



# Marine carbon cycle response to a warmer Southern Ocean: the case of the Last Interglacial

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Abstract. Recent studies investigating future warming scenarios have shown that the ocean carbon sink will weaken over the coming century due to ocean warming and changes in oceanic circulation. However, significant uncertainties remain regarding the magnitude of the oceanic carbon cycle response to warming. Here, we investigate the Southern Ocean's (SO, south of  $40^{\circ}$ S) carbon cycle response to warmer conditions, as simulated under Last Interglacial boundary conditions (LIG, 129-115

- 5 thousand years ago, ka). We find a  $\sim$ 150% increase in carbon dioxide (CO<sub>2</sub>) outgassing over the SO at the LIG compared to pre-industrial (PI), due to a 0.5°C increase in SO sea surface temperatures. This is partly compensated by an equatorward shift of the Southern Hemisphere westerlies and weaker Antarctic Bottom Water formation, which lead to an increase in dissolved inorganic carbon (DIC) in the deep ocean at the LIG compared to PI. These deep ocean DIC changes arise from increased deep and bottom water residence times, and higher remineralization rates due to higher temperatures. While our LIG simulation
- 10 features a large reduction in SO sea-ice compared to PI, we find that changes in sea ice extent exert a minor control on the marine carbon cycle. Our results thus suggest that the projected poleward intensification of the SH westerlies, coupled with warmer conditions at the surface of the SO would weaken the SO carbon uptake over the coming century.

# 1 Introduction

- Future increases in atmospheric carbon dioxide (CO<sub>2</sub>) concentration are unequivocally projected to further warm the Southern Ocean (SO) and reduce sea ice concentrations (Bracegirdle et al., 2020). The current state of knowledge suggests the mitigating effects of carbon cycle feedbacks on global warming to be less efficient under future scenarios (Pachauri et al., 2014). This primarily stems from the changes in carbon uptake by the terrestrial biosphere and ocean under a changing climate. Both land and ocean presently act as sinks of anthropogenic carbon, absorbing about 25% of anthropogenic emissions (Friedlingstein et al., 2019), of which 40% has been attributed to the SO (Caldeira and Duffy, 2000; DeVries, 2014). Recent studies have
- shown the CO<sub>2</sub> uptake by the SO to have weakened over the 1990s (Gruber et al., 2019a, b), primarily due to a strengthening and poleward shift of the Southern Hemisphere (SH) mid-latitude westerlies (Lovenduski et al., 2007; Le Quéré et al., 2007;





Zickfeld et al., 2007), and to have strengthened after 2000 due to cooling over the Pacific and increased stratification over the Atlantic and Indian sectors of the SO (Landschuetzer et al., 2016; Gruber et al., 2019b).

- In addition, the consensus amongst studies analysing future climate simulations points towards amplified warming over high latitudes (Holland and Bitz, 2003; Smith et al., 2019; Fan et al., 2020). For example, the SO annual mean sea surface temperature (SST) anomaly is projected to exceed 0.5°C for Shared Socioeconomic Pathway scenario (SSP) 245 and 1.5°C for SSP 585 at 2100 relative to 2015 (Bracegirdle et al., 2020; Tonelli et al., 2021), with reduced sea ice during austral spring (Roach et al., 2020). At the same time, SH westerlies are projected to strengthen and shift poleward over the coming century (Goyal et al., 2021). Overall, climate change will reduce the amount of anthropogenic carbon taken up by the ocean (Plattner
- 30 et al., 2001; Bernardello et al., 2014; Wang et al., 2014; Howes et al., 2015) by reducing CO<sub>2</sub> solubility (Bernardello et al., 2014), weakening the efficiency of the biological pump (Boyd, 2015), increasing outgassing from upwelling of Circumpolar Deep Waters (CDW) due to stronger and poleward shifted SH westerlies (Lovenduski et al., 2007; Zheng et al., 2013; Gruber et al., 2019a), and reducing sea-ice extent (Shadwick et al., 2021). However, uncertainties still remain in regards to the response of the SO carbon cycle under a warmer climate.
- The Last Interglacial (LIG, 129-115 thousand years ago, ka) was the warmest interglacial of the last million years (Masson-Delmotte et al., 2013; PAGES, 2016). The warmer climate at the LIG is primarily attributed to a stronger northern hemispheric summer insolation (Laskar et al., 2004) owing to the orbital configuration of higher eccentricity and obliquity (Berger, 1978), rather than higher greenhouse gas concentrations as projected for the future. The LIG is associated with sea levels 6-9 m higher than pre-industrial (PI) (Dutton et al., 2015) and annual mean SSTs around 0.5°C higher than PI (Capron et al., 2014; Hoffman
- 40 et al., 2017). The area-weighted summer warming is estimated to be 1.1-1.9°C for the North Atlantic and 1.6-1.8°C for the SO compared to PI (Govin et al., 2012; Capron et al., 2014; Hoffman et al., 2017). Summers over land areas are reconstructed to be 4-5°C warmer at high latitudes in the Northern Hemisphere (CAPE-Last Interglacial Project Members, 2006), 3-11°C over Greenland (Dahl-Jensen et al., 2013) and 2.2°C over Antarctica (Masson-Delmotte et al., 2013). Reconstructions of sea ice are mostly focused on the Arctic, and suggest ice-free conditions south of 78°N (Van Nieuwenhove et al., 2008, 2011, 2013;
- 45 Kageyama et al., 2021). The Atlantic Meridional Overturning Circulation (AMOC) has been suggested to have weakened at the peak of the LIG (127ka) and strengthened afterwards (Galaasen et al., 2014), with strong evidence pointing towards periods of reduced Antarctic Bottom Water (AABW) formation during the early part of the LIG, due to discharges from the Antarctic ice-sheet (Hayes et al., 2014; Rohling et al., 2019).

A few studies have investigated the terrestrial carbon response to LIG conditions (Kleinen et al., 2015; Brovkin et al., 2016),

- 50 but the marine carbon cycle response at the LIG, and particularly in the SO, has received little attention. A comprehensive study of the changes in the SO carbon cycle at the LIG compared to PI can enhance our understanding of the processes involved in the SO carbon cycle, and their sensitivity to changes in boundary conditions. It can also help us, to a certain extent, to better quantify the impact of expected future physical and dynamical changes in a warming SO on the marine carbon cycle, bearing in mind that there was no additional anthropogenic carbon released during the LIG.
- 55 Here, we study the impact of a warmer climate, in particular at high latitudes, on the oceanic carbon cycle by analysing an equilibrium last interglacial simulation (lig127k, Otto-Bliesner et al. (2017)) performed with the ACCESS-ESM1.5 model



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(Ziehn et al., 2020; Yeung et al., 2021). The paper is structured as follows: In Section 2, components of the ACCESS-ESM1.5 model are described, followed by a framework for decomposing variables relevant for carbon cycle changes. The main results are presented in Section 3, discussing the changes in climate and the carbon cycle. These include differences in ocean properties, such as temperature, sea ice cover, stratification and circulation, and their effects on the carbon cycle, including air-sea fluxes, net carbon storage, and the efficiency of the biological pump. Finally in Section 4, we conclude and discuss the limitations of the current study including possible sources of uncertainty, and what inferences can be drawn for future global warming scenarios.

#### 2 Methods

# 65 2.1 Model description and experimental design

An equilibrium last interglacial simulation (lig127k) is performed with the Australian Community Climate and Earth System Simulator Earth System Model, ACCESS-ESM1.5 (Ziehn et al., 2020), which includes interactive land and ocean carboncycles, and is Australia's submission to the Paleoclimate Modeling Intercomparison Project 4 (PMIP4) - Coupled Model Intercomparison Project (CMIP6) (Yeung et al., 2021). ACCESS-ESM1.5 differs from the previous version, ACCESS-ESM1

- 70 (Law et al., 2017), mostly in the land and ocean components. ACCESS-ESM1.5 is built upon the ACCESS1.4 physical model, and includes the UK Met Office Unified Model (UM) version 7.3 (Martin et al., 2010; Bellouin et al., 2011) as atmospheric component, which is directly coupled to the updated land Community Atmosphere Biosphere Land Exchange model (CABLE) version 2.4 (Kowalczyk et al., 2013), both with a horizontal resolution of 1.875° × 1.25°. UM has 38 vertical levels. The ocean model is the NOAA/GFDL Modular Ocean Model (MOM) version 5 (Griffies, 2014), which is coupled to the Los Alamos
- 75 National Laboratory sea ice model (LANL CICE) version 4.1 (Hunke et al., 2010). MOM5's resolution is 1° × 1° with 50 vertical levels. The coupler is the Ocean Atmosphere Sea Ice Soil Model Coupling Toolkit (OASIS-MCT) (Craig et al., 2017). The current version of the land model CABLE also includes biogeochemistry (BGC) implemented using the CASA-CNP module (Wang et al., 2010). The ocean carbon cycle is simulated using the Whole Ocean Model of Biogeochemistry And Trophic-dynamics (WOMBAT) model (Oke et al., 2013).
- 80 WOMBAT is a nutrient–phytoplankton-zooplankton–detritus (NPZD) model, with one class of phytoplankton and one class of zooplankton. It includes DIC, alkalinity (ALK), phosphate ( $PO_4$ ), oxygen ( $O_2$ ) and iron. The biogeochemical tracers are coupled using the stoichiometric C (carbon):N (nitrogen):P (phosphorous): $O_2$  ratio of 106:16:1:172. The air-sea gas exchange is based on the square of the wind speed (Wanninkhof, 1992), and the seawater partial pressure of  $CO_2$  ( $pCO_2$ ) is calculated following the third phase of Ocean Carbon-Cycle Model Intercomparison Project (OCMIP) protocol using temperature, salinity,
- B5 DIC, ALK and PO<sub>4</sub>. WOMBAT simulates production in and export from the photic zone, and remineralization and dissolution at depth for both organic and inorganic (CaCO<sub>3</sub>) particulate matter, with parameters adjusted for inorganic export to be  $\sim$ 8% of organic export. The remineralization of organic matter is calculated based on depth and temperature dependent parameters, while the dissolution of CaCO<sub>3</sub> uses a constant dissolution rate. ACCESS-ESM1.5 does not include a sediment component,



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and all particulates reaching the bottom ocean layer are instantly remineralized following the relevant remineralization rates. Other specific details of the model can be found in Oke et al. (2013), Law et al. (2017) and Ziehn et al. (2020).

WOMBAT includes two DIC tracers, one being forced by the prescribed atmospheric  $CO_2$  concentration (PI: 284.3 ppm, LIG: 275 ppm), and the other forced by a constant atmospheric  $CO_2$  concentration of 280 ppm. Unless mentioned otherwise, all of the analyses presented here are based on the tracers forced with  $CO_2$  concentrations of 280 ppm. This allows quantification of the effects of the LIG climate on the carbon cycle independently of the difference in background  $CO_2$  concentrations.

- A pre-industrial 1850 simulation (piControl) is run in accordance with the CMIP6 protocol (Eyring et al., 2016), with a constant atmospheric CO<sub>2</sub> forcing of 284.3 ppm, but using CMIP5 solar irradiance (1365.65  $Wm^{-2}$ , as explained in Ziehn et al. (2020)). This simulation is run for 1000 years. Initialized from this PI control run, a LIG simulation (lig127k) is performed using orbital parameters following the PMIP4 protocol (Otto-Bliesner et al., 2017; Yeung et al., 2021), but with the solar constant adjusted to the CMIP5-PMIP3 value to be comparable to the piControl simulation (1365.65  $Wm^{-2}$ ). CO<sub>2</sub> concentration
- 100 for the LIG is set at 275 ppm. Vegetation is kept constant at 1850 PI conditions. The lig127k experiment is run for 650 years. The run is at equilibrium over the SO, although there is a small drift in globally averaged DIC of  $\sim 1 \mu mol/kg/100$  years below 3 km. The analysis presented here is based on the average of the last 100 years of the lig127k simulation compared to the last 100 years of the piControl simulation. Further details on these experiments can be found in Otto-Bliesner et al. (2017), Ziehn et al. (2020), and Yeung et al. (2021).

## 105 2.2 pCO<sub>2</sub> decomposition

Changes in surface  $pCO_2$ , which ultimately control the direction and magnitude of air-sea fluxes, can further be decomposed into contributions from SST, sea surface salinity (SSS), DIC, and ALK (Sarmiento and Gruber, 2006) using the following equations:

$$\Delta p CO_2^{SST} = e^{(\gamma_{SST} \times \Delta SST)} \times p CO_2^{PI} - p CO_2^{PI} \tag{1}$$

110 where  $\Delta pCO_2^{SST}$  represents the contribution of SST change to surface pCO<sub>2</sub> change,  $\gamma_{SST}$ =0.0423°C<sup>-1</sup> is the Revelle factor (Sarmiento and Gruber, 2006),  $\Delta SST$  is the difference in SST between LIG and PI, and  $pCO_2^{PI}$  is the surface seawater pCO<sub>2</sub> from the piControl simulation; and

$$\Delta p CO_2^X = \gamma_X \times \frac{\Delta X}{\overline{X}} \times p CO_2^{PI} \tag{2}$$

where  $\Delta pCO_2^X$  represents the contribution of change in variable 'X' (SSS, DIC, and ALK) to surface pCO<sub>2</sub> change,  $\Delta X$  is 115 the change in 'X' between LIG and PI, and  $\overline{X}$  is the mean value of 'X' at PI,  $\gamma_X$  the Revelle factors, with  $\gamma_{SSS} = 1$ ,

$$\gamma_{DIC} = \begin{cases} 15, & \text{if } lat \ge 60\\ 13, & lat < 60^{\circ} \end{cases}$$

and

$$\gamma_{ALK} = \begin{cases} -12, & \text{if } lat \ge 60^\circ \\ -11, & lat < 60^\circ \end{cases}$$





These values are based on previous estimates of the meridional profiles of DIC and ALK buffer factors (Sarmiento and Gruber,
2006; Smith and Marotzke, 2008; Egleston et al., 2010; Jiang et al., 2019), although uncertainties still remain regarding these estimates. This decomposition helps shed light on the independent effects of different physical variables to the net surface pCO<sub>2</sub> change.

# 2.3 Carbon partitioning

To better quantify changes in the carbon cycle between the LIG and PI experiments, we split the total DIC into its remineralized  $(C_{org})$ , dissolved carbonate  $(C_{CaCO_3})$ , and preformed  $(C_{pre})$  components using the equations listed below.

 $C_{org}$  can be estimated from regenerated phosphate (PO<sub>4</sub><sup>Reg</sup>) using the C:P stoichiometric ratio ( $r_{C:P}$ =106) (Ito and Follows, 2005):

$$\Delta C_{org} = r_{C:P} \times \Delta PO_4^{Reg} \tag{3}$$

 $PO_4^{Reg}$  can be approximated using Apparent Oxygen Utilization (AOU) and the P:O<sub>2</sub> stoichiometric ratio ( $r_{P:O_2}=1/172$ ) (Ito 130 and Follows, 2005; Duteil et al., 2012):

$$\Delta PO_4^{Reg} = r_{P:O_2} \times \Delta AOU \tag{4}$$

AOU estimates the oxygen consumed during respiration and can be calculated as the difference of dissolved oxygen concentration ( $O_2$ ) from the saturated concentration of oxygen ( $O_2^{sat}$ ) (Weiss, 1970):

$$AOU = O_2^{sat}(T, S) - O_2 \tag{5}$$

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This can then be used to infer the efficiency of the biological pump ( $BP_{Eff}$ ) as per Ito and Follows (2005):

$$\overline{BP_{Eff}} = \overline{PO_4^{Reg}} / \overline{PO_4} \tag{6}$$

where  $\overline{X}$  is the mean value of X.

The contribution to DIC from the carbonate pump is estimated by:

$$\Delta C_{CaCO_3} = 0.5 \times [\Delta ALK + r_{N:P} \times \Delta PO_4^{Reg}] \tag{7}$$

140 where the term  $r_{N:P} \times \Delta PO_4^{Reg}$ , accounts for the reduction in ALK from production of nitrate (NO<sub>3</sub><sup>-</sup>) (Sarmiento and Gruber, 2006), which is estimated using PO<sub>4</sub><sup>Reg</sup> and the N:P stoichiometric ratio ( $r_{N:P}$ =16). Finally, the preformed carbon concentration ( $C_{pre}$ ) is obtained as:

$$\Delta C_{pre} = \Delta DIC - \Delta C_{org} - \Delta C_{CaCO_3} \tag{8}$$

## 3 Results

145 Changes in the climate system are first presented (Section 3.1), followed by their effects on air-sea gas exchange in the SO (Section 3.2). To understand these, we quantify the different contributors to changes in surface partial pressure of  $CO_2$  next





(Section 3.3), and finish by analysing deep ocean changes and the global oceanic carbon inventory (Section 3.4). Unless otherwise mentioned, the SO is defined as the ocean area south of  $40^{\circ}$ S.

#### 3.1 Ocean dynamics and sea ice cover

- As a result of the insolation anomalies and associated feedbacks, the global mean annual SST anomaly at the LIG compared to PI as simulated by the ACCESS-ESM1.5 equals to  $0.17^{\circ}$ C, with a pronounced warming at high latitudes (not shown, Yeung et al. (2021)). A mean  $0.53^{\circ}$ C warming is simulated over the SO south of  $40^{\circ}$ S (Fig. 1a). Warmer conditions are simulated everywhere south of  $50^{\circ}$ S apart from a  $\sim 1^{\circ}$ C cooling centered at  $58^{\circ}$ S in the South Atlantic, and up to a  $1.5^{\circ}$ C cooling in the subantarctic eastern Pacific. The strongest warming is simulated over the southeast Atlantic and Indian Ocean sectors (up to
- 155 3°C). Regional SSTs up to 4°C higher are simulated around 60°S for both austral spring (Fig. A1b) and summer (not shown). The higher SSTs over the SO compared to PI are accompanied by a marked reduction in sea-ice extent over both austral summer and winter (Fig. 1a), peaking at 41% reduction in austral winter (Fig. A1a).

A 1.5° equatorward shift of the SH westerlies is simulated at the LIG (Fig. 1e), with a 10% weakening of the winds south of 50°S. This leads to weaker upwelling south of 55°S and stronger upwelling north of 55°S (Fig. 1f). Seasonally, the largest changes in upwelling are found for the winter and spring seasons (Fig. A1c and d). The equatorward shift of the westerlies reduces the northward Ekman transport south of 55°S leading to warming, while the higher Ekman transport north of 55°S induces a cooling around 50°S (Fig. 1a and b). The Antarctic Circumpolar Current is weaker, and the transport through Drake passage is reduced from PI by ~55Sv (not shown).

#### 3.2 Response of the air-sea gas exchange

- 165 These physical changes in the SO impact the carbon cycle and the air-sea  $CO_2$  exchange (Fig. 2). It is worth reiterating here that we analyse changes in the carbon cycle using tracers for a constant atmospheric  $CO_2$  concentration of 280 ppm for both simulations (Section 2.1), which enables us to solely analyse the impact of climatic and oceanic circulation changes on the carbon cycle. The  $CO_2$  flux over the SO at PI shows a carbon uptake near the Antarctic coast, an outgassing band between 65-45°S, and another uptake zone further north (Fig. 2a and blue line in d). This  $CO_2$  outgassing is due to the upwelling
- 170 of DIC-rich deep waters (Fig. 1f). At the LIG, the upwelling region widens over the Atlantic and Indian ocean sectors, and narrows over the eastern Pacific sector (Fig. 2). There is a strong increase in outgassing in the Atlantic and western Indian, as well as at  $\sim 67^{\circ}$ S in the eastern Pacific sectors (Fig. 2c and red line in d). An increase in CO<sub>2</sub> uptake is simulated in the Amundsen, Bellinghausen, Weddell, Lazarev, Riiser-Larsen and Ross Seas, and the sub-antarctic east Pacific (Fig. 2c). The zonal mean CO<sub>2</sub> flux in Fig. 2d shows a small increase in uptake south of 62°S, possibly due to a reduction in convection.
- 175 Overall, there is a net increase in the mean SO outgassing by  $\sim 150\%$  at the LIG compared to PI (Fig. 2e). This increase in outgassing mostly occurs during the austral winter and spring seasons (Fig. A1e and f). To better understand these changes, we decompose the oceanic pCO<sub>2</sub> changes into their different components.





#### 3.3 Changes in surface carbon dioxide partial pressure

- The air-sea gas exchange is primarily controlled by the seawater partial pressure of CO<sub>2</sub> (pCO<sub>2</sub>). Using the equations presented
  in Section 2.2, Fig. 3 shows a decomposition of the changes in pCO<sub>2</sub> into their SST, SSS, DIC and ALK contributions. In line with the net increase in SO outgassing at the LIG (Fig. 2), the net pCO<sub>2</sub> over the SO is 1.2 µatm higher at the LIG compared to PI (black circle) (Fig. 3a). In agreement with the changes in air-sea CO<sub>2</sub> flux, surface pCO<sub>2</sub> is higher in a zonal band centered at ~55°S, while it is lower in coastal regions (contours in Fig. 3b and black line in g). The contributions from individual components add up to reflect the simulated differences reasonably well both in terms of magnitude (1.4 µatm, gray square in Fig. 3a) and spatial distribution (Fig. 3b), affirming the validity of the decomposition method. The overall slightly positive
- pCO<sub>2</sub> anomaly at the LIG compared to PI results from the competing effects of lower CO<sub>2</sub> solubility (red triangle, +5.65  $\mu$ atm) and changes in surface DIC and ALK (blue diamond, -4.25  $\mu$ atm).

The largest contributor to solubility is SST (brown triangle), while the largest contributor of the combined DIC and ALK effect is DIC (cyan diamond in Fig. 3a). Higher SSTs in the SO at the LIG lead to a 5.8  $\mu$ atm pCO<sub>2</sub> increase (brown triangle

- in Fig. 3a). This SST induced increase is present over most of the SO, and is highest over the south Atlantic, Indian and west Pacific regions (Fig. 3c). Changes in SSS do not contribute significantly to the pCO<sub>2</sub> anomalies (magenta triangle in Fig. 3a and d). Changes in DIC cause the largest single contribution to the overall pCO<sub>2</sub> change with the net decrease in surface DIC leading to a -10.9  $\mu$ atm pCO<sub>2</sub> change (cyan diamond in Fig. 3a). An exception are the higher DIC concentrations in the Atlantic Ocean around 55°S and in the eastern Pacific sector between 60°S-70°S (Fig. A1g,h, and A2d), leading to higher pCO<sub>2</sub> (Fig.
- 195 3e). This higher DIC is due to increased upwelling in these regions (Fig. 1b and A1c,d), resulting from the equatorward shift of the SH westerlies. The contributions based on changes in ALK and DIC have very similar patterns, albeit of opposite signs, with a small difference over the west Pacific around 50°S. The combined effect of ALK and DIC changes leads to a total net decrease in pCO<sub>2</sub> by 4.25  $\mu$ atm due to the higher impact of changes in DIC (-10.9  $\mu$ atm, cyan diamond) compared to ALK (+6.65  $\mu$ atm, green diamond). Overall, south of 45°S, changes in SST lead to an overall increase in pCO<sub>2</sub> with the highest

200 contributions over the Indian and West Pacific sectors of the SO, while changes in DIC lead to an overall decrease of pCO<sub>2</sub>, with strongest reductions in the southern Atlantic, Indian and West Pacific sectors of the SO but an increase in the Amundsen Sea sector (Fig. 3).

The zonal mean patterns of these contributions are presented in Fig. 3g. Coastal regions around Antarctica, south of  $\sim$ 75°S, show that the lower  $\Delta$ pCO<sub>2</sub>, both simulated (black line) and calculated (gray line), primarily arise from the combined DIC

- and ALK component (blue line) that is mainly attributed to the lower DIC (cyan dashed line) compensated by changes in ALK (green dashed line). Contributions from changes in SST are negligible (brown dashed line), however the net solubility contribution (red line) is slightly negative, due to a freshening (magenta dashed line). Between 70°S and 60°S,  $\Delta pCO_2$  anomalies are close to zero as the reduced CO<sub>2</sub> solubility due to higher SSTs (red line) is compensated by the combined DIC and ALK changes (blue line). Between 60°S and 45°S, the simulated surface ocean  $\Delta pCO_2$  is positive (~2 µatm, black line) as the
- 210 effect of higher SSTs dominates over the combined ALK and DIC components. North of  $\sim 45^{\circ}$ S, the positive  $\Delta pCO_2$  signal arises from the DIC+ALK component while solubility mitigates the anomaly (Fig. 3g).





Figure 3g shows that while solubility is mostly controlled by changes in SST, contributions from SSS changes are important south of 72°S. Similarly, DIC has the dominant control over the DIC+ALK component. The contributions from both the solubility and (DIC+ALK) components reach their maxima (solubility being positive and DIC+ALK being negative) around 62°S, which corresponds to the maximum divergence in wind-driven surface currents (Fig. 1f). To summarise, the higher overall pCO<sub>2</sub> (and hence lower air-sea CO<sub>2</sub> flux) at the LIG compared to PI can be attributed to higher SSTs south of 45°S, and the combined DIC-ALK component between 45°S and 35°S due to the equatorward shift of the upwelling regions (Fig. 3g). The SST patterns were already discussed in Section 3.1. DIC patterns (Fig. A2d) can result from changes in circulation, and the biological pump. These are investigated in the next section.

#### 220 3.4 Deep ocean changes and carbon inventory

Figure 1c and d show that North Atlantic Deep Water (NADW) formation is  $\sim$ 5 Sv higher in our LIG simulation compared to the PI simulation, leading to colder waters at depths of 1-2 km between 0-60°N, accompanied by higher SSTs and SSSs in the Labrador Sea (not shown). We also simulate a  $\sim$ 4 Sv reduction in AABW formation (Fig. 1c and d). This reduction in AABW formation results in a warming of SO deep waters by up to 2°C compared to PI. Due to northward shifted winds (Fig. 1e,f),

225 the Antarctic Intermediate Water (AAIW) formation regions also shift northward (Downes et al., 2017). The changes in SST and upwelling (Fig. 1), alongside the reduced sea ice extent, lead to an increase in net primary production (NPP) and export production over the SO by ~17 % and ~11 % respectively (Fig. A2a and b).

These circulation changes impact the DIC distribution in the ocean (Fig. 4b). For instance, the northward shift of AAIW (Fig. 1d) leads to lower oxygen (Fig. 4a), higher DIC (Fig. 4b), and higher  $PO_4$  concentrations (Fig. 4c) at intermediate depths

230 of the SO north of 55°S. Similarly, the increased accumulation of nutrients and DIC, as well as oxygen depletion in deep and abyssal waters (Fig. 4a, b and c), can be attributed to a weaker AABW formation rate (Fig. 1d), which results in a higher efficiency of the biological pump, as detailed below.

The ACCESS-ESM1.5 simulates a  $\geq$  50  $\mu$ mol/kg decrease in oxygen everywhere below 3 km at the LIG compared to PI. A decrease in oxygen can also be seen in the SO across all depths, including a northward reaching tongue attributable to AAIW

at  $\sim$ 1 km depth extending to the Equator (Fig. 4a). DIC anomalies follow the patterns of dissolved oxygen, albeit with the opposite sign. The simulated DIC concentrations are  $\sim$ 50  $\mu$ mol/kg higher across all depths in the SO and globally below 3.5 km depth (Fig. 4b).

DIC anomalies are decomposed into contributions from remineralized organic carbon, dissolved calcium carbonate, and preformed carbon components (Fig. 4d, f and h, Section 2.3). 60% of the DIC increase in the SO and abyssal ocean can

- 240 be attributed to an increase in remineralised carbon (Fig. 4d), resulting from a 10% more efficient biological pump (Eq. 6, Section 2.3). This increase in remineralised carbon can be attributed to increased residence times, due to weaker bottom and intermediate water formation rates (Fig. 1 c and d). Weaker AABW and AAIW indeed lead to positive apparent oxygen utilization (AOU) and regenerated PO4 anomalies (Fig. 4e). In addition, a 2°C warming of bottom waters leads to a  $\sim$ 6% increase in remineralization rates, thus further contributing to the higher remineralised carbon. Around 25% of the abyssal
- 245 increase in DIC is attributed to the carbonate pump (Fig. 4f). This reflects changes in NPP, export production (Fig. A2) and





water mass residence times, given that carbonate production is a constant percentage of total NPP in this model setup, and carbonate dissolution is constant and independent of water chemistry or temperature. Preformed DIC represents only  $\sim 5\%$  of the DIC changes (Fig. 4h). This increased sequestration of DIC in the deep ocean reduces the surface DIC concentration (Fig. A2 and A1g, h), thus contributing to a lowering of surface pCO<sub>2</sub> (Fig. 3e and g).

For the Northern Hemisphere, stronger NADW (Fig. 1c and d) results in decreased DIC concentrations in intermediate and deep waters of the North Atlantic by up to 75 μmol/kg (Fig. 4b). This negative DIC anomaly can be explained by a decrease in both remineralised and preformed DIC. Stronger NADW subducts more DIC depleted surface waters into the deep ocean (Duteil et al., 2012), and reduces residence times, thus leading to a decrease in remineralized carbon. This is also associated with a ~40 µmol/kg increase in dissolved oxygen content (Fig. 4a), and up to a 0.75 µmol/kg decrease in PO<sub>4</sub> concentration (Fig. 4c). The reduction in preformed carbon (Fig. 4h) is most likely due to a new NADW formation site in the Labrador Sea along with a slight northward shift of the deep water formation regions in the Norwegian and Greenland Seas.

## 4 Discussion and Conclusion

Understanding how the marine carbon cycle responds to a warmer climate is of vital importance to better constrain future climate projections. To this end, we analyse the SO marine carbon cycle response to warmer conditions as simulated in an equilibrium simulation of the LIG. The lig127k simulation performed with the ACCESS-ESM1.5 presented here displays an annual mean warming of 0.53°C at the surface of the SO compared to PI. This simulated southern high latitude warming (Fig. 1a) is in agreement with the multi-model mean of the PMIP4 lig127k simulations, although it is at the higher end of the spectrum (Otto-Bliesner et al., 2021). Seasonally, we simulate regional SSTs up to 4°C higher around 60°S for both the austral spring and summer, in line with both, proxy records (Capron et al., 2017; Hoffman et al., 2017) and the PMIP3 125ka
experiment performed with the NORESM-1ME model (Kessler et al., 2018), while the PMIP4 lig127k multi-model mean for experiment displays then 0.5%C merening even the SO (Otta Plicener et al., 2021). The simulated SO are inplaced by the summer displays then 0.5%C merening even the SO (Otta Plicener et al., 2021). The simulated SO are included by the summer displays then 0.5%C merening even the SO (Otta Plicener et al., 2021). The simulated SO are included by the summer displays then 0.5%C merening even the SO (Otta Plicener et al., 2021).

- for austral summer displays less than 0.5°C warming over the SO (Otto-Bliesner et al., 2021). The simulated SO sea ice concentration at the LIG for austral winter (Fig. 1 and A1) is also in good agreement with those inferred in previous studies (Holloway et al., 2016, 2017), even though the changes in Antarctic sea ice area of the ACCESS-ESM1.5 are the largest amongst all the CMIP6-PMIP4 models (Otto-Bliesner et al., 2021).
- A 1.5° equatorward shift of the westerlies, resulting in a 10% weakening of the westerlies south of 50°S, is simulated at the LIG (Fig. 1e), leading to significant changes in SO upwelling (Fig. 1b and f). There is no clear consensus for position and strength of the SH westerlies during the LIG (Fogwill et al., 2014), although the higher obliquity might have led to a weakening of the westerlies compared to PI (Timmermann et al., 2014). These changes in winds lead to a northward shift of the AAIW formation regions. Our LIG simulation is also characterised by a weakening of AABW formation rates, which might
- 275 be due to changes in surface density. Significant weakening of AABW during the LIG due to Antarctic meltwater discharge has previously been inferred from paleo-proxy records (Hayes et al., 2014; Rohling et al., 2019). A shift in westerlies might also contribute to a weakening of the AABW (Menviel et al., 2008; Huiskamp et al., 2016; Glasscock et al., 2020).





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DIC in the deep ocean, through an increase in the efficiency of the biological pump (Fig. 4). This reduces the surface DIC concentrations, leading to a net reduction in outgassing over the SO, as has been previously hypothesised (Toggweiler et al., 2006). However, while the combined DIC and ALK contributions would lead to a lower pCO2 at the surface of the SO, this change is overcompensated by the warmer conditions (Fig. 3). Reduced solubility due to higher SSTs leads to an increase in outgassing over most of the SO, while the reduced sea-ice cover does not seem to significantly impact the CO2 fluxes. This results in a net outgassing of CO<sub>2</sub> over the SO, with a  $\sim$ 150% increase at the LIG compared to PI (Fig. 2), and the largest increase over the austral winter and spring seasons (Fig. A1). The simulated NPP and export production are ~17% 285 and  $\sim 11\%$  higher, respectively, over the SO at the LIG compared to PI (Fig. A2), providing a negative feedback to the higher

The reduced upwelling south of 55°S and weaker AABW transport simulated here lead to an increased sequestration of

- outgassing. While our analysis is based on a constant atmospheric CO<sub>2</sub> concentration of 280 ppm, the lower CO<sub>2</sub> at LIG (275 ppm) compared to PI (284.3 ppm) would suggest the LIG SO to be an even greater CO<sub>2</sub> source to the atmosphere, implying a stronger sink somewhere else in the ocean or on land (Brovkin et al., 2016).
- 290 Numerical studies have unequivocally shown the impact of changes in the magnitude of the westerlies on the carbon cycle, with stronger westerlies leading to increased upwelling and SO outgassing, and vice-versa (e.g., Menviel et al., 2008; d'Orgeville et al., 2010; Lauderdale et al., 2013; Huiskamp et al., 2016; Lauderdale et al., 2017; Menviel et al., 2018; Gottschalk et al., 2019), however, the impact of changes in the latitudinal position of the westerlies on the carbon cycle is less certain (e.g., d'Orgeville et al., 2010; Völker and Köhler, 2013; Lauderdale et al., 2013; Huiskamp et al., 2016). Given the simulated in-

crease in DIC content in the deeper SO and the reduced DIC at the surface of the SO, our simulations suggest that the simulated 295 equatorward shift of the westerlies reduces the SO CO<sub>2</sub> outgassing.

SH westerlies are projected to strengthen and shift poleward over the coming century (Goyal et al., 2021), which should increase CO<sub>2</sub> outgassing in the SO (Lovenduski et al., 2007; Zickfeld et al., 2007; Menviel et al., 2018; Gruber et al., 2019a). Projections also unequivocally suggest the SO to be warming for all emission scenarios (Bracegirdle et al., 2020), which would further contribute to a weaker SO sink of  $CO_2$ . Finally, while we find a more efficient biological pump at the LIG, Boyd 300 (2015) suggested the biological pump to be less efficient under a future warming scenario. Such combined effects of warming, a poleward shift and strengthening of the westerlies, and a less efficient biological pump would suggest a reduction in carbon uptake by the SO in the future.

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Code and data availability. Outputs of the physical variables from the lig127k simulation are archived on the CMIP6 ESGF website at https://doi.org/10.22033/ ESGF/CMIP6.13703 (Yeung et al., 2019). Outputs of the carbon cycle model have already been CMORised, and will be available on the ESGF website by the end of August 2021.





*Author contributions.* DC performed all of the analysis and writing of the results. LM and KJM provided support for interpretation and writing of the results. NKHY performed the lig127k simulation. MC and TZ contributed to the model setup and troubleshooting. All authors contributed towards the final manuscript.

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**Figure 1.** Annual mean (a) SST anomalies (°C) between LIG and PI overlaid with 15% sea ice concentration (black for PI, magenta for LIG, solid lines for DJF, dashed lines for JJA). Value in the box shows the SO (40-90°S) mean  $\Delta$ SST. (b) Anomalies of the Ekman pumping velocities between LIG and PI ( $ms^{-1}$ ) overlaid with sea ice concentration (same as (a)). (c) Global mean meridional streamfunction (Sv) for PI and (d) LIG. Positive values indicate clockwise water mass transport and negative values counterclockwise transport. Zonal mean (e) winds ( $ms^{-1}$ ), and (f) Ekman pumping velocities ( $ms^{-1}$ ) over SO for PI (blue) and LIG (red).







**Figure 2.** Annual mean air-sea CO<sub>2</sub> flux (molCm<sup>-2</sup>y<sup>-1</sup>) for (a) PI, (b) LIG, and (c) LIG-PI. Red colors indicate outgassing of CO<sub>2</sub> from the ocean, and blue uptake by the ocean. Thick black lines show the zero line contour. (d) Zonal mean CO<sub>2</sub> flux (molCm<sup>-2</sup>y<sup>-1</sup>) over the SO for PI (blue) and LIG (red). (e) Mean CO<sub>2</sub> flux over the SO (molCm<sup>-2</sup>y<sup>-1</sup>) for PI (blue) and LIG (red).







**Figure 3.** Attribution of changes in annual mean surface partial pressure of  $CO_2$  (p $CO_2$ ) over SO ( $\mu$ atm). (a) Summary of decomposition of p $CO_2$  over SO. Simulated difference in p $CO_2$  (black circle) and sum of contributions from all components (gray square, Section 2.2). p $CO_2$  change from solubility (SST+SSS, red triangle), and sum of DIC and ALK components (blue diamond). Individual contributions from SST (brown triangle), SSS (magenta triangle), DIC (cyan diamond), and ALK (green diamond). (b) Map of the sum of p $CO_2$  contributions from all components (corresponding to gray square in (a)) in shading, overlaid with the p $CO_2$  change simulated by the model as contours (corresponding to black circle in (a)). Maps of individual p $CO_2$  contributions from (c) SST (corresponding to the brown triangle in (a)), (d) SSS (corresponding to the magenta triangle in (a)), (e) DIC (corresponding to the cyan diamond in (a)), and (f) ALK (corresponding to the green diamond in (a)). (g) Zonal mean contributions to p $CO_2$  over the SO ( $\mu$ atm) for simulated  $\Delta$ p $CO_2$  (black), sum of contributions from all components (gray), solubility (red), sum of DIC and ALK components (blue), SST (dashed brown), SSS (dashed magenta), DIC (dashed cyan), and ALK changes (dashed green). Note the non-linear scale on the top and bottom thirds of (g).







**Figure 4.** Global marine carbon budget and decomposition. Global zonal and annual mean anomalies of (a) dissolved oxygen concentration  $(\mu mol/kg)$ , (c) phosphate concentration  $(\mu mol/kg)$ , (e) regenerated phosphate concentration  $(\mu mol/kg)$ , (g) preformed phosphate concentration  $(\mu mol/kg)$ , (b) total DIC  $(\mu mol/kg)$ , (d) remineralized carbon  $(\mu mol/kg)$ , (f) dissolved carbonate  $(\mu mol/kg)$ , and (h) preformed DIC  $(\mu mol/kg)$ . Note that the phosphate components (c, e, and g), and the DIC components (b, d, f and h) each share common colorbars.







**Figure A1.** Seasonal anomalies for austral winter, JJA (1st column, a, c, e, g), and austral spring, SON (2nd column, b, d, f, h) of (a,b) SST (°C) overlaid with 50% sea ice concentration (dashed for PI and solid for LIG), (c,d) Ekman pumping velocities ( $ms^{-1}$ ) overlaid with 50% sea ice concentration (dashed for LIG), (e,f) sea-air CO<sub>2</sub> flux (mol C m<sup>-2</sup>y<sup>-1</sup>, positive indicating outgassing from and negative uptake by the ocean), and (g,h) surface DIC concentrations ( $\mu$ mol/kg).







**Figure A2.** Changes in annual mean productivity and related variables. (a) Anomalies of depth integrated annual mean NPP (as cholorphyll in seawater, kgm<sup>-3</sup>) overlaid with 15% sea ice concentration (black for PI, red for LIG, solid lines for DJF, dashed lines for JJA). (b) Anomalies of annual mean detrital concentration at 200m depth (kgm<sup>-3</sup>) overlaid with sea ice (same as (a)). (c) Anomalies of annual mean surface phosphate concentrations ( $\mu$ mol/kg) overlaid with sea ice (same as (a)). (d) Anomalies of annual mean surface DIC concentrations ( $\mu$ mol/kg) overlaid with sea ice (same as (a)).