

# Ensemble simulation of the Last Glacial Maximum marine biogeochemistry and atmospheric pCO<sub>2</sub> drawdown due to the soft-tissue biological carbon pump

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## Abstract.

During the Last Glacial Maximum (LGM), atmospheric pCO<sub>2</sub> was approximately 90 ppm lower than in the pre-industrial era. Several hypotheses have been proposed to explain this difference, including changes in nutrient supply, increased iron input to the ocean, reduced air-sea gas exchange due to changes in sea-ice coverage, and variations in overturning circulation strength driven by differences in wind stress and atmospheric moisture diffusivity. Current modeling approaches that simulate LGM marine biogeochemistry typically use parameter sets calibrated under pre-industrial conditions, assuming that these parameter values are generic and independent of environmental conditions. This could introduce uncertainty due to the imperfect knowledge of the values that should be assigned to the parameters for the LGM environment. The extent to which this uncertainty affects the simulated LGM marine biogeochemistry remains unclear. In this study, we employ an optimality-based variable stoichiometry plankton ecosystem model coupled with a 3D Earth system model to simulate LGM conditions. We conduct sensitivity analyses with 24 combinations of marine biogeochemical (reduced benthic denitrification rate, decreased sedimentary iron input, a higher PO<sub>4</sub><sup>3-</sup> inventory, and increased atmospheric iron deposition) and physical boundary conditions (changes in wind stress pattern and reduced meridional moisture diffusivity over the Southern Ocean). For each combination, we perform 20 simulations using 20 biogeochemical parameter sets selected out of 600—each calibrated against present-day observations and representing pre-industrial biogeochemistry about equally well—resulting in a total of 480 simulations. We aim to quantify the uncertainty in simulated LGM marine biogeochemistry and atmospheric pCO<sub>2</sub> arising from uncertainties in parameter settings and boundary conditions. Our results show that changes in iron input, including increased aeolian dust deposition and decreased sedimentary input, exert the most profound influence on marine biogeochemistry and pCO<sub>2</sub> drawdown. Changes in macro-nutrients alone have limited effects, owing to co-limitation effects and the variable stoichiometry in our model. The impact of physical conditions on biogeochemical tracers varies, depending on the specific biogeochemical settings. We found that the changes in carbon to nutrient ratios in particulate organic matter are positively correlated with the changes in Fe supply, and could amplify the effect of Fe availability on changes in the atmospheric pCO<sub>2</sub>. Compared to pre-industrial reference conditions, atmospheric pCO<sub>2</sub> under full LGM conditions decreases by 36 to 58 ppm across the 20 simulations. The spread between the maximum and minimum decrease in simulated glacial pCO<sub>2</sub> is approximately 50% of the mean decrease (43 ppm). These findings highlight that although the 20 parameter sets similarly reproduce pre-industrial

marine biogeochemistry, significant variance remains in the marine biogeochemical and atmospheric pCO<sub>2</sub> responses to LGM forcings.

## 1 Introduction

The Last Glacial Maximum (LGM) was characterised by substantially lower atmospheric pCO<sub>2</sub>, around 90 ppm lower than the pre-industrial (Monnin et al., 2001), making it an important case study for understanding the mechanisms influencing changes in pCO<sub>2</sub>, with implications for future climate projections. A robust quantitative understanding of the reduced atmospheric CO<sub>2</sub> is still lacking, and several hypotheses have been proposed to explain the lower atmospheric pCO<sub>2</sub> during the LGM. Those include lower sea surface temperatures and higher solubility of CO<sub>2</sub> (Jaccard and Galbraith, 2012), partly counteracted by reduced ocean volume and increased salinity, changes in orbital parameters (Weaver et al., 1998), a higher global nitrate inventory due to reduced benthic denitrification (McElroy, 1983), and a higher global phosphate inventory driven by enhanced terrestrial erosion (Broecker, 1982; Wallmann, 2010; Wallmann et al., 2016), both associated with sea-level retreat. Other factors include changes in the AMOC (Atlantic Meridional Overturning Circulation) (Muglia and Schmittner, 2015), atmospheric moisture diffusivity (Sigman et al., 2007), reduced air-sea gas exchange due to changes in Antarctic sea-ice cover (Stephens and Keeling, 2000), changes in air-sea disequilibrium of carbon (Khatiwala et al., 2019), and enhanced iron (Fe) input to the ocean (Martin, 1990; Martínez-García et al., 2014). Iron is a critical micro-nutrient for marine phytoplankton, particularly in high-nutrient, low-chlorophyll (HNLC) regions, where its scarcity limits productivity (Martin et al., 1987). Two major sources of Fe to the ocean are atmospheric dust deposition and sedimentary inputs. While atmospheric deposition has long been recognised as a major source of Fe, recent studies have shown that sedimentary inputs could have a much larger impact on the Fe cycle in the global ocean (Tagliabue et al., 2016; Somes et al., 2021).

Model simulations are suitable tools for studying the effects of these factors on marine biogeochemistry and their potential contribution to reduced pCO<sub>2</sub> during the LGM. Models that have been applied to study the LGM range from simple box model approaches (Broecker, 1982; Sarmiento and Toggweiler, 1984) to 3D Earth system models that consider ocean circulation and marine biogeochemical cycles (Bopp et al., 2003; Brovkin et al., 2007; Tagliabue et al., 2009; Somes et al., 2017; Kempainen et al., 2019; Ödalen et al., 2020; Matsumoto et al., 2020a; Kageyama et al., 2021). The biogeochemical components of these Earth system models contain various representations of marine ecosystems, transport and sinking of particles, and cycling of macro- and micro-nutrients.

The calibration of poorly known parameters is an important part of setting up reliable biogeochemical models. In principle, calibration efforts look for a set of optimal parameter estimates that minimises a metric that quantifies the data-model misfit, such as the root mean square error. Some challenges exist in the optimisation process, particularly for Earth system models that often have to cope with sparse data availability with uneven spatial and temporal distribution (Schartau et al., 2017). Also, the computational cost for spinning up Earth system models is often high, particularly for those with higher functional complexity and higher spatial resolution. In the case of high computational costs, the number of parameter sets for which model solutions can be evaluated is limited and for this reason 'optimal' parameters are often chosen pragmatically (Séférian et al.,

2016). Apart from the number of possible model runs, the choice of an appropriate metric is also crucial. This choice depends  
60 objectively on the available data considered, but it is also subjective with regard to the error model employed (Jolliff et al.,  
2009; Schartau et al., 2017; Kriest et al., 2020). Calibration becomes particularly challenging when it comes to simulating the  
LGM: Data-model misfits are usually evaluated for pre-industrial or present-day conditions. To what extent a parameter set  
calibrated against pre-industrial or present-day observations could similarly well represent the LGM is as yet unknown.

Here we employ an optimality-based variable stoichiometry plankton ecosystem model (OPEM) coupled to the University of  
65 Victoria (UVic) Earth system model of intermediate complexity (UVic-OPEM) to simulate LGM marine biogeochemistry and  
atmospheric  $p\text{CO}_2$ . The model has been devised employing with variable stoichiometry of particulate organic matter (POM),  
and it provides a relatively reliable representation of the most important biogeochemical processes in today's ocean (Pahlow  
et al., 2020; Chien et al., 2020; Li et al., 2024). In addition, by resolving elemental stoichiometry in UVic-OPEM, it is possible  
to investigate how elemental ratios of particulate organic matter may have varied during the LGM, and how those differences  
70 affect the changes in the atmospheric  $p\text{CO}_2$ . We conduct sensitivity analyses with 24 combinations of LGM boundary condi-  
tions (Table 2), namely biogeochemical (reduced benthic denitrification rate, decreased sedimentary iron input, a higher  $\text{PO}_4^{3-}$   
level, and increased atmospheric iron deposition) and physical (changes in wind stress pattern and reduced meridional moisture  
diffusivity over the Southern Ocean, Muglia and Schmittner, 2015; Sigman et al., 2007). For each combination, the subset of  
20 of the 600 parameter sets was considered that exhibit the best model agreement with the observations, resulting in a total  
75 of 480 simulations. The objective of this study is to examine the potential uncertainty in LGM simulations due to the typically  
employed selection of one specific parameter set calibrated for pre-industrial-climate boundary conditions. We also investigate  
the effects of variable stoichiometry of particulate organic matter (POM) along with different biogeochemical and physical  
boundary conditions, on atmospheric  $p\text{CO}_2$  and marine biogeochemistry during the LGM. We are particularly interested in  
identifying uncertainties that might play a role in estimating  $p\text{CO}_2$  changes in the LGM. The biogeochemical mechanisms  
80 driving glacial carbon drawdown serve as critical natural analogs for evaluating ocean-based carbon dioxide removal (CDR)  
strategies. For instance, uncertainties in how the glacial ocean responded to increased dust deposition mirror uncertainties in  
assessing the efficacy of ocean iron fertilization. Our results may provide insights not only regarding the interpretation of past  
 $p\text{CO}_2$  variations, but also into the effectiveness of potential CDR approaches.

## 2 Materials and Methods

### 85 2.1 The UVic-OPEM Earth system model

The Optimality-based Plankton Ecosystem Model was originally implemented in UVic2.9 (Chien et al., 2020; Pahlow et al.,  
2020), which has since been updated to UVic2.10 (Mengis et al., 2020) for this study. In the current version, we also apply the  
temperature-independent mortality that has been used for ordinary phytoplankton to diazotrophs. This modification reduces  
nitrogen fixation in the Arctic region, where it was considered too high in Chien et al. (2020), but it has only a marginal effect  
90 on the global nitrogen distribution and fluxes.

## 2.2 Model calibration and selection of the best 20 parameter sets

We constructed 600 parameter sets, each representing a unique combination of values assigned to 19 model parameters, including detritus remineralisation rate, the linear increase of sinking velocity with depth, and 17 parameters related to plankton physiology (Table 1). The leakage (implicit remineralisation of organic matter by bacteria) and mortality terms for non-N<sub>2</sub>-fixing ordinary phytoplankton and diazotrophs are assumed identical to reduce the number of possible parameter combinations. The respective 600 simulations were restarted from a previously-calibrated simulation (Chien et al., 2023) and spun up with a prescribed pre-industrial atmospheric pCO<sub>2</sub> of 284.3 ppm for over 10,000 years.

**Table 1.** Parameter names, variational ranges, the parameter set that yields the lowest cost, the lowest 20 cost parameter sets, units and descriptions. Note that ordinary phytoplankton and diazotrophs share the same temperature dependent leakage rate and temperature independent mortality rate.

| Symbol                                              | Range       | Lowest cost | Lowest 20 costs range | Units                                                                    | Definition                                    |
|-----------------------------------------------------|-------------|-------------|-----------------------|--------------------------------------------------------------------------|-----------------------------------------------|
| $A_{0, \text{phy}}$                                 | 100–400     | 254.3       | 113.6–391.5           | $\text{m}^3 (\text{mol C})^{-1} \text{d}^{-1}$                           | phytoplankton potential nutrient affinity     |
| $A_{0, \text{dia}}$                                 | 50–300      | 115.9       | 72.4–264.8            | $\text{m}^3 (\text{mol C})^{-1} \text{d}^{-1}$                           | diazotroph potential nutrient affinity        |
| $\alpha_{\text{phy}}$                               | 0.3–0.6     | 0.48        | 0.36–0.60             | $\text{m}^2 \text{W}^{-1} (\text{mol C}) \text{gChl}^{-1} \text{d}^{-1}$ | phytoplankton potential light affinity        |
| $\alpha_{\text{dia}}$                               | 0.6–1.1     | 0.73        | 0.61–1.02             | $\text{m}^2 \text{W}^{-1} (\text{mol C}) \text{gChl}^{-1} \text{d}^{-1}$ | diazotroph potential light affinity           |
| $Q_{0, \text{phy}}^{\text{N}}$                      | 0.045–0.065 | 0.061       | 0.048–0.064           | $\text{mol} (\text{mol C})^{-1}$                                         | phytoplankton subsistence N quota             |
| $Q_{0, \text{dia}}^{\text{N}}$                      | 0.11–0.14   | 0.114       | 0.112–0.138           | $\text{mol} (\text{mol C})^{-1}$                                         | diazotroph subsistence N quota                |
| $Q_{0, \text{phy}}^{\text{P}}$                      | 1.4–2.3     | 1.97        | 1.46–2.28             | $\text{mmol} (\text{mol C})^{-1}$                                        | phytoplankton subsistence P quota             |
| $Q_{0, \text{dia}}^{\text{P}}$                      | 2–3.5       | 2.54        | 2.06–3.30             | $\text{mmol} (\text{mol C})^{-1}$                                        | diazotroph subsistence P quota                |
| $k_{\text{Fe}, \text{phy}}$                         | 0.02–0.08   | 0.031       | 0.021–0.076           | $\mu\text{mol m}^{-3}$                                                   | phytoplankton half-saturation constant for Fe |
| $k_{\text{Fe}, \text{dia}}$                         | 0.10–0.16   | 0.11        | 0.10–0.15             | $\mu\text{mol m}^{-3}$                                                   | diazotroph half-saturation constant for Fe    |
| $g_{\text{max}}$                                    | 1–1.5       | 1.24        | 1.03–1.43             | $\text{d}^{-1}$                                                          | zooplankton maximum specific ingestion rate   |
| $\phi_{\text{phy}}$                                 | 90–300      | 174         | 119–226               | $\text{m}^3 (\text{mol C})^{-1}$                                         | capture coefficient of phytoplankton          |
| $\phi_{\text{dia}}$                                 | 80–300      | 155         | 117–206               | $\text{m}^3 (\text{mol C})^{-1}$                                         | capture coefficient of diazotrophs            |
| $\phi_{\text{det}}$                                 | 20–130      | 62          | 30–92                 | $\text{m}^3 (\text{mol C})^{-1}$                                         | capture coefficient of detritus               |
| $\phi_{\text{zoo}}$                                 | 45–250      | 175         | 80–175                | $\text{m}^3 (\text{mol C})^{-1}$                                         | capture coefficient of zooplankton            |
| $M_{0, \text{phy}} = M_{0, \text{dia}}$             | 0.01–0.04   | 0.013       | 0.012–0.037           | $\text{d}^{-1}$                                                          | temperature-independent mortality             |
| $\lambda_{0, \text{phy}} = \lambda_{0, \text{dia}}$ | 0.01–0.03   | 0.011       | 0.010–0.026           | $\text{d}^{-1}$                                                          | temperature-dependent leakage                 |
| $\nu_{\text{det}}$                                  | 0.05–0.08   | 0.072       | 0.052–0.078           | $\text{d}^{-1}$                                                          | remineralisation rate at 0 °C                 |
| $w_{\text{dd}}$                                     | 0.04–0.07   | 0.064       | 0.042–0.066           | $\text{d}^{-1}$                                                          | linear increase in sinking speed with depth   |

We adopt a likelihood-based cost function for evaluating the biogeochemical model performance (Equations 4 – 9 in Chien et al., 2020). Our cost function considers mismatches in means, and spatial and temporal variabilities of  $\text{PO}_4^{3-}$ , excess nitrate with respect to phosphate ( $\text{N}^* = \text{NO}_3^- - 16 \cdot \text{PO}_4^{3-} + 2.9 \text{ mmol m}^{-3}$ , Gruber and Sarmiento, 1997; Mills et al., 2015), and modified apparent oxygen utilisation ( $\text{AOU}^* = \text{AOU} - 2 \cdot \text{PO}_4^{3-}$ ), which eliminates the covariation between AOU and  $\text{PO}_4^{3-}$ ,

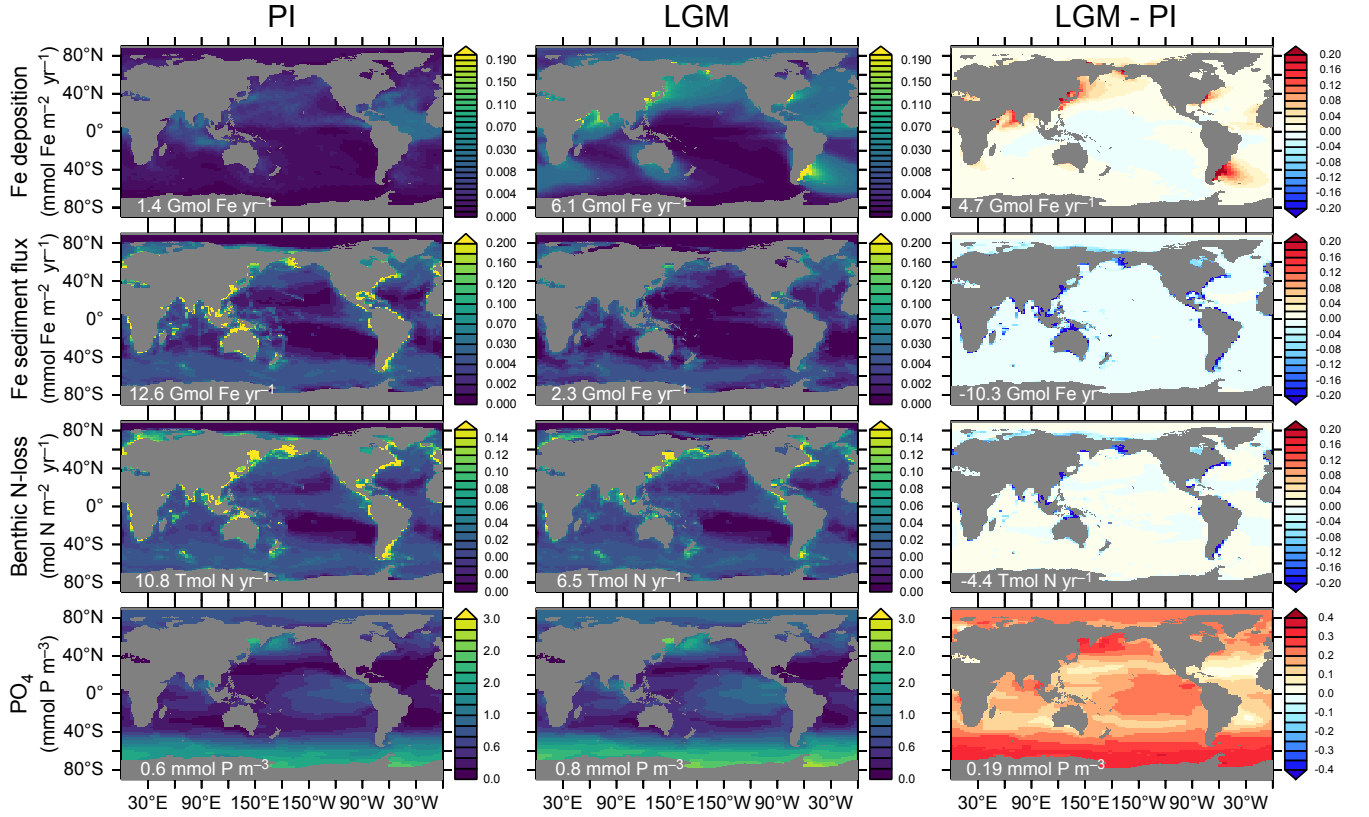
in 17 biomes (Fay and McKinley, 2014). Interestingly, most of the best-20 ranges cover about 80 % of the total ranges, except for the four grazing-related parameters  $\phi$ s, where the range is 'only' about 40–60 %. This indicates that parameter values are relatively poorly constrained by currently available observations. Modern observations may include anthropogenic signals, which could bias parameter selection. Thus, we assume here that these effects are small relative to spatial variability and note this as a source of uncertainty we cannot quantify. The 20 best parameter sets (with the lowest cost function values) are employed for analysing model behaviour under pre-industrial (PI) and last-glacial-maximum (LGM) conditions.

### 2.3 Boundary conditions

We set-up a generic LGM configuration with LGM-specific orbital parameters of the Earth. Besides the different orbital parameters, LGM conditions, including the strength of the AMOC, likely also deviated from the pre-industrial era by a lower moisture diffusivity over the Southern Ocean and a different wind pattern (Muglia and Schmittner, 2015; Muglia et al., 2018). We adopt these forcings from Somes et al. (2017) and Muglia and Schmittner (2015) to investigate their influence on the marine biogeochemistry and carbon cycle. Specifically, meridional moisture diffusivity over the Southern Ocean was reduced by a factor of 2, and the wind stress patterns are from monthly averages of models which have participated in the PMIP3 (Fig. S1). In addition, we configure a 120 m lower sea level compared to PI conditions for the changes in biogeochemical fluxes. The lower sea level was not implemented through an explicit modification of model bathymetry. Instead, it was represented diagnostically in the parameterizations of benthic processes. Specifically, we excluded benthic denitrification and sedimentary Fe input in regions shallower than 120 m relative to the pre-industrial sea level, thereby mimicking the exposure of continental shelves under LGM conditions. This results in reduced benthic denitrification (Somes et al., 2017), reduced sedimentary input of Fe (Tagliabue et al., 2010; Muglia et al., 2017), and a 15 % increase in the  $\text{PO}_4^{3-}$  inventory (Wallmann et al., 2016), which is conserved in the model. We also include a higher LGM Fe deposition, which was obtained from the LGM dust deposition estimate of Albani et al. (2014, their case C4fn-lgm). Applying a constant iron content of 3.5 % and 1 % solubility of deposited Fe (Kobayashi et al., 2021; Saini et al., 2023) yields an annual Fe deposition of  $6.1 \text{ Gmol Fe yr}^{-1}$ , which is about four times the pre-industrial flux (Fig. 1). All other physical and biogeochemical processes were computed using the same bathymetry as in the pre-industrial control simulation. This approach allows us to isolate the impact of shelf exposure on benthic fluxes without introducing additional changes to ocean circulation or ecosystem structure that would arise from a fully modified LGM bathymetry.

### 2.4 Experiment setup

We define four sets of physical and six sets of biogeochemical model configurations, two of which represent pre-industrial climate conditions, PIctl (physical) and PIallbgc (biogeochemical), and the others are different combinations of our generic LGM configuration with PI and LGM boundary conditions (Table 2): LGMatmctl and LGMallbgc refer to the full set of physical and biogeochemical LGM boundary conditions, LGMatmws has PI moisture diffusivity and LGM winds, LGMatmpi has PI moisture diffusivity and winds, and LGMFedep, LGMFesed, LGM $\text{PO}_4$  and LGMbdeni combine PI biogeochemical boundary conditions with one of LGM Fe deposition, Fe sedimentary release,  $\text{PO}_4^{3-}$  inventory, and benthic denitrification, re-



**Figure 1.** Fe deposition and sedimentary fluxes, benthic denitrification (benthic N-loss), and surface  $\text{PO}_4^{3-}$  in the PIallbgc and LGM biogeochemical configurations and their differences. PI results are the means of 20 simulations in PIctl\_PIallbgc, and LGM Fe deposition, Fe sediment flux, benthic denitrification, and surface  $\text{PO}_4^{3-}$  are from PIctl\_LGMFedep, PIctl\_LGMFesed, PIctl\_LGMBdeni, and PIctl\_LGMPO<sub>4</sub> simulation means. Values on the bottom left of each panel indicate the global annual Fe and N fluxes or  $\text{PO}_4^{3-}$  concentration.

spectively. These result in a total of 24 (4 physical x 6 biogeochemical) combinations of different physical and biogeochemical boundary conditions. For example, PIctl\_LGMallbgc stands for simulations with pre-industrial physics and full LGM biogeochemistry (Table 2). The combination of LGM moisture diffusivity and PI wind stress resulted a shut-down of the AMOC in the UVic\_ESCM and hence is not discussed here. From the calibration stage, we selected the 20 parameter sets that best reproduce present-day observations, each corresponding to a fully spun-up pre-industrial simulation. These simulations serve as initial conditions for the 24 combinations of physical and biogeochemical boundary conditions.

For each of the 24 configurations, we restarted simulations using the 20 selected parameter sets, resulting in an ensemble of 480 simulations. During the spin-up, radiative forcing was prescribed according to a fixed pCO<sub>2</sub> level (284.3 ppm for PI and 190 ppm for LGM), i.e., the feedback between atmospheric CO<sub>2</sub> and radiative forcing was switched off, while atmospheric pCO<sub>2</sub> was allowed to evolve freely in response to changes in the global carbon cycle.

This physically and biogeochemically coupled but radiatively uncoupled configuration allows us to isolate the effects of boundary conditions on pCO<sub>2</sub>, while also accounting for terrestrial carbon responses and ocean–atmosphere carbon exchange. All simulations in the ensemble were spun up for more than 10,000 years under their respective boundary conditions until the marine biogeochemistry approached a steady state. We use the results from the final year of the spin-up simulations for our analysis. The experimental setup and workflow are illustrated in Fig. 2.

To evaluate whether different physical, biogeochemical, or both conditions result in ensembles that are significantly different from the 20 simulations in PIctl, PIallbgc, or PIctl\_PIallbgc, we calculate their p-values for each tracer evaluated with Student’s t-Test in R package "Stats". To understand if changes in individual parameters significantly influence the changes in atmospheric pCO<sub>2</sub> from PIctl\_PIallbgc to LGMatmctl\_LGMallbgc, we obtain p-values using the lm (Fitting Linear Models function) in R package "Stats".

**Table 2.** Boundary conditions applied in the ensemble simulations for our (A) physical and (B) biogeochemical configurations .

| (A)                    |                       |                          | (B)                          |                |                                     |                                                      |                                      |
|------------------------|-----------------------|--------------------------|------------------------------|----------------|-------------------------------------|------------------------------------------------------|--------------------------------------|
| Physical configuration | Moisture diffusivity* | Wind stress <sup>†</sup> | Biogeochemical configuration | Fe deposition* | Fe sedimentary release <sup>†</sup> | PO <sub>4</sub> <sup>3-</sup> inventory <sup>§</sup> | Benthic denitrification <sup>‡</sup> |
| PIctl                  | PI                    | PI                       | PIallbgc                     | PI             | PI                                  | PI                                                   | PI                                   |
| LGMatmctl              | LGM                   | LGM                      | LGMallbgc                    | LGM            | LGM                                 | LGM                                                  | LGM                                  |
| LGMatmws               | PI                    | LGM                      | LGMFedep                     | LGM            | PI                                  | PI                                                   | PI                                   |
| LGMatmpi               | PI                    | PI                       | LGMFesed                     | PI             | LGM                                 | PI                                                   | PI                                   |
|                        |                       |                          | LGMPO <sub>4</sub>           | PI             | PI                                  | LGM                                                  | PI                                   |
|                        |                       |                          | LGMbdeni                     | PI             | PI                                  | PI                                                   | LGM                                  |

\*50% lower LGM moisture diffusivity over the Southern Ocean in LGM

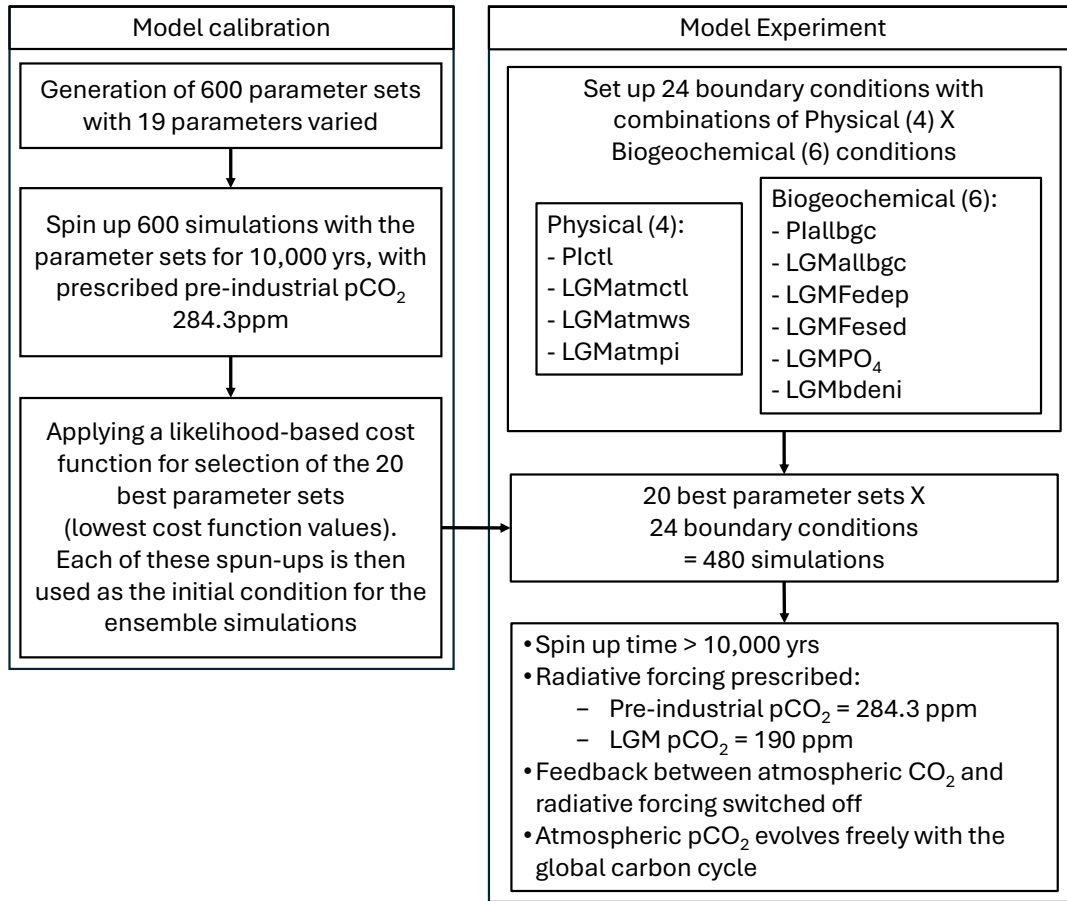
<sup>†</sup>LGM wind pattern from Somes et al. (2017); Muglia and Schmittner (2015)

\* ~4 times higher during LGM

<sup>†</sup> ~5 times lower during LGM

<sup>§</sup> ~15% higher during LGM

<sup>‡</sup> ~40% lower during LGM



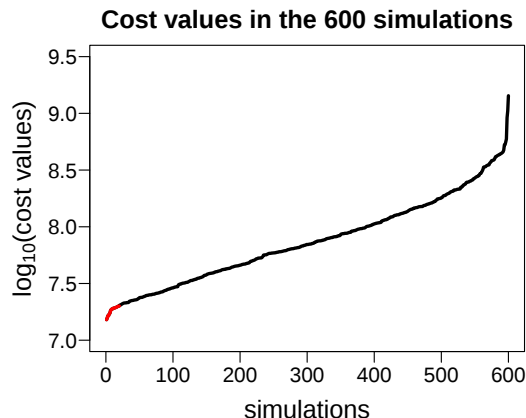
**Figure 2.** Experimental setup and workflow.

## 155 3 Results

### 3.1 Model-data misfit costs of all 600 and the selected 20 parameter sets

The 600 parameter sets yield a wide range of model-data misfit costs spanning 2 orders of magnitude (Fig. 3). When the initial 600 pre-industrial simulations are ordered according to increasing cost, the 600 cost values overall show an approximately exponential increase from the low to the high end, with around 8 simulations on both ends deviating from the trend. The 20 simulations with the lowest costs comprise 3.3 % of all simulations, and the cost function values amongst these 20 best are no more than 1.3-fold higher than the overall lowest cost value.

Globally-averaged vertical profiles of NO<sub>3</sub><sup>-</sup>, PO<sub>4</sub><sup>3-</sup>, O<sub>2</sub>, and dissolved inorganic carbon (DIC) in the 20 simulations of Plctl\_Plallbgc are compared with observational data from the World Ocean Atlas 2013 (WOA 2013, Garcia et al., 2013a, b) and GLODAPv2 (Key et al., 2015; Lauvset et al., 2016), as well as the original UVic results (Mengis et al., 2020) in Fig. 4.



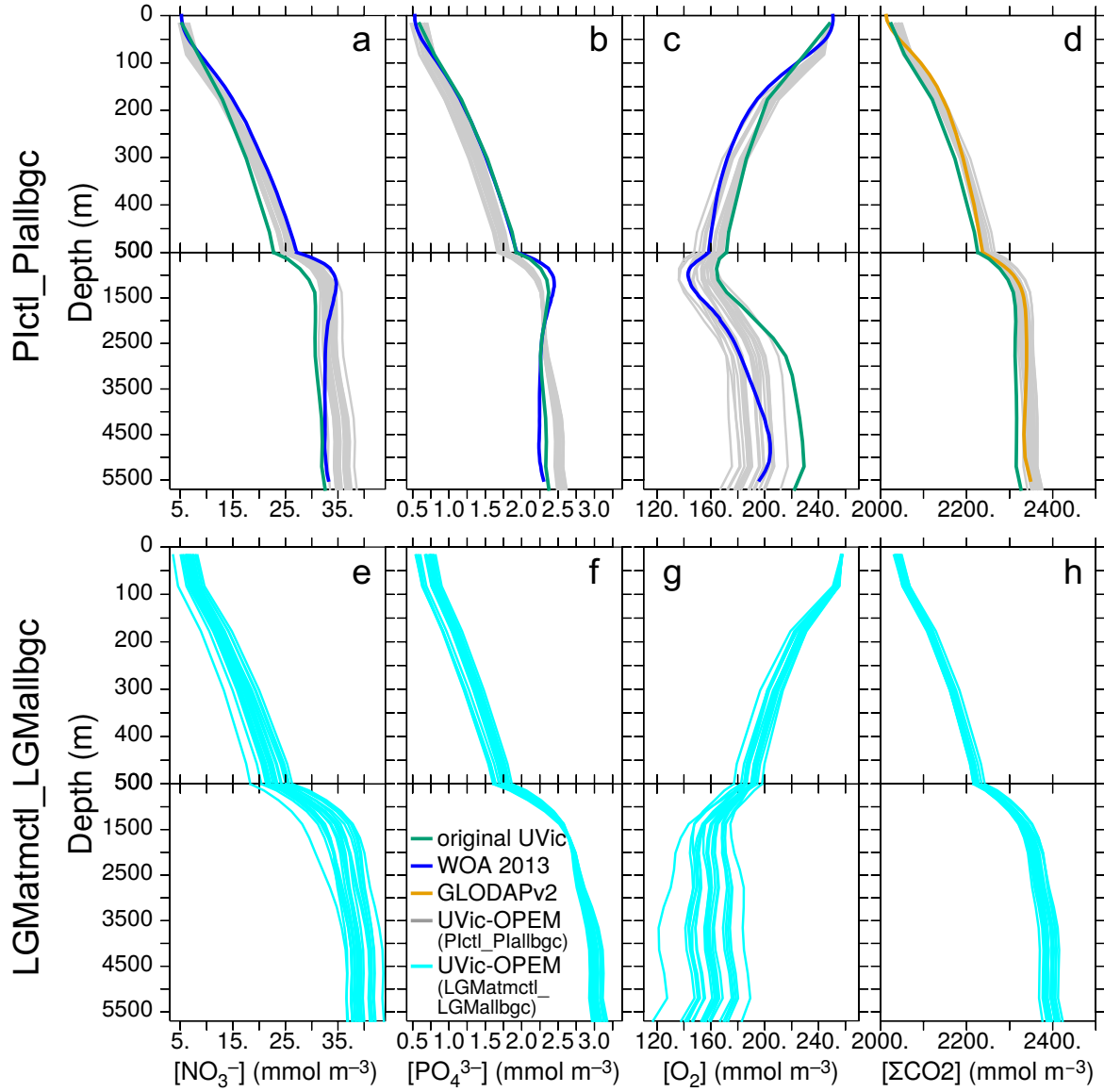
**Figure 3.** Cost values among all 600 simulations used for calibrating UVic-OPEM ordered from low to high. The red part denotes the 20 simulations with the lowest costs.

165 Among the 20 best simulations,  $\text{NO}_3^-$  in the upper 500 m of the model is slightly lower than in WOA 2013 but higher than in the original UVic standard solution. In the deep ocean (below 2500 m), simulated  $\text{NO}_3^-$  concentrations are generally higher than WOA 2013. The globally-averaged concentrations are close to the WOA 2013 average.  $\text{PO}_4^{3-}$  concentrations are slightly lower in the upper ocean but higher in the deep ocean than WOA 2013. Our 20  $\text{O}_2$  profiles scatter around the WOA 2013 profile and are lower and closer to the observations than the original UVic simulation (Mengis et al., 2020). The globally-averaged  
170 concentrations are about 1.7 % higher than WOA 2013. The DIC profiles are higher than in the original UVic and closer to the GLODAPv2 data. The global mean concentrations are 0.1 % higher than GLODAPv2 on average.

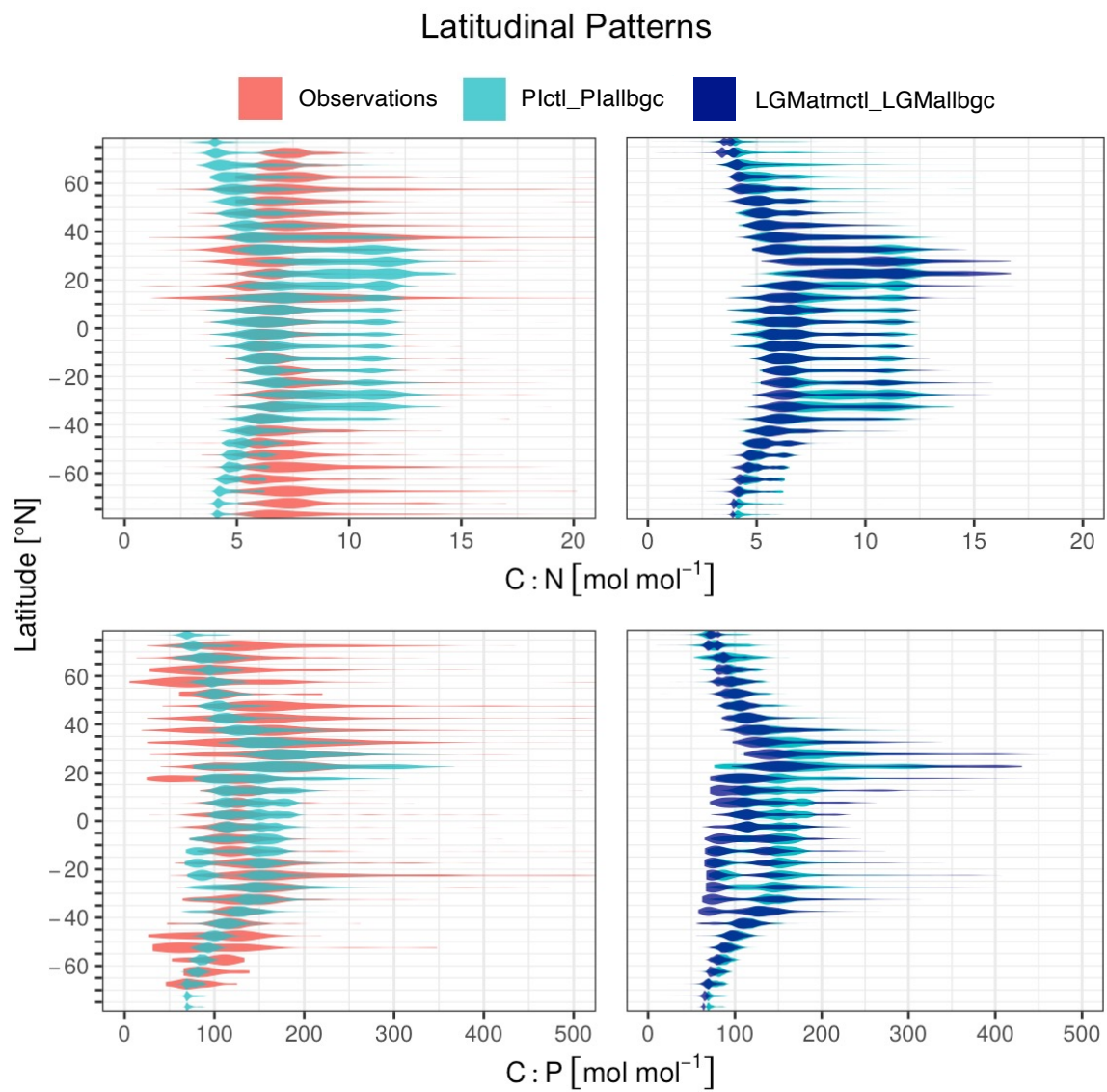
Simulated latitudinal patterns of carbon to nitrogen and carbon to phosphorus ratios in particulate organic matter (pC:N and pC:P, respectively) are also compared with observational data (Tanioka et al., 2022). Overall, the modeled ratios show good agreement with observations (Fig. 5), although both the observational data and model results exhibit substantial variability.  
175 Modeled pC:N is slightly underestimated at high latitudes ( $> 40^\circ$ ), and overestimated at low latitudes, while pC:P in general is well reproduced across latitudes.

### 3.2 Effects of different physical and biogeochemical conditions

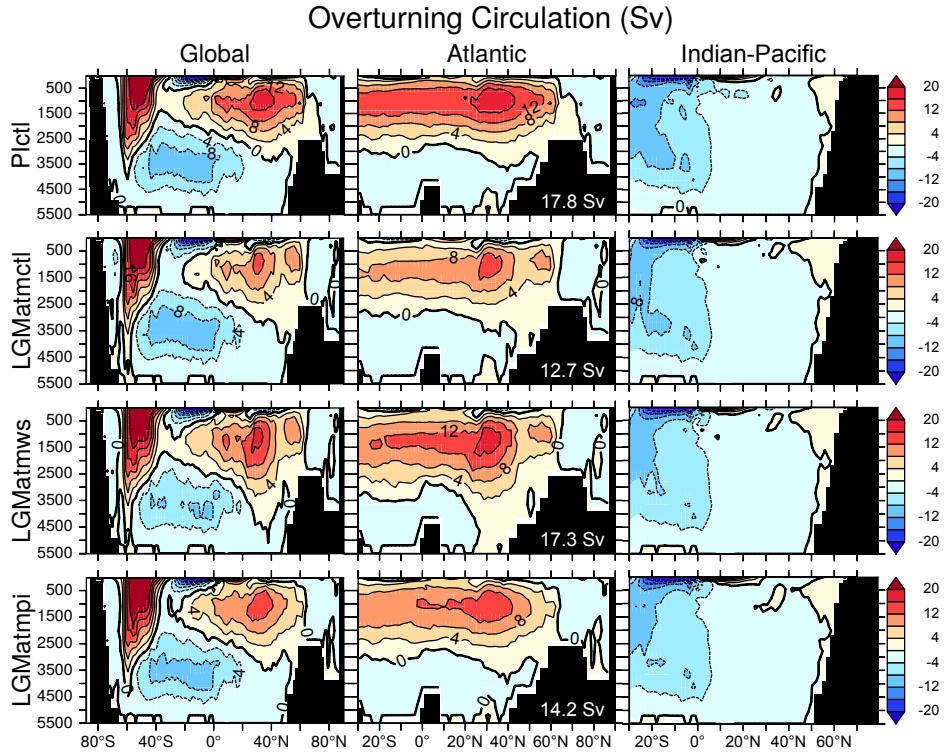
We first compare the simulations with LGM and PI physics, each combined with PI biogeochemistry (PIallbgc) to assess the glacial-interglacial changes in global temperature and the Atlantic Meridional Overturning Circulation (AMOC). We then eval-  
180 uate the effects of different biogeochemical conditions across physical conditions, and vice versa, on marine biogeochemical inventories and atmospheric  $\text{pCO}_2$ .



**Figure 4.** Globally-averaged vertical profiles of  $\text{NO}_3^-$ ,  $\text{PO}_4^{3-}$ ,  $\text{O}_2$ , and  $\text{DIC}$  ( $\Sigma\text{CO}_2$ ) concentrations.  $\text{NO}_3^-$ ,  $\text{PO}_4^{3-}$ , and  $\text{O}_2$  are considered in the cost function. Black lines are results from the 20 Pictl\_PIallbgc simulations, and green lines indicate model results from the original UVic (UVic 2.10, Mengis et al., 2020). Blue and purple lines represent  $\text{NO}_3^-$ ,  $\text{PO}_4^{3-}$ , and  $\text{O}_2$  observational data from the World Ocean Atlas 2013 (WOA 2013, Garcia et al., 2013a, b) and  $\Sigma\text{CO}_2$  data from GLODAPv2 (Key et al., 2015; Lauvset et al., 2016), respectively. Light blue lines in the bottom panels show results from the LGMatmctl\_LGMallbgc simulations.



**Figure 5.** Observed and simulated latitudinal patterns of carbon-to-nitrogen (C:N) and carbon-to-phosphorus (C:P) ratios in particulate organic matter (POM). Observations in the left panels refers to the compilation by Tanioka et al. (2022). Model results represent the mean values from the 20 simulations in Pictl\_Plallbgc and LGMatmctl\_LGMallbgc.



**Figure 6.** Meridional overturning streamfunction in the Global, Atlantic, and Indian-Pacific Oceans for Pictl, LGMatmctl, LGMatmws, and LGMatmpi, all combined with the PIallbgc biogeochemical configuration. Results are ensemble means over the 20 simulations in each of the configurations. Values in white indicate the strength of maximum AMOC at 26.5°N under each condition.

### 3.2.1 Global temperature and ocean circulation

The simulated global surface air temperature in LGMatmctl\_PIallbgc is 4.3°C lower than in Pictl\_PIallbgc. Although studies based on proxy records indicate a larger cooling of approximately 5–6 °C (Tierney et al., 2020; Seltzer et al., 2021), the discrepancy is consistent with known limitations of intermediate-complexity models and does not affect the relative differences analysed in this study. In the ocean, the different physical boundary conditions result in distinct thermohaline circulation patterns (Fig. 6). The strength and depth of the maximum value of the AMOC in Pictl\_PIallbgc at 26.5°N is  $17.82 \pm 0.04$  Sv at 1100m, which agrees well with the observational value from the RAPID array of  $17.2 \pm 0.9$  Sv at the same depth (McCarthy et al., 2015). The strength and depth of the maximum AMOC at 26.5°N in LGMatmctl\_PIallbgc is reduced by 29% compared to Pictl\_PIallbgc (Fig. 6), which is close to a multi-proxy constrained weakening by 36% (Pöppelmeier et al., 2023).

### 3.2.2 Dissolved iron

For the pre-industrial control (Pictl) with PI biogeochemistry (Pictl\_PIallbgc), the ensemble average (employing the 20 lowest-cost parameter sets) of globally-averaged dissolved Fe (dFe) concentration is  $596 \pm 14 \text{ nmol m}^{-3}$ . This value is only slightly lower than the observational estimate of  $624 \text{ nmol m}^{-3}$  for the modern ocean (Huang et al., 2022) (Fig. S2a and Table S1). In the \*\_LGMFedep (\*\_ stands for all four physical conditions combined with LGMFedep) configurations, the increased Fe input from enhanced dust deposition results in higher dFe concentrations than all other configurations. The average concentration is 15 % higher than in the \*\_PIallbgc simulations. In the \*\_LGMFesed simulations, where sedimentary iron fluxes are reduced due to the lower LGM sea level, average dFe concentration is 35 % lower than in the \*\_PIallbgc simulations, and the lower Fe availability has a strong impact on productivity ( $-44 \%$ ) and export of particulate organic carbon (POC,  $-39 \%$ ). The dFe concentrations in the \*\_LGMPO<sub>4</sub> and \*\_LGMbdeni simulations are all similar to those of the Pictl\_PIallbgc baseline, indicating that changes in phosphorus and nitrogen cycling have minimal impact on dFe inventories, with Fe availability remaining relatively stable. In \*\_LGMallbgc simulations, the increased Fe deposition and reduced sedimentary input result in a 1 % decrease in dFe concentration (Fig. S2a and Table S1). Differences in dFe between physical configurations for the same biogeochemistry are much smaller than between biogeochemical configurations (Fig. S2a and S3a). Interestingly, in all LGM physical configurations, dFe is lower than in the corresponding Pictl\_\* configurations (Fig. S3i). In the model, sedimentary Fe input is associated with the POC flux at the sea floor. Since the POC export declines under LGM conditions due to the lower temperature and lower surface nutrients, sedimentary Fe inputs and consequently the dFe inventories are lower in the LGM than in the Pictl simulations.

Surface dFe (sdFe, 0–50 m) shows similar but more accentuated pattern changes compared to globally-averaged dFe, except that the LGMatmctl\_\* configurations have higher sdFe than the other LGM physical configurations for the same biogeochemistry (Fig. S2b and Table S1). The overall variation in sdFe is also similar to that of globally-averaged dFe. In \*\_LGMFedep, the sdFe is 54 % higher than in \*\_PIallbgc, and this difference is greater than for the globally-averaged dFe (Fig. S3a, b and Table S1). These results highlight the significant role of atmospheric Fe deposition in controlling sdFe concentrations.

### 3.2.3 Nitrate

The average NO<sub>3</sub><sup>−</sup> concentration in the \*\_LGMFedep simulations is slightly higher than in \*\_PIallbgc. Nevertheless, it is significantly lower in the \*\_LGMFesed simulations than in \*\_PIallbgc (Figs. S2c and S3c). This strong reduction in \*\_LGMFesed simulations is likely due to a lower iron availability in \*\_LGMFesed, which limits N<sub>2</sub> fixation. Indeed, the global N<sub>2</sub> fixation rates are also significantly lower than for \*\_PIallbgc (Fig. S2k, S4c and Table S2).

The enhanced N<sub>2</sub> fixation due to the greater phosphate availability in \*\_LGMPO<sub>4</sub> and the reduced benthic denitrification in \*\_LGMbdeni both produce higher NO<sub>3</sub><sup>−</sup> inventories than \*\_PIallbgc (Fig. S3c and Table S1). The NO<sub>3</sub><sup>−</sup> inventories in \*\_LGMallbgc on average is  $2.0 \text{ mmol m}^{-3}$  higher than in \*\_PIallbgc across all physical configurations (Fig. S3c and Table S1). Although NO<sub>3</sub><sup>−</sup> inventories in \*\_LGMFesed are reduced, mainly due to a lower sedimentary flux of dFe, dFe supply from atmospheric deposition in \*\_LGMallbgc seems to compensate that effect.

In LGMatmctl\_LGMallbgc simulations, the average  $\text{NO}_3^-$  concentration is 6 % higher than in Pictl\_PIallbgc (Fig. S3s and Table S1), and an increase in deep-water  $\text{NO}_3^-$  concentrations is also observed (Fig. 4). The increase in  $\text{NO}_3^-$  from Pictl\_PIallbgc to LGMatmctl\_LGMallbgc is lower than the estimates based on paleo proxies (>10 %, Glock et al., 2018; Wallmann et al., 2016; Deutsch et al., 2004). Nevertheless, the increase in 5 out of the 20 individual simulations is larger than 10 %, and the maximum increase reaches 16 %.

In addition to the counteracting effects of \*\_LGMallbgc on  $\text{N}_2$  fixation and denitrification, surface  $\text{NO}_3^-$  ( $\text{sNO}_3^-$ ) concentrations are also affected by the strength of the ocean circulation and biological utilisation. Therefore, changes in  $\text{sNO}_3^-$  do not necessarily follow the same pattern as changes in the  $\text{NO}_3^-$  inventory. Across all physical configurations, while the  $\text{NO}_3^-$  inventory in the \*\_LGMFedep simulations is only about 7 % lower than in \*\_PIallbgc, the difference in  $\text{sNO}_3^-$  is 24 % (Fig. S3c and d). On the other hand, the  $\text{NO}_3^-$  inventory in \*\_LGMFesed is about 30 % lower on average than in \*\_PIallbgc, but  $\text{sNO}_3^-$  is only 18 % lower. This apparent inconsistency between changes in  $\text{NO}_3^-$  inventory and the surface concentration reflects different responses to changes in iron supply.

### 3.2.4 Surface phosphate

The mean surface  $\text{PO}_4^{3-}$  ( $\text{sPO}_4^{3-}$ ) in Pictl\_PIallbgc is  $0.62 \pm 0.07 \text{ mmol m}^{-3}$ , which is close to the  $0.56 \text{ mmol m}^{-3}$  in WOA2013 (Garcia et al., 2013b) (Fig. S2e and Table S1). The average  $\text{sPO}_4^{3-}$  in the \*\_LGMFedep simulations is about 17 % lower than in the \*\_PIallbgc simulations, which can be attributed to a higher biological  $\text{PO}_4^{3-}$  utilisation for elevated dFe supply. The average  $\text{sPO}_4^{3-}$  in the \*\_LGMFesed simulations is about 62 % higher than in \*\_PIallbgc, i.e., even higher than in \*\_LGMPO<sub>4</sub> (Fig. S3e and Table S1).

### 3.2.5 Dissolved oxygen

The average dissolved marine oxygen ( $\text{O}_2$ ) concentration in the Pictl\_PIallbgc simulations is  $179 \pm 9 \text{ mmol m}^{-3}$ . The  $\text{O}_2$  inventories in the LGM simulations show considerable variability, with some simulations exhibiting significant deviations from \*\_PIallbgc (Fig. S3f).  $\text{O}_2$  in the \*\_LGMFedep simulations is  $22 \text{ mmol m}^{-3}$  lower than in the \*\_PIallbgc simulations. This decrease is primarily driven by higher oxygen consumption due to the increased remineralisation of particulate organic carbon (POC), a shift that is also consistent with increased water column and benthic denitrification. Conversely, the \*\_LGMFesed configurations, which feature reduced iron input, result in a remarkable increase by  $62 \text{ mmol m}^{-3}$  in  $\text{O}_2$  concentrations (Fig. S3f).

The \*\_LGMPO<sub>4</sub> and \*\_LGMbdeni simulations produce oxygen concentrations only slightly below those in \*\_PIallbgc. Hence, changes in  $\text{PO}_4^{3-}$  and  $\text{NO}_3^-$  availability alone have limited effects on the  $\text{O}_2$  inventory. Overall, the average  $\text{O}_2$  in \*\_LGMallbgc simulations is about 2 % lower than in \*\_PIallbgc.

Among the LGM physical configurations,  $\text{O}_2$  levels are higher in the upper ocean due to a higher solubility under colder conditions, but are lower in the deep water due to a more sluggish circulation (Fig. 4c and g), and the globally-averaged concentrations generally are lower than in the respective Pictl\_\* simulations. For example, the average concentration for LGMatmctl\_\* is 2.8 % lower than for Pictl\_\*. Comparing LGMatmctl\_LGMallbgc to Pictl\_PIallbgc, global  $\text{O}_2$  decreases by  $10 \text{ mmol m}^{-3}$ , which is due to the depletion in the deep water, while the upper ocean concentrations are typically higher

(Fig. 4). This agrees with estimated LGM O<sub>2</sub> levels based on paleo proxies (Jaccard and Galbraith, 2012; Anderson et al., 2019) and model simulations (Bopp et al., 2017; Somes et al., 2017). An exception is the \*\_LGMFesed simulations, where O<sub>2</sub> levels are higher than in the Pictl\_\* simulations, which indicates a strong coupling with dFe reduction and changes in physical boundary conditions (Fig. S3n).

### 3.2.6 Dissolved inorganic carbon and atmospheric pCO<sub>2</sub>

In the \*\_LGMFedep simulations, globally-averaged DIC is 3.4 mmol m<sup>-3</sup> higher than in the corresponding \*\_PIallbgc simulations (Fig. S3g). In contrast, DIC in the \*\_LGMFesed simulations is 7.8 mmol m<sup>-3</sup> lower than in the \*\_PIallbgc simulations (Fig. S3g). The \*\_LGMPO<sub>4</sub> and \*\_LGMbdeni simulations yield DIC concentrations similar to those of the pre-industrial control. Our LGM simulations yield higher DIC concentrations than the Pictl\_\* simulations throughout the different biogeochemical configurations (Fig. S3o). Also, the increase in DIC is mostly observed in the deep water (Fig. 4). Further, DIC concentrations within each of the LGM physical configurations are similar, despite the different strength of overturning circulation (Fig. 6 and S3g).

The glacial-interglacial changes in the strength of the solubility pump caused by a reduced temperature, for example, can be obtained by comparing simulated results from Pictl\_PIallbgc and LGMatmpi\_PIallbgc. The latter shows an average increase of 0.6 % in DIC, which is equivalent to 238 Pg of carbon.

Atmospheric pCO<sub>2</sub> in the biogeochemically coupled but radiatively uncoupled 20 Pictl\_PIallbgc simulations is 284.4 ± 0.1 ppm, consistent with the radiatively prescribed 284.3 ppm that was used for the calibration stage (Fig. S2h and Table S1). This indicates equilibrated carbon fluxes between air, land and ocean carbon pools at the end of the spin-ups. Switching to LGM Fe fluxes has a relatively strong effect on pCO<sub>2</sub>, when compared to the effects of the LGM PO<sub>4</sub><sup>3-</sup> inventory or benthic denitrification. For each of the 4 physical configurations (Pictl, LGMatmctl, LGMatmws, and LGMatmpi, Table 2a), the difference in pCO<sub>2</sub> of employing the biogeochemical LGM forcing conditions with respect to the biogeochemical \*\_PIallbgc is most pronounced for \*\_LGMFesed, where pCO<sub>2</sub> increases by 60 ppm, and \*\_LGMFedep, showing a decrease by 26 ppm (Fig. S3h and Table S1).

The average atmospheric pCO<sub>2</sub> in the physical LGM configurations, LGMatmctl, LGMatmws, and LGMatmpi, are all lower than in the respective simulations for Pictl across all biogeochemical configurations (Fig. S2h and S3p). The pCO<sub>2</sub> in the 20 LGMatmctl\_LGMallbgc simulations is 241 ppm on average, which is significantly higher than the LGM atmospheric pCO<sub>2</sub> of approximately 190 ppm recorded in ice cores (Monnin et al., 2001). Thus, the specific mechanisms isolated in these configurations can account for only a portion of the full glacial-interglacial pCO<sub>2</sub> drawdown.

Atmospheric pCO<sub>2</sub> shows different levels of variability among the 20 simulations within each of the biogeochemical configurations. The spread in pCO<sub>2</sub> (i.e., the difference between the maximum and minimum values across simulations) is largest for \*\_LGMFesed, ranging from 26 ppm to 31 ppm (Fig. S2h). The atmospheric pCO<sub>2</sub> is also influenced by changes in the terrestrial carbon pool. While all LGM physical configurations lead to reduced non-glaciated terrestrial carbon (35% lower than in Pictl condition) due to changes in radiative forcing, the three LGM atmospheric configurations result in different surface temperature distributions (Fig. S5), which in turn affect the amount of carbon released to the atmosphere and ocean. The combined

carbon loss from land and the atmosphere to the ocean in the LGMatmctl, LGMatmws, and LGMatmpi simulations is  $251 \pm 15$ ,  $234 \pm 13$ , and  $239 \pm 9$  Pg C, respectively, relative to Pictl. Of these, changes in atmospheric carbon ( $1 \text{ ppm} = 2.123 \text{ PgC}$ ) account for approximately 30 %, 18 %, and 17 % in the LGMatmctl, LGMatmws, and LGMatmpi simulations, respectively.

### 3.3 Biogeochemical rate estimates and elemental composition of POM

#### 3.3.1 Marine NPP and POC export

Net primary production (NPP) in Pictl\_Plallbgc ranges from 44 to 80 Pg C yr<sup>-1</sup>, with mean and standard deviation of  $64 \pm 8$  Pg C yr<sup>-1</sup> (Fig. S2i and Table S2), in line with observational estimates ( $36 - 77 \text{ Pg C yr}^{-1}$ , Carr et al., 2006; Honjo et al., 2008; Buitenhuis et al., 2013). Increasing Fe deposition enhances NPP by about  $2.8 \text{ Pg C yr}^{-1}$  in the \*\_LGMFedep simulations when compared with \*\_Plallbgc. A strong reduction in NPP is associated with the reduced sedimentary Fe flux in the \*\_LGMFesed simulations, where NPP is about 44 % lower on average than in \*\_Plallbgc (Fig. S4a). The NPP increases by  $1.5 \text{ Pg C yr}^{-1}$  in \*\_LGMPO<sub>4</sub> and  $1.8 \text{ Pg C yr}^{-1}$  in \*\_LGMbdeni (Fig. S4a). When combined with the biogeochemical conditions, NPP in \*\_LGMallbgc is 11 % lower than in \*\_Plallbgc. Due to the lower temperature, NPP decreases by 18 % in LGMatmpi\_\* compared to Pictl\_\* (Fig. S4i). Compared with the effect of the lower temperature, switching to LGM moisture diffusivity and wind patterns has only relatively small effects on NPP. Compared with LGMatmpi\_\*, NPP increases by 3 % and 1 % in LGMatmws\_\* and LGMatmctl\_\* simulations, respectively.

Particulate organic carbon (POC) export shows a similar pattern to that of NPP. The average flux in the Pictl\_Plallbgc simulations is  $9.2 \pm 0.6 \text{ Pg C yr}^{-1}$  (Fig. S2j and Table S2). POC export increases by 9 % for \*\_LGMFedep and decreases by 39 % for \*\_LGMFesed, with only small changes in \*\_LGMPO<sub>4</sub>, \*\_LGMbdeni, and \*\_LGMallbgc when compared with \*\_Plallbgc (Fig. S4b). The reduction in POC export due to a cooler climate is 5 % among the LGM physical configurations. This temperature-driven reduction reflects the close coupling between global NPP and export production in the model, while regional changes in iron supply and nutrient utilisation can modulate export production. Proxy-based studies indicate that changes in export production during the LGM were spatially heterogeneous with decreases in some low-latitude regions but increases in areas affected by enhanced iron supply, such as the Southern Ocean (Cartapanis et al., 2018; Kohfeld et al., 2005; Toyos et al., 2022).

#### 3.3.2 Marine N<sub>2</sub> fixation and water-column and benthic denitrification

N<sub>2</sub> fixation rates in Pictl\_Plallbgc range from 132 to 281 Tg N yr<sup>-1</sup> (Fig. S2k), agreeing with observations ( $223 \pm 30 \text{ Tg N yr}^{-1}$ , Shao et al., 2023), and inverse-model estimates ( $126 - 223 \text{ Tg N yr}^{-1}$ , Wang et al., 2019). WC denitrification (WC N-loss) rates vary strongly among the Pictl\_Plallbgc simulations, ranging from 8 to 119 Tg N yr<sup>-1</sup>, and benthic denitrification rates range from 116 to 169 Tg N yr<sup>-1</sup> (Fig. S2l and m), also in line with previous estimates (39 to 66 and 68 to 122 Tg N yr<sup>-1</sup> for WC and benthic denitrification rates, respectively, Eugster et al., 2013).

In the model, N<sub>2</sub> fixation is the counterpart of WC and benthic denitrification, thus N<sub>2</sub> fixation equals the sum of WC and benthic denitrification at equilibrium. Across our physical configurations, N<sub>2</sub> fixation increases by 37 % on average in

\*\_LGMFedep simulations compared with \*\_PIallbgc, due to the extra input of Fe relieving iron limitation of diazotrophs (Fig. S4c). WC denitrification is 197 % higher (Fig. S4d), compensating for most of the increase in N<sub>2</sub> fixation, and benthic  
325 denitrification only increases by 5 % (Fig. S4e). N<sub>2</sub> fixation drops by 53 % on average in the \*\_LGMFesed simulations, while benthic denitrification decreases by 43 % and WC denitrification shuts down entirely (Fig. S2l).

In the \*\_LGMPO<sub>4</sub> simulations, N<sub>2</sub> fixation and the sum of denitrification increase by 5 % compared to \*\_PIallbgc (Fig. S4c). Benthic denitrification decreases by 31 % on average in the \*\_LGMbdeni simulations, while WC denitrification increases by 52 %, and N<sub>2</sub> fixation is 18 % lower than in \*\_PIallbgc. In \*\_LGMallbgc, N<sub>2</sub> fixation and WC and benthic N denitrification  
330 decrease by 31 %, 22 %, and 33 %, respectively (Fig. S4c, d and e).

### 3.3.3 Elemental composition of POM

The elemental ratios of particulate organic matter (POM) provide insights into nutrient utilisation efficiency and the coupling between carbon, nitrogen, and phosphorus in marine ecosystems. Here we analyse the particulate carbon to nitrogen (pC:N), carbon to phosphorus (pC:P), and nitrogen to phosphorus (pN:P) ratios. The ecological elemental ratios do not follow normal  
335 distributions, and we calculate medians of the elemental ratios without biomass weighting in the ocean grid cells that are not covered by ice, rather than mean values to avoid unnecessary bias (Isles, 2020). The global median of surface (0 – 50 m) pC:N, pC:P, and pN:P in the PIctl\_PIallbgc simulations are  $8.0 \pm 0.5 \text{ mol mol}^{-1}$ ,  $134 \pm 8 \text{ mol mol}^{-1}$ , and  $16.5 \pm 0.7 \text{ mol mol}^{-1}$ , respectively (Fig. S2n–p and Table S2). When comparing different biogeochemical conditions to \*\_PIallbgc, the change in globally-averaged pC:N is negatively correlated with the change in  $\text{sNO}_3^-$  (Figs. S3d and S4f). Nevertheless, the pC:N is  
340 lower despite lower  $\text{sNO}_3^-$  in \*\_LGMFesed (Tables S1 and S2) because of the shrinking area with low  $\text{sNO}_3^-$ , caused by low  $\text{NO}_3^-$  utilisation under strong iron limitation, such as the South Pacific Ocean, and hence the area with high pC:N also becomes smaller (Fig. S6). In general, pC:N in the model is mainly affected by two factors. One is the Fe limitation of carbon fixation and  $\text{NO}_3^-$  utilisation, and the other is the supply of  $\text{NO}_3^-$  to the surface ocean.

The relation between pC:P and surface  $\text{PO}_4^{3-}$  ( $\text{sPO}_4^{3-}$ ) is similar to the relation between pC:N and  $\text{sNO}_3^-$  (Fig. S2o). Since  
345 the  $\text{PO}_4^{3-}$  inventory is constant in the model,  $\text{sPO}_4^{3-}$  is largely affected by the supply of dFe. In LGMatmctl\_LGMallbgc simulations,  $\text{sPO}_4^{3-}$  and  $\text{sNO}_3^-$  are higher than in the PIctl\_PIallbgc (Fig. S3t and u), and the pC:N and pC:P are generally lower, particularly in low latitude regions where the surface nutrients are higher (Fig. 5). In \*\_LGMFesed simulations  $\text{sPO}_4^{3-}$  increases with lower iron supply. As a result, pC:P in \*\_LGMFesed simulations decreases by  $54 \text{ mol mol}^{-1}$  (39 %), while pC:N decreases by only  $1.3 \text{ mol mol}^{-1}$  (16 %) when compared with \*\_PIallbgc (Fig. S4g and f).

350 Compared with \*\_PIallbgc, the average  $\text{sPO}_4^{3-}$  increases by 31% in \*\_LGMPO<sub>4</sub> and by 62% in \*\_LGMFesed. Moreover, the effects of Fe limitation on carbon fixation and nitrogen assimilation are weaker in \*\_LGMPO<sub>4</sub> than in \*\_LGMFesed. As a result, pC:P and pN:P decrease by 10% and 6%, respectively, in \*\_LGMPO<sub>4</sub>, which is substantially smaller than the reductions found in \*\_LGMFesed.

Due to the stronger increase in pC:P relative to pC:N, pN:P decreases by 31 % in \*\_LGMFesed (Fig. S4h), while other  
355 biogeochemical LGM configurations have smaller effects. Physical boundary conditions also contribute to the changes in the

elemental ratios of POM via effects on  $\text{sNO}_3^-$  and  $\text{sPO}_4^{3-}$ . The pC:N and pC:P in general are highest for LGMatmctl and lowest for PIctl and LGMatmws, and pN:P is highest in the LGMatmctl configuration (Fig. S4n–p).

## 4 Discussion

### 4.1 Physical boundary conditions and ocean circulation

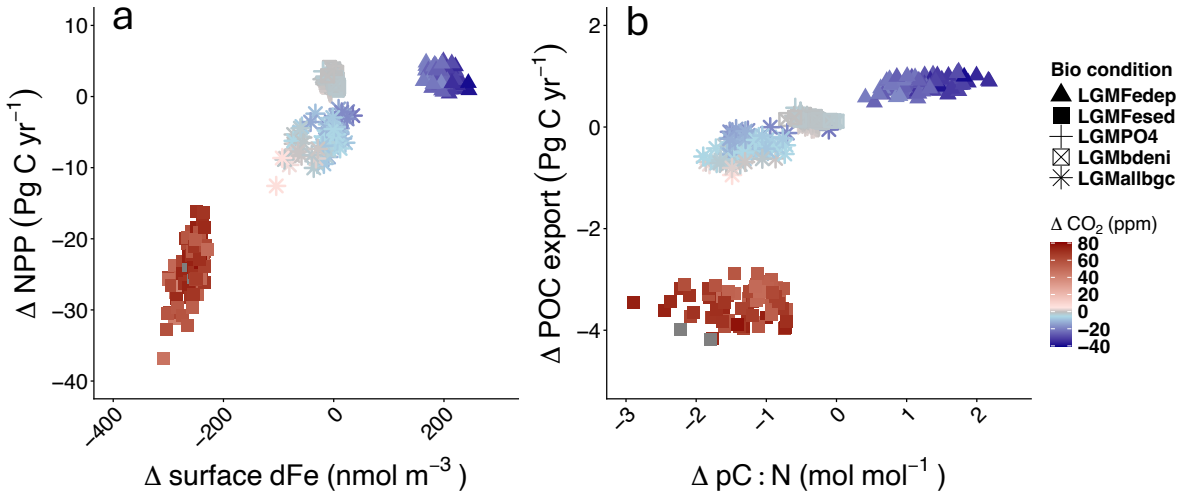
360 The UVic-ESCM responds to LGM wind stress changes with an increased northward salt transport in the North Atlantic, which enhances surface salinity and density, thereby strengthening North Atlantic Deep Water formation while intensifying and deepening the AMOC. In contrast, reduced atmospheric moisture diffusivity decreases meridional freshwater transport to the Southern Ocean, leading to changes in surface buoyancy and enhanced vertical stratification. This suppresses deep water formation, particularly of Antarctic Bottom Water, which weakens the deep return flow, resulting in a weaker and shallower  
365 global overturning circulation, including the AMOC (Sigman et al., 2007; Somes et al., 2017).

The model results show only small changes in the MOC of the Indian and Pacific Oceans, with close agreement between LGMatmctl\_Plallbgc and PIctl\_Plallbgc. This indicates that the effect of the LGM lower surface temperature is almost compensated by the LGM winds and atmospheric moisture diffusivity. As a consequence of the virtual insensitivity of the Indo-Pacific MOC to a switch from pre-industrial to LGM boundary conditions, changes in the global MOC essentially mirror the changes  
370 in the AMOC (Fig. 6).

### 4.2 Effects of biogeochemical boundary conditions on biogeochemistry

Among the four glacial-interglacial changes in biogeochemical boundary conditions investigated, iron supply has the strongest impact on the inventories and fluxes. Clearly, it is the changes in Fe availability within the surface layer that alter iron limitation and affect NPP and POC export (Fig. 7). The lower dFe inventory during the LGM seems in conflict with the general  
375 understanding of Fe fertilisation during the LGM (Martin, 1990; Martínez-García et al., 2014). Interestingly, although NPP and POC export are lower in the \*\_LGMallbgc relative to the \*\_Plallbgc simulations, the corresponding atmospheric  $\text{pCO}_2$  in general is lower as well (Fig. 7 and Table. S1). The atmospheric  $\text{pCO}_2$  in the \*\_LGMallbgc simulations is lower compared to \*\_Plallbgc, and is primarily driven by the redistribution of surface DIC and the associated changes in global air–sea  $\text{CO}_2$  fluxes, just before equilibrium is re-established (Fig. S7).

380 Glacial-interglacial changes in sedimentary Fe input were not considered in several earlier studies (Somes et al., 2017; Ödalen et al., 2020; Matsumoto et al., 2020a, b; Vollmer et al., 2022). Nevertheless, the strong sensitivity of the biogeochemical inventories and fluxes to changes in sedimentary influx of Fe demonstrates its importance. The reduction in sedimentary input of Fe from \*\_Plallbgc to \*\_LGMFesed ( $10.3 \text{ Gmol Fe yr}^{-1}$ ) is greater than the increase in atmospheric Fe deposition from \*\_Plallbgc to \*\_LGMFedep ( $4.7 \text{ Gmol Fe yr}^{-1}$ ), resulting in smaller dFe inventories in \*\_LGMallbgc. At first glance, one  
385 might expect that the magnitude of sedimentary Fe input in the model is relatively large and that its reduction could therefore be overestimated. However, the simulated sedimentary Fe flux in PIctl\_Plallbgc ( $12.6 \text{ Gmol Fe yr}^{-1}$ ) falls close to the lower



**Figure 7.** Differences in (a) NPP and surface dFe and (b) POC export and pC:N with respect to default biogeochemical conditions (Pllallbgc). Color represents atmospheric pCO<sub>2</sub> and symbols indicate the different biogeochemical configurations. The cluster of points with little deviation from \*\_Pllallbgc simulations is from \*\_LGMPO4 and \*\_LGMbdeni simulations.

end of estimates from other models (0.6–194 Gmol Fe yr<sup>-1</sup>, Tagliabue et al., 2016). A recent sensitivity study also shows that a much higher present-day sedimentary Fe release (117 Gmol Fe yr<sup>-1</sup>) than assumed previously (15 Gmol Fe yr<sup>-1</sup>) yields better model-data misfit in global and surface dFe distribution (Somes et al., 2021). This indicates that our representation is relatively conservative, and does not impose an unrealistically large reduction in sedimentary Fe supply.

While dFe availability affects N<sub>2</sub> fixation, simulations with higher dFe supply and concentrations do not necessarily result in higher NO<sub>3</sub><sup>-</sup> inventories. For example, the average NO<sub>3</sub><sup>-</sup> concentration for \*\_LGMFedep is 2.0 mmol m<sup>-3</sup> lower than for \*\_Pllallbgc. This is because changes in Fe supply in our simulations do not only affect N<sub>2</sub> fixation, but can induce changes in denitrification, which in turn depends on the level and consumption of oxygen, in association with the attenuation of the exported particulate organic matter (POM). Two NO<sub>3</sub><sup>-</sup> sinks can occur in the model solutions. One sink is due to the water-column (WC) denitrification, which is affected by WC O<sub>2</sub> supply and consumption via POM remineralisation. The other is benthic denitrification, which is sensitive to the amount of POC reaching the sea floor. Thus, both water column and benthic denitrification are affected by the POM export. The higher Fe supply in the surface ocean increases not only N<sub>2</sub> fixation but also NPP and POM export, thereby promoting both WC and benthic denitrification. The net effect on the NO<sub>3</sub><sup>-</sup> inventory depends on the balance of these counteracting fluxes. Whenever NO<sub>3</sub><sup>-</sup> concentrations are lower in a simulation in \*\_LGMFedep than in \*\_Pllallbgc, the increase in denitrification is higher than the increase in N<sub>2</sub> fixation at the beginning of the spin-up, because the increase in N<sub>2</sub> fixation due to a higher Fe supply is hampered by the supply of the other limiting macronutrient, PO<sub>4</sub><sup>3-</sup>. Under such conditions, the model yields results with a lower NO<sub>3</sub><sup>-</sup> inventory once the simulation reaches equilibrium despite an

increase in  $N_2$  fixation. This explains that while global  $N_2$  fixation in \*\_LGMFedep simulations is higher than for \*\_Plallbgc  
405 (Fig. S4c and Table S2), the  $NO_3^-$  inventories in \*\_LGMFedep show a mixed response (Fig. S3c and Table S1).

Since the global  $PO_4^{3-}$  inventory is fixed in the model, the higher  $PO_4^{3-}$  levels in the LGM $PO_4$  configuration directly  
translate into higher surface concentrations. In the \*\_LGM $PO_4$  simulations,  $sPO_4^{3-}$  on average is about 31 % higher than  
in \*\_Plallbgc, which is more than the 15 % increase in the inventory, indicating a non-linear relationship between  $sPO_4^{3-}$   
and its global distribution (Figs. 4, S3e and Table S1). Changes in ocean circulation also affect  $sPO_4^{3-}$ . For example, for all  
410 the biogeochemical configurations,  $sPO_4^{3-}$  for LGMatmws, LGMatmpi and particularly LGMatmctl are lower than for Plctl  
(Fig. S3m).

While earlier works have hypothesised that increased  $sPO_4^{3-}$  and  $sNO_3^-$  during the LGM could have benefited NPP and  
decreased  $pCO_2$  (Broecker, 1982; Wallmann, 2010; Wallmann et al., 2016), the changes of  $sPO_4^{3-}$  in \*\_LGM $PO_4$  and  $sNO_3^-$   
in \*\_LGMbdeni have only limited effects on the NPP and  $pCO_2$  in our simulations. The relatively weak effects of the increases  
415 in the two major nutrients can be ascribed to (1) co-limitation effects and (2) the variable stoichiometry in UVic-OPEM.  $PO_4^{3-}$ ,  
 $NO_3^-$ , and Fe are the three limiting nutrients for phytoplankton growth in the model, and nutrient co-limitation is observed  
in many regions of the ocean (Browning et al., 2017; Browning and Moore, 2023). An increase in  $PO_4^{3-}$  or  $NO_3^-$  alone  
would trigger or aggravate limitation by the other two nutrients. The flexibility in the carbon to nitrogen (C:N) and carbon  
to phosphorous (C:P) ratios of phytoplankton in UVic-OPEM further complicates the relationship between NPP and nutrient  
420 limitation. When  $sPO_4^{3-}$  or  $sNO_3^-$  increase, pC:P and pC:N decrease in response (Fig. S2 and Table S2), and this partly offsets  
the effect of increasing major nutrients on NPP, POC export, and  $pCO_2$  (Fig. 7).

As a consequence, macronutrient availability alone exerts only a limited control on export production and associated oxygen  
consumption in our simulations. Instead, iron availability emerges as the more influential limiting factor, in particular in regions  
where Fe supply limits productivity. Changes in Fe supply therefore have a strong impact on POC export, remineralisation,  
425 and ultimately the  $O_2$  inventory.

A stronger Fe supply from atmospheric deposition, such as in the LGMFedep condition, could have enhanced the utilisation  
of  $PO_4^{3-}$  and  $NO_3^-$ , leading to an expansion of ocean regions depleted in those major nutrients. This 'nutrient robbing' effect,  
whereby stimulated productivity in one area depletes macronutrients in downstream regions, is a primary criticism of Ocean  
Iron Fertilisation as a marine CDR strategy, as it could limit long-term global net carbon sequestration. Nevertheless, our results  
430 suggest that the variable stoichiometry may partially mitigate this effect. By increasing pC:P and pC:N in response to nutrient  
stress, our model solutions sustain NPP even when macronutrient concentrations decline. This negative feedback mechanism  
does not exist in models that assume fixed stoichiometry, and warrants further investigation to determine its global significance  
in CDR scenarios.

The higher DIC in the \*\_LGMFedep simulations can be attributed to increased primary production and carbon export driven  
435 by the Fe fertilisation effect, leading to greater uptake of  $CO_2$  from the atmosphere and increased carbon storage in the ocean.  
In contrast, the lower DIC in the \*\_LGMFesed simulations is likely linked to reduced primary production and carbon export,  
as Fe availability is a critical limiting factor for these processes. While global average DIC concentrations appear similar across  
the LGM physical configurations, the vertical distribution tells a more complex story (Fig. S8). Specifically, deep-water DIC

concentrations vary with the circulation state; the LGMatmctl configuration, which exhibits the most sluggish AMOC in our  
440 ensemble, shows the highest deep-water DIC inventory. This suggests that the relationship between circulation strength and  
DIC storage is not straightforward and likely involves additional factors such as circulation geometry and deep ocean residence  
time.

Our results emphasise the importance of iron as a limiting nutrient, especially during the LGM, when changes in dust  
deposition and sea level had significant impacts on Fe availability. We also demonstrate that the lower temperatures during  
445 the LGM should have reduced NPP and POC export and thus the sedimentary flux of Fe, which also affects the oceanic dFe  
inventory.

### 4.3 Interactions between physical and biogeochemical boundary conditions

Lower temperatures, different wind patterns, and reduced moisture diffusivity over the Southern Ocean in the three LGM  
physical configurations result in different general circulation patterns, which affect biogeochemical cycles, as demonstrated  
450 in previous modelling studies (Somes et al., 2017; Muglia and Schmittner, 2015). The LGM boundary conditions affect in-  
ventories and fluxes differently. For example, DIC is  $14.7 \text{ mmol m}^{-3}$  higher for LGMatmpi\_\* than for PIctl\_\*, while the dif-  
ferences between LGMatmws\_\* and LGMatmpi\_\*, and between LGMatmctl\_\* and LGMatmpi\_\* are only  $-0.3 \text{ mmol m}^{-3}$   
and  $0.8 \text{ mmol m}^{-3}$ , respectively (Fig. S3o). On the other hand,  $\text{N}_2$  fixation shows contrasting responses across the simulations.  
Relative to PIctl\_\*, it decreases by  $19 \text{ Tg N yr}^{-1}$  in LGMatmpi\_\*, but increases by 20 and  $18 \text{ Tg N yr}^{-1}$  in LGMatmws\_\* and  
455 LGMatmctl\_\*, respectively. This indicates that, in our experiments, the glacial-interglacial changes in the DIC inventory are  
dominated by the lower temperature, while  $\text{N}_2$  fixation is also sensitive to the changes in the physical conditions.

Across all biogeochemical conditions, our LGM configurations have lower  $\text{pCO}_2$  by  $-35$ ,  $-21$ , and  $-19$  ppm in LGMatmctl\_\*,  
LGMatmws\_\*, and LGMatmpi\_\*, respectively. Thus, the LGM wind pattern and moisture diffusivity over the Southern Ocean  
contribute about 16 ppm, similar to the 19 ppm drawdown of  $\text{pCO}_2$  owing to the lower temperature.

460 We also notice that the effects of the biogeochemical configurations depend on which physical configuration they are com-  
bined with and vice versa. For example, the  $\text{NO}_3^-$  deviations from PIctl\_\* are more strongly expressed in the \*\_LGMFedep  
simulations than in the other biogeochemical configurations (Fig. S3k and Table S1). This highlights how an increase in dFe  
supply enhances the sensitivity of  $\text{N}_2$  fixation and denitrification to changes in physical boundary conditions. The results  
underscore the importance of the interplay between the biogeochemical and physical boundary conditions in controlling the  
465 balance of  $\text{NO}_3^-$  gains and losses in the ocean. Also,  $\text{pCO}_2$  decreases by 0.8 ppm from PIctl\_PIallbgc to PIctl\_LGMallbgc, by  
9.3 ppm from LGMatmctl\_PIallbgc to LGMatmctl\_LGMallbgc, by 34.2 ppm from PIctl\_PIallbgc to LGMatmctl\_PIallbgc, and  
by 42.7 ppm from PIctl\_LGMallbgc to LGMatmctl\_LGMallbgc. This non-linear interaction between physical circulation and  
biogeochemical processes is consistent with previous studies showing that the glacial  $\text{pCO}_2$  drawdown arises from the com-  
bined and often synergistic effects of ocean circulation changes and biological processes such as Fe fertilisation (Tagliabue  
470 et al., 2009; Buchanan et al., 2016).

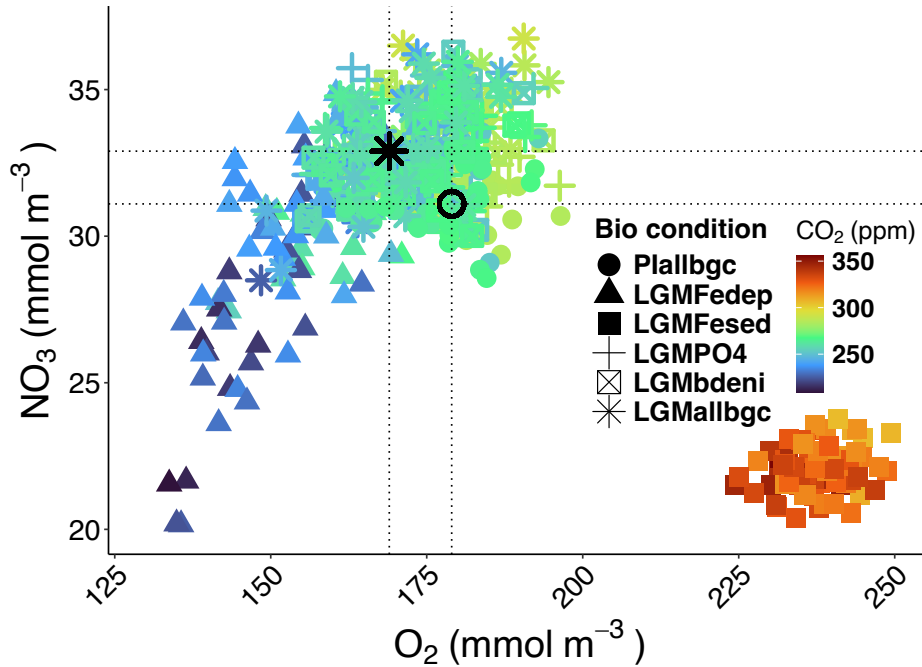
#### 4.4 $\text{NO}_3^-$ and $\text{O}_2$ levels as constraints for simulating LGM conditions

During the LGM,  $\text{O}_2$  and  $\text{NO}_3^-$  levels and distributions in the ocean were different from the pre-industrial era. According to paleo proxy data,  $\text{O}_2$  was lower during the LGM (Jaccard and Galbraith, 2012; Jacobel et al., 2020; Anderson et al., 2019), and  $\text{NO}_3^-$  was higher (Deutsch et al., 2004; Wallmann et al., 2016; Glock et al., 2018). Globally-averaged  $\text{O}_2$  and  $\text{NO}_3^-$  in LGMatmctl\_LGMallbgc simulations are  $10 \text{ mmol m}^{-3}$  lower and  $1.8 \text{ mmol m}^{-3}$  higher on average than in PIctl\_PIallbgc (Table S1). In some of the \*\_LGMFedep simulations,  $\text{O}_2$  and  $\text{NO}_3^-$  levels both become lower than in PIctl\_PIallbgc (Fig. 8). In these simulations, high iron input from the atmosphere increases POM production and export, and the ensuing consumption of  $\text{O}_2$  causes more widespread oxygen deficient zones, where denitrification occurs.

Such a decrease in  $\text{NO}_3^-$  as simulated by some of the experiments is in conflict with the observational records. Thus, a low  $\text{NO}_3^-$  inventory indicates that a strong decrease in  $\text{pCO}_2$  might be due to the wrong reasons in the model, e.g., overestimation of the marine biological carbon pump. On the other hand, the strong Fe limitation in \*\_LGMFesed simulations results in a decrease in  $\text{NO}_3^-$  due to suppressed  $\text{N}_2$  fixation, an increase in  $\text{pCO}_2$  due to reduced primary production, and higher  $\text{O}_2$  concentrations as a result of lower  $\text{O}_2$  consumption from POM export (Fig. 8). This increase in  $\text{O}_2$  is also inconsistent with proxy-based reconstructions. Those different  $\text{NO}_3^-$ ,  $\text{O}_2$ , and  $\text{pCO}_2$  values highlight the critical role of Fe supply for the LGM model experiments.

#### 4.5 Changes in atmospheric $\text{pCO}_2$ and elemental ratio shifts in POM

In the LGMatmctl\_LGMallbgc configuration, the average  $\text{pCO}_2$  is 43.5 ppm lower than in PIctl\_PIallbgc (Fig. S3x and Table S1). The biogeochemical boundary conditions combined contribute less than the physical boundary conditions. The changes in  $\text{pCO}_2$  from \*\_PIallbgc to \*\_LGMallbgc (biogeochemical) with PIctl and LGMatmctl configurations are  $-0.8$  and  $-9.3$  ppm, respectively, while the changes from PIctl to LGMatmctl (physical) with \*\_PIallbgc and \*\_LGMallbgc conditions are  $-34.2$  and  $-42.7$  ppm, respectively (Table S1). The decreasing  $\text{pCO}_2$  due to the LGM physical configuration with \*\_PIallbgc ( $-34.2$  ppm, PIctl\_PIallbgc to LGMatmctl\_PIallbgc) agrees with a 33 ppm decrease in a previous model experiment considering changes in physical boundary conditions (Ödalen et al., 2020). Nevertheless, the contribution from the LGM biogeochemical boundary conditions (\*\_PIallbgc to \*\_LGMallbgc) to the decrease in  $\text{pCO}_2$  is small, and the 43.5 ppm decrease from PIctl\_PIallbgc to LGMatmctl\_LGMallbgc are only about 50 % of the observed ( $\approx 90$  ppm). The main goal of our experiments is to evaluate changes in some biogeochemical and physical aspects that could affect  $\text{pCO}_2$ , and not all processes, such as brine-induced stratification (Bouttes et al., 2011) and air-sea disequilibrium (Khatiwala et al., 2019), are considered. These missing processes might also contribute to the ocean receiving less carbon from terrestrial sources than suggested by observational estimates. In our simulations, terrestrial carbon input into the ocean-atmosphere system ranges from  $177 \pm 1$  Pg C in LGMatmctl to  $198 \pm 1$  Pg C in LGMatmpi simulations. These simulated values are slightly lower than the lower bound of the observational estimate of  $511 \pm 289$  Pg C, which is based on  $\delta^{13}\text{C}$  measurements from benthic foraminifera (Peterson et al., 2014).



**Figure 8.** Globally-averaged  $\text{NO}_3^-$  vs.  $\text{O}_2$  in the 480 simulations. Color represents atmospheric  $\text{pCO}_2$  and symbols indicate the different biogeochemical configurations. The black open circle and asterisk indicate mean  $\text{NO}_3^-$  and  $\text{O}_2$  in Plctl\_Plallbgc and LGMatmctl\_LGMallbgc simulations, respectively. The cluster of points with high  $\text{pCO}_2$  and low  $\text{NO}_3^-$  is from \*\_LGMFesed simulations.

Some earlier experiments using Earth system models were able to generate a larger decline in the  $\text{pCO}_2$  (−64 to −84 ppm Ödalen et al., 2020; Vollmer et al., 2022; Matsumoto et al., 2020a). However, these model experiments only consider the increase in atmospheric Fe deposition but not the decline in the supply from the sediment, which was included in our \*\_LGMallbgc conditions. When considering only the effects of Fe deposition, the average  $\text{pCO}_2$  in the LGMatmctl\_LGMFedep simulations is 63 ppm lower than in Plctl\_Plallbgc, i.e., 20 ppm more than for LGMatmctl\_LGMallbgc. Further, different parameter combinations also affect the changes in  $\text{pCO}_2$ , e.g., the maximum  $\text{pCO}_2$  drawdown in LGMatmctl\_LGMFedep is 75 ppm (Fig. S3x), which is close to the observations. It is worthwhile to mention that increased Fe deposition alone reduces atmospheric  $\text{pCO}_2$  by 22.7 ppm from Plctl\_Plallbgc to Plctl\_LGMFedep, which is close to the changes in other model experiments considering only LGM dust deposition under pre-industrial climate conditions (14 – 22.8 ppm; Nickelsen and Oeschles, 2015; Ödalen et al., 2020; Matsumoto et al., 2020a).

The effects of variable stoichiometry on atmospheric  $\text{pCO}_2$  changes during the LGM have been examined in several previous studies. These include an empirical relationship between  $\text{pC:P}$  and ambient  $\text{PO}_4^{3-}$ , and between  $\text{pC:N}$  and ambient  $\text{NO}_3^-$  (Galbraith and Martiny, 2015), as well as a power law formulation for  $\text{pC:P}$  and  $\text{pC:N}$  that considers the influences of ambient  $\text{PO}_4^{3-}$  and  $\text{NO}_3^-$ , temperature, and light intensity (Matsumoto et al., 2020a). For models that adopt the empirical relationship between  $\text{pC:P}$  and ambient  $\text{PO}_4^{3-}$ , the magnitude of the  $\text{pCO}_2$  decline increases by 13 – 16 ppm (Ödalen et al., 2020; Fillman

et al., 2023). When the relationship between pC:N and ambient  $\text{NO}_3^-$  is also considered, the increase reaches 11 ppm (Matsumoto et al., 2020a). The power-law formulation of Matsumoto et al. (2020a) leads to an additional drawdown of 20 ppm compared to a fixed stoichiometry scheme using the Redfield ratio C:N:P = 106:16:1.

To facilitate comparison with studies neglecting the changes in sedimentary Fe flux, benthic N-loss, and  $\text{PO}_4^{3-}$  inventory, we calculate the changes in pC:N (26%) and pC:P (28%) from P1ctl\_P1allbgc to LGMatmctl\_LGMFedep, which isolates the effect of enhanced atmospheric Fe deposition in our simulations. Multiplying the relative changes by the total pCO<sub>2</sub> decline yields an estimated additional drawdown of approximately 16–17 ppm attributable to variable stoichiometry.

While this simple calculation neglects spatial variations, it provides a first-order estimate of the impact of stoichiometric flexibility on atmospheric pCO<sub>2</sub> drawdown, showing that the effects of changes in elemental ratios on pCO<sub>2</sub> in our simulations agree with previous studies considering the effects of Fe deposition.

In our experiments, pC:N and pC:P are lower under full LGM conditions (LGMatmctl\_LGMallbgc) by 8% and 10%, respectively, than in P1ctl\_P1allbgc. In LGMatmctl\_LGMallbgc, the higher surface nutrient concentrations, resulting from the weaker biological utilisation due to a lower global temperature and reduced Fe supply from the sediment, largely influence the elemental ratios of POM, as shown in the \*\_LGMFesed simulations. Nevertheless, the lower pC:N and pC:P could be simply linked to extra N and P incorporated into the POM, and do not necessarily indicate that the effects of variable stoichiometry of POM on atmospheric pCO<sub>2</sub> are reversed in LGMatmctl\_LGMallbgc. We note, however, that regional variations, particularly in regions such as the Southern Ocean, may differ from the global mean response and could play a more important role locally.

#### 4.6 Effects of parameter variations

The ranges of variation in the model's parameter values, which generally reflect ranges of uncertainty, contribute significantly to the variability of the simulated carbon cycle responses during the Last Glacial Maximum (LGM) and form a central motivation for this study. This variability arises because the response of biogeochemical tracers and fluxes to changes in boundary conditions depends strongly on the choice of the combination of parameter values.

The changes in Fe supply greatly affect the mean changes in the tracers and fluxes when compared with the pre-industrial simulations, and also contribute to higher variability among the 20 simulations with different parameter settings for each model configuration (Fig. S3 and S4). It is not surprising that the variations among the 20 ensemble members are often stronger when the median values deviate more from the pre-industrial simulations, but that is not always the case. For example,  $\text{NO}_3^-$  shows a different behaviour. When compared with \*\_P1allbgc, the deviation of the median is highest in \*\_LGMFesed but the variation is much larger in \*\_LGMFedep (Fig. S3c). That is because the  $\text{NO}_3^-$  level is affected by N<sub>2</sub> fixation, WC and benthic denitrification, and the shut-down of WC denitrification in \*\_LGMFesed subdues the variations in  $\text{NO}_3^-$ .

Compared to the pre-industrial reference condition (P1ctl\_P1allbgc), atmospheric pCO<sub>2</sub> under full LGM conditions (LGMatmctl\_LGMallbgc) decreases by 36 to 58 ppm among the 20 simulations. The difference between the minimum and maximum pCO<sub>2</sub> changes amounts to 50 % of the 43.5 ppm average decrease. We apply the lm (fitting linear model) function in R package “stats” to calculate P-values for the correlations between changes in pCO<sub>2</sub> from P1ctl\_P1allbgc to LGMatmctl\_LGMallbgc ( $\Delta\text{pCO}_2$ ) and in the perturbed parameters to better understand how individual parameters contribute to the pCO<sub>2</sub> variation (P

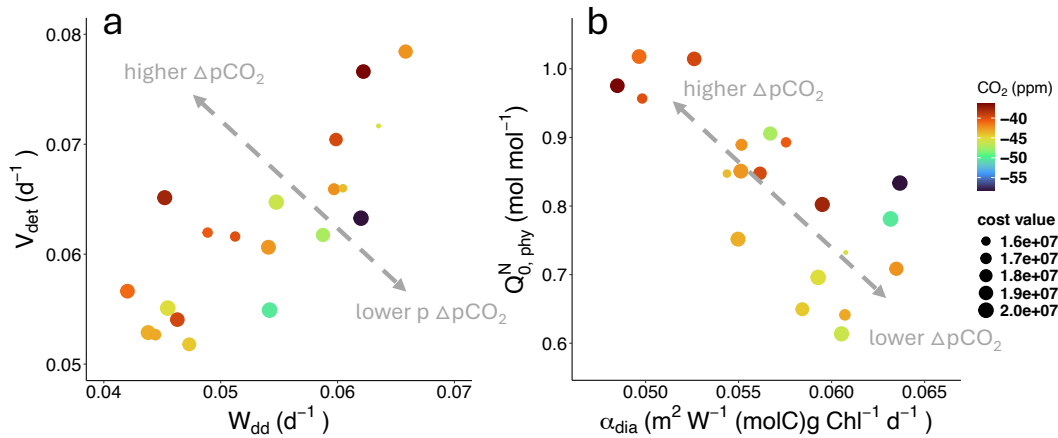
555 < 0.05, Fig. S9). Among the 19 parameters, the nitrogen subsistence quota ( $Q_{0, \text{phy}}^N$ ), temperature-dependent mortality ( $\lambda_{0, \text{phy}}$ ), and zooplankton maximum specific ingestion rate ( $g_{\text{max}}$ ) are significantly associated with the  $\Delta\text{pCO}_2$  (Fig. S9b, e, and m). Thus, not only phytoplankton- but also zooplankton (top-down)-related parameters affect the variations in  $\text{pCO}_2$  in our simulations.

Overall, the selected 20 parameter sets yield strongly different biogeochemical fluxes, inventories, and atmospheric  $\text{pCO}_2$  in our various LGM model configurations. The spread in simulated changes relative to pre-industrial conditions (PIctl, PIallbgc, and PIctl\_PIallbgc) across the 20 ensemble members arises from differences in how the selected parameter sets control model sensitivity to LGM boundary conditions in each configuration. Interestingly, the  $\Delta\text{pCO}_2$  appears unrelated to cost values based on the likelihood-based cost function we employed for the parameter selection (Fig. S9t). Indeed, the  $\Delta\text{pCO}_2$  is not necessarily correlated with cost values, which may be explained by different correlations (a) among parameters within the best 20 parameter sets, (b) between parameters and cost values, and (c) between parameters and  $\Delta\text{pCO}_2$ . This is easiest to see when the best selection of values of a pair of parameters is highly correlated. It means that collinearities exist, with effects on the cost function that are strongly interconnected. For example,  $\nu_{\text{det}}$  and  $w_{\text{dd}}$  are the strongest positively-correlated pair among the 19 parameters (Fig. S10), and because changes in  $\nu_{\text{det}}$  and  $w_{\text{dd}}$  have opposite effects on  $\Delta\text{pCO}_2$  (Fig. S9), they tend to compensate each other's effect on  $\Delta\text{pCO}_2$  in the LGMatmctl\_LGMallbgc simulations. Thus, the pairs in the best 20 parameter sets yield limited changes in the  $\Delta\text{pCO}_2$ , and a strong deviation in the  $\Delta\text{pCO}_2$  can only occur when  $\nu_{\text{det}}$  and  $w_{\text{dd}}$  change in opposite directions (Fig. 9a). The other case would be  $\alpha_{\text{dia}}$  and  $Q_{0, \text{phy}}^N$ , which is the strongest negatively-correlated pair among the 19 parameters (Fig. S10). Since changes in  $\alpha_{\text{dia}}$  and  $Q_{0, \text{phy}}^N$  also have opposite effects on the  $\Delta\text{pCO}_2$ , opposing changes in  $\alpha_{\text{dia}}$  and  $Q_{0, \text{phy}}^N$  could result in a larger deviation in the  $\Delta\text{pCO}_2$  but the effects on the cost values remains limited (Fig. 9b), owing to a weak correlation of either parameter with the cost value among the best 20 simulations. The weak relationship between the  $\Delta\text{pCO}_2$  and cost values demonstrates the complexity of LGM model simulations with respect to the parameter settings, and emphasises the importance of perturbed parameter ensemble simulations.

## 5 Conclusion and Future Directions

575 We investigate the role of several physical and biogeochemical boundary conditions in the face of parameter uncertainty in an Earth system model for simulating marine biogeochemistry and atmospheric  $\text{pCO}_2$  under pre-industrial and LGM conditions. We find that persistent changes in Fe supply are the most critical factor, while changes in major nutrients ( $\text{NO}_3^-$  or  $\text{PO}_4^{3-}$ ) alone have limited effects, at least partly due to co-limitation effects and the variable stoichiometry of POM in the model. The results from simulations with different combinations of physical and biogeochemical boundary conditions show that physical boundary conditions also affect marine biogeochemical cycles and atmospheric  $\text{pCO}_2$ , and the variable  $\text{pP:C}$  and  $\text{pN:C}$  can contribute about 16 – 17 ppm additional drawdown of  $\text{pCO}_2$  when considering LGM Fe deposition alone.

Due to the decline in sedimentary Fe input, productivity decreases and leads to higher surface  $\text{NO}_3^-$  and  $\text{PO}_4^{3-}$  in the LGM than the pre-industrial simulations. This mechanism is often ignored in LGM modelling studies. Nevertheless, we show that it



**Figure 9.** Effects of parameter pairs on the  $\Delta p\text{CO}_2$  between Pictl\_Piallbgc and LGMatmctl\_LGMallbgc. (a)  $\nu_{\text{det}}$  and  $w_{\text{dd}}$  and (b)  $\alpha_{\text{dia}}$  and  $Q_{0, \text{phy}}^N$  are the most positively and negatively correlated pairs in the 19 parameters, respectively.

could have strong impact on marine biogeochemical cycles, elemental ratios of POM, and the changes in the  $p\text{CO}_2$ . This Fe source to the ocean thus requires further understanding and examination.

The variation of  $p\text{CO}_2$  changes among the 20 LGM simulations highlights the uncertainty when applying parameter sets calibrated using pre-industrial conditions. We argue that understanding the uncertainty introduced by different boundary conditions and parameter sets is critical for accurately simulating LGM  $p\text{CO}_2$  dynamics and the interactions between physical and biogeochemical factors in Earth system models.

**Code availability.** All model codes and data used for the analyses are available at <https://doi.org/10.5281/zenodo.20816740> (Chien, 2026). All observational data in the study are publicly available, including the World Ocean Atlas 2013 ([www.ncei.noaa.gov/products/ocean-climate-laboratory](http://www.ncei.noaa.gov/products/ocean-climate-laboratory)), GLODAPv2 (<https://glodap.info/>) and POM (Tanioka et al., 2022).

**Author contributions.** CTC designed the study, performed the model simulations, conducted the analyses, and wrote the original draft. MP contributed to the development and implementation of the optimality-based plankton ecosystem model (OPEM) and provided guidance on model configuration and interpretation. CJS assisted in the design of the glacial boundary conditions. MS contributed to the parameter calibration framework and uncertainty analysis. AO contributed to the interpretation of the results, and assisted in manuscript revision. All authors discussed the results and contributed to improving the final manuscript.

**Competing interests.** The authors declare that they have no competing interest.

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