# Sensitivity of asymmetric Oxygen Minimum Zones to mixing intensity and stoichiometry– in the tropical Pacific using a basin-scale model (OGCM-DMEC V1.4)

Kai Wang<sup>1</sup>, Xiujun Wang<sup>1,2\*</sup>, Raghu Murtugudde<sup>2</sup>, Dongxiao Zhang<sup>3</sup>, Rong-Hua Zhang<sup>4</sup>

<sup>5</sup> <sup>1</sup>College of Global Change and Earth System Science, Beijing Normal University, Beijing 100875, China <sup>2</sup>Earth System Science Interdisciplinary Center, University of Maryland, College Park, Maryland 20740, USA <sup>3</sup>JISAO, University of Washington and NOAA, Pacific Marine Environmental Laboratory, Seattle, Washington 98115, USA <sup>4</sup>Institute of Oceanology, Chinese Academy of Sciences, Qingdao, Shandong 266071, China

Correspondence to: Xiujun Wang (xwang@bnu.edu.cn)

- 10 Abstract. The tropical Pacific Ocean holds the world's two largest Oxygen Minimum Zones (OMZs), showing a prominent hemispheric asymmetry, with a much stronger and broader OMZ north of the equator. However, many models have difficulties in reproducing the observed asymmetric OMZs in the tropical Pacific. Here, we apply a fully coupled basin-scale model to evaluate the impacts of stoichiometry and the intensity of vertical mixing on the dynamics of OMZs in the tropical Pacific. We first utilize observational data of dissolved oxygen (DO) to calibrate and validate the basin-scale model. Our model experiments demonstrate that enhanced vertical mixing combined with a reduced O:C utilization ratio can significantly
- improve our model capability of reproducing the asymmetric OMZs. Our study shows that DO is more sensitive to biological processes over 200-700400 m but to physical processes overbelow 400-1000 m. Enhanced vertical mixing causes a largemodest increase in physical supply (1-2 mmol m<sup>-3</sup> yr<sup>-1</sup>) and a small increase (<0.5 mmol m<sup>-3</sup> yr<sup>-1</sup>) in biological consumption over 200-1000 m whereas applying a reduced O:C utilization ratio leads to a large decrease in(2-8 mmol m<sup>-3</sup> yr<sup>-1</sup>)
- 20 <sup>1</sup>) in both biological consumption, and a small decrease in physical supply in the OMZs. Our analyses suggest that biological consumption (greater rate to the south than to the north) cannot explain the asymmetric featuredistribution of mid-depth DO in the tropical Pacific OMZs, but physical processes supply (stronger supplyvertical mixing to the south) play a major role in regulating the asymmetry of the tropical Pacific's OMZs. This study also highlights the important roles of physical and biological interactions/ and feedbacks in contributing to the asymmetry of OMZs in the tropical Pacific.

### 25 **1 Introduction**

Photosynthesis and respiration are important processes in all ecosystems on the Earthearth, with carbon and oxygen being the two main elements. The carbon cycle has garnered much attentions attention, with significant progress advances in both the observations (Feely et al., 1999; Takahashi et al., 2009) and modelling (DeVries et al., 2019; Le Quéré et al., 2010; McKinley et al., 2016) of biological processes (e.g., uptake of CO<sub>2</sub> and respiration); and physical/chemical processes (e.g., carbon fluxes

30 between the atmosphere, land and ocean). However, the oxygen cycle has received much less attention despite its large role in the earth system (Breitburg et al., 2018; Oschlies et al., 2018).

Dissolved oxygen (DO) is a sensitive indicator of physical and biogeochemical processes in the ocean thus a key parameter for understanding the ocean's role in the climate system (Stramma et al., 2010). In addition to photosynthesis and respiration,

- 35 the distribution of DO in the world's oceans is also regulated by air-sea gas exchange, ocean circulation and ventilation (Bettencourt et al., 2015; Bopp et al., 2002; Levin, 2018). Unlike most dissolved nutrients that display an increase in concentration with depth, DO concentration is generally low at mid-depth of the oceanoceans. The most remarkable feature in the oceanic oxygen dynamics is the so-called Oxygen Minimum Zone (OMZ) that is often present below 200 m in the open oceans (Karstensen et al., 2008; Stramma et al., 2008). Previous studies have used the isoline of 20 mmol m<sup>-3</sup> as the boundary
- 40 of the OMZ for the estimation of OMZ volume (Bettencourt et al., 2015; Bianchi et al., 2012; Fuenzalida et al., 2009), and also as an up limit to determine the suboxic water (Wright et al., 2012).

The world's two largest OMZs are observed in the Eastern Tropical North Pacific (ETNP) and South Pacific (ETSP), showing a peculiar asymmetric structure across the equator, i.e., a much larger volume of suboxic water (<20 mmol m<sup>-3</sup>) to the north than to the south (Bettencourt et al., 2015; Paulmier and Ruiz-Pino, 2009). It is known that OMZs are caused by the biological consumption associated with remineralization of organic matter (OM), and weak physical supply of DO due to sluggish subsurface ocean circulation and ventilation (Brandt et al., 2015; Czeschel et al., 2011; Kalvelage et al., 2015). Although there

45

have been a number of observation-based analyses addressing the dynamics of OMZs in the tropical Pacific during the past decade (Czeschel et al., 2012; Garçon et al., 2019; Schmidtko et al., 2017; Stramma et al., 2010), our understanding is limited
on the underlying mechanisms that regulate DO dynamics at mid-depth (Oschlies et al., 2018; Stramma et al., 2012).

Large-scale physical-biogeochemical models have become a useful tool to investigate the potential sensitivity of OMZs to climate change (Duteil and Oschlies, 2011; Ward et al., 2018; Williams et al., 2014). However, many models have beenare unable to reproduce the observed patterns of asymmetric OMZs in the tropical Pacific (Cabre et al., 2015; Shigemitsu et al., 2017), which may be due to "unresolved ocean transport processes, unaccounted for variations in respiratory oxygen demand, or missing biogeochemical feedbacks" (Oschlies et al., 2018). A common problem is that the two asymmetric OMZs merge into one in most models due to overestimated OMZ volume in the tropical Pacific, which may be related to the regulation of physical supply and/or biological respiration demand (Cabre et al., 2015; Shigemitsu et al., 2017). RecentWhile some studies have also indicatedsuggest that a realistic representation of circulation and ventilation processes with a high-resolution ocean
60 model is critical to predictsimulation of the asymmetric OMZs in the tropical Pacific (Berthet et al., 2019; Busecke et al., 2019). Hence, it's necessary to carry out model-data integrative, other modelling studies have demonstrated that physical

processes (e.g., vertical mixing) play a major role in regulating the distribution of mid-depth DO (Brandt et al., 2015; Llanillo et al., 2018). On the other hand, there have been advances in understanding of biogeochemical regulation on DO consumption, i.e., oxygen-restricted remineralization of organic matters (Kalvelage et al., 2015). Hence, it's necessary to carry out integrative
 model-data studies to improve model capacity of simulating the dynamics of the tropical OMZs, and to better understand the
 relative roles of physical and biological processes. Without such process understandings, it is unclear a priori whether simply
 increasing resolution will render better simulations and predictions in regulating the asymmetry of the tropical Pacific OMZs.

A basin-scale ocean general circulation model coupled with a dynamic marine ecosystem-carbon model (OGCM-DMEC) washas been developed for the tropical Pacific, which showedhas demonstrated capability of reproducing observed spatial and temporal variations of physical, nutrient and carbon fields in the upper ocean (Wang et al., 2008; Wang et al., 2015; Wang et al., 2009b), and nitrate, iron, particulate organic carbon (POC/detritus) and export production below 200 m (Yu et al., 2021). In this study, we conduct model sensitivity experiments and evaluation onevaluations to examine the responses of mid-depth DO and the sources/sinks to parameterizations of two relevantkey processes (i.e., oxygen-restricted remineralization and vertical mixing). We first carry out model calibration and validation using observational data of basin-scale DO concentration and oxygen consumption in the water column of the southernsouth tropical Pacific to improve the simulation of OMZs in the tropical Pacific. Then, we analyse the impacts of new parameterizations on biological consumption and physical supply and their relative contributions to the dynamics of mid-depth DO. The objective of this study is to advance our model capacity to simulate the oceanic oxygen cycle, and to identify the mechanisms driving the asymmetric OMZs in the tropical Pacific.

#### 80 2. Model description

#### 2.1 Ocean physical model

The basin-scale OGCM, OGCM a reduced-gravity, primitive-equation, sigma-coordinate model, and it is coupled to an advective atmospheric model (Murtugudde et al., 1996). There are 20 layers with variable thicknesses and a total depth of ~1200 m in the OGCM. The mixed layer (the upper-most layer) depth is determined by the Chen mixing scheme (Chen et al., 85 1994), which varies from 10 m to 50 m on the equator. The remaining layers in the euphotic zone are approximately 10 m in thickness. The vertical resolution is approximately 30-50 m in the core OMZ (at ~300-500 m). The model domain is between 30°S and 30°N for the Pacific, and zonal resolution is 1°. Meridional resolution varies between 0.3° and 0.6° over 15°S-15°N (1/3° over 10°S-10°N), and increases to 2° in the southern and northern "sponge layers" (the 25°-30° bands) where temperature, salinity, nutrients and DO are gradually relaxed back towards the observed climatological seasonal means. The model closes 90 the western boundary and no representation of the Indonesian throughflow is included. The boundary conditions of temperature, DO and from the World Atlas, 2013 (WOA2013: salinity, nitrate are Ocean http://www.nodc.noaa.gov/OC5/woa13/pubwoa13.html), and boundary condition for dissolved iron is based on limited field data, and given is represented by a linear regression against temperature (see details in Christian et al., 2001). Such model configuration may have a disadvantage for longer simulations and analyses, but has the advantage in reproducing the spatial

95 patterns of most physical and biogeochemical fields.

The model is forced by atmospheric conditions the surface momentum, heat and freshwater fluxes: climatological monthly means of solar radiation and cloudiness, and interannual 6-day means of precipitation and surface wind stress. Precipitation is from ftp://ftp.cdc.noaa.gov/Datasets/gpcp. Wind stresses are from the National Centers for Environmental Prediction (NCEP)

100 reanalysis (Kalnay et al., 1996). Air temperature and humidity above the ocean surface are computed by the atmospheric mixed layer model. Initial conditions were obtained from the outputs of an interannual hindcast simulation over 1948-2000, which itself is initialized from a 30-year spin up with climatological forcing, followed by two 40-year interannual simulations. The initial conditions for the spin up are specified from the WOA2013, iron concentration for the spin up was initialized from limited field data collected in the tropical Pacific (Johnson et al., 1997). We carry out an interannual simulation for the period of 1978-2010, and analyse the mean states from model simulations over the period of 1991-2010.

#### 105

#### 2.2 Ocean biogeochemical model

The DMEC model consists of eleven components: small (S) and large (L) sizes of phytoplankton ( $P_S$  and  $P_L$ ), zooplankton ( $Z_S$ and  $Z_L$ ) and detritus ( $D_S$  and  $D_L$ ), dissolved organic nitrogen (DON), ammonium, nitrate, dissolved iron, and DO (Figure 1). Phytoplankton growth is co-limited by nitrogen and iron, which is critical in the tropical Pacific. The model simulates the iron

110 cycle using variable Fe:N ratios, and incorporates atmospheric iron input. All biological components use nitrogen as their unit, in which sources/sinks are determined by biological and chemical processes in addition to the physical processes (circulation and vertical mixing) that are computed by the OGCM.

In this model, net community production (NCP) is computed as:

 $NCP = 6.625(\mu_{s}P_{s}\mu_{s}P_{s} + \mu_{L}P_{L} - \frac{r_{s}}{r_{s}}r_{s}Z_{s} - r_{L}Z_{L} - c_{DON}DON - c_{DS}\frac{D_{s}}{D_{s}}D_{s} - c_{DL}D_{L})$ 115 (1)

where 6.625 is the C:N ratio,  $\mu$  the rate constant of phytoplankton growth, r the rate constant of zooplankton respiration, c the rate constants of detritus decomposition and DON remineralization. The equations for biogeochemical processes and model parameters are given in Appendix A and B. There were changes in some parameters comparing with those in Wang et al. (2008), which were based on our model calibration and validation for chlorophyll (Wang et al., 2009a), nitrogen cycle (Wang 120 et al., 2009b) and carbon cycle (Wang et al., 2015).

- Recently, we have made further improvements in the parameterizations of detritus decomposition and DON remineralization (eq. B21-B23), which result from the first round of model calibration on DO distribution using WOA2013. In brief,  $c_{DON}$ decreases with depth over 100-1000 m, following an exponential function in this study. The differences in the related
- 125 parameters are given in Appendix C.

### 2.3 Computation of oxygen sources and sinks

The time evolution of DO is regulated by physical, biological and chemical processes:

145

 $O_{mix}$  is the vertical mixing term that is calculated by three subroutines. Briefly, the first one computes convection to remove instabilities in the water column, and the second one determines the mixed layer depth-<u>based on the available surface turbulent</u> kinetic energy. The third one computes partial vertical mixing (K<sub>z</sub>) between two adjacent layers to relieve gradient Richardson (R<sub>i</sub>) number instability, which is calculated as follows:

150 
$$K_z = \left(1 - \left(\frac{Ri}{0.7}\right)^{\lambda}\right) (Ri < 0.7)$$
  
 $K_z = 0 \ (Ri \ge 0.7)$ 
(3(8)  
(4(9))

where the mixing parameter  $\lambda$  is set to 1. Clearly, partial vertical mixing is the dominant process influencing physical supply of DO in the intermediate waters.

### 155 Monthly rate of total physical supply (*O<sub>sup</sub>*) below the euphotic zone is computed as:

$$O_{sup} = \frac{\Delta O}{\Delta t} + O_{cons}$$
(10)  
Total physical supply consists of meridional, zonal and vertical advection, and vertical mixing. The advection terms are computed from the corresponding velocity and DO gradient, and the vertical mixing term is calculated as the residue.

The biological source/sink term Obie is computed as follows:

160 O<sub>bto</sub> = R<sub>OC</sub>NCP (5) where R<sub>OC</sub>-is the O:C utilization ratio (set to 1.3 in reference simulation, according to the Redfield ratio). Below the euphotic zone, DO concentration is determined by physical supply and biological consumption that results from detritus decomposition and DON remineralization, in which DON remineralization is dominant because DON pool is several times greater than detritus (Wang et al., 2008).
165 The flux of O<sub>2</sub> from the atmosphere to the surface ocean is computed as: O<sub>gas</sub> = (O<sub>Sat</sub> - O)K<sub>0</sub> (6) where O<sub>sat</sub> is the O<sub>2</sub> saturation, a function of temperature and salinity (Weiss, 1970), and K<sub>0</sub> the gas transfer velocity that is a function of wind speed (u<sub>s</sub>) and SST according to Wanninkhof (1992):

70 
$$K_0 = 0.31 u_s^2 \sqrt{\frac{S_c}{S_{c20}}}$$
 (7)  
where  $S_e$  and  $S_{e2\ell}$  are the Schmidt number at SST and  $20^\circ$  C, respectively:  
 $S_c = 1953 - 128T + 3.99T^2 - 0.05T^3$  (8)

#### 3. Model experiments

#### 3.1 Evaluation of DO distribution from the reference run

- 175 We first evaluate simulated DO for the tropical Pacific Ocean using the outputs from OGCM-DMEC V1.2 (hereafter reference run), which use the same set of parameters as Yu et al. (2021)4 (hereafter reference run), which use the same set of parameters as Yu et al. (2021)4 (hereafter reference run), which use the same set of parameters as Yu et al. (2021)4 (hereafter reference run), which use the same set of parameters as Yu et al. (2021)4 (hereafter reference run), which use the same set of parameters as Yu et al. (2021)4. We focus on model-data comparisons over 200-400 m, 400-700 m and 700-1000 m, that broadly represent the upper OMZ, lower OMZ and beneath the OMZ, respectively. The WOA2013 data shows a much larger area of suboxic waters (<20 mmol m<sup>-3</sup>) in the ETNP than in the ETSP over 200-400 m and 400-700 m (Figure 2a and 2c), but no suboxic water
- 180 over 700-1000 m (Figure 2e). Although the reference run produces two OMZs off the equator over 200-400 m (Figure 2b), the sizes of suboxic water are much larger in the reference run than those in the WOA2013 data. The reference run significantly over-estimates the size of suboxic water and underestimates DO concentration over 400-700 m (Figure 2d). The difference between WOA2013 and the reference run is small over 700-1000 m, except in the eastern tropical Pacific (Figure 2f).

#### 3.2 Sensitivity experiments

185 Given that the mid-depth DO concentration is influenced by physical supply and biological consumption, the underestimated DO at mid-depth would be a result of overestimation of consumption associated with DON remineralization and/or underestimation of supply. The reference run applied a zero value for background diffusion (see eq. 9). However, a previous modelling study demonstrated that vertical background diffusion was an important process for DO supply at mid-depth (Duteil and Oschlies, 2011). Accordingly, we conduct a sensitivity experiment to test a set of values for background diffusion (Kb as

- 190 0.1, 0.25 and 0.5 cm<sup>2</sup> s<sup>-1</sup>). The addition of background diffusion is only applied to the two key variables (DO and DON) in this analysis to eliminate any potential interactions and feedbacks between various physical and biogeochemical processes (note: our model experiments showed no significant effects on modelled DO dynamics with the background diffusion applied to the nutrients).
- 195 There have been <u>several</u> advances in understanding of oxygen consumption. For example, recent studies have <u>showedshown</u> that <u>the</u> O:C utilization ratio varies <u>largelygreatly</u> across different basins, e.g., from 0.6 to 2.1 in the Pacific (Moreno et al., 2020; Tanioka and Matsumoto, 2020), and rates of DOM remineralization or oxygen consumption are influenced by oxygen level, i.e., a reduction under low DO conditions (Beman et al., 2020; Bertagnolli and Stewart, 2018; Sun et al., 2021). Based on the field data at mid-depth (~350 m) in the Peruvian OMZ (Kalvelage et al., 2015), we derive a kinetics function between
- 200 oxygen consumption rate and DO concentration, which yields <u>two values for</u> the half saturation constant (Km-<u>being) as</u> 6.9 and 18.7 mmol m<sup>-3</sup> (Figure 3). By <u>addingapplying</u> this <u>functional form to-function</u>, O:C <u>utilization ratio in</u> equation 5, one would get a varying <u>7 becomes variable</u> and <u>is</u> also reduced O:C <u>utilization ratio</u>, (i.e.,  $R_{oc} = 1.3 \frac{DO}{DO+Km}$ ), with lower ratios in in-low-DO waters.
- 205 The reference run applied a zero value for background diffusion. However, a previous modelling study demonstrated that vertical background diffusion was an important process for DO supply at mid depth (Duteil and Oschlies, 2011). Accordingly, we conduct Therefore, the sensitivity experiments consist of a few more simulations (Table S1) to investigate how applying with a reduced O:C utilization ratio (setting Km as 6.9 and 18.7 mmol m<sup>-3</sup>) and adding background diffusion (setting Kb as 0.25 and -0.5 cm<sup>2</sup>-s<sup>-1</sup>) affect the simulated DO distribution and asymmetry of OMZs in the tropical Pacific. To eliminate complex interactions and feedbacks, the addition of background diffusion is only applied to the key variables (DO and DON)
- in this study.added background diffusion (Table S1).

Figure 4 illustrates that based on WOA2013 database, there is a larger volume of suboxic water located north of ~5°N and a smaller volume of suboxic water over 12°S-4°S, which are separated by relatively higher DO (>20 mmol m<sup>-3</sup>) water along the equator. There is an improvement in simulated DO with <u>a</u> reduced O:C utilization ratio (Figure 4b and 4c) and enhanced vertical mixing (Figure 4d and 4h4g). Clearly, the combination of <u>a</u> reduced O:C utilization ratio and enhanced vertical mixing leads to a further improvement in simulated mid-depth DO (Figure 4e, 4f, 4i4h and 4j4i). In particular, the combination of a stronger background diffusion with a smaller O:C utilization ratio (i.e., the Km18.7Kb0.5 run) results in the best simulation that reproduces the observed spatial distribution of mid-depth DO, especially the asymmetric featurehemispheric asymmetry (i.e., a larger volume of suboxic water to the north but a smaller size of suboxic water to the south).

To further evaluatedemonstrate the performanceimpact of experimentsparameter choices, a few statistical measures are applied over 200-400 m, 400-700 m and 700-1000 m in the ETNP (165°W-90°W, 5°-20°N) and ETSP (110°W-80°W, 10°S-3°S). As shown in Table 1, compared with the reference run, bias and root mean square error (RMSE) are reduced in all newsensitivity simulations, with the smallest values from Km18.7Kb0.5 run except over 700-1000 m in the ETNP. For example, both bias and RMSE in the Km18.7Kb0.5 run are smallest over 200-700 m in the ETNP (<7.8 and 10.2 mmol m<sup>-3</sup>). Many current models showhave much largelarger RMSE (~20-80 mmol m<sup>-3</sup>) with respect to observed DO from mixed layer to 1000 m (Bao and Li, 2016; Cabre et al., 2015). Figure 5 also illustrates that the Km18.7Kb0.5 run produces the best outputs, withaccording to the largest correlation coefficients (0.77 0.94) and also combined assessments (the smallest-distance to 1 in the observation) of the 230 correlation coefficient and normalized standard deviation (0.54 1.81 in NSD). The distance is shortest over 400-700 m and 700-100 m in both the ETNP and 0.33 1.63ETSP in the Km18.7Kb0.5 simulation. Clearly, the correlation coefficient was largest (0.38-0.99) in ETSP-all sections; and the NSD is closest to 1 in the core OMZ of ETNP.

We also compare the sizes of suboxic water and hypoxic waterwaters between the model simulations and WOA2013 (Table 2). Based on WOA2013, sizes of suboxic water and hypoxic waterwaters are 5.97x10<sup>15</sup> m<sup>3</sup> and 19.98x10<sup>15</sup> m<sup>3</sup> in the north, and 1.43×10<sup>15</sup> m<sup>3</sup> and 7.12x10<sup>15</sup> m<sup>3</sup> in the south, respectively. While a reduced O:C utilization ratio and enhanced vertical mixing can lead to an improvement in simulated OMZ volume, a significant improvement is obtained with the combination of a reduced O:C utilization ratio and enhanced vertical mixing. Overall, the Km18.7Kb0.5 simulation has the best performance for reproducing the OMZ volumes, showing similar volumes for the suboxic water (5.55x10<sup>15</sup> m<sup>3</sup> to the north and 1.12x10<sup>15</sup>
240 m<sup>3</sup> to the south) and the hypoxic water (20.91x10<sup>15</sup> m<sup>3</sup> and 7.39x10<sup>15</sup> m<sup>3</sup>).

We then further validate the modelled DO from the best run (Km18.7Kb0.5), using the time series of the observed DO data (<u>https://cchdo.ucsd.edu/</u>). Figure 6 illustrates that the model can generally reproduce the vertical-zonal distributions of DO along 10°N and 17°S, spanning from-1989 to 2009, particularly in the eastern tropical Pacific. For example, cruise data from the P04 line during April-May, 1989 show a large area of low DO water spanning from ~200 m to ~800 m (Figure 6a), and our model also predicts low DO water over ~200-700 m (Figure 6b).

4 Model results and discussions

245

In this section, we further compare the improved model simulations (Km18.7, Kb0.5 and Km18.7Kb0.5) with the reference run to diagnose the influences of improved parameterizations on the distribution of mid-depth DO, and biological consumption and physical supply. We then analyse the interactions of physical and biogeochemical processes, and the impacts on the source and sink for the mid-depth DO. In the endFinally, we explore the underlying mechanisms regulating the asymmetry of OMZs in the tropical Pacific.

#### 4.1 Changes of mid-depth DO due to a reduced O:C utilization ratio and enhanced vertical mixing

We first compare the changes in DO concentrations between the three model simulations over 200-400 m, 400-700 m, and

- 255 700-1000 m (Figure 7). Clearly, applying a reduced O:C utilization ratio causes an increase of DO in all three layers, with the greatest increase (>6 mmol m<sup>-3</sup>) in the 200-400 m layer (Figure 7a), followed by a modest increase (~3-6 mmol m<sup>-3</sup>) over 400-700 m (Figure 7d). Although DO increase is generally smaller in the 700-1000 m layer (Figure 7h7g) than in the 400-700 m layer (Figure 7d), the increase is greater in the north OMZ over 700-1000 m than over 400-700 m. Enhanced vertical mixing results in a small increase of DO (~2-5 mmol m<sup>-3</sup>) in the 10°S-10°N band over 200-400 m (Figure 7b), but a large increase
- 260 $(\sim 5-15 \text{ mmol m}^{-3})$  in majority of the basin over 400-700 m and 700-1000 m (Figure 7e and  $\frac{747}{10}$ ).

Overall, the mid-depth DO shows an increase with the combination of a reduced O:C utilization ratio and enhanced vertical mixing (Figure 7c, 7f & 7i7i). A great increase of DO (>15 mmol m<sup>-3</sup>) occurs in majority over most of the basin over 400-700 m, mainly in the central tropical Pacific over 200-400 m, but in a few small areas over 700-1000 m. The spatial pattern and 265 magnitude of DO increase resulting from the combination of a reduced O:C utilization ratio and enhanced vertical mixing, have a large similarity to those with a reduced O:C utilization ratio for the 200-400 m layer (Figure 7a), but to those under enhanced vertical mixing below 400 m (Figure 7e & 7i7h). For example, the relative increase of DO is similarly larger in the northermorth OMZ over 200-400 m under a reduced O:C utilization ratio with and without the addition of background diffusion, and over 700-1000 m under enhanced vertical mixing (i.e., with additional background diffusion) with and without 270 the change in the O:C utilization ratio. Our analyses suggest that the dominant process regulating the DO dynamics is biological

consumption over 200-700400 m, but physical supply over 400-1000 m.

#### 4.2 Effects of a reduced O:C utilization ratio and enhanced vertical mixing on consumption and supply

To better understand the effects of changes in the biological and/or physical parameters on the DO dynamics, we then evaluate the responses of biological consumption and physical supply. As illustrated in Figure 8, changes in biological consumption are almost identical undercaused by a reduced O:C utilization ratio are almost identical with or without background diffusion. In 275 particular, biological consumption shows a large decrease ( $\sim 2-8 \text{ mmol m}^{-3} \text{ yr}^{-1}$ ) over 200-400 m (Figure 8b), and a relatively small decrease (~0.2-1.0 mmol m<sup>-3</sup> yr<sup>-1</sup>) over 400-700 m, with the largest decrease in the northernnorth OMZ (Figure 8e); (there). There is a very small change in biological consumption over 700-1000 m, i.e., a decrease of <0.1 mmol m<sup>-3</sup> yr<sup>-1</sup> over majoritymuch of the basin but an increase of <0.1 mmol m<sup>-3</sup> yr<sup>-1</sup> in some parts of subtropical region (Figure Si8h). On the other hand, enhanced vertical mixing leads to a small increase (<0.2 mmol m<sup>-3</sup> yr<sup>-1</sup>) in biological consumption in all three layers,

280

with a relatively larger increase in the northermorth OMZ (Figure 8c, 8f and 8<del>i). 8i</del>).

Figure 9 shows the effects of a reduced O:C utilization ratio and enhanced vertical mixing on physical supply. With the combination of a reduced O:C utilization ratio and enhanced vertical mixing, physical supply shows a small increase (by ~0.21.0 mmol m<sup>-3</sup> yr<sup>-1</sup>) in the whole basin over 700-1000 m (Figure 9h9g) and only outside the OMZs over 400-700 m (Figure 9d), but a relatively larger decrease (by ~0.2-6 mmol m<sup>-3</sup> yr<sup>-1</sup>) in the OMZs over 200-700 m (by ~0.2 6 mmol m<sup>-3</sup> yr<sup>-1</sup>) (Figure 9a and 9d). Clearly, enhanced vertical mixing leads to an increase of physical supply over majority of the basin, with <u>a</u> greater increase over 400-1000 m (~0.2-1.0 mmol m<sup>-3</sup> yr<sup>-1</sup>) than over 200-400 m (~0-0.4 mmol m<sup>-3</sup> yr<sup>-1</sup>) (Figure 9c, 9f and 9j9i). However, applying a reduced O:C utilization ratio causes a large decrease of physical supply above 700 m, with <u>a</u> greater decrease over 400-700 m in the OMZs (~0.2-6 mmol m<sup>-3</sup> yr<sup>-1</sup>), and very small changes (<0.2 mmol m<sup>-3</sup> yr<sup>-1</sup>) over 700-1000 m (Figure 9b, 9e and 9i9h). Overall, the rate of physical supply is largely determined largely by vertical mixing over 700-1000 m, by both vertical mixing and biological consumption over 400-700 m, but by consumption over 200-400 m, implying complex physical-biological interactions and feedbacks in the tropical Pacific OMZs.

#### 4.3 Interactive effects of physical and biological processes on the source and sink of mid-depth DO

295 There is evidence that enhanced mixing can have large influences not only on physical processes (e.g., the strength of water mixing) but also on biological processes (e.g., transport of organic materials), which have direct or indirect effects on the evolution of mid depth DO (Andrews et al., 2017; Duteil and Oschlies, 2011; Stramma et al., 2012). Our Our further analyses show an increase in physical supply under enhanced vertical mixing in most parts of the 200-1000 m layer-in the eastern tropical Pacific (over 120°W 90°W) (Figure 10). Interestingly, the greater increase (>1 mmol m<sup>-3</sup> yr<sup>-1</sup>) is, with larger values 300 below the OMZs over 15°S 10°N using 1.3 as the O:C utilization ratioparticularly to the south (Figure 10a), but occurs over a much larger area (i.e., over 15°S 20°N) and within the OMZs using a reduced (and also varying) O:C utilization ratio (Figure 10d). Enhanced vertical mixing also results in a generally small increase in biological consumption, with greater increases in OMZs using a reduced O:C utilization ratio (Figure 10e) than using a constant Redfield ratio of 1.3 (Figure 10b). The small increase in consumption outside of OMZs is largely is attributable to increased DON concentration (data not shown Figure 10c) 305 that results from the enhanced vertical mixing whereas the increase of consumption inside the OMZs would be a result of the interactions and feedbacks of various physical, biological and chemical processes. Clearly, there is an overall increase in net flux, with the largest increases occurring mainly outside the OMZs below  $\sim 400$  m (Figure 10c and 10f10d).

To further investigate the interactive effects of a reduced O:C utilization ratio and enhanced mixing, we then compare the responses of biological consumption and physical supply to changes in the O:C utilization ratio with and without background diffusion (Figure 11). While a reduced O:C utilization ratio can result in a decrease in consumption above 600 m, the decrease is slightly less in the OMZs with background diffusion (Figure 11d) than without background diffusion (Figure 11a). Similarly, physical supply also shows a decrease in the OMZs under a reduced O:C utilization ratio (Figure 11b), with a lesser decrease under the addition of background diffusion (Figure 11e). The greatest difference is found in the core OMZs for both biological consumption (Figure 11h) and physical supply (Figure 11i), but larger differences are found in supply. A previous modeling

study also demonstrates that physical contribution to the changes of DO is much greater than biogeochemical contribution

(Montes et al., 2014). However, a reduced O:C utilization ratio results in a clear increase in net flux in the whole water column over 200 1000 m, with a great increase above the core OMZs within the 10°S 10°N band (Figure 11c and 11f).

As expected, applying a reduced O:C utilization ratio results in a decrease in consumption in the suboxic waters (Figure 10f), with a greater decrease in the north OMZ than in the south OMZ. Interestingly, physical supply shows an overall decrease in the water column under a reduced O:C utilization ratio, with a greater decrease in the upper OMZs (Figure 10e). A decreased rate of consumption leads to a large increase in DON concentration, with a greater increase in the north OMZ than in the south OMZ (Figure 10g). Net flux shows a small increase in the whole basin under a reduced O:C utilization ratio, with a greater

325 increase over ~200-400 m (Figure 10h).

The combination of enhanced vertical mixing and a reduced O:C utilization ratio results in an increase of supply below the OMZs but a decrease of supply inside of OMZs (Figure 10i). There is an overall decrease of biological consumption in the water column, with a greater decrease in the upper OMZs (Figure 10j), which is similar to the changes under a reduced O:C

utilization ratio (Figure 10f). DON concertation shows a greater increase in the north OMZ than in the south OMZ under the combination of a reduced O:C utilization ratio and enhanced vertical mixing (Figure 10k), which is similar to the changes in DON under a reduced O:C utilization ratio (Figure 10g). Applying a reduced O:C utilization ratio combined with enhanced vertical mixing leads to an increase in net flux over 200-1000 m, with a larger increase outside of OMZs (Figure 10l), which is much greater than that under a reduced O:C utilization ratio (Figure 10h), and also greater than that by enhanced vertical mixing particularly in the lower part of OMZs (Figure 10d).

There is evidence that physical and biogeochemical processes have multiple interactions with impacts on various physical, chemical and biological fields which in turn have implications for the DO dynamics (Breitburg et al., 2018; Duteil and Oschlies, 2011; Oschlies et al., 2018). For example, observational and modelling studies show that changes in vertical mixing intensity
 can affect the distribution of DOM thus oxygen consumption at mid-depth (Duteil and Oschlies, 2011; Talley et al., 2016). On the other hand, applying a smaller O:C utilization ratio leads to lower consumption rates (Moreno et al., 2020), thus to a relatively higher DO concentration in the OMZs. Therefore, changes in the consumption caused by enhanced vertical mixing and/or a reduced O:C utilization ratio would alter the gradients of DO concentration in the water column thus change the intensity of vertical mixing inside and around the OMZs.

Our analyses also show that the changes in both the supply and consumption under improved parameterizations of both vertical mixing and remineralization of DOM (i.e., Km18.7Kb0.5) are quite different from the sums of changes caused by single parameter change, particularly in the OMZs (Figure 10m and 10n), indicating strong physical and biological interactions with positive feedbacks in the low-DO waters. Clearly, the interactions have a relatively larger effect on physical supply because

<sup>345</sup> 

#### 350 of its sensitivity to changes in both physical and biological parameters. As a result, the interactive effects result in an overall increase in net flux in the OMZs (Figure 10p).

Physical supply could can be divided into horizontal advection, vertical advection, and vertical mixing. Our Figure S1 shows that the dominant process for DO supply is vertical mixing particularly above ~600 m in the OMZs. Other modelling studies

- 355 have also demonstrated that vertical mixing is the dominant process supplying oxygen from the thermocline to OMZs (Duteil et al., 2020; Llanillo et al., 2018). By comparing with previous model performs wellresults (Duteil et al., 2020; Shigemitsu et al., 2017), our model simulation with the combination of enhanced vertical mixing and a reduced O:C utilization ratio shows a good performance in simulating the meridional and zonal advections, and vertical mixing processes of for DO transport (see Figure S2), which allows us to evaluate the responses of different supply physical components to the enhanced vertical mixing 360 and a reduced O:C utilization ratio.

As shown in Figure  $\frac{1211}{12}$ , there is no clear pattern in the responses of advective supply transport, with very small values (<  $\sim 1$ mmol  $m^{-3}$  yr<sup>-1</sup>) over the entire basin (Figure 12h and 12i). However, the DO supply by vertical mixing shows a strong response to different model parameterizations, with similar patterns to as those of in total supply and (see Figure 10). Enhanced 365 background diffusion leads to a large decrease in increase of vertical mixing (>1 mmol m<sup>-3</sup> yr<sup>-1</sup>) over most of the suboxic waters (Figure 12c and 12f). Whilebasin, with greater increase mainly below the OMZs (Figure 11c). On the other hand, applying a reduced O:C utilization ratio causes a decrease in the DO supply ( $-1.6 \text{ mmol m}^3 \text{ yr}^-$ ) by vertical mixing, the decrease is larger in the OMZs without the addition of background diffusion. On the other hand, there is an increase in the supply by vertical mixing below the OMZs under a reduced O:C utilization ratio, in particular with the addition of background 370 diffusion (Figure 12f). The largest difference (~1 2 mmol m<sup>-3</sup> yr<sup>4</sup>) is found within the hypoxic waters (Figure 12i), which reflects the strong feedback between physical and biological processes in the OMZs, small increase in vertical mixing of DO  $(<0.5 \text{ mmol m}^{-3} \text{ yr}^{-1})$  outside of suboxic waters but a large decrease  $(\sim 2-8 \text{ mmol m}^{-3} \text{ yr}^{-1})$  inside of the suboxic waters (Figure

11g). A significant decrease of vertical mixing (>3 mmol  $m^{-3}$  yr<sup>-1</sup>) is mainly found above ~400 m in the OMZs, which corresponds to the decrease in vertical gradient of DO concentration (Figure 11h).

375

There is evidence that the physical and biogeochemical processes have multiple interactions with impacts on various physical, chemical and biological fields and implications for DO dynamics (Breitburg et al., 2018; Duteil and Oschlies, 2011; Oschlies et al., 2018). For example, observational and modelling studies show that changes in vertical mixing intensity can affect the distributions of organic matter thus oxygen consumption at mid depth (Duteil and Oschlies, 2011: Talley et al., 2016), and 380 vertical distributions of DOM concentration and its remineralization around the OMZ in turn can alter the intensity of vertical mixing for DO (Loginova et al., 2019). Recent studies have demonstrated that a changing O:C utilization ratio (or respiration quotient) has various impacts on biological and chemical processes, with an impact on microbial respiration thus oxygen consumption (Moreno et al., 2020; Tanioka and Matsumoto, 2020). In particular, applying a smaller O:C utilization ratio leads to lower consumption rates, thus higher DO levels (Moreno et al., 2020), which would have large effects on DO gradients thus

385

vertical mixing particularly in low DO waters (e.g., in the OMZs).

#### 4.4 Impacts of biological and physical processes on asymmetric OMZs

There is evidence of asymmetric features in many biogeochemical parameters Vertical mixing of DO shows an increase (~1-2 mmol  $m^{-3} vr^{-1}$ ) outside of the OMZs and a decrease (~2-8 mmol  $m^{-3} vr^{-1}$ ) inside of the OMZs in response to the combination of enhanced background diffusion and a reduced O:C utilization ratio (Figure 11k), which is similar to the net response of

- 390 vertical mixing to the changes caused by individual parameters (see Figure 11c and 11g). However, the combined effects exceed the sum of two individual responses in the south OMZ and the lower part of the north OMZ (Figure 11o). An early study has demonstrated that enhanced background diffusion can lead to an increase not only in vertical mixing of DO directly, but also in biological consumption caused by enhanced export production in the tropical Pacific OMZs (Duteil and Oschlies, 2011), which in turn changes the vertical gradient of DO concentration, thus affects the intensity of vertical mixing.
- 395

#### 4.4 Impacts of biological and physical processes on asymmetric OMZs

There is evidence of asymmetric features in many biogeochemical fields in the tropical Pacific. For example, POC flux at 500 m is greater in the northern tropical Pacific (~4 mmol C m<sup>-2</sup> d<sup>-1</sup>) (Van Mooy et al., 2002) than in the southern tropical Pacific (<1 mmol C m<sup>-2</sup> d<sup>-1</sup>) (Pavia et al., 2019). Similarly, our regional model reproduces an asymmetric pattern for POC flux, with 400 larger values to the north than to the south. Field studies have reported an asymmetry in DOM distribution over ~200-1000 m in the central-eastern tropical Pacific, i.e., higher levels of DON and DOC to the north than to the south (Hansell, 2013; Libby and Wheeler, 1997; Raimbault et al., 1999). Our best model simulation (i.e., the Km18.7Kb0.5 simulation) also reveals an asymmetric DON at mid-depthbelow 300 m, i.e., ~6-78 mmol m<sup>-3</sup> in the ETNP and ~43-5 mmol m<sup>-3</sup> in the ETSP (data not showFigure S3a). However, an earlier field study reported higher rates of organic carbon remineralization over 200-1000 m to the south ( $\sim 2-10 \text{ mmol m}^{-3} \text{ yr}^{-1}$ ) than to the north ( $\sim 1-6 \text{ mmol m}^{-3} \text{ yr}^{-1}$ ) in the eastern/central tropical Pacific (Feely et al., 405 2004). Similarly, our model simulation also shows such an asymmetric feature of in biological consumption below 200 over <u>300-600</u> m in the tropical Pacific, i.e.,  $\sim 2-84$  mmol m<sup>-3</sup> yr<sup>-1</sup> in the ETSP and  $\sim \leq 1-6$  mmol m<sup>-3</sup> yr<sup>-1</sup> in the ETNP. (Figure S3b).

It appears that the asymmetric distributions differ largely between biological parameters, and fields in the tropical Pacific. In 410 particular, there are almost opposite patterns between oxygen consumption (or DOM remineralization) and DOM concentration. This discrepancy, which may be attributed to the rates difference in the rate of DOM remineralization in the water column, which between the north and south. The rate of DOM remineralization is determined not only by DOM concentrationmicrobial activity, but also by the stoichiometry associated with microbial respiration (Wang et al., 2008; Zakem and Levine, 2019). Recent studies on the respiration quotient demonstrate that the O:C utilization ratio is lower to the north

than to the south in the tropical Pacific (Tanioka and Matsumoto, 2020; Wang et al., 2019), which primarily reflects the 415

difference in oxygen limitation on microbial respiration (Kalvelage et al., 2015). Apparently, <u>suchthe</u> asymmetry in biological consumption <u>(lower rate in the north than in the south)</u> cannot explain the asymmetry in the tropical Pacific OMZs (i.e., lower DO levels to the north than to the south), indicating that other processes are responsible for the asymmetry.

- 420 Numerous studies have indicated that physical mixing is the only source of DO for the tropical OMZs (Brandt et al., 2015; Czeschel et al., 2012; Duteil et al., 2020). For example, There is evidence that turbulent diffusion is argued to accountaccounted for 89% of the net DO supply for the core OMZ of souththe southern tropical Pacific (Llanillo et al., 2018). There is evidence that larger scale mass transport due to circulation and ventilation is more efficient in the south Pacific than in the north Pacific (Kuntz and Schrag, 2018), and the transit time from the surface to the OMZ is much longer in the ETNP than in the ETSP (Fu
- 425 et al., 2018). Both our analyses and other modeling studies (Duteil, 2019; Shigemitsu et al., 2017) demonstrate that DO supply via vertical mixing is much weaker in the northern OMZ than in the southern OMZ in the tropical Pacific. All these analyses indicate that physical processes play a major role in shaping the asymmetry of the OMZs over the tropical Pacific.
  <u>. Our model simulations show that zonal, meridional and vertical advections for DO supply are relatively weak (<2 mmol m<sup>-3</sup> yr<sup>-1</sup>). However, the intensity of vertical mixing is much stronger (~2-6 mmol m<sup>-3</sup> yr<sup>-1</sup>) at mid-depth, indicating that vertical</u>
- 430 mixing plays a bigger role in supplying DO into the OMZs.

Our further analyses show that the intensity of vertical mixing over 200-700 m is stronger to the south than to the north of the equator (Figure S2), which is consistent with some other modelling studies that reported stronger DO supply via vertical mixing in the south OMZ than in the north OMZ in the tropical Pacific (Duteil, 2019; Shigemitsu et al., 2017). There is evidence that

435 larger-scale mass transport due to circulation and ventilation is more efficient in the South Pacific than in the North Pacific (Kuntz and Schrag, 2018), and the transit time from the surface to the OMZ is much longer in the ETNP than in the ETSP (Fu et al., 2018). All these analyses indicate that vertical mixing is largely responsible for asymmetric distribution of mid-depth DO, and physical processes play a major role in shaping the asymmetry of the OMZs in the tropical Pacific.

#### 440 **4.5 Implications and limitations of the current research**

There are inter-dependencies between the physical and biogeochemical processes <u>at mid-depth</u> (Duteil and Oschlies, 2011; Gnanadesikan et al., 2012; Niemeyer et al., 2019), which <u>maycan</u> have <u>influencesan influence</u> on the asymmetry of OMZs in the tropical Pacific. Our study shows that <u>the</u> rate of physical supply is sensitive to changes in <u>both physical-vertical mixing</u> <u>below 400 m</u> and biological <u>parameterizations</u>, <u>particularly in low DO waters</u>.consumption over 200-400 m. Since the <u>physical</u>

445 contribution <u>of physical supply to mid-depth DO flux</u> exceeds <u>thethat of</u> biological <u>contribution to mid-depth DO-consumption</u> in the tropical Pacific (Llanillo et al., 2018; Montes et al., 2014), and the physical processes are more dominant in the ETSP, one may expect that physical-biological feedbacks are stronger to the south, which can lead to <u>a</u> relatively larger- net flux into the south OMZ.

- 450 Physical and biogeochemical interactions are complex over spaceand region-specific, which haveproduce direct and indirect effects on the sourcesources and sinksinks of DO (Levin, 2018; Oschlies et al., 2018). Our study demonstrates that there is a much greater increase in net DO flux in the core OMZ to the south than to the north that results from these interactions and feedbacks (Fig. 10p). On the one hand, supply of DO is greater under stronger physical transport in the south tropical Pacific. On the other hand, stronger physical processes can also lead to higher levels of nutrients and biological production and thus
- enhanced export production and oxygen consumption at mid-depth (Duteil and Oschlies, 2011), which can offset the rate of physical supply. In addition, stronger physical processes can also result in strengthened transport of DO and OM out to other regions (Gnanadesikan et al., 2012; Yu et al., 2021), which hashave complex impacts on DO balance in the southernsouth OMZ.
- 460 ThereTo date, most regional to global models have difficulty in reproducing the observed asymmetric OMZs in the tropical Pacific (Cabre et al., 2015; Duteil, 2019), which is probably caused by misrepresentations of physical processes such as background diffusion at mid-depth and ocean circulation (Cabre et al., 2015; Duteil and Oschlies, 2011). In addition, model configuration such as vertical and horizontal resolutions can also influence physical transportations of DO (Busecke et al., 2019; Duteil et al., 2014), and distributions of nutrients (with impacts on biological production and DO consumption)(Berthet
- 465 <u>et al., 2019</u>). Other possible causes may be associated with the ocean-atmosphere interactions and feedbacks due to the <u>uncertainties in atmospheric forcing fields</u> (Duteil, 2019; Ridder and England, 2014; Stramma et al., 2012).
- Although there have been advances in our understanding of the regulation of DO consumption by biogeochemical processes, large scale models do not have representative processes due to various reasons. For example, there is evidence of DO depletion 470 at mid-depth caused by zooplankton migration (Bianchi et al., 2013; Kiko et al., 2017), and there are strong interactions and feedbacks between carbon, nitrogen and oxygen cycles in marine ecosystemecosystems. Limited studies indicate that O:C:N utilization ratios during microbial respiration vary largely in the water column (Moreno et al., 2020; Zakem and Levine, 2019), and nitrogen. Nitrogen cycling (e.g., oxidation, nitrification and denitrification) not only has impacts on oxygen consumption/production but is also is influenced by the oxygen level (Beman et al., 2021; Kalvelage et al., 2013; Oschlies et 475 al., 2019; Sun et al., 2021). However, little attention has been paid to understandunderstanding the coupling of carbon and oxygen cycles. It should be noted that; the available data are also not sufficient for the parameterizations of relevant processes, which has hampered our ability to assess the impacts of those biogeochemical processes associated with the nitrogen cycle on oxygen fields. Future observational and modelling studies are needed not only to improve our knowledge on the coupling of carbon, nitrogen and oxygen cycles in the ocean, but also to advance our understanding on the physical and biogeochemical 480 interactions and feedbacks associated with the marine stoichiometry.

### 5. Conclusion

In this paper, we use a basin scale model to investigate the impacts of parameterizations of vertical mixing and DOM remineralization on the dynamics of mid-depth DO, and analyse the underlying mechanisms for asymmetric OMZs in the tropical Pacific. Our study showsresults show that the model is capable of reproducing the observed DO distributions and asymmetric OMZs with the combination of enhanced vertical mixing and <u>a</u> reduced O:C utilization ratio that causes an increase in DO concentration (or net flux) at mid-depth. Overall, enhanced vertical mixing makes a <u>greaterlarger</u> contribution to the increase <u>over-of DO below ~400-1000</u> m, and the contribution from <u>a</u> reduced O:C utilization ratio is greater over 200-700400 m.

490 Our analyses demonstrate that there is a largemodest increase in physical supply and a small increase in biological consumption under enhanced vertical mixing, and the increase in consumption is a result of redistribution of DOM in the water column. On the other hand, applying a reduced O:C utilization ratio leads toresults in a large decrease in both biological consumption, and a small decrease in physical supply in the OMZs (due to the vertical changes in vertical DO gradients). These findings point to strong physical-biological interactions and feedbacks at mid-depth in the tropical Pacific-OMZs.

495

485

This study suggestsuggests that biological consumption (i.e., greater rate to the south) cannot explain the asymmetric feature in the tropical Pacific OMZs (i.e., lower DO levels to the north), but physical processes (i.e., stronger supplyvertical mixing to the south) play a major role in shaping the asymmetric OMZs of the tropical Pacific. In addition, the interactions between physical and biological processes are also stronger in the southernsouth OMZ than in the northernnorth OMZ, probablylikely because physical supply is sensitive to changes in boththe parameterizations of both vertical mixing and DOM remineralization. Further studies with improved approaches will enable toat better understandunderstanding of the interactions and feedbacks between physical and biogeochemical processes.

### Phytoplankton equations

$$\frac{\partial P_S}{\partial t} = \mu_S P_S - g_{P_S} (1 - e^{-\Lambda P_S}) Z_S - m_S P_S$$
(B1)  

$$\frac{\partial P_L}{\partial t} = \mu_L P_L - g_{P_L 1} (1 - e^{-\Lambda P_L}) Z_L - g_{P_L 2} (1 - e^{-\Lambda P_L}) Z_S - m_L P_L$$
(B2)

510

### **Zooplankton equations**

$$\frac{\partial Z_S}{\partial t} = [\lambda(g_{P_S}(1 - e^{-\Lambda P_S}) + g_{P_L 2}(1 - e^{-\Lambda P_L})) + g_{D_S}(1 - e^{-\Lambda D_S}) + g_{D_L 2}(1 - e^{-\Lambda D_L}) - (r_s + \delta_s)]Z_S - g_{Z_S}(1 - e^{-\Lambda Z_S})Z_L$$
(B3)

$$\frac{\partial Z_L}{\partial t} = [\lambda(g_{P_L 1}(1 - e^{-\Lambda P_L}) + g_{Z_S}(1 - e^{-\Lambda Z_S})) + g_{D_L 1}(1 - e^{-\Lambda D_L}) - (r_L + \delta_L)]Z_L$$
(B4)

515

# **Detritus equations**

$$\frac{\partial D_S}{\partial t} = (m_S P_S + m_L P_L + (r_s Z_S + r_L Z_L)\chi)(1 - \gamma) - g_{D_S}(1 - e^{-\Lambda D_S})Z_S - (c_{D_S} + \omega_{D_S} h^{-1})D_S$$
(B5)  

$$\frac{\partial D_L}{\partial t} = (1 - \lambda)[\left(g_{P_S}(1 - e^{-\Lambda P_S}) + g_{P_L 2}(1 - e^{-\Lambda P_L})\right)Z_S + \left(g_{P_L 1}(1 - e^{-\Lambda P_L}) + g_{Z_S}(1 - e^{-\Lambda Z_S})\right)Z_L] + \delta_S Z_S + \delta_L Z_L - (c_{D_L} + \omega_{D_L} h^{-1})D_L - g_{D_L 2}(1 - e^{-\Lambda D_L})Z_S - g_{D_L 1}(1 - e^{-\Lambda D_L})Z_L$$
(B6)

520

### **DON equations**

$$\frac{\partial DON}{\partial t} = (m_S P_S + m_L P_L + (r_S Z_S + r_L Z_L)\chi)\gamma + (c_{D_S} D_S + c_{D_L} D_L)\zeta - c_{DON} DON$$
(B7)

### **Nutrients equations**

$$525 \quad \frac{\partial NO_{3}}{\partial t} = -\mu_{S}P_{S}\frac{N_{S\_UP}}{N_{S\_UP}+A_{UP}} - \mu_{L}P_{L}\frac{N_{L\_UP}}{N_{L\_UP}+A_{UP}} + \varphi NH_{4}$$
(B8)  
$$\frac{\partial NH_{4}}{\partial t} = -\mu_{S}P_{S}\frac{A_{up}}{N_{S\_UP}+A_{UP}} - \mu_{L}P_{L}\frac{A_{up}}{N_{L\_UP}+A_{UP}} + (r_{S}Z_{S} + r_{L}Z_{L})(1-\chi) + c_{DON}DON + (c_{D_{S}}D_{S} + c_{D_{L}}D_{L})(1-\zeta) - \varphi NH_{4}$$
(B9)

$$\frac{\partial Fe}{\partial t} = -(\mu_S P_S R_S + \mu_L P_L R_L - s_{Fe} D_L Fe) + R_S [(r_S Z_S + r_L Z_L)(1 - \chi) + c_{DON} DON + c_{D_S} D_S + c_{D_L} D_L (1 - \zeta)]$$
(B10)

### 530 Nitrogen uptake

$$N_{S\_UP} = \frac{NO_3}{K_{S\_NO_3} + NO_3} \left(1 - \frac{NH_4}{K_{NH_4} + NH_4}\right)$$
(B11)

$$N_{L_{\_}UP} = \frac{NO_3}{K_{L_{\_}NO_3} + NO_3} \left(1 - \frac{NH_4}{K_{NH_4} + NH_4}\right)$$
(B12)

$$A_{UP} = \frac{NH_4}{K_{NH_4} + NH_4}$$

### 535 Other equations

### Phytoplankton growth rate

$$\mu_{S} = \mu_{S0} e^{k_{T}T} f(I) \psi_{S}(N, Fe)$$
(B14)

$$\mu_L = \mu_{L0} e^{k_T T} f(I) \psi_L(N, Fe)$$
(B15)

540

### Nutrient limitation

$$\psi_{S}(N,Fe) = \min\left(\frac{NO_{3}+NH_{4}}{K_{S_{N}}+NO_{3}+NH_{4}}, \frac{Fe}{K_{S_{F}}e+Fe}\right)$$
(B16)

$$\psi_L(N, Fe) = \min\left(\frac{NO_3 + NH_4}{K_{L_N} + NO_3 + NH_4}, \frac{Fe}{K_{L_Fe} + Fe}\right)$$
(B17)

### 545 Light limitation

$$f(I) = 1 - e^{-\frac{\alpha I}{\eta P_{MAX}}}$$
(B18)

### Light attenuation

$$I(z) = I_0 exp^{-k_A Z}$$
(B19)

550 
$$k_A = k_W + k_C \text{Chl} + k_D (D_S + D_L)$$
 (B20)

### Detritus decomposition and DON remineralization

$$c_{DS} = c_{DS0} e^{k_B (T - T_0)}$$
(B21)  
$$c_{DL} = c_{DL0} e^{k_B (T - T_0)}$$
(B22)

555 
$$c_{DON} = c_{DON0} e^{k_B (T - T_0)}$$
 (B23)

### Phytoplankton carbon to chlorophyll ratio $(\eta)$

$$Chl = \left(\frac{P_S}{\eta_S} + \frac{P_L}{\eta_L}\right) R_{C:N}$$
(B24)

$$\eta_{S} = \eta_{S0} - (\eta_{S0} - \eta_{MIN}) \frac{\ln I_0 - \ln I}{4.605}$$
(B25)

560 
$$\eta_L = \eta_{L0} - (\eta_{L0} - \eta_{MIN}) \frac{\ln I_0 - \ln I}{4.605}$$
 (B26)

$$\eta_{S0} = \eta_{S\_MAX} - k_{PS}\mu_S^* \tag{B27}$$

$$\eta_{L0} = \eta_{L\_MAX} - k_{PL} \mu_L^* \tag{B28}$$

$$\mu_{S}^{*} = \mu_{S0} e^{k_{T}T} \min\left(\frac{NO_{3}}{K_{S\_N} + NO_{3}}, \frac{\text{Fe}}{K_{S\_\text{Fe}} + \text{Fe}}\right)$$
$$\mu_{L}^{*} = \mu_{L0} e^{k_{T}T} \min\left(\frac{NO_{3}}{K_{L\_N} + NO_{3}}, \frac{\text{Fe}}{K_{L\_\text{Fe}} + \text{Fe}}\right)$$

565

## Appendix B: Model biogeochemical parameters

Symbol	Parameter	Unit	Value
m <sub>S</sub>	Small phytoplankton mortality rate	d-1	0.15
$m_{ m L}$	Large phytoplankton mortality rate	d <sup>-1</sup>	0.35
r <sub>S</sub>	Small zooplankton excretion rate	d <sup>-1</sup>	0.53
r <sub>L</sub>	Large zooplankton excretion rate	d <sup>-1</sup>	0.44
δs	Small zootoplankton mortality rate	d <sup>-1</sup>	0.12
$\delta_{ ext{L}}$	Large zootoplankton mortality rate	d <sup>-1</sup>	0.12
$g_{\rm PS}$	Maximum grazing rate for small phytoplanktion	d <sup>-1</sup>	2.6
$g_{\rm PL1}$	Maximum grazing rate for large phytoplanktion	d <sup>-1</sup>	1.2
gzs	Maximum grazing rate for small zootoplanktion	d <sup>-1</sup>	1.7
$g_{\rm PL2}$	Maximum grazing rate for large phytoplanktion	d <sup>-1</sup>	0.9
$g_{\rm DS}$	Maximum grazing rate for small detritus	d <sup>-1</sup>	1.0
$g_{DL1}$	Maximum grazing rate for large detritus	d <sup>-1</sup>	3.0
$g_{\rm DL2}$	Maximum grazing rate for large detritus	d-1	1.5
Λ	Ivlev coefficient	(mmol m <sup>-3</sup> ) <sup>-1</sup>	0.5
λ	Zootoplankton assimilation coefficient	%	75
χ	Excretion coefficient	%	55
γ	Dissolution coefficient	%	90
ξ	Dissolution coefficient	%	90
R <sub>C:N</sub>	C:N ratio	mol:mol	6.625
R <sub>s</sub>	Fe:N ratio for small phytoplankton	$\mu$ mol:mol	15
R <sub>L</sub>	Fe:N ratio for large phytoplankton	$\mu$ mol:mol	40
$\eta_{S_{MIN}}$	Minimum PhyC:Chl ratio in small phytoplanktion	g:g	30
$\eta_{L\_MIN}$	Minimum PhyC:Chl ratio in large phytoplanktion	g:g	15
$\eta_{S_{MAX}}$	Maximum PhyC:Chl ratio in small phytoplanktion	g:g	200
$\eta_{L\_MAX}$	Maximum PhyC:Chl ratio in large phytoplanktion	g:g	120
$k_{\rm PS}$	Photoacclimation coefficient for small phytoplanktion	(g:g)d	95
$k_{\rm PL}$	Photoacclimation coefficient for large phytoplanktion	(g:g)d	70

(B29)

(B30)

WDS	Sinking velocity for small detritus	m d <sup>-1</sup>	1
WDL	Sinking velocity for large detritus	m d <sup>-1</sup>	3.5
arphi	Nitrification rate (when I<5 $\mu$ molEm <sup>-2</sup> s <sup>-1</sup> )	d <sup>-1</sup>	0.04
SFe	Iron scavenge coefficient	d <sup>-1</sup> (nmol Fe m <sup>-3</sup> ) <sup>-1</sup>	0.00001
$\mu_{ m S0}$	Maximum growth rate at 0°C for small phytoplankton	d <sup>-1</sup>	0.58
$\mu_{ m L0}$	Maximum growth rate at 0°C for large phytoplankton	d <sup>-1</sup>	1.16
$k_{\mathrm{T}}$	Temp. Dependent coefficient for $\mu$	°C-1	0.06
$K_{S_N}$	Half saturation constant for N limitation	mmol m <sup>-3</sup>	0.3
$K_{L_N}$	Half saturation constant for N limitation	mmol m <sup>-3</sup>	0.9
$K_{S\_Fe}$	Half saturation constant for iron limitation	mmol m <sup>-3</sup>	14
$K_{L_{Fe}}$	Half saturation constant for iron limitation	mmol m <sup>-3</sup>	150
$K_{S_NO3}$	Half saturation constant for nitrate uptake	mmol m <sup>-3</sup>	0.3
K <sub>L_NO3</sub>	Half saturation constant for nitrate uptake	mmol m <sup>-3</sup>	0.9
$\mathbf{K}_{\mathrm{NH4}}$	Half saturation constant for ammonium uptake	mmol m <sup>-3</sup>	0.05
α	Initial slope of the P – I curve	mg C mg chl <sup>-1</sup> ( $\mu mol \to m^{-2} s^{-1}$ ) <sup>-1</sup>	0.02
$P_{\rm MAX}$	Maximum carbon specific growth rate	h-1	0.036
$k_{ m W}$	Light attenuation constant for water	m <sup>-1</sup>	0.028
$k_{\rm C}$	Light attenuation constant for chlorophyll	$m^{-1} (mg chl m^{-3})^{-1}$	0.058
k <sub>D</sub>	Light attenuation constant for detritus	$m^{-1} (mg chl m^{-3})^{-1}$	0.008
$c_{\rm DS0}$	Small detritus decomposition rate at 0°C	d <sup>-1</sup>	0.001
$c_{\rm DL0}$	Large detritus decomposition rate at 0°C	d <sup>-1</sup>	0.008

Appendix C: Comparisons in biogeochemical paran	eters
---	-------

Symbol	Parameter	Unit	Yu et al. (2021)	This study		
T <sub>0</sub>	Limit temperature	°C	10	0		
$k_{\rm B}$	Temperature dependent coefficient	-	0.002	0.001		
C <sub>DON0</sub>	DON remineralization constant	<b>d</b> <sup>-1</sup>				
	0-100 m		0.001	0.00075		
	100-600 m		0.0002-0.001	0.00013-0.00075*		
	600-1000 m		0.0002	0.00003-0.00013*		

\*  $C_{DON0}$  decreases with depth by an exponential function.

575 *Code and data availability.* The exact version of the software code used to produce the results presented in this paper is archived on Zenodo (https://doi.org/10.5281/zenodo.5148146, Wang et al., 2021). Other code and data are available upon request from the authors. Request for materials should be addressed to X.J.W. (xwang@bnu.edu.cn).

*Author contributions.* X.J.W. and K.W. designed the study, performed the simulations and prepared the manuscript. R.M., 580 D.X.Z. and R.H.Z. contributed to analysis, interpretation of results and writing.

Competing interests. The authors declare that they have no conflict of interest.

*Acknowledgements.* This work was supported by the Chinese Academy of Sciences' Strategic Priority Project (XDA1101010504). The authors wish to acknowledge the use of the Ferret (http://ferret.pmel.noaa.gov/Ferret/).

### References

- Andrews, O., Buitenhuis, E., Le Quere, C., and Suntharalingam, P.: Biogeochemical modelling of dissolved oxygen in a changing ocean, Philosophical transactions. Series A, Mathematical, physical, and engineering sciences, 375, 2017.
- Bao, Y. and Li, Y.: Simulations of dissolved oxygen concentration in CMIP5 Earth system models, Acta Oceanologica Sinica, 35, 28-37, 2016.

 Beman, J. M., Vargas, S. M., Vazquez, S., Wilson, J. M., Yu, A., Cairo, A., and Perez-Coronel, E.: Biogeochemistry and hydrography shape microbial community assembly and activity in the eastern tropical North Pacific Ocean oxygen minimum zone, Environmental microbiology, 23, 2765-2781, 2020.

- Beman, J. M., Vargas, S. M., Vazquez, S., Wilson, J. M., Yu, A., Cairo, A., and Perez-Coronel, E.: Biogeochemistry and hydrography shape microbial community assembly and activity in the eastern tropical North Pacific Ocean oxygen minimum zone, Environmental microbiology, 23, 2765-2781, 2021.
- Beman, J. M., Vargas, S. M., Vazquez, S., Wilson, J. M., Yu, A., Cairo, A., and Perez Coronel, E.: Biogeochemistry and
   hydrography shape microbial community assembly and activity in the eastern tropical North Pacific Ocean oxygen minimum zone, Environmental microbiology, 23, 2765–2781, 2020.

Bertagnolli, A. D. and Stewart, F. J.: Microbial niches in marine oxygen minimum zones, Nature reviews. Microbiology, 16, 723-729, 2018.

Berthet, S., Séférian, R., Bricaud, C., Chevallier, M., Voldoire, A., and Ethé, C.: Evaluation of an Online Grid - Coarsening

Algorithm in a Global Eddy - Admitting Ocean Biogeochemical Model, Journal of Advances in Modeling Earth Systems, 11, 1759-1783, 2019.
 Bettencourt, J. H., Lopez, C., Hernandez-Garcia, E., Montes, I., Sudre, J., Dewitte, B., Paulmier, A., and Garcon, V.: Boundaries of the Peruvian oxygen minimum zone shaped by coherent mesoscale dynamics, Nature Geoscience, 8, 937-

U967, 2015.

- Bianchi, D., Dunne, J. P., Sarmiento, J. L., and Galbraith, E. D.: Data-based estimates of suboxia, denitrification, and N2O production in the ocean and their sensitivities to dissolved O2, Global Biogeochemical Cycles, 26, 1-13, 2012.
   <u>Bianchi, D., Galbraith, E. D., Carozza, D. A., Mislan, K. A. S., and Stock, C. A.: Intensification of open-ocean oxygen depletion by vertically migrating animals, Nature Geoscience, 6, 545-548, 2013.</u>
- Bopp, L., Le Quere, C., Heimann, M., Manning, A. C., and Monfray, P.: Climate-induced oceanic oxygen fluxes:
  Implications for the contemporary carbon budget, Global Biogeochemical Cycles, 16, 1-13, 2002.
  Brandt, P., Bange, H. W., Banyte, D., Dengler, M., Didwischus, S. H., Fischer, T., Greatbatch, R. J., Hahn, J., Kanzow, T., Karstensen, J., Krortzinger, A., Krahmann, G., Schmidtko, S., Stramma, L., Tanhua, T., and Visbeck, M.: On the role of

circulation and mixing in the ventilation of oxygen minimum zones with a focus on the eastern tropical North Atlantic, Biogeosciences, 12, 489-512, 2015.

620 Breitburg, D., Levin, L. A., Oschlies, A., Gregoire, M., Chavez, F. P., Conley, D. J., Garcon, V., Gilbert, D., Gutierrez, D., Isensee, K., Jacinto, G. S., Limburg, K. E., Montes, I., Naqvi, S. W. A., Pitcher, G. C., Rabalais, N. N., Roman, M. R., Rose, K. A., Seibel, B. A., Telszewski, M., Yasuhara, M., and Zhang, J.: Declining oxygen in the global ocean and coastal waters, Science, 359, 2018.

Busecke, J. J. M., Resplandy, L., and Dunne, J. P. P.: The Equatorial Undercurrent and the Oxygen Minimum Zone in the Pacific, Geophysical Research Letters, doi: 10.1029/2019GL082692, 2019. 6716–6725, 2019.

- 625 Pacific, Geophysical Research Letters, doi: 10.1029/2019GL082692, 2019. 6716–6725, 2019. Cabre, A., Marinov, I., Bernardello, R., and Bianchi, D.: Oxygen minimum zones in the tropical Pacific across CMIP5 models: mean state differences and climate change trends, Biogeosciences, 12, 5429-5454, 2015. Chen, D., Rothstein, L. M., and Busalacchi, A. J.: A Hybrid Vertical Mixing Scheme and Its Application to Tropical Ocean Models, Journal of Physical Oceanography, 24, 2156-2179, 1994.
- 630 Christian, J. R., Verschell, M. A., Murtugudde, R., Busalacchi, A. J., and McClain, C. R.: Biogeochemical modelling of the tropical Pacific Ocean. I: Seasonal and interannual variability, Deep Sea Research Part II: Topical Studies in Oceanography, 49, 509-543, 2001.

Czeschel, R., Stramma, L., and Johnson, G. C.: Oxygen decreases and variability in the eastern equatorial Pacific, J Geophys Res-Oceans, 117, 1-12, 2012.

635 Czeschel, R., Stramma, L., Schwarzkopf, F. U., Giese, B. S., Funk, A., and Karstensen, J.: Middepth circulation of the eastern tropical South Pacific and its link to the oxygen minimum zone, J Geophys Res-Oceans, 116, 2011.

DeVries, T., Le Quere, C., Andrews, O., Berthet, S., Hauck, J., Ilyina, T., Landschuetzer, P., Lenton, A., Lima, I. D., Nowicki, M., Schwinger, J., and Seferian, R.: Decadal trends in the ocean carbon sink, Proceedings of the National Academy of Sciences of the United States of America, 116, 11646-11651, 2019.

640 Duteil, O.: Wind Synoptic Activity Increases Oxygen Levels in the Tropical Pacific Ocean, Geophysical Research Letters, 46, 2715-2725, 2019.

Duteil, O., Frenger, I., and Getzlaff, J.: Intermediate water masses, a major supplier of oxygen for the eastern tropical Pacific ocean, Ocean Science, doi: 10.5194/os-2020-17, 2020. 2020.

Duteil, O. and Oschlies, A.: Sensitivity of simulated extent and future evolution of marine suboxia to mixing intensity, 645 Geophysical Research Letters, 38, 2011.

Duteil, O., Schwarzkopf, F. U., Böning, C. W., and Oschlies, A.: Major role of the equatorial current system in setting oxygen levels in the eastern tropical Atlantic Ocean: A high-resolution model study, Geophysical Research Letters, 41, 2033-2040, 2014.

### Feely, R. A., Sabine, C. L., Schlitzer, R., Bullister, J. L., Mecking, S., and Greeley, D.: Oxygen utilization and organic

- 650 carbon remineralization in the upper water column of the Pacific Ocean, Journal of Oceanography, 60, 45-52, 2004.
   <u>Feely, R. A., Wanninkhof, R., Takahashi, T., and Tans, P.: Influence of El Niño on the equatorial Pacific contribution to atmospheric CO2 accumulation, Nature, 398, 597-601, 1999.</u>
   Fu, W. W., Bardin, A., and Primeau, F.: Tracing ventilation source of tropical pacific oxygen minimum zones with an
- adjoint global ocean transport model, Deep-Sea Research Part I: Oceanographic Research Papers, 139, 95-103, 2018.
  Fuenzalida, R., Schneider, W., Garces-Vargas, J., Bravo, L., and Lange, C.: Vertical and horizontal extension of the oxygen minimum zone in the eastern South Pacific Ocean, Deep-Sea Res Pt Ii, 56, 1027-1038, 2009.
  Garçon, V., Karstensen, J., Palacz, A., Telszewski, M., Aparco Lara, T., Breitburg, D., Chavez, F., Coelho, P., Cornejo-D'Ottone, M., Santos, C., Fiedler, B., Gallo, N. D., Grégoire, M., Gutierrez, D., Hernandez-Ayon, M., Isensee, K., Koslow, T., Levin, L., Marsac, F., Maske, H., Mbaye, B. C., Montes, I., Naqvi, W., Pearlman, J., Pinto, E., Pitcher, G., Pizarro, O.,
- 660 Rose, K., Shenoy, D., Van der Plas, A., Vito, M. R., and Weng, K.: Multidisciplinary Observing in the World Ocean's Oxygen Minimum Zone Regions: From Climate to Fish — The VOICE Initiative, Frontiers in Marine Science, 6, 2019. Gnanadesikan, A., Dunne, J. P., and John, J.: Understanding why the volume of suboxic waters does not increase over centuries of global warming in an Earth System Model, Biogeosciences, 9, 1159-1172, 2012. Hansell, D. A.: Recalcitrant dissolved organic carbon fractions, Annual review of marine science, 5, 421-445, 2013.
- Johnson, K. S., Gordon, R. M., and Coale, K. H.: What controls dissolved iron concentrations in the world ocean?, Marine Chemistry, 57, 137-161, 1997.
  Kalnay, E., Kanamitsu, M., Kistler, R., Collins, W., Deaven, D., Gandin, L., Iredell, M., Saha, S., White, G., Woollen, J., Zhu, Y., Chelliah, M., Ebisuzaki, W., Higgins, W., Janowiak, J., Mo, K. C., Ropelewski, C., Wang, J., Leetmaa, A., Reynolds, R., Jenne, R., and Joseph, D.: The NCEP/NCAR 40-year reanalysis project, B Am Meteorol Soc, 77, 437-471,
- 670 1996. Kalvelage, T., Lavik, G., Jensen, M. M., Revsbech, N. P., Loscher, C., Schunck, H., Desai, D. K., Hauss, H., Kiko, R., Holtappels, M., LaRoche, J., Schmitz, R. A., Graco, M. I., and Kuypers, M. M.: Aerobic microbial respiration In oceanic oxygen minimum zones, PloS one, 10, 2015.

Kalvelage, T., Lavik, G., Lam, P., Contreras, S., Arteaga, L., Löscher, C. R., Oschlies, A., Paulmier, A., Stramma, L., and Kuypers, M. M. M.: Nitrogen cycling driven by organic matter export in the South Pacific oxygen minimum zone, Nature

Kuypers, M. M. M.: Nitrogen cycling driven by organic matter export in the South Pacific oxygen minimum zone, Nature Geoscience, 6, 228-234, 2013.
Karstensen, J., Stramma, L., and Visbeck, M.: Oxygen minimum zones in the eastern tropical Atlantic and Pacific oceans, Progress in Oceanography, 77, 331-350, 2008.

Kiko, R., Biastoch, A., Brandt, P., Cravatte, S., Hauss, H., Hummels, R., Kriest, I., Marin, F., McDonnell, A. M. P.,

680 Oschlies, A., Picheral, M., Schwarzkopf, F. U., Thurnherr, A. M., and Stemmann, L.: Biological and physical influences on marine snowfall at the equator, Nature Geoscience, 10, 852-858, 2017.

Kuntz, L. B. and Schrag, D. P.: Hemispheric asymmetry in the ventilated thermocline of the Tropical Pacific, Journal of Climate 31, 1281-1288, 2018.

Le Quéré, C., Takahashi, T., Buitenhuis, E. T., Rödenbeck, C., and Sutherland, S. C.: Impact of climate change and

685 variability on the global oceanic sink of CO2, Global Biogeochemical Cycles, 24, n/a-n/a, 2010.

Levin, L. A.: Manifestation, Drivers, and Emergence of Open Ocean Deoxygenation, Annual review of marine science, 10, 229-260, 2018.

Libby, P. S. and Wheeler, P. A.: Particulate and dissolved organic nitrogen in the central and eastern equatorial Pacific, Deep-Sea Research Part I: Oceanographic Research Papers, 44, 345-361, 1997.

- 690 Llanillo, P. J., Pelegri, J. L., Talley, L. D., Pena-Izquierdo, J., and Cordero, R. R.: Oxygen Pathways and Budget for the Eastern South Pacific Oxygen Minimum Zone, Journal of Geophysical Research: Oceans, 123, 1722-1744, 2018.
   Loginova, A. N., Thomsen, S., Dengler, M., Ludke, J., and Engel, A.: Diapycnal dissolved organic matter supply into the upper Peruvian oxycline, Biogeosciences, 16, 2032 2047, 2019.
- McKinley, G. A., Pilcher, D. J., Fay, A. R., Lindsay, K., Long, M. C., and Lovenduski, N. S.: Timescales for detection of trends in the ocean carbon sink, Nature, 530, 469-472, 2016.
- Montes, I., Dewitte, B., Gutknecht, E., Paulmier, A., Dadou, I., Oschlies, A., and Garcon, V.: High-resolutionmodeling of the Eastern Tropical Pacific oxygen minimum zone: Sensitivity to the tropical oceanic circulation, Journal of Geophysical Research: Oceans, 119, 5515–5532, 2014.
- Moreno, A. R., Garcia, C. A., Larkin, A. A., Lee, J. A., Wang, W. L., Moore, J. K., Primeaub, F. W., and Martiny, A. C.:
  Latitudinal gradient in the respiration quotient and the implications for ocean oxygen availability, PNAs, 117, 22866–22872, 2020.

Murtugudde, R., Seager, R., and Busalacchi, A.: Simulation of the tropical oceans with an ocean GCM coupled to an atmospheric mixed-layer model, Journal of Climate, 9, 1795-1815, 1996.

Niemeyer, D., Kriest, I., and Oschlies, A.: The effect of marine aggregate parameterisations on nutrients and oxygen minimum zones in a global biogeochemical model, Biogeosciences, 16, 3095-3111, 2019.

Oschlies, A., Brandt, P., Stramma, L., and Schmidtko, S.: Drivers and mechanisms of ocean deoxygenation, Nature Geoscience, 11, 467-473, 2018.

Oschlies, A., Koeve, W., Landolfi, A., and Kahler, P.: Loss of fixed nitrogen causes net oxygen gain in a warmer future ocean, Nature communications, 10, 2805, 2019.

710 Paulmier, A. and Ruiz-Pino, D.: Oxygen minimum zones (OMZs) in the modern ocean, Progress in Oceanography, 80, 113-128, 2009.

Pavia, F. J., Anderson, R. F., Lam, P. J., Cael, B. B., Vivancos, S. M., Fleisher, M. Q., Lu, Y., Zhang, P., Cheng, H., and Edwards, R. L.: Shallow particulate organic carbon regeneration in the South Pacific Ocean, Proceedings of the National Academy of Sciences of the United States of America, 116, 9753-9758, 2019.

715 Raimbault, P., Slawyk, G., Boudjellal, B., Coatanoan, C., Conan, P., Coste, B., Garcia, N., Moutin, T., and Pujo-Pay, M.: Carbon and nitrogen uptake and export in the equatorial Pacific at 150°W: Evidence of an efficient regenerated production cycle, Journal of Geophysical Research: Oceans, 104, 3341-3356, 1999. <u>Ridder, N. N. and England, M. H.: Sensitivity of ocean oxygenation to variations in tropical zonal wind stress magnitude.</u>

<u>Global Biogeochemical Cycles, 28, 909-926, 2014.</u>
Schmidtko, S., Stramma, L., and Visbeck, M.: Decline in global oceanic oxygen content during the past five decades, Nature, 542, 335-339, 2017.

Shigemitsu, M., Yamamoto, A., Oka, A., and Yamanaka, Y.: One possible uncertainty in CMIP5 projections of low-oxygen water volume in the Eastern Tropical Pacifi, Geophysical Research Letters, 31, 804-820, 2017.

Stramma, L., Johnson, G. C., Firing, E., and Schmidtko, S.: Eastern Pacific oxygen minimum zones: Supply paths and multidecadal changes, J Geophys Res-Oceans, 115, 2010.

Stramma, L., Johnson, G. C., Sprintall, J., and Mohrholz, V.: Expanding oxygen-minimum zones in the tropical oceans, Science, 320, 655-658, 2008.

Stramma, L., Oschlies, A., and Schmidtko, S.: Mismatch between observed and modeled trends in dissolved upper-ocean oxygen over the last 50 yr, Biogeosciences, 9, 4045-4057, 2012.

- 735 Olsen, A., Bellerby, R., Wong, C. S., Delille, B., Bates, N. R., and de Baar, H. J. W.: Climatological mean and decadal

change in surface ocean pCO2, and net sea-air CO2 flux over the global oceans, Deep Sea Research Part II: Topical Studies in Oceanography, 56, 554-577, 2009.

Talley, L. D., Feely, R. A., Sloyan, B. M., Wanninkhof, R., Baringer, M. O., Bullister, J. L., Carlson, C. A., Doney, S. C., Fine, R. A., Firing, E., Gruber, N., Hansell, D. A., Ishii, M., Johnson, G. C., Katsumata, K., Key, R. M., Kramp, M.,

- 740 Langdon, C., Macdonald, A. M., Mathis, J. T., McDonagh, E. L., Mecking, S., Millero, F. J., Mordy, C. W., Nakano, T., Sabine, C. L., Smethie, W. M., Swift, J. H., Tanhua, T., Thurnherr, A. M., Warner, M. J., and Zhang, J. Z.: Changes in ocean heat, carbon content, and ventilation: a review of the first decade of go-ship global repeat hydrography, Annual review of marine science, 8, 185-215, 2016.
- Tanioka, T. and Matsumoto, K.: Stability of Marine Organic Matter Respiration Stoichiometry, Geophysical Research Letters, 47, 2020.

Van Mooy, B. A. S., Keil, R. G., and Devol, A. H.: Impact of suboxia on sinking particulate organic carbon: Enhanced carbon flux and preferential degradation of amino acids via denitrificatio, Geochimica et Cosmochimica Acta, 66, 457–465, 2002.

Wang, W. L., Moore, J. K., Martiny, A. C., and Primeau, F. W.: Convergent estimates of marine nitrogen fixation, Nature, 566, 205-211, 2019.

Wang, X. J., Behrenfeld, M., Le Borgne, R., Murtugudde, R., and Boss, E.: Regulation of phytoplankton carbon to chlorophyll ratio by light, nutrients and temperature in the Equatorial Pacific Ocean: a basin-scale model, Biogeosciences, 6, 391-404, 2009a.

Wang, X. J., Le Borgne, R., Murtugudde, R., Busalacchi, A. J., and Behrenfeld, M.: Spatial and temporal variations in

755 dissolved and particulate organic nitrogen in the equatorial Pacific: biological and physical influences, Biogeosciences, 5, 1705-1721, 2008.

Wang, X. J., Murtugudde, R., Hackert, E., Wang, J., and Beauchamp, J.: Seasonal to decadal variations of sea surface pCO2 and sea-air CO2 flux in the equatorial oceans over 1984–2013: A basin-scale comparison of the Pacific and Atlantic Oceans, Global Biogeochemical Cycles, 29, 597-609, 2015.

Wang, X. J., Murtugudde, R., and Le Borgne, R.: Nitrogen uptake and regeneration pathways in the equatorial Pacific: a basin scale modeling study, Biogeosciences, 6, 2647-2660, 2009b.
 Wanninkhof, R.: Relationship between wind speed and gas exchange over the Ocean, J Geophys Res-Oceans, 97, 7373-7382, 1992.

Ward, B. A., Wilson, J. D., Death, R. M., Monteiro, F. M., Yool, A., and Ridgwell, A.: EcoGEnIE 1.0: plankton ecology in the cGEnIE Earth system model, Geoscientific Model Development, 11, 4241-4267, 2018.

Weiss, R. F.: The solubility of nitrogen, oxygen and argon in water and seawater, Deep-Sea Research, 17, 721-735, 1970.
Williams, J. H. T., Totterdell, I. J., Halloran, P. R., and Valdes, P. J.: Numerical simulations of oceanic oxygen cycling in the FAMOUS Earth-System model: FAMOUS-ES, version 1.0, Geoscientific Model Development, 7, 1419-1431, 2014.
Wright, J. J., Konwar, K. M., and Hallam, S. J.: Microbial ecology of expanding oxygen minimum zones, Nature Reviews
770 Microbiology, 10, 381-394, 2012.

Yu, J., Wang, X., Murtugudde, R., Tian, F., and Zhang, R. H.: Interannual - to - Decadal Variations of Particulate Organic Carbon and the Contribution of Phytoplankton in the Tropical Pacific During 1981 - 2016: A Model Study, Journal of Geophysical Research: Oceans, 126, 2021.

Zakem, E. J. and Levine, N. M.: Systematic Variation in Marine Dissolved Organic Matter Stoichiometry and Remineralization Ratios as a Function of Lability, Global Biogeochemical Cycles, 33, 1389-1407, 2019.

# Tables

Table 1. Bias and root mean square error (RMSE) for DO (mmol m<sup>-3</sup>) comparisons between WOA2013 and model simulations <u>averaged</u> over 1991-2010 in the780Eastern Tropical North Pacific (ETNP) and Eastern Tropical South Pacific (ETSP).

Layers	Statistics	Ref	Kb0.1	Kb0.25	Kb0.5	Km18.7	Km6.9	Km18.7	Km6.9 Kb0.25	Km6.9 Kb0.5	Km18.7 Kb0.25	Km18.7 Kb0.5
ETNP (165°W-90°W, 5°N-20°N)												
200-400 m	Bias	-17.44	-16.99	-16.34	-14.87	-11.32	-14.84	-11.32	-13.51	-11.85	-9.71	-7.8
	RMSE	16.35	16.21	15.73	14.91	12.43	14.63	12.43	13.83	12.84	11.4	10.2
400-700 m	Bias	-16.35	-14.37	-11.85	-7.5	-12.51	-14.95	-12.51	-9.98	-5.39	-6.88	-2.04
	RMSE	10.6	9.54	8.26	6.73	8.45	9.83	8.45	7.49	6.38	6.5	6.78
700-1000 m	Bias	-9.22	-6.61	-3.58	0.62	-5.99	-8.32	-5.99	-2.71	1.38	-5.75	3.27
	RMSE	5.1	3.06	2.93	6.52	2.64	4.29	2.64	3.59	7.19	5.39	9.08
ETSP (110°W	-80°W, 10°S	-3°S)										
200-400 m	Bias	-7.09	-6.84	-6.43	-5.39	0.19	-3.91	0.19	-2.84	-1.13	2.09	4.85
	RMSE	7.39	7.20	6.83	5.98	2.36	4.46	2.36	3.69	2.86	3.27	5.51
400-700 m	Bias	-11.3	-8.83	-5.94	-0.88	-7.94	-10.43	-7.94	-4.51	1.34	-1.21	5.23
	RMSE	12.98	10.79	8.52	6.03	10.06	12.15	10.06	7.41	5.65	5.81	7.38
700-1000 m	Bias	-7.3	-4.18	-0.97	3.38	-5.13	-7.08	-5.13	-0.62	3.94	1.05	5.46
_	RMSE	12.82	10.67	8.98	8.63	11.22	12.49	11.22	8.76	8.68	8.59	9.34

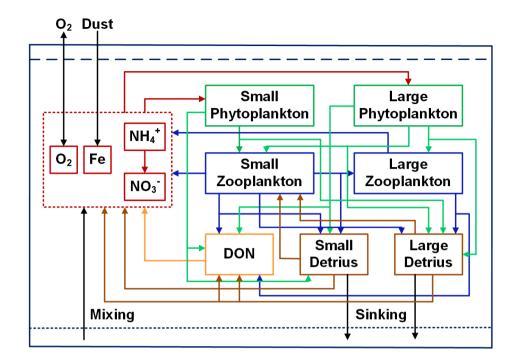
**Table 2.** Volumes (10<sup>15</sup> m<sup>3</sup>) of suboxic and hypoxic waterwaters from WOA2013 and model simulations.

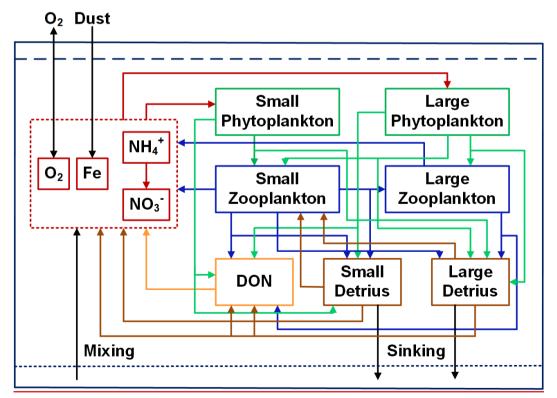
Suboxic: DO <20 mmol m<sup>-3</sup>; Hypoxic: DO <60 mmol m<sup>-3</sup>.

Regions	Waters	WOA2013	Reference	Kb0.1	Kb0.25	Kb0.5	Km6.9	Km18.7	Km6.9 Kb0.25	Km6.9 Kb0.5	Km18.7 Kb0.25	Km18.7 Kb0.5
North Pacific	Suboxic	5.97	10.61	9.74	8.73	7.33	9.98	8.83	8.08	6.68	6.88	5.55
	Hypoxic	19.98	22.67	22.58	22.32	21.61	22.5	22.17	22.11	21.35	21.71	20.91
South	Suboxic	1.43	3.78	3.34	2.86	2.15	3.39	2.78	2.42	1.71	1.81	1.12
Pacific	Hypoxic	7.12	10.42	9.85	9.19	8.17	10.21	9.8	8.94	7.88	8.49	7.39

785

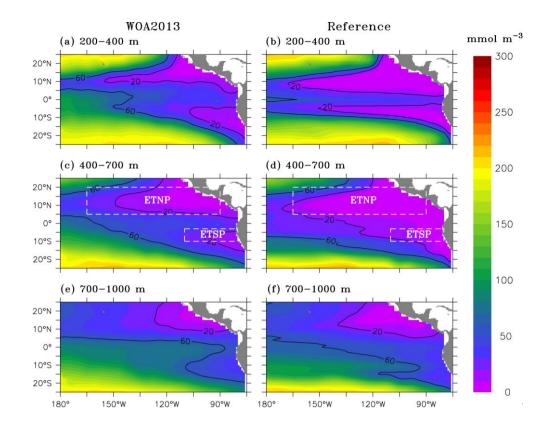
Figures





### 

**Figure 1.** Flow diagram of <u>the</u> ecosystem model. Red, green, blue, yellow and brown lines and arrows denote fluxes originating from inorganic forms, phytoplankton, zooplankton, DON and detritus, respectively.



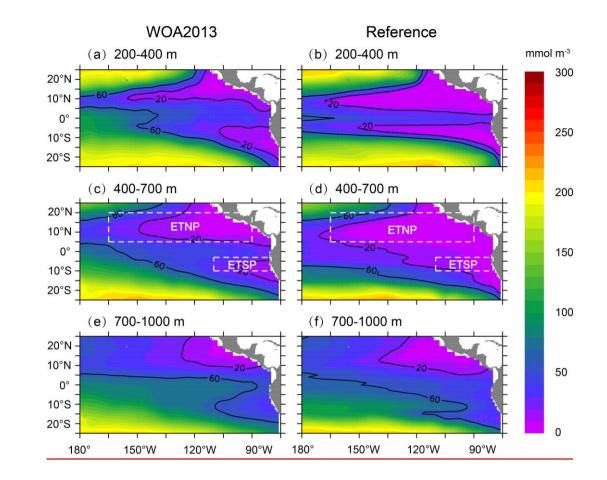


Figure 2. Comparisons of DO concentration between WOA2013 (left panel) and <u>the</u>reference run (right panel) <u>duringover</u> 1991-2010.
 White dash lines in (c) and (d) <u>denotesdenote</u> two boxes for ETNP (165°W-90°W, 5°N-20°N) and ETSP (110°W-80°W, 10°S-3°S).

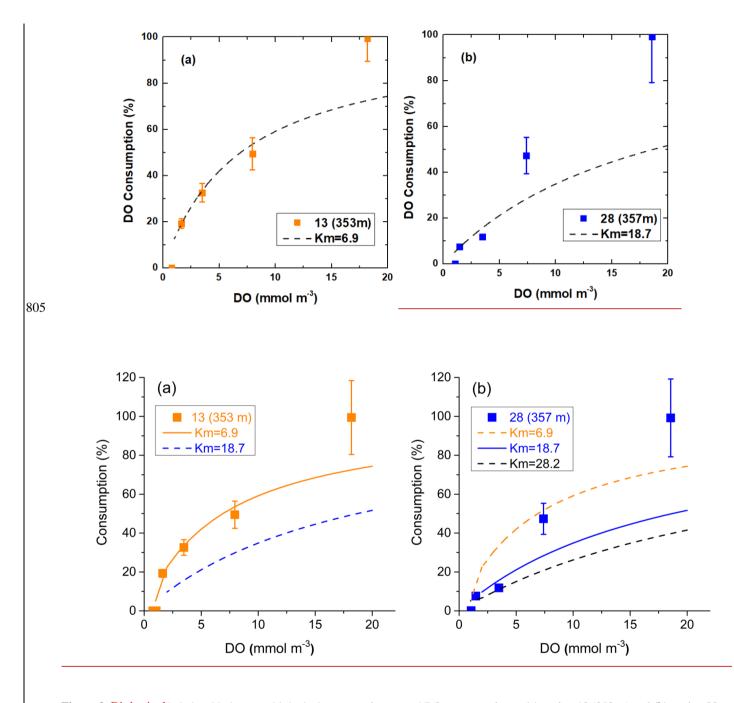


Figure 3. BiologicalRelationship between biological consumption vs.and DO concentration at (a) station 13 (353 m) and (b) station 28
 (357 m) in the Peruvian OMZ. Data are Km is the half saturation constant from the fitting curve derived from data of Kalvelage et al.
 (2015). For station 28, two Km values are derived using four (excluding the smallest value, blue curves) and five (black curve) data points,

respectively. Since the curve with Km=28.2 is too far away from the most data points, the value 18.7 is selected as the half saturation

constant.

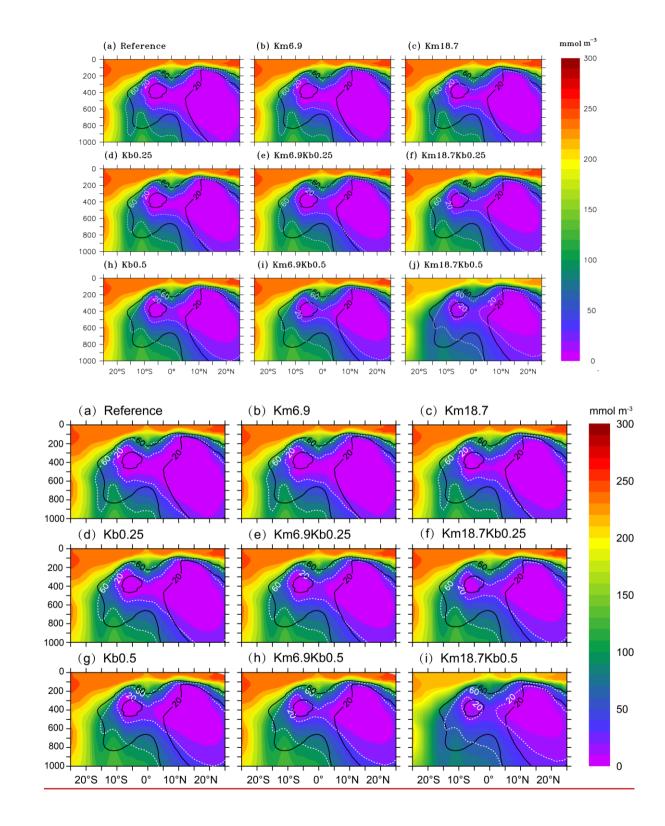
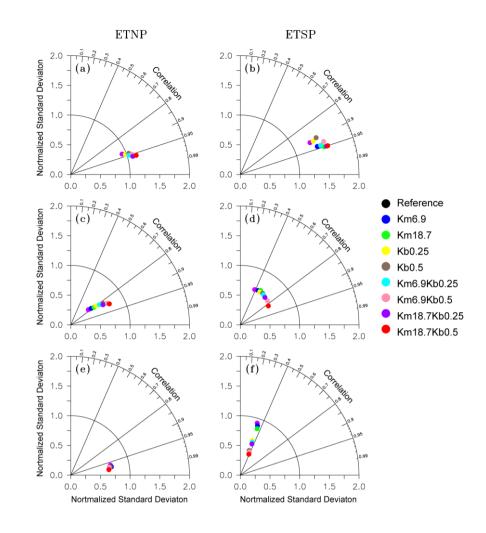


Figure 4. Vertical distribution of DO and asymmetric OMZsconcentration over 120°W-90°W from different model simulations-for: (a)
 the reference run, (b and c) with a reduced O:C utilization ratio, (d and h)g) with enhanced vertical mixing, and (e, f, h, and i, and j)) the combination of a reduced O:C utilization ratio and enhanced vertical mixing. Black lines denote contours of DO concentrations of 20 mmol m<sup>-3</sup> and 60 mmol m<sup>-3</sup> from WOA2013 data.



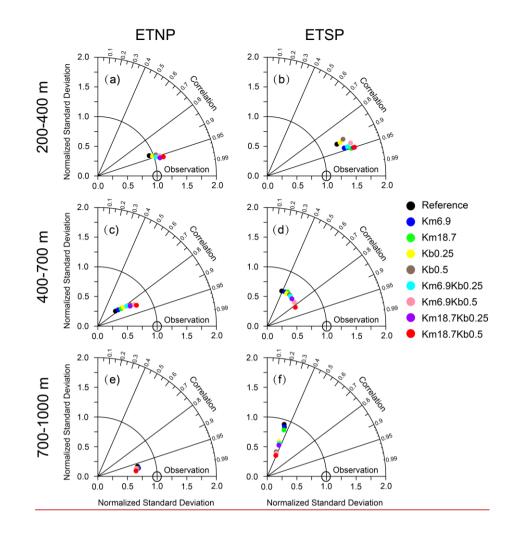
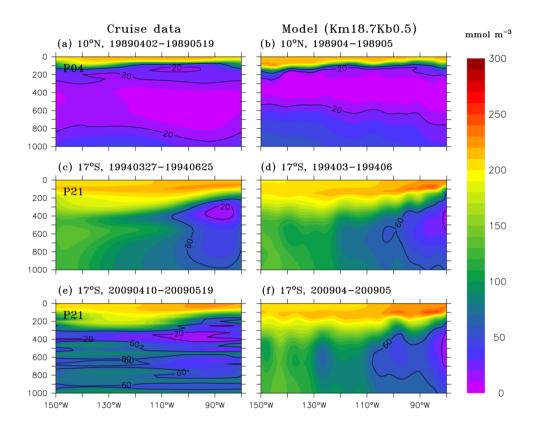


Figure 5. Taylor diagrams for the performance of simulated DO concentration (against WOA2013) from model simulations for ETNP (165°W-90°W, 5°N-20°N, left panel) and ETSP (110°W-80°W, 10°S-3°S, right panel) over (a and b) 200-400 m, (c and d) 400-700 m, and (e and f) 700-1000 m.



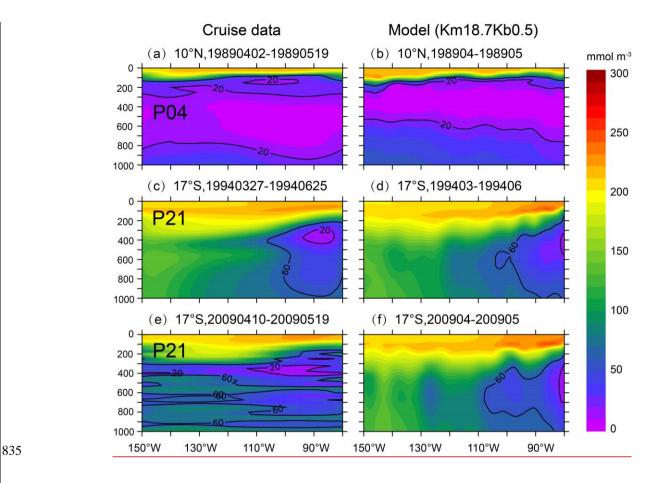
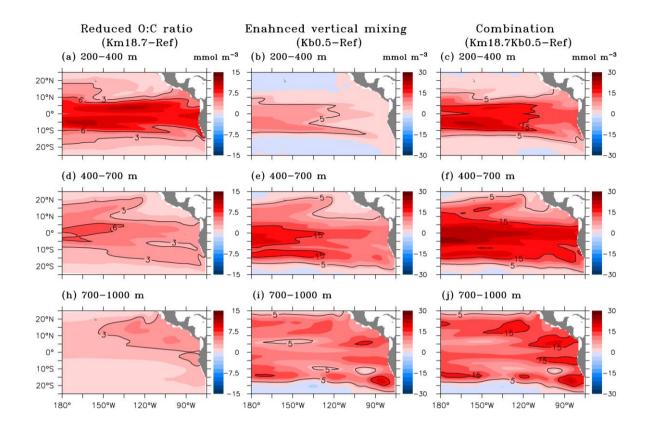
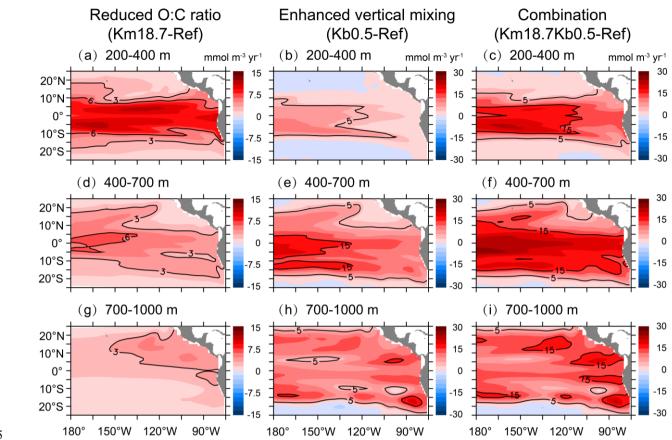


Figure 6. Distribution of DO from cruise data (left panel) and model simulation from the-Km18.7Kb0.5 (see text for explanation;-right panel). Observed DO along the P04 and P21 linesdata are from CCHDO (https://cchdo.ucsd.edu/), (https://cchdo.ucsd.edu/), along (a) P04 (10°N) during April 02 - May 19, 1989, (c) P21 (17°S) during March 27 - June 25, 1994, and (e) P21 (17°S) during April 10 - May 19, 2009.

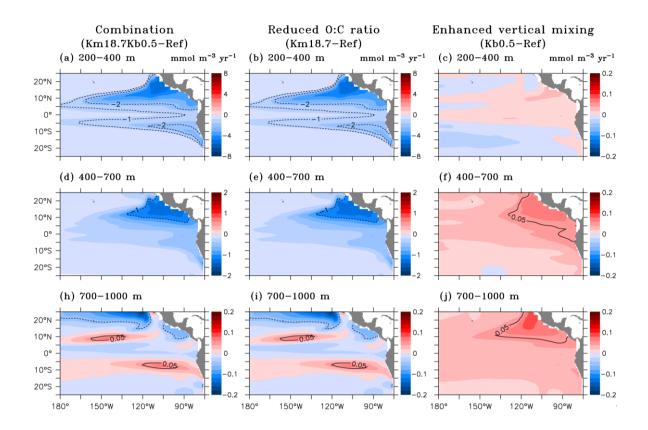


## **4 Model evaluation and discussions**



845

**Figure 7.** Changes of <u>in</u> DO concentration averaged over (**a**, **b** and **c**) 200-400 m, (**d**, **e** and **f**) 400-700 m, and ( $\underline{\mathbf{g}}$ ,  $\underline{\mathbf{h}}_{\underline{\mathbf{s}}}$  and  $\underline{\mathbf{i}}$ ) 700-1000 m due to <u>a</u> reduced O:C utilization ratio (left panel), enhanced vertical mixing (middle panel), and the combination of <u>a</u> reduced O:C utilization ratio and enhanced vertical mixing (right panel).



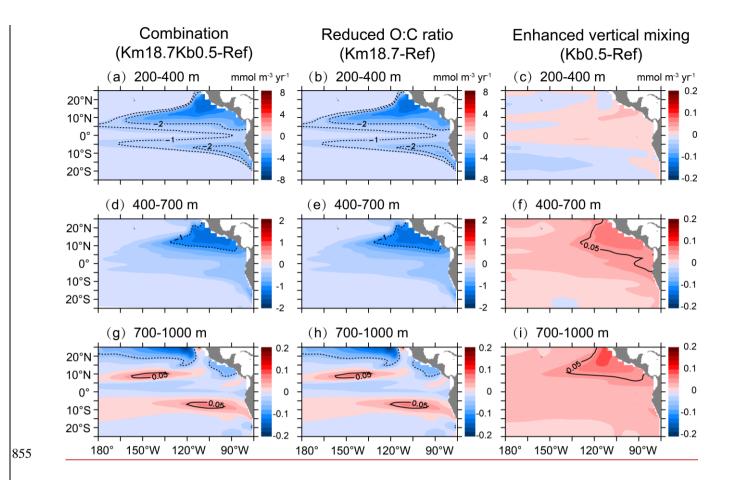
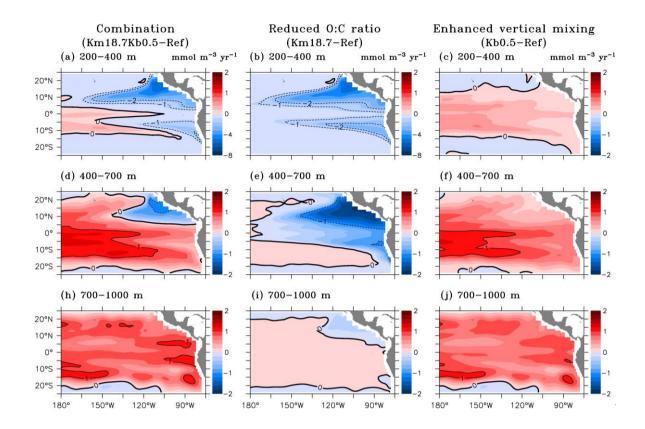


Figure 8. Changes in biological consumption over (a, b and c) 200-400 m, (d, e and f) 400-700 m, and ( $\underline{g}$ ,  $\underline{h}$ ,  $\underline{i}$  and  $\underline{ji}$ ) 700-1000 m due to the combination of <u>a</u> reduced O:C utilization ratio and enhanced vertical mixing (left panel), <u>a</u> reduced O:C utilization ratio (middle panel), and enhanced vertical mixing (right panel).



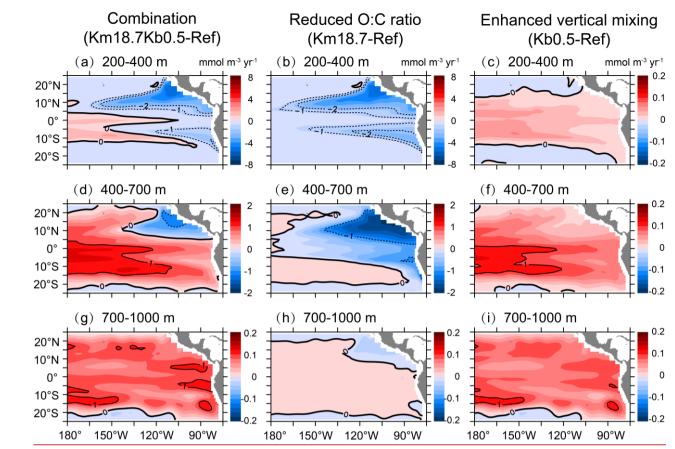
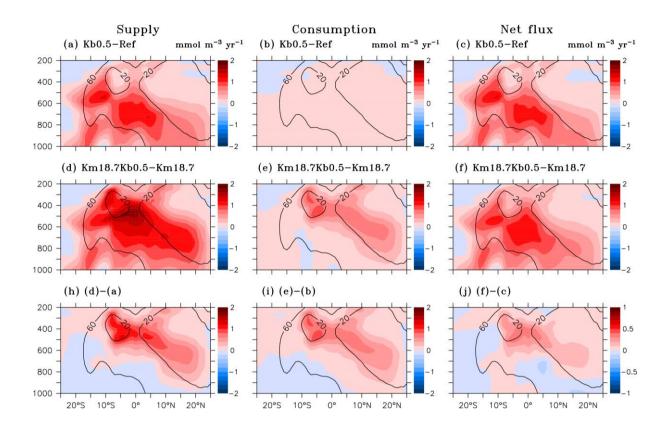
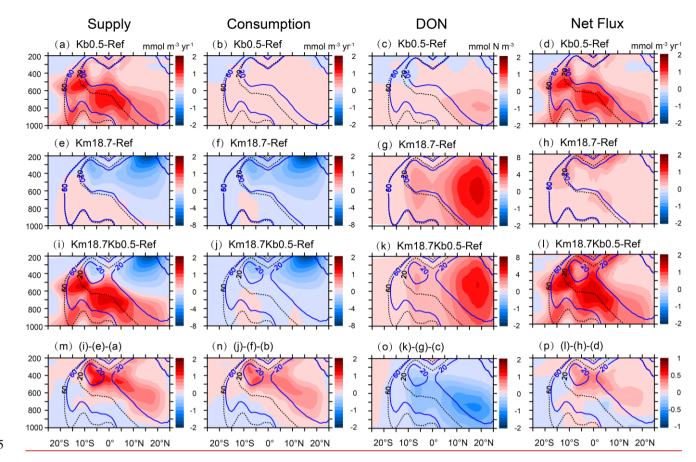
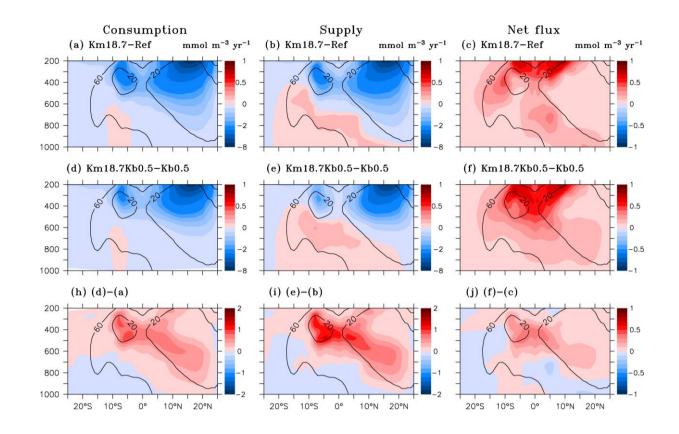


Figure 9. Changes in physical supply due to over (a, b and c) 200-400 m, (d, e and f) 400-700 m, and ( $\underline{g}$ ,  $\underline{h}$ ,  $\underline{i}$  and  $\underline{j}$ ) 700-1000 m due to the combination of  $\underline{a}$  reduced O:C utilization ratio and enhanced vertical mixing (left panel),  $\underline{a}$  reduced O:C utilization ratio (middle panel), and enhanced vertical mixing (right panel).





**Figure 10.** Changes in physical supply <u>(left panel)</u>, biological consumption <u>(middle panel)</u>, <u>DON concentration</u> and net flux (right <u>panel)</u><u>over 120°W-90°W</u> under enhanced vertical mixing with ((a, b, c, and d, top row), a reduced O:C utilization ratio (e, and f, <u>middle row)g</u>, and without (a, b, <u>h</u>, second row), combination effects (i, j, k, and c, top row) reduced O:C utilization ratio, <u>l</u>, <u>third row)</u>, and the differences between them (h, i, residuals or interactions (m, n, o, and jp, bottom row). Black and blue lines denote contours of DO concentrations of 20 mmol m<sup>-3</sup> and 60 mmol m<sup>-3</sup> from the reference run (dashed lines) and other simulations (solid lines).



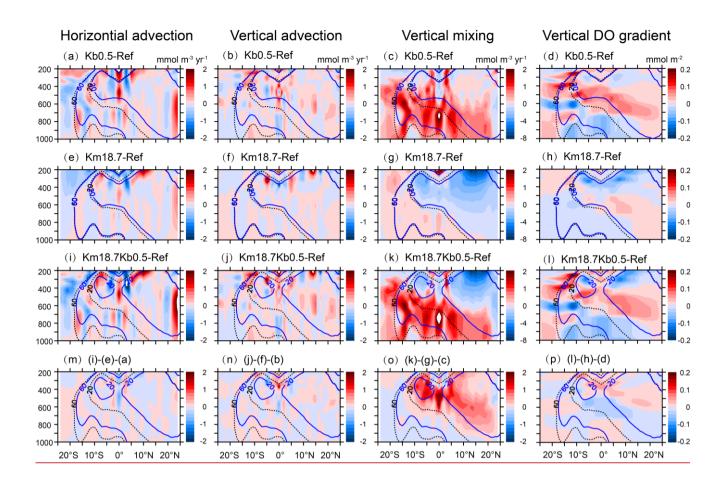
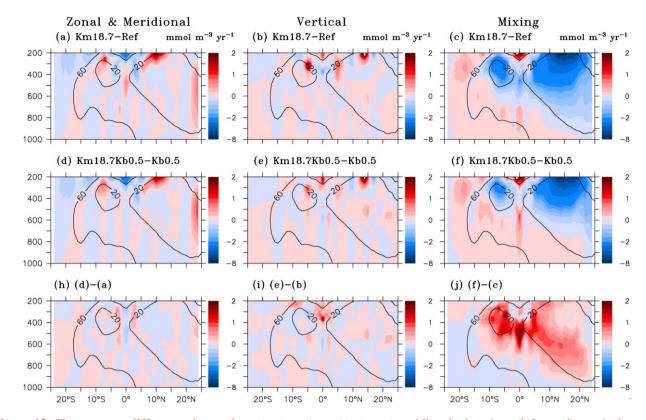


Figure 11. Changes in biological consumption (left panel ), physical supply (middle panel), and net flux (right panel)horizontal
 advection, vertical advection, vertical mixing and vertical DO gradient over 120°W-90°W under enhanced vertical mixing (a, b, c, and d, top row), a reduced O:C utilization ratio with (d, (e, and f, middle row) and without enhanced vertical mixing (a, b, and c, top row), and the differences between them (h, i, g, and j, bottomh, second row).



895 Figure 12. Changes), and differences in zonalcombination effects (i, j, k, and meridional advections (left panel), vertical advection (middle pane), and vertical mixing (right panel) under a reduced O:C utilization ratio with (d, e, and f, middle row) and without enhanced vertical mixing (a, b, and c, topl, third row), and the differences between them (h, i, and j, bottom row). residues or interactions (m, n, o, and p, bottom row). Black and blue lines denote contours of DO concentrations of 20 mmol m<sup>-3</sup> and 60 mmol m<sup>-3</sup> from the reference run (dashed lines) and other simulations (solid lines).