the other three biomes. No significant correlation was found between the simulated and observed SPOC/TPOC fraction in the mesopelagic.

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Transfer efficiency (Teff; Fig. 10) was computed as the ratio between the SPOC or LPOC concentrations in the depth bins centered at 697 m (range 662–734 m) and 200 m (range 190–210 m). The shallowest bin corresponds to the bottom of the epipelagic layer, and the 500 m interval was chosen following previous studies (Lam et al., 2011; Dall'Olmo and Mork, 2014). Different depth ranges between 180 and 800 m gave similar Teff patterns. Analysis of Teff yielded some important insights:

445   1. SPOC and LPOC showed similar Teff, both in the observations (medians of 0.39–0.41) and in the model (medians of 0.27–0.29). Similar patterns were found when each biome was considered separately.

2. BGC-Argo T_{eff} usually exceeded PISCES T_{eff} and spanned a wider range. The best agreement was generally found in the STG and the Subantarctic biomes, and the poorest agreement occurred in the Mediterranean where BGC-Argo T_{eff} was typically twice the modelled T_{eff}.

3. Distinct patterns were found for four CATS in the Labrador Sea (NASPG) characterized by high epipelagic TPOC (>440 mmol C m^{-2}). In this subset, PISCES and BGC-Argo T_{eff} were in good agreement for SPOC (median 0.40 for both) whereas, for LPOC, PISCES T_{eff} (0.36–0.45) doubled BGC-Argo T_{eff} (0.12–0.28), which had some of the lowest values of the dataset. These CATS showed a LPOC minimum at 600–800 m in fall, which PISCES could not reproduce (section 3.2.1).

4. PISCES and BGC-Argo T_{eff} showed a weak positive correlation for SPOC (r = 0.26, p = 0.07), with some improvement when the Labrador Sea "outliers" were removed (r = 0.34, p = 0.02). A negative correlation between simulations and observations was found in the case of LPOC T_{eff} (r = -0.29, p = 0.04), which became non-significant without the Labrador Sea "outliers" (r = 0.04, p = 0.81).

Our observational estimates are sensitive to the choice of b_{bp700}-POC conversion factors. These factors are especially uncertain in the mesopelagic due to data scarcity, which prompted us to use a constant global value for "c" in Eq 1. (Fig. 2); c = 1000 mmol C m^{-2} is the asymptotic POC-b_{bp700} conversion factor below 1000 m, taken from Cetinić et al. (2012). To address this uncertainty, we conducted sensitivity tests where c was halved or doubled. The resulting range of 500–2000 mmol C m^{-2} is probably generous, as indicated by our indirect estimates based on the studies of Bishop (1999) and Bishop et al. (1999) (Appendix A). As expected, changing c had little effect on epipelagic POC, a larger effect on mesopelagic POC and T_{eff}, and no effect on the SPOC/TPOC fraction (because the same conversion factor is used to estimate SPOC and LPOC). Halving c resulted in steeper POC profiles, which brought mesopelagic SPOC closer to the 1:1 line in the STG and Mediterranean, at the expense of increasing the SPOC bias in the subpolar biomes and that of LPOC everywhere (Fig. S11). Doubling c caused a less steep vertical decrease of BGC-Argo POC, which overall worsened the model-observations agreement in the mesopelagic (Fig. S12), except for LPOC stocks in the NASPG.

4. Discussion

4.1 Towards a globally consistent picture of POC fields in observations and models

Global quantification of POC stocks through the water column has been elusive until recently because of data sparseness and limitations of model parameterizations, both of which are especially severe below the epipelagic. Our joint analysis of PISCES simulations, satellite observations and over 70,000 BGC-Argo vertical profiles reveals a globally consistent picture across the epi- and mesopelagic layers (Fig. 3-10). The global ocean stock of TPOC simulated by PISCES amounts to 4 Pg C, shared in a proportion of 39% (1.6 Pg C), 25% (1 Pg C) and 36% (1.4 Pg C) between the epi-, meso- and bathypelagic layers (Table 3; the bathypelagic is defined here as 1000–5000 m to include the entire model domain in the stock calculation).

480   The PISCES-simulated TPOC concentration is on average within a factor of 1.56 (1.42) of BGC-Argo estimates for the median (mean) seasonal biome profiles shown in Fig. 4 (Mediterranean excluded). Aumont et al. (2017) reported a similar reliability index of 1.6 for the comparison between PISCES and in situ chemically determined POC profiles. Thus, our evaluation lends further confidence to the POC reactivity continuum parameterization implemented in PISCES, which represents both SPOC and LPOC as a mixture of fractions with different lability (Aumont et al., 2017), in globally representative biomes.

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The epipelagic TPOC stock simulated by PISCES, 1.6 Pg C, is comparable to previous observational estimates. Gardner et al. (2006) estimated the global POC stock within the first optical attenuation depth using a compilation of in-situ POC and c_p measurements leveraged by ocean color satellite data. They obtained a global stock of 0.43 Pg C, and invoked some scaling arguments to estimate that the total POC stock down to middle-mesopelagic "background levels" would range 1–2 Pg C.

490   Stramska (2009) obtained a larger global epipelagic POC stock of 1.8–2.3 Pg C using the satellite algorithm of Stramski et al. (2008). Our match-up analysis indicates that satellite TPOC exceeds BGC-Argo estimates by up to seven-fold at high latitudes outside of the summer season (Fig. 3). This deviation is well beyond the nominal uncertainty of the satellite POC product (<30%) and the range of observed POC/b_{bp700} ratios at the sea surface (Fig. 2). Therefore, we conclude that the algorithm of Stramski et al. (2008) overestimates POC at high latitudes in winter, an issue that deserves further investigation. Potential

495   explanations are the satellite algorithm being calibrated mostly against samples from lower latitudes (50°N–30°S), or its sensitivity to the differential atmospheric attenuation of blue and green wavelengths at low solar elevation.

Recently, Evers-King et al. (2017) calculated the global POC stock in the upper mixed layer using several satellite algorithms (including that of Stramski et al., 2008) and indicated that 0.77–1.3 Pg C was a plausible range. Our PISCES-based estimate

500   for the mixed-layer POC stock, using the same MLD climatology (Schmidtko et al., 2013), is 0.58 Pg C. The lower PISCES-derived estimate may arise from the combination of POC overestimation by some satellite algorithms, discussed above, with PISCES' tendency to underestimate mixed-layer POC in oligotrophic areas (Fig. 4 and 7). This negative bias in PISCES is compensated by a positive POC bias in deep Chla maxima (below the MLD), resulting in smaller deviations between BGC Argo and PISCES over the entire epipelagic layer in the STG biome (Fig. 8 and S13). Given the limited coverage of in situ

505   seawater sampling and satellite observations in some regions and seasons (Evers-King et al., 2017), further intercomparison between those observations, BGC-Argo data and models is needed to better constrain epipelagic POC stocks.

POC in the lower mesopelagic and below has been traditionally treated as a "background" signal (Bellacicco et al., 2019; Gardner et al., 2006; Loisel and Morel, 1998). This approach is convenient for studies that focus solely on upper-ocean

510   processes because POC concentration decreases exponentially with depth. Yet, our global estimates and several previous studies highlight the need to turn our attention to the large POC stocks (> 2 Pg C) that reside in the meso- and bathypelagic layers, whose dynamics are still poorly understood. In line with previous studies (Dall'Olmo and Mork, 2014; Poteau et al.,

2017), we showed that mesopelagic POC exhibits clear seasonal cycles in productive regions (Fig. 6, S5–S7 and S10), owing to their connection with the upper-ocean through numerous biological and physical processes (Boyd et al., 2019; Briggs et al.,
515   2020). Despite being less reactive on average than upper-ocean POC, meso- and bathypelagic organic particles are microbial hotspots that host key biogeochemical functions, from enzymatic decomposition of macromolecules (Arnosti et al., 2012; Baltar et al., 2010a, 2010b) to aerobic and anaerobic respiration (Bianchi et al., 2018; Karthäuser et al., 2021) and chemosynthesis (Arístegui et al., 2009; Herndl and Reinthaler, 2014; Pachiadaki et al., 2017). Moreover, mesopelagic particles are consumed by upper trophic levels that sustain fisheries (Bode et al., 2021; Woodstock et al., 2021).
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In situ bio-optical measurements are poised to play a key role in monitoring marine POC stocks in layers that cannot be accessed by remote sensing. For example, Sauzède et al. (2020) merged BGC-Argo and satellite observations to obtain a dynamic 3D view of particle backscattering. Using a data-driven machine learning approach, they were able to predict the profiles of $\log_{10}b_{bp700}$ measured by two BGC-Argo floats in the NASPG and STG biomes (R^2 of ~0.85 and mean absolute
525   percentage deviation of ~12%) from the sole knowledge of physical properties of the water column and surface ocean color (remote sensing reflectance). Their estimates were recently extended to POC (Sauzède et al., 2021), which can be of great utility for constraining biogeochemical models. Here we took an entirely different approach, based on converting available b_{bp700} data to POC with a simple empirical algorithm (Fig. 2) and then comparing it to the outputs of the PISCES model. Our PISCES-based estimates obtained a median $R^2 = 0.86$ and mean absolute percentage deviation of 38% (5-day depth-binned
530   $\log_{10}$POC; 50 CATS from 28 globally distributed floats). This good skill is remarkable because neither the empirical POC estimates nor PISCES were tuned to maximize their mutual agreement. Still, our study shows that the comparison of bio-optics-derived POC measurements and PISCES is affected by different types of uncertainty that we analyze in the following sections.

4.2 Bio-optical underpinnings of POC fields based on BGC-Argo observations

535   Accurate interconversion between bio-optical variables and concentrations is key for constraining ocean particle dynamics and their model representation (Bishop et al., 2004; Gardner et al., 2006). Here we discuss the processes that appear to drive, to first order, the variability of the POC/b_{bp700} ratios embodied in Eq. 1 and 2 (Fig. 2; Appendix A), and the main strengths and weaknesses of our scheme.

540   Changes in the trophic status appear as the primary driver POC/b_{bp700} variability in the epipelagic layer (Cetinić et al., 2012; Figure A1). Productive waters host greater absolute and relative abundance of diatoms (Uitz et al., 2006) (see also Table 3), which have lower POC per cell volume (Menden-Deuer and Lessard, 2000) and are covered with silica frustules that may scatter light more efficiently than naked cells (Twardowski et al., 2001), altogether resulting in lower POC content per unit b_{bp700} (Cetinić et al., 2012; Oubelkheir et al., 2005). The proportions of different autotrophic and heterotrophic organisms and
545   detritus are also likely to vary with upper ocean productivity (see 4.3). If the mass-specific backscattering coefficients of these

components were better known, their systematic variation patterns could be used to develop a continuous formulation for POC/b_{bp700}, rather than the regionalized conversion factors used here. However, POC/b_{bp700} is influenced by other seawater constituents whose occurrence is less predictable. Foremost, biogenic calcite (e.g. from coccolithophores) and desert dust (Claustre, 2002; Loisel et al., 2011), both of which enhance b_{bp700}. In our data set, we detected a CATS (float WMO 6901647, year 2016) that was strongly affected by coccolith backscattering in the Iceland Basin, an area known for its massive coccolithophore blooms (Moore et al., 2012). This CATS was an obvious outlier in our model-observations scatterplots (Fig. S13), likely because the enhancement of b_{bp700} caused by coccoliths had no translation in TPOC, and was therefore removed from the analysis.

The decrease in POC/b_{bp700} along the vertical axis (Fig. 2) reflects the increase in the particles' index of refraction, hence the backscattering ratio (Cetinić et al., 2012; Nencioli et al., 2010). This change is likely caused by the remineralization of organic materials (Martin et al., 1987) that leaves a higher mineral fraction (Honjo et al., 2008; Lam et al., 2011), and the increase in the structural complexity of aggregates with depth (Organelli et al., 2018). According to our sensitivity analysis (Fig. S11 and S12), the prescribed exponential decrease of POC/b_{bp700} towards a constant POC/b_{bp700} (c = 1000 mmol C m^{-2}) at depths >1000 m provides a good compromise globally, given the limited knowledge of mesopelagic POC/b_{bp700} and its variability across regions. However, the comparison between simulated and observed mesopelagic SPOC (and thus TPOC) is more favorable in subpolar than in subtropical biomes. In the latter, better model-observations agreement was found when POC/b_{bp700} was halved (c = 500 mmol C m^{-2}). A lower c would also bring the simulated Teff for SPOC and LPOC closer to BGC-Argo estimates. It is tempting to hypothesize that lower latitudes have lower mesopelagic POC/b_{bp700} owing to the greater proportion of calcite (Francois et al., 2002; Honjo et al., 2008; Lam et al., 2011, 2015). These hypotheses need further verification.

Finally, the decrease of surface POC/b_{bp700} with deeper vertical mixing imposed by Eq. 2 partially reflects the dilution of surface particle assemblages by entrainment of deeper waters (Lacour et al., 2019) with lower POC/b_{bp700} (Bol et al., 2018). Modulation of POC/b_{bp700} by vertical mixing improves the agreement between PISCES and BGC-Argo data in regions with wide seasonal MLD amplitude such as the NASPG. Still, our approach should be seen as a simplistic first-order approximation, and alternative formulations should be further evaluated when more data become available. For example, POC/b_{bp700} might be kept constant regardless of the MLD, or estimated for each profile as the average within the mixed layer of the pre-computed POC/b_{bp700} profile (eq. 1). In both cases, however, POC/b_{bp700} would have to decrease abruptly below the mixed layer to meet the low POC/b_{bp700} in the mesopelagic, producing sharp discontinuities that have not been observed. On the other hand, the behavior prescribed by Eq. 2 may not be appropriate for situations when vertical mixing cannot erode the seasonal thermocline or pycnocline, because in such cases it will entrain water from the deep Chla maximum, and not mesopelagic water, to the upper mixed layer.

Our POC-b_{bp700} conversion algorithm omits several other sources of uncertainty. Foremost, it assumes that SPOC and LPOC

580 have the same POC/b_{bp700} ratio, which largely corresponds to that of the more abundant SPOC (Fig. 6–8). In addition, the in-situ data used to parameterize Eq. 1 and 2 are not free of uncertainty (Cetinić et al., 2012; Lam et al., 2011; Morán et al., 1999; Organelli et al., 2018; Strubinger Sandoval et al., 2021). Despite these limitations, the POC-b_{bp700} conversion scheme used here provides POC estimates that are generally consistent with PISCES (Fig. 8) and with previous assessments (Aumont et al., 2017). Moreover, this scheme allowed us to identify potential shortcomings of satellite-based assessments (Evers-King et al.,

585 2017; Stramski et al., 2008). The development of more sophisticated POC-b_{bp700} interconversion schemes is desirable and would greatly benefit from the measurement of mass-specific bio-optical properties of various seawater constituents across different ocean basins and depths.

4.3 Correspondence between observed and simulated POC fractions

An additional source of uncertainty in our analysis is the imperfect match between the PISCES tracers and the observable POC

590 fractions (Table 2). A positive aspect of our evaluation is the reasonable agreement between the simulated SPOC/TPOC fraction and the BGC-Argo estimates based on the b_{bp700} spike signal (Fig. 9). Our estimates converge to a global median value of 85%, with 95% of the data between 69–92%, fully within the range of size-fractionated chemical POC determination (Aumont et al., 2017 and references therein). Our SPOC/TPOC estimates can also be compared to the suspended POC estimated with the marine snow catcher during spring in subpolar epipelagic and upper mesopelagic waters (Baker et al., 2017).

595 The slow-sinking POC measured by the marine snow catcher should not be included in this comparison because it sinks at around 18 m d^{-1}, a range more typical of particles > 100 μm (Cael et al., 2021). The median 94% of suspended POC reported by Baker et al. is higher than the 86% (81%) SPOC/TPOC estimated here from BGC-Argo (PISCES) in the subpolar biome, suggesting further comparison between different approaches is needed. The tendency of SPOC/TPOC to decrease between the epipelagic and the mesopelagic, found in both PISCES and BGC-Argo data (Fig. 9, Table 3), was also reported in previous

600 studies (Aumont et al., 2017 and references therein; Organelli et al., 2020), lending further confidence to our estimates. On the other hand, the simulated partitioning of POC into different living and detrital compartments is probably less realistic, as discussed in greater detail below.

An aspect that deserves further attention is the partitioning between phytoplanktonic (autotrophic), heterotrophic and detrital

605 POC in the upper ocean. Unfortunately, this partitioning is far from being well established from observations. Using bio-optical data, Oubelkheir et al. (2005) estimated that detritus accounted for around 60% of POC across a wide range in ocean productivity, and 70–85% of POC was assigned to non-phytoplanktonic material (detritus + heterotrophs). These percentages accord well with those found by Claustre et al. (1999) in the tropical Pacific (their Table 2; see also Organelli et al., 2020). By contrast, Graff et al. (2015) found wide latitudinal variations of the non-phytoplanktonic POC, increasing between ~20 and

610 ~80% from the tropical to the temperate Atlantic. Bellacicco et al. (2019) analyzed the covariation between b_{bp700} and Chla in the (much larger) BGC-Argo dataset. They concluded that the "background" b_{bp700}, defined as the b_{bp700} that does not covary with Chla, decreases with productivity from >80% in subtropical gyres to <20% in the NASPG, with a mean contribution of

65% in oligotrophic areas. Although the "background" b_{bp700} fraction cannot be entirely assigned to non-phytoplanktonic material, the biogeographic patterns found by Bellacicco et al. (2019) are hardly compatible with those found by Graff et al. (2015) or with a nearly-constant detrital fraction across biomes. The detrital and non-phytoplanktonic POC fractions in PISCES near the sea surface range, respectively, between 15–31% and 42–59% (annual mean values by biome; Table 3). Although they are within the full range of observations, these values suggest that PISCES underestimates the percentage of detrital POC, especially in oligotrophic waters. A deeper analysis, properly matching observations and simulations over time and space, should be undertaken to obtain mechanistic understanding of upper-ocean POC partitioning, with potential consequences for remotely-sensed POC export estimates (e.g., Siegel et al., 2014).

PISCES bias may also arise from inappropriate representation of some POC reservoirs, such as heterotrophic prokaryotes (bacteria and archaea, *BACT* in PISCES), which have long been recognized as important contributors to the suspended POC (Morel and Ahn, 1990; Gasol et al., 1997). However, *BACT* are not explicitly modelled in PISCES as prognostic tracers, meaning they are not interacting fully with other tracers. Instead, they are diagnosed in the productive surface layer from zooplankton biomass, based on an old model version that had interactive *BACT*. Below this layer, *BACT* biomass is propagated downwards with a power function based on Arístegui et al. (2009), which resembles a Martin curve (Martin et al., 1987) and is therefore very sensitive to the reference depth (Z_0) (Buesseler et al., 2020). We find two main arguments against the inclusion of PISCES-estimated *BACT* in our POC estimates with the current model configuration. First, the empirical *BACT* estimation in PISCES has not been validated, to our knowledge, and may introduce noise in the comparisons. Second, PISCES-simulated POC already includes heterotrophic prokaryotes, because their biomass was not removed from the in situ POC measurements used to adjust the POC parameters in PISCESv2_RC (Aumont et al., 2017). In consequence, adding *BACT* to SPOC causes overestimation of mesopelagic POC (Fig. S14–S16) and can produce unrealistic temporal patterns (Fig. S15). Nevertheless, we believe that inclusion of prognostic bacteria would enable more realistic simulation of POC stocks, with the potential side effect of improving the simulation of element fluxes in PISCES.

The comparison between simulated and observed LPOC is a novel contribution of our study. The method of Briggs et al. (2011, 2020), originally developed to study intense POC export events, was here extended to estimate SPOC and LPOC separately over the full annual cycle through the epi- and mesopelagic domains. Our analysis shows that, despite the mismatch in terms of concentration, the LPOC derived from the spikes of high-resolution bio-optical profiles is strongly correlated to the PISCES-simulated LPOC ($r = 0.78$, $r^2 = 0.61$) along a wide trophic gradient (Fig. 8). This result is encouraging and supports the more widespread deployment of instruments that perform high-resolution bio-optical sampling to shed light on the spatiotemporal dynamics of large aggregates and particles (Briggs et al., 2020; Lampitt et al., 1993; Stemmann et al., 2008). On the other hand, it is unclear to what extent the LPOC inferred from the b_{bp700} spike signal is capturing mesozooplankton biomass, in addition to aggregates. Exclusion of PISCES mesozooplankton (*ZOO2*) from the comparison increases model-observations mismatch, with BGC-Argo LPOC exceeding PISCES estimates by around twofold, although the high correlation

remains (r = 0.76, p < 10^{-10}). Imaging devices mounted on BGC-Argo floats may provide more accurate quantification of LPOC, allowing for the separation of detrital LPOC (Trudnowska et al., 2021) from mesozooplankton and micronekton (Haëntjens et al., 2020) and the separation and quantification of particle classes contributing to flux across the complete particle spectrum (Bourne et al., 2021).

4.4 Importance of realistic physics and model evaluation across scales

In Fig. 5, PISCES simulations and BGC-Argo observations are compared using an array of skill metrics computed on the seasonal vertical profiles of TPOC between 0–1000 m. Starting with the 3D seasonal climatology, we observed that the correlation between the median (aggregated) profiles was generally better than the correlation between the spatially collocated (non-aggregated) profiles within a given biome and season (Fig. 5a), whereas no differences were found in terms of dispersion metrics (Fig. 5b and c). The difference in correlation was larger in subpolar biomes, suggesting that the model-observations spatial mismatch was magnified in regions with more energetic ocean dynamics and sharper physical and biogeochemical gradients, whose real-world location may not be well reproduced by the ocean circulation model used to force PISCES. For PISCES 1D simulations, their correlation coefficients with their CATS counterparts was usually in the high range of the correlation coefficients obtained by the biome-median PISCES 3D profiles. In terms of dispersion metrics, the ensemble of PISCES 1D simulations showed wider dispersion, but the best 1D simulations clearly outperformed the 3D simulation in a given region and season. The better skill of 1D simulations was more evident during spring, a season characterized by the onset of stable stratification after deep winter vertical mixing in middle and high latitudes. The greatest difference between 3D and 1D simulations was found in the Mediterranean, highlighting the more realistic vertical mixing and upper ocean productivity in the 1D simulations.

Our cross-scale evaluation indicates how crucial it is to evaluate model physics before extracting conclusions on biogeochemical model performance (Doney et al., 2004; Kriest et al., 2020; Löptien and Dietze, 2019). In our 1D CATS approach, the skill of PISCES simulations was maximized by carefully matching observed and modelled vertical mixing (Fig. S1–S4), which is a key driver of upper ocean ecosystems. This approach has a subjective component, and may also suffer from the idealized assumption that BGC-Argo profiles reflect mostly vertical-scale processes, disregarding horizontal advection (Alonso-González et al., 2009). Yet, the similar misfit patterns encountered for different CATS within a given biome support the robustness of the 1D matching approach (compare Fig. 6 versus S6, and Fig. 7 versus S9). To further evaluate this issue, we matched different neighboring NEMO grid cells to the same in situ CATS. Again, this exercise indicated that our main conclusions are not sensitive to the choices made for 1D model-observations matching (compare Fig. 6 and Fig. S5). Indeed, alternative matching approaches can be devised, each of which with advantages and pitfalls, for example: (1) sampling the outputs of biogeochemical models at the locations visited by BGC-Argo floats, which may require high resolution models; (2) deploying virtual BGC-Argo floats (van Sebille et al., 2018) and comparing them statistically to observations; or (3) forcing 1D biogeochemical simulations with observed physical fields, e.g. vertical mixing (Llort et al., 2015) or light (Terzić et al.,

680 2019). As a general rule, the good skill of the best PISCES 1D simulations (Fig. 5) indicates that our CATS approach can be used to tease apart model-observations misfits caused by the physical and biogeochemical components, opening up new avenues for parameter optimization (Falls et al., *submitted*) and model development.

Our cross-scale evaluation is also informative as to the spatiotemporal scales that can be addressed with a given model setup.
685 This matter was recently tackled by Bisson et al. (2019) using the export production model of Siegel et al. (2014), which is forced by satellite-derived primary production. In particular, they showed that this diagnostic model could be optimized to reproduce global climatological patterns but exhibited poor skill when faced with non-climatological datasets, which reflect local snapshots of ecosystem functioning. Model evaluation at climatological scales provides an incomplete picture, especially in productive regions where much POC export can take place during intense but short-lived events (Briggs et al., 2020). Such
690 events are smoothed out when climatologies are computed, and their coherence with the physical forcing is lost. Our work shows that a prognostic model like PISCES can afford both event-scale and climatological scale predictions. This capability is important to test our process-level understanding, which underpins climate change projections.

4.5 Joint use of BGC-Argo and models for process-level understanding

Properly representing POC stocks is crucial for constraining epi- and mesopelagic carbon budgets and, ultimately, estimating
695 the strength of the BCP and predicting its future evolution. The mismatch patterns between simulated and observed POC profiles (Fig. 4, 6 and 7) indicate different types of model shortcomings in subpolar and subtropical latitudes. The poorest agreement between PISCES and BGC-Argo data is found when their respective estimates of mesopelagic Teff are compared (Fig. 10), which should prompt further research on this key descriptor of the BCP, both in terms of POC fluxes (Buesseler et al., 2020) and stocks (Lam et al., 2011; this study), and their relationship with the structure and productivity of the upper ocean
700 ecosystem.

In the subpolar biomes, we find discrepancies between the patterns of SPOC and LPOC vertical export triggered by the intense surface phytoplankton blooms typical of these waters (Fig. 6, S5–S7 and S10). The observed rapid SPOC increase at depth cannot be explained by the SPOC gravitational sinking, and fragmentation of rapidly sinking LPOC has to be invoked (Briggs
705 et al., 2020). This fragmentation may be caused by zooplankton feeding (Mayor et al., 2020; Stemmann et al., 2004a, 2004b; Stukel et al., 2019) and swimming (Goldthwait et al., 2004), combined with bacterial hydrolysis of aggregate-binding polymers (Arnosti et al., 2012; Baltar et al., 2010a), and turbulence at high kinetic energy dissipation rate (Takeuchi et al., 2019). In relative terms, mesopelagic SPOC increases less strongly than LPOC during blooms (Fig. 6i), which is consistent with SPOC being a byproduct of the transformation of less abundant LPOC. Fragmentation processes may supply fresher SPOC to the
710 mesopelagic, enhancing the coexistence of suspended particles with variable freshness (Alonso-Gonzalez et al., 2010; Aumont et al., 2017) and overall contributing to POC remineralization (Giering et al., 2014; Mayor et al., 2020).

In the subtropical gyres and the most oligotrophic Mediterranean waters, PISCES underestimates TPOC between 0–1000 m except for the deep Chla maximum, where it overestimates TPOC. The prominent deep POC maximum simulated by PISCES is generally not found in observations from the STG biome (Fig. 7 and S9), where deep Chla maxima generally reflect phytoplankton photoacclimation, not enhanced phytoplankton biomass (Cornec et al., 2021). Thus, PISCES possibly overestimates the productivity of deep Chla maxima globally, indirectly causing stronger nutrient limitation at the surface. Between the deep Chla maximum and 200 m, POC pools decrease more steeply in PISCES than in BGC-Argo observations. Between 200 and 700 m, by contrast, simulated SPOC and LPOC Teff are only 10% lower than observations, well within observation uncertainty. Thus, the mesopelagic SPOC deficit simulated by PISCES in the STG originates mostly through the insufficient vertical POC export at 200 m depth. The Mediterranean represents a different case, whereby the large disagreement between PISCES and BGC-Argo Teff may result from either poorly constrained POC/b_{bp700}, simulated POC dynamics, or both. Thus, this region may provide a good testbed for studying the role of the mineral fraction in the BCP, including the controversial ballast hypothesis (François et al., 2002; Klaas and Archer, 2002; Passow, 2004).

In summary, the mismatch with observations suggests the need to improve the representation of SPOC–LPOC interconversion in PISCES. The size distribution of POC along the vertical axis is a key variable for constraining POC budgets, because it reflects the interplay between gravitational sinking, remineralization, trophic transfer and 3D dynamics including horizontal POC advection (Alonso-González et al., 2009; Boyd et al., 2019). In many instances (Fig. 9), our analysis suggests that better model performance in the mesopelagic may be achieved by increasing the net transfer of LPOC to SPOC, e.g. through LPOC fragmentation, and the Teff of both fractions. The vertical model-observations mismatch patterns observed here emphasize that POC budgets have to be computed with the highest vertical resolution affordable, or otherwise an apparent POC budget balance may result from compensating imbalances in different horizons (Giering et al., 2014; Marsay et al., 2015). The detailed level of information available from BGC-Argo floats may prove to be extremely valuable to help improve the POC schemes embedded in models such as PISCES.

5. Conclusions and outlook

In this study we compared globally distributed POC observations between 0–1000 m made by BGC-Argo floats to the predictions made by the PISCES model (PISCESv2_RC). A subset of BGC-Argo floats profiling at high vertical resolution enabled us to analyze small and large POC separately. The comparisons rely on a proposed new scheme for converting a bio-optical measurement (b_{bp700}) to POC. Although PISCES recreates with good skill the main features observed in subpolar and subtropical biomes, the comparison is still hampered by: (1) spatial and temporal variability in POC/b_{bp700} conversion factors, (2) mismatches in observed and simulated physics, and (3) imperfect correspondence between observed and simulated POC fractions. Evaluation of these uncertainties allowed us to detect limitations of the biogeochemical model parameterizations. Some limitations may be tackled by optimizing model parameters (e.g., particle sinking and remineralization rates), whereas

745 others may require changes in model structure, for example the representation of zooplankton feeding on, and transformation of, mesopelagic particles (Mayor et al., 2020). The descriptive work and model-data matching strategies presented here pave the way towards the use of BGC-Argo observations for data assimilation, parameter optimization (Falls et al., *submitted*) and, ultimately, model development.

750 Widespread use of BGC-Argo data for understanding POC budgets and the BCP can complement classical model constraints based on vertical POC fluxes and ocean-interior nutrient remineralization. Merging of BGC-Argo and satellite data streams through data-driven approaches —which allow for great flexibility—, and mechanistic models —which provide process-level understanding—, can soon provide us with a high-resolution 4D view of the oceanic carbon cycle. Further work is granted to investigate POC dynamics through combination of PISCES, autonomous observations, and ship-based observation programs

755 (e.g., GEOTRACES, Lam et al. (2015)) and data compilations (Mouw et al. ,2016; Evers-King et al., 2017).

Below we list several research priorities, whose implementation would advance the study of the biological carbon pump through the synergies between BGC-Argo observations and modelling.

- Observations:

760
  - High-resolution bio-optical profiles are indispensable for process-level understanding. Combination of low- and high-resolution sampling in separate or individual floats (e.g. burst sampling) may provide a good compromise between float lifetime and observation capabilities.
  - More in situ measurements are needed to constrain POC estimation from bio-optical variables, especially in meso- and bathypelagic waters. Addition of transmissometers may enable more accurate POC quantification

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    (Bishop, 1999) and particle characterization (Boss et al., 2015).
  - Inclusion of new types of sensors (Claustre et al. 2020) and extension of measurements into the bathypelagic (Deep-Argo; Roemmich et al., 2019) hold high potential for advancing BCP research.
  - Further developments in BGC-Argo data processing are needed, with the final goal of supplying a wide public with user-friendly data products in near real time.

770 - Models:
  - Evaluating model skill at resolving POC stocks, in addition to fluxes, is key to ensure that models reproduce observed fluxes for the right reasons.
  - Evaluation against globally consistent datasets is critical to avoid model overtuning towards small, sparse datasets, such as vertical POC fluxes.

775
  - Continuous development of schemes representing particle dynamics across the entire size spectrum is needed to constrain ecologically, climatically and economically relevant element fluxes.

- o Extension of prognostic modelling of bio-optical properties (Dutkiewicz et al., 2019) into the meso- and bathypelagic layers would enable direct matching with measurements made from autonomous platforms, facilitating their assimilation by models.

780 - • Joint planning of field observation and modelling projects, from their very conception and through their entire development, is key to fully exploit the capabilities of each approach.

785 *Data availability*. The simulated and observed datasets analyzed in this article are available at https://doi.org/10.5281/zenodo.5139602. The code and documentation of NEMO and PISCES are available at https://www.nemo-ocean.eu/. The authors can provide the code used to process the datasets on reasonable request.

Supplement. A supplement with 16 figures (Fig. S1–S16) and one table (Table S1) is available.

Author contributions. M.G. and R.B. designed the study. M.F. produced and/or reprocessed global climatological datasets. 790 R.B. produced the NEMO dynamical fields used to force PISCES 1D offline simulations. O.A. provided the global PISCES simulation. M.G processed BGC-Argo coherent annual time series, ran PISCES 1D simulations, analyzed data and produced the figures, and wrote the manuscript with contributions from all coauthors.

Competing interests. The authors declare no conflict of interest.

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Appendix A: Calculation of POC/b_{bp700} ratios and related optical considerations

805 POC/b_{bp700} ratios at the sea surface were obtained from the slope of the linear regression between POC and b_{bp700} (Table A1). Our approach essentially follows the literature compilation made by Cetinić et al. (2012). The linear regressions generally yielded small and non-significant y intercepts for sea-surface data. Therefore, we assumed the slopes were equal to the POC/b_{bp700} ratio. When the b_{bp} measurements were made at another wavelength, we converted them to b_{bp700} assuming an exponent $\eta = 0.41$ (Cetinić et al., 2012), according to the equation:

810 $$b_{bp}(\lambda_1) = b_{bp}(\lambda_0) \cdot (\lambda_1/\lambda_0)^{\eta} \tag{A1}$$

For measurements taken at 555 nm, which was the wavelength used in two studies (Table A1), this resulted in 10% lower b_{bp700} compared to b_{bp555}, thus higher POC/b_{bp700} ratios.

The dataset of Cetinić et al. (2012) was the only one showing a significant negative intercept of the POC vs. b_{bp700} linear
815 regression. Unlike the other datasets, where the POC vs. b_{bp700} relationship reflected mostly sea-surface variability in a given biome, this was the only dataset that included data collected between the sea surface and 600 m. Bol et al. (2018) reprocessed this dataset by computing the linear regression between POC and b_{bp700} in different depth bins, forcing the intercept through zero. The slope of the POC vs. b_{bp700} regressions, and therefore the POC/b_{bp700} ratio, decreased by more than twofold between the surface and the 600 m bins. These results suggest that the POC vs. b_{bp700} relationship may be better modelled with nonlinear
820 equations, as done for surface data in some previous studies (Balch et al., 2010; Johnson et al., 2017; Stramski et al., 1999). However, one must keep in mind that the ecosystem processes that define POC/b_{bp700} ratios in the surface layer may be different from those occurring in the mesopelagic (see Discussion, 4.2).

The relationship between POC and particulate beam attenuation at 660 nm, c_p, has been analyzed on more occasions than the
825 POC–b_{bp700} relationship. The backscattering ratio, b_{bp}/b_p, relates particulate backscattering to the total particulate scattering, and is directly related to the refractive index of the particle assemblage (Babin et al., 2003; Dall'Olmo et al., 2009; Loisel et al., 2011; Organelli et al., 2018; Stramski and Kiefer, 1991; Stramski et al., 1999; Twardowski et al., 2001; Ulloa et al., 1994). Light absorption by particles is negligible in the 650-700 nm spectral region, such that total beam attenuation c_p is a good approximation of the scattering coefficient b_p. Thus, once the known absorption and scattering coefficients of seawater are
830 removed, b_{bp}/c_p is a good approximation of the backscattering ratio and can be used to compare the relationships between POC vs. c_p and POC vs. b_{bp700}.

The observed POC/b_{bp700} variability in the surface layer across biomes, reflected in our POC estimation algorithm, seems analogous to that found for the relationship between POC and c_p by Cetinić et al. (2012) (Fig. A1). The POC content per unit
835 b_{bp700} decreases with the maximum b_{bp700} of each dataset, which may be related to the structure and species composition of

upper-ocean ecosystems (see Discussion, 4.3). It is also plausible that the fraction of POC (in terms of particle size or type) that contributes to backscattering varies across sites, further enhancing POC/b_{bp700} variability.

The number of studies that tackled the interconversion between POC and bio-optical proxies in the mesopelagic layer is much
840   smaller than those that focused on the epipelagic layer. Besides Cetinić et al. (2012) in the subpolar North Atlantic, we are only aware of the studies of Bishop (1999) and Bishop et al. (1999). The latter two studies found a POC vs. c_p slope of around 16 mmol C m^{-2} (= mmol C m^{-3} / m^{-1}). Given that modern transmissometers accept more forward scattered light that those used by Bishop (1999), the corresponding slope would be approximately 27 mmol C m^{-2} (J. Bishop, *personal communication*). Bishop (1999) found this slope to be nearly constant in contrasting areas over the 0–1000 m depth range, as depicted in Fig.
845   A1 with the datasets labelled "Bishop1999eqpac" (Equatorial Pacific) and "Bishop1999all" (Equatorial Pacific, NE Pacific and North Atlantic together).

The linear relationship between POC and c_p is compatible with the nonlinear relationship between POC and b_{bp700} along the vertical profile if the backscattering ratio also increases with depth, as found by Cetinić et al. (2012). The latter study found
850   an increase in the backscattering ratio (b_{bp700}/c_{p653}) from around 1.2% at the surface to around 1.5% in the upper mesopelagic. The b_{bp700}/c_{p653} ratio was more variable in deeper layers and values > 2% were not rare (see also Nencioli et al., 2010; Organelli et al., 2018; Twardowski et al., 2001). The POC/b_{bp700} ratio of 1000 mmol C m^{-2} used here for deep waters (Eq. 1), based on the analysis of Bol et al. (2018), would correspond to a POC vs. c_p slope of 27 mmol C m^{-2} if the backscattering ratio was 2.7% in the lower mesopelagic (600–1000 m, Fig. 2). Analogously, the range of POC/b_{bp700} used in our sensitivity tests, 500 to 2000
855   mmol m^{-2}, would correspond to backscattering ratio between 5.4% (likely too high) and 1.35% (closer to available estimates). More collocated measurements of POC, b_{bp700} and c_p in the mesopelagic and below are needed to reduce these uncertainties.

Beyond the natural variability, the interconversion between POC and bio-optical proxies is also confounded by methodological issues (Cetinić et al., 2012; Strubinger-Sandoval et al., 2021), most of which were not fully addressed in the studies compiled
860   here. In particular, filtration of large sample volumes in the m^3 range with in situ pumps yielded much stronger POC–c_p relationships than small volume sampling (Bishop, 1999).

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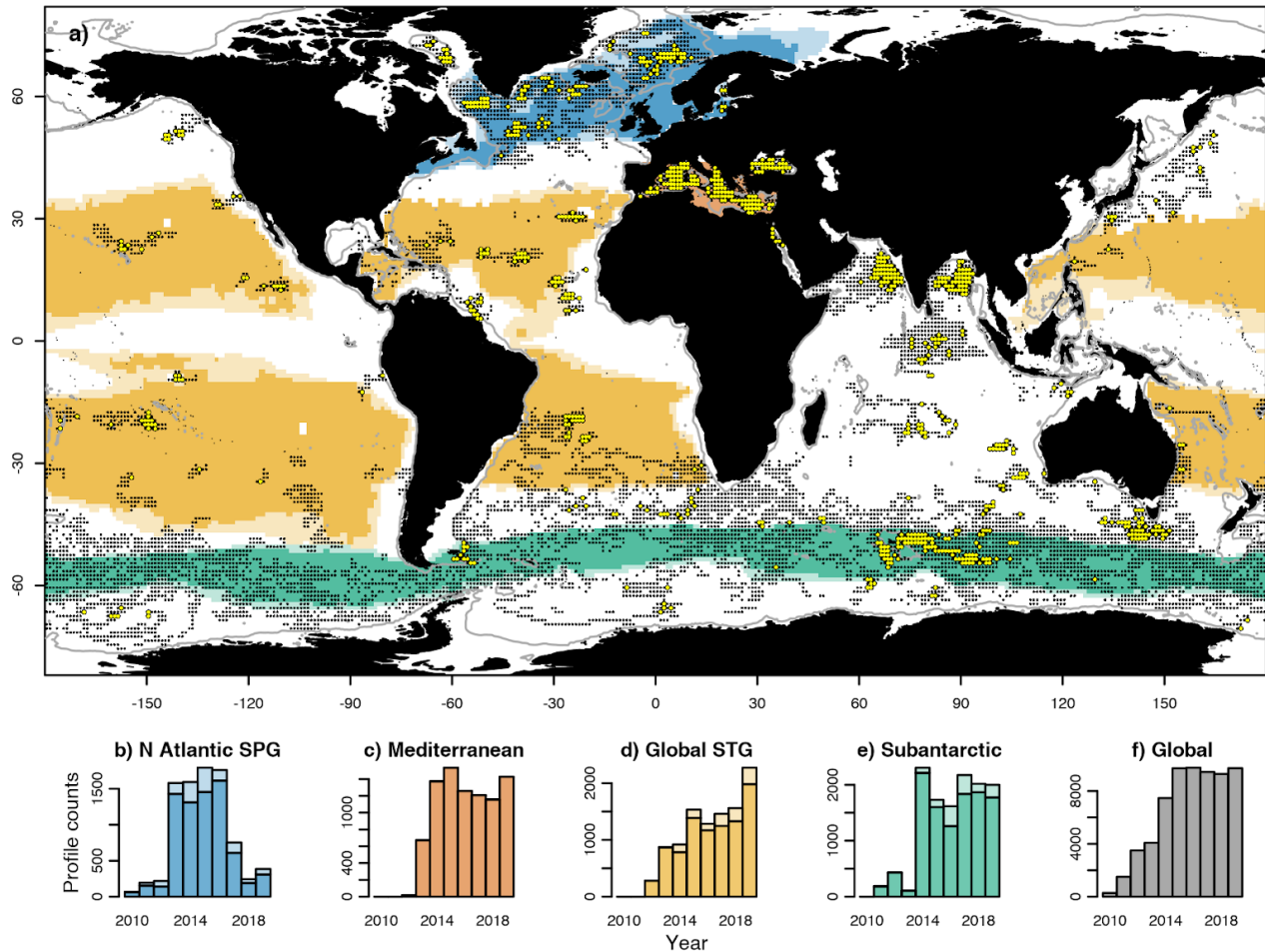
Figures

Figure 1. Global distribution and abundance of BGC-Argo profiles between 2010 and 2019. Grid cells (1°x1°) with at least 1 profile of the backscattering coefficient at 700 nm (b_{bp700}) are marked with black dots, and those with at least 20 profiles are marked with yellow-filled circles. The gray contours indicate the 1000 m isobath. Color shading indicates ocean biomes (see text), whose names are indicated on top of the bottom histograms. For each biome, light color indicates its average extent over 13 years of satellite observations (1998-2010), whereas the darker color indicates the "core" grid cells that never changed biome classification during the same period. Bottom panels (b-f) show the number of BGC-Argo bbp700 profiles per year in the four selected biomes and in the global ocean.
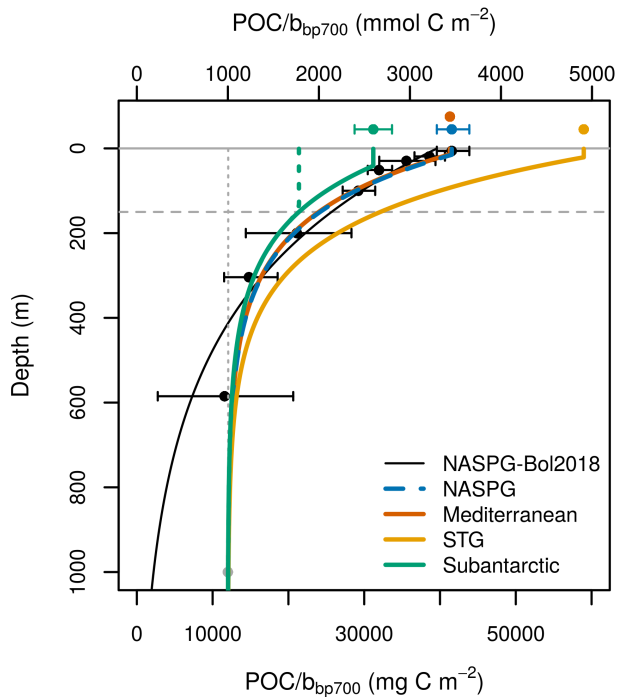
Figure 2. Empirical model used to convert the backscattering coefficient at 700 nm (bbp700) to particulate organic carbon (POC). Black dots and error bars show the dataset of Cetinić et al. (2012) as binned by Bol et al. (2018). The black curve shows the exponential fit of Bol et al. (2018) to the Cetinić dataset. The blue curve shows our fit to the same dataset (NASPG: North Atlantic Subpolar Gyre), forced to converge to a nonzero minimum value at depth. The remaining curves show similar functions with the same exponential slope ($b = -6.57$) as the NASPG fit, but with different surface values derived from the following studies: Loisel et al. (2001) for the Mediterranean Sea; Stramski et al. (2008) for the ensemble of subtropical and tropical areas excluding equatorial upwellings (STG); and Johnson et al. (2017) for the Southern Ocean (Subantarctic). The depth of the homogeneous surface layer, $z_{surf,biome}$ in Eq. 1 and 2, corresponds to the 5% quantile of the climatological MLD in summer in a given biome: 15 m in the NASPG, 14 m in the MED, 21 m in the STG and 41 m in the SO. The dotted green line, shown for the SO case only, illustrates the behavior of the algorithm for an hypothetical mixed layer depth of 150 m.
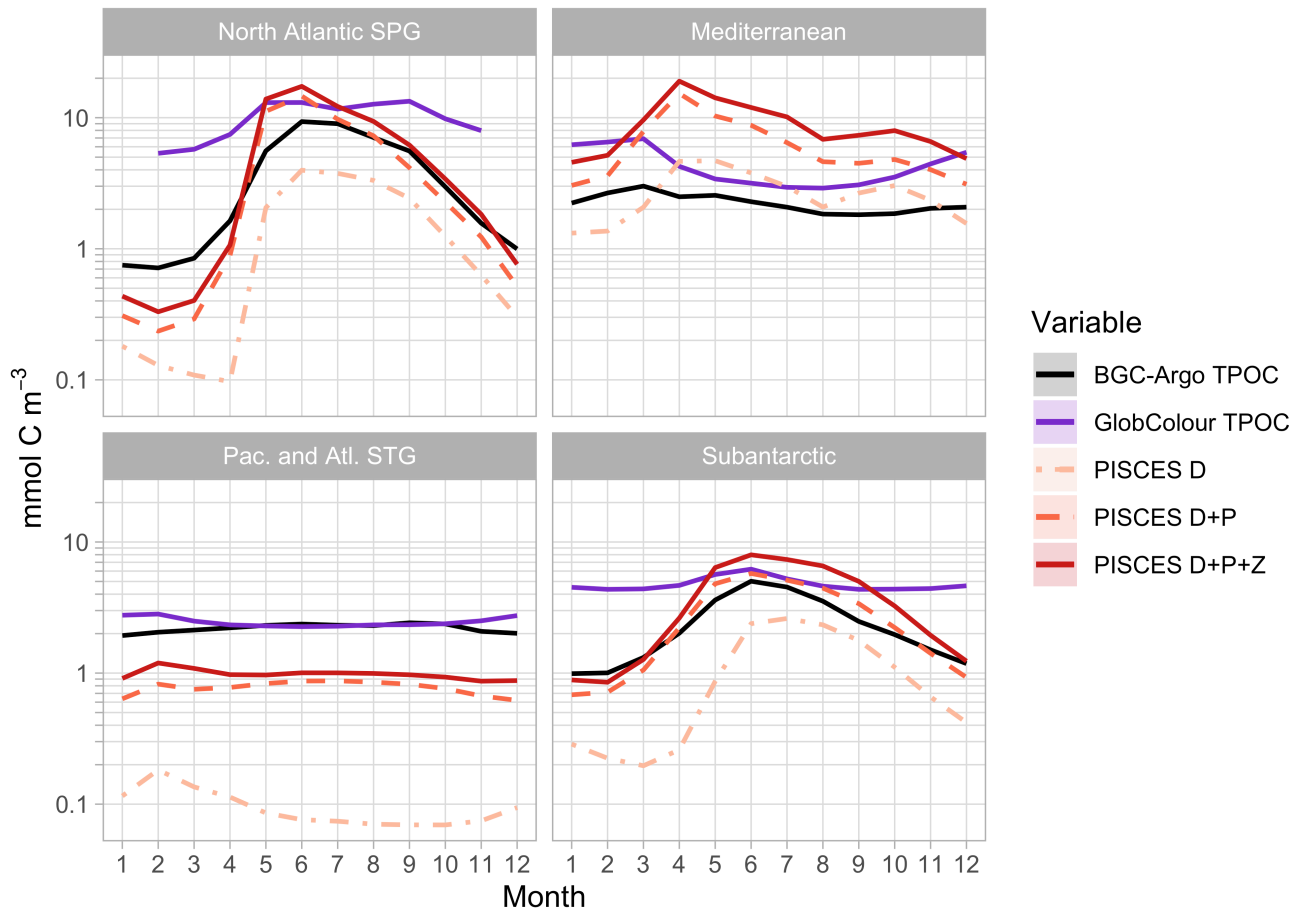
Figure 3. Monthly sea-surface particulate organic carbon (POC) concentration at the sea surface. Shown are POC estimates based on BGC-Argo (derived from the backscattering coefficient at 700 nm, b_{bp700}), satellite (GlobColour) and PISCES. PISCES TPOC results from the addition of the detritus (D), phytoplankton (P) and zooplankton (Z) tracers (Table 2), which are shown here as cumulative sums.

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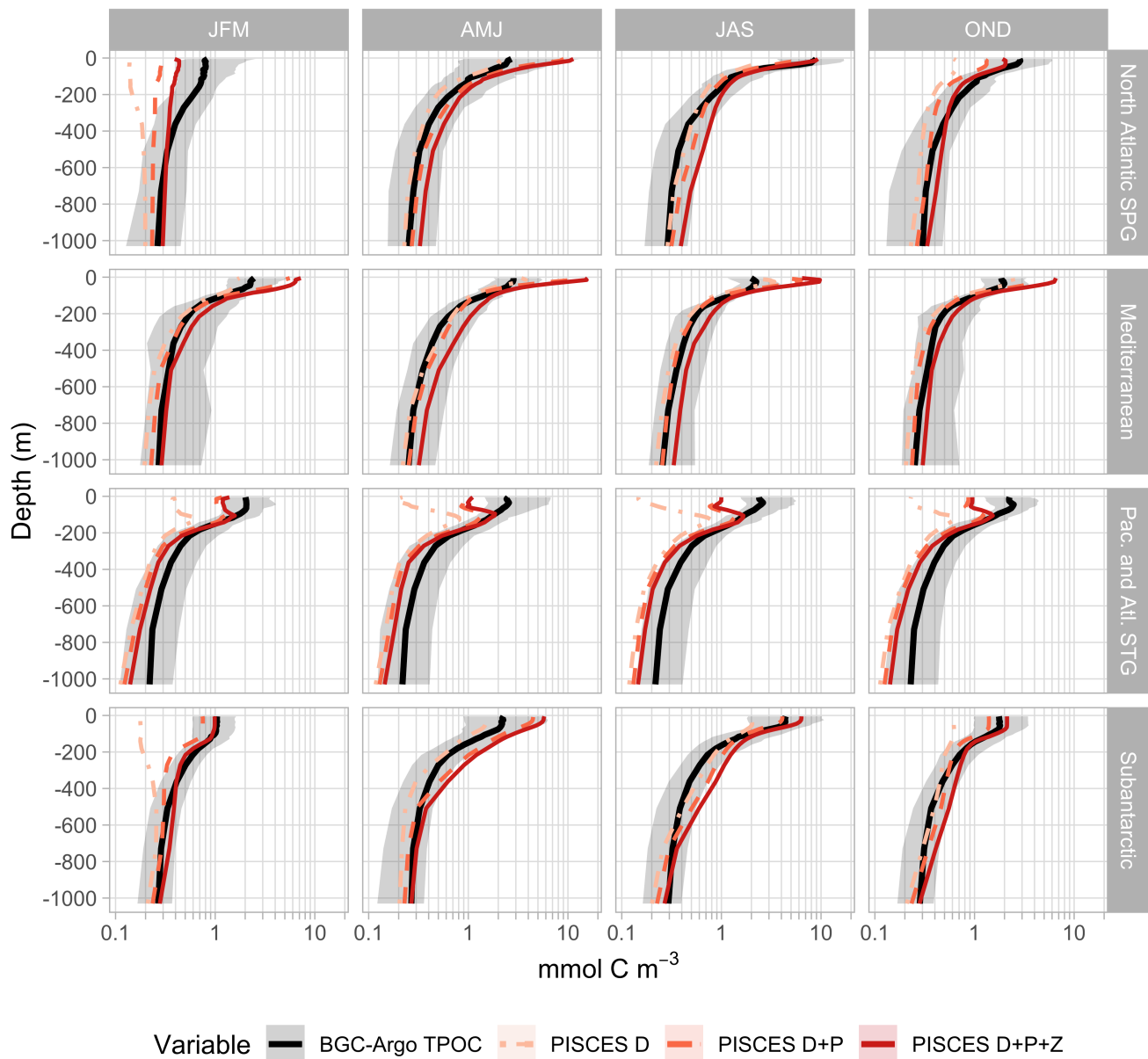
Figure 4. Seasonal 0–1000 m profiles of particulate organic carbon (POC). BGC-Argo estimates of TPOC (based on the backscattering coefficient at 700 nm, b_{bp700}) are represented with the median and the 0.025–0.975 quantiles within each biome. PISCES TPOC results from the addition of the detritus (D), phytoplankton (P) and zooplankton (Z) tracers (Table 2), which are shown here as cumulative sums.
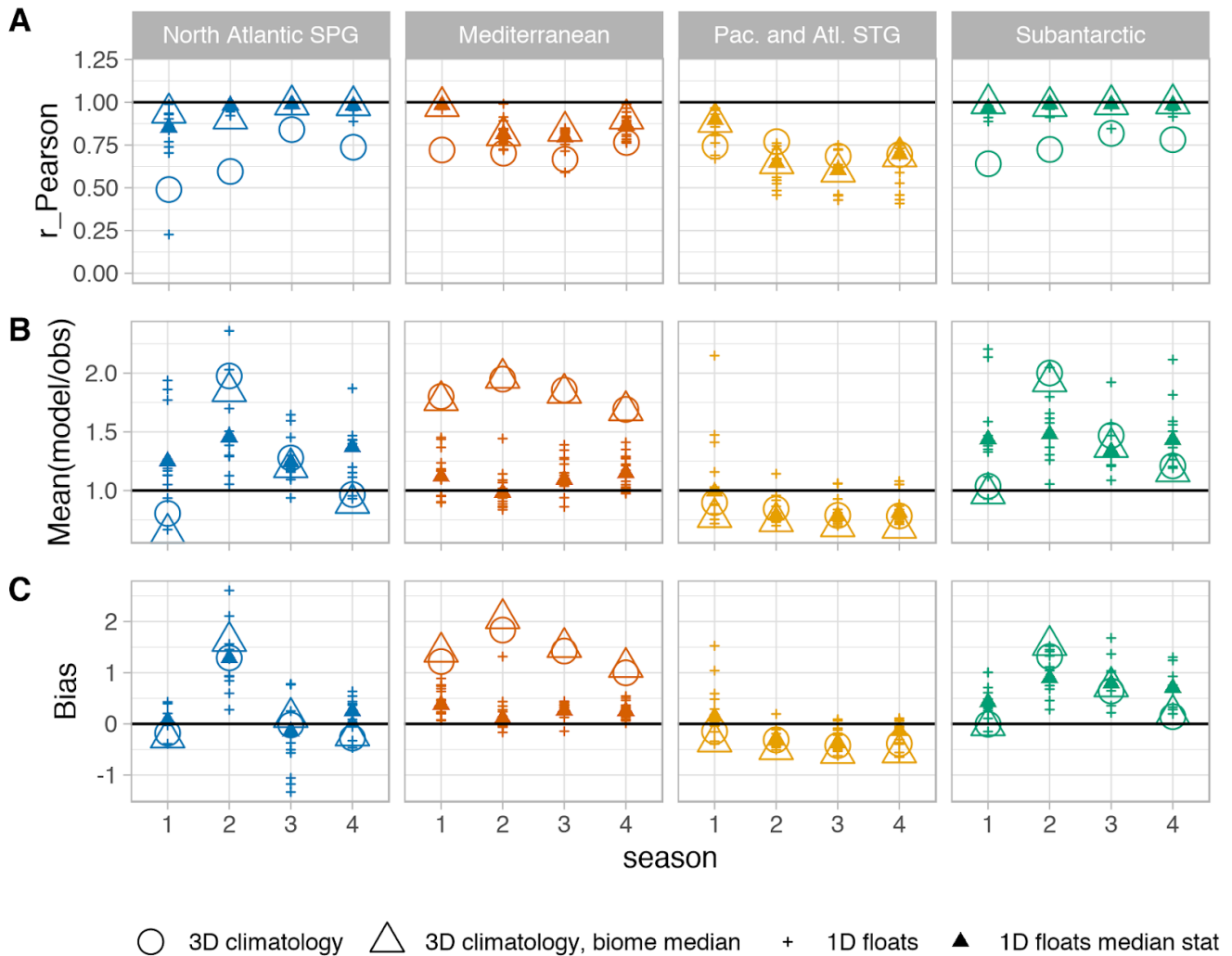
Figure 5. Summary of skill metrics for the comparison between PISCES and BGC-Argo particulate organic carbon (POC) profiles in different biomes and seasons. Pearson's correlation coefficient, mean model/data quotient, and mean bias (mmol m⁻³) are computed in linear space (untransformed data). Horizontal black lines show the benchmark values for each statistic. Symbols represent different types of simulations and data aggregation levels. In the case of the PISCES 3D climatological simulation, the statistics are computed at two spatial aggregation levels: using all the grid cells with matching BGC-Argo profiles in a given biome (circles); and after averaging all the documented grid cells within the biome (empty triangles), as displayed in Fig. 4. In the case of the PISCES 1D simulations, statistics are computed for each individual CATS separately (small crosses), and the biome-season median is also shown (small filled triangles).
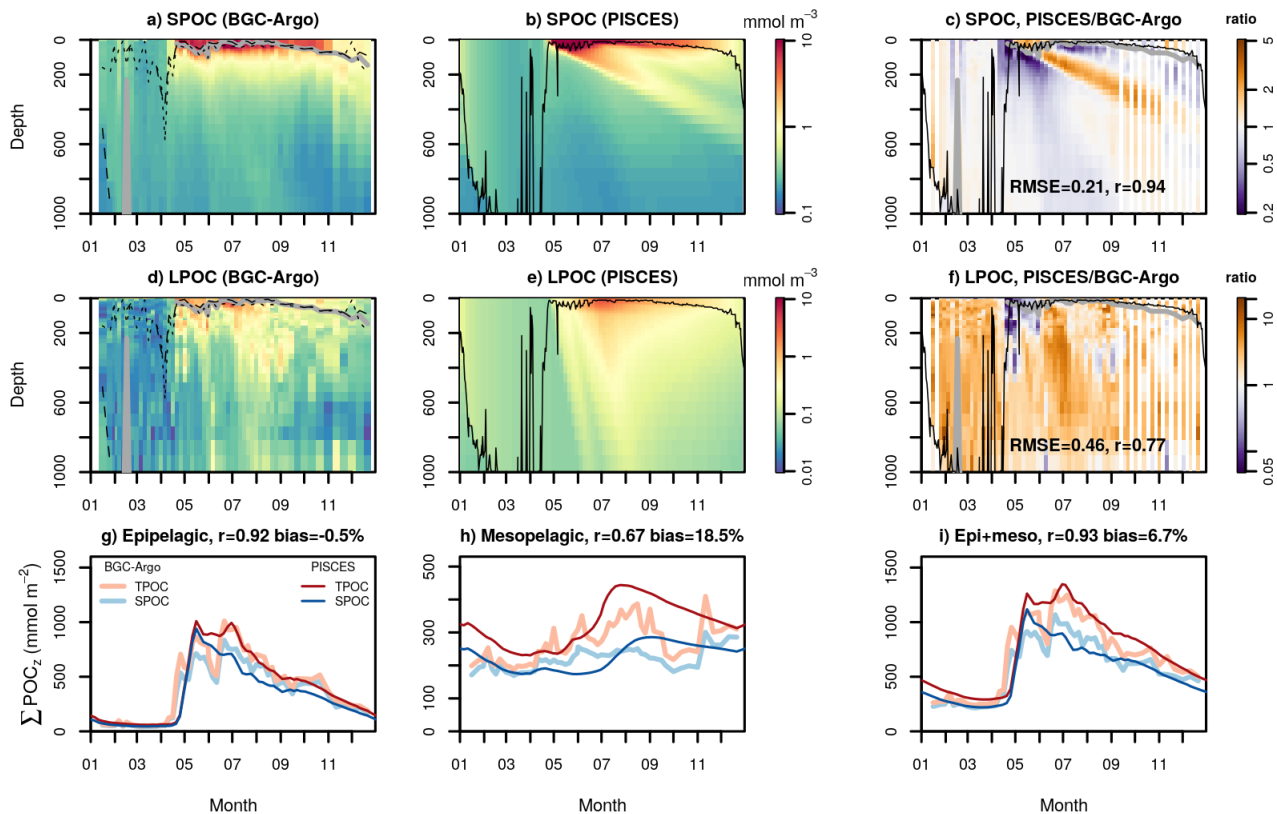
Figure 6. Temporal evolution of small and large POC concentrations (SPOC and LPOC, respectively) in the North Atlantic subpolar gyre (NASPG). Panels a and d show observations from the BGC-Argo float WMO 6901486, which drifted in the Labrador Sea during the year 2015. Panels b and e show the corresponding PISCES 1D simulation. Panels c and f show the ratio between the model and the observations and the corresponding correlation coefficient and RMSE in $\log_{10}$ scale. Solid lines depict the observed MLD (a and d, in gray) and the simulated turbocline depth (b and e, in black). In panels a and d, the dotted lines show the depths of the maximum Chla fluorescence gradient and the dashed lines an alternative MLD estimated with a 0.005°C temperature threshold that captures weak stratification. The bottom panels show the vertically integrated SPOC and LPOC stocks in different layers: 0–200 m or epipelagic (g), 200–1000 m or mesopelagic (g) and 0–1000 m (i). On top of panels g–i we show the correlation and mean bias between BGC-Argo and PISCES total POC (TPOC).

Figure 7. Temporal evolution of small and large POC concentrations (SPOC and LPOC, respectively) in the South Pacific subtropical gyre (STG). Panels a and d show observations from the BGC-Argo float WMO 6901660, which drifted westwards near Tahiti, during the year 2016. Panels b and e show the corresponding PISCES 1D simulation. Panels c and f show the ratio between the model and the observations and the corresponding correlation coefficient and RMSE in $\log_{10}$ scale. Solid lines depict the observed MLD (a and d, in gray) and the simulated turbocline depth (b and e, in black). In panels a and d, the dotted lines show the depths of the maximum Chla fluorescence gradient and the dashed lines an alternative MLD estimated with a 0.005°C temperature threshold that captures weak stratification. The bottom panels show the vertically integrated SPOC and LPOC stocks in different layers: 0–200 m or epipelagic (g), 200–1000 m or mesopelagic (g) and 0–1000 m (i). On top of panels g–i we show the correlation and mean bias between BGC-Argo and PISCES total POC (TPOC).

Figure 8. Mean annual POC stocks. BGC-Argo versus PISCES scatterplots are shown for the standing stocks (vertical integrals) of small POC (SPOC) and large POC (LPOC) in the epipelagic (0–200 m) and mesopelagic (200–1000 m) layers. Reference lines indicate a range of model:data ratios, from 1:1 (perfect correspondence) to a factor of 3 or its inverse. Biomes are distinguished with different colors, and the size of the circles is proportional to the temporal correlation between BGC-Argo and PISCES stocks at each location (as shown in the bottom panels of Figures 6 and 7).
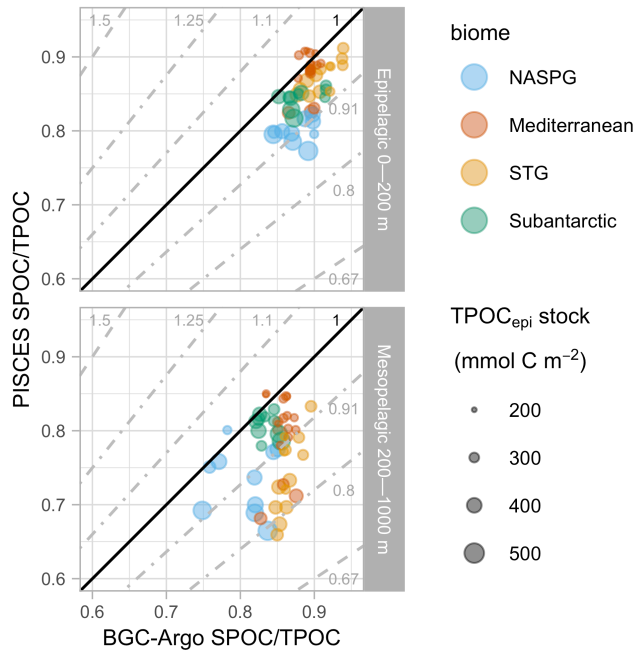
Figure 9. Mean annual SPOC/TPOC fractions. BGC-Argo versus PISCES scatterplots are shown for the epipelagic (0–200 m) and mesopelagic (200–1000 m) layers. Reference lines indicate a range of model:data ratios, from 1:1 (perfect correspondence) to a factor of 1.5 or its inverse. Biomes are distinguished with different colors, and the size of the circles is proportional to the annual mean TPOC stock in the epipelagic layer, used as an indicator of productivity.
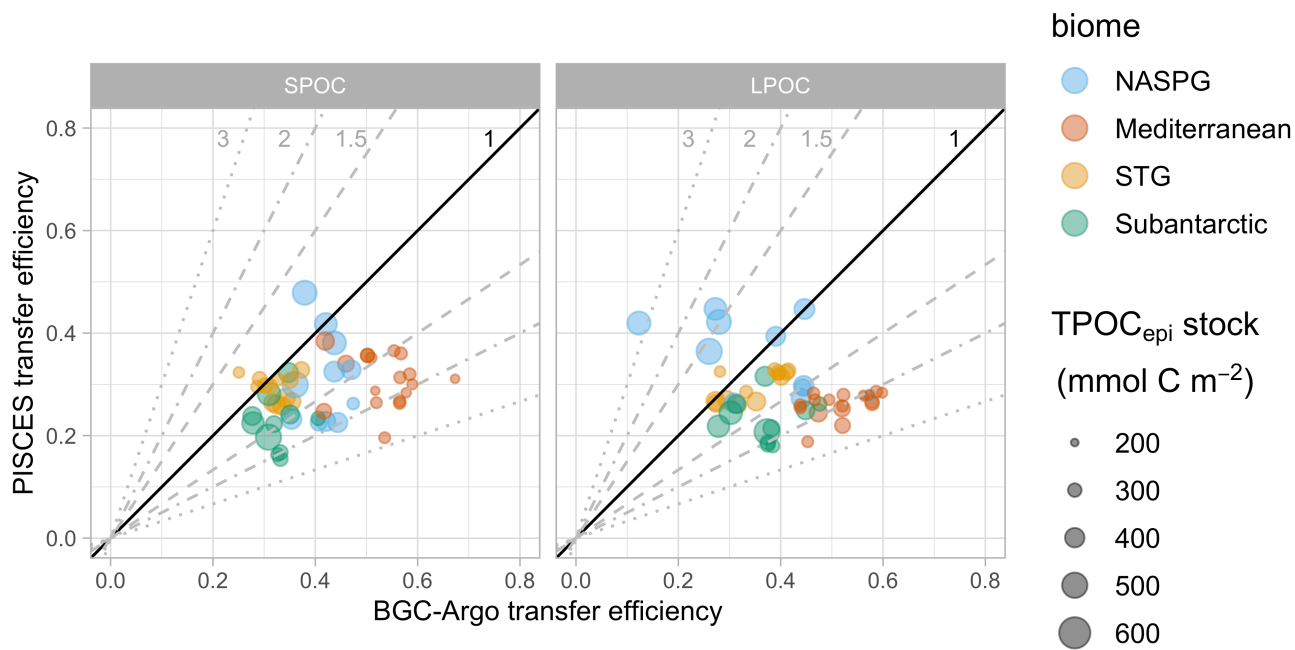
Figure 10. Mean annual POC transfer efficiency between 200 and 700 m. BGC-Argo versus PISCES scatterplots are shown for small POC (SPOC) and large POC (LPOC). Reference lines indicate a range of model:data ratios, from 1:1 (perfect correspondence) to a factor of 3 or its inverse. Biomes are distinguished with different colors, and the size of the circles is proportional to the annual mean TPOC stock in the epipelagic layer (0–200 m), used as an indicator of productivity.

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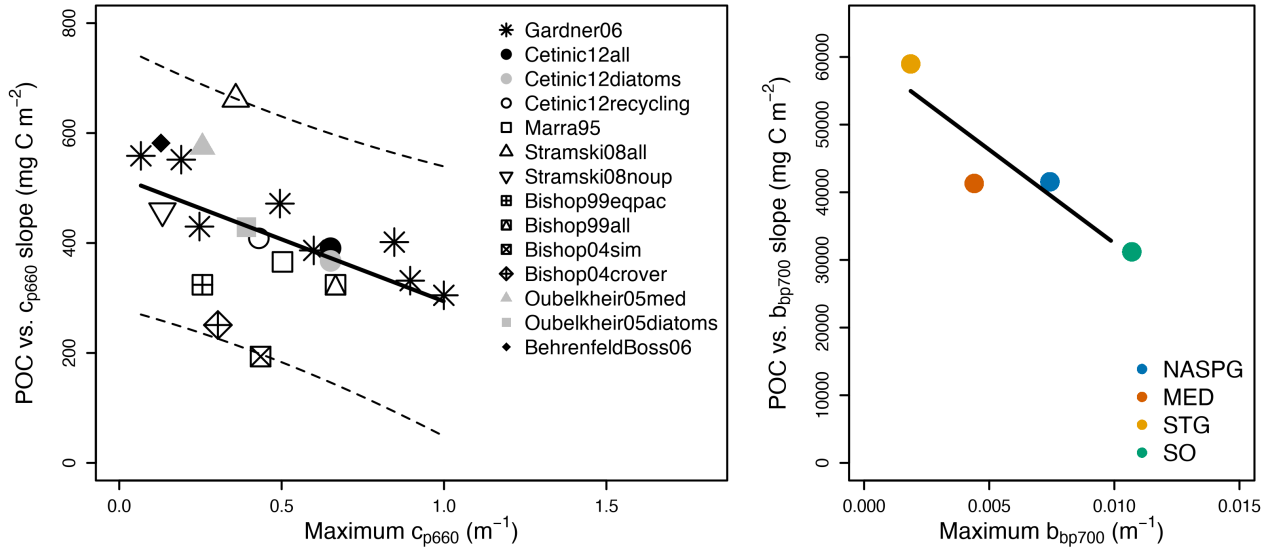
Figure A1. Coherent patterns in the conversion factors between POC and bio-optical variables. Left: relationship between the POC versus c_p slope and the maximum c_p in a given dataset, based on the figure 8 from Cetinić et al. (2012) with the addition of Bishop (1999) datasets. Right: analogous relationship for b_{bp700}, as used in our study. Linear fits are only illustrative.

Tables

Table 1. List of acronyms.

Type	Acronym	Definition
Region name	NASPG	North Atlantic Subpolar Gyre
	STG	Subtropical Gyres of the Atlantic and Pacific Oceans
Process or concept	BCP	Biological carbon pump
	CATS	Coherent annual time series
	MLD	Mixed layer depth
	Teff	Vertical transfer efficiency of POC
Operationally defined chemical compartment[a]	POC	Particulate organic carbon (used here as a generic name)
	SPOC	Small particulate organic carbon
	LPOC	Large particulate organic carbon
	TPOC	Total particulate organic carbon (here equivalent to SPOC + LPOC)
Bio-optical variable	b_{bp700}	Particulate backscattering coefficient at 700 nm wavelength
	Chla	Chlorophyll-*a* concentration estimated from fluorescence[b]
Numerical model or parameterization	NEMO	Nucleus for European Modelling of the Ocean
	PISCES	Pelagic Interactions Scheme for Carbon and Ecosystem Studies
	RC	Reactivity continuum parameterization for POC degradation
Project name	NAOS	Novel Argo Ocean Observing System
	remOCEAN	Remotely-Sensed Biogeochemical Cycles in the Ocean

[a] These variables may be estimated directly from seawater sampling, indirectly from in-situ or remote sensors, and from biogeochemical models. See Table 2.

[b] See text section 2.2 for details.

Table 2. Match between BGC-Argo observations, PISCES tracers (Aumont et al., 2017), real-world particulate organic carbon pools, and particle size ranges[a].

POC fraction	BGC-Argo observation	PISCES tracer (carbon currency)	Closest real-world correspondence
SPOC	b_{bp700} vertical profiles, despiked signal. Most sensitive to 0.5–30 μm particles (Dall'Olmo et al., 2009; Organelli et al., 2018) Calibrated as POC with Eq. 1 and 2	PHY: "Nanophytoplankton", includes calcifiers (default fraction 30%). Nominally 1–10 μm.	Picocyanobacteria and non-diatom phytoeukaryotes, generally 0.5–20 μm.
		PHY2: Silicifying microphytoplankton. Nominally 10–50 μm.	Diatoms, between 1.5–μm cells (Vaulot et al., 2008) and mm–scale chains. Mean ESD[b] generally < 50 μm in seawater (Snoejis et al., 2002; Ciotti et al., 2002; Bricaud et al., 2004).
		POC: detrital particulate organic carbon. Nominally < 100 μm. In practice, it includes heterotrophic prokaryotes' biomass with current parameter values (see 4.3).	Detrital particles between ~0.2[c] μm and 100 μm. In practice, measurements may include particle-attached and free-living organisms, viruses, colloids and adsorbed DOC.
		ZOO: Microzooplankton. Nominally 10–200 μm.	Microzooplankton. Mostly ciliates and flagellates with size similar to their prey, down to around 2 μm (Calbet 2008)
		Heterotrophic prokaryotes (BACT): currently not a prognostic tracer in PISCES. Not considered *explicitly* in this study (see 4.3).	Free-living heterotrophic prokaryotes (bacteria and archaea), < 1 μm.
LPOC	b_{bp700} vertical profiles, spike signal. Particle size between ~100 μm (b_{bp700} spike of $2.3 \cdot 10^{-5}$ m^{-1}) and ~2 mm (b_{bp700} spike of $8 \cdot 10^{-3}$ m^{-1}) (Briggs et al., 2020) Calibrated as POC with Eq. 1 and 2	GOC: detrital particulate organic carbon. Nominally > 100 μm	Detrital particles > 100 μm (aggregates, fecal pellets). Includes attached microbes.
		ZOO2: Mesozooplankton. Includes flux feeders[d]. Nominally 0.2–2 mm.	Mesozooplankton[d]

[a] See Stemmann and Boss (2012) for typical seawater particle size spectra.

[b] Equivalent spherical diameter

[c] Particles in the 0.2–0.8 µm size range and DOC are retained with variable efficiency by the filters commonly used to determine POC (Bishop, 1999; Cetinić et al., 2012; Graff et al., 2015; Lee et al., 1995; Morán et al., 1999; Strubinger Sandoval et al., 2021).

[d] PISCES mesozooplankton represents mostly copepods in the euphotic layer and flux feeders below it. The fraction of flux feeders is diagnosed in PISCES from the proportion between flux feeding rates and total mesozooplankton ingestion rates. By construction, flux feeding becomes the dominant mode of mesozooplankton feeding below the

euphotic layer under productive surface waters in PISCES. In reality, a wide variety of feeding strategies and organisms are found in the mesopelagic (Ikenoue et al., 2019; Kiørboe, 2011; Mayor et al., 2020; Stukel et al., 2019).

Table 3. Global and regional POC stocks, concentrations and fractions in different layers as simulated by PISCESv2_RC.

Variable	Depth range (m)	Open-ocean biomes					Global[d]
		Ice	SPSS[a]	STSS[b]	STPS[c]	Equatorial	
Area (%)		6.5	14.3	12.0	46.0	9.2	100
TPOC (Tg C)	0–200	73	281	263	263	154	1590
	200–1000	57	221	173	397	90	1006
	1000–5000	80	274	237	633	144	1438
TPOC (mmol C m^{-3})	0–200	1.18	2.22	2.57	1.66	1.99	1.85
	200–1000	0.29	0.48	0.44	0.25	0.29	0.33
	1000–5000	0.14	0.18	0.17	0.10	0.13	0.13
SPOC[e] (%)	0–200	72	81	79	89	84	81
	200–1000	75	75	70	79	71	75
	1000–5000	81	80	77	79	75	79
Phyto[f] (%)	0–20	45	45	41	58	49	50
Diatoms[g] (%)	0–20	28	21	14	3	4	14
Non-phyto (%)	0-20	55	55	59	42	51	50
Detritus[h] (%)	0–20	15	25	31	26	31	24
	0–200	36	43	48	44	50	42

| | 200–1000 | 62 | 67 | 70 | 78 | 78 | 70 |

[a] Subpolar seasonally stratified

[b] Subtropical seasonally stratified

[c] Subtropical permanently stratified

[d] Larger than the sum of biomes because the latter do not include coastal areas.

[e] SPOC / TPOC.

[f] $(PHY + PHY2)$ / TPOC.

[g] $PHY2$ / TPOC.

[h] $(POC + GOC + ZOO + ZOO2)$ / TPOC.

[i] PISCES detritus $(POC + GOC)$ divided by TPOC. Heterotrophic prokaryotes are *de facto* included in TPOC.

Table A1. Compilation of studies that reported linear regressions between POC and b_{bp700} in the euphotic layer of the oceans.

Location	Reference	N	Depth range	Slope (mg C m^{-2} m)	Intercept	Comments
North Atlantic Subpolar Gyre	Cetinić et al., 2012	321	0-600	35422 ± 1754	- 14.4 ± 5.8	Downcast
		321	0-600	43317 ± 2092	- 18.4 ± 5.8	Upcast
	Bol et al., 2018	n.a. [a]	0-10	41550	forced through 0	Subset of Cetinić et al., 2012 at the surface
Mediterranean	Loisel et al., 2001	N/A	N/A	41305	1.43	Original measurements at 555 nm
Subtropical and tropical Pacific and Atlantic (upwellings excluded)	Stramski et al., 2008	54	4-8	58968	2.75	Original measurements at 555 nm
Subantarctic	Johnson et al., 2017	67	0-100	31200 ± 2470	3.0 ± 6.8	

[a] n.a. = not available