

Abstract

Sustaining biological export over the open ocean requires a physical supply of nutrients to the mixed layer and thermocline. The relative importance of diapycnal mixing, diapycnal advection and isopycnal stirring by mesoscale eddies in providing this nutrient supply is explored using a field campaign in oligotrophic waters in the subtropical North Atlantic, consisting of transects over and off the mid-Atlantic ridge. Eddy stirring rates are estimated from the excess temperature variance dissipation relative to the turbulent kinetic energy dissipation, and using eddy statistics from satellite observations combined with 9-month-long mooring data. The vertical nutrient fluxes by diapycnal mixing, diapycnal advection and isopycnal mesoscale eddy stirring are assessed using nitrate measurements from observations or a climatology. Diapycnal mixing and advection provide a nutrient supply within the euphotic zone, but a loss of nutrients within the upper thermocline. Eddy stirring augments, and is comparable to, the diapycnal transfer of nutrients within the summertime upper thermocline, while also acting to replenish nutrients within the deeper parts of the thermocline. The eddy supply of nitrate is relatively small in the centre of the subtropical gyre, reaching up to $0.06 \text{ mol N m}^{-2}\text{yr}^{-1}$, but is likely to be enhanced on the flanks of the gyre due to larger isopycnal slopes and lateral nitrate gradients. The nutrient supply to the euphotic zone is achieved via a multi-stage mechanism: a diapycnal transfer of nutrients by small-scale turbulence to the euphotic zone, and an isopycnal stirring of nutrients by mesoscale eddies replenishing nutrients in the upper thermocline.

Plain Language Summary

Phytoplankton growth requires a supply of nutrients to the base of the euphotic zone, which is usually provided by a combination of vertical mixing or vertical upwelling of nutrients. However, in the oligotrophic waters of the central North Atlantic, it is unclear how the vertical supply of nutrients is sustained. Here we use field data to explore the roles of mixing across density surfaces, advection across density surfaces and mesoscale eddy stirring along density surfaces in supplying nutrients to some of the most nutrient-depleted surface waters in the central North Atlantic. Diapycnal mixing and advection are found to be important in supplying nutrients to the euphotic zone during summer, but at the expense of eroding the nutrients in the upper thermocline. In contrast, mesoscale eddy stirring augments the diapycnal supply of nutrients to the euphotic zone, and replenishes nutrients in the upper thermocline.

1 Introduction

Biological export of organic matter in the open ocean is usually viewed as being sustained by a vertical supply of nutrients to the euphotic zone from the upper thermocline. This vertical transfer of nutrients is easily provided in regions of climatological wind-driven upwelling, but is more difficult to achieve in downwelling areas, where a combination of diapycnal transfer and time-dependent upwelling needs to take place.

Maintaining biological export is particularly challenging in the oligotrophic waters of the North Atlantic subtropical gyre, where tracer-based estimates of export production range from 0.42 to $0.65 \text{ mol N}^{-2}\text{m}^{-2}\text{y}^{-1}$ in the Sargasso Sea (Jenkins & Goldman, 1985; Jenkins, 1988; Jenkins & Wallace, 1992; Stanley et al., 2015). These estimates of export production are at least twice as large as those based on conventional estimates of nitrate supply, combining together the contributions from atmospheric deposition (Knap et al., 1986), convective entrainment (Michaels et al., 1994), diapycnal mixing (Lewis et al., 1986; Dietze et al., 2004) and horizontal Ekman transfers (Williams & Follows, 1998).

Time-varying circulations involving mesoscale eddies and fronts have been invoked to partly explain the mismatch between estimates of nutrient supply and export (McGillicuddy Jr

67 & Robinson, 1997; McGillicuddy et al., 1998; Lévy et al., 2001). Such mesoscale circulations may affect both vertical and horizontal transfers of nutrients (Lee & Williams, 68 2000; Williams & Follows, 2003; Resplandy et al., 2011; Lévy et al., 2012). In particular, time-varying eddy flows have been argued to provide a rectified pumping of nutrients 69 into the euphotic zone: an upward intrusion of nutrients into the euphotic zone leads to biological consumption, while a downward intrusion of nutrients into the dark ocean 70 interior leads to no biological response (McGillicuddy Jr & Robinson, 1997). While this rectification process is appealing in potentially sustaining an enhanced nutrient supply 71 to the euphotic zone, there is a price that remains unresolved from this process: an underlying loss of nutrients from the thermocline. 72 73 74 75 76

77 This difficulty in sustaining nutrient levels in the upper thermocline is highlighted in two contrasting model experiments. McGillicuddy Jr et al. (2003) illustrated how time-varying eddies may provide a dominant supply of nitrate to the euphotic zone over the 78 subtropical gyre, overcoming the effects of large-scale wind-driven downwelling, in an eddy-resolving model study including an artificial restoring of nutrients below the euphotic 79 zone. In contrast, Oschlies (2002) found a much weaker eddy supply of nutrients to the euphotic zone in model integrations without this interior restoring. Thus, the maintenance 80 of the nutrient concentrations in the thermocline is central to whether time-varying, 81 vertical motions may account for a sustained nutrient supply to the euphotic zone. 82 83 84 85

86 In our work, the question of how nutrients in the thermocline are sustained against the action of processes supplying nutrients to the overlying euphotic zone is explored in 87 terms of the competing effects of nutrient transfers by small-scale mixing across density surfaces, referred to as diapycnal mixing, advection across density surfaces, referred to 88 as diapycnal advection, and mesoscale eddy stirring along density surfaces, referred to as isopycnal stirring. First, a theoretical context is provided for how the nutrient budget 89 for a density layer is affected by advective and diffusive nitrate fluxes, including diapycnal transfers from microstructure and the isopycnal effects of mesoscale eddy advection 90 and diffusion (Section 2). Then, observations from a summer field campaign conducted in some of the most oligotrophic waters of the subtropical North Atlantic, in the 91 vicinity of the Mid-Atlantic Ridge (Tuerena et al., 2019), are considered (Section 3) to estimate the rates of diapycnal mixing, diapycnal advection and isopycnal stirring by mesoscale 92 eddies (Section 4). The supply of nutrients by diapycnal mixing, diapycnal advection and isopycnal stirring is then assessed, for different density classes, along two transects over 93 and off the Mid-Atlantic Ridge (Section 5). The relevance of our findings is then discussed 94 for our field site and for the rest of the North Atlantic subtropical gyre (Section 6). 95 96 97 98 99 100 101

2 Theoretical context

Our aim is to focus on the transfers of nutrients by both microscale turbulence and by mesoscale eddies using an isopycnal formulation. The layer-integrated tracer equation is firstly discussed for a nutrient, then the relevant diffusive transfer terms identified, and closures applied to estimate these terms.

2.1 Thickness-weighted tracer equation

Following Bleck and Boudra (1981), Bleck (1998) and McDougall (1984), the tracer equation integrated over the thickness of the density layer may be written as

$$\underbrace{\frac{\partial}{\partial t}(hN)}_{\text{tendency}} + \underbrace{\nabla_{\sigma} \cdot (h\mathbf{u}N)}_{\text{isopycnal advection}} + \underbrace{(w^*N)|_{bot} - (w^*N)|_{top}}_{\text{diapycnal advection}} = \underbrace{\nabla_{\sigma} \cdot (\nu h \nabla_{\sigma} N)}_{\text{isopycnal diffusion}} + \underbrace{F_{dia}|_{bot} - F_{dia}|_{top}}_{\text{diapycnal diffusion}} + \underbrace{hS}_{\text{biological source}}, \quad (1)$$

where h is the vertical thickness of the isopycnal layer, N is the nutrient tracer, \mathbf{u} is the velocity along the density layer, ∇_{σ} is the gradient along a density surface, w^* is the diapycnal velocity across the bounding density surfaces, denoted bottom and top, ν is the isopycnal diffusivity from microscale turbulence, F_{dia} is the diapycnal tracer flux across the bounding density surfaces, and S is the biological source. The gradient ∇_{σ} is evaluated with the distance taken from a projection along a horizontal plane, which avoids the inclusion of metric terms (Bleck & Boudra, 1981). The diapycnal velocity, w^* , follows the notation of McDougall (1984) and is equivalent to $\dot{s}\partial z/\partial\sigma$ in the notation of Bleck and Boudra (1981). The diffusive and diapycnal transfers are all taken to be representative of microscale turbulence.

Following Gent et al. (1995), consider the effect of time-varying mesoscale eddies, which are assumed to act to transfer tracers along density surfaces. Take a prime to represent a mesoscale eddy deviation and an overbar to represent a time average over the lifetime of many mesoscale eddy events, so that all variables are separated into mean and mesoscale eddy contributions, such as $\mathbf{u} = \bar{\mathbf{u}} + \mathbf{u}'$. Applying this partitioning to the thickness-weighted flux of tracer along the density layer, $h\mathbf{u}N$, and then applying a time average over the lifetime of many mesoscale eddies, obtains

$$\underbrace{\overline{h\mathbf{u}N}}_{\text{isopycnal tracer flux}} = \underbrace{(\bar{h}\bar{\mathbf{u}} + \overline{h'\mathbf{u}'})\bar{N}}_{\text{advective tracer flux}} + \underbrace{\overline{(h\mathbf{u})'N'}}_{\text{diffusive tracer flux}}, \quad (2)$$

where (i) the time-mean tracer \bar{N} is advected along the density layer by the time-mean velocity $\bar{\mathbf{u}}$ and the time-varying mesoscale eddies, $h'\mathbf{u}'/\bar{h}$, involving a correlation in velocity and layer thickness, and (ii) the tracer is diffused along the density layer by the mesoscale eddy correlations in the volume flux, $(h\mathbf{u})'$, and tracer concentration, N' .

Combining (1) and (2), the tracer equation integrated over a density layer and including a time average over the lifetime of eddy events is given by

$$\underbrace{\frac{\partial}{\partial t}(\overline{hN})}_{\text{tendency}} + \underbrace{\nabla_{\sigma} \cdot ((\bar{h}\bar{\mathbf{u}} + \overline{h'\mathbf{u}'})\bar{N})}_{\text{isopycnal advection}} + \underbrace{(\overline{w^*N})|_{bot} - (\overline{w^*N})|_{top}}_{\text{diapycnal advection}} = \underbrace{-\nabla_{\sigma} \cdot (\overline{(h\mathbf{u})'N'}) + \nabla_{\sigma} \cdot (\nu \overline{h \nabla_{\sigma} N})}_{\text{isopycnal diffusion}} + \underbrace{\overline{F_{dia}|_{bot}} - \overline{F_{dia}|_{top}}}_{\text{diapycnal diffusion}} + \underbrace{\overline{hS}}_{\text{biological source}}, \quad (3)$$

The evolution of the nutrient in the layer is then determined by (i) the isopycnal advection by the time-mean flow and the mesoscale eddies; (ii) the diapycnal advection

135 of the tracer across the bounding density surfaces; (iii) an isopycnal diffusive transfer by
 136 a combination of mesoscale eddies and microscale turbulence; (iv) a diapycnal diffusion
 137 by microscale turbulence; and (iv) a biological source including a rectification by eddy
 138 changes in layer thickness.

139 The mesoscale and microscale turbulent processes affect the tracer balance in distinct
 140 ways. The microscale turbulence provides a diapycnal diffusion and isopycnal diffusion
 141 of tracer together with a diapycnal advection of tracer (McDougall, 1984, 1987;
 142 Groeskamp, Griffies, et al., 2019). Mesoscale eddies provide an isopycnal transfer of tracer,
 143 involving both an isopycnal advection defining a bolus velocity given by $\overline{h'\mathbf{u}'}/\bar{h}$, and an
 144 isopycnal diffusion given by $\overline{(h\mathbf{u})'N'}$. There may also be an additional eddy advection
 145 of the tracer due to a skew flux contribution linked to the tracer distribution (Canuto
 146 & Dubovikov, 2011), which we ignore here. These isopycnal advective and diffusive effects
 147 of mesoscale eddies have distinctive tracer signals (Lee et al., 1997) and are parameterised
 148 separately in ocean models (Gent & McWilliams, 1990; Redi, 1982).

149 2.2 Diffusive nutrient supply

150 Using field data, we wish to compare the magnitude of the different nutrient supply
 151 terms within the seasonal boundary layer and thermocline in the tracer equation (3)
 152 involving the convergences of the diapycnal diffusion by microstructure, the diapycnal
 153 advection by microstructure, and the isopycnal diffusion by mesoscale eddies over an isopyc-
 154 nal layer of thickness h , which are represented respectively by

$$\underbrace{(\overline{F_{dia}|_{bot}} - \overline{F_{dia}|_{top}})/\bar{h}}_{\text{diapycnal diffusion}}, \quad \underbrace{((\overline{w^*N})|_{bot} - (\overline{w^*N})|_{top})/\bar{h}}_{\text{diapycnal advection}} \quad \text{and} \quad \underbrace{-(1/\bar{h})\nabla_\sigma \cdot \overline{(h\mathbf{u})'N'}}_{\text{isopycnal diffusion}},$$

155 which are in units of $\text{mol N m}^{-3}\text{s}^{-1}$. The isopycnal diffusion from mesoscale eddies is
 156 much larger than the contribution from microscale turbulence (Tennekes & Lumley, 2018),
 157 so the microscale turbulence contribution is ignored here.

158 2.2.1 Closure for diapycnal diffusion

159 The diapycnal diffusive flux of nutrients associated with microscale turbulence is
 160 parameterised here by a down-gradient closure,

$$\overline{F_{dia}} = -\kappa_{dia} \frac{\partial \overline{N}}{\partial z}, \quad (4)$$

161 where κ_{dia} is a diapycnal diffusivity. Hence, the convergences in the diapycnal diffusive
 162 tracer fluxes in (3) may be re-expressed as

$$\frac{\overline{F_{dia}|_{bot}} - \overline{F_{dia}|_{top}}}{\bar{h}} = \frac{1}{\bar{h}} \left(- \left(\kappa_{dia} \frac{\partial \overline{N}}{\partial z} \right) \Big|_{bot} + \left(\kappa_{dia} \frac{\partial \overline{N}}{\partial z} \right) \Big|_{top} \right). \quad (5)$$

163 2.2.2 Closure for diapycnal advection

164 Diapycnal advection is generated by contrasts in the diapycnal diffusive density flux.
 165 The diapycnal velocity, w^* , is defined here by

$$w^* = \frac{\partial F_{dia,\gamma}}{\partial \gamma} = \frac{\partial}{\partial \gamma} \left(-\kappa_{dia} \frac{\partial \gamma}{\partial z} \right), \quad (6)$$

166 where $F_{dia,\gamma}$ is the diapycnal diffusive flux of density, γ is the neutral density and κ_{dia}
 167 is the diapycnal diffusivity (Nurser et al., 1999; de Lavergne et al., 2016). The diapyc-
 168 nal velocity is directed towards the regions of enhanced mixing. Here we define the di-
 169 apycnal velocity as positive towards lighter classes due to our focus on upward nutrient

170 fluxes towards the euphotic zone, even though previous studies often define it as posi-
 171 tive towards denser classes. This closure excludes the effective advection driven by the
 172 non-linearity of the equation of state, however, this effect is typically small at low and
 173 mid latitudes (Klocker & McDougall, 2010).

174 The diapycnal advection of nitrate is found to be important on annual time scales
 175 when integrated over the entire water column (Groeskamp, Griffies, et al., 2019), although
 176 it is unclear how important this diapycnal advection of nitrate is within the thermocline.

177 *2.2.3 Closure for isopycnal diffusion by mesoscale eddies*

178 The isopycnal diffusive transfer of nutrients is viewed here as being provided by mesoscale
 179 eddies, involving quasi-geostrophic circulations stirring nutrients along density surfaces
 180 in an adiabatic manner. The thickness-weighted diffusive flux of nutrients, $\overline{(h\mathbf{u})'N'}$, is
 181 assumed to be mainly determined by the eddy correlations in velocity and nutrient con-
 182 centration, such that $\overline{(h\mathbf{u})'N'} \simeq \bar{h} \overline{\mathbf{u}'N'}$, which is parameterised in terms of the isopy-
 183 cnal gradient in nutrients, so that the isopycnal flux of nutrients, $\bar{\mathbf{F}}_{iso}$, is given by

$$\bar{\mathbf{F}}_{iso} = \overline{(h\mathbf{u})'N'}/\bar{h} = -\kappa_{iso}\nabla_{\sigma}\bar{N}, \quad (7)$$

184 where κ_{iso} is an isopycnal diffusivity and ∇_{σ} denotes the gradient along density surfaces.
 185 Hence, the convergences in the isopycnal diffusive tracer flux from mesoscale eddies in
 186 (3) may be re-expressed as

$$-\nabla_{\sigma} \cdot \overline{(h\mathbf{u})'N'} = -\nabla_{\sigma} \cdot (\bar{h} \bar{\mathbf{F}}_{iso}) = \nabla_{\sigma} \cdot (\bar{h}\kappa_{iso}\nabla_{\sigma}\bar{N}). \quad (8)$$

187 The convergence in the diffusive isopycnal flux of nutrients in (8) can be re-written
 188 using a co-ordinate transformation where the isopycnal gradient is made up of the sum
 189 of a horizontal gradient at a constant depth and the isopycnal slope multiplied by a ver-
 190 tical gradient at the density surface,

$$\nabla_{\sigma} \cdot \overline{(h\mathbf{u})'N'} = \nabla_z \cdot \overline{(h\mathbf{u})'N'} + \mathbf{S} \cdot \frac{\partial}{\partial z} \overline{(h\mathbf{u})'N'} \quad (9)$$

191 where ∇_z is the horizontal gradient operator and the vector representing the isopycnal
 192 slope is given by $\mathbf{S} = \mathbf{i}\partial z/\partial x|_{\sigma} + \mathbf{j}\partial z/\partial y|_{\sigma}$ with \mathbf{i} and \mathbf{j} unit vectors in the x and y di-
 193 rections.

194 Next, we apply this theoretical framework to field observations to assess the rel-
 195 ative roles in sustaining nutrients in the upper thermocline of the North Atlantic sub-
 196 tropical gyre from the diapycnal diffusive nutrient supply from microscale turbulence in
 197 (5), the diapycnal advective nutrient supply from microscale turbulence (6), and the isopy-
 198 cnal diffusive nutrient supply from mesoscale eddies in (8).

199 **3 Field programme**

200 The observations analysed here were obtained by a field study in some of the most
 201 oligotrophic waters of the North Atlantic subtropical gyre, between 24°N to 36°N, as part
 202 of the RidgeMix programme to investigate the role of internal tides in diapycnal nutri-
 203 ent supply to the euphotic zone (Fig. 2a). The sampling campaign was conducted on the
 204 *RRS James Clark Ross* (cruise JR15007) in May to June 2016. The field programme in-
 205 volved 67 stations sited either along the crest of the Mid-Atlantic Ridge or in the ad-
 206 jacent deep-ocean basin (Fig. 2a, blue and red dots, respectively). There is a marked dif-
 207 ference in the intensity of small-scale turbulence and associated nutrient fluxes between
 208 the on- and off-ridge regions (Tuerena et al., 2019), so that our subsequent analyses are
 209 accordingly separated into on- and off-ridge areas.

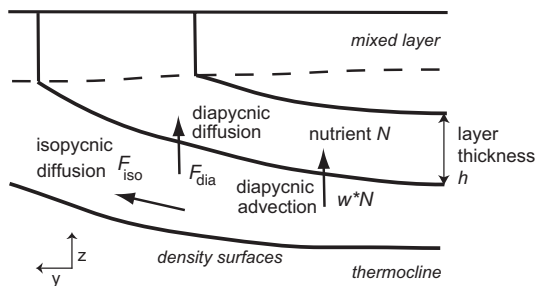


Figure 1. The diffusive nutrient flux, \mathbf{F} , may be separated into diapycnal and isopycnal components for a density layer with a nutrient concentration N and layer thickness h , the diffusive nutrient flux, \mathbf{F} , may be viewed as the sum of a diapycnal component directed across density surfaces (driven by microscale turbulence), F_{dia} , and an isopycnal component induced by mesoscale eddy stirring, F_{iso} . Diapycnal advection, w^* , is formed by contrasts in the diapycnal diffusive flux of density and is directed towards regions of turbulence, and so also provides a diapycnal nutrient flux, w^*N . The diapycnal and isopycnal transfers are potential mechanisms to supply nutrients, N , to the upper thermocline and mixed layer, and so contribute to sustaining export production from the euphotic zone.

210 At all of the CTD stations, temperature, salinity and water samples were measured
 211 for the full depth or the upper 1000 m of the water column. Micro-molar nutrient mea-
 212 surement was carried out at all CTD stations, for the analysis of nitrate, nitrite, phos-
 213 phosphate and silicate, using a four-channel Bran and Luebbe AAIII segmented flow, colori-
 214 metric, autoanalyzer. Two internal standards for nitrate, phosphate, and silicate were
 215 analysed in each run, covering the range of concentrations in the deep and surface lay-
 216 ers. Certified reference materials (Kanso) were analysed every 2-3 runs to ensure con-
 217 tinued precision throughout the cruise, and cruise averages within the accepted range
 218 for each nutrient and with a 99% precision.

219 In this region, the mixed layer extends to a depth of less than 50 m during sum-
 220 mer, and increases to respective depths of 150 m to 100 m at the end of winter at 36°N
 221 and 24°N, based upon climatology (Fig. 2b,c, red and blue dashed lines). The density
 222 surfaces, strictly Neutral Density (Jackett & McDougall, 1997), outcropping in the win-
 223 ter mixed layer over the region include $\gamma = 26$ to 26.5 (kg m^{-3}). The cruise sections re-
 224 veal the expected southward deepening of density surfaces within the subtropical gyre
 225 (Fig. 2c, black lines). The euphotic zone ranges from depths of 50 m to 150 m, and con-
 226 tains waters that are relatively depleted in nitrate (Fig. 2c, dashed magenta line and green
 227 dots). Below the euphotic zone, there are relatively enriched nutrients, reaching typically
 228 $5 \mu\text{mol kg}^{-1}$ at 250 m both along and off the ridge (Fig. 2c, green dots). The deep chloro-
 229 phyll maximum deepens southward from a depth of 150 m at 36°N to 200 m at 26°N (Fig.
 230 2c, dashed magenta line and blue shading).

231 Measurements of microstructure temperature variance and velocity shear were made
 232 at 30 of the stations using a free-falling Vertical Microstructure Profiler (VMP-6000, Rock-
 233 land Scientific). The microstructure was measured on the length scales of dissipation of
 234 turbulent flows, typically a few millimetres to tens of centimetres. The rates of dissipa-
 235 tion of turbulent kinetic energy, ϵ (W kg^{-1}), and of temperature variance, χ ($^{\circ}\text{C}^2\text{s}^{-1}$),
 236 were estimated following the standard methods after Oakey (1982).

237 A mooring provided measurements of horizontal velocity for the full water column
 238 for 9 months between September 2015 and July 2016 at 36.23°N, 32.75°W (Fig. 2a). The
 239 mooring included two Teledyne RD Instruments 75-kHz Long Ranger acoustic Doppler

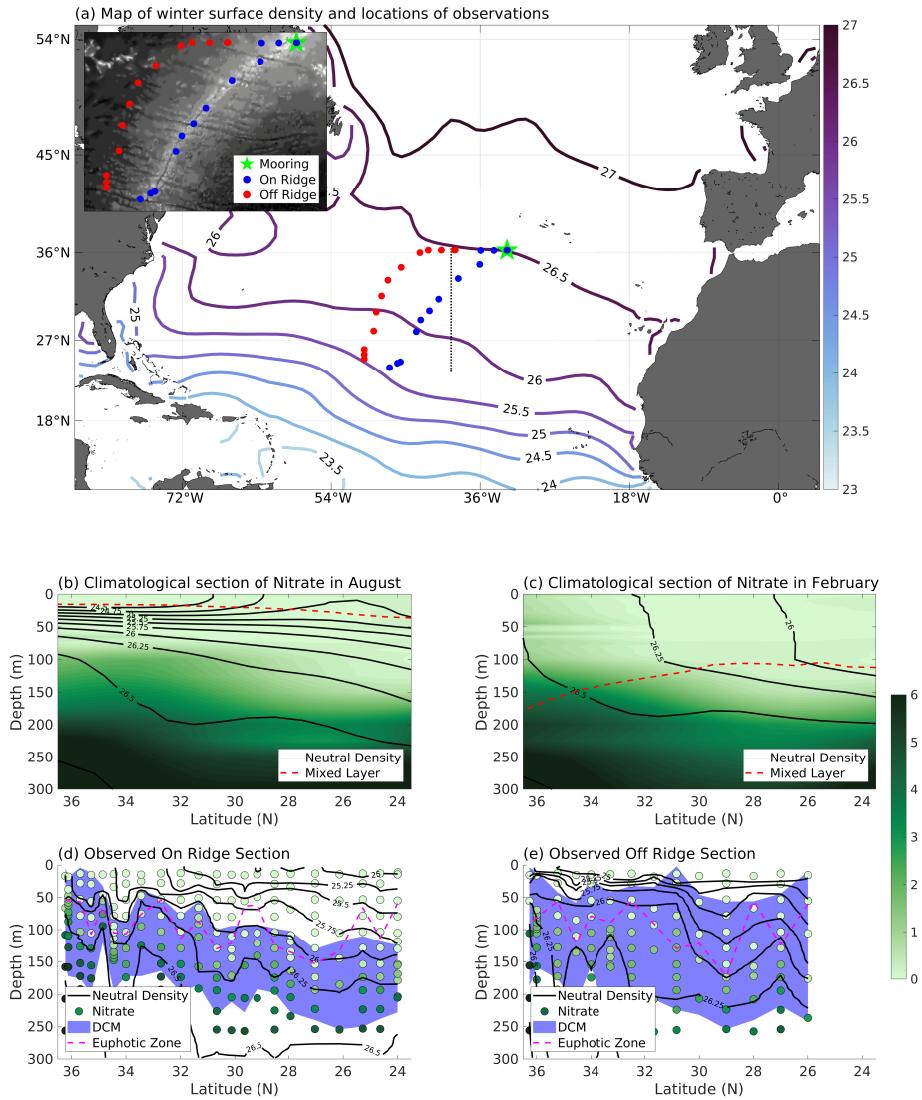


Figure 2. Context of the observational study showing: (a) map of the February surface density from climatology, with the locations of: stations marked with circles, separated into on-ridge (blue) and off-ridge (red); the mooring (green star); and the location of climatology sections (grey lines). The inset shows the positions of the observations against the backdrop of topography. Climatological meridional sections for (b) August and (c) February for nitrate ($\mu\text{mol kg}^{-1}$, background colours), neutral density (black lines), and the base of the mixed layer (red dashed line). Observed meridional sections for (d) on-ridge or (e) off-ridge, showing nitrate (green circles), neutral density (black lines), deep chlorophyll maximum (DCM, blue shading), and the base of the euphotic zone (dashed magenta line). The DCM is defined by the depth of a 10% variation in the maximum fluorescence, and the base of the euphotic zone is defined by 1% of photosynthetic radiation measured at a depth of 6 m.

240 current profilers (ADCPs) and two Flowquest 75-kHz ADCPs. All ADCPs recorded hourly-
241 averaged horizontal velocity in 8 m vertical bins (Vic et al., 2018).

242 **4 Estimating diapycnal mixing, diapycnal advection and isopycnal eddy** 243 **stirring from observational data**

244 The rates of diapycnal mixing and advection by small-scale turbulence and isopyc-
245 nal stirring by mesoscale eddies are now assessed using the field observations. Although
246 the diapycnal diffusivity associated with small-scale turbulence is typically 8 orders of
247 magnitude smaller than the isopycnal diffusivity linked to mesoscale eddy stirring, the
248 nutrient gradient across density surfaces is much greater than the nutrient gradient along
249 density surfaces, such that diapycnal and isopycnal diffusive fluxes of nutrients are broadly
250 comparable in magnitude.

251 **4.1 Diapycnal diffusivity**

252 The diapycnal diffusivity associated with small-scale turbulence is diagnosed using
253 a turbulence closure (Osborn, 1980), which assumes a balance between the rates of
254 turbulent kinetic energy dissipation and the turbulent buoyancy flux,

$$\kappa_{dia} = \Gamma \frac{\epsilon}{N^2}, \quad (10)$$

255 where, Γ is the dissipation ratio indicating the proportion of the turbulent kinetic en-
256 ergy being used to mix tracers, N^2 is the buoyancy frequency defined as $N^2 = -\frac{g}{\rho} \frac{\partial \sigma}{\partial z}$
257 (where σ is the local potential density and g is the gravitational acceleration), and the
258 empirical choice of $\Gamma = 0.2$ is representative for stratified shear turbulence (Gregg et
259 al., 2018). The diapycnal diffusivity, κ_{dia} , is diagnosed in neutral density bins of 0.12 kg
260 m^{-3} calculated using the routines of Jackett and McDougall (1997).

261 The diapycnal diffusivity is systematically enhanced over the Mid-Atlantic Ridge
262 relative to off the ridge by a factor of typically 2 to 6, ranging from $5 \times 10^{-5} \text{m}^2 \text{s}^{-1}$ to
263 $4 \times 10^{-6} \text{m}^2 \text{s}^{-1}$ for neutral surfaces between $\gamma = 25.5$ and 28 (Fig. 3a, blue and red lines).
264 In both cases these diffusivities are much larger than molecular diffusivity, typically $10^{-7} \text{m}^2 \text{s}^{-1}$,
265 implying the diffusivity can reasonably be applied to any tracer.

266 **4.2 Diapycnal advection**

267 The diapycnal advection, w^* , is generated by a convergence in the diapycnal dif-
268 fusive density flux, $F_{dia,\gamma}$ from (6). The diapycnal diffusivity is larger on the lighter and
269 denser surfaces with a minimum at $\gamma = 26.2$, and the vertical gradient in the neutral den-
270 sity is larger for density layers lighter than $\gamma = 26.2$, but relatively weak for denser lay-
271 ers (Fig. 4a,b). The resulting diapycnal diffusive density flux, $F_{dia,\gamma}$, is directed towards
272 denser surfaces, with larger values along lighter surfaces and a minimum value close to
273 $\gamma = 26.4$ (Fig. 4c). The diapycnal advection, w^* , is then directed towards the regions
274 of elevated turbulence, i.e. from dense to light surfaces for densities lighter than $\gamma = 26.4$,
275 but towards larger densities from $\gamma = 27$ (Fig. 4d). There is the same vertical structure
276 for the diapycnal advection both on and off ridge, although w^* is typically an order of
277 magnitude higher on the ridge due to the stronger tidal mixing. The errors in the esti-
278 mate of w^* are large due to taking the diapycnal gradient of the diapycnal diffusivity,
279 which has large uncertainties, combined with the vertical gradient of neutral density.

280 Hence, this structure of the diapycnal velocity, w^* , directed towards the density
281 extremes, is consistent with the most intense turbulence in the water column occurring
282 near the surface and bottom boundaries, most likely driven by wind forcing and flow-
283 topography interaction, and a mid-water column minimum. Such vertical structure in

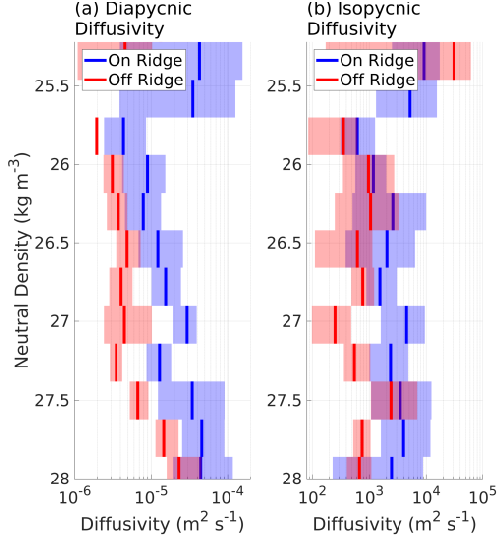


Figure 3. Profiles in neutral density of (a) the average diapycnal diffusivity and (b) the average isopycnal diffusivity for the on-ridge (blue) and off-ridge (red) stations. Shading indicates the 95% bootstrap confidence interval.

284 the intensity of turbulent mixing and resultant diapycnal velocities is compatible with
 285 previous observations on a global scale (Kunze et al., 2006; Waterhouse et al., 2014).

286 4.3 Isopycnal diffusivity from a tracer variance approach

287 The isopycnal diffusivity associated with mesoscale eddy stirring is diagnosed us-
 288 ing a temperature variance budget approach (Ferrari & Polzin, 2005),

$$\underbrace{\frac{\partial \overline{\theta'^2}}{\partial t}}_{\text{tendency}} + \underbrace{\nabla \cdot (\overline{\mathbf{u}\theta'^2} + \overline{\mathbf{u}'\theta'^2} - \kappa_\theta \nabla \overline{\theta'^2})}_{\text{advective and diffusive fluxes}} + \underbrace{2\overline{\mathbf{u}'\theta'} \cdot \nabla \overline{\theta}}_{\text{production}} = \underbrace{-\chi}_{\text{dissipation}}, \quad (11)$$

289 where θ is the potential temperature, χ is the rate of dissipation of temperature vari-
 290 ance, κ_θ is the molecular diffusivity of temperature, the velocity is a full three-dimensional
 291 velocity and ∇ is a full three-dimensional gradient operator here, and again the overbar
 292 indicates a time average over many eddy events and the primes indicate the deviation
 293 from this average. There is a balance between the tendency of temperature variance, the
 294 divergence of the advective and diffusive fluxes of variance, production of temperature
 295 variance, and the dissipation of temperature variance.

296 In our observations, the root mean square temperature anomalies on density sur-
 297 faces are $\overline{\theta'^2} = 0.1$ °C at depth, increasing to 0.3 °C towards the surface. By combin-
 298 ing these estimates with temperature variance dissipation rates, typically $\chi = 10^{-9}$ °C²s⁻¹
 299 at depth to 10^{-8} °C²s⁻¹ near the surface, then a lifetime for the temperature variance
 300 can be estimated of typically 100 days (see Naveira Garabato et al. (2016)). Assuming
 301 a background mean flow for this region of 0.01 ms⁻¹, then the temperature variance is
 302 likely to be advected over a horizontal scale of 100 km before being dissipated, which is
 303 a much smaller scale than the horizontal extent of our sections spanning 1000 to 2000 km.
 304 Hence, the tendency, advection and diffusion terms in (11) are neglected, so that an ap-
 305 proximate local balance is assumed to hold between the production and dissipation of

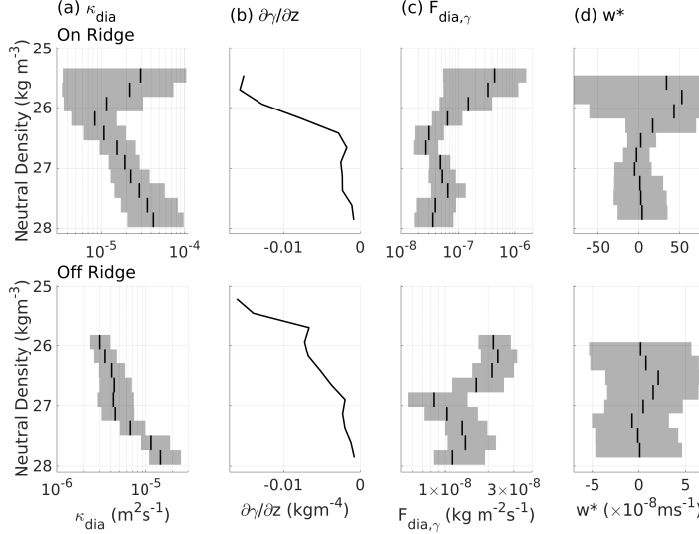


Figure 4. The diapycnal advection for the on-ridge (upper row) and off-ridge (lower row) stations. The profiles: (a) diapycnal diffusivity, κ_{dia} (m^2s^{-1}), (b) vertical density gradient, $\partial\gamma/\partial z$ (kg m^{-4}), (c) the diapycnal diffusive flux of density, $F_{dia,\gamma}$ ($\text{kg m}^{-2}\text{s}^{-1}$), positive directed towards denser surfaces and (d) diapycnal advection, w^* (m s^{-1}), positive directed upwards. The black lines show the mean values derived from the microstructure data and the cruise-observed gradients. The shaded regions indicate 95% confidence ranges using a bootstrap method.

306 temperature variance,

$$\overline{\mathbf{u}'\theta'} \cdot \nabla\bar{\theta} = -\chi/2. \quad (12)$$

307 Following Naveira Garabato et al. (2016), the production of temperature variance
 308 is decomposed into (i) microscale turbulence providing a diapycnal transfer and (ii) mesoscale
 309 eddy stirring providing an isopycnal transfer:

$$\overline{\mathbf{u}'_t\theta'_t} \cdot \frac{\partial\bar{\theta}}{\partial z} + \overline{\mathbf{u}'_e\theta'_e} \cdot \nabla_\sigma\bar{\theta} = -\chi/2, \quad (13)$$

310 where contributions of the mesoscale (e) and microscale (t) are separated from the long-
 311 term average (denoted by the overbar). Employing down-gradient closures for the di-
 312 apycnal and isopycnal eddy stirring terms, (4) and (7), then

$$-\kappa_{dia} \left| \frac{\partial\bar{\theta}}{\partial z} \right|^2 - \kappa_{iso} |\nabla_\sigma\bar{\theta}|^2 = -\chi/2, \quad (14)$$

313 which allows the isopycnal diffusivity, κ_{iso} , to be diagnosed using

$$\kappa_{iso} = \frac{\chi/2 - \kappa_{dia} \left| \frac{\partial\bar{\theta}}{\partial z} \right|^2}{|\nabla_\sigma\bar{\theta}|^2}. \quad (15)$$

314 An example of this diagnostic method applied to a single station is shown in Appendix A.

315 The isopycnal diffusivity, κ_{iso} , is estimated within different neutral density bins, using
 316 χ and ϵ from 30 VMP-6000 casts, N^2 from the CTD casts adjacent to the VMP deploy-
 317 ments, and large-scale potential temperature gradients estimated either across density
 318 surfaces from the CTD sections or along density surfaces in the World Ocean Atlas cli-
 319 matology (Locarnini et al., 2013; Zweng et al., 2013). Unphysical values of κ_{iso} are re-
 320 moved from the data, based upon the occurrence of either negative κ_{iso} (where χ is less

321 than the production of temperature variance) or near-zero isopycnal temperature gra-
 322 dents (less than $5 \times 10^{-7} \text{ }^\circ\text{C m}^{-1}$).

323 The resulting estimates of isopycnal diffusivity typically range from 1000 to 4000 m^2s^{-1}
 324 along the ridge (Fig. 3b, blue line). Isopycnal stirring rates are weaker off the ridge, typ-
 325 ically by a factor of 2 to 4 (Fig. 3b, red line). There is a similar enhancement in diapyc-
 326 nal and isopycnal diffusivity on the ridge for densities greater than 26, which is prob-
 327 ably a result of the enhanced tidal mixing there (Vic et al., 2018; Tuerena et al., 2019).

328 4.4 Isopycnal diffusivity from a mixing-length approach

329 The isopycnal diffusivity, κ_{iso} , is also estimated using a mixing-length approach based
 330 upon assumed quasi-geostrophic dynamics (Ferrari & Nikurashin, 2010). This estimate
 331 entails a two-stage process. First, the isopycnal diffusivity in the absence of a mean flow,
 332 $\kappa_{iso,0}$, is estimated from the horizontal scale over which mesoscale eddy stirring is re-
 333 dicted to operate, k^{-1} , related to perturbations in sea surface height, h' , and to eddy
 334 kinetic energy, EKE ,

$$\kappa_{iso,0} = d_1 \frac{g}{|f|} (\overline{h'^2})^{1/2} = \frac{1}{2} d_0 k^{-1} EKE^{1/2}, \quad (16)$$

335 where d_0 and d_1 are empirically-derived coefficients, linked to the decorrelation time scale,
 336 which have been applied in different dynamical regimes of the Southern Ocean (Ferrari
 337 & Nikurashin, 2010). Second, the isopycnal diffusivity in the presence of a mean flow,
 338 κ_{iso} , is expressed as a modification to the diffusivity without a mean flow (Naveira Gara-
 339 bato et al., 2011),

$$\kappa_{iso} = \frac{\kappa_{iso,0}}{1 + d_2 U_0^2 EKE^{-1}}, \quad (17)$$

340 where d_2 is an empirically-derived coefficient, linking the phase speed of generated ed-
 341 dies to the mean flow, and U_0 is the background mean flow velocity. An alternative ac-
 342 count for the suppression entails the replacement of the mean flow speed (U_0) with the
 343 Rossby wave speed (c_w) (Klocker & Abernathy, 2014).

344 The isopycnal diffusivity in (17) is evaluated using a combination of altimetric data
 345 and mooring observations, by following this procedure:
 346 (i) The mixing length scale k^{-1} is estimated from altimetry by comparing the diffusiv-
 347 ity derived from sea surface height to the EKE (16) based upon a daily-gridded altimet-
 348 ric product with 1/4 degree horizontal resolution between September 2015 and June 2017;
 349 (ii) A profile of the isopycnal diffusivity in the absence of a mean flow, $\kappa_{iso,0}$, (16) is di-
 350 agnosed from a profile of EKE (derived from mooring-based horizontal velocity data pro-
 351 cessed with a 40-hour low-pass Butterworth filter) combined with the mixing-length scale
 352 in (i) averaged in a 1-degree box around the mooring;
 353 (iii) A profile of the isopycnal diffusivity in the presence of the mean flow, κ_{iso} , is cal-
 354 culated (17) using the output of (ii), plus the mooring-based estimates of the mean flow
 355 and EKE, and coefficients derived by Ferrari and Nikurashin (2010).

356 At the site of the mooring, EKE reaches just over $0.007 \text{ m}^2\text{s}^{-2}$ close to the surface,
 357 where the mean flow speed is 2.9 cm s^{-1} . Both EKE and mean flow speed decrease rapidly
 358 over a depth scale of 500 m to $0.0036 \text{ m}^2\text{s}^{-2}$ and 0.5 cm s^{-1} , respectively (Fig. 5a,b).
 359 The first baroclinic Rossby wave speed for this location is given by $c_w = -\beta L_d^2$, where
 360 β is the meridional gradient in the Coriolis parameter and L_d is the first baroclinic mode
 361 Rossby deformation radius. Taking typical values for these inputs at the mooring site,
 362 $\beta = 1.8 \times 10^{-11} \text{ m}^{-1}\text{s}^{-1}$ and $L_d = 30 \text{ km}$ (Tulloch et al., 2009), gives a wave speed
 363 of $c_w = 1.6 \text{ cm s}^{-1}$. This speed is comparable to the mean flow speeds shown by the
 364 mooring (Fig. 5b), implying that the suppression is not strongly sensitive to the choice
 365 between the mean flow and Rossby wave speeds. The isopycnal diffusivity peaks at $1550 \text{ m}^2\text{s}^{-1}$
 366 at the surface and reduces to typically $630 \text{ m}^2\text{s}^{-1}$ at a depth of 1000 m (Fig. 5c, black
 367 line).

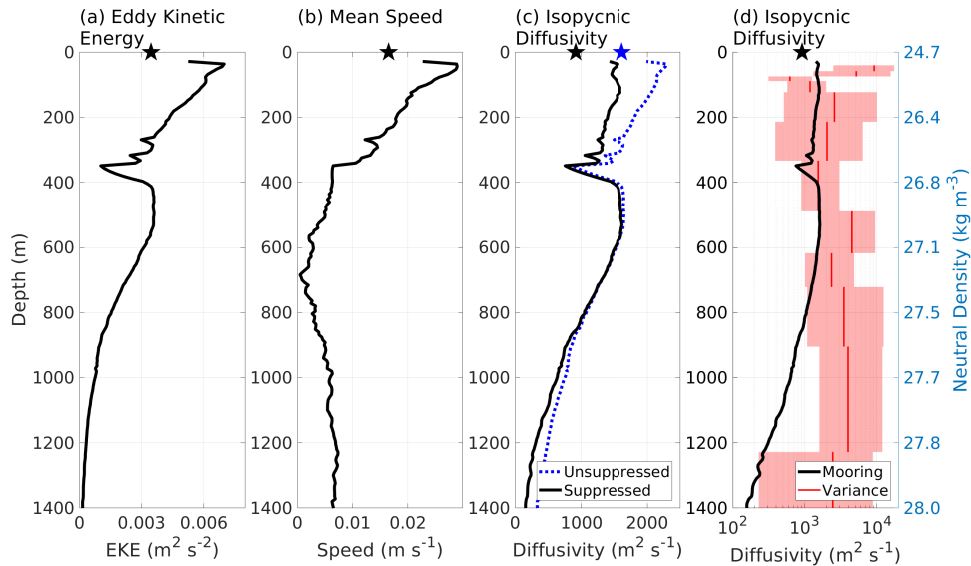


Figure 5. Profiles from the mooring (lines) and surface estimates from altimetry (stars) of: (a) eddy kinetic energy, (b) the time-mean velocity, (c) the isopycnal diffusivity in the absence (blue dashed line) and presence (black line) of the mean flow, and (d) the isopycnal diffusivity from the mooring with mean flow (black line) and the isopycnal diffusivity on the ridge calculated using the temperature variance approach (red lines and boxes). The diffusivity from temperature variance is remapped using the median depth of the density surface across the survey.

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4.5 Comparison with other estimates of isopycnal diffusivity from eddy stirring

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In this study, the effect of mesoscale eddies in providing an isopycnal diffusivity has been estimated for the North Atlantic subtropical gyre using two independent methods, one utilising a tracer variance approach and the other founded on mixing length theory. Both estimates indicate a maximum isopycnal diffusivity near the surface of approximately 2000 to 4000 $\text{m}^2 \text{s}^{-1}$, characteristically decaying with depth to 500 to 2000 $\text{m}^2 \text{s}^{-1}$ off and on ridge, respectively. The mixing length estimates are typically slightly lower than the temperature variance estimates, although they are indistinguishable within error envelopes for most depth bins (Fig. 5d).

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There have been a range of investigations of isopycnal diffusivity in the Northeast Atlantic, particularly in the region of the Mediterranean outflow, that yield diffusivity estimates at 250 m of 800 to 1400 $\text{m}^2 \text{s}^{-1}$ based on measurements of thermohaline variability (Joyce et al., 1998) and of order 1000 $\text{m}^2 \text{s}^{-1}$ from a tracer release experiment (Ledwell et al., 1998). Similar estimates of the diffusivity, 1100 $\text{m}^2 \text{s}^{-1}$ near the surface and 100 to 300 $\text{m}^2 \text{s}^{-1}$ at depth, have been made for the region on the basis of conservation of salt and temperature (Zika & McDougall, 2008; Zika et al., 2010).

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There have also been a range of global estimates of the isopycnal diffusivity that are broadly in line with our diagnostics. Theoretical arguments applied to a global climatology at the location of our mooring imply an isopycnal diffusivity that reaches a surface maximum of 1000 $\text{m}^2 \text{s}^{-1}$ and decays to 400 $\text{m}^2 \text{s}^{-1}$ at 1000 m depth (Groeskamp et al., 2020). Global surface calculations give an approximate surface diffusivity of 1000 to 2000 $\text{m}^2 \text{s}^{-1}$ for our region, using separation of surface floats (Roach et al., 2018) and

altimetry (Abernathey & Marshall, 2013; Klocker & Abernathey, 2014). Isopycnal diffusivities of 500 to 1500 m² s⁻¹ at 1000 m depth are given by Argo float observations, both from float separation statistics (Roach et al., 2018) and thermohaline distributions in combination with model velocities (Cole et al., 2015).

Thus, the results presented in this study are broadly consistent, in terms of both magnitude and vertical structure, with isopycnal diffusivity estimates from a range of previous studies.

5 Nutrient supply by diapycnal mixing, diapycnal advection and isopycnal eddy stirring

The estimates of the diapycnal and isopycnal diffusivities, respectively associated with microscale turbulence and mesoscale eddies, and the diapycnal advection, are next combined with the nitrate data to assess their relative roles in nutrient supply to the upper layers of the North Atlantic subtropical gyre. Here we focus on the contributions of three terms in the nutrient budget: the vertical convergences in the diapycnal diffusion, diapycnal advection, and isopycnal diffusion (see Section 2.2):

$$\underbrace{(\overline{F_{dia}}|_{bot} - \overline{F_{dia}}|_{top})/\overline{h}}_{\text{diapycnal diffusion}}, \quad \underbrace{((\overline{w^*N})|_{bot} - (\overline{w^*N})|_{top})/\overline{h}}_{\text{diapycnal advection}}, \quad \text{and} \quad \underbrace{(\mathbf{S}/\overline{h}) \cdot \left(\frac{\partial}{\partial z}(\overline{h}\kappa_{iso}\nabla_{\sigma}\overline{N})\right)}_{\text{isopycnal diffusion}}. \quad (18)$$

These analyses were also applied to phosphate and lead to broadly similar inferences (not shown here) and the details of the calculation of the gradients are provided in Appendix B.

5.1 Diapycnal diffusive nitrate fluxes

For the diapycnal contribution to the diffusive nitrate flux, F_{dia} , the diapycnal diffusivity, κ_{dia} , is estimated from cruise measurements along the ridge, ranging from an upper bound of 4 to 5 × 10⁻⁵ m² s⁻¹ at $\gamma = 25.4$ and 27.8, to a lower bound of 4 × 10⁻⁶ m² s⁻¹ at $\gamma = 25.8$ (Fig. 6a). The diapycnal nitrate gradient, $\partial N/\partial z$, varies in the opposite sense, from relatively low values at $\gamma = 25.4$ and 27.8, to higher gradients between $\gamma = 26.2$ and 27.2 (Fig. 6b, black lines for CTD stations). The diapycnal contribution of the diffusive nitrate flux, F_{dia} , thus varies from lower values of 6 × 10⁻⁵ and 8 × 10⁻⁴ mol N m⁻² yr⁻¹ at $\gamma = 25.4$ and $\gamma = 28$, to higher values of 2 × 10⁻² to 3 × 10⁻² mol N m⁻² yr⁻¹ at $\gamma = 26.5$ to 27.2 (Fig. 6c). This density variation of F_{dia} resembles the variation of $\partial N/\partial z$ more closely than that of κ_{dia} .

Estimates of the diapycnal component of the vertical nitrate flux, F_{dia} , decrease in magnitude off the ridge by a factor of typically 2 to 3, due to the smaller diapycnal diffusivity, κ_{dia} , there (Tuerena et al., 2019). The estimates of F_{dia} increase slightly in magnitude if the nitrate gradients are taken from climatology (Fig. 6b, red lines) rather than from the cruise observations.

5.2 Diapycnal advective nitrate fluxes

On the ridge, the diapycnal advection, w^* , is relatively large and positive, directed towards lighter surfaces at $\gamma = 25.6$, and becomes smaller and negative, directed towards denser surfaces at $\gamma = 27$ (Fig. 7a). The accompanying nitrate concentrations are close to zero on the density surfaces lighter than $\gamma = 26$ and become largest around $\gamma = 27.4$ (Fig. 7b). Consequently, the diapycnal advection of nutrients is small on lighter surfaces and becomes negative at $\gamma = 27.2$ (Fig. 7c). There are, however, large errors in this estimate arising from the estimate of w^* and the dependence on vertical gradients in κ_{dia} .

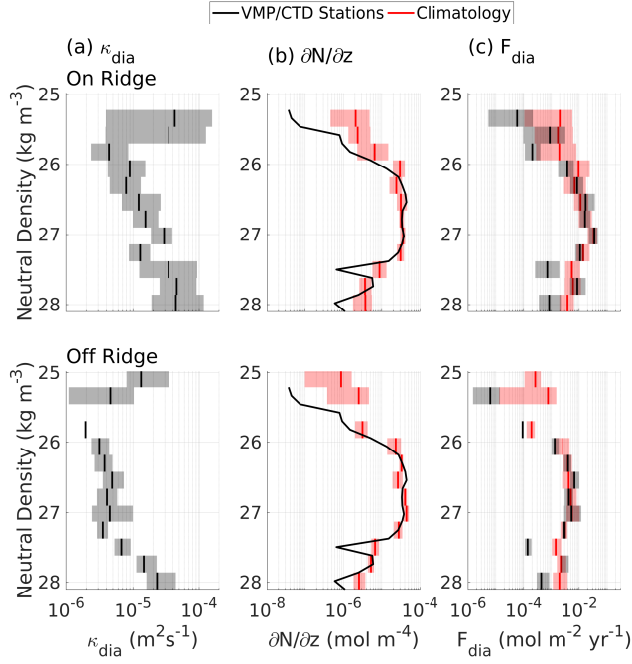


Figure 6. The diapycnal component of the vertical diffusive nitrate flux, F_{dia} , plotted versus neutral density, for the on-ridge (upper row) and off-ridge (lower row) stations: (a) diapycnal diffusivity, κ_{dia} (m^2s^{-1}), (b) nitrate gradient directed across density surfaces, $\partial N/\partial z$ (mol N m^{-4}) and (c) F_{dia} ($\text{mol N m}^{-2}\text{yr}^{-1}$). The black lines show the mean values derived from the microstructure data and the cruise-observed gradients. The red lines are estimates for which the gradients have been derived from climatological data. The shaded regions indicate 95% confidence ranges using a bootstrap method.

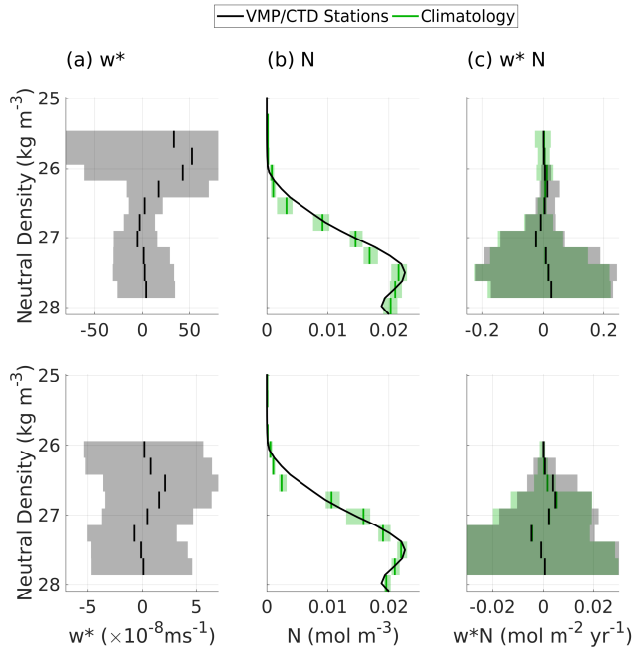


Figure 7. The diapycnal advection contribution to the vertical nitrate supply plotted versus neutral density, for the on-ridge (upper row) and off-ridge (lower row) stations. The profiles of (a) diapycnal advection, w^* (m s^{-1}) directed positive upwards, (b) nitrate concentration on density surfaces, N (mol m^{-3}), and (c) diapycnal advective nitrate flux, w^*N ($\text{mol N m}^{-2}\text{yr}^{-1}$). The black lines show the mean values derived from the microstructure data and the cruise-observed gradients. The green lines are estimates for which the gradients have been derived from climatological data. The shaded regions indicate 95% confidence ranges using a bootstrap method.

433 Off the ridge, the diapycnal advection, w^* , is again positive on lighter surfaces at
 434 $\gamma = 26.5$ and negative on denser surfaces at $\gamma = 27.2$. The diapycnal advection is an or-
 435 der of magnitude weaker off the ridge, and the upwelling extends onto denser surfaces
 436 compared with on the ridge. The vertical structure of the nitrate is similar to that on
 437 the ridge. The diapycnal advection of nutrients is then small and positive on lighter sur-
 438 faces at $\gamma = 26.6$ and changes to negative at $\gamma = 27.2$ (Fig. 7c). There are again large
 439 errors associated with the diapycnal velocity calculation, which is carried through to the
 440 nitrate flux.

441 5.3 Isopycnal diffusive nitrate fluxes

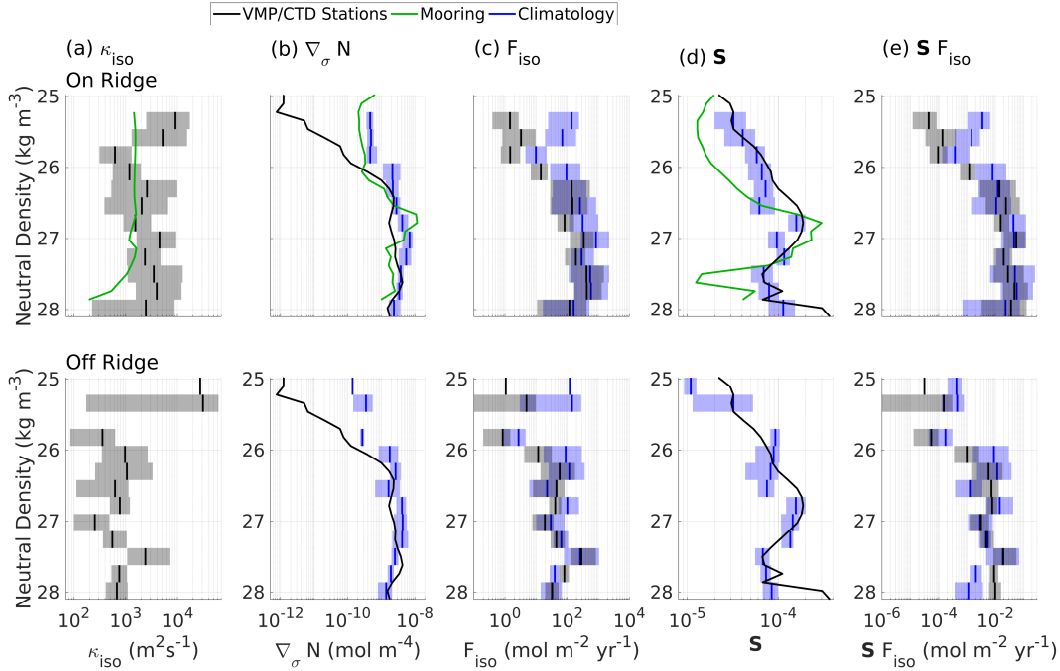


Figure 8. The isopycnal stirring contribution to the vertical diffusive nitrate flux plotted versus neutral density, for the on-ridge (upper row) and off-ridge (lower row) stations. The profiles: (a) isopycnal diffusivity, κ_{iso} (m^2s^{-1}), (b) nitrate gradient along density surfaces, $\nabla_{\sigma}N$ (mol N m^{-4}), (c) isopycnal nitrate flux, $-\kappa_{iso}\nabla_{\sigma}N$ ($\text{mol N m}^{-2}\text{yr}^{-1}$), (d) slope parallel to density surfaces, S , and (e) vertical component of the isopycnal nitrate flux, SF_{iso} ($\text{mol N m}^{-2}\text{yr}^{-1}$). The black lines show the mean values derived from the microstructure data and the cruise-observed gradients. The blue lines are estimates for which the gradients have been derived from climatological data. The green lines show the mixing length-estimated isopycnal diffusivity based on climatological gradients. The shaded regions indicate 95% confidence ranges using a bootstrap method.

442 For the isopycnal contribution to the diffusive nitrate flux, F_{iso} , the isopycnal dif-
 443 fusivity, κ_{iso} , is estimated from cruise measurements along the ridge, ranging from an
 444 upper bound of $8 \times 10^3 \text{m}^2\text{s}^{-1}$ at $\gamma = 25.4$ to a lower bound of $6 \times 10^2 \text{m}^2\text{s}^{-1}$ at $\gamma =$
 445 25.8 (Fig. 8a). The nitrate gradient along density surfaces, $\nabla_{\sigma}N$, is very low along $\gamma =$
 446 25 (less than $10^{-11} \text{mol N m}^{-4}$), and increases to a higher value of typically 1×10^{-9}
 447 to $4 \times 10^{-9} \text{mol N m}^{-4}$ for $\gamma = 26.2$ to 27.8 (Fig. 8b).

448 The resulting isopycnal contribution to the vertical diffusive nitrate flux, F_{iso} , ranges
 449 from a lower value of $1.4 \text{ mol N m}^{-2}\text{yr}^{-1}$ along $\gamma = 25.3$ to an upper value of $430 \text{ mol N m}^{-2}\text{yr}^{-1}$
 450 along $\gamma = 27.7$ (Fig. 8c). Most of this isopycnal stirring of nitrate is effectively hori-
 451 zontal. However, a small fraction of the stirring is directed vertically, and this fraction
 452 is given by the isopycnal slope, \mathbf{S} , which typically varies from 1×10^{-5} to 2×10^{-4} (Fig. 8d).
 453 The resulting isopycnal contribution to the vertical diffusive flux, F_{iso} , varies between
 454 lower values of 5×10^{-5} at $\gamma = 25.3$ and higher values of 1.5×10^{-2} to $5.5 \times 10^{-2} \text{ mol N m}^{-2}\text{yr}^{-1}$
 455 at $\gamma = 26.5$ to 27.2 (Fig. 8e).

456 Estimates of F_{iso} decrease in magnitude off the ridge by a factor of typically 2, due
 457 to the smaller isopycnal diffusivity, κ_{iso} , there, linked to the smaller dissipation of ther-
 458 mal variance off the ridge from (15). The estimates of F_{iso} increase slightly in magni-
 459 tude if the nitrate gradients are taken from climatology, especially on lighter surfaces.

460 5.4 Vertical structure of the nitrate fluxes and the seasonal boundary 461 layer

462 The relative importance of the diapycnal and isopycnal contributions to the verti-
 463 cal diffusive nitrate fluxes, F_{dia} and F_{iso} , and the diapycnal advection of nitrate, w^*N ,
 464 vary with the proximity to the mixed layer and euphotic zone along the ridge, irrespec-
 465 tive of whether the nutrients are from climatology or the cruise sections (Fig. 9a,b).

466 For light density surfaces that intersect the summer mixed layer and euphotic zone
 467 from $\gamma = 25.2$ to 26.0 , the vertical component of the diapycnal diffusive nitrate flux, F_{dia} ,
 468 is generally larger than that of the isopycnal diffusive nitrate flux, F_{iso} . The diapycnal
 469 advection is small in the summer surface mixed layer, where the nutrients are generally
 470 deplete. For surfaces that intersect the winter mixed layer between $\gamma = 26.2$ and 26.8 ,
 471 the nutrient transfer components are comparable to each other, sometimes with the di-
 472 apycnal diffusive component, F_{dia} , or the isopycnal diffusive component, F_{iso} , dominat-
 473 ing. The diapycnal advective flux, w^*N , is directed towards lighter surfaces, with this
 474 upward transfer being due to the enhanced mixing in the surface boundary layer.

475 For denser surfaces below the winter mixed layer from $\gamma = 26.8$, the isopycnal dif-
 476 fusive nitrate flux, F_{iso} , is generally larger than the diapycnal diffusive nitrate flux, F_{dia} ,
 477 with a maximum value of $0.055 \text{ mol N m}^{-2}\text{yr}^{-1}$ at $\gamma = 27$. The diapycnal advection
 478 of nitrate, w^*N , on denser surfaces close to $\gamma = 27$, is negative, and so opposes the di-
 479 apycnal diffusive and eddy stirring transfers. The diapycnal advection of nitrate is weakly
 480 positive on denser surfaces, $\gamma = 27.5$, towards the bottom boundary. Hence, the ver-
 481 tical nitrate flux is controlled, by reinforcing contributions from diapycnal diffusion, di-
 482 apycnal advection and isopycnal eddy stirring at the base of the summer mixed layer,
 483 and by a loss from diapycnal advection and a supply from isopycnal eddy stirring be-
 484 low the seasonal boundary layer.

485 In comparison, off the ridge, there are smaller diapycnal diffusive nitrate fluxes, F_{dia} ,
 486 and smaller diapycnal advective fluxes, w^*N , due to the weaker tidal mixing (Tuerena
 487 et al., 2019). There is a more varied structure, with the vertical component of the isopy-
 488 cnal nitrate flux, F_{iso} , usually being comparable to or larger than both the diapycnal dif-
 489 fusive nitrate flux and the diapycnal advective flux, with a maximum value of $0.02 \text{ mol N m}^{-2}\text{yr}^{-1}$
 490 at $\gamma = 27.5$ (Fig. 9c,d).

491 In terms of the seasonality, in winter and into early spring, the nitrate supply to
 492 a density layer is likely to be dominated by the isopycnal nitrate transfer as winter mixed
 493 layers intersect denser surfaces and there are stronger nutrient gradients. In summer, the
 494 diapycnal nitrate transfer instead may dominate as there are weaker isopycnal nitrate
 495 gradients. The diapycnal advective flux is likely to increase in autumn and winter as the
 496 strong turbulence extends deeper into the water column and the depletion of surface nu-
 497 trients is reduced.

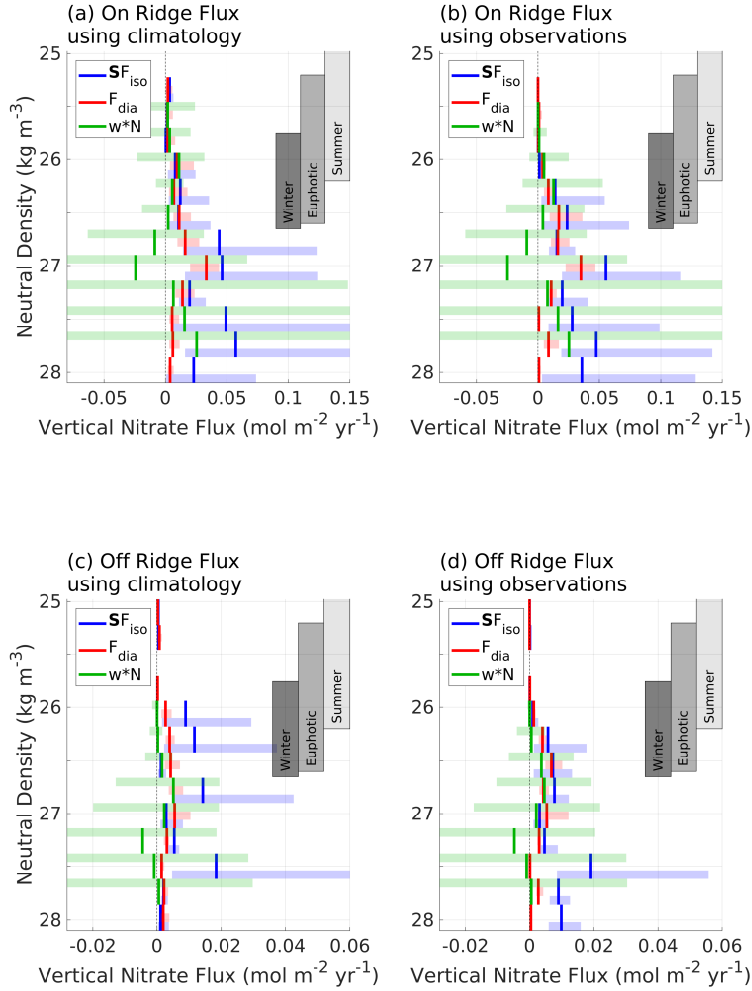


Figure 9. Profiles, in neutral density, of the diapycnal diffusion, diapycnal advection, and isopycnal diffusion contributions to the vertical nitrate flux for the on-ridge stations (a,b) and off-ridge stations (c,d), taken from microstructure-derived diffusivities and (a,c) the climatology-derived nutrient gradients or (b,d) the cruise-derived nutrient gradients. Blue shades indicate the isopycnal diffusion contributions, green shades indicate the diapycnal advection contributions, and red shades the diapycnal diffusion contributions. The shaded areas are the 95% confidence intervals derived from a bootstrap method. The grey bars indicate the range of neutral density surfaces in the cruise region representing, from left to right: the base of the summer mixed layer, the base of the summer euphotic zone, and the base of the winter mixed layer.

498 5.5 Vertical convergence of nitrate fluxes

499 Whether there is a nitrate supply to a density class is determined by the vertical
500 convergence of the vertical nitrate flux. On the ridge, there is generally a positive nitrate
501 supply from the sum of the diapycnal diffusion, diapycnal advection and isopycnal dif-
502 fusion of nitrate (Fig. 10a,b, black crosses).

503 Within the seasonal boundary layer extending over the summer and winter mixed
504 layers between $\gamma = 25.6$ and 26.4 , there is generally a reinforcing supply of nitrate from
505 the vertical convergence of diapycnal diffusive, diapycnal advective and isopycnal diffu-
506 sive components of the vertical nitrate flux (Fig. 10a,b, red, green and blue dots).

507 Below the winter mixed layer within the thermocline, there is instead a loss of ni-
508 trate from diapycnal advection, w^*N , between $\gamma = 26.8$ and 27.2 (Fig. 10a,b, green dots)
509 and a loss of nitrate from diapycnal mixing, F_{dia} , at $\gamma = 27.2$ (Fig. 10a,b, red dots). These
510 losses of nitrate are though offset by a supply of nitrate from the isopycnal diffusive com-
511 ponent, F_{iso} at $\gamma = 26.8$ to 27.2 (Fig. 10a,b, blue dots) and by diapycnal advection, w^*N ,
512 on denser surfaces.

513 Off the ridge, the overall nitrate supply is positive over most γ surfaces, but is neg-
514 ative for $\gamma = 27.2$ for the climatology, and for $\gamma = 26.9$ for the cruise section (Fig. 10c,d,
515 black crosses). Within the seasonal boundary layer for $\gamma = 25.4$ to 26.5 , there is again
516 a supply of nitrate from reinforcing contributions from the diapycnal diffusive and isopy-
517 cnal components of the vertical nitrate flux, F_{dia} and F_{iso} , and the convergence in the
518 diapycnal advection, w^*N (Fig. 10c,d, red, blue and green dots). Below the winter mixed
519 layer, for $\gamma = 27.2$ in the climatology and from $\gamma = 26.9$ to 27.3 in the cruise section, there
520 is instead a loss of nitrate from the convergences of the diapycnal nitrate flux, F_{dia} and
521 the diapycnal advection of nitrate, w^*N , which is partly offset by a nitrate supply by
522 the convergence of the isopycnal nitrate flux, F_{iso} .

523 Hence, the vertical nitrate supply changes from (i) a nitrate supply from reinforc-
524 ing diapycnal and isopycnal transfers within the seasonal boundary layer, to (ii) a loss
525 of nitrate below the seasonal boundary layer from either the diapycnal diffusion or di-
526 apycnal advection of nitrate, which is partly offset by an isopycnal diffusive supply in-
527 volving the stirring by mesoscale eddies.

528 6 Discussion and conclusions

529 There is a problem of understanding how export production is sustained in olig-
530 otrophic waters within the extensive downwelling zones of the subtropical gyres. In the
531 Sargasso Sea, tracer-based estimates of export production (Jenkins & Goldman, 1985;
532 Jenkins, 1988; Jenkins & Wallace, 1992; Stanley et al., 2015) exceed by a factor of two
533 or more the combined estimates from nitrate supply from entrainment, Ekman trans-
534 ports, atmospheric deposition and diapycnal mixing (McGillicuddy et al., 1998; Williams
535 & Follows, 2003). This conundrum might be solved by an enhanced physical supply of
536 nutrients through the effects of transient eddy and frontal upwelling of nutrients into the
537 euphotic zone (McGillicuddy Jr & Robinson, 1997; McGillicuddy et al., 1998; Lévy et
538 al., 2001, 2012) and regionally-enhanced diapycnal mixing over rough topography (Tuerena
539 et al., 2019). Both these possible physical solutions lead to a new issue, as the time-varying
540 upwelling and diapycnal supply will cease to be effective when the nutrient concentra-
541 tions in the upper thermocline become eroded (Oschlies, 2002; Williams & Follows, 2003).
542 The resolution of this conundrum might involve the lateral supply of nutrients to the up-
543 per thermocline, thereby sustaining nutrient concentrations below the euphotic zone.

544 The lateral supply of nutrients may be achieved through the time-mean geostrophic
545 flow, although this transfer becomes weak at the gyre boundaries as geostrophic stream-
546 lines align with nutrient contours along density surfaces. The Ekman-induced horizon-

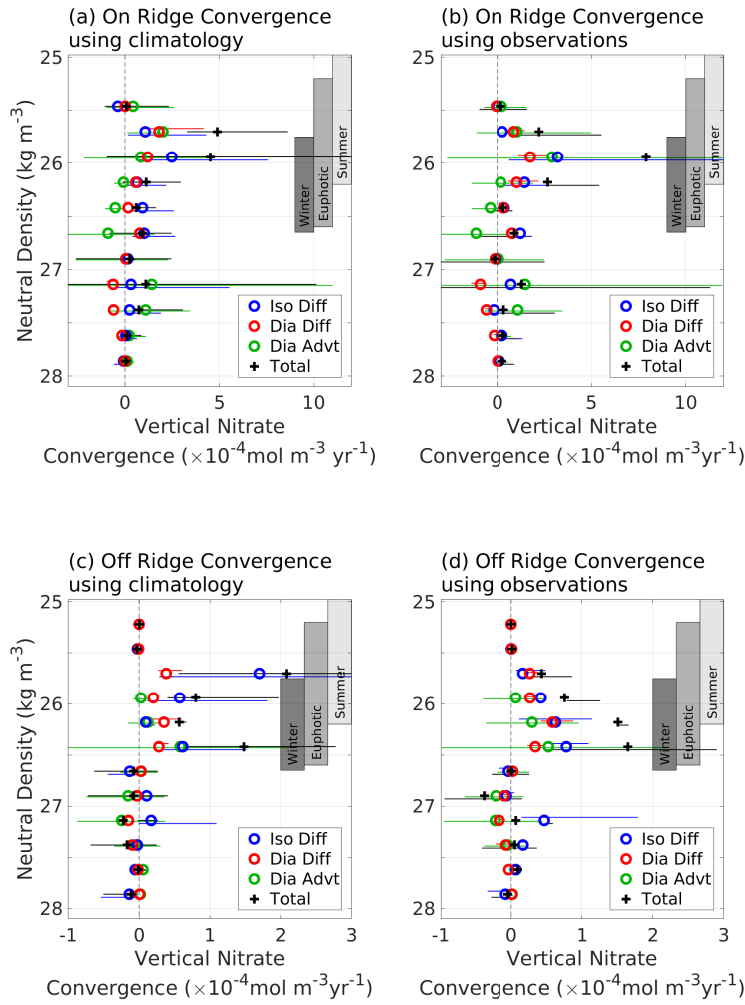


Figure 10. Profiles, in neutral density, of the diapycnal diffusion, diapycnal advection and isopycnal diffusion contributions to the vertical nitrate convergence for the on-ridge stations (a,b) and off-ridge stations (c,d), taken from the microstructure-derived diffusivities and (a,c) the climatology-derived nutrient gradients or (b,d) the cruise-derived nutrient gradients. Blue shades indicate the isopycnal diffusion contributions, green shades indicate the diapycnal advection contributions, and the red shades the diapycnal diffusion contributions. The horizontal lines indicate the 95% confidence intervals derived from a bootstrap method. The grey bars show the range of neutral density surfaces in the cruise region representing, from left to right: the base of the summer mixed layer, the base of the summer euphotic zone, and the base of the winter mixed layer. Convergences of vertical nitrate fluxes are calculated using the median separation of the neutral surfaces over the relevant subset of data, with fluxes that have been smoothed using a 3-span moving average.

547 tal transport of nutrients does provide an effective supply across time-mean geostrophic
 548 streamlines along the flanks of the subtropical gyres (Williams & Follows, 1998), but this
 549 supply becomes weak in the upper thermocline and towards the central parts of the gyre.
 550 There may also be a transfer of nutrients across the time-mean geostrophic streamlines
 551 via a combination of eddy-induced advection and diffusion along density surfaces, as il-
 552 lustrated in idealised eddy-resolving model studies (Lee & Williams, 2000; Lévy, 2008).

553 In this field-based investigation, which targeted some of the most oligotrophic wa-
 554 ters of the North Atlantic subtropical gyre, the relative importance of diapycnal mix-
 555 ing, diapycnal advection and eddy stirring in supplying nutrients is assessed over the sea-
 556 sonal boundary layer and the thermocline. This region is far from the complicating ef-
 557 fects of boundary current flows or areas of strong air-sea interaction, but does lie over
 558 the mid-Atlantic ridge, where there is enhanced diapycnal mixing from the spring-neap
 559 cycle of the tides interacting with the topography (Vic et al., 2018; Tuerena et al., 2019).
 560 In our data analysis, we find that the combination of diapycnal diffusion and diapycnal
 561 advection leads to a supply of nutrients to the euphotic zone, but at the same time to
 562 a loss of nutrients in the upper thermocline. Isopycnal diffusion from the stirring by mesoscale
 563 eddies augment the diapycnal supply of nitrate to the lighter density surfaces intersect-
 564 ing the euphotic zone. This eddy stirring crucially provides nutrients to the upper ther-
 565 mocline and so acts to offset the loss of nutrients associated with the diapycnal trans-
 566 fer. Thus, the eddy-induced lateral transfer of nutrients along density surface may be
 567 part of the solution to the long-standing question of how the supply of nutrients to the
 568 euphotic zone is sustained.

569 While our study reveals clear signals of how the eddy stirring acts to augment and
 570 partly sustain the diapycnal supply of nutrients, our combined estimates of diapycnal
 571 mixing and eddy stirring-driven supply of nitrate at 300 m only reaches 0.03 and 0.05 mol N m⁻² yr⁻¹
 572 on the ridge for climatological and observed nutrients. Our estimates are thus smaller
 573 by a factor of 3 to 4 than estimates of nitrate export of 0.17 mol N m⁻²yr⁻¹ from oxy-
 574 gen utilisation rates (Jenkins, 1987) at a nearby 1000 km triangular site centered at 26.5°N ,
 575 32°W . Hence, there is a possibility that the eddy stirring is only providing a modest, back-
 576 ground contribution to the nutrient supply to the euphotic zone and upper thermocline.

577 However, our estimates of the nutrient supply role of eddy stirring are based on a
 578 field programme at the centre of a subtropical gyre and the eddy stirring is expected to
 579 provide a larger contribution toward the flanks of the gyre. The eddy stirring supply of
 580 nitrate, evaluated from the vertical component of the convergence of the diffusive isopyc-
 581 nal nitrate flux, $(\mathbf{S}/\bar{h}) \cdot \partial/\partial z (\bar{h} \kappa_{iso} \nabla_\sigma \bar{N})$ in (18), is proportional to the product of the
 582 isopycnal diffusivity, κ_{iso} , the isopycnal nitrate gradient, $\nabla_\sigma \bar{N}$ and the slope of the den-
 583 sity surfaces, \mathbf{S} . Both the nitrate gradient and isopycnal slope are relatively low over our
 584 field site in the centre of the subtropical gyre (Fig. 11a,b), as revealed by the climato-
 585 logical distributions of nitrate and density during summer for the $\gamma = 26.5$ surface, which
 586 spans depths of 50 m to 300 m (Garcia et al., 2013). Both of these contributions to the
 587 vertical nitrate flux are amplified by a factor of at least 10 on the boundaries of the sub-
 588 tropical gyre (Fig. 11c,d). The lateral diffusivity associated with eddy stirring is though
 589 inhibited toward the surface by the presence of strong flows (Ferrari & Nikurashin, 2010;
 590 Groeskamp et al., 2020), which leads to a reduction in κ_{iso} from 500 to 1000 m²s⁻¹ (Groeskamp
 591 et al., 2020) in the gyre interior, to 200 m²s⁻¹ over the Gulf Stream (Bower et al., 1985;
 592 Abernathey & Marshall, 2013).

593 While our field study has by necessity focussed on the vertical component of the
 594 mesoscale eddy-induced diffusive transfer of nutrients, eddies also provide a lateral ad-
 595 vective transfer of nutrients, which may be important in transferring nutrients across gyre
 596 boundaries (Lee & Williams, 2000; Williams & Follows, 2003) and augment the lateral
 597 nutrient supply by the horizontal Ekman transport (Williams & Follows, 1998; Letscher
 598 et al., 2016).

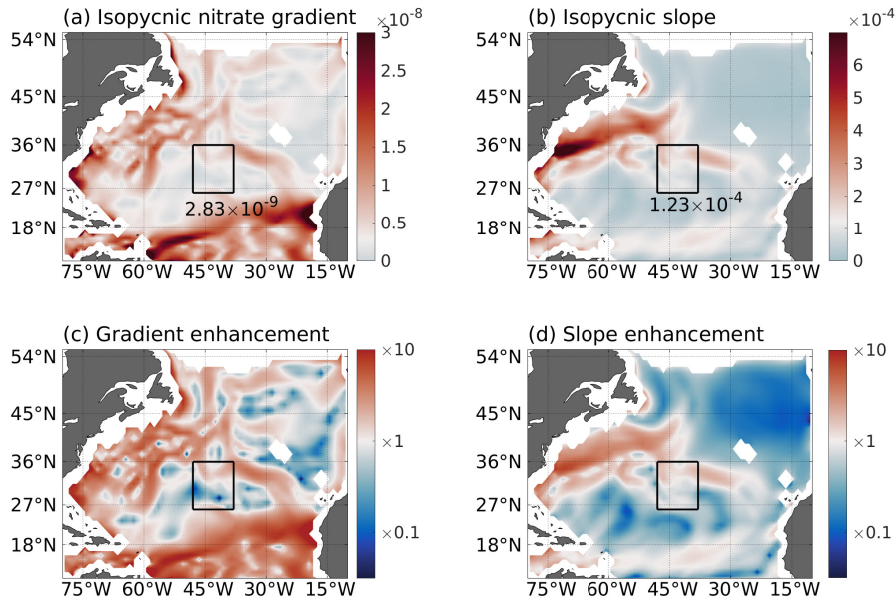


Figure 11. Maps showing (a) the magnitude of the isopycnal gradients of nitrate, $|\nabla_{\sigma} N|$ (mol m^{-4}), (b) the slope of the isopycnal surfaces, \mathbf{S} , and in (c) and (d), the enhancement relative to the study region for (a) and (b). The nitrate and density fields are taken from the summer World Ocean Atlas climatology along the $\gamma = 26.5$ surface. The black box represents the region of our observations and the numbers are the average within that box.

599 In summary, the nutrient supply to the euphotic zone in the centre of the oligotrophic
 600 subtropical gyre may be achieved via a multi-stage mechanism: a diapycnal transfer of
 601 nutrients by microscale turbulence to the euphotic zone from the upper thermocline, and
 602 an isopycnal transfer of nutrients by mesoscale eddies acting to both augment the sup-
 603 ply to the euphotic zone and replenish upper thermocline nutrients. The wider gener-
 604 ality of our central result is suggested by the similar multi-stage process uncovered in
 605 a modelling study of the Southern Ocean supply of trace metals (Uchida et al., 2020),
 606 whereby eddy stirring was found to replenish the thermocline and diapycnal mixing to
 607 provide the transfer to the euphotic zone.

608 Appendix A Estimates of diffusivity from station data

609 Following Naveira Garabato et al. (2016), the production and the dissipation of the
 610 temperature variance is estimated from the CTD/VMP stations, allowing the isopycnal
 611 diffusivity to be diagnosed as a residual,

$$\kappa_{iso} = \frac{\langle \chi \rangle / 2 - \Gamma \epsilon N^{-2} \left| \frac{\partial \theta_m}{\partial z} \right|^2}{|\nabla_{\sigma} \theta_m|^2}. \quad (\text{A1})$$

612 The dissipation of the temperature variance for a single station usually exceeds the pro-
 613 duction term from the microscale turbulence (Fig. A1c), taken from the combination of
 614 the diapycnal diffusivity and temperature gradients (Fig. A1a,b). With the addition of
 615 the isopycnal temperature gradients (Fig. A1d), the diffusivity for a single station can
 616 then be calculated (Fig. A1e).

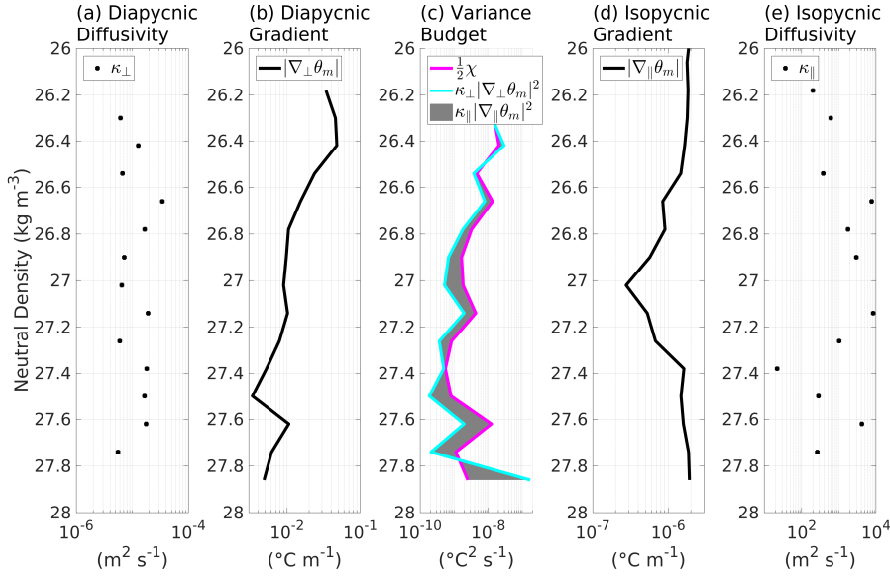


Figure A1. Profiles at a single station in neutral density space of: (a) the diapycnal diffusivity; (d) absolute temperature gradients across density surfaces; (c) temperature variance dissipation (cyan line) and temperature variance production by diapycnal mixing (magenta line) with the shaded area indicating the production by isopycnal stirring; (d) absolute temperature gradients along density surfaces; and (e) the diffusivity along density surfaces.

Appendix B Calculation of isopycnal slopes and gradients

Isopycnal gradients of potential temperature and nutrient are taken both from observations and from a climatology. The methodology is set out due to complexities of evaluating these gradients on isopycnal or neutral surfaces (Groeskamp, Barker, et al., 2019).

For the observations, each cast is linearly interpolated in the vertical onto a consistent neutral density grid (Jackett & McDougall, 1997) with a spacing of 0.12 kg m^{-3} , such that each cast has a profile of depth, potential temperature and nitrate in neutral density space. This interpolation avoids making a local assumption in calculating gradients from the combination of horizontal and vertical gradients. All subsequent calculations are then performed on the same neutral density grid. For each value of neutral density, a flat surface (linear in x and y directions) is fitted using distance north and east from a fixed position for each variable (Fig. B1). The zonal and meridional gradients in depth, potential temperature and nitrate, are then extracted from these fitted surfaces. These surfaces effectively smooth over the local gradients driven by mesoscale processes and provides gradients representative of the size of the survey of the order of 1000 km.

For the climatology, individual profiles are again remapped onto a consistent neutral density grid using the routines of (Jackett & McDougall, 1997). It is assumed that in the construction of the climatology, the gradients associated with mesoscale features are already removed, so that local gradients are used. The gradients are taken on the same grid as the climatology using centered differencing, such that the resulting gradients are taken over twice the horizontal spacing of the climatology resolution. These gradients are then applied to the observations using the closest position in the climatology grid.

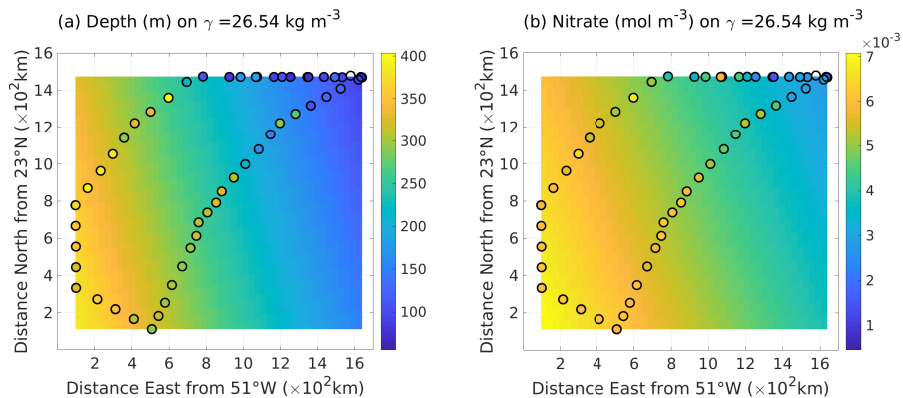


Figure B1. Maps showing the fitted surfaces of depth (slope) and nitrate for the surface $\gamma = 26.54 \text{ kg m}^{-3}$. The observations, interpolated onto the density surface, are shown by the circles. The background colour shows the fitted surface from which the gradients are extracted.

641 Acknowledgments

642 RidgeMix was supported by the U.K. Natural Environment Research Council (NE/L004216/1).
 643 We are grateful to the technicians, officers, and crew of RRS James Clark Ross for their
 644 invaluable role in data collection. The hydrographic, turbulence and inorganic nutrient
 645 data are freely available via the British Oceanographic Data Centre: Vic et al. (2018)
 646 has published the hydrographic and turbulence data, and the nutrient data is available
 647 in Tuerena R. and Mahaffey C. (2021), doi: 10.5285/b9bd0df2-f8de-21fe-e053-6c86abc0cb35.
 648 We thank Claire Mahaffey and Clare Davis for support with biogeochemical measure-
 649 ments and Malcolm Woodward for use of the nutrient analyzer at sea. We are grateful
 650 to Trevor McDougall for pointing out the importance of the diapycnal advection term,
 651 which had been omitted in a previous version of this work, and to Ryan Abernathey and
 652 an anonymous referee for positive comments, which all have strengthened the study.

653 References

- 654 Abernathey, R. P., & Marshall, J. (2013). Global surface eddy diffusivities derived
 655 from satellite altimetry. *Journal of Geophysical Research: Oceans*, *118*, 901–
 656 916.
- 657 Bleck, R. (1998). Ocean modeling in isopycnic coordinates. In *Ocean modeling and
 658 parameterization* (pp. 423–448). Springer.
- 659 Bleck, R., & Boudra, D. B. (1981). Initial testing of a numerical ocean circulation
 660 model using a hybrid(quasi-isopycnic) vertical coordinate. *Journal of Physical
 661 Oceanography*, *11*(6), 755–770.
- 662 Bower, A. S., Rossby, H. T., & Lillibridge, J. L. (1985). The Gulf Stream—barrier or
 663 blender? *Journal of Physical Oceanography*, *15*, 24 - 32.
- 664 Canuto, V., & Dubovikov, M. (2011). Comparison of four mixed layer mesoscale
 665 parameterizations and the equation for an arbitrary tracer. *Ocean Modelling*,
 666 *39*(1-2), 200–207.
- 667 Cole, S. T., Wortham, C., Kunze, E., & Owens, W. B. (2015). Eddy stirring and
 668 horizontal diffusivity from Argo float observations: Geographic and depth

- 669 variability. *Geophysical Research Letters*, *42*, 3989–3997.
- 670 de Lavergne, C., Madec, G., Le Sommer, J., Nurser, A. G., & Garabato, A. C. N.
671 (2016). On the consumption of Antarctic bottom water in the abyssal ocean.
672 *Journal of Physical Oceanography*, *46*, 635 – 661.
- 673 Dietze, H., Oschlies, A., & Kähler, P. (2004). Internal-wave-induced and double-
674 diffusive nutrient fluxes to the nutrient-consuming surface layer in the oligo-
675 trophic subtropical North Atlantic. *Ocean Dynamics*, *54*(1), 1–7.
- 676 Ferrari, R., & Nikurashin, M. (2010). Suppression of eddy diffusivity across jets in
677 the Southern Ocean. *Journal of Physical Oceanography*, *40*, 1501 - 1519.
- 678 Ferrari, R., & Polzin, K. L. (2005). Finescale structure of the T–S relation in the
679 eastern North Atlantic. *Journal of Physical Oceanography*, *35*(8), 1437–1454.
- 680 Garcia, H. E., Locarnini, R. A., Boyer, T. P., Antonov, J. I., Baranova, O. K.,
681 Zweng, M. M., . . . Johnson, D. R. (2013). World Ocean Atlas 2013 volume
682 4: Dissolved inorganic nutrients (phosphate, nitrate, silicate). In S. Levitus &
683 A. V. Mishonov (Eds.), .
- 684 Gent, P. R., & McWilliams, J. C. (1990). Isopycnal mixing in ocean circulation
685 models. *Journal of Physical Oceanography*, *20*(1), 150–155.
- 686 Gent, P. R., Willebrand, J., McDougall, T. J., & McWilliams, J. C. (1995). Parame-
687 terizing eddy-induced tracer transports in ocean circulation models. *Journal of*
688 *Physical Oceanography*, *25*(4), 463–474.
- 689 Gregg, M., D’Asaro, E., Riley, J., & Kunze, E. (2018). Mixing efficiency in the
690 ocean. *Annual Review of Marine Science*, *10*, 443–473.
- 691 Groeskamp, S., Barker, P. M., McDougall, T. J., Abernathey, R. P., & Griffies,
692 S. M. (2019). Venn: An algorithm to accurately calculate neutral slopes and
693 gradients. *Journal of Advances in Modeling Earth Systems*, *11*, 1917–1939.
- 694 Groeskamp, S., Griffies, S. M., Iudicone, D., Marsh, R., Nurser, A. G., & Zika, J. D.
695 (2019). The water mass transformation framework for ocean physics and
696 biogeochemistry. *Annual review of marine science*, *11*, 271–305.
- 697 Groeskamp, S., LaCasce, J. H., McDougall, T. J., & Rogé, M. (2020). Full-depth
698 global estimates of ocean mesoscale eddy mixing from observations and theory.
699 *Geophysical Research Letters*, *47*(18), e2020GL089425.
- 700 Jackett, D. R., & McDougall, T. J. (1997). A neutral density variable for the
701 World’s Oceans. *Journal of Physical Oceanography*, *27*, 237 – 263.
- 702 Jenkins, W. (1987). ^3H and ^3He in the Beta Triangle: Observations of gyre venti-
703 lation and oxygen utilization rates. *Journal of Physical Oceanography*, *17*(6),
704 763–783.
- 705 Jenkins, W. (1988). Nitrate flux into the euphotic zone near Bermuda. *Nature*,
706 *331*(6156), 521–523.
- 707 Jenkins, W., & Goldman, J. (1985). Seasonal oxygen cycling and primary produc-
708 tion in the Sargasso Sea. *Journal of Marine Research*, *43*(2), 465–491.
- 709 Jenkins, W., & Wallace, D. (1992). Tracer based inferences of new primary produc-
710 tion in the sea. In *Primary productivity and biogeochemical cycles in the sea*
711 (pp. 299–316). Springer.
- 712 Joyce, T. M., Luyten, J. R., Kubryakov, A., Bahr, F. B., & Pallant, J. S. (1998).
713 Meso- to large-scale structure of subducting water in the subtropical gyre of
714 the eastern North Atlantic Ocean. *Journal of Physical Oceanography*, *28*,
715 40–61.
- 716 Klocker, A., & Abernathey, R. (2014). Global patterns of mesoscale eddy properties
717 and diffusivities. *Journal of Physical Oceanography*, *44*(3), 1030–1046.
- 718 Klocker, A., & McDougall, T. J. (2010). Influence of the nonlinear equation of state
719 on global estimates of dianeutral advection and diffusion. *Journal of Physical*
720 *Oceanography*, *40*(8), 1690–1709.
- 721 Knap, A., Jickells, T., Pszenny, A., & Galloway, J. (1986). Significance of
722 atmospheric-derived fixed nitrogen on productivity of the Sargasso Sea. *Na-*
723 *nature*, *320*(6058), 158–160.

- 724 Kunze, E., Firing, E., Hummon, J. M., Chereskin, T. K., & Thurnherr, A. M.
725 (2006). Global abyssal mixing inferred from lowered ADCP shear and CTD
726 strain profiles. *Journal of Physical Oceanography*, *7*, 1553–1576.
- 727 Ledwell, J. R., Watson, A. J., & Law, C. S. (1998). Mixing of a tracer in the pycno-
728 cline. *Journal of Geophysical Research*, *103*, 21499–21529.
- 729 Lee, M.-M., Marshall, D. P., & Williams, R. G. (1997). On the eddy transfer of tracers:
730 Advective or diffusive? *Journal of Marine Research*, *55*(3), 483–505.
- 731 Lee, M.-M., & Williams, R. G. (2000). The role of eddies in the isopycnal transfer
732 of nutrients and their impact on biological production. *Journal of Marine Re-
733 search*, *58*(6), 895–917.
- 734 Letscher, R. T., Primeau, F., & Moore, J. K. (2016). Nutrient budgets in the sub-
735 tropical ocean gyres dominated by lateral transport. *Nature Geoscience*, *9*(11),
736 815–819.
- 737 Lévy, M. (2008). The modulation of biological production by oceanic mesoscale tur-
738 bulance. *Transport and mixing in geophysical flows*, 219–261.
- 739 Lévy, M., Ferrari, R., Franks, P. J., Martin, A. P., & Rivière, P. (2012). Bringing
740 physics to life at the submesoscale. *Geophysical Research Letters*, *39*(14).
- 741 Lévy, M., Klein, P., & Treguier, A.-M. (2001). Impact of sub-mesoscale physics on
742 production and subduction of phytoplankton in an oligotrophic regime. *Jour-
743 nal of marine research*, *59*(4), 535–565.
- 744 Lewis, M. R., Hebert, D., Harrison, W. G., Platt, T., & Oakey, N. S. (1986). Verti-
745 cal nitrate fluxes in the oligotrophic ocean. *Science*, *234*(4778), 870–873.
- 746 Locarnini, R. A., Mishonov, A. V., Antonov, J. I., Boyer, T. P., Garcia, H. E., Bara-
747 nova, O. K., ... Seidov, D. (2013). World Ocean Atlas 2013 volume 1: Tem-
748 perature. In S. Levitus & A. V. Mishonov (Eds.), .
- 749 McDougall, T. J. (1984). The relative roles of diapycnal and isopycnal mixing on
750 subsurface water mass conversion. *Journal of Physical Oceanography*, *14*(10),
751 1577–1589.
- 752 McDougall, T. J. (1987). Thermobaricity, cabbeling, and water-mass conversion.
753 *Journal of Geophysical Research: Oceans*, *92*(C5), 5448–5464.
- 754 McGillicuddy, D., Robinson, A., Siegel, D., Jannasch, H., Johnson, R., Dickey, T.,
755 ... Knap, A. (1998). New evidence for the impact of mesoscale eddies on
756 biogeochemical cycling in the Sargasso Sea. *Nature*, *394*, 263–266.
- 757 McGillicuddy Jr, D., Anderson, L., Doney, S., & Maltrud, M. (2003). Eddy-driven
758 sources and sinks of nutrients in the upper ocean: Results from a 0.1 resolution
759 model of the North Atlantic. *Global Biogeochemical Cycles*, *17*(2).
- 760 McGillicuddy Jr, D., & Robinson, A. (1997). Eddy-induced nutrient supply and new
761 production in the Sargasso Sea. *Deep Sea Research Part I: Oceanographic Re-
762 search Papers*, *44*(8), 1427–1450.
- 763 Michaels, A. F., Knap, A. H., Dow, R. L., Gundersen, K., Johnson, R. J., Sorensen,
764 J., ... others (1994). Seasonal patterns of ocean biogeochemistry at the US
765 JGOFS Bermuda Atlantic Time-series Study site. *Deep Sea Research Part I:
766 Oceanographic Research Papers*, *41*(7), 1013–1038.
- 767 Naveira Garabato, A. C., Polzin, K. L., Ferrari, R., & Forryan, J. D. Z. A. (2016).
768 A microscale view of mixing and overturning across the Antarctic Circumpolar
769 Current. *Journal of Physical Oceanography*, *46*, 233 - 254.
- 770 Naveira Garabato, A. C., Ferrari, R., & Polzin, K. L. (2011). Eddy stirring in the
771 Southern Ocean. *Journal of Geophysical Research: Oceans*, *116*(C9).
- 772 Nurser, A., Marsh, R., & Williams, R. G. (1999). Diagnosing water mass forma-
773 tion from air–sea fluxes and surface mixing. *Journal of Physical Oceanography*,
774 *29*(7), 1468–1487.
- 775 Oakey, N. S. (1982). Determination of the rate of dissipation of turbulent energy
776 from simultaneous temperature and velocity shear microstructure measure-
777 ments. *Journal of Physical Oceanography*, *12*, 256 – 271.

- 778 Osborn, T. R. (1980). Estimates of the local rate of vertical diffusion from dissipa-
779 tion measurements. *Journal of Physical Oceanography*, *10*, 83 - 89.
- 780 Oschlies, A. (2002). Can eddies make ocean deserts bloom? *Global Biogeochemical*
781 *Cycles*, *16*(4), 53–1.
- 782 Redi, M. H. (1982). Oceanic isopycnal mixing by coordinate rotation. *Journal of*
783 *Physical Oceanography*, *12*(10), 1154–1158.
- 784 Resplandy, L., Lévy, M., Madec, G., Pous, S., Aumont, O., & Kumar, D. (2011).
785 Contribution of mesoscale processes to nutrient budgets in the arabian sea.
786 *Journal of Geophysical Research: Oceans*, *116*(C11).
- 787 Roach, C. J., Balwada, D., & Speer, K. (2018). Global observations of horizontal
788 mixing from argo float and surface drifter trajectories. *Journal of Geophysical*
789 *Research: Oceans*, *123*, 4560–4575.
- 790 Stanley, R., Jenkins, W., Doney, S., & III, D. L. (2015). The ^3He flux gauge in the
791 Sargasso Sea: a determination of physical nutrient fluxes to the euphotic zone
792 at the Bermuda Atlantic Time-series Site. *Biogeosciences*, *12*(17).
- 793 Tennekes, H., & Lumley, J. L. (2018). *A first course in turbulence*. MIT press.
- 794 Tuerena, R. E., Williams, R. G., Mahaffey, C., Vic, C., Green, J. A. M., Naveira-
795 Garabato, A., ... Sharples, J. (2019). Internal tides drive nutrient fluxes into
796 the deep chlorophyll maximum over mid-ocean ridges. *Global Biogeochemical*
797 *Cycles*, *33*, 995–1009.
- 798 Tulloch, R., Marshall, J., & Smith, K. S. (2009). Interpretation of the propagation
799 of surface altimetric observations in terms of planetary waves and geostrophic
800 turbulence. *Journal of Geophysical Research: Oceans*, *114*, C02005.
- 801 Uchida, T., Balwada, D., Abernathey, R. P., McKinley, G. A., Smith, S. K., & Lévy,
802 M. (2020). Vertical eddy iron fluxes support primary production in the open
803 Southern Ocean. *Nature Communications*, *11*(1), 1–8.
- 804 Vic, C., Garabato, A. C. N., Green, J. A. M., Spingys, C., Forryan, A., Zhao, Z.,
805 & Sharples, J. (2018). The lifecycle of semidiurnal internal tides over the
806 northern mid-Atlantic ridge. *Journal of Physical Oceanography*, *48*, 61–80.
- 807 Waterhouse, A. F., MacKinnon, J. A., Nash, J. D., Alford, M. H., Kunze, E., Sim-
808 mons, H. L., ... Lee, C. M. (2014). Global patterns of diapycnal mixing from
809 measurements of the turbulent dissipation rate. *Journal of Physical Oceanogra-*
810 *phy*, *44*, 1854–1872.
- 811 Williams, R. G., & Follows, M. J. (1998). The Ekman transfer of nutrients and
812 maintenance of new production over the North Atlantic. *Deep Sea Research*
813 *Part I: Oceanographic Research Papers*, *45*(2-3), 461–489.
- 814 Williams, R. G., & Follows, M. J. (2003). Physical transport of nutrients and the
815 maintenance of biological production. In *Ocean biogeochemistry* (pp. 19–51).
816 Springer.
- 817 Zika, J. D., & McDougall, T. J. (2008). Vertical and lateral mixing processes de-
818 duced from the mediterranean water signature in the North Atlantic. *Journal*
819 *of Physical Oceanography*, *38*, 164 – 176.
- 820 Zika, J. D., McDougall, T. J., & Sloyan, B. M. (2010). Weak mixing in the eastern
821 North Atlantic: An application of the tracer-contour inverse method. *Journal*
822 *of Physical Oceanography*, *40*, 1881 – 1893.
- 823 Zweng, M. M., Reagan, J. R., Antonov, J. I., Locarnini, R. A., Mishonov, A. V.,
824 Boyer, T. P., ... Biddle, M. M. (2013). World Ocean Atlas 2013 volume 2:
825 Salinity. In S. Levitus & A. V. Mishonov (Eds.), .