

Abstract

In stratified shelf seas, oxygen depletion beneath the thermocline is a result of a greater rate of biological oxygen demand than the rate of supply of oxygenated water. Suitably equipped gliders are uniquely placed to observe both the supply through the thermocline and the consumption of oxygen in the bottom layers. A Seaglider was deployed in the shallow (≈ 100 m) stratified North Sea in a region of known low oxygen during August 2011 to investigate the processes regulating supply and consumption of dissolved oxygen below the pycnocline. The first deployment of such a device in this area, it provided extremely high resolution observations, 316 profiles (every 16 min, vertical resolution of 1 m) of CTD, dissolved oxygen concentrations, backscatter and fluorescence during a three day deployment.

The high temporal resolution observations revealed occasional small scale events that supply oxygenated water into the bottom layer at a rate of $2 \pm 1 \mu\text{mol dm}^{-3} \text{day}^{-1}$. Benthic and pelagic oxygen sinks, quantified through glider observations and past studies, indicate more gradual background consumption rates of $2.5 \pm 1 \mu\text{mol dm}^{-3} \text{day}^{-1}$. This budget revealed that the balance of oxygen supply and demand is in agreement with previous studies of the North Sea. However, the glider data show a net oxygen consumption rate of $2.8 \pm 0.3 \mu\text{mol dm}^{-3} \text{day}^{-1}$ indicating a localised or short-lived increase in oxygen consumption rates. This high rate of oxygen consumption is indicative of an unidentified oxygen sink. We propose that this elevated oxygen consumption is linked to localised depocentres and rapid remineralisation of resuspended organic matter.

The glider proved to be an excellent tool for monitoring shelf sea processes despite challenges to glider flight posed by high tidal velocities, shallow bathymetry, and very strong density gradients. The direct observation of these processes allows more up to date rates to be used in the development of ecosystem models.

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1 Introduction

Mooring observations from 2007 and 2008 (Greenwood et al., 2010), historical data and a hydrographic survey of the North Sea in August 2010 (Queste et al., 2013) revealed repeated incidents of seasonal oxygen depletion in offshore waters of the central North Sea. In August 2011, a Seaglider was deployed in the region with the lowest recorded bottom mixed layer (BML) oxygen saturation from the 2010 survey to further investigate the mechanisms causing this seasonal oxygen depletion.

The North Sea is a relatively shallow (15–200 m) shelf sea situated between the British Isles and northwestern continental Europe. It gradually deepens from south to north with the exception of the shallow Dogger Bank (Fig. 1). Dogger Bank effectively separates the North Sea into two regions of different physical, chemical and biological properties (Otto et al., 1990; Lenhart et al., 1997). Water properties in the northern half of the North Sea are largely dominated by North Atlantic inflow. For their study of carbon dynamics, Thomas et al. (2005) determine the waters of the North Sea to be 90 % sourced from the North Atlantic (9000 km³ yr⁻¹ via the Pentland Firth and Fair Isle, 42 000 km³ yr⁻¹ via the Shetlands), 8 % from the English Channel (4900 km³ yr⁻¹), and 2 % from a combination of Baltic (500 km³ yr⁻¹) and riverine flow (200 km³ yr⁻¹). Water entering the North Sea from the northern boundary generally circulates anticlockwise, following the Scottish coast southward before turning east and crossing the North Sea north of Dogger Bank (Brown et al., 1999; Turrell et al., 1992; Hill et al., 2008). The strength of this anticlockwise circulation is strongly correlated with the North Atlantic Oscillation Index (NAOI); positive NAOI is associated with strong anticlockwise circulation while negative NAOI is associated with greatly reduced anticlockwise circulation in the northern North Sea (Lenhart et al., 1995, 2004; Rodwell et al., 1999). The northern half of the North Sea is seasonally stratified through surface heating. This stratification breaks down completely in winter (Ducrottoy et al., 2000; Sharples et al., 2006; Greenwood et al., 2010). Despite representing only 2 % of the inflow, Baltic waters can play a disproportionate role in defining the biochemical properties of the central North Sea

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as under specific conditions of circulation or surface wind stress, the fresh water run-off issuing from the Baltic, Kattegat and Skagerrak can be advected into the central region (Otto et al., 1990). This not only affects stratification by increasing the salinity difference between surface mixed layer (SML) and BML but also provides supplementary nutrients to the SML.

The August 2010 North Sea survey (Queste et al., 2013) recorded bottom waters with oxygen saturations below 70 % in two regions; north of Dogger Bank (ND) and the Oyster Grounds (OG) (Fig. 2) exhibited dissolved oxygen saturations circa 65 % ($200 \mu\text{mol dm}^{-3}$) and 68 % ($180 \mu\text{mol dm}^{-3}$) respectively. Historical data revealed a similar distribution of low dissolved oxygen (DO) in the BML during summer in the ND and OG regions. The historical data also showed a sharp decline in summer oxygen saturations at these two sites since 1990. Queste et al. (2013) suggested that the same mechanisms likely lead to the depletion of oxygen at OG and ND. It is thought that the replenishment of oxygenated waters through advective processes is limited by local topography. ND is situated away from the fronts associated with the north side of the shallow Dogger Bank; it is characterised by variable, weak wind-driven currents (Otto et al., 1990). This slow circulation limits the horizontal supply of DO while weak tidal currents also promote settling of organic matter. Weak winds and strong surface heat fluxes promote stratification and keep the BML isolated from air-sea exchange. There is also weak inflow of oxygenated waters as tides and topography lead to the formation of bottom fronts to the south and east of the OG site (Van Raaphorst et al., 1998; Weston et al., 2008). Organic matter produced at the pycnocline along the deep chlorophyll maximum (DCM), particularly at ND, is exported directly into the BML (Thomas et al., 2004; Weston et al., 2008; Fernand et al., 2013). This organic matter, when remineralised, leads to the consumption of DO in the BML. At both the OG and ND sites, it has been suggested that mixing events may cause resuspension of bottom sediment. Transfer of this organically rich sediment into an oxic water column could cause rapid increases in pelagic DO consumption (Van Raaphorst et al., 1998; Greenwood et al., 2010).

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The intensity and duration of oxygen depletion depend on the relative magnitudes of oxygen consumption and oxygen supply. To accurately predict potential seasonal oxygen consumption under future climate scenarios, it is necessary to identify the relative magnitude of sources of organic matter that lead to the consumption of DO in the BML and the amount of DO supply through the pycnocline on a finer scale. There is growing interest in the issue of low oxygen in coastal waters around the world (Diaz and Rosenberg, 2008) and further research is required to effectively manage the impact of hypoxia in coastal waters. This study aims to improve our understanding of the key short term mixing processes that regulate oxygen supply at high temporal frequency by quantifying through Seaglider observations the major inputs and sinks of dissolved oxygen.

2 Seaglider observations

A Seaglider from the University of East Anglia, SG510, was deployed at 56°41.96' N, 2°26.37' E on the 19 August 2011 (Fig. 1) in a water depth of ≈ 75 m. Seagliders are buoyancy-driven autonomous underwater vehicles that move through the water in a sawtooth pattern between the surface and the seabed (or 1000 m in deeper water; Eriksen et al., 2001). Typically, the glider recorded data between 2 m from the surface and 7 m off the seabed; sample points beyond this range were discarded because of poor flow conditions due to the glider turning around. Seagliders sample every 5 s using an on-board sensor suite composed of a Seabird unpumped conductivity and temperature sensor, a Wetlabs Triplet EcoPuck and an Aanderaa 4330F optode. SG510 sampled along a 32 km northward transect during a 3 day period (158 dives, 316 vertical profiles) providing a full depth cast every 16 min.

Temperature and salinity were calibrated against an in situ ship-based profile at launch; dissolved oxygen and chlorophyll *a* were calculated using recent manufacturer calibrations. Data were corrected for sensor lag and time offsets; further details are provided by Queste (2014). Due to the sharp temperature gradient, significant hystere-

Apparent oxygen utilisation (AOU, defined as the difference between the solubility of dissolved oxygen and observed oxygen concentration; Fig. 3f) shows a pattern typical of oxygen change within the central stratified North Sea (Queste et al., 2013). The SML is homogeneously saturated. At the pycnocline, supersaturation (negative AOU) is visible and correlated with elevated chlorophyll *a* concentration (Fig. 4f); this supersaturation is evidence of strong primary production occurring at the DCM (Weston et al., 2005). The BML exhibits increasing AOU throughout the survey; this oxygen depletion is caused by the remineralisation of organic matter within the BML and isolation from air–sea exchange by the strong density gradient.

3 Dissolved oxygen budget

To constrain the balance of DO consumption and supply, we quantify the various sources and sinks of DO and compare these with the observed change in DO over the survey period. Pena et al. (2010) review the processes that affect DO (Eq. 1). Advective-dispersive transport, air–sea exchange, and biological production/consumption are the three dominant processes which can be described mathematically as:

$$\frac{\delta O_2}{\delta t} = \mathbf{u} \frac{\delta O_2}{\delta x} + \mathbf{v} \frac{\delta O_2}{\delta y} + \mathbf{w} \frac{\delta O_2}{\delta z} + K_Z \frac{\delta^2 O_2}{\delta z^2} + K_H \left(\frac{\delta^2 O_2}{\delta x^2} + \frac{\delta^2 O_2}{\delta y^2} \right) + \text{ase} + \text{bio} \quad (1)$$

where O_2 is the concentration of DO, $(\mathbf{u}, \mathbf{v}, \mathbf{w})$ is the water velocity, z is depth, K_Z and K_H are vertical and horizontal eddy diffusivity respectively, ase is air–sea exchange and bio biological processes (photosynthesis, respiration, bacterial remineralisation of organic matter).

In the context of a strongly stratified two-layer system, air–sea exchange (ase) does not affect the BML and can be neglected. In this region and on short time scales (3 days), the \mathbf{u} and \mathbf{v} components (horizontal advection by currents), K_Z (vertical eddy

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diffusivity) and K_H (horizontal eddy diffusivity) can be neglected for the BML due to the homogeneity of the bottom mixed layer across the central North Sea (Queste et al., 2013). Therefore, for the BML in this region, the w component (vertical cross-thermocline transport) and biological processes (bio) dominate (Greenwood et al., 2010).

As evidenced by the chlorophyll *a* and supersaturation signatures (Fig. 3), the majority of the production occurs within the DCM and SML. Consequently, the dominant biological processes within the BML are (i) benthic oxygen demand and (ii) the remineralisation of organic matter which may derive from (ii.a) sinking from the DCM and (ii.b) organic matter still in suspension issuing from the spring bloom. We can therefore simplify the equation describing supply and consumption of DO within the BML to:

$$\frac{\delta O_2}{\delta t} = w \frac{\delta O_2}{\delta z} + R_{\text{Benthic}} + R_{\text{DCM}} + R_{\text{Bloom}} \quad (2)$$

where R is respiration of (i) Benthic organic matter and originating from (ii.a) the DCM and from (ii.b) past production (Bloom).

4 Observed change in dissolved oxygen in the BML

Figure 6 illustrates AOU at different depths throughout the BML. AOU is shown at depths beginning several metres below the pycnocline to account for tidal vertical displacement. There is a clear trend of uniformly increasing AOU over time, indicating oxygen depletion, throughout the entire BML. A linear regression of mean AOU throughout the BML indicates a rate of $2.8 \mu\text{mol dm}^{-3} \text{ day}^{-1}$ of DO consumption with a standard error of the regression of 0.3. Three sharp decreases in AOU are indicated by the vertical dotted line. These relate to vertical mixing events identified in Fig. 7 and will be discussed in the following section.

An oxygen consumption rate of $2.8 \pm 0.3 \mu\text{mol dm}^{-3} \text{ day}^{-1}$ is very large; to reach the observed AOU of $76 \mu\text{mol dm}^{-3}$ from saturation would require a month. However in this

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derived from MODIS Aqua data (Fig. 5) reveal significant horizontal variability near the Seaglider transect in surface waters. This surface variability is clearly shown in Fig. 3b where the Seaglider observed a freshening by 0.3 of the SML at the end of the survey. This change was not reflected in the BML, where the glider observed a weak but opposing trend. In the BML, the Seaglider observed a 0.1 °C cooling and 0.03 salinification (together equivalent to a densification of 0.035 kg m⁻³), likely due to increasing Atlantic water influence.

Although both 2010 and 2011 show an overall similar spatial pattern with depleted oxygen in the northern half, areas of lowest oxygen saturation are located in different regions (Fig. 2). Given the potentially very high consumption rates, these patterns of oxygen saturation are mainly a snapshot set of observations of late summer and may not be representative of the entire season or the previous month. This illustrates the interannual variability and importance of horizontal transport processes on broad scales in the North Sea, and also potentially in other shelf seas, linked to other dominating climate modes. Horizontal transport processes likely affect seasonal oxygen depletion by influencing BML and SML temperatures thereby affecting stratification while also providing nutrients to sustain productivity and potentially supplying labile organic matter to the BML.

On time scales similar to this survey, it is likely that advective processes play a limited role in regulating seasonal oxygen depletion; instead, vertical mixing is assumed to be the largest potential source of oxygen input into the BML. Seaglider observations reveal a highly stratified water column with a strong pycnocline spanning approximately 3 m with a density difference of 1.5–2 kg m⁻³ (Fig. 3c). Evidence of mixing can be found in changing properties of the BML as observed by the glider. Here, temperature is used as a tracer to assess mixing across the pycnocline on short time scales through the large temperature difference between the SML and BML (Fig. 7). Temperature data from below the thermocline are used to derive the vertical temperature gradient in the BML (Fig. 7, black vertical lines). A small value indicates a homogenous BML while a large value indicates a strong vertical gradient in the BML. Figure 7 shows changes

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in vertical temperature gradients on the order of a few hours (or hundreds of metres) horizontally. The BML temperature gradient increases simultaneously to mean BML temperature. This indicates warming of BML water by mixing across the thermocline and injection of warm SML water into the cooler BML.

Referring back to Fig. 6, we observe similar fluctuations in AOU. The three principal mixing events, illustrated by the grey vertical lines, correspond to sudden decreases in AOU indicating sudden and rapid reoxygenation of the BML. After these reoxygenation events, net oxygen consumption rates increase within a few hours. The three events over the three days show increases in BML DO (decreases in AOU) of approximately 1 to 3 $\mu\text{mol dm}^{-3}$ per event ($2 \pm 1 \mu\text{mol dm}^{-3} \text{ day}^{-1}$).

Figure 8 shows wind and tidal velocities along with bathymetry at the glider's position. There is no obvious correlation between the mixing events identified in Fig. 7 and the speed or direction of the tidal velocities. The majority of mixing events tend to occur during periods of peak wind velocities, but the duration of mixing events does not seem to match the time scale of wind velocity changes; furthermore, two mixing events occur in periods of low wind velocities. Mixing events do not seem to correlate with bathymetry in the region either (Fig. 8).

These mixing events are short scale (a few hours or a couple of kilometres) processes that the Seaglider is able to reveal through its very high resolution observations. Since wind events do not seem to be responsible, these events may be linked to a variety of physical processes occurring in the region. Van Haren et al. (1999) highlighted the importance of internal waves as a potential source of mixing around the Dogger Bank. Another possibility is the generation of shear spikes as described by Burchard and Rippeth (2009) due to the interaction of wind and tides, explaining the lack of correlation with either one in particular.

Regardless of the cause, we observe short lived or very localised occasional oxygen resupply events leading to an approximate oxygen supply of $2 \pm 1 \mu\text{mol dm}^{-3} \text{ day}^{-1}$.

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Relating this back to the overall oxygen budget gives:

$$2.8 \mu\text{mol dm}^{-3} \text{ day}^{-1} = w \frac{\delta O_2}{\delta z} + R_{\text{Benthic}} + R_{\text{DCM}} + R_{\text{Bloom}}$$

$$2.8 \mu\text{mol dm}^{-3} \text{ day}^{-1} = -2 + R_{\text{Benthic}} + R_{\text{DCM}} + R_{\text{Bloom}} \quad (4)$$

6 Oxygen consumption

6.1 Benthic respiration

We now assess the relative magnitude of the different oxygen sinks within the BML. As the North Sea is a shallow region, vertical transport of organic matter from the SML and BML to the benthic layer is fairly rapid. On a whole North Sea scale, Van Raaphorst et al. (1998) report that 17 to 45 % of primary production is remineralised in the sediments, although the bulk of this occurs in the Skagerrak and Norwegian Trench area (50 to 70 %) due to transport processes. On a more local scale, tidal currents resuspend and transport this organic matter to temporary depocentres. This makes estimating benthic respiration across the North Sea difficult due to the consequent spatial heterogeneity. Van Raaphorst et al. (1998) report that particulate matter from the East Anglian plume and the southern North Sea may settle in the OG and the southern flank of the Dogger Bank.

Neubacher (2009) directly measured nutrient and oxygen dynamics at the benthic-pelagic interface at the ND site. DO uptake by the sediment was determined to be approximately $250 \mu\text{mol m}^{-2} \text{ h}^{-1}$ with variations linked to organic matter export to the benthic layer. Assuming a uniformly mixed BML with a height of 38 m (Fig. 3), this amounts to a decline in oxygen concentration of $1.6 \mu\text{mol dm}^{-3} \text{ day}^{-1}$ consumed throughout the BML from respiration in the sediment.

6.2 Remineralisation of DCM-originating organic matter

Primary production estimates for the North Sea range from 40 to 300 g C m⁻² yr⁻¹ depending on area and technique (modelling or empirical), with maxima in the central North Sea area (north of Dogger Bank; Weston et al., 2005). A transect by Weston et al. (2005) north of Dogger Bank showed whole water column-integrated primary production values of 167, 370 and 270 g C m⁻² yr⁻¹ for areas classified as Dogger Bank, the front (located along the northern edge of Dogger Bank) and the stratified area (the ND site) respectively.

More recently, the importance of the DCM has been reviewed. The DCM contribution to total primary production for the stratified North Sea during the summer has been estimated at between 58 and 60 % (Fernand et al., 2013; Weston et al., 2005). Values for new production range from 37 to 66 % produced at the DCM for the whole water column (Fernand et al., 2013; Weston et al., 2005). As the DCM relies on small scale mixing across the pycnocline to provide nutrients from below, it is necessarily located very near to the pycnocline. Consequently, any mixing also injects highly labile organic matter into the BML. Furthermore, sinking organic matter produced at the DCM is exported to the BML immediately; therefore any remineralisation of organic matter occurs in the BML. This means that nearly all export production from the DCM is consumed within the BML.

The Seaglider observed DCM chlorophyll *a* concentrations of 2 to 3 mg m⁻³ in 2011 (Fig. 3). These values are similar to those found by Fernand et al. (2013) but lower than concentrations observed by Weston et al. (2005) (ca. 5–10 mg m⁻³). This is likely explained by the systematic variation in DCM chlorophyll *a* concentrations across the North Sea. The site considered here is located further north than the site investigated in Weston et al. (2005) study and is not subject to as much mixing from internal waves fuelling the DCM in the ND region (Ducrotoy et al., 2000; Weston et al., 2005). Chlorophyll *a* concentrations at the DCM fluctuate on a near diurnal cycle (Fig. 3). This may be linked to either fluctuations in light availability, or a diurnal change in wind velocities.

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Increased wind speed could lead to an increase in chlorophyll *a* fluorescence a few hours later by encouraging nutrient supply to the DCM. Fluctuations in chlorophyll *a* at the DCM (Fig. 3) are not due to quenching as periods of high DCM chlorophyll *a* concentrations are present when surface chlorophyll *a* fluorescence is affected by high light intensity. Instead, it is likely that these fluctuations in chlorophyll *a* fluorescence at the DCM are caused by diurnal increases in nutrient supply linked to stronger winds in the mornings (Fig. 8).

We now investigate the export of DCM organic matter to the BML. Figure 3d show a similar pattern for BML chlorophyll *a* concentration and DCM chlorophyll *a* concentrations. We observe a delay between peaks in DCM chlorophyll and BML chlorophyll with BML chlorophyll *a* concentrations peaking a few hours later. As we have already observed a delay between peaks in wind velocities and DCM chlorophyll *a* concentrations, we can assume that wind-driven mixing is not responsible for injecting these large amounts of organic matter into the BML. Neither are these increases in BML chlorophyll *a* correlated to tidal velocities or changes in BML salinity, thus ruling out the influence of horizontal transport processes or tidal mixing on export of DCM organic matter to the BML. It seems likely that the wind drives production at the DCM but export mechanisms are driven by biological processes.

The Seaglider data show pulses, or short-lived increases, in BML chlorophyll *a* concentrations of 0.15 mg m^{-3} occurring on daily cycles. We assume these are events where DCM organic matter sinks to the BML. With an approximate BML height of 38 m, these pulses equate to $5.7 \text{ mg m}^{-2} \text{ day}^{-1}$ of chlorophyll *a* export to the BML. Using a particulate organic carbon (POC) to chlorophyll *a* ratio of 50 : 1 (Sathyendranath et al., 2009; Weston et al., 2004), this amounts to approximately $0.285 \text{ g C m}^{-2} \text{ day}^{-1}$ export to the BML. If all this organic matter were to be remineralised within the BML, this would equate to $33.58 \text{ mmol DO m}^{-2} \text{ day}^{-1}$ or $0.9 \text{ } \mu\text{mol dm}^{-3} \text{ day}^{-1}$.

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By summing the benthic ($1.6 \mu\text{mol dm}^{-3} \text{ day}^{-1}$) and pelagic ($0.9 \mu\text{mol dm}^{-3} \text{ day}^{-1}$) oxygen sinks within the budget, we obtain:

$$2.8 \mu\text{mol dm}^{-3} \text{ day}^{-1} = w \frac{\delta O_2}{\delta z} + R_{\text{Benthic}} + R_{\text{DCM}} + R_{\text{Bloom}}$$
$$2.8 \mu\text{mol dm}^{-3} \text{ day}^{-1} = w \frac{\delta O_2}{\delta z} + 2.5 \quad (5)$$

7 Balance of consumption and supply

Oxygen consumption arising in the water column from recently exported matter ($0.9 \mu\text{mol dm}^{-3} \text{ day}^{-1}$) and benthic processes ($1.6 \mu\text{mol dm}^{-3} \text{ day}^{-1}$) provides a potential for oxygen consumption of $2.5 \mu\text{mol dm}^{-3} \text{ day}^{-1}$. This is described as “potential” as the value is likely an overestimate of oxygen consumption rates because it is unlikely that all of the sinking organic matter is remineralised pelagically. Much of this sinking organic matter is deposited before being remineralised and therefore a portion of the DCM originating consumption ($0.9 \mu\text{mol dm}^{-3} \text{ day}^{-1}$) is already accounted for in the benthic consumption ($1.6 \mu\text{mol dm}^{-3} \text{ day}^{-1}$).

Furthermore, no POC:chlorophyll *a* ratio could be obtained from North Sea DCM phytoplankton communities in this region. It is likely that in reality the ratio is in fact lower than 50 : 1 as DCM phytoplankton communities are adapted to lower light conditions and contain greater concentrations of photopigments. If this is the case, it would only reduce the estimate of oxygen depletion potential from DCM export, reinforcing the implication that there are unresolved mechanisms contributing to the oxygen depletion.

Our estimates can also be skewed due to aliasing from the glider’s travel and advection. Distinguishing vertical processes such as injection events from horizontal advection or spatial aliasing from glider travel is critical when assessing oxygen dynamics. In this study, the temperature and salinity data allow different watermasses to be distin-

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guished; horizontal advection does occur throughout the bottom mixed layer. Such an event occurs at the end of the record and is excluded from our study.

Even by accounting for additional organic matter issuing from the DCM and SML (faecal pellets, lysed cells, zooplankton) and doubling the potential for oxygen consumption from the DCM, the sum of oxygen consumption potential from the DCM and the benthic compartment remain only marginally larger than the total net observed rate of dissolved oxygen consumption within the BML ($2.8 \mu\text{mol dm}^{-3} \text{day}^{-1}$). This either implies oxygen resupply rates much lower than the $2 \mu\text{mol dm}^{-3} \text{day}^{-1}$ identified in Fig. 7, or more likely, the existence of an additional oxygen sink relevant to the time and location of the survey which has not been accounted for.

$$2.8 \mu\text{mol dm}^{-3} \text{day}^{-1} = w \frac{\delta O_2}{\delta z} + R_{\text{Benthic}} + R_{\text{DCM}} + R_{\text{Bloom}}$$

$$2.8 \mu\text{mol dm}^{-3} \text{day}^{-1} = -2 + 2.5 + \text{Unidentified Sink} \quad (6)$$

In the results of this study, observed net consumption is significantly greater than previously reported values in the literature. As previously stated, the observed rate of apparent oxygen utilisation would lead to lower oxygen saturations than observed if maintained from the start of the stratified season until the glider deployment. The implication here is that the Seaglider surveyed the region during a short-lived or localised increase in apparent oxygen utilisation.

If we look at the budget in a wider seasonal context and ignore the glider-observed net consumption rate ($2.8 \mu\text{mol dm}^{-3} \text{day}^{-1}$), the estimates of oxygen supply and consumption derived above ($-2 + 2.5 \mu\text{mol dm}^{-3} \text{day}^{-1}$) are in agreement with past studies of the North Sea where net seasonal consumption rates of $0.4 \mu\text{mol dm}^{-3} \text{day}^{-1}$ were observed (Greenwood et al., 2010). This also points to the unknown sink being either short-lived or very localised and explains why the observed net consumption rate does not correlate with previous studies.

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8 Potential dissolved oxygen sinks

Several processes could account for this unidentified oxygen sink. Bacterial recycling within the BML is rarely accounted for and could contribute to this unaccounted for oxygen depletion. In addition, recent work has begun focusing more on the importance of resuspension events in seasonal oxygen depletion (van der Molen et al., 2012; Couceiro et al., 2013; Thompson et al., 2011; Greenwood et al., 2010).

Background benthic DO consumption is limited by oxygen penetration depth and Red-Ox levels (Neubacher, 2009). When deep benthic organic matter is resuspended into an oxic water column, bioavailable surface area is greatly increased and oxygen is readily available leading to very rapid and short-lived oxidation of reductants such as nitrite, ammonium and sulphide. Organic matter is likely also briefly degraded more rapidly when resuspended and more surface area is exposed to the oxic BML.

Three particular causes have been identified as providing enough turbulent energy to the seafloor to resuspend organic matter: tidal resuspension, large wind mixing events and trawling (van der Molen et al., 2012; Couceiro et al., 2013; Thompson et al., 2011; Jago et al., 2006, 2002; Van Raaphorst et al., 1998; Greenwood et al., 2010). Current speeds and storm events sufficient to cause resuspension have been recorded at both the OG and ND sites (de Jonge et al., 1996; Kröncke and Knust, 1995; Greenwood et al., 2010; Weston et al., 2008). The influence of trawling has only recently been investigated but has been shown to potentially lead to a small (0.5 %) decrease in BML DO (van der Molen et al., 2012) in modelling studies. A strong resuspension event leading to an even greater decrease in DO was documented by Greenwood et al. (2010) at the Oyster Grounds.

In the 2011 survey data, glider observations of optical backscatter at 650 nm in the BML show the existence of resuspension events (Fig. 3e). Glider optical backscatter measurements showed fluctuations similar to that of chlorophyll *a* concentrations with strong increases and rapid declines in BML backscatter. The distance from the seafloor at the deepest point of the dive prevents the glider from observing fluctua-

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tions of backscatter in the near bottom fluff layer. Despite this, there is a clear gradient present with increasing backscatter nearer the seabed showing that bottom stress does resuspend some organic matter (Fig. 4e). The peaks in BML optical backscatter do not correlate with observed cross-pycnocline mixing (Fig. 7) but seem to occur on a diurnal cycle, about 4 to 6 h after daily peak wind speeds (Fig. 8).

Correctly understanding and quantifying resuspension becomes increasingly important when one considers temporary depocentres. These sites accumulate organic matter originating from much wider regions through slowing of currents. These sites have the capacity to rapidly sequester large amounts of organic matter, not only during peak production but also throughout the winter. This creates a reservoir of highly labile organic matter which, when resuspended, leads to very rapid consumption of DO. Depocentres are not exclusive to the North Sea; they are driven by local hydrography and topography, acting as hotspots of oxygen depletion in shelf sea environments around the world.

9 Conclusions

This study, through the use of an autonomous underwater ocean glider, has given new insights into the time scales of processes that regulate the supply and consumption of oxygen across thermoclines and into the BML. The average observed rate of oxygen consumption rate of $2.8 \pm 0.3 \mu\text{mol dm}^{-3} \text{day}^{-1}$ is high; this would have resulted in the observed AOU of $76 \mu\text{mol dm}^{-3}$ within only a month of the onset of stratification. The glider observations' high temporal and vertical resolution reveals the periods when resupply of oxygenated water occurs, and enables these vertical processes to be distinguished from horizontal processes. Long term trends in net oxygen consumption hide a more complex short term variability reflecting variations in oxygen supply and consumption. During a period when the BML was isolated, the AOU was shown to increase significantly over a short 3 day period. The observed vertical mixing events have no apparent correlation with winds or tides, and remain a process to be further

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explored. Optical backscatter readings from the Seaglider in the BML showed varying suspended sediment, which generally increased near the bed and varied on diurnal time scales.

Greenwood et al. (2010) recorded average (since stratification) oxygen depletion rates in 2007 of $0.43 \mu\text{mol dm}^{-3} \text{ day}^{-1}$ at ND and $0.75 \mu\text{mol dm}^{-3} \text{ day}^{-1}$ at the OG. We observed export of organic matter from the DCM with the potential of consuming $0.9 \mu\text{mol dm}^{-3} \text{ day}^{-1}$, and the study by Neubacher (2009) found rates of $1.6 \mu\text{mol dm}^{-3} \text{ day}^{-1}$ for the sediment-water oxygen flux in incubation experiments. These rates highlight two particular aspects of the mechanisms governing seasonal oxygen depletion in the North Sea. The first is that consumption along the seafloor plays a predominant role in the consumption of BML oxygen. This supports the idea that depocentres, such as the OG, are particularly prone to seasonal oxygen depletion. Advected organic matter will add to the consumption budget, increasing the ratio of consumption to supply, whilst the reduced water flow that promotes this deposition also promotes stronger stratification. The second is that supply and consumption of organic matter are tightly coupled. Net oxygen depletion rates averaged over a season are an order of magnitude smaller than gross oxygen consumption and supply rates. A small shift in either gross DO consumption or supply would have a large impact on net oxygen consumption rates.

This study has shown the potential of new AUVs in regions such as the North Sea. This deployment was cut short and only 5% of battery capacity was utilised; a full duration glider deployment within the North Sea could last upwards of two months. Despite the deployment lasting only three days, the data revealed small scale mixing mechanisms that would not be identifiable with traditional oceanographic methods. A longer glider deployment would provide more confidence in these estimates of rates and would reveal the temporal variability of biological processes across an entire season. A Seaglider performing repeat transects in and out of known depocentres could provide critical high resolution observations of subsurface primary production and organic matter transport in regions of both low and high biomass accumulation.

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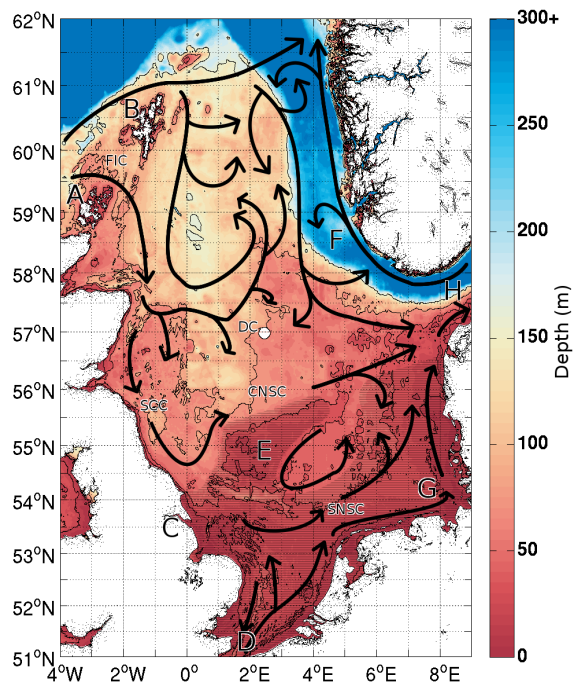


Figure 1. Bathymetry of the study area in metres (GEBCO, 2010). Depth contours have been added for the 20, 40, 80 and 160 m isobaths. The region where the glider was deployed is indicated by the large white dot at approximately 57° N 2°30' E. Major land marks are indicated as A and B: Shetland and Orkney Isles. C: Humber estuary. D: the Channel. E: Dogger Bank. F: Norwegian Trench. G: German Bight. H: Skagerrak. The general circulation of the North Sea (adapted from Turrell et al., 1992; Hill et al., 2008) is overlaid as arrows for FIC: Fair Isle Current, DC: Dooley Current, SCC: Scottish Coastal Current, CNSC: Central North Sea Current, SNSC: Southern North Sea Current. Hatch marks cover areas not subject to thermal stratification in summer.

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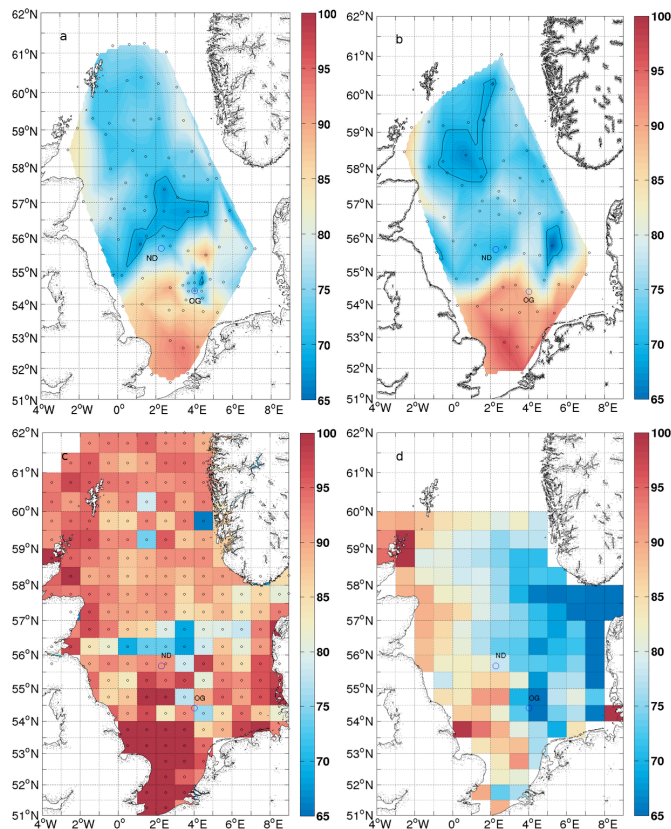


Figure 2. BML oxygen saturation (%) and with the 70 % saturation contour during the August 2010 (a) and August 2011 surveys (b), mean summer bottom oxygen saturation values (%) from 1900 to 2010 from the ICES database (c) and mean summer bottom oxygen saturation values (%) from 1958 to 2008 from the GETM-ERSEM model output using ERA interim reanalysis data (d).

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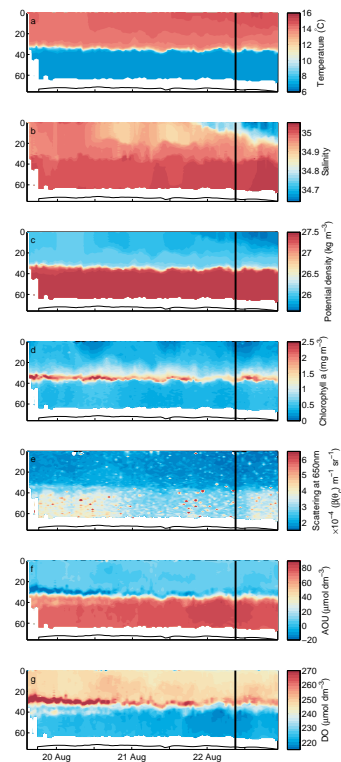


Figure 3. Seaglider section of temperature **(a)** ($^{\circ}\text{C}$), salinity **(b)**, potential density **(c)** (kg m^{-3}), chlorophyll *a* concentration **(d)** (mg m^{-3}), optical scattering at 650 nm **(e)** ($\times 10^{-4} \beta(\theta_c) \text{ m}^{-1} \text{ sr}^{-1}$), apparent oxygen utilisation **(f)** ($\mu\text{mol dm}^{-3}$) and oxygen concentration **(g)** ($\mu\text{mol dm}^{-3}$) sampled along the transect. Bathymetry as detected by the onboard altimeter is indicated along the bottom. The thick black line indicates the transition to a different water mass; data after are excluded from the oxygen calculations.

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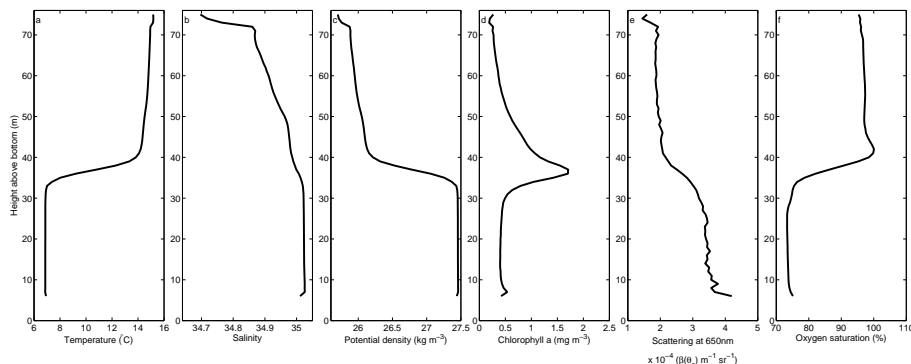


Figure 4. Mean profiles as observed by the Seaglider of temperature **(a)** ($^{\circ}\text{C}$), salinity **(b)**, potential density **(c)** (kg m^{-3}), chlorophyll *a* concentration **(d)** (mg m^{-3}), optical scattering at 650 nm **(e)** ($\times 10^{-4} \beta(\theta_c) \text{ m}^{-1} \text{ sr}^{-1}$) and oxygen saturation **(f)** (%) as height above the seabed.

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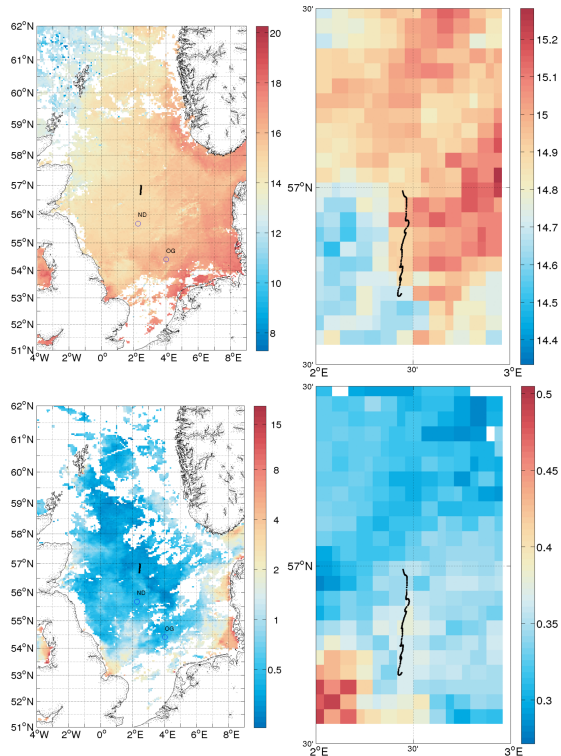


Figure 5. 3 day composite of MODIS Aqua daytime sea surface temperature at $11 \mu\text{m}$ ($^{\circ}\text{C}$, **a** and **b**) and chlorophyll *a* concentration (mg m^{-3} , **c** and **d**) across the entire North Sea (left) and around the Seaglider deployment area (right) from the 20 to 22 August 2011. The colour scale for the North Sea map of chlorophyll *a* is logarithmic to highlight chlorophyll *a* distribution in the central North Sea where concentrations are low. Northward Seaglider track is indicated by the black line.

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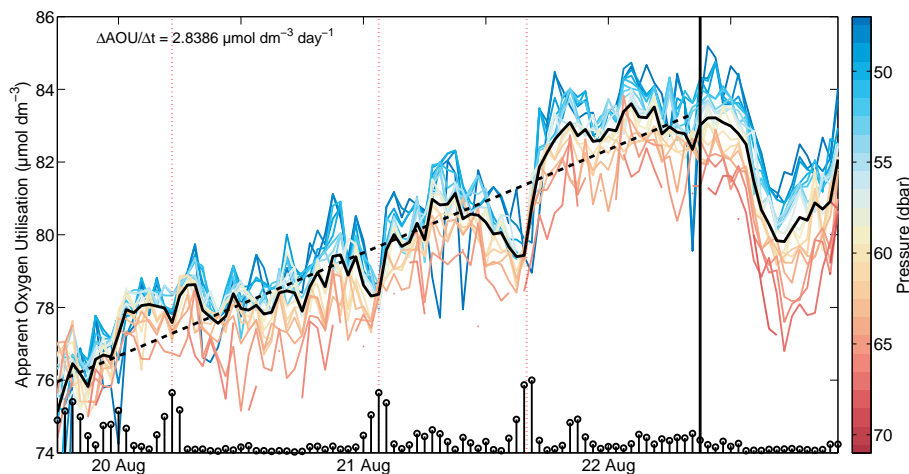


Figure 6. Seaglider apparent oxygen utilisation ($\mu\text{mol dm}^{-3}$) along isobars at 1 dbar intervals from 45 to 71 dbar, coloured by pressure. Black vertical lines indicate the maximum difference in temperature at each time step as shown in Fig. 7. Binned data from the BML (45 dbar and below) are plotted against time and coloured by pressure to highlight any vertical gradients. The dotted vertical lines indicate when vertical mixing events were evident. The thick black line indicates the transition to a different water mass.

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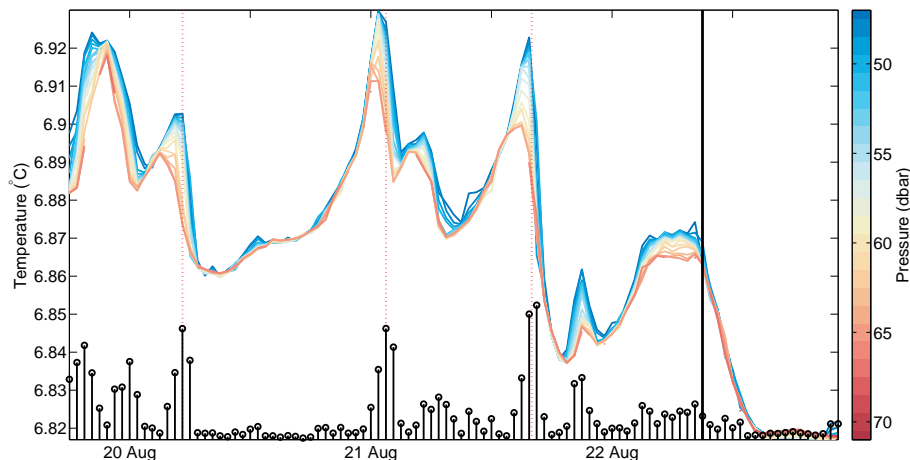


Figure 7. Seaglider temperature ($^{\circ}\text{C}$) along isobars at 1 dbar intervals from 45 to 71 dbar, coloured by pressure. Vertical black stems indicate the difference between minimum and maximum values at each time step. These illustrate changes in vertical gradients; a small value indicates a homogeneous BML while a large value indicates a strong vertical gradient in the BML. We can clearly see events where the temperature gradient increases in the BML, showing injection of warmer surface water across the thermocline.

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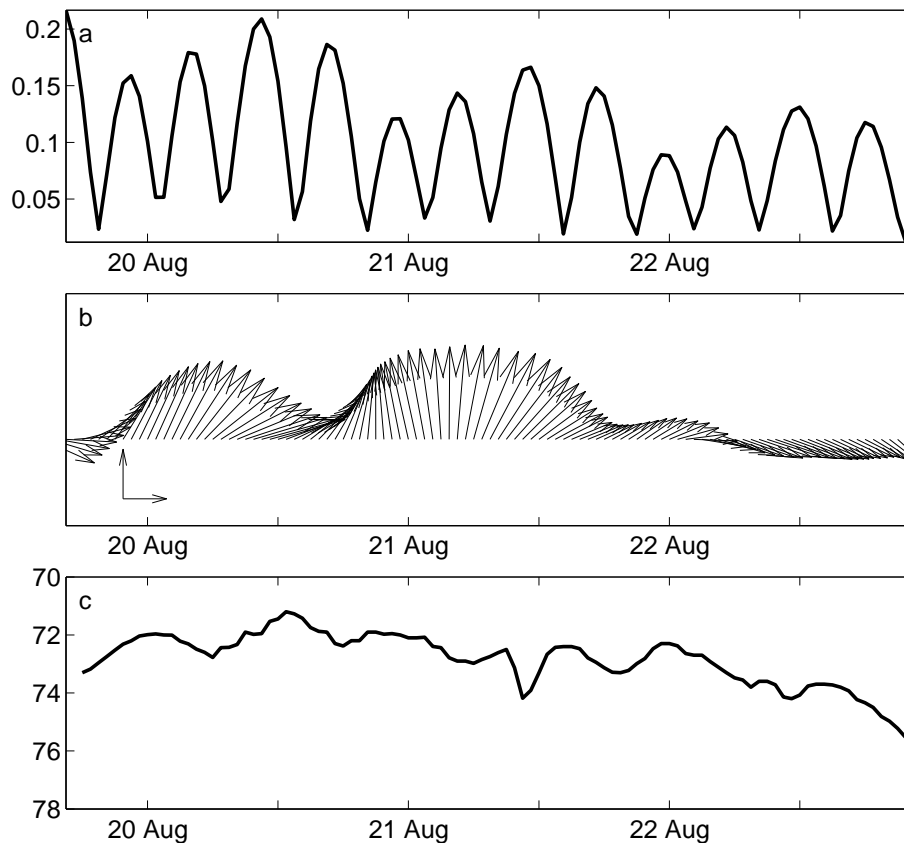
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Figure 8. Tidal velocities (ms^{-1} , **a**), 10m wind speed with scale arrows representing 5 ms^{-1} (**b**) and depth (m, **c**) at the Seaglider's location during the survey. Tidal data was obtained from the TMD tide toolbox and OTIS European Shelf Model (Egbert et al., 2010). Wind data was obtained from ECMWF ERA-Interim reanalysis data. Bathymetry was gathered by the Seaglider's on-board altimeter.