Testing the reconstruction of modelled particulate organic carbon from surface ecosystem components using PlankTOM12 and Machine Learning

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14 Abstract. Understanding the relationship between surface marine ecosystems and the export of carbon to depth by 15 sinking organic particles is key to represent the effect of ecosystem dynamics and diversity, and their evolution under 16 multiple stressors, on the carbon cycle and climate in models. Recent observational technologies have greatly 17 increased the amount of data available, both for the abundance of diverse plankton groups and for the concentration 18 and properties of particulate organic carbon in the ocean interior. Here we use synthetic model data to test the potential 19 of using Machine Learning (ML) to reproduce concentrations of particulate organic carbon within the ocean interior 20 based on surface ecosystem and environmental data. We test two machine learning methods that differ in their 21 approaches to data-fitting, the Random Forest and XGBoost methods. The synthetic data is sampled from the 22 PlankTOM12 global biogeochemical model using the time and coordinates of existing observations. We test 27 23 different combinations of possible drivers to reconstruct small (POCs) and large (POCL) particulate organic carbon 24 concentrations. We show that ML can successfully be used to reproduce modelled particulate organic carbon over 25 most of the ocean based on ecosystem and modelled environmental drivers. XGBoost showed better results compared 26 to Random Forest thanks to its gradient boosting trees architecture. The inclusion of Plankton Functional Types (PFTs) 27 in driver sets improved the accuracy of the model reconstruction by 58% on average for POCs, and by 22% for POCL. 28 Results were less robust over the Equatorial Pacific and some parts of the high latitudes. For POCs reconstruction, the 29 most important drivers were the depth level, temperature, microzooplankton and PO₄, while for POC_L it was the depth 30 level, temperature, mixed-layer depth, microzooplankton, phaeocystis, PO_4 and chlorophyll *a* averaged over the 31 mixed-layer depth. These results suggest that it will be possible to identify linkages between surface environmental 32 and ecosystem structure and particulate organic carbon distribution within the ocean interior using real observations, 33 and to use this knowledge to improve both our understanding of ecosystem dynamics and of their functional 34 representation within models. 35

1. Introduction.

38 Progress in numerical ocean modelling over multiple decades coupled with fundamental knowledge of fluid dynamics 39 have led to an explicit representation of ocean dynamics in Earth System Models and of most of its key features, apart 40 from small-scale features which are parametrized. In contrast, ecosystem dynamics in ocean biogeochemical models 41 are much more reliant on empirical data for growth and loss processes, with the theoretical basis limited to the dynamic 42 representation of interactions among lower trophic levels (zooplankton and smaller organisms) and their influence on 43 carbon pools and fluxes (Le Quéré et al., 2005; Hood et al., 2006). The recent advances in observational technologies 44 including imaging data (Guidi et al., 2016), genomics (Kirchman et al., 2016), and field study (Mutshinda et al., 2017; 45 Batten et al., 2019, Lombard et al., 2019), offer new opportunities to improve our understanding of marine ecosystem 46 dynamics, and to better represent its influence on carbon pools and fluxes in models that are used to project future 47 climate change and associated impacts on ecosystems.

48 One strategy to represent lower trophic interactions in global biogeochemical models is to combine different species
 49 into Plankton Functional Types (PFTs) based on their unique influence on global biogeochemical cycles (Le Quéré et al., 2005; Hood et al., 2006). This approach enables the representation of plankton types that are unique, have an

- 51 influence on other PFTs within the ecosystem and are of quantitative importance for carbon flux and other
- 52 biogeochemical fluxes. The PlankTOM12 model is among the most detailed in this category of models with its
- 53 inclusion of an explicit representation of twelve PFTs: six phytoplankton, five zooplankton, and bacteria.
- 54 PlankTOM12 builds on the published version PlankTOM10 (Le Quéré et al., 2016) that has been extended to include
- 55 gelatinous zooplankton (Wright et al., 2021) and pteropods (Buitenhuis et al., 2019). Much effort has been put into 56 the development of PFTs and associated representation of surface ecosystem dynamics, which has led to the

57 demonstration that: (1) the representation of trophic levels was a key determinant of the low chlorophyll *a*

- 58 concentration observed in the Southern Ocean summer (Le Quéré et al., 2016); (2) CaCO₃ dissolution above the
- 19 lysocline is needed to reproduce observations of both biomass and export of PFT calcifiers, and (3) gelatinous
- 60 zooplankton plays an important role in determining surface biomass of other PFTs (Wright et al., 2021).
- 61 In contrast, the transfer of organic matter resulting from surface ecosystem dynamics into carbon exported to the deep 62 ocean via the sinking of particulate organic matter has received much less attention, so that improvements in the 63 representation of the PFTs do not necessarily translate into improvements in sinking of particulate matter (Wright et 64 al., 2021). The export flux of particulate organic carbon from the surface ocean to depth is around 10 PgC yr⁻¹ 65 (Schlitzer, 2002), which is as large as the CO₂ emitted to the atmosphere by human activities and nearly four times 66 larger than the mean oceanic CO₂ sink in recent decades (Friedlingstein et al., 2022). Changes in carbon exported to 67 depth can have a large impact on air-sea CO₂ fluxes and on the amount of CO₂ emissions that remain in the atmosphere 68 where they cause climate change.
- 69 The growing amount of observations provides the opportunity to develop a new approach to explore the linkages
- between surface ecosystem dynamics and the distribution of particulate organic carbon in the ocean, and to improve

71 the representation of particle sinking fluxes in models. However, there is a risk of over-interpreting the data by 72 applying Machine Learning (ML) methods directly to link the observed surface environment and ecosystem structure

applying Machine Learning (ML) methods directly to link the observed surface environment and ecosystem structure
 with the observed particulate organic carbon distribution. The use of synthetic observations based on model data

75 with the observed particulate organic carbon distribution. The use of synthetic observations based on model 74 therefore provides a minimum test to assess the likely success and usefulness of such an approach.

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ML has been widely used in biogeochemical and geophysical applications and provided efficient results in
reconstructions of ocean surface pCO₂ (Friedrich and Oschlies, 2009; Telszewski et al., 2009; Landschützer et al.,
2013; Denvil-Sommer et al., 2019) and of particulate organic carbon (Sauzède et al., 2016, 2017) as well as in the
analysis of driver importance (Sauzède et al., 2020).

79Here we use model data to verify the hypothesis that the composition of surface ecosystems and environmental80conditions are indeed reflected in the abundance and size of the organic particles in the ocean interior. We reconstruct81the concentration of organic particles as represented by small (POCs, particles < 256µm) and large (POCL, particles >82256µm) particulate carbon in the PlankTOM12 model. Using this information alongside with modelled environmental83and ecosystem conditions we develop a ML method to reproduce POCs and POCL over the global ocean and verify84the hypothesis. This constitutes a necessary although not sufficient test that the approach can subsequently be used to85reveal linkages using real observations and to inform model developments.

86 2. Data and Methods.

87 In this section we describe a set of variables that will be used to test the ML method's ability to reconstruct particulate 88 organic carbon concentrations based on ocean model data. We create a set of synthetic data by sampling a model at 89 the time and location of real-world observations. We discuss the availability and distribution of real-world 90 observations and their limitations. In this section we also describe the PlankTOM12 global ocean biogeochemical 91 model and how we use it to develop a ML method and test its ability to reconstruct small and large particulate organic 92 carbon with a limited number of observations. To provide resemblance to the real data availability we focus on the 93 period 2009-2013 which guarantees additional sampling of co-located biological, chemical, and environmental 94 variables from the Tara expeditions (Sunagawa et al., 2020).

95 Two sets of data are needed to test the Machine Learning method: a set of targets and a set of drivers. The drivers 96 represent the input variables to the ML method (here the biological, chemical, and environmental variables). The 97 targets represent the variables we are trying to reconstruct (here the particulate organic matter POCs and POCL). The 98 ML will then determine the relationship between the drivers and targets, which can then be applied in regions where 99 drivers are available to infer targets where the later data do not exist.

2.1.1. Measurements of particle size distributions and concentrations (the targets).

We use observations of particle distribution in two ways. First to determine the time and location of the observations,
and second to verify that the ocean model is of sufficient quality to be used in this analysis. The sampling of the
particulate organic carbon concentration is based on the data from an Underwater Vision Profiler 5 (UVP5) (Gorsky
et al., 2000, 1992; Picheral et al., 2010; Kiko et al., 2022). UVP5 measures particles of size from 50 µm to a few mm.
For the purpose of comparing the UVP5 data with the PlankTOM12 model data, we converted measured biovolume
concentration (mm³/L) of particles to carbon biomass concentrations (µmol/L) using the empirical equation from
Alldredge (1998) for particulate organic carbon:

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Figure 1. Location of the observations from the UVP5 database over the period 2009-2013. Green dots correspond to Tara
 expeditions, and were included in the global UVP5 database.

We summed size classes from 50.8 µm to 256 µm for the small particulate organic carbon (POC_s) and from 256 µm to 5.16 mm for large particulate organic carbon (POC_L). POCs below 100 µm are not well captured by the UVP sensor, which therefore underestimates this size-class of aggregated particles. We extrapolated the total size of particles up to 0.001 mm by using the size spectra theory to provide a better estimate of POC biomass concentration in line with the model. Following Guidi et al. (2008) we used the abundance of particles sized from 0.250 to 1.5 mm excluding rare particles to estimate the coefficients of logarithmic relationship between the size of particles and its abundance:

$$log(abundance) = a * log(size) + b$$

121 Using this equation we estimated the abundance of particles of size less than 100 μm.

122 There are 2603 vertical profiles of UVP5 measurements during 2009-2013, including 752 profiles which are colocated with the stations from the Tara expeditions that provide the environmental and ecosystem variables (Figure 1; Section 2.1.2). The measurements are sparse in time and space. There are no measurements in the Southern Ocean, Western Pacific Ocean and Eastern Indian Oceans.

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2.1.2. Measurements of environmental and ecosystem variables (the drivers).

127 We use observations of environmental and ecosystem variables to determine the time and location of the observations 128 that are colocated with the target variables. To represent the main physical and chemical drivers responsible for the 129 concentration and variability of POC_s and POC_L we use measurements of ocean temperature, chlorophyll *a*, phosphate

130 PO₄, nitrates NO₃, mixed-layer depth (MLD). These variables were measured during Tara expeditions along with the

particle size distributions and concentrations using UVP instruments onboard these cruises. However, chlorophyll *a*, PO4, and nitrates were not measured systematically at each depth level. Thus, their averages over MLD are tested as

133 possible drivers as well. To represent the biological drivers, we use information on PFTs.

134 2.1.3. The NEMO-PlankTOM12 Global biogeochemical model.

135 We used the output from the NEMO-PlankTOM12 coupled physical-biogeochemical model of the global ocean at

daily and monthly time resolution. NEMO represents physical transport processes and is used in its v3.6-ORCA2 version, with a horizontal resolution 2° longitude and 0.3° to 1.5° latitude, and 31 vertical levels. It is forced by daily

version, with a horizontal resolution 2° longitude and 0.3° to 1.5° latitude, and 31 vertical levels. It is forced by daily
 meteorological data from NCEP reanalysis (Kalnay et al., 1996) over the period 1948-2020, with output for 2009-

139 2013 used here. This model version is identical to that used to estimate the ocean CO₂ sink in the Global Carbon

140 Budget 2021 annual update (Friedlingstein et al. 2021).





Figure 2. Schematic representation of the flow of matter in and out of the two particulate organic carbon (OC) components
of the PlankTOM12 marine ecosystem model. The various boxes represent: Phyto - phytoplankton that includes diatoms
(DIA), mixed phytoplankton (MIX), coccolithophore (COC), picophytoplankton (PIC), phaeocystis (PHA) and N₂-fixers
(FIX); PRO - protozooplankton, PTE - pteropod, MES - mesozooplankton, MAC - macrozooplankton, GEL - gelatinous

146 zooplankton, BAC - bacteria.

PlankTOM12 represents ecosystem dynamics based on the representation of 12 PFTs: diatoms (DIA), mixed
phytoplankton (MIX), coccolithophore (COC), picophytoplankton (PIC), phaeocystis (PHA), N₂-fixers (FIX), microor protozooplankton (PRO), pteropod (PTE), mesozooplankton (MES), gelatinous zooplankton (GEL), and bacteria
(BAC). PlankTOM12 keeps track of the carbon biomass (µmol/L) of these PFTs over model depth levels resulting

151 from environmental and ecosystem processes and their interactions (Le Quéré et al. 2016).

152 PlankTOM12 represents sinking processes through the explicit representation of two organic particle of different size,

with small particles sinking at a constant speed of 3 m/d, and larger particles sinking at a variable speed between 3 and 150 m/d depending on the ballast effect of their mineral content (Buitenhuis et al., 2013). In addition, a dissolved

154 and 150 m/d depending on the ballast effect of their mineral content (Buitenhuis et al., 2013). In addition, a dissolved 155 organic carbon component is transported via ocean currents. Particles are generated through mass flux from the PFTs

organic carbon component is transported via ocean currents. Particles are generated through mass flux from the PFTs resulting from mortality and egestion and from aggregation through differential sinking or turbulent coagulation, and

destroyed through grazing by zooplankton and remineralisation by bacteria and through disaggregation from shear

158 currents. Large PFTs contribute mostly to POC_L, while small PFTs contribute mostly to POC_S. (Le Quéré et al. 2016;

159 Fig. 2).

The NEMO-PlankTOM12 model output was sampled at the time and location identified from the observations mentioned above to create a synthetic dataset. The model grid-coordinate closest to the real geographical position was chosen. If several measurements were co-localised at the same grid coordinate and same time step (day for daily PlankTOM12 and month for monthly PlankTOM12 outputs), it is counted as one measurement. This model sampling produced 400 positions when using the daily or monthly PlankTOM12 outputs. All drivers and targets were taken from the model output at the corresponding coordinates up to 1400 m depth. These outputs served as the reference for validation and evaluation of the ML methods and for establishing the sets of the most important drivers.

167 2.2. Method.

168 We tested 2 ML methods that are widely used in target's reconstruction based on tabular data sets: the Random Forest 169 regressor and the XGBoost (Extreme Gradient Boosting) regressor. The Random Forest (RF) regressor is an ensemble 170 algorithm that contains a number of decision trees on various subsets of the given dataset and takes as output the 171 average of prediction from each tree estimator. RF can run several trees at the same time allowing a use of a large 172 number of input variables, and it is robust to overfitting (Biau, 2012). XGBoost (XGB) regressor is an effective tree-173 based ensemble learning algorithm (Chen and Guestrin, 2016). It builds several models sequentially where each new 174 model attempts to correct errors from the previous one. XGBoost uses the gradient descent algorithm to minimise the 175 loss function of the model. Using RF and XGBoost we can estimate the driver importance to identify which driver has 176 the greatest impact on the predictions. To check the driver importance, we use drop col feat imp python function 177 (https://gist.github.com/erykml/6854134220276b1a50862aa486a44192). This method estimates how the accuracy of 178 the ML output changes if one of the drivers is dropped off from a driver set (DS) based on the training dataset.

179 Effective ML algorithm requires sets of training, validation and test data. The training data builds up the ML model. 180 Model evaluates training data repeatedly to learn about the relationship between inputs (driver set) and known outputs 181 (target set) and adjusts itself to better represent the target. The purpose of validation data is to evaluate the model 182 during its training by introducing new unseen data. It allows us to evaluate how a developed model works on a new 183 dataset and to optimise hyperparameters. The test data evaluate the final accuracy of the ML model and confirm that 184 the model works correctly on any unseen data. It is new data that did not participate in the training algorithm. The 185 accuracy is worse on validation and test data compared to training data set. The difference in model performance on 186 training and validation data can signal an overfitting, while this difference between validation and test data can 187 demonstrate an effect of data mismatch. It is worth noting that RF does not necessarily need validation data set as they 188 perform internal validation. During the training algorithm each tree is constructed from a random subset of original 189 data, usually it represents two thirds of data and one third of data is used to estimate out-of bag error to assess model 190 performance. XGB uses a validation data set to evaluate the model during training and to prevent overfitting by 191 applying an early stopping. In the present study the available data were split into training and validation data sets (Fig. 192 3a). Validation data is not included in RF training, however we use it to test the performance of trained RF and tune 193 hyperparameters afterwards. The test data are taken from the regions where there are no observations (Fig. 3b): 3 194 months for each year from the period 2009-2013 and 6 positions for each month were chosen randomly. This will 195 allow us to identify the possible accuracy of reconstruction that can be reached in these regions when we will apply a 196 developed method to real observations. However, when POCs and POCL will be reconstructed using only real-world

197 observations, we will need to split all available data into training, validation and test data sets.



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 Figure 3. The spatial distribution of: (a) - training (blue) and validation (green) data sets; (b) - test data set; based on PlankTOM12 monthly outputs.

201 We RandomForestRegressor function from scikit-learn (https://scikituse 202 learn.org/stable/modules/generated/sklearn.ensemble.RandomForestRegressor.html) with its default parameters and 203 min sample leaf equals 20. To apply XGBoost regressor we use XGBRegressor from xgboost 204 (https://xgboost.readthedocs.io/en/stable/python/python intro.html). Parameters were follows set as 205 n estimators=2000, max depth=7, eta=0.01, subsample=0.7, colsample bytree=0.8, gamma=0.01 for POC_L and 206 gamma = 0.3 for POCs, early stopping rounds = 10.

We tested 27 driver sets (DSs) that are summarised in Table 1. For each DS we identify the most important drivers that influenced the reconstruction of small (POCs) and large (POCL) particulate organic carbon concentration. The drivers include geographic variables (depth, sin(latitude), cos(longitude)), physical variables (incident light, MLD, co-located temperature), chemical variables (PO4, NO3, including co-located values and averages over the MLD), and biological variables (chlorophyll *a*, 12 PFTs listed above: DIA, MIX, COC, PIC, PHA, FIX, PRO, PTE, MES, GEL, BAC, including co-located values and averages over the MLD).



Driver set	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27
depth																											
Sin(lat)																											
Sin(long)																											
Cos(long)																											
Incident light																											
MLD																											
Temperature vp																											
CHL vp																											
CHL mean																											
NO ₃ vp																											
PO ₄ vp																											
NO ₃ mean																											
PO ₄ mean																											
BAC vp																											
MES vp																											
PTE vp																											
DIA vp																											
COC vp																											
PIC vp																											
PHA vp																											
GEL vp																											
PRO vp																											
MAC vp																											
MIX vp																											
FIX vp																											
BAC mean																											
MES mean																											
PTE mean																											
DIA mean																											

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COC mean														
PIC mean														
PHA mean														
GEL mean														
PRO mean														
MAC mean														
MIX mean														
FIX mean														
Big Zoopl														
Big Zoopl2														
CHL back vp														
CHL back mean														
GEL back vp														
PRO back vp														
MAC back vp														
COC back mean														

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The driver sets can be split into 9 thematic groups which together test the role of PFTs and sub-classes within, the role of surface versus depth profiles for some variables, and the role of information from the previous month:

- I. <u>No PFTs</u> (short name (sh.n.) 'No PFT'): Driver sets 1 and 2 do not include any PFTs and focus on the influence of temperature, MLD, chlorophyll *a*, NO₃ and PO₄ on POC₅ and POC_L reconstruction.
- 219 220 Introduction of PFTs (sh.n. 'PFT introduction'): DSs 3, 4 and 5 are dedicated to the investigation of the II. 221 introduction of PFTs in the reconstruction. In DS 3 we introduced 12 PFTs vertical profiles, even though this 222 information will be challenging to reproduce with observations due to the lack of data. Nevertheless, it is 223 important to test the capacity of ML if all 12 PFTs were available over the depth. DS 4 includes the vertical 224 profiles of 6 heterotrophs (zooplanktons and bacteria) because they contribute to influencing the vertical 225 distribution of POCs and POCL, and 6 phytoplankton averaged over MLD because they are responsible for 226 primary production. In DS 5 we added averages over MLD of the 6 heterotrophs that were not included in 227 DS 4.
- III. Big zooplankton (sh.n. 'Zooplankton combined'): In DSs 6 and 7 we tested the influence of big zooplanktons summed into one variable to account for their combined effect rather than the distinctions among PFTs. The big zooplankton is represented by the sum of mesozooplankton, gelatinous zooplankton and macrozooplankton in DS 6, with the addition of pteropod in DS 7.
- IV. <u>Exclusion of bacteria</u> (sh.n. 'No vertical BAC'): DS 8 does not have a bacteria (BAC) vertical profile compared to set 5.
- V. <u>Individual zooplankton types</u> (sh.n. 'Individual PFT'): DSs 9, 10, 11, 12, 13 and 14 test the influence of individual types of heterotrophs, bacteria (BAC), microzooplankton (PRO), pteropod (PTE), mesozooplankton (MES), gelatinous zooplankton (GEL), microzooplankton (MAC), respectively.
- VI. <u>Geographical position and seasons</u> (sh.n. 'Lat-Long' and 'Incident light'): DS 15 is based on DS 5 (which showed the most promising results) and includes geographical coordinates as additional drivers in the form of sin(lat), sin(long), cos(long). DS 16 includes in addition to the DS 5 the role of incident light.
- VII. <u>Use of only PFTs and chlorophyll *a* (sh.n. 'PFT only + CHL'): DS 17 is based on only the 12 PFTs, while
 DS 18 is formed from DS 17 plus information on chlorophyll *a* averaged over the MLD. DSs 19 and 20 are
 based on DS 6. To form the DS 19 we exclude temperature, NO₃ and PO₄ from the list of drivers in DS 6.
 DS 20 is an extended version of DS 19 with all 12 PFTs concentration averaged over the MLD.
 </u>
- VIII. <u>Chlorophyll *a* and chemical variables</u> (sh.n. 'Biochemical variables'): DSs 21, 22, 23, 24 are based on DS 5 and test the individual influence of chlorophyll *a* (DS 21), NO₃ (DS 22), PO₄ (DS 23) vertical profiles and its ensemble (DS 24).
- IX. Previous time step (sh.n. 'Month 1'): DSs, 25, 26 and 27 investigate the role of chlorophyll a (DS 27) and some zooplanktons from the previous time step: gelatinous zooplankton and microzooplankton (DS26); gelatinous zooplankton, micro- and macrozooplankton, averaged over MLD chlorophyll a and coccolithophore (DS25).

The evaluation of the method is based on the mean correlation coefficient, total root-mean square errors (RMSE), and total absolute bias between the ML outputs and PlankTOM12 POC_s and POC_L components. Moreover, we provide the global maps of correlation coefficient and RMSE to vertical profiles of POC_s and POC_L at each grid point. Global maps help to identify zones where the large errors can be hidden in the mean diagnostics due to the error compensation.

255 **3.** Results.

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257 3.1. Data analysis.

258 In this study we test the capacity to reconstruct particulate organic carbon from sparse observations by using ML and

a synthetic data set based on the PlankTOM12 model output. We compare observations and the output of the ocean model to provide a minimum of validation for the model data and to help explain differences in ML results when

applied to real observations in the future.

262 Figure 4 shows the vertical profile of small (POCs) (Fig.4a) and large (POCL) particulate organic carbon (Fig.4b)

based on the median from observations (green) and from daily PlankTOM12 model output (blue). Shading

corresponds to values between 0.25 and 0.75 percentiles.



(a) (b)
 Figure 4. Comparison of the vertical distribution of particulate organic carbon concentrations (μmol/l) from UVP5
 measurements (green), PlankTOM12 daily model (blue) and extrapolated UVP5 measurements (red): (a) - small particulate
 organic carbon concentrations; (b) - large particulate organic carbon concentrations. The median is shown in dark and the
 shading corresponds to values between the 0.25 and 0.75 percentiles. The size of the particles does not correspond
 completely between the observations and the model, for POC_L the UVP particle range is chosen as 0.256-5.16 mm that
 corresponds approximately to the POC_L in the model.

272 PlankTOM12 overestimates POC_S up to 3 µmol/L in the first 200m (Fig.4a, green and blue curves). UVP5 does not

273 capture all small particles that is why we extrapolated the size range of UVP measurements (red curve, see details in 274 2.1.1). The extrapolated measurements show an increase in POCs in the first 100m, however this increase still results 275 in the lower concentration compared with PlankTOM12. These results indicate that PlankTOM12 overestimates the 276 concentration of small particulate organic carbon. PlankkTOM12 also overestimates POC_L by up to 0.08 µmol/L in 277 the first 200m and does not catch the increase in POCL between 300 and 500m. Observations show an increase in 278 POC_L concentration in the first 50m while PlankTOM12 reproduces it lower, at 100m. The RMSE between modelled 279 and observed POCs is 0.33 µmol/L, with correlation coefficient equals 0.083. RMSE equals 0.23 µmol/L with 280 correlation coefficient 0.061 for POCL. The exclusion of isolated large values of POCL (>2 µmol/L) from the 281 observation data set reduces the RMSE of POC_L to 0.062 μ mol/L with correlation 0.18. We believe that these 282 differences result from differences in space and time resolution of observations and ocean model outputs. In-situ 283 measurements are obtained at a particular time of the day and a particular latitude-longitude position while the model

provides estimations over the day (or month) and on the model grid (2° longitude and mean 1.1° latitude resolution).

285 We concluded that observed and modelled POCs and POCL have a common tendency in their vertical distributions.

However, among other things, differences in amplitudes may affect our findings in this work when we develop a ML

287 method based on observations only.

- 288 Due to the constraint in data availability further we use monthly PlankTOM12.
- 289 Before developing a ML method, we investigate the interactions between targets and drivers in the model. Table 2
- shows the correlation coefficients between the POCs and POCL and corresponding drivers that can influence POCs
- and POC_L variability. Correlation between drivers could also provide valuable information to minimise the number of

- driver but they are not shown here where the focus is on discovering the effect of a large set of drivers on POC
- distribution, and because driver correlations could also result from the physics as well as from the model construction. POC_s correlates with gelatinous zooplankton (GEL, r=0.66), microzooplankton (PRO, r=0.63), coccolithophore
- (COC, r=0.56), as well as with their values from previous time step (GEL, r=0.67; PRO, r=0.51; COC, r=0.59).
- 296 Coccolithophore is one of the most abundant phytoplankton types in this version of the PlankTOM model (similar to
- 297 Wright et al., 2021). The growth of phytoplankton transfers dissolved inorganic carbon into dissolved organic carbon
- 298 which further aggregates into POCs and POCL. Also, POCs is generated from microzooplankton egestion and
- excretion (Fig. 2). In addition to the mentioned above PFTs, POCs shows a correlation 0.44 with temperature vertical
- 300 profile at both the considered time step and at the previous time step. POCs has a negative correlation with NO₃ (r=-
- **301** 0.46) and PO₄ (r=-0.41).

302 POC_L does not show a high correlation with any of the proposed drivers individually and is therefore most likely the 303 result of multiple processes and/or multiple drivers, including for its production and destruction. The ML approach 304 should be able to identify combinations of drivers beyond straight correlations that are investigated directly here. 305 POC_L has the highest correlation with chlorophyll a (r=0.42), gelatinous zooplankton at the considered time step 306 (r=0.37), and at previous time step (r=0.36). Gelatinous zooplankton contribute to POC_L formation through egestion 307 and excretion mainly from mucus (Fig. 2). As explained in Wright et al. (2021), mucus forms a large low-density mass 308 through aggregation with other particles. It can explain a correlation of gelatinous zooplankton with POCL in PlankTOM12. 309

Table 2. Correlation coefficient between small (POCs) and large (POCL) particulate organic carbon concentration and
 possible drivers. Estimation is based on monthly PlankTOM12 output at the position of real-world observations from Fig.
 1. 'vp' - vertical profile, 'mean' - average over MLD, 'back' - values from previous month.

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Driver	POCs	POCL	Driver	POCs	POCL	Driver	POCs	POCL
POC	1.00	0.33	BAC vp	-0.14	0.15	BAC back vp	-0.10	0.09
GOC	0.33	1.00	MES vp	-0.09	0.07	MES back vp	-0.09	-0.07
Depth	-0.32	-0.24	PTE vp	-0.07	0.17	PTE back vp	-0.08	0.08
Temperature vp	0.44	0.17	DIA vp	-0.04	0.15	DIA back vp	-0.03	0.09
Temp back vp	0.44	0.17	COC vp	0.56	0.31	COC back vp	0.60	0.31
MLD	-0.01	-0.07	PIC vp	0.00	0.07	PIC back vp	0.06	0.06
NO3 vp	-0.46	0.01	PHA vp	0.27	0.15	PHA back vp	0.30	0.17
PO ₄ vp	-0.41	0.04	GEL vp	0.66	0.37	GEL back vp	0.68	0.36
NO3 back vp	-0.46	0.03	PRO vp	0.63	0.16	PRO back vp	0.51	0.14
PO ₄ back vp	-0.41	0.05	MAC vp	0.07	0.14	MAC back vp	0.08	0.13
CHL vp	0.18	0.42	MIX vp	0.07	0.17	MIX back vp	0.03	0.05
CHL back vp	0.11	0.22	FIX vp	-0.00	0.23	FIX back vp	-0.00	0.23

^{3.2.} Development of the Machine Learning method.



Figure 5. Comparison of the performance of the Random Forest (RF) and XGBoost methods and their fit to data for small
 (POC_s) particulate organic carbon concentration; (a, b, c) - RMSE in µmol/l, (d, e, f) - absolute bias in µmol/, (g, h, i) correlation coefficient; (a, d, g) - training data set, (b, e, h) - the validation data set, (e, f, i) - the test data set. Results
 compare data from the original (sampled) PlankTOM12 model output and POC_s reconstructed using RF (blue) and XGB
 (orange). The low RMSE and absolute biases indicate better performance of the ML method.

We tested 27 sets of drivers (Table 1) and two ML methods, Random Forest (RF) and XGBoost regression (XGB).

323 Figure 5 shows the statistics of POCs reconstruction using RF and XGB. XGB (orange) generally overperforms RF

324 (blue). The statistics are slightly worse for the validation and test data sets, as expected. For reconstructions using

325 XGB, the RMSE and absolute bias are about 0.05 µmol/L and 0.03 µmol/L on the training data set and vary around

326 0.1 µmol/L and 0.05 µmol/L, on the validation and test data, respectively. Correlation coefficients (Fig. 5g, h, i) have

- 327 high values on all datasets showing that the vertical profiles of POCs have a correct shape. These results show that the
- available spatial and temporal coverage of *in situ* observations can be sufficient to reconstruct POCs with an
- appropriate accuracy over the global ocean. The analysis of global maps (shown below) will help to identify areas
 - 330 with low accuracy and their differences with training regions.

The worse results (highest RMSE, highest absolute bias, lowest correlation) are produced when there are no PFTs in the driver set (DS1 and DS2; Figure 5): for XBoost, RMSEs are $0.24 \mu mol/L$, absolute biases equal to $0.12 \mu mol/L$ with correlation coefficient 0.67 on the test data sets. Poor results are also obtained for DS9, 11, 12, 13 and 14: these

with correlation coefficient 0.67 on the test data sets. Poor results are also obtained for DS9, 11, 12, 13 and 14: these 5 driver sets do not have any information on microgrouplankton (PRO) and show high PMSEs and absolute biases

5 driver sets do not have any information on microzooplankton (PRO) and show high RMSEs and absolute biases,

335 around 0.16 µmol/L and 0.074 µmol/L, with low correlation, 0.83, compared with other driver sets which include PRO. These results indicate that microzooplankton plays an important role in POCs variability in the PlankTOM12

- 336
- 337 model.



338 339 Figure 6. Comparison of the performance of the Random Forest (RF) and XGBoost methods and their fit to data for large 340 (POC_L) particulate organic carbon concentration; (a, b, c) - RMSE in µmol/l, (d, e, f) - absolute bias in µmol/l, (g, h, i) -341 correlation coefficient; (a, d, g) - training data set, (b, e, h) - the validation data set, (e, f, i) - the test data set. Results 342 compare data from the original (sampled) PlankTOM12 model output and POC_L reconstructed using RF (blue) and XGB 343 (orange). The low RMSE and absolute biases indicate better performance of the ML method.

344 Figure 6 shows the statistics of POCL reconstruction using RF and XGB. XGBoost again slightly overperforms RF on

- 345 most driver sets. Results for driver sets with PFTs show lower RMSEs and absolute biases, and higher correlation 346 coefficients. Except for the effect of PFTs on the POC_L reconstruction, we did not observe a clear influence of one
- 347 driver or group of drivers. Using XGBoost the reconstruction of POCL shows the RMSE in DS1 is high at 0.03 µmol/L,
- 348 while it is in the range of 0.021-0.026 µmol/L in DS3-DS27, with absolute bias in DS1 of 0.02 µmol/L and 0.015-
- 349 0.018 µmol/L for DS3-DS27 based on test data (Fig. 6c, f). Likewise, a correlation coefficient of 0.56 for DS1, and
- 350 between 0.7 and 0.77 for DS3-DS27 based on the training data set (Fig. 6g).

351 We estimated the ranking of importance for each driver averaged over 27 driver sets (Table 1) for RF and XGB (Fig.

352 7). Both, RF and XGB, show that microzooplankton (PRO), depth level, temperature, NO3 and PO4 play a dominant

353 role in reconstruction of POCs. The absence of gelatinous zooplankton (GEL) can slightly improve the reconstruction.

- Also, latitude and longitude do not affect POCs reconstruction. The depth level, temperature, MLD, microzooplankton
- 355 (PRO) and phaeocystis (PHA), PO₄, and chlorophyll a averaged over MLD play a dominant role in POC_L 356 reconstruction.
- 357 The sinus of latitude is in the top ten drivers that most affect POC_L using XGBoost method: POC_L distribution has a
- 358 lot of meridional variability that results in the sinus of latitude being in the top 10 drivers. As for POCs, gelatinous
- 359 zooplankton (GEL) shows a negative rank of driver importance and its removal from the list of drivers can improve
- 360 the statistics of reconstruction. Also, chlorophyll a concentration from the previous month shows a similar effect on
- **361** POC_L (Fig. 7c, d).
- 362 It is worth noting that any driver that shows negative importance in the reconstruction has only a small influence on 363 the accuracy (Fig. 5 and 6). Thus, its removal does not improve the reconstruction significantly.
- Based on Figures 5, 6 and 7 we have chosen 10 driver sets with low RMSEs and absolute biases, and high correlation
- 365 coefficients (based on test data set) for POC_s and POC_L to provide global maps of these statistics and to see their
- regional distributions. DS 5, 15, 16, 21, 22, 23, 24, 25, 26, 27 were chosen for further investigation of POCs reconstruction; DS 5, 8, 15, 16, 17, 21, 23, 25, 26, 27 for POC_L reconstruction. Common for POCs and POC_L driver
- **368** sets 5, 15, 16, 21, 23, 25, 26, 27 include all PFTs and their average over MLD, geographical positions and incident
- 369 light as well as chlorophyll a, PO₄, and gelatinous zooplankton and microzooplankton from the previous time step
- 370 (Table 1). Also, we found that POCs reconstructions rest on biochemical conditions (DSs 21 and 24), while POCL
- 371 reconstruction mostly depends on the composition of the PFTs in the driver set (DSs 8 and 17). Additionally, we keep
- 372 DS1 to demonstrate a global effect of PFTs on reconstruction.



(C)
 Figure 7. Ranking of importance for each driver averaged over 27 driver sets: (a) - Random Forest (RF) for reconstruction of small (POC_s) particulate organic carbon concentration; (b) - XGBoost (XGB) for small (POC_s) particulate organic carbon concentration; (d) - XGB for POC_L concentration. 'vp' – vertical profile, 'mean' – average over MLD, 'back' – values from previous month.

3.3. POC_S and POC_L vertical profile reconstruction over the global ocean

378

379 In the previous section we showed that XGBoost provides the best results for the reconstructions of POCs and POCL.

Further we use this ML method. Here we will discuss the regional results of DS1 without PFTs and 10 best driver sets chosen for each target separately.



80°I

60°

XGB, DS1

382

Large particulate organic carbon







(d)





387 reconstruction of POC_L based on DS25 using XGBoost.

388 Figure 8 shows POCs and POCL concentration averaged over the depth and period 2009-2013 for PlankTOM12 (Fig. 389 8a, b), XGboost reconstruction based on DS1 (Fig. 8c, d) and XGBoost reconstruction based on DS25 (Fig.8 e, f). 390 XGBoost captures well the spatial patterns: the high concentration of POCs in the Equatorial Eastern Pacific and its 391 low concentration at high latitudes, as well as the high concentration of POCL in the Equatorial Eastern Pacific and in 392 the North of the Indian Ocean and its low concentration in the Subtropical North and South Atlantic and in the 393 Subtropical North Pacific. The presence of PFTs in driver sets (Fig. 8e, f) improves the reconstruction: the spatial 394 patterns and its amplitude are visually close to ones from PlankTOM12 (Fig. 8a, b). The high concentration of POCs 395 in the Equatorial Eastern Pacific is represented better using DS25 compared with DS1 where the concentration in the 396 latitude band 0°S-20°S along the Peru is overestimated. Also, small decreases of POCs in the Subtropical North and South Atlantic are captured better when we use DS25. Similar for POCs, the high concentration in the Equatorial 397 398 Eastern Pacific is represented better using DS25 compared with DS1 where the concentration misses the small

 $\begin{array}{ll} \textbf{399} \\ \textbf{400} \\ \textbf{as in the Subtropical North Pacific are pronounced better with DS25.} \end{array}$

401 Figure 9 shows regional correlation coefficients and RMSEs between PlankTOM12 and XGBoost reconstruction over 402 the global ocean for 2009-2013. We averaged correlation coefficient and RMSEs over 7 latitude zones: 90°N-60°N, 403 60°N-40°N, 40°N-20°N, 20°N-20°S, 20°S-40°S, 40°S-60°S, 60°S-90°S. In POCs reconstruction, the DS1 shows the 404 lowest correlation across latitude bands (between 0.22 and 0.9), and highest RMSEs (0.05-0.34 µmol/L; Fig.9a, b). 405 DSs 25 and 26 show the highest correlations in the range of 0.68 (in region 60°S-90°S) and 0.97 (in region 20°N-20°S) 406 and the lowest RMSEs in the range of 0.021 (in region 60°S-90°S) and 0.14 µmol/L (in region 90°N-60°N). DS25 407 contains information on the previous-month distribution for micro-, macrozooplankton and gelatinous zooplankton 408 vertical profiles as well as coccolithophores and chlorophyll a averaged over the MLD. DS26 is like DS25 but the 409 drivers which bring information from the previous month are microzooplankton and gelatinous zooplankton vertical 410 profiles.

411 10 driver sets (excluding DS1) show their highest RMSEs in POCs reconstruction in the region 90°N-60°N, with 412 values up to 0.14 µmol/L in DS27 (Fig. 9b). Figure 10 shows maps of RMSEs (a, b) and correlation coefficients (c, 413 d) between PlankTOM12 and reconstructed small particulate organic carbon (POCs) by XGBoost using driver sets 1 414 (a, c) and 25 (b, d). The region 90°N-60°N shows improvement in RMSEs and absolute biases in DS25 compared with 415 DS1, with RMSEs decreasing from 0.2 µmol/L to 0.03 µmol/L in Norwegian Sea, Baffin Bay, and the Arctic Ocean. 416 However, errors stay high in the coastal regions, Northwestern passage and Hudson Bay that contribute to the high 417 total RMSEs in this region. Results are similar for the region 60°N-40°N, where correlation coefficients increased 418 from 0.3 to 0.87 on average over these zones (Fig. 10c, d). The tropical region 20°N-20°S shows correlation coefficient 419 up to 0.97 for all driver sets except DS1. However, RMSEs are high in the tropical region, about 0.11µmol/L on 420 average (Fig. 9b), with RMSEs values of 0.2 µmol/L in the Tropical Eastern Pacific and Bay of Bengal in DS25 (Fig. 421 10b). The high RMSEs in the Tropical Eastern Pacific can indicate insufficient data in a region of high interannual 422 variability to correctly reconstruct POCs distribution. The region of the Southern Ocean ($>60^{\circ}S$) shows the lowest 423 correlation coefficients (in the range of 0.64-0.69) and RMSEs (in the range 0.023-0.044 µmol/L) for POCs (Fig. 9a, 424 b). The inclusion of PFTs in the driver set significantly improves the RMSE in the region around 40°S for small 425 (POCs) particulate organic carbon. The statistics are improved by about 75% in the region 40°S-60°S with RMSE 426 decreasing from 0.18 (DS1) to 0.03 (DS25) and the correlation coefficient increasing from 0.22 (DS1) to 0.84 (DS25), 427 on average (Fig. 9a, b; Fig. 10). The improvements in the Southern region are related to the role of zooplankton in the 428 carbon flux in this area (Le Quéré et al., 2016; Wright et al., 2021).





Figure 9. Correlations and RMSE averaged over latitude zones between PlankTOM12 and XGBoost reconstruction over 431 the global ocean for 2009-2013: (a, c) - correlation coefficient, (b, d) - RMSE in µmol/l (b, d);, (a, b) - small particulate 432 organic carbon (POC_s), (c, d) - large particulate organic carbon (POC_L).

433 In POCL reconstruction, DS1 also shows the lowest correlation coefficients (0.35-0.75) and the highest RMSEs (0.027-434 0.47 µmol/L) (Fig. 9c, d). DS25 shows the best results on average, with the correlation coefficient varying between 435 0.43 (in the region 60°S-90°S) and 0.84 (in the region 20°N-20°S), and RMSE varying between 0.021 (in the region 436 20°S-40°S) and 0.046 (in the region 90°N-60°N) µmol/L. POC_L are reconstructed better in subtropical and tropical 437 regions compared to high latitude zones (Fig. 9c, d).

438 As for POCs, 10 driver sets (excluding DS1) show their highest RMSEs in POCL reconstruction in the region 90°N-439 60°N, with values up to 0.05 µmol/L in DS27 (Fig. 9d). Figure 11 shows maps of RMSEs (a, b) and correlation 440 coefficients (c, d) between PlankTOM12 and reconstructed large particulate organic carbon (POC_L) by XGBoost using driver sets 1 (a, c) and 25 (b, d). Contrast to POC_s reconstruction, the region 90°N-60°N does not show improvement in RMSEs for POC_L reconstruction (Fig. 11b) in DS25 compared with DS1, with still high RMSEs in Norwegian Sea, Baffin Bay, and the Arctic Ocean, and additionally for POC_L in Greenland Sea, where the algorithm did not have data for training. Similar to POC_s , errors stay high in the coastal regions, Northwestern passage and Hudson Bay that contribute to the high total RMSEs in this region.

446 Global maps of statistics suggest that the most sensible region to driver set's composition for POC_L is the Southern 447 Ocean, as for POCs (Fig. 11). In the 40°S-60°S region, RMSE is reduced from 0.037 µmol/L in DS1 to 0.024 µmol/L 448 in DS25 (Fig. 9d), and the correlation coefficient is increased from 0.42 to 0.66 (Fig. 9c) on average, respectively. In 449 the Southern region 60°S-90°S, RMSE is reduced from 0.047 µmol/L in DS1 to 0.033 µmol/L in DS25, and the 450 correlation coefficient is increased from 0.33 to 0.42 (Fig. 9c) on average, respectively. The average correlation 451 coefficients in this zone were found to be less than 0.5 in all tests with the highest value 0.5 in DS21. DS21 contains 452 all PFTs and chlorophyll a vertical profile as drivers. The RMSE for DS21 in this region is close to the one of DS25, 453 $0.34 \mu mol/L$ and $0.33 \mu mol/L$, respectively. It identifies the importance of chlorophyll a in the Southern Ocean as 454 driver of POC_L variability.



455

Figure 10. RMSE and correlation between monthly PlankTOM12 and results of POC_s reconstruction using XGBoost over
 the period 2009-2013 for POC_s. (a, b) – RMSEs, (c, d) – correlation coefficients; (a, c) – reconstruction based on DS1
 (NoPFT); (b, d) – reconstruction based on DS25 (vertical profiles of zooplanktons, and zooplankton and phytoplankton

459 averaged over MLD).



Figure 11. RMSE and correlation between monthly PlankTOM12 and results of POC_L reconstruction using XGBoost
 over the period 2009-2013 for POC_L. (a, b) – RMSEs, (c, d) – correlation coefficients; (a, c) – reconstruction based on
 DS1 (NoPFT); (b, d) – reconstruction based on DS25 (vertical profiles of zooplanktons, and zooplankton and

464 phytoplankton averaged over MLD).

465 The statistics of POC_s and POC_L reconstruction do not vary significantly between driver sets in all regions except in 466 the Southern Ocean. This region is most sensitive to the composition of driver sets for both POC_s and POC_L.

467 4. Conclusion.

468

The aim of this work was to test the potential of using Machine Learning to reproduce modelled concentrations of
 particulate organic carbon within the ocean using the distribution of available observations. We co-localised outputs
 of the PlankTOM12 global biogeochemical ocean model with the positions of observations of small (POCs) and large
 (POCL) particulate organic carbon concentrations. Using PlankTOM outputs as references we could identify the best

473 ML method for POC reconstruction and estimate method's accuracy in regions with poor observational cover.

We tested two ML methods to reconstruct POCs and POCL: the XGBoost regressor and Random Forest. Both methods are algorithms based on decision trees. XGBoost overperformed Random Forest by about 9% on average for POCs reconstruction and by about 3% on average for POCL reconstruction. XGBoost regressor builds the model sequentially improving it at each iterative step. At each iteration, XGBoost regressor analyses the prediction and gives more weight to the data where the fit is still wrong. It is a good tool for an unbalanced data set, like in our case where the data of particulate organic carbon concentration are sparse in time and space.

We tested the influence of a wide range of environmental and ecosystem drivers on POC_s and POC_L reconstruction.
 The introduction of Plankton Functional Types (PFTs) in the driver set greatly improves the fit and shows a linkage
 between surface ecosystem structure and particulate organic carbon distribution within the ocean interior. We

improved the accuracy of POCs reconstruction by 59% on RMSE, 63% on absolute bias and by 52% on correlation
by introducing Plankton Functional Types (PFTs) in the driver sets (from the comparison of DS1 and DS25). The
presence of PFTs in the driver sets also improved the accuracy of POC_L reconstruction by 22% on RMSE, absolute
bias and correlation (from the comparison of DS1 and DS25). POCs variability mostly depends on the depth level,
vertical profiles of microzooplankton, temperature and PO4. POC_L variability depends on the depth level, MLD,
chlorophyll *a* averaged over MLD, vertical profiles of temperature, microzooplankton, phaeocystis and PO4.
Additionally, we identified that chlorophyll *a* in driver sets improves the POC_L reconstruction in the Southern Ocean.

+0.5 Automation Automation and the souther sets improves the POUL reconstruction in the Souther

490 Despite the good accuracy over the global ocean on average, the statistics are worse in the coastal regions and in the 491 Tropical Eastern Pacific. The coastal regions suffer from the lack of data to represent the coastal dynamics. Therefore 492 the ML reconstructions assign open-ocean processes to coastal regions, leading to significant biases. The Tropical 493 Eastern Pacific is a region of strong interannual variability and the sparse measurements in time make it harder to 494 capture this variability correctly. Other regions with poor coverage by observations - the Eastern Indian Ocean, the 495 Western Pacific Ocean and the Southern Ocean - show the statistics of reconstruction comparable to one from regions 496 with a good cover - regions in the Atlantic Ocean. However, we found that the Southern Ocean is a more sensible region to the driver set's composition. The observational data is particularly sparse in this region and our analysis 497 498 suggests that identifying the drivers of importance based on real dataset will be difficult.

499 Here we showed that the XGBoost regressor and Random Forest are suitable for this problem and can reconstruct 500 modelled POCs and POC₁ with appropriate accuracy. This is evidenced from the globally averaged correlation 501 coefficient up to 0.88 for POCs and 0.68 for POCL, and the globally averaged RMSE up to 20 % (0.08 µmol/L) of 502 standard deviation of PlankTOM12 POCs, and 65% (0.028 µmol/L) of standard deviation of PlankTOM12 POCL. ML 503 outputs represent well the spatial patterns of POCs and POCL distribution. However, the validity of the approach on 504 observations is dependent on the availability of co-located information on the drivers of importance. For some drivers 505 this should be possible (e.g. environmental conditions and chlorophyll a), while for other drivers information is more 506 sparse (e.g. the PFTs). Our analysis suggests that additional PFT observations would help provide broader insights 507 into the distribution of POC in the ocean. The next step of this work is to apply ML to real data using methods from 508 the present study. Testing the present ML approach on observations will also help provide suggestions for an optimal 509 set of drivers that can be measured specifically for POC reconstruction. For example, based on model results only, 510 our results suggest that microzooplankton concentration is particularly important and should be measured more 511 systematically, especially in the regions of high interannual variability. Likewise, this work provides information on 512 the variables that are less important in POC variability, like vertical profiles of gelatinous zooplankton, or mixed 513 phytoplankton for POC_s and coccolithophore for POC_L, and, thus, less important to be measured in this context. These 514 results will need to be tested with observations before firmly confirming the validity of the drivers. The validated 515 driver sets can help guide observational programs. In addition, recent advances in plankton imaging (Irisson et al., 516 2022; Lombard et al., 2019; Orenstein et al., 2022) and omics (Faure et al., 2021) will soon provide a new global set 517 of data to estimate PFT concentrations across ocean basins allowing to better identify potential biological drivers of 518 POC variability. The new available data of PFTs will significantly facilitate the application of ML methods, such as 519 the one developed here, to observational data.

520 The relationships between key variables and surrounding conditions based on Machine Learning can provide a new way for establishing parameters in ocean model parameterisation. The parameters can be time and space dependent and, thus, vary from one region to another better representing the physics. Relationship between POC concentration and environmental and ecosystem conditions can help to replace parameters in parameterised sinking velocity in PlankTOM. The reconstructed POC concentration over the global ocean will contribute to the reconstruction of porosity and opacity of particles that are key variables in the sinking matter velocity.

This study provides insights on the drivers that may be responsible for POC_s and POC_L variability and regional dependencies. However, the dependencies are simply returning the outcome of complex ecosystem processes among the drivers as represented in the PlankTOM12 model. Although these processes are based on current understanding and a broad range of observations (Le Quéré et al., 2016; Wright et al., 2021; Buitenhuis et al., 2019), they remain results from a model output. Observations could reveal different drivers that are important for POC_s and POC_L. Depending on data availability and its time and space resolution, the final product based on observations should provide new insights on the drivers that govern particulate organic carbon concentration in the real ocean.

534 Data and code availability. PlankTOM12 data used within this study are available at 535 https://doi.org/10.5281/zenodo.7324781. UVP5 data can be found at https://doi.org/10.1594/PANGAEA.924375 (R. 536 Kiko et al., 2021). Codes for data preparation, development of machine learning methods and tests of different driver 537 sets as well as codes that provide figures shown in the article can be found at https://doi.org/10.5281/zenodo.7326992.

538 Author contribution. All authors contributed to the development of the methodology. ADS, CLQ, ETB designed the 539 experiments, and ADS carried them out. ADS developed codes and performed the simulations. ADS prepared the 540 paper with contributions from all coauthors.

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