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Multi-timescale control of Southern Ocean diapycnal mixing over Atlantic tracer budgets

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Abstract

Oceanic cross-density (diapycnal) mixing helps sustain the ocean density stratification and its Meridional Overturning Circulation (MOC) and is key to global tracer distributions. The Southern Ocean (SO) is a key region where different overturning cells connect, allowing nutrient and carbon rich Indian and Pacific deep waters, and oxygen rich Atlantic deep waters to resurface. The SO is also rife with localized intense diapycnal mixing due to breaking of internal waves induced by the interaction of energetic eddies and currents with rough topography. SO diapycnal mixing is believed to be of secondary importance for the MOC. Here we show that changes to SO mixing can cause significant alterations to Atlantic biogeochemical tracer distributions over short and long timescales in an idealized model of the MOC. While such alterations are dominated by the direct impact of changes in diapycnal mixing on tracer fluxes on annual to decadal timescales, on centennial timescales they are dominated by the mixing-induced variations in the advective transport of the tracers by the Atlantic MOC. This work suggests that an accurate representation of spatio-temporally variable local and non-local mixing processes in the SO is essential for climate models' ability to i) simulate the biogeochemical cycles and air sea carbon fluxes on decadal timescales, ii) represent the indirect impact of mixing-induced changes to MOC on biogeochemical cycles on longer timescales.

Keywords: Southern Ocean, Turbulent Mixing, Carbon cycle, Tracer budgets

1 Introduction

In the ocean interior and away from boundaries, internal waves are mainly radiated by winds at the surface and by tidal and deep geostrophic flows interacting with rough bottom topography [1]. Breaking of internal waves induces turbulent mixing across density surfaces (i.e. diapycnally), which modifies water mass properties, maintains ocean stratification, and contributes to sustaining the deep branches of the global Meridional Overturning Circulation (MOC). Mixing is also key to redistribution of heat, carbon, biological nutrients, and other tracers globally and in depth [2–5]. Ocean circulation models do not resolve the processes responsible for diapycnal mixing and parameterize diapycnal mixing in terms of a turbulent effective diffusivity, \mathcal{K} [6–9].

1.1 Southern Ocean dynamics

Unbounded by continents, the SO is a crucial link between waters formed in different ocean basins, and inter basin exchanges of heat, carbon, nutrients and other tracers [10]. The lack of boundaries generates unique regional dynamics by allowing for the formation of the Antarctic Circumpolar Current system (ACC), an uninterrupted deep-reaching eastward flow driven by strong westerly winds. Within the ACC and in the upper ocean, the wind-driven Ekman transport is balanced by the resurfacing of deep waters. The Ekman transport is northward within the latitude band of the westerly winds, but southward towards Antarctica where easterly winds blow. In the deep SO, the zonal pressure gradient sustained by deep topography allows for northward transport of the abyssal waters that form and sink around Antarctica [11–13]. While the Ekman transport primarily induces isopycnal (i.e. along density surfaces) water transports, in the SO interior diapycnal mixing allows for further vertical mixing of waters and tracers [14–16]; see Fig. 1.

Within the SO, deep waters are upwelled adiabatically (i.e. along isopycnals) and are modified by surface buoyancy forcing upon turbulent entrainment into the mixed layer [17, 18]. Indian and Pacific deep waters predominantly upwell into a region of positive buoyancy forcing and form the northward-flowing Antarctic Intermediate Water (AAIW) and South Antarctic Mode Water (SAMW) [18, 19]. These waters eventually return to the SO as North Atlantic Deep Waters (NADW), completing the *upper* MOC cell (Red arrows in Fig. 1). Alternatively, waters can upwell into a region of negative buoyancy forcing in the SO, as happens for up to 75% of the NADW, forming the denser Antarctic Bottom Waters (AABW) [10, 18–22]. These abyssal waters sink and move northward along topographic ridges and canyons, and are then converted to deep waters through diapycnal mixing above the rough seafloor [12, 23] (Orange arrows in Fig. 1), forming the *lower* MOC cell (Blue arrows).

The SO is known to have high rates of diapycnal mixing from direct observational estimates, inverse bulk budgets, and numerical models [14, 16, 24–27]. This mixing is generated by strong ACC bottom currents interacting with

rough topography, causing internal wave breaking over abyssal ridges, mountains and rises [24, 28, 29] (Orange arrows in Fig. 1). Studies have reported strong variations in diapycnal mixing rates for different regions of the SO, ranging from $10^{-5}m^2s^{-1}$ to $10^{-3}m^2s^{-1}$.

1.2 Southern Ocean biogeochemistry

65% of interior ocean waters make first contact with the atmosphere in the SO [30, 31], with Pacific and Indian deep waters bringing waters rich in dissolved inorganic carbon (DIC) and nutrients to the surface [21]. Thus, the SO connects the vast reservoir of nutrients and carbon below the mixed layer with the surface [19] (DIC⁺ and N⁺ grey waters in Fig. 1). Vigorous vertical diapycnal mixing also enhances the supply of nutrients from the deep ocean to the surface [31]. Globally, the supply of surface nutrients is dominated by their upwelling in the SO [31, 32]. Patterns of SO circulation are therefore key to controlling the global biogeochemical cycles, the exchange of carbon dioxide (CO₂) between the atmosphere and the deep ocean, and the response of the ocean and atmosphere to climate change [33, 34]. The upwelling branch of the MOC, in particular, is key to controlling the global climate from both dynamical and biochemical perspectives [19].

The global distribution of conservative and non conservative tracers have been shown to be sensitive to ocean circulation and ventilation [35–37]. Enhanced mixing increases the deep ocean ventilation via the SO and reduces ocean carbon storage through the biological and solubility carbon pumps [21, 38]. Enhancement in SO diapycnal mixing is known to be able to alter tracer distributions through increased ventilation of intermediate waters [36]. Such changes are believed to be indirect since they are due to the interior diapycnal mixing altering the MOC over centennial to millennial timescales.

Theoretical studies have highlighted that diapycnal mixing rates in the deep SO are not to be of leading-order importance for regulating the global MOC, secondary to the role of winds, eddies, and surface buoyancy forcing. However, mixing in the ocean basins to the north of the SO play a primary role in regulating the ocean circulation, and close the MOC by upwelling the abyssal waters that form and sink at high latitudes [8, 8, 12, 39–41]. Limited work has been carried out to assess the sensitivity of biogeochemical tracer distributions to SO diapycnal mixing rates.

In this work, we examine the role of diapycnal mixing in the SO on Atlantic tracer budgets over varying timescales, and whether these changes are due to altered tracer diffusion or advection. We show that diapycnal mixing in the SO is crucial in setting the distribution of tracers on a range of timescales through two mechanisms. On long time scales, circulation and tracer advection changes dominate, especially when looking at changes to distribution on a global scale, whereas over short time scales, changes to tracer diffusion dominate, especially for changes within the SO only.

2 Model Description

This work uses the zonally averaged model of the Atlantic MOC (AMOC) by [41] (hereafter NV12), extended in [42] to include a simple biochemical model based on that introduced in [43]. The model, and similar ones, have been used extensively over the past decade to study various aspects of the role of the MOC in the climate system from both dynamical and biochemical aspects (e.g. [42, 44–47]). Thus, we simply provide a brief phenomenological description of the model here and refer the reader to the original works for more details.

2.1 The physical model

In the NV12 model, the Atlantic Ocean is split into three key areas: a SO circumpolar channel, a North Atlantic deep water formation zone, and the ocean basin that connects the two (consistent with the schematic in Fig. 1). While NV12 employed a constant \mathcal{K} value throughout the water column in all three zones, here we consider vertically-variable mixing and distinguish between the mixing in the basin from that in the SO. It is the impact of variations in the latter that we explore herein to study the sensitivity of tracer distributions to SO diapycnal mixing.

2.2 The Biochemical model

The simple biogeochemical component that was added to NV12 in [42] allows for representation of the coupled evolution of nutrients (N) and carbon (C), where N represents the ultimate limiting nutrient phosphate. Organic material is parameterized to sink downward, measured in units of phosphate, with the biological export timescale set to 360 days. A well-mixed atmospheric box is coupled to the ocean carbon cycle such that the partitioning of C between the atmosphere and the ocean is explicitly calculated in the model. The model conserves the total amount of C in the atmosphere and the ocean; if C is gained in the ocean, it must be lost in the atmosphere.

The biogeochemical tracers (C, N) are transported by advection and diffusion and have sources and sinks via biological activity (F_p) of uptake and remineralization. The concentration of C is able to change via surface fluxes driven by $p\text{CO}_2$ differences to account for air-sea gas transfer. More explicitly, the equations for C and N are

$$\delta_t N + J(\psi, N) = \delta_z(\mathcal{K}\delta_z N) + \delta_z F_p, \quad (1)$$

$$\delta_t C + J(\psi, C) = \delta_z(\mathcal{K}\delta_z C) + R\delta_z F_p + C_{flux}|_{z=0}, \quad (2)$$

$$C_{flux} = -(kw \times kH) \times (p\text{CO}_2a - p\text{CO}_2o) \times (1 - icecover), \quad (3)$$

where ψ is the meridional overturning streamfunction representing the AMOC (Fig.1, Fig.3a,b), kw is the air sea CO_2 exchange constant and kH is the fugacity of CO_2 . The second term on the LHS represents advection of N and C by the MOC, the first term on the RHS represents turbulence diapycnal

151 mixing, and the final term represents biological activity [43]. Biological activity
 152 in surface waters removes both nutrients and carbon at the ratio set by R , with
 153 carbon and nutrients both released by respiration as organic matter sinks.

154 2.3 Experiment Design

155 Our goal is to explore the sensitivity of the MOC and tracer budgets to SO
 156 mixing. Figure 2a shows the diapycnal mixing, \mathcal{K} , in the deep Atlantic region
 157 based on the tidal mixing estimates of [48] and the lee wave mixing esti-
 158 mates of [29] combined. While lee-wave-induced mixing is dominant in the
 159 SO due to strong eddy/current interaction with topography, tidal mixing is
 160 dominant in the basin to the north due to tidal flow interaction with the mid-
 161 Atlantic ridge. Panel b shows the global zonally-averaged diapycnal mixing in
 162 the SO from a combination of internal tides, lee waves and surface-enhanced,
 163 wind-induced mixing based on the observationally-constrained Southern Ocean
 164 Estate Estimate [49]

165 In all experiments (soon to be discussed) the mixing in the ocean basin will
 166 be fixed to the Atlantic basin averaged \mathcal{K}_B shown by the blue curve in figure
 167 2c. All surface forcings are also kept constant in all experiments, and are as
 168 prescribed in [29]. In the SO, we consider three paradigms: the Low mixing
 169 scenario of $\mathcal{K}_L = 10^{-5}$ m²/s, the High mixing scenario of $\mathcal{K}_H = 10^{-4}$ m²/s,
 170 and the Realistic mixing scenario motivated by figure 2b, referred to as \mathcal{K}_R .
 171 It should be noted that 10^{-5} and 10^{-4} m²/s are canonical values in physical
 172 oceanography, respectively representing the background mixing in the ocean
 173 and the mixing required for closure of the MOC [2, 46, 50, 51].

174 Since \mathcal{K}_B will be fixed in all experiments, hereafter we will use \mathcal{K}_R , \mathcal{K}_L ,
 175 and \mathcal{K}_H to name the experiments. While in some cases the same SO diffusivity
 176 will be applied to density, N , and C , in others we will apply a mixing profile
 177 to N and C that is different from that applied to density. This will be done to
 178 distinguish between the direct impact of mixing on tracers from the indirect
 179 effect on tracers because of the changes to the MOC induced by mixing of
 180 density. Thus, in naming of the experiments we will use superscripts 't' and
 181 ' ρ ' to refer to tracer and density mixing. For example, an experiment named
 182 [$\mathcal{K}_H^t, \mathcal{K}_L^\rho$] refers to one in which high SO mixing acts on tracers while low SO
 183 mixing acts on density whereas \mathcal{K}_L simply refers to an experiment in which
 184 low SO mixing is applied to both tracers and density.

185 The \mathcal{K}_L experiment, hereafter referred to as the 'control', is run until an
 186 equilibrated overturning circulation is achieved (4000 yrs) and then extended
 187 until the ocean biogeochemistry reaches an equilibrated state as well (another
 188 4000 yrs). C and N are initialized with uniform concentrations ($2.2e^{-3}$ mol N
 189 m^{-3} and 2.2 mol C m^{-3} respectively [42]). The atmospheric carbon concentra-
 190 tion is set to 278 part per million (ppm), representative of the pre-industrial
 191 level [43]. Starting from this equilibrated \mathcal{K}_L solution, we run a host of configu-
 192 rations varying the SO density and tracer diffusivities. Each run is 4000yr long
 193 so that a new equilibrium is achieved. In this work we examine changes to both
 194 transient model solutions and the equilibrated changes for each experiment.

2.4 Model Verification

Figure 3 compares the model solution for the control case to the circulation inferred from a climatologically-constrained ocean state estimate (ECCO [52]). In the control run (\mathcal{K}_L), the upper cell extends to $\sim 2000\text{m}$ deep with a maximum strength of 17 Sv while the lower cell transport is about 4 Sv (Fig.3b), both well within the range of inverse models [10, 53] and ocean state estimates [9, 52]. Panel *c* compares the model stratification in the Atlantic basin (for experiments $\mathcal{K}_L, \mathcal{K}_H, \mathcal{K}_R$) with the World Ocean Circulation Experiment climatology (WOCE; [54]), showing a slight under-prediction by the model in the deep ocean. Since the basin diffusivity is fixed at \mathcal{K}_B for all experiments, unsurprisingly the model stratifications in the basin north of the SO is similar in all three experiments. This implies the secondary influence of the SO mixing on the basin stratification, consistent with theoretical arguments discussed earlier [41].

Altered SO mixing causes up to a 30% change in the strength of circulation (3d,e) after 4000 years. Increasing SO mixing to \mathcal{K}_H and \mathcal{K}_R increases AMOC strength by up to ~ 3 Sv in the upper branch and by ~ 2 Sv in the deep branch. The maximum transport is increased from 16.9 Sv in \mathcal{K}_L , to 17.8 Sv in \mathcal{K}_H , and to 18.8 in \mathcal{K}_R . Previous studies have similarly found increased strength of convection with higher SO vertical mixing [21]. The upper cell is also deepened with enhanced mixing, as in [41, 55] when mixing was increased globally, as the increased downward mixing of buoyant surface waters in the SO is able to deepen the pycnocline, altering the structure of the AMOC. As we will show, this change is effective on timescales of equilibration of the circulation (centennial to millennial). We will also show that the direct impact of mixing on tracers can be more significant and effective on shorter timescales.

The distribution of C and N produced by the model are also qualitatively similar to observations and reproduce many of the key features (Fig.4): the surface waters are nutrient depleted due to phytoplankton productivity. As waters circulate and organic matter sinks, the N and C concentrations in old deep waters get enhanced (Fig.1). The short residence time of these waters in the surface SO means that biota are unable to utilize all of the nutrients [43]. Therefore, AABWs and AAIWs have high (preformed) N content. The concentration of nutrients in AAIWs are rapidly depleted as waters travel north due to productivity. Regions with high (low) N levels also have high (low) C concentrations, as photosynthesis takes in both, and remineralisation releases both.

3 Results: sensitivity of C & N distribution to Southern Ocean mixing

3.1 Multiscale temporal changes

Figure 5 shows the N distribution anomaly between various mixing scenarios and the control simulation (\mathcal{K}_L), over the first 100 years after mixing in the SO

is perturbed for C & N and/or density. The first row shows the total change when both \mathcal{K}^t (which acts on N and C) and \mathcal{K}^ρ get enhanced. The net change in the first row can be interpreted as due to two contributions: (I) the indirect impact because of the change in the MOC caused by alteration of the density field by higher \mathcal{K}^ρ , and (II) the direct impact due to action of enhanced \mathcal{K}^t directly on the C & N gradient.

The indirect impact is isolated in the second row where \mathcal{K}^t is unchanged from the control but \mathcal{K}^ρ is enhanced. Not surprisingly, the anomaly is large north of the SO channel, reflecting the change in the Atlantic MOC due to the modification of the Atlantic density stratification thanks to enhanced SO density mixing. The direct impact is isolated in the third row where \mathcal{K}^ρ is unchanged from the control, but \mathcal{K}^t is enhanced. This anomaly is expectedly SO-focused and evolves much faster than the former signal in second row.

The last row bears similarity to the first row as it too captures the impact of enhanced mixing (on both C & N and density), but for a more realistic mixing profile than the simple $\mathcal{K}_H = 10^{-5}$ m²/s. The SO change is more dramatic for the realistic mixing profile, since it includes the large surface turbulent mixing induced by the SO storm tracks (Fig. 2c). The higher \mathcal{K}^t rates in the first, third, and fourth rows act to disperse the C & N more rapidly from tC & N -rich regions such as the AAIW and lower cell waters. C & N are lost from SO upper AAIW waters, and mixed into the C & N depleted regions above and below. Such changes are the greatest in \mathcal{K}_R , showing the importance of the enhanced surface water mixing rates.

To further quantify the changes to SO and basin N inventories by mixing perturbations, we calculate the absolute changes between various experiments and the control run (\mathcal{K}_L). The results are shown in Fig. 6 and are broken down into different basin as well as into upper ocean versus full depth. Since the results based on \mathcal{K}_R are similar to those experiments with \mathcal{K}_H , we don't include the former in this figure to avoid further overcrowding it.

The mean N concentrations increase over the 4000 years of experiments in all regions of the Atlantic. The changes to N concentrations in the surface waters of \mathcal{K}_H show the greatest changes at all times, changing by a mean of 20% once a new steady state is reached (Fig. 6 solid red line). The SO shows initially higher changes in N distributions than the basin. By 500 years, however, changes to N distributions in the basin and the SO are comparable at around 13% by the time a new steady state is reached at 4000 years. This emphasizes the global impact of altered SO diapycnal mixing on N distributions.

Over the first 5 years, changes to N distributions are only felt in the SO and SO & basin surface waters in experiments where \mathcal{K}^t is enhanced. It takes over 100 years for the enhanced \mathcal{K}^t in the SO to have an impact on the distribution of N in the Atlantic basin, and even on millennial time scales, this change is less than 2%. By 5 years, enhanced \mathcal{K}^ρ alters N distributions in just the SO, but only by 1%, while changes to \mathcal{K}^t alter the N distributions by 4% in the same region. It takes 20 years for alterations to the MOC to alter the N distributions in the basin. Changes to N distributions in the basin are

282 always greater for enhanced \mathcal{K}^ρ as compared to cases with \mathcal{K}^T . In the SO, it
 283 takes at least 200 years for the indirect effect of altered MOC to alter tracer
 284 distributions more significantly than just increasing \mathcal{K}^t .

285 We also carried out a similar analysis for passive tracers, released with
 286 initial distributions similar to that of the N distribution in the \mathcal{K}_L experiment.
 287 The results, which are not discussed here, looked qualitatively similar to what
 288 was described above for N , albeit with lower magnitudes in the concentration
 289 anomalies due to the lack of biological processes.

290 **3.2 Long-term equilibrated changes**

291 While mixing is spatio-temporally variable, and so a steady-state solution does
 292 not necessarily describe the real world, it is insightful to consider the long-term
 293 equilibrated solutions of the perturbation experiments discussed above to gain
 294 an understanding of order of magnitude importance of the changes. To this
 295 end, in Fig. 7 we plot the changes in the steady-state C and N distributions
 296 between the various experiments and the control experiment. In all cases, the
 297 C and N concentrations are increased in surface waters, as found previously
 298 with enhanced SO deep water upwelling [21]. Changes to N are overall more
 299 prominent than those to C , as shown by the percentage change contours of
 300 fig.7. The greatest changes for both occur in the \mathcal{K}_R experiment, highlighting
 301 the conservative nature of our choice of high diffusivity in \mathcal{K}_H .

302 Consistent with the transient solutions, in experiments with enhanced mix-
 303 ing only acting on ρ (panels c,g), the signal extends beyond the SO channel
 304 since it comes from changes to the whole AMOC. As a result, on the long-term
 305 (millennial) equilibration timescale, the indirect impact of mixing on tracers
 306 due to changes it induces in the MOC can overwhelm the effect of direct impact
 307 of mixing on tracers. However, as was discussed in relation to Figs 5, 6, on
 308 inter-annual to decadal timescales, the direct impact is dominant and perhaps
 309 more readily relevant to the short timescale climate change.

310 Again consistent with the transient solutions, in the experiments with
 311 enhanced mixing only applied to N & C , the signal is strongest in the SO due
 312 to the transfer of tracers from the C & N rich lower cell to the upper cell of the
 313 MOC (panels b,f). The subsequent increase in the surface N concentration is
 314 significant for the uptake of C by biological productivity, and for the sinking
 315 of organic matter containing C and N as part of the biological carbon pump.
 316 An increase in surface nutrients concentrations with enhanced SO upwelling
 317 has been observed previously [21].

318 The climatic implications of the changes to N and C budgets may be encapsu-
 319 lated through the net change in atmospheric pCO_2 concentrations between
 320 the various experiments and the control run in which weak mixing acts on N ,
 321 C , and ρ . This is summarized in table 1. The direct impact of enhanced mixing
 322 on tracers, as in $[\mathcal{K}_H^t, \mathcal{K}_L^\rho]$, cause the redistribution of C to the upper surface
 323 waters. This is highly important for atmospheric C concentrations. These sur-
 324 face waters allow C to be exchanged between the atmosphere and the ocean.
 325 Therefore, enhanced \mathcal{K}^t results in only a very slight increase in C content of

326 the upper cell, but an increase in atmospheric C reservoir of 12.3ppm (Table1).
327 Although increased \mathcal{K}^ρ caused a stronger redistribution of C to the upper cell,
328 this C was not directed to the very surface waters of the upper cell. There-
329 fore the redistribution was not as important for ocean atmosphere fluxes of
330 C , but caused more drastically observable changes to the C content of the
331 upper cell. Enhanced \mathcal{K}^t and \mathcal{K}^ρ both cause a steady state atmospheric CO_2
332 concentration greater than in \mathcal{K}_L (Table1). Increased model mixing is known
333 to raise predicted atmospheric CO_2 concentrations, and to make partitioning
334 CO_2 into the deep ocean harder [4]

335 4 Discussion

336 This work highlights the significant sensitivity of Atlantic nutrient and carbon
337 distributions to enhanced vertical mixing with the Southern Ocean, demon-
338 strating the significance of SO diapycnal mixing on biogeochemical tracer
339 budgets. By distinguishing between the impact of SO mixing on tracers
340 through direct tracer mixing and the indirect impact through changing the
341 Atlantic stratification and circulation, we were able to show that both effects
342 are significant, albeit the former acting on much shorter timescales than the
343 latter. The direct impact due to altered tracer diapycnal fluxes manifests on
344 inter-annual to decadal timescales. On longer timescales, the tracer concentra-
345 tions are affected by the indirect impact of mixing through the change in the
346 Atlantic MOC. The change to the Atlantic MOC by SO mixing is due to the
347 importance of the SO density stratification to setting the Atlantic deep ocean
348 stratification.

349 Changes to nutrient concentrations within the SO occur on timescales of
350 inter-annual to decadal, while the surface water nutrient concentrations begin
351 to be modified by the change in the MOC beyond 50 years. This timescale cor-
352 responds to transport and utilisation of nutrients by the northward SAMW [31]
353 to be utilized for downstream productivity, as the resurfacing of deep nutrient
354 rich waters at around 30 °S ranges from multi-decadal to a centennial. [13].
355 Given that between 30°S and 30°N upper ocean nutrient supply is dominated
356 by waters originating in the SO, [31], the global nutrient and carbon budgets
357 are therefore hyper-sensitive to SO mixing. Our results agree with the sugges-
358 tion of [36] that flux divergences are likely to be much smaller in comparison
359 to changes to the advective fluxes of tracers, and that increasing the diapyc-
360 nal mixing values in just the SO causes minimal changes to vertical diapycnal
361 nutrient fluxes, with changes to advection and convection much larger.

362 The steady state atmospheric $p\text{CO}_2$ concentration was increased by 25ppm
363 when a 'realistic' mixing profile (admittedly within our idealized framework) is
364 considered relative to \mathcal{K}_L . Given that glacial inter glacial cycles in atmospheric
365 $p\text{CO}_2$ were around 80 - 100ppm [56], and that present day to pre-industrial
366 atmospheric carbon changes are around 130ppm this is a highly significant
367 result for climate modeling. This demonstrates the importance of correctly
368 representing the strongly surface enhanced diapycnal mixing in surface waters

369 of the SO for climate modelling. It also shows the importance of the choice of
370 diapycnal mixing parameterisation in the SO in more advanced coupled climate
371 models. Its significance can be further realized by noting that our analyses
372 only included the Atlantic basin.

373 Fig. 8 summarizes our key messages by showing a zoomed in view of the
374 zonally-averaged SO (from the model), and comparing the SO-branch of the
375 upper and lower MOC between the low mixing control run and the high mixing
376 case in which a realistic SO-averaged diffusivity was applied. The background
377 colour, from the control run, shows the richness of nutrients in the AABW
378 within the SO. The arrows at mid-depths represent the interior mixing induced
379 by breaking of waves within the vigorous wave field within the SO which
380 result in diapycnal diffusion of nutrients across density surfaces. It is the direct
381 impact of enhancement to such mixing, which increases the rate of diapycnal
382 upwelling of nutrients from high concentration AABW into the nutrient poor
383 water masses that introduces the largest changes in Atlantic C and N budgets,
384 as well as in pCO_2 levels on decadal to centennial timescales. These water masses
385 experience wind driven along isopycnal upwelling at the boundary of the upper
386 and lower cells within the ACC (see Fig. 1). On longer timescales, alterations
387 to nutrient distributions arise from the change in Atlantic MOC structure from
388 enhanced SO mixing. Such mixing changes the density stratification within the
389 SO manifesting in the form of change to density layer slopes (Fig. 8 coloured
390 contours). Given the global connectivity of the ocean basins through the SO
391 (thanks to the ACC system), any change in SO isopycnal slopes translates to
392 a change to the global deep stratification, thereby to alteration of the global
393 MOC.

394 Given that mixing is induced by turbulent events that are highly intermit-
395 tent in time and space, this work highlights the inevitable need for realistic
396 spatio-temporal representation of small scale SO mixing in climate models. In
397 a recent complementary work, we reported a hyper-sensitivity of SO air-sea
398 carbon fluxes to moderate changes in upper ocean diapycnal mixing in a real-
399 istic ocean state estimate[57]. While that work was focused on the SO alone,
400 as opposed to our Atlantic focus here, it shows that our idealized framework
401 likely significantly underestimates the sensitivity of the global budgets to the
402 SO mixing. Even with such underprediction, the changes reported herein are
403 significant. While some parameterization of internal wave generation within
404 the SO have been implemented in models previously (albeit with many caveats
405 of their own), to our knowledge, no explicit parameterization of small scale
406 mixing within the SO has been attempted to date.

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663 **Supplementary information.**

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666 **Declarations**

667 The authors have no conflict of interest to report.

Experiment	change in atmospheric pCO ₂ from K _L (ppm)
K _H	11.2
[K _L ^t , K _L ^b]	12.3
[K _L ^t , K _H ^p]	3.9
K _R	24.6

Table 1 Change in steady state atmospheric pCO₂ from experiment K_L

5 Tables

6 Figures

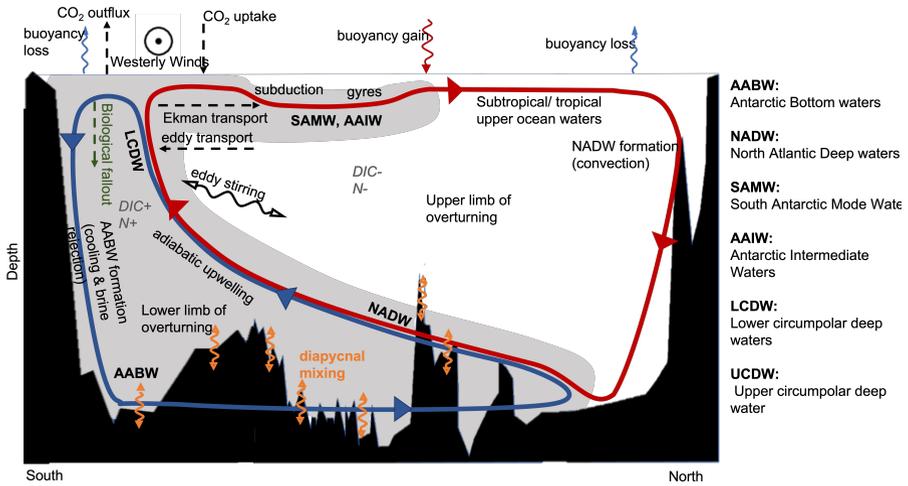


Fig. 1 A schematic of the Atlantic ocean circulation with an emphasis on the SO dynamics, watermasses, and tracers. Adapted from [58] and [10]

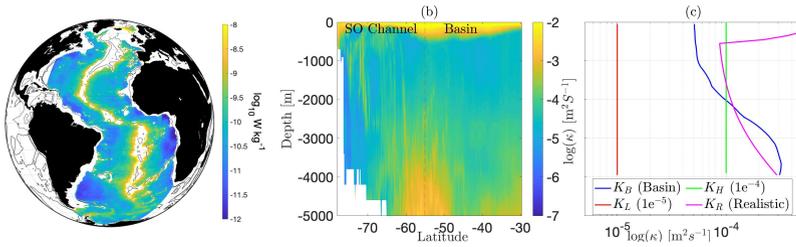


Fig. 2 a) Turbulent diffusivity in the deep Atlantic basin (3000m) based on the tidal mixing estimate of [48] and lee-wave mixing estimate of [29]. b) Zonally-averaged diapycnal mixing in the SO from a combination of the interior mixing shown in panel a and wind-induced surface mixing based on the Southern Ocean Estate Estimate (SOSE; [49]). c) Basin-averaged diffusivities employed in the model: \mathcal{K}_B is prescribed in the basin (and held the same in all experiments) whereas \mathcal{K}_R , \mathcal{K}_L , and \mathcal{K}_H are employed in the SO for various perturbations. \mathcal{K}_B is averaged over the basin (32S to 48N) based on the estimate in panel a. \mathcal{K}_R is the SO-averaged diffusivity based on the estimate in panel b. \mathcal{K}_L and \mathcal{K}_H are the two canonical values representing the lower and upper bounds of ocean interior mixing.

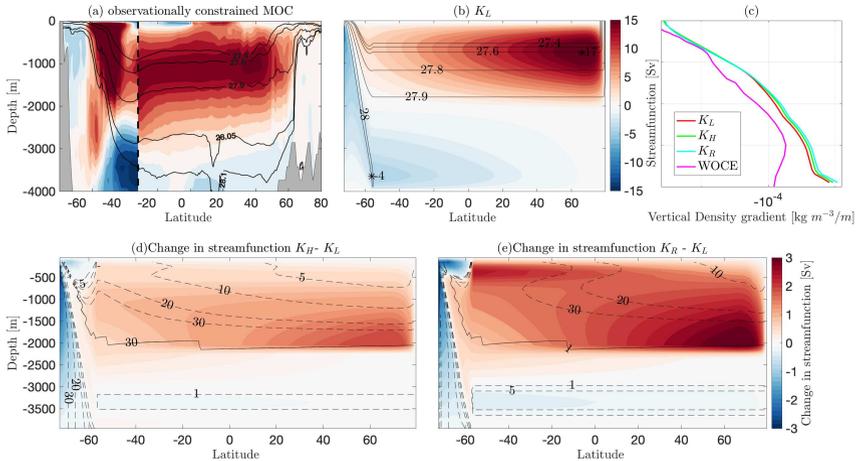


Fig. 3 a) Zonally-averaged Atlantic MOC streamfunction (ψ in Eqs. 1.2) from a climatologically-constrained ocean estimate (ECCO; [52]). Each $\text{Sv} = 10^6 \text{ m}^3/\text{s}$. The streamfunction is calculated in the Atlantic basin up to 32°S and globally between 70°S – 32°S . b) The overturning streamfunction for the control case (\mathcal{K}_L) with density contours overlain (black lines). c) Vertical gradient of neutral density [59] in the model basin for \mathcal{K}_L (red), \mathcal{K}_H (green), and \mathcal{K}_R (blue), compared to density gradient from World Ocean Circulation Experiment (WOCE [54]) climatology averaged between 48°N to 32°S in the Atlantic basin (pink). d) Change in overturning circulation between \mathcal{K}_H and \mathcal{K}_L experiments. Dashed lines indicate percentage change in overturning streamfunction from \mathcal{K}_L . e) Similar to d but for the difference between \mathcal{K}_R and \mathcal{K}_L .

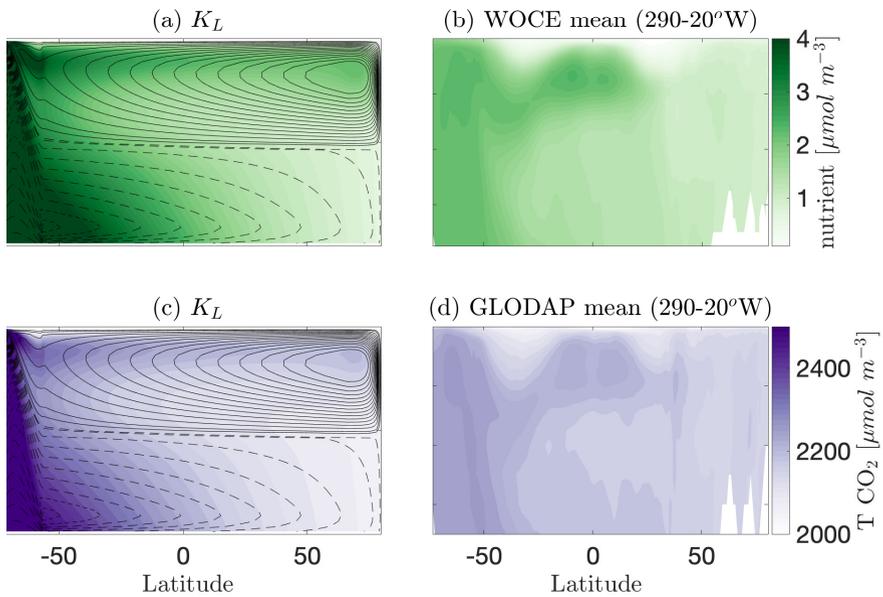


Fig. 4 Concentration of N from experiment \mathcal{K}_L , compared to the climatological zonally-integrated phosphate in the Atlantic basin from WOCE[54] in (b). Concentration of C from experiment \mathcal{K}_L in (c) is also compared to the climatological zonally-integrated C from the Global Ocean Data Analysis Project (GLODAP[60]) in (d).

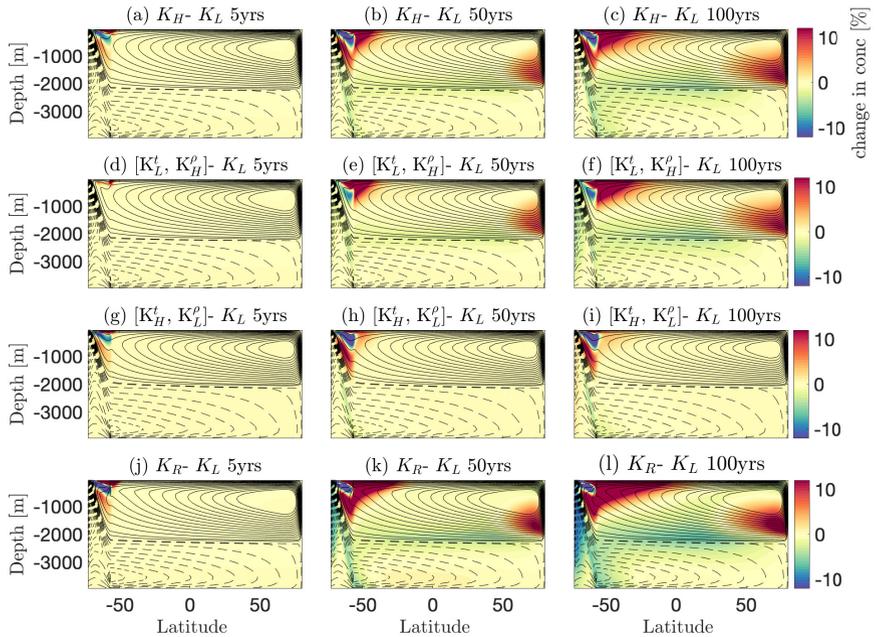


Fig. 5 Change in N concentration at $t=5, 50$ and 100 years. First Row: change between \mathcal{K}_H and \mathcal{K}_L experiments. Second Row: between $[\mathcal{K}_L^t, \mathcal{K}_H^\rho]$ and \mathcal{K}_L experiments. Third row: between $[\mathcal{K}_L^t, \mathcal{K}_H^\rho]$ and \mathcal{K}_L experiments. Fourth row: between \mathcal{K}_R and \mathcal{K}_L experiments. Red/blue implies increased/decreased tracer concentration. Black overlain lines indicate zonally averaged residual overturning circulation for each experiment. Note that the N concentration for the \mathcal{K}_L experiment is in steady-state, as was shown in figure 4a.

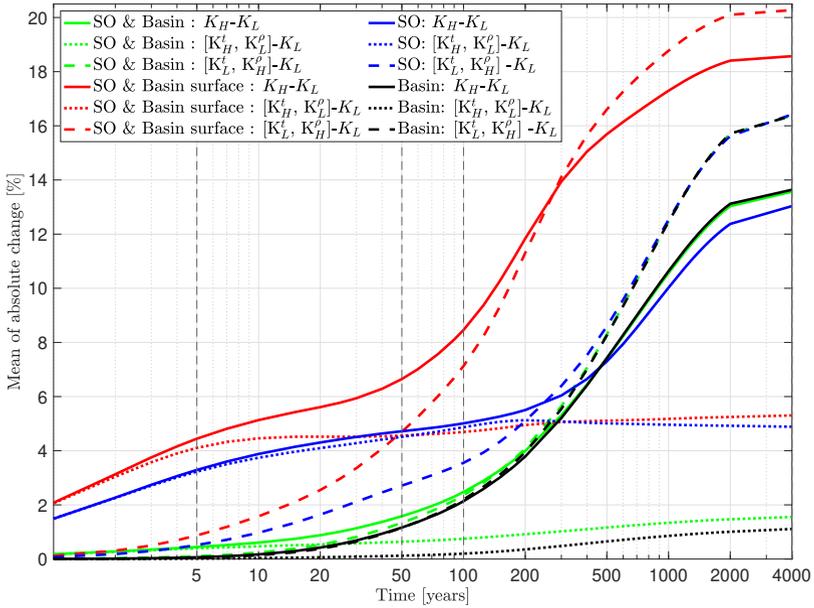


Fig. 6 Mean of absolute change in integrated N concentration for various inventories: the entire ocean (SO+basin) in green, the surface ocean (top 200m of SO+basin) in red, only the basin (full depth) in black, and only the SO (full depth) in blue. Plain lines denote the difference between \mathcal{K}_H and \mathcal{K}_L experiments, dotted lines represent differences between $[\mathcal{K}_H^t, \mathcal{K}_L^\rho]$ and \mathcal{K}_L experiments, and dashed lines denote differences between $[\mathcal{K}_L^t, \mathcal{K}_H^\rho]$ and \mathcal{K}_L experiments. Vertical dashed lines indicate the time slices considered in Fig. 5.

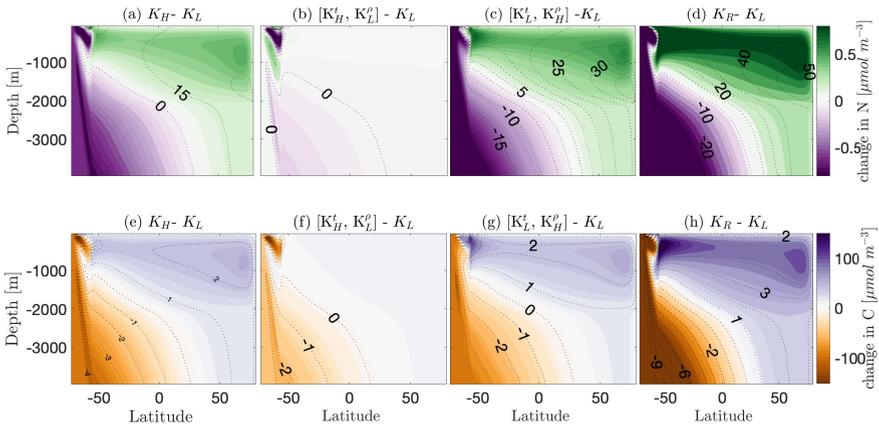


Fig. 7 Equilibrated changes to N distributions. Top row: change in N concentration between \mathcal{K}_L and a) \mathcal{K}_H , b) $\mathcal{K}_H^t, \mathcal{K}_L^\rho$, c) $\mathcal{K}_L^t, \mathcal{K}_H^\rho$, and d) \mathcal{K}_R . Green/purple denotes regions of increased/decreased N concentration. Black-labelled overlain lines indicate percentage changes from \mathcal{K}_L . Bottom row: same as above but for C , with purple/orange denoting regions of increased/decreased C concentration.

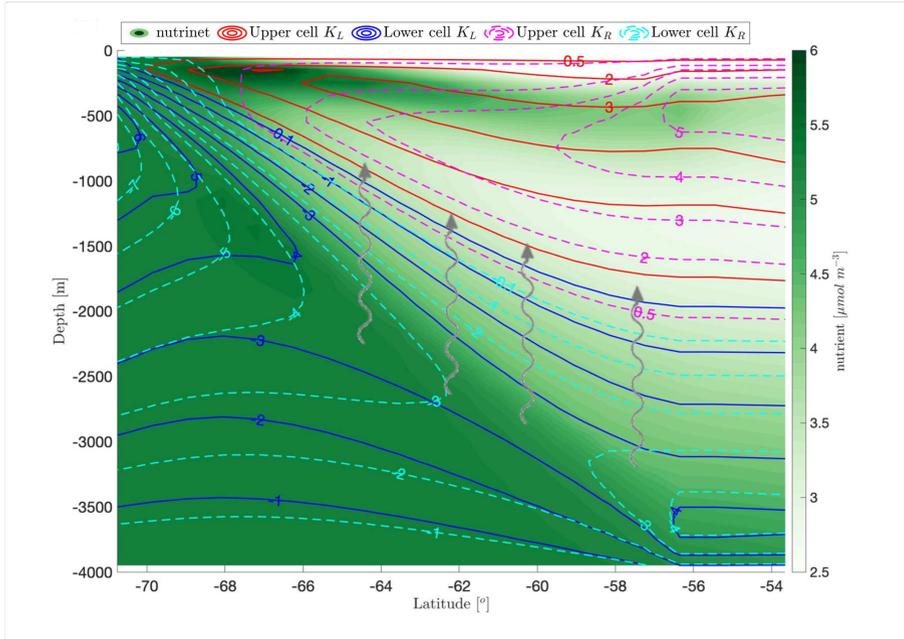


Fig. 8 Tracer distributions in the SO are altered via two processes: firstly, through the direct impact of mixing (grey arrows) on tracer gradients, and secondly by changing the SO density stratification and thereby the global MOC. Red and Blue contours show the overturning stream functions for the upper and lower MOC cells for the control \mathcal{K}_L experiment (solid lines) vs the \mathcal{K}_R experiment (dashed lines) in Sv. The background green colour represents the nutrient concentration from the equilibrated control experiment, \mathcal{K}_L .