

SHIRSHOV INSTITUTE OF OCEANOLOGY

CRUISE REPORT No. 39

RV *AKADEMIK IOFFE* CRUISE 10 - 30 September 2012

**North Atlantic Repeat Hydrography of sill sections
between Greenland - Iceland - Faroe Islands and Shetlands**

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2013

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ABSTRACT

RV *Akademik Ioffe* Cruise 39 was a contribution to the RUSSIA CLIVAR Community Research Programme. The sections over North Atlantic sills between Greenland - Iceland - Faroe Islands and Shetlands were designed to estimate variability of the meridional fluxes and water mass exchange between the North Atlantic and the Arctic Ocean. The Denmark Strait section was repeated three times during six days to study short-term variability of the water mass properties circulation and meridional transport.

KEYWORDS

CRUISE 39 2012, AKADEMIK IOFFE, CLIVAR, DENMARK STRAIT, NORTH ATLANTIC SILLS, INTEROCEAN EXCHANGE, CTD OBSERVATIONS, VMADCP, LADCP

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1. CRUISE NARRATIVE

1.1 Cruise Details

Expedition Designation: R/V *Akademik Ioffe* Cruise 39, RUSSIA CLIVAR

Principal Scientists: Dr Sergey V. Gladyshev (Shirshov).

Ship: RV *Akademik Ioffe*.

Ports of Call: Kangerlussuaq (Greenland) to Rotterdam (Netherlands).

Cruise Dates: 10th September to 30th September 2012.

1.2 Cruise Summary

1.2.1 Cruise Track and Stations

The cruise track with station positions is shown in **Fig. 1**. Only small volume samples were taken, details are listed in **Table 1**.

1.2.2. Equipment

The principal instruments used during the cruise were a SBE 9P-743 CTD with dual temperature and conductivity sensors (SBE 3 SN 03P5642, SBE 4 SN 042827, SBE 3 SN 03P4401, SBE 4 SN 042925), oxygen sensor (SBE 43, SN 430699), turbidity sensor (SeaPoint Turbidity, SN 10218 STM), Benthos altimeter model PSA-900D, LADCP WHM-300 kHz down-looking (S/N 12312), LADCP WHM-300 kHz up-looking (S/N 12367). These were mounted together with a multisampler Carousel SBE 32 equipped with 22 5-litre Niskin bottles. Upon recovery each bottle was sampled in turn for dissolved oxygen, nutrients, salinity. All sampling was done on deck. Currents were measured using vessel mounted ADCP (VMADCP) TRDI OS38 kHz (S/N 1185) installed at the central point of the ship hall.

Navigation information was provided by a Trimble SPSx50/SPSx51 - Modular GPS receiver and every second was recorded on the PC. Additional measurements were made with an ELAC 12 kHz, Aanderaa meteorological package.

1.2.3 Sampling

Nominal depths sampled were: bottom, 1100, 1000, 900, 800, 700, 600, 500, 400, 300, 200, 150, 100, 50, 30, 20, 10m. On deep casts fewer shallow and intermediate bottles were fired. The actual bottle depths are shown in **Fig. 2**.

1.2.4 Number of Stations Occupied

103 stations (103 casts) were occupied during the cruise that includes sections between Greenland and Iceland (3 repeats), Faroe and Iceland, Shetlands and Faroe (3 repeats) (**Fig. 1**).

1.3 Scientific Objectives

The cruise objectives were to:

1. To complete a CTD section from the Great Britain to Greenland.
2. To survey the North Atlantic sills with high-resolution CTD and LADCP/VMADCP data to determine the circulation and meridional fluxes.

1.4 Narrative

1.4.1 Introduction

The Meridional overturning circulation (MOC) in the North Atlantic is one of the main drivers of the widely known global oceanic “conveyor belt” – an important element of the Earth’s climate system [e.g., [van Aken, 2007](#)]. Warm upper-ocean waters transported northward by the North Atlantic Current release heat to the atmosphere, gain density due to cooling and eventually sink in the subpolar North Atlantic and adjacent Arctic seas thereby generating the return southward flow of colder waters at depths (**Fig. 3**) [[Dickson and Brown, 1994](#); [Koltermann et al., 1999](#)]. Temporal variability of the large-scale circulation and associated heat transport in the subpolar North Atlantic is one of the principal factors behind the high-latitude climate anomalies in the Northern Hemisphere.

Progress in understanding the causes of the ongoing climate change and forecasting climate variability in the Arctic and over European part of Russia for the next decades require reliable observation-based estimates of the variability of the North Atlantic circulation and the

Atlantic–Arctic heat and freshwater fluxes, as well as elucidation of the underlying mechanisms. In a number of recent studies, radical changes in the thermohaline regime and large-scale circulation in the Atlantic Ocean have been suggested to occur under global warming. For instance, the long-term freshening of the subpolar North Atlantic deep waters since the mid-1960s [Dickson et al., 2002] has been (cautiously) attributed to climate change-related factors [Curry et al., 2003; Hansen et al., 2004]. Hypothetically, under global warming, an increased evaporation in the tropics and increased precipitation at high latitudes, coupled with an intensified melting of Arctic ice, lead to the upper-ocean freshening in the regions of deep water formation and, hence, to the deep water freshening in the Atlantic Ocean. At the same time, milder winters along with the upper-ocean freshening lead to a decrease in the deep water production rates, which results in slowing of the Atlantic Meridional Overturning Circulation [e.g., Hansen et al., 2004; Bryden et al., 2005].

To better understand the past and present changes in the ocean-atmosphere dynamical system, as well as their causes and consequences, data on the full-depth oceanic variability are needed. An indispensable effective tool for assessing the large-scale circulation and thermohaline changes in the deep ocean and investigating mechanisms governing these changes are repeated full-depth transoceanic observations.

Since 1997, the P.P. Shirshov Institute of Oceanology has carried out the long-term monitoring of the North Atlantic circulation and water mass properties in the 59.5°N hydrographic section between Cape Farewell (Greenland) and Scotland (**Fig. 3**). Since 2002, the section has been repeated yearly on board the Russian research vessels, providing high precision data on temperature, salinity, oxygen and nutrients concentrations, and current velocities in the entire water column – “from shore to shore”, from the sea surface to the bottom. In 2011, in addition to annual repeat measurements at 59.5°N, the P.P. Shirshov Institute of Oceanology started full-depth repeat observations of the oceanic exchange between the Atlantic and Arctic oceans through the straits between Greenland, Iceland, Faeroe and Shetland Islands (**Fig. 3**). The full-depth observations – of the same oceanic quantities as at 59.5°N – are performed in the straits from research vessels twice a year, in spring and fall. Based on the unique data set thus collected, a number of fundamental findings have already been achieved. Below, we briefly summarize the main subjects and results of our research.

The 59.5°N transatlantic section (**Fig. 3**) was designed for monitoring the large-scale circulation and thermohaline / chemical properties of oceanic waters at the northern periphery of the NA – the region where the warm upper-ocean waters are transformed by deep convection and mixing into the colder intermediate and deep waters – the Labrador Sea Water (LSW), Iceland Scotland Overflow Water (ISOW) and Denmark Strait Overflow Water (DSOW) (**Fig. 3**) – transported southward in the lower limb of the Atlantic MOC. Hydrographic data collected at 59.5°N along with those obtained within the framework of the kindred projects, primarily the French OVIDE (<http://www.ifremer.fr/lpo/ovide>), and historical data sets have been used for studying the dense water production [Falina et al., 2007; Falina et al., 2012], decadal temperature and salinity changes in the intermediate–deep water column [Sarafanov et al., 2007; Sarafanov et al., 2008; Sarafanov et al., 2010b], causes of these changes [Sarafanov, 2009; Sarafanov et al., 2010b], the mean state [Sarafanov et al., 2012] and long-term variability of the large-scale circulation in the region [Sarafanov et al., 2009; Sarafanov et al., 2010a; Våge et al., 2011].

1.4.2 Deep convection in the Irminger Sea

The oxygen data collected in 1997 in the northern North Atlantic in several sections ending nearby the southern tip of Greenland provided the observation-based support for the hypothesis [Pickart et al., 2003] that winter convection in the Irminger Sea may penetrate deep into the LSW layer (1000 – 2000 m) thus causing local renewal of this water mass. A separate lateral maximum of oxygen concentrations in the deep LSW layer was detected east of Cape Farewell (59.5°N, 36–40°W): the concentrations increased (by ~0.1 ml/l) from the Labrador Sea eastern edge toward the Irminger Sea (**Fig. 4**) rather than the reverse, as would be expected if LSW observed in the Irminger Sea interior in 1997 were solely of advective origin [Falina et al., 2007].

1.4.3. Reversal of the deep-water freshening

The LSW and Nordic Seas overflow-derived deep waters, ISOW and DSOW, freshened in the northern North Atlantic during the last three–four decades of the 20th century [Dickson et al., 2002]. Between the 1960s and 1990s, the water column in the region freshened on average by about 0.03 [Curry et al., 2003].

The long-term freshening reversed in the mid-1990s [Sarafanov et al., 2007; Sarafanov et al., 2008; Sarafanov et al., 2010b]. The salinification (and warming) of the intermediate and deep waters since the mid-1990s (**Fig. 5**) was much more intense than the preceding freshening. Over nearly a decade (1997–2006), temperature / salinity in the intermediate–deep water column ($\sigma_0 \geq 27.45$, depths > 500 – 1000 m) at 59.5°N increased by $\sim 0.3^\circ\text{C} / 0.03$ – 0.04 [Sarafanov et al., 2008].

In the Irminger Sea, the long-term freshening in the deep water column ($\sigma_0 > 27.80$, depths $> \sim 2000$ m) reversed in the early 2000s [Sarafanov et al., 2010b]. The observed freshening reversal was a lagged consequence of the persistent ISOW salinification that occurred upstream, in the Iceland Basin, after 1996 due to salinification of the northeast Atlantic waters entrained into the overflow. It was demonstrated [Sarafanov et al., 2010b] that the entrainment salinity increase was associated with the North Atlantic Oscillation (NAO)-induced weakening and contraction of the Subpolar Gyre and corresponding northwestward advance of subtropical waters that followed the NAO decline in the mid-1990s and continued through the mid-2000s. Remarkably, the deep water freshening reversal was not related to changes in the overflow water salinity.

1.4.4. Deep-ocean salinity changes and the NAO

Close relationship between the thermohaline properties of the northern North Atlantic intermediate and deep waters and the winter NAO index on a decadal time scale ($r^2 \approx 0.65$, 1950s–2000s, **Fig. 6b** and **6c**) was revealed [Sarafanov, 2009] from the observation-based salinity time series for LSW in the Labrador Sea [Yashayaev, 2007] and ISOW in the Iceland basin [Boessenkool et al., 2007; Sarafanov et al., 2007]. Persistent NAO decline (amplification) leads to warming and salinification (cooling and freshening) in the intermediate–deep water column.

An explanation for the close link between the NAO and the coherent decadal changes in the intermediate and deep water properties in the region was proposed [Sarafanov, 2009]. The two factors dominate this link (**Fig. 6d**): (i) intensity of convection in the Labrador Sea controlling injection of relatively cold fresh waters into the intermediate layer and (ii) zonal extent of the Subpolar Gyre that regulates the relative contributions of cold fresh subpolar waters

and warm saline subtropical waters to the entrainment into the Norwegian Sea overflow south of the Iceland–Scotland Ridge and to the Atlantic inflow to the Nordic Seas. These factors act in phase leading to the observed coherent thermohaline changes in the intermediate–deep water column.

Due to weakening of the surface forcing associated with the NAO transition into neutral to low phase (1950s to mid-1960s, mid-1990s to mid-2000s), convection in the Labrador Sea weakens diminishing cold fresh water penetration into the intermediate layer. This results in warming and salinification at the intermediate depths in the Subpolar Gyre. Concurrently, the Subpolar Gyre contracts allowing northward advance of warm saline upper-ocean and intermediate subtropical waters in the northeastern North Atlantic. Northward progression of subtropical waters increases temperature and salinity at the upper intermediate levels and, correspondingly, increases temperature and salinity of the northeast Atlantic waters entrained into the Iceland–Scotland overflow along its pathway to the deep Iceland basin. As a result, temperature and salinity at the deep levels increase. The contrary changes – intensification of deep convection in the Labrador Sea and expansion of the Subpolar Gyre – caused by amplifying surface forcing (mid-1960s to mid-1990s) lead to cooling and freshening at the intermediate–deep levels. Additionally, under high-NAO conditions, deep convection may occur in the Irminger Sea potentially contributing to cooling and freshening at the intermediate (LSW) levels. The two regimes of convection and large-scale circulation corresponding to stronger (early 1990s) and weaker (mid-1960s, mid-2000s) NAO-related atmospheric forcing are schematically visualized in **Fig. 7**.

1.4.5 Deep-ocean salinity changes and climate change

There are increasing concerns that in the warmer climate, the MOC may substantially decline due to a decrease in the convective activity in the northern North Atlantic and Nordic Seas [e.g., [Meehl et al., 2007](#)]. The long-term freshening in the Nordic Seas and freshening of the northern North Atlantic deep waters in the 1960s–1990s have been considered as a likely indicator or precursor of the dramatic change in the MOC [e.g., [Hansen et al., 2004](#)]. The freshening has been attributed to a combination of factors potentially associated with the global warming: the increasing ice melt and net precipitation at high latitudes [e.g., [Curry et al., 2003](#)]. A probable causality between the climate change and the decreasing North Atlantic deep water

salinity has supported the concerns and unfavorable predictions, thus ‘warming up’ the reasonable scientific debate on climate change and overblown speculations in media.

Despite the long-term increase in freshwater input to the Arctic, freshening in the northern North Atlantic had reversed in the mid-1990s, as we demonstrated above. This reversal forces us to revise the hypotheses on the mechanisms behind the deep-water thermohaline anomalies. It seems doubtful that the persistent global temperature growth may lead to the opposite decadal trends (positive-then-negative-then-positive, **Fig. 6**) in the deep water salinity.

Our results [[Sarafanov et al., 2008](#); [Sarafanov, 2009](#); [Sarafanov et al., 2010b](#)] suggest that natural atmospheric variability over the North Atlantic plays the major role in the deep-water thermohaline variability on a decadal time scale. There are no reasons to associate the deep-water freshening in the 1960s–1990s with climate change, unless the 3-decade-long surface forcing amplification is evidently shown to be a consequence of the latter. Having said that, the net 1950s–2000s trends in the water mass salinities are negative implying that the global factors (e.g., probable intensification of hydrological cycle [[Curry et al., 2003](#)]) may act on longer time scales.

1.4.6 Decadal variability of the Deep Western Boundary Current at Cape Farewell

Recent decadal changes in the Deep Western Boundary Current (DWBC) transport southeast of Cape Farewell were assessed from hydrographic data (1991–2007, **Fig. 7a**), direct velocity measurements (2002–2006) and satellite altimetry (1992–2007). Following the approach used in earlier studies [e.g., [Bacon, 1998](#)], we first determined that the DWBC ($\sigma_0 > 27.80$) baroclinic transport (T_{BC}) referenced to 1000 m depth increased by ~ 2 Sv between the mid-1990s (1994–1997) and 2000s (2000–2007) (**Fig. 8b**) [[Sarafanov et al., 2009](#)]. In the next step, we quantified velocity changes at the reference level (1000 m) by combining estimates of the hydrography-derived velocity changes in the water column and the altimetry-derived velocity changes at the sea surface [see [Sarafanov et al., 2010a](#)]. The inferred increase in the southward velocity at 1000 m above the DWBC in 1994–2007 indicates that the increase in the DWBC absolute transport was larger but very close to the 2-Sv increase in the DWBC T_{BC} . This result along with the observed coherence of the DWBC absolute and baroclinic transport changes between individual observations [[Sarafanov et al., 2010a](#)] imply that the DWBC absolute

transport variability in the region is well represented by its baroclinic component on decadal and shorter time scales.

The historical record of the DWBC T_{BC} (1955–2007, **Fig. 8c**) updated after Bacon [1998] shows distinct decadal variability (± 2 – 2.5 Sv) with the transport minima in the 1950s and mid-1990s, maximum in the early 1980s and moderate-to-high transport in the 2000s. The DWBC T_{BC} decadal variability is consistent with the general pattern of the recent decadal hydrographic and circulation changes in the northern North Atlantic. The DWBC T_{BC} anomalies negatively correlate ($R = -0.80$, 1955–2007) with thickness anomalies of the Labrador Sea Water (LSW) at its origin implying a close link between the DWBC transport southeast of Cape Farewell and the LSW production in the Labrador Sea (**Fig. 8d**). During the recent three decades (late 1970s – late 2000s), the DWBC T_{BC} changes were also in-phase with changes in the strength and zonal extent of the Subpolar Gyre [see Sarafanov et al., 2010a]. In particular, the Gyre weakening at shallow levels in the mid-1990s – mid-2000s was accompanied by the DWBC strengthening in the Irminger Sea [Sarafanov et al., 2009; Sarafanov et al., 2010a; Våge et al., 2011]. The results imply that the decadal changes in the (i) LSW production, (ii) SPG strength and (iii) DWBC transport in the Irminger Sea are linked, representing a complex coherent oceanic response to the decadal variability of the surface forcing.

1.4.7 Mean state of the full-depth circulation in the 2000s

A mean state of the full-depth summer circulation in the Atlantic Ocean in the region in between Cape Farewell (Greenland), Scotland and the Greenland-Scotland Ridge (see **Fig. 3**) was assessed by combining 2002–2008 yearly hydrographic measurements at 59.5°N , mean dynamic topography, satellite altimetry data and available estimates of the Atlantic–Nordic Seas exchange [see Sarafanov et al., 2012]. The mean absolute transports by the upper-ocean, mid-depth and deep currents and the MOC ($\text{MOC}_{\sigma} = 16.5 \pm 2.2$ Sv, at $\sigma_0 = 27.55$) at 59.5°N were quantified in the density space. Inter-basin and diapycnal volume fluxes in between the 59.5°N section and the Greenland-Scotland Ridge were then estimated from a box model.

The estimated meridional and diapycnal volume fluxes contributing to the MOC are schematically visualized in **Fig. 9**. The dominant components of the meridional exchange across 59.5°N are the North Atlantic Current (NAC, 15.5 ± 0.8 Sv, $\sigma_0 < 27.55$) east of the Reykjanes Ridge, the northward Irminger Current (IC, 12.0 ± 3.0 Sv) and southward Western Boundary

Current (WBC, 32.1 ± 5.9 Sv) in the Irminger Sea and the deep water export from the northern Iceland Basin (3.7 ± 0.8 Sv, $\sigma_0 > 27.80$). About 60% (12.7 ± 1.4 Sv) of waters carried in the MOC σ upper limb ($\sigma_0 < 27.55$) by the NAC/IC across 59.5°N (21.1 ± 1.0 Sv) recirculates westwards south of the Greenland-Scotland Ridge and feeds the WBC. 80% (10.2 ± 1.7 Sv) of the recirculating NAC/IC-derived upper-ocean waters gains density of $\sigma_0 > 27.55$ and contributes to the MOC σ lower limb. Accordingly, the contribution of light-to-dense water conversion south of the Greenland-Scotland Ridge (~ 10 Sv) to the MOC σ lower limb at 59.5°N is one and a half times larger than the contribution of dense water production in the Nordic Seas (~ 6 Sv).

1.4.8 Cascading of dense shelf waters in the Irminger Sea

Based on the hydrographic data collected at 59.5°N , 64.3°N and $65\text{--}66^\circ\text{N}$ in the western Irminger Sea in the 1990s – 2000s, an observational evidence for the deep-reaching cascading of dense shelf waters south of the Denmark Strait was found [Falina et al., 2012]. The data collected in the northwestern Irminger Sea ($65\text{--}66^\circ\text{N}$) indicate that the East Greenland Current ~ 200 km south of the Denmark Strait occasionally carries shelf waters as dense as the overflow-derived deep waters transported by the DWBC ($\sigma_0 > 27.80$). Hydrographic traces of cascading of dense shelf waters down the East Greenland slope were found from repeat measurements at 64.3°N , where the densest fresh plumes were observed within the DWBC ($\sigma_0 > 27.80$) (Fig. 10). Using the data collected at 59.5°N , we showed that the fresh ‘signals’ originating from the shelf can be traced in the DWBC as far downstream as the latitude of Cape Farewell, where the anomalously fresh oxygenated plumes are repeatedly observed in the ISOW and DSOW density classes.

The results of our analysis along with the results from earlier studies [e.g., Rudels et al., 1999; Rudels et al., 2002] indicate that shelf water cascading in the northern Irminger Sea is an intermittent process occurring in all seasons of the year. This implies that, despite the apparent short duration of a particular cascading event, the cumulative contribution of such events to the thermohaline variability and southward export of the deep waters in the WBC can be considerable. Our tentative estimate based on data from two synoptic surveys at $\sim 59.5^\circ\text{N}$ suggests that the transient contribution of a cascading event in the northern Irminger Sea to the DWBC transport at Cape Farewell can be as large as $\sim 25\%$.

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1.5 Preliminary Results

The upper-ocean, mid-depth and deep water circulation patterns, merging the results of the present analysis with those from the earlier studies [e.g., *Macrander et al.*, 2005; *Østerhus et al.*, 2005, 2008; *Schott and Brandt*, 2007; *Sutherland and Pickart*, 2008; *Lherminier et al.*, 2010; *Våge et al.*, 2011], are schematically visualized Figures 12–14. A schematic diagram of the meridional overturning circulation in the Atlantic Ocean north of 59.5°N is displayed in Fig. 9.

The results provide the following conceptual view of the gyre / overturning circulation at the northern periphery of the Atlantic Ocean in the 2000s.

The NAC and IC collectively carry 21.1 ± 1.0 Sv of warm upper-ocean waters across 59.5°N northwards within the MOC σ upper limb ($\sigma_0 < 27.55$). About 40% of this flow forms the Atlantic Inflow to the Nordic Seas, and 60% (12.7 ± 1.4 Sv) recirculates westwards in the subpolar gyre northern limb south of Iceland to feed the WBC in the Irminger Sea. Only 20% (2.4 ± 1.2 Sv) of the recirculating NAC/IC-derived waters exits the Irminger Sea in the WBC at shallow levels ($\sigma_0 < 27.55$), while 80% (10.2 ± 1.7 Sv, a half of the NAC/IC northward flow across 59.5°N) gains density of $\sigma_0 > 27.55$ and enters the MOC σ lower limb. The resulting net southward transport in the MOC σ lower limb at the latitude of Cape Farewell is 16.5 ± 2.2 Sv, of which ~60% (~10.2 Sv) is due to light-to-dense water transformation south of the GSR.

As no dense-to-light water re-conversion is expected to occur in the subpolar gyre, the NAC/IC-derived waters, once entering the MOC σ lower limb in the Irminger Sea, will eventually contribute to the MOCz lower limb (~11 Sv at 59.5°N) at the southern margin of the subpolar region. There, at ~48°N, the MOC σ and MOCz are of nearly the same magnitude, 16 ± 2 Sv, as estimated from data collected in the 1990s [see *Schott and Brandt*, 2007; *Lumpkin et al.*, 2008]. This is very close to our estimate of the mean MOC σ at 59.5°N. The comparison is tentative, though, because it does take into account the decadal variability of the MOC [Koltermann et al., 1999; Willis, 2010]. With this caveat in mind, our results imply a minor contribution to the MOC σ by the net dense water formation in the subpolar gyre between ~48°N and 59.5°N. This inference concurs with the results by *Pickart and Spall* [2007] suggesting a

minor contribution to the Atlantic MOC by the net water mass transformation in the Labrador Sea.

To conclude, the results of the present study, verified with independent estimates where possible, provide the first observation-based quantitative view of a mean state of the gyre / overturning circulation at the northern periphery of the Atlantic Ocean. The most interesting features of the obtain circulation pattern are as follows:

- Nearly half of volume of the upper-ocean waters transported northward across 59.5°N in the eastern limb of the subpolar gyre (NAC and IC, $\sigma_0 < 27.55$) overturns in the density plane south of the GSR and feeds the lower limb of the Atlantic MOC σ .
- The contribution to the MOC σ lower limb at 59.5°N by overturning (light-to-dense transformation) of the NAC / IC-derived upper-ocean waters south of the GSR is one and a half times as large as the contribution of the Nordic Seas overflows.
- The net southward flow in MOC σ lower limb at 59.5°N is associated primarily with the deep water ($\sigma_0 > 27.80$) export. Nearly half of the net southward flow of deep waters across 59.5°N is due to entrainment of the Atlantic waters in the Irminger Sea.
- The DWBC at 59.5°N is fed primarily by the Denmark Strait Overflow and by the diapycnal flux / entrainment from the mid-depth layer, while the contribution to the DWBC transport from the ISOW flow is minor. A major part of the ISOW transported into the Irminger Sea from the Charlie-Gibbs Fracture Zone recirculates southward in the eastern Irminger Sea and exits the basin via an interior pathway rather than along the western boundary. The results can be used for validation of numerical models. From this perspective, multi-year mean transports have an obvious advantage over individual section-based synoptic estimates, which bear the impress of vigorous variability occurring on a variety of spatial and temporal scales. The methodological outcome is that the combined use of repeat hydrography, the MDT by *Rio and Hernandez* [2004] and satellite altimetry data can provide a useful estimate of the mean full-depth circulation across a transatlantic section without imposing *a priori* constraints.

1.6 Major Problems and Goals Not Achieved

The CTD secondary pump S/N 053794 was replaced with S/N 053792 after sta. 2829 due to its bad performance (noisy data of the secondary sensors). Bottle # 7 did not close on sta. 2837 - 2845. To fix the problem we removed pylon to tighten screws. Bottle # 17 did not close on sta. 2919 – 2925 because of cock release mechanism problem. We had to stop cast during sta. 2911 because of ship maneuvering and because of large cable angle. There was a large ship drift about 0.5 nm on station. There was no connection to SLAVE after a few casts. To set up connection we always re plug the star cable in.

2. CONTINUOUS MEASUREMENTS (on station and underway)

2.1 Navigation

Navigation data from Trimble SPSx50/SPSx51 GPS was recorded every 1 second and was stored on the PC in binary format.

2.2 Meteorological Measurements

The standard mean meteorological measurements were stored in the separate files on the same PC with navigation data. Recording were running immediately after departure from Kangerlussuaq (Greenland) on 10th September, and worked reliably until completion of the cruise in Rotterdam (Netherlands) on 30th September.

2.3 Thermosalinograph

SBE 21 S/N 3254 data were collected along the section line starting on September 16th.

2.4 Echosounding

The bathymetric equipment aboard during RV Akademik Ioffe Cruise 38 consists of an ELAC 12 kHz hydrographic echosounder. Data were collected for most of the cruise. The Hull mounted transducer is located 5.8 metres below the sea surface and this value was entered to estimate the depth.

Depth was indicated on the echosounder display and stored on the PC together with the navigation.

Two files with extension NAV and MET with maximum size 256032 b were created. File name corresponded to GMT time when the file was opened for records.

2.5 Vessel Mounted Acoustic Doppler Current Profiler (VMADCP) OS 38 kHz

The Ocean Surveyor 38 kHz is designed for vessel-mount current profile measurement in the upper ocean water from depths greater than 40-50 meters. The system consists of a transducer and electronics chassis connected to PC. Data are transmitted in binary format through the I/O cable. GPS data in NMEA format are transmitted separately to another PC COM – port. The VMADCP can operate in two regimes (Narrow Bandwidth and Broad Bandwidth Profiling). Its main specifications are shown below.

To collect OS 38 kHz data we used *VmDas* software (version 1.46). The NMEA messages *VmDas* reads are standard GGA, HDG, HDT, VTG messages.

	Bin size	Maximum range	Accuracy (cm/s) ²
NarrowBand (long-range mode)	16 m	800 - 1000 m	30
	24 m	900 - 1200 m	23
BroadBand (high-precision mode)	16 m	520 - 730 m	12
	24 m	730 – 780 m	9

We mainly used a following configuration to collect the data.

NP00001 – Narrow Bandwidth profiling

NN060 – number of bins 60

NS1600 – cell size 16 m

NF1600 – blanking size 16 m

BP00 – no bottom track (BP),

We also used the Broad Bandwidth regime with cell size 24 m and bottom track mode at the beginning of data collection to refine bottom topography near Greenland. This regime was switched to NB16 regime after 3 hours of work.

VmDas saves data in a few files with extension ENX, ENS, ENR (raw data with and without navigation), NR – NMEA messages, STA and LTA averaged data. Misalignment angle was introduced in configuration file and was used by VmDas for data correction.

Data processing performed STA files with 40-profile averaging. Taking into account that single ping takes about 3 seconds, one 40-profile ensemble lasts near 120 seconds in Narrow Bandwidth regime.

Data processing consists of data conversion in NetCDF format with extension NC and further cleaning, filtering, tide removing (using barotropic tidal model TPXO 7.2) and averaging. The standard averaging was 3 km. IFREMER software was used to process OS 38 kHz data.

3. ON-STATION MEASUREMENTS

3.1 CTD

3.1.1 Equipment

The deep profiler system used during the cruise included the following components: SBE 32 painted aluminum 24 bottle multisampler frame, SBE 9P-0743 CTD, Up and Down looking RD Instruments WHM – 300 kHz Acoustic Doppler Current Profiler (LADCP), Separate Battery pack pressure case ext. 6000 m connected to LADCPs with star cable, 24 x 5 litre Test Oceanic Niskin bottles, Benthos altimeter PSA-900D.

Lab equipment for data acquisition and archiving of CTD/LADCP data consisted of the following items mounted on the deck.

Pentium IV – Intel 2.2 GHz, PC Intel Core 2 Duo 2.4GHz Personal Computers. APC Back-UPS 550VA/330W, SBE 11p Deck Unit.

Cruise Preparation

Equipment and sensors were assembled when the ship turned around Greenland (11-14th September). Water bottles were checked for integrity of seals, taps, stoppers and lanyards before being fitted and roped to the multisampler frame.

Deployment

The CTD performed well during the cruise with little evident instrument drift and good accuracy.

The CTD was deployed with a lowering rate of 60 metres/min (30-40 metres/min in the upper 200 metres or deeper if the conditions are rough). It is recovered at a rate of 60 metres/min.

The LADCPs fitted within the frame with a separate battery pressure case performed well. These units contain a compass and tilt sensors which could possibly provide useful information on the attitude and rotation of the whole profiler package throughout deployments.

Bottle firing using the deck unit and pylon was very reliable during the cruise.

Operationally this has been a successful cruise with virtually no time being lost due to mechanical or equipment failure.

3.1.2 Data processing and calibration

CTD data were logged at 24 scans per second and passed from the CTD deck unit to the PC.

The CTD data was recorded onto disk by the PC using SEABIRD SEASOFT-Win 32: Seasave 7, Software Release 7.21d. A screen display of temperature, oxygen, salinity and density profiles vs pressure are used to decide the depths at which bottles are to be tripped on the up cast. The bottles are tripped using the enable and fire buttons on the PC screen. During post-processing, the SEASAVE software stores 35 scans at each bottle trip within a separate file. At the end of the station, all the data and header files associated with the station are transferred immediately via ethernet to the second PC. The SBE data processing software is used to create 1 dbar processed data files.

The data processing takes the following steps:

DATCNV Converts the raw data to physical parameters.

WILDEDIT For every block of 100 scans, flags all scans whose pressure, temperature, conductivity and oxygen values differ from the mean by more than 2 standard deviations. Recomputes mean from unflagged data then marks as bad all scans exceeding 20 standard deviations from these new values.

FILTER Low pass filter pressure channel with time constant used for pressure 0.150 seconds.

ALIGNCTD Aligns the oxygen values relative to the pressure values accounting for the time delays in the system. Time offsets of 4.000 secs for oxygen are used.

CELLTM A recursive filter used to remove the thermal mass effects from the conductivity data. Thermal anomaly amplitude and time constants of 0.0300 and 7.0000 were used.

LOOPEDIT Marks as bad, all cycles on the down trace for which the vertical velocity of the CTD unit is less than 0.25 metres/sec.

WINDOW FILTER cosine filter temperature and conductivity, window size 23 scans.

DERIVE Computes salinity, potential temperature, sigma-t, sigma theta and oxygen values.

BINAVG Averages the down cast into 1 dbar pressure bins.

SPLIT Splits the data into DOWN and UP cast.

Calibration data

The CTD calibrations used during this cruise were supplied by Sea Bird Electronics and are as follows:

Pre-cruise calibration:

CALIBRATION DATE: 27-Jul 2012 (all stations)

Conductivity Sensor S/N 042827

$g = -1.0034748\ 3\ e\ +001$
 $h = 1.3698533\ 5\ e\ +000$
 $i = 4.6731011\ 0\ e\ -004$
 $j = 2.9400787\ 5\ e\ -005$
 $CPcor = -9.5700e-008$
 $CTcor = 3.2500e-006$

Post-cruise calibration:

CALIBRATION DATE: 13-Apr 2013 (all stations)

Conductivity Sensor S/N 042827

$g = -1.0036105\ 3\ e\ +001$
 $h = 1.3704675\ 1\ e\ +000$
 $i = 2.5341485\ 3\ e\ -004$
 $j = 4.8047158\ 8\ e\ -005$
 $CPcor = -9.570\ 0\ e\ -008$ (nominal)
 $CTcor = 3.250\ 0\ e\ -006$ (nominal)
 Average drift between *pre* and *post-cruise* calibrations: +0.0000 ms/m

Pre-cruise calibration:

CALIBRATION DATE: 22-Mar-12 (all stations)

Temperature Sensor S/N 035642 (loaned for one cruise from SBE)

Temperature ITS-90 = $1/\{g + h[\ln(f_0/f)] + i[\ln^2(f_0/f)] + j[\ln^3(f_0/f)]\} - 273.15$ (°C)
 Following the recommendation of JPOTS: T68 is assumed to be $1.00024 * T90$ (-2 to 35°C)
 f is the frequency
 $g = 4.3826941\ 2\ e\ -003$
 $h = 6.2902700\ 6\ e\ -004$
 $i = 1.9981456\ 2\ e\ -005$
 $j = 1.5264191\ 2\ e\ -006$
 $f_0 = 1000.0$

Pressure Sensor S/N 89105 (all stations) no drift

CALIBRATION DATE: 27-June-11

$C1 = -4.9053\ 7\ 1\ e+004$
 $C2 = -1.2105\ 9\ 4\ e+000$
 $C3 = 1.4283\ 5\ 0\ e-002$
 $D1 = 3.9016\ 0\ 0\ e-002$
 $D2 = 0.0000\ 0\ 0\ e+000$
 $T1 = 3.0010\ 1\ 7\ e+001$
 $T2 = -5.7583\ 8\ 4\ e-004$
 $T3 = 4.2101\ 2\ 0\ e-006$
 $T4 = 2.2654\ 0\ 0\ e-009$
 $T5 = 0.0000\ 0\ 0\ e+000$
 $AD590M = 1.28912e-002$
 $AD590B = -8.43097e+000$
 Slope = 0.99995
 Offset = 1.4284 (dbars)

Oxygen Sensor 430699

CALIBRATION DATE: 31-May-2013 (All Stations)

$Oxygen(ml/l) = \{Soc * (V + Voffset)\} * Oxsat(T,S) * e^{(Tcorr*T)} * e^{(Pcor*P)}$

Where:

V = SBE 43 output voltage signal (volts)

T = CTD temperature ($^{\circ}\text{C}$)

S = CTD salinity (psu)

P = CTD pressure (dbars)

$O_{xsat}(T,S)$ = oxygen saturation (ml/l)

Soc , V_{offset} , t_{cor} , and p_{cor} are calibration coefficients

Let:

$$\phi = O_{xsat}(T,S) * e^{(T_{corr}*T)} * e^{(P_{cor}*P)}$$

$$Oxygen(ml/l) = Soc * (V + V_{offset}) * \phi$$

$$Oxygen(ml/l) / \phi = Soc * (V + V_{offset}) = M * V + B$$

Where:

$$Soc = M$$

$$V_{offset} = B / M$$

In our case for sill sections

$$Soc = 4.0237e-001$$

$$Tau = 0.0$$

$$Boc = 0.0000$$

$$V_{offset} = -0.3524$$

$$t_{cor} = 0.001700$$

$$p_{cor} = 1.35e-004$$

3.1.3 Final Post-Cruise CTD Calibrations

Temperature Calibration Temperature Sensor S/N 035642, 034401

We applied 0.0000 correction for the primary sensor **S/N 035642** and -0.0001 for the secondary one **S/N 034401**. (Because of the small sensor drift between *pre-* and *post-*cruise calibrations we used *pre-calibration coefficients* for all stations of the cruise).

Pressure Calibration Pressure Sensor S/N 89105

Final CTD pressure correction: Since no drift for pressure sensor was defined by SeaBird Electronics pressure was corrected for atmospheric pressure only. With offset in *.con* or *.xmlcon* file set to -0.0026 db, pressure measured by CTD should equal barometric pressure

- Calculate offset (db) = barometer reading – CTD reading
- Conversion of psia to decibars: decibars = (psia - 14.7) * 0.6894759
- Enter calculated offset in *.con* or *.xmlcon* file
- Example:
 - CTD reads -2.5 dbars

– Barometer reads 14.65 psia.

Converting to decibars, barometer reads $(14.65 - 14.7) * 0.6894759 = -0.034$ dbars

– offset (db) = barometer reading – CTD reading = $-0.034 - (-2.5) = 2.466$

Salinity Calibration Conductivity Sensor S/N 042827

We used *pre-cruise calibration coefficients* for all stations of the cruise corrected by -0.0008 PSU (sensor slope correction 1.0000204) and 0.0001 PSU correction for the secondary sensor **S/N 042925**. This drift was defined by taking 327 salinity samples from depth deeper than 300 m between 16 and 27 September 2012. These samples were measured with Autosal 8400 B laboratory salinometer (S/N 67465) between 17 and 28 September. The results of these measurements are shown in **Fig. 11**.

Description of Equipment and Technique

Salinity samples are analyzed on Guildline Autosal model 8400 B salinometer. Samples are drawn in 150 ml dark glass medicine bottles. All bottles were equipped with plastic liners and caps. The salinometer cell is filled and rinsed three times with sample water before readings are recorded. Minimum three salinometer readings were recorded for every sample and standardization. If the values are fluctuating, more readings are taken.

Sampling Procedure and Data Processing Technique

Salinity samples are drawn into 150 ml medicine bottles after three rinses. The bottles are filled up to the shoulders and then capped with caps with plastic liners. Files for each separate run are prepared. These files consist of various metadata (date, cruise, lab temperature, geographic location, operator, etc.) and sample specific data such as the bath temperature, sample ID number, and average conductivity ratio. A PC based program computes the salinity using average conductivity ratio of the runs and the standard IAPSO formula. Any changes in the salinometer readings between successive standardizations are assumed to have occurred as a linear drift of the instrument. Thus, the program applies a correction to the ratios, which varies linearly with the samples analyzed. The salinity data is then placed in the water sample database.

Laboratory and Sample Temperatures

Full cases of samples are taken from the winch room to the salinometer lab where they are left for a period of at least 24 hours to equilibrate to laboratory temperature before being analyzed.

The bath in the salinometer was kept at 21°C.

Standards Used

The salinometer was standardized using IAPSO standard water, Batch P152, dated May, 2013. Standardization with a new bottle was carried out at the beginning and end of the run as well as after 50 bottle samples.

3.1.4 SBE 43 Dissolved Oxygen Sensor Calibration using Winkler Titrations

We use a method for statistically estimating calibration coefficients for calculating dissolved oxygen in milliliters per liter from SBE 43 output voltage. The technique requires dissolved oxygen concentration in ml/l (determined from Winkler titration of water samples) and SBE 43 oxygen voltage outputs at the times the water samples were collected. Sea-Bird's data processing software, SBE Data Processing, is used to produce a data table suitable for the analysis.

Background

The equation used in Sea-Bird's software for calculating dissolved oxygen in ml/l from SBE 43 output voltage is a form of that given in Owens-Millard (1985):

$$Oxygen(ml/l) = \{Soc * (V + Voffset + tau * \partial V/\partial t) + Boc * e^{(-0.03*T)}\} * Oxsat(T,S) * e^{(Tcorr*T)} * e^{(Pcor*P)} \quad eqn 1$$

Where:

V = SBE 43 output voltage signal (volts)

$\partial V/\partial t$ = time derivative of SBE 43 output signal (volts/second)

T = CTD temperature (°C)

S = CTD salinity (psu)

P = CTD pressure (dbars)

$Oxsat(T,S)$ = oxygen saturation (ml/l)

Soc , Boc , $Voffset$, tau , $tcor$, and $pcor$ are calibration coefficients

Characterization of the SBE 43 in the laboratory and ocean suggest that the most accurate results are obtained by setting Boc and tau to zero. Equation 1 then reduces to:

$$Oxygen(ml/l) = \{Soc * (V + Voffset)\} * Oxsat(T,S) * e^{(Tcorr*T)} * e^{(Pcor*P)} \quad eqn 2$$

The SBE 43 is expected to provide an output voltage that is linear with respect to oxygen concentration. Normal calibration drift manifests itself as a loss of sensitivity and is evident as a change of slope and offset in the linear relationship between oxygen concentration and voltage output. The coefficients $tcor$ and $pcor$ correct for small secondary responses to temperature and pressure. Because these coefficients change very slowly over time, the values given on the SBE 43 calibration certificate are used in this analysis.

Setting Boc and tau to zero, we will rearrange equation 2 into a linear form and perform a linear regression to obtain a new Soc and $Voffset$.

Let:

$$\phi = Oxsat(T,S) * e^{(Tcorr*T)} * e^{(Pcor*P)}$$

The oxygen equation then reduces to:

$$Oxygen(ml/l) = Soc * (V + Voffset) * \phi$$

This may be expressed in a linear form as shown in equation 3 below. A linear regression is calculated using Winkler oxygen concentration divided by ϕ as the dependent variable and SBE 43 output voltage as the independent variable.

$$Oxygen(ml/l) / \phi = Soc * (V + Voffset) = M * V + B \quad eqn 3$$

Where:

$$Soc = M$$

$$Voffset = B / M$$

Winkler oxygen divided by ϕ versus SBE 43 output voltage for this cruise is shown in

Fig. 12 and includes a linear regression line calculated from the data.

The Soc value is 4.0237e-001.

The $Voffset$ value is -0.3524 for the sill sections

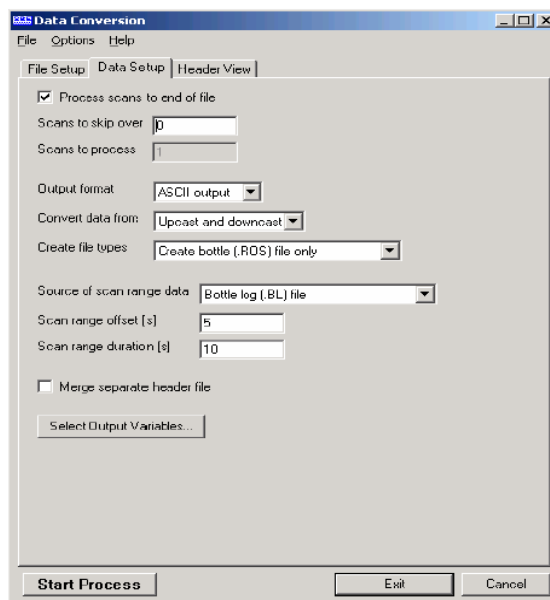
871 oxygen samples were used to build this linear fits.

Procedure

As a first step, extract pressure, temperature, salinity, oxygen saturation, and SBE 43 voltage from the parts of your CTD data collected when the water sampler closures occurred.

We run SBE Data Processing, and select Data Conversion in the Run menu. Select the appropriate configuration (.con) and data (.dat) files on the *File Setup* tab. In the *Data Setup* tab we set *Convert data from* to *Upcast and downcast* and *Create file types* to *Create bottle (.ros) file only*.

To extract CTD data concurrent to the water sampler closures, Data Conversion must know when the closures occurred. Select an appropriate *Source of scan range data*, depending on your instrument type and how the sampler was commanded to close bottles:

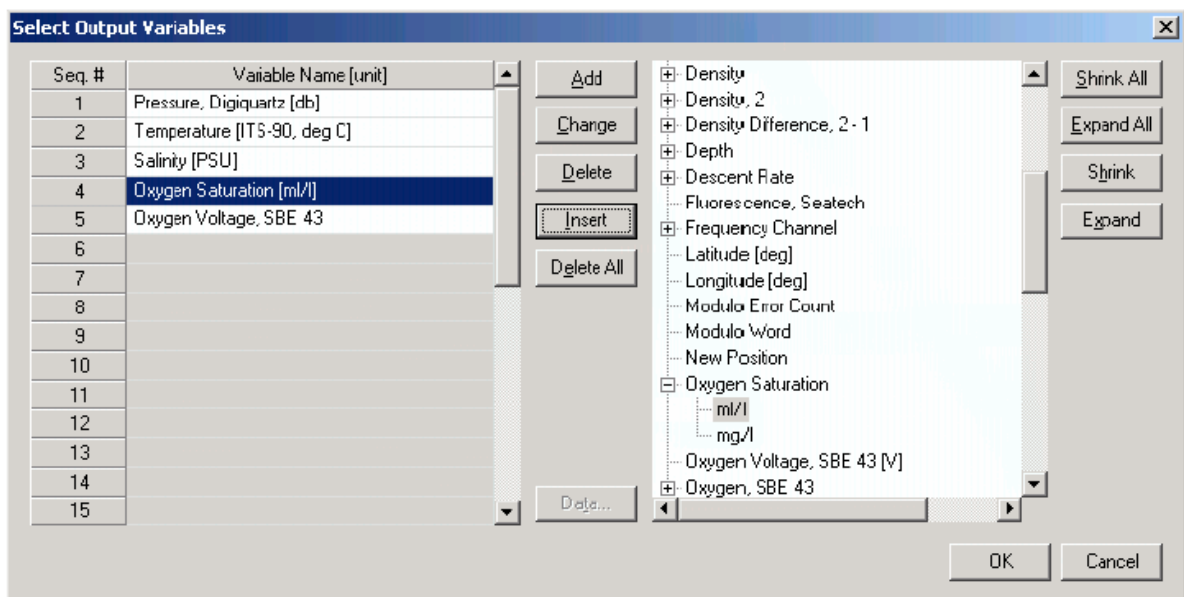


- SBE 9plus with SBE 11plus - The data stream is marked with a *bottle confirm* bit each time a closure occurred.
- Using SEASAVE to operate the water sampler - A .bl file, with scan ranges corresponding to closures, is created during the cast.

Like all sensors, the SBE 43 has a finite response time to a change in dissolved oxygen concentration. This response time is usually on the order of 6 seconds. For this reason, good sampling procedure dictates that the instrument package should be stopped in the water column long enough for the SBE 43 and all other sensors to completely equilibrate before closing the water sampler. An equilibration time of 5 to 6 response times, or 30 to 36 seconds, is adequate. We used to wait 30 seconds.

Data Conversion is extracting data 5 seconds before each water sampler closure and will extract 10 seconds of data. Note that 10 seconds is longer than the SBE 43 response time. Because we are extracting data for 5 seconds after the water sampler closure, the instrument package must remain stopped for at least this long.

To estimate *Soc* and *Voffset*, we need pressure, temperature, salinity, oxygen saturation (ml/l), and SBE 43 Oxygen Voltage to go with each Winkler titration data value. We use *Select Output Variables* and add each of the required parameters; the dialog box is shown below.



After selecting all the variables, click *OK* to return to the Data Conversion Data Setup tab. Then click *Start Process* to create the *.ros* file.

In our case, the *.ros* file contains 10 seconds of data centered on the moment the bottle closure occurred for every bottle closure. To make a useful table, we select Rosette Summary from SBE Data Processing's Run menu. Rosette Summary calculates averages and standard deviations for the variables selected in Data Conversion. Select the appropriate *.con* and *.ros* files on the *File Setup* tab. To average the data in the *Data Setup* tab we click the *Select Averaged Variables* button; after selecting all the variables, click *OK* to return to the Rosette Summary Data Setup tab. Then click *Start Process* to create a data table file with the *.btl* extension.

Further, we run a program READBTLDATA to create a file with average pressure, temperature, salinity, oxygen saturation, and SBE 43 output voltage for each water sampler closure depth, by importing the *.btl* file and the Winkler titration dissolved oxygen values from our titration log, matching water sampler closures to number of station, bottle number and

pressures. The program also calculates ϕ , using $tcor$ and $pcor$ from the SBE 43 calibration sheet.

Then, calculate *Winkler O₂* / ϕ .

Using the table we perform a linear regression, with:

- *Winkler O₂* / ϕ (shown as Winkler/phi in the table) as the *Y* data
- SBE 43 output voltages as the *X* data

Reference

Owens, W. B., and R. C. Millard Jr., 1985: A new algorithm for CTD oxygen calibration. *J. Physical Oceanography*, 15, 621-631.

(NOTE: calibration expressed as ml/l)

3.2 Oxygen Bottle Samples

Oxygen samples were drawn first from every bottle. Duplicate samples were taken on each cast, usually from the first two bottles. Samples were drawn into clear, wide necked calibrated glass bottles and fixed on deck with reagents dispensed using Aquastep bottle top dispensers. A test station used to check on the oxygen bottle calibrations and as an opportunity to train a number of people to take the samples. The samples were shaken on deck and again in the laboratory 1/2 hour after collection, when the bottles were checked for the tightness of the stoppers and presence of bubbles. The samples were then stored under water until analysis.

Bottle temperatures were taken, following sampling for oxygen, using a hand held electronic thermometer probe. The temperatures were used to calculate any temperature-dependent changes in the sample bottle volumes.

Samples were analyzed in the constant temperature laboratory, starting three hours after sample collection, following the Winkler whole bottle titration with an amperometric method of endpoint detection, as described by Culberson (1991). The equipment used was supplied by Metrohm and included the Titrino unit and control pad, exchange unit with 10 ml burette to dispense the thiosulphate in increments of 2 μ l, with an electrode for amperometric end point detection.

The difference for the duplicate pairs sampled on each station was in a range 0.00-0.02 ml/l (Table 2).

The thiosulphate normality was checked on each run and recalculated every time the reservoir was topped up against potassium iodate. The exact weight of this standard, the calibrated 5 ml exchange unit driven by a Metrohm Dosimat and the 1L glass volumetric flask used to dispense and prepare the standard.

The introduction of oxygen with the reagents and impurities in the manganese chloride were corrected for by blank measurements made on each run, as described in the WOCE Manual of Operations and Methods (Culberson, 1991).

Collected data shows that dissolved oxygen concentrations varied from 5.18 to 8.96 ml/l. In order to control the accuracy of the oxygen measurements at each cast were taken parallel samples from the 1-2 bottles or duplicate samples.

Reproducibility of measurements

3393 samples were taken during the cruise; in addition, 441 duplicates were analyzed. Statistics on the duplicates are given in Table 2. These include both duplicates taken from the same bottle (replicates) and those taken from different bottles fired at the same depth. The data gave a standard deviation of 0.005 ml/l, or 0.081%.

3.3 Nutrient Bottle Samples

Samples for nutrient measurements were collected following oxygen samples from each Niskin bottle. Water was collected in clean plastic containers that had been rinsed three times by seawater through the latex tube.

Concentrations of silicate and phosphate were determined by photometric methods with spectrophotometer Cary 100 Seam Varian. All samples were analyzed immediately after sampling.

Silicate determined by Korolev's method based on colorimeter of blue silicomolybdic complex (methodology described in Modern methods..., 1992). The ascorbic acid used as a restorative. The absorbance was read at 810 nm. Relative error of this method on concentration of dissolved silicate at 4.5 μM is $\pm 4\%$, on concentration at 45 μM - $\pm 2,5\%$. Measured concentrations were in a range from 0.51 to 11.77 μM .

Phosphates determined according to the method Murphy and Raily (Modern methods..., 1992). Phosphate, dissolved in sea water, react with ammonium molybdate in a presence of sulfuric acid and tartrate potassium-antimony. The generated complex aggregate of phosphomolybdic heteropolyacid and trivalent antimony restorative by the ascorbic acid, and then determined the absorbance at 885 nm (we use the cavity 10cm). Relative error of this method $\pm 1\%$.

In order to ensure accuracy and increase precision of determination 3-8 duplicate samples were analyzed at each run. The mean difference for the duplicate pairs sampled on each station was in a error limits of the methods (Table 2).

References:

Culberson, C.H. 1991. 15 pp in the WOCE Operations Manual (WHP Operations and Methods) WHPO 91/1, Woods Hole.

Modern methods of hydrochemical research of the ocean, 1992. IO RAS, Moscow (in Russian).

(O. Lukyanova)

3.4 Lowered Acoustic Doppler Current Profiler (LADCP)

The TRDI WHS 300 kHz ADCPs consists of a pressure case rated to 6000 metres with 4 transducers at one end in a convex arrangement and the beams diverging at 20 degrees from the vertical. At the opposite end to the transducers is a connector that enables downloading of data and connects it to other pressure cases containing another ADCP and the power supply pack. This arrangement allowed the ADCPs and the battery pack to be mounted vertically as up and down-looking on the CTD frame. Connection amongst all units was established using star cable with three male and two female terminations. Two male cable ends were always attached to the frame, this enabled comms leads to be readily connected pre and post deployment.

Communications: The 20-m communication leads (which also allow external power to be supplied to the ADCP) were sufficiently long to route it through to the port side of the deck lab where it was connected to a dedicated PC and external power supply. The latter was set at 48+ volts and was left on whilst the ADCP was on deck. 5 minutes prior to deployment the external power supply was shut off, the instrument checked and the configuration file sent to the ADCP as described in the manual instructions. The free end of the fly leads was greased and the end cap refitted, this was then taped to the frame for security.

Post deployment: When the CTD/LADCP was brought inboard, the fly-lead connectors were dried and the comms leads were connected to them. This stopped undue bending of the cables and kept them clear of the water bottles, aiding sampling. External power was applied again and the cast data downloaded as per the manual with a baud rate of 57600. The processing is accomplished using software developed by Visbeck after transferring the data to the PC.

Battery power was supplied to the ADCP in the form of 42 volts from 28 x 1.5 volt alkaline cells. Four of these packs were available for the cruise, as the ADCP will function at a minimum of 32 volts this was deemed an adequate stock for the duration.

Data quality: The data quality from the ADCP was good throughout. Due to the bad weather instrument titles sometimes exceeded 12° and this data was rejected during processing.

The LADCPs seem to function well and generates useful information on currents. The battery supply has its limitations though and thought should be given to alternatives to the present set-up.

3.4.1 LADCP Processing for Current Profiles

A brief account of the LADCP current data processing, file nomenclature and directory structure is provided in the following lines. Little emphasis is put into a detailed description of the main programming tools used, since these are part of a standard software package developed by Gerd Krahnemann (version 10.13).

Outline of LADCP current calculation method

The Broad Band LADCP used during AI39 cruise was designed to measure the instantaneous relative velocities of scatterers in the water column by taking advantage of the Doppler frequency shift, phase changes and correlation between coded pulses transmitted and received by the LADCP's four transducers. Conversion of this raw data stream to a profile of absolute currents involved an elaborate calculation method.

Firstly, Doppler shifts needed to be scaled to velocity units by taking into account the depth-dependent sound velocity (estimated from CTD T and S measurements). Directions could be inferred from trigonometric calculations based on the geometry of the transducer set, the orientation of the package (measured with a flux gate compass) and the local magnetic declination. The depth of the instrument was calculated from the integration of the measured vertical velocity and later adjusted to match the depth given by the CTD's pressure sensor.

The velocities corresponding to each single ensemble (or, in effect, to each transducer ping) were gridded in bins of depth set 10 meters. Statistical rejection of spiky measurements within each of these bins followed.

In order to reject the unwanted motion of the instrument (but also the barotropic component of the current), shear profiles were calculated for each ensemble. A complicated editing scheme preceded this shear calculation. A final shear profile (baroclinic current) was derived by real- depth gridding of the shear profiles calculated for individual ensembles. It was hoped that any relative velocities introduced by the high-frequency motion of the CTD package would be smoothed out by this repeated averaging.

The barotropic component of the flow was finally calculated from bottom-tracking measurements (bottom-track mode) or, in most occasions, in an integral sense from differential GPS positions of the ship (water-track mode).

The definitive velocity profile was hence obtained as the sum of the baroclinic and barotropic components.

During AI39 cruise, no specific error calculation was performed. Profiles of shear standard deviation were included in the cast log sheet folder. Internal wave signals were obvious throughout the cruise.

Relevant PC files

The raw data were downloaded from the LADCP into a devoted PC after each cast and stored as a binary file called vNNNNm_01.000 for Master and vNNNNs_01.000 for Slave the c:\ladcp\AI39\dNNNN directory, where NNNN stands for the CTD cast number, e.g. raw data from cast 2827 were stored in the files d:\AI39\data\ladcp\v2827m_01.000 and v2827s_01..000.

The configuration files (named Mconf.txt and Sconf.txt) containing the operating instructions (setting of track mode, bin depth, etc.) given to the LADCP previously to deployment was stored in the same directory.

Text files of the form NNNNm.log and NNNNs.log are the log of the 'bbtalk' session (testing the state and functioning of the instrument) previous to deployment. The details of the sessions for every single cast in the cruise are to be found in the cast log sheets.

A whole variety of files were created and manipulated during the different processing stages, and no mention will be made of the majority of them for reasons of clarity. The processing procedure may be summarised in two steps:

- 1- create CTD pressure, temperature and salinity data file as well as navigation collected every second in order to obtain the best possible estimates of depth and sound velocity. This is done using 'SBE Data Processing software and ConvLADCP Fortran program.
- 2- use the Gerd Krahnmann's standard matlab package (v. 10.13) with P. Lherminier's improvements (IFREMER) to process LADCP and CTD data

References

M. Visbeck 1994: Deep Velocity Profiling using Lowered Acoustic Doppler Current Profiler: Bottom Track and Inverse Solutions J. Atmos. Oceanic Technol. 10, 764-773.

4. CRUISE LOGISTICS

Mobilization

Mobilization for the cruise took place on the way from Kangerlussuaq (Greenland) to the first station of the cruise. It took six days. The scientific team arrived at the ship on September 10th.

ACKNOWLEDGEMENTS

The principal scientists would like to thank the Master, officers, crew and scientists of the RV Akademik Ioffe for making this such an enjoyable, as well as successful cruise.

TABLES

Table 1. CTD casts

Table 2. Performance of chemical analysis along the sill sections in 39 cruise R/V *Akademik Ioffe*

FIGURES

Fig. 1 Station location and ship track (in red). The shelf area with depth less than 200 m is shaded

Fig. 2 Vertical distribution of samples (a) along the sill sections, (b) along the 59.5 section.

Fig. 3. Schematic diagram of the large-scale circulation in the northern North Atlantic compiled from [Schmitz and McCartney, 1993; Schott and Brandt, 2007; Sutherland and Pickart, 2008; Lherminier et al., 2010]. Abbreviations for the main topographic features, currents and water masses are explained in the legend. The nominal locations of the 59.5°N hydrographic section (1997 – present) and sections across the straits between Greenland, Iceland, Faeroe and Shetland Islands (2011 – present) are shown with the solid green lines.

Fig. 4. Oxygen concentrations (ml/l) in the water column (lower panel) as observed in March–October 1997 in four hydrographic sections (upper panel) ending nearby the southern tip of Greenland. A separate oxygen maximum in the LSW layer (1000–2000 m) in the Irminger Sea at 59.5°N strongly implies local convective renewal of LSW before 1997. Adapted from [Falina et al., 2007].

Fig. 5. Warming and salinification in the northern North Atlantic between the mid-1990s and mid-2000s, as observed at 59.5°N. The figure shows the 2006–1997 temperature (°C, left) and salinity (right) differences on isobaric surfaces in the Irminger Sea and Iceland Basin. Adapted from [Sarafanov et al., 2007].

Fig. 6. Coherence of the decadal salinity changes (1950s – 2000s) of the intermediate (LSW) and deep (ISOW) waters in the northern North Atlantic and their link to the North Atlantic Oscillation (NAO) index. **(a)** Schematic representation of the LSW and ISOW pathways and locations of the Icelandic Low (L) and Azores High (H) centers constituting the NAO dipole pattern. The red dotted line indicates the 59.5°N transatlantic section. **(b)** Salinity time series for LSW in the Labrador Sea [Yashayaev, 2007] and ISOW in the Iceland basin [Boessenkool et al., 2007; Sarafanov et al., 2007] overlaid by the third order polynomial fits. **(c)** Time series of the winter NAO index, after [Hurrell, 1995], overlaid by 7-year running mean and third order polynomial fit. **(d)** Mechanism of the NAO effect on the decadal changes in temperature (T) and salinity (S) of the northern North Atlantic intermediate and deep waters. Positive / negative links shown with the dark / light grey arrows mean that changes in ‘causative’ and ‘consequential’ characteristics have the same / opposite sign(s). The overall effect of the NAO on T and S of the in the water column is negative: persistent NAO decline leads to warming and salinification of the water masses and vice versa, as shown in (b) and (c). Adapted from [Sarafanov, 2009].

Fig. 7. Schematic representation of the upper-ocean circulation and convection intensity in the northern North Atlantic under high (left) and low (right) NAO conditions. Blue (magenta) solid arrows indicate the upper-ocean flows with higher fraction of colder fresher subpolar (warmer saltier subtropical) waters. The main pathways of the Nordic overflow-derived deep waters are shown with the dotted curves. “C” and “E” symbols are used to denote, respectively, the deep convection sites and the domain, where the Atlantic waters are entrained into ISOW. Larger (smaller) circles indicate stronger (weaker) convection. SPG and STG – the subpolar and subtropical gyres, respectively. Adapted from [Sarafanov, 2009].

Fig. 8. The Deep Western Boundary Current (DWBC) transport variability and its link to the convection intensity in the Labrador Sea. **(a)** Locations of the hydrographic sections (1991–2007) and schematic of the deep water circulation in the Irminger Sea. **(b)** The DWBC transport anomalies at Cape Farewell in 1991–2007, $1 \text{ Sv} = 10^6 \text{ m}^3 \text{ s}^{-1}$. The 1994–1997 and 2000–2007 mean anomalies and the 1994–2007 linear trend are shown. **(c)** Anomalies of the DWBC transport at Cape Farewell and the Labrador Sea Water (LSW) thickness in the Labrador Sea in the 1950s–2000s. **(d)** Correlation coefficient (R^2) for the two times series shown in **(c)** at the 0–5-year lag, the LSW thickness leads. The correlation maximum is achieved at the 1–3-year lag. The DWBC transport anomalies in the southern Irminger Sea are foregone by the convection intensity anomalies in the Labrador Sea. Adapted from [Sarafanov et al., 2009].

Fig. 9. Schematic diagram of the Meridional Overturning Circulation (MOC) at the northern periphery of the Atlantic Ocean, northeast of Cape Farewell. The dotted lines refer to the σ_0 isopycnals 27.55 and 27.80. The arrows denote the integral meridional and diapycnal volume fluxes. Where the signs are specified, the positive (negative) transports are northward (southward). The NAC and EGIC transports in the upper layer ($\sigma_0 < 27.55$) at 59.5°N are the throughputs accounting for the recirculations. EGIC – the East Greenland / Irminger Current – refers to the upper part of the Western Boundary Current. Other abbreviations are explained in the legend to **Fig. 3**. Adapted from [Sarafanov et al., 2012].

Fig. 10. Salinity observed in the northwestern Irminger Sea at 64.3°N in February 1998. The σ_0 isopycnals 27.55, 27.70, 27.80 and 27.88 are plotted as the thick black lines; the station locations are marked with the ticks on the top axis. The plot shows fresh dense waters descending (cascading) down the continental slope of Greenland down to the LSW layer ($27.70 < \sigma_0 < 27.80$) and the layer of the Nordic Seas overflow-derived deep waters ($\sigma_0 > 27.80$). Adapted from [Falina et al., 2012].

Fig. 11 Autosal 8400B CTD salinity difference for primary (in black) and secondary (in red) conductivity sensors based on 860 salinity samples measurements (16 runs) during 38 cruise of R/V *Akademik Ioffe*. Vertical bars show standard deviation of each run.

Fig. 12 Regression line for Winkler oxygen divided by ϕ versus SBE 43 output voltage the sill sections, (b) 59.5 section.

Fig. 13 The vertical distribution of (a) potential temperature and (b) salinity and (c) dissolved oxygen between Greenland and Iceland 16-18 September 2012. Density is shown in black.

Fig. 14 The vertical distribution of (a) potential temperature and (b) salinity and (c) dissolved oxygen between Greenland and Iceland 18-19 September 2012. Density is shown in black.

Fig. 15 The vertical distribution of (a) potential temperature and (b) salinity and (c) dissolved oxygen between Greenland and Iceland 19-21 September 2012. Density is shown in black.

Fig. 16 The vertical distribution of (a) potential temperature and (b) salinity and (c) dissolved oxygen between Iceland and Faroe Islands 24-25 September 2012. Density is shown in black.

Fig. 17 The vertical distribution of (a) potential temperature and (b) salinity and (c) dissolved oxygen between Shetlands and Faroe Islands 26-27 September 2012. Density is shown in black.

SHIP/CRS	CLIVAR	CAST	UTC	POSITION								
STNNB R	CASTN O	TYPE	DATE	TIME	CODE	LATITUD E	LONGITUD E	NAV	DEPTH	BOTTO M	BOTTLES	COMMENTS
2827	1	ROS	091612	1532	BE	67 24.3 N	033 04.9 W	GPS	306.88	4	12	CTD,LADCP,O2,SiO3,PO4
2827	1	ROS	091612	1541	BO	67 24.3 N	033 04.9 W	GPS	306.88	4	12	CTD,LADCP,O2,SiO3,PO4
2827	1	ROS	091612	1557	EN	67 24.2 N	033 05.0 W	GPS	306.88	4	12	CTD,LADCP,O2,SiO3,PO4
2828	1	ROS	091612	1704	BE	67 19.8 N	032 44.0 W	GPS	436.89	7	14	CTD,LADCP,O2,SiO3,PO4
2828	1	ROS	091612	1716	BO	67 19.7 N	032 43.9 W	GPS	436.89	7	14	CTD,LADCP,O2,SiO3,PO4
2828	1	ROS	091612	1735	EN	67 19.7 N	032 43.9 W	GPS	436.89	7	14	CTD,LADCP,O2,SiO3,PO4
2829	1	ROS	091612	1846	BE	67 15.1 N	032 22.4 W	GPS	362.88	4	13	CTD,LADCP,O2,SiO3,PO4
2829	1	ROS	091612	1858	BO	67 15.1 N	032 22.6 W	GPS	362.88	4	13	CTD,LADCP,O2,SiO3,PO4
2829	1	ROS	091612	1915	EN	67 15.2 N	032 22.7 W	GPS	362.88	4	13	CTD,LADCP,O2,SiO3,PO4
2830	1	ROS	091612	2033	BE	67 09.9 N	031 58.9 W	GPS	300.98	5	13	CTD,LADCP,O2,SiO3,PO4
2830	1	ROS	091612	2050	BO	67 09.9 N	031 58.9 W	GPS	300.98	5	13	CTD,LADCP,O2,SiO3,PO4
2830	1	ROS	091612	2107	EN	67 09.9 N	031 58.9 W	GPS	300.98	5	13	CTD,LADCP,O2,SiO3,PO4
2831	1	ROS	091612	2223	BE	67 04.9 N	031 35.6 W	GPS	295.25	4	14	CTD,LADCP,O2,SiO3,PO4
2831	1	ROS	091612	2234	BO	67 04.8 N	031 35.5 W	GPS	295.25	4	14	CTD,LADCP,O2,SiO3,PO4
2831	1	ROS	091612	2250	EN	67 04.7 N	031 35.0 W	GPS	295.25	4	14	CTD,LADCP,O2,SiO3,PO4
2832	1	ROS	091612	2358	BE	67 00.1 N	031 12.7 W	GPS	353.76	5	15	CTD,LADCP,O2,SiO3,PO4
2832	1	ROS	091712	0012	BO	66 59.9 N	031 12.1 W	GPS	353.76	5	15	CTD,LADCP,O2,SiO3,PO4
2832	1	ROS	091712	0031	EN	66 59.7 N	031 11.6 W	GPS	353.76	5	15	CTD,LADCP,O2,SiO3,PO4
2833	1	ROS	091712	0142	BE	66 55.0 N	030 49.7 W	GPS	571.74	4	17	CTD,LADCP,O2,SiO3,PO4
2833	1	ROS	091712	0157	BO	66 55.0 N	030 49.5 W	GPS	571.74	4	17	CTD,LADCP,O2,SiO3,PO4
2833	1	ROS	091712	0223	EN	66 54.7 N	030 49.1 W	GPS	571.74	4	17	CTD,LADCP,O2,SiO3,PO4
2834	1	ROS	091712	0337	BE	66 49.9 N	030 27.1 W	GPS	408.75	5	15	CTD,LADCP,O2,SiO3,PO4
2834	1	ROS	091712	0354	BO	66 49.8 N	030 26.7 W	GPS	408.75	5	15	CTD,LADCP,O2,SiO3,PO4
2834	1	ROS	091712	0414	EN	66 49.6 N	030 26.2 W	GPS	408.75	5	15	CTD,LADCP,O2,SiO3,PO4
2835	1	ROS	091712	0530	BE	66 45.0 N	030 02.8 W	GPS	326.32	4	14	CTD,LADCP,O2,SiO3,PO4
2835	1	ROS	091712	0540	BO	66 44.9 N	030 02.8 W	GPS	326.32	4	14	CTD,LADCP,O2,SiO3,PO4
2835	1	ROS	091712	0557	EN	66 44.8 N	030 02.7 W	GPS	326.32	4	14	CTD,LADCP,O2,SiO3,PO4
2836	1	ROS	091712	0717	BE	66 39.9 N	029 38.8 W	GPS	299.57	6	11	CTD,LADCP,O2,SiO3,PO4

2836	1	ROS	091712	0726	BO	66 39.8 N	029 38.8 W	GPS	299.57	6	11	CTD,LADCP,O2,SiO3,PO4
2836	1	ROS	091712	0740	EN	66 39.8 N	029 38.7 W	GPS	299.57	6	11	CTD,LADCP,O2,SiO3,PO4
2837	1	ROS	091712	0854	BE	66 35.2 N	029 16.4 W	GPS	308.69	5	11	CTD,LADCP,O2,SiO3,PO4
2837	1	ROS	091712	0909	BO	66 35.1 N	029 16.2 W	GPS	308.69	5	11	CTD,LADCP,O2,SiO3,PO4
2837	1	ROS	091712	0923	EN	66 35.2 N	029 16.3 W	GPS	308.69	5	11	CTD,LADCP,O2,SiO3,PO4
2838	1	ROS	091712	1037	BE	66 30.1 N	028 54.1 W	GPS	317.64	6	12	CTD,LADCP,O2,SiO3,PO4
2838	1	ROS	091712	1049	BO	66 30.0 N	028 53.9 W	GPS	317.64	6	12	CTD,LADCP,O2,SiO3,PO4
2838	1	ROS	091712	1104	EN	66 30.0 N	028 53.6 W	GPS	317.64	6	12	CTD,LADCP,O2,SiO3,PO4
2839	1	ROS	091712	1213	BE	66 25.1 N	028 31.6 W	GPS	296.45	6	12	CTD,LADCP,O2,SiO3,PO4
2839	1	ROS	091712	1225	BO	66 24.8 N	028 31.4 W	GPS	296.45	6	12	CTD,LADCP,O2,SiO3,PO4
2839	1	ROS	091712	1242	EN	66 24.7 N	028 31.0 W	GPS	296.45	6	12	CTD,LADCP,O2,SiO3,PO4
2840	1	ROS	091712	1354	BE	66 20.0 N	028 08.5 W	GPS	339.91	5	13	CTD,LADCP,O2,SiO3,PO4
2840	1	ROS	091712	1404	BO	66 19.9 N	028 08.5 W	GPS	339.91	5	13	CTD,LADCP,O2,SiO3,PO4
2840	1	ROS	091712	1421	EN	66 19.8 N	028 08.4 W	GPS	339.91	5	13	CTD,LADCP,O2,SiO3,PO4
2841	1	ROS	091712	1538	BE	66 15.0 N	027 45.3 W	GPS	470.30	6	14	CTD,LADCP,O2,SiO3,PO4
2841	1	ROS	091712	1556	BO	66 14.7 N	027 45.4 W	GPS	470.30	6	14	CTD,LADCP,O2,SiO3,PO4
2841	1	ROS	091712	1617	EN	66 14.5 N	027 45.8 W	GPS	470.30	6	14	CTD,LADCP,O2,SiO3,PO4
2842	1	ROS	091712	1752	BE	66 09.0 N	027 14.9 W	GPS	522.86	6	17	CTD,LADCP,O2,SiO3,PO4
2842	1	ROS	091712	1806	BO	66 08.9 N	027 14.9 W	GPS	522.86	6	17	CTD,LADCP,O2,SiO3,PO4
2842	1	ROS	091712	1832	EN	66 08.7 N	027 14.6 W	GPS	522.86	6	17	CTD,LADCP,O2,SiO3,PO4
2843	1	ROS	091712	1919	BE	66 05.0 N	027 02.8 W	GPS	658.52	7	20	CTD,LADCP,O2,SiO3,PO4
2843	1	ROS	091712	1934	BO	66 04.9 N	027 02.6 W	GPS	658.52	7	20	CTD,LADCP,O2,SiO3,PO4
2843	1	ROS	091712	2004	EN	66 04.8 N	027 02.7 W	GPS	658.52	7	20	CTD,LADCP,O2,SiO3,PO4
2844	1	ROS	091712	2104	BE	66 01.0 N	026 48.0 W	GPS	452.30	6	17	CTD,LADCP,O2,SiO3,PO4
2844	1	ROS	091712	2117	BO	66 01.1 N	026 48.0 W	GPS	452.30	6	17	CTD,LADCP,O2,SiO3,PO4
2844	1	ROS	091712	2140	EN	66 01.4 N	026 47.8 W	GPS	452.30	6	17	CTD,LADCP,O2,SiO3,PO4
2845	1	ROS	091712	2256	BE	65 55.9 N	026 29.1 W	GPS	285.35	5	15	CTD,LADCP,O2,SiO3,PO4
2845	1	ROS	091712	2306	BO	65 56.0 N	026 29.1 W	GPS	285.35	5	15	CTD,LADCP,O2,SiO3,PO4
2845	1	ROS	091712	2325	EN	65 56.1 N	026 29.2 W	GPS	285.35	5	15	CTD,LADCP,O2,SiO3,PO4
2846	1	ROS	091812	0109	BE	65 49.9 N	026 00.0 W	GPS	220.76	5	13	CTD,LADCP,O2,SiO3,PO4
2846	1	ROS	091812	0119	BO	65 49.9 N	025 59.9 W	GPS	220.76	5	13	CTD,LADCP,O2,SiO3,PO4
2846	1	ROS	091812	0133	EN	65 49.9 N	025 59.8 W	GPS	220.76	5	13	CTD,LADCP,O2,SiO3,PO4
2847	1	ROS	091812	0251	BE	65 44.9 N	025 38.8 W	GPS	270.25	4	14	CTD,LADCP,O2,SiO3,PO4

2847	1	ROS	091812	0301	BO	65 44.8 N	025 38.7 W	GPS	270.25	4	14	CTD,LADCP,O2,SiO3,PO4
2847	1	ROS	091812	0316	EN	65 44.6 N	025 38.7 W	GPS	270.25	4	14	CTD,LADCP,O2,SiO3,PO4
2848	1	ROS	091812	0433	BE	65 39.9 N	025 15.9 W	GPS	082.13	5	8	CTD,LADCP,O2,SiO3,PO4
2848	1	ROS	091812	0440	BO	65 39.7 N	025 16.0 W	GPS	082.13	5	8	CTD,LADCP,O2,SiO3,PO4
2848	1	ROS	091812	0448	EN	65 39.5 N	025 16.0 W	GPS	082.13	5	8	CTD,LADCP,O2,SiO3,PO4
2849	1	ROS	091812	0600	BE	65 34.8 N	024 54.8 W	GPS	067.18	4	6	CTD,LADCP,O2,SiO3,PO4
2849	1	ROS	091812	0607	BO	65 34.8 N	024 54.7 W	GPS	067.18	4	6	CTD,LADCP,O2,SiO3,PO4
2849	1	ROS	091812	0615	EN	65 34.8 N	024 54.8 W	GPS	067.18	4	6	CTD,LADCP,O2,SiO3,PO4
2850	1	ROS	091812	0728	BE	65 40.0 N	025 15.7 W	GPS	086.30	3	8	CTD,LADCP,O2,SiO3,PO4
2850	1	ROS	091812	0739	BO	65 39.9 N	025 16.2 W	GPS	086.30	3	8	CTD,LADCP,O2,SiO3,PO4
2850	1	ROS	091812	0746	EN	65 39.9 N	025 16.2 W	GPS	086.30	3	8	CTD,LADCP,O2,SiO3,PO4
2851	1	ROS	091812	0857	BE	65 44.8 N	025 38.2 W	GPS	257.84	5	12	CTD,LADCP,O2,SiO3,PO4
2851	1	ROS	091812	0910	BO	65 45.0 N	025 38.8 W	GPS	257.84	5	12	CTD,LADCP,O2,SiO3,PO4
2851	1	ROS	091812	0924	EN	65 45.2 N	025 38.9 W	GPS	257.84	5	12	CTD,LADCP,O2,SiO3,PO4
2852	1	ROS	091812	1035	BE	65 49.8 N	025 59.7 W	GPS	222.30	6	12	CTD,LADCP,O2,SiO3,PO4
2852	1	ROS	091812	1046	BO	65 49.9 N	025 59.7 W	GPS	222.30	6	12	CTD,LADCP,O2,SiO3,PO4
2852	1	ROS	091812	1059	EN	65 50.0 N	025 59.6 W	GPS	222.30	6	12	CTD,LADCP,O2,SiO3,PO4
2853	1	ROS	091812	1234	BE	65 55.9 N	026 28.7 W	GPS	285.64	6	13	CTD,LADCP,O2,SiO3,PO4
2853	1	ROS	091812	1245	BO	65 56.0 N	026 28.6 W	GPS	285.64	6	13	CTD,LADCP,O2,SiO3,PO4
2853	1	ROS	091812	1301	EN	65 56.0 N	026 28.4 W	GPS	285.64	6	13	CTD,LADCP,O2,SiO3,PO4
2854	1	ROS	091812	1411	BE	66 00.8 N	026 47.6 W	GPS	437.52	4	17	CTD,LADCP,O2,SiO3,PO4
2854	1	ROS	091812	1428	BO	66 01.0 N	026 47.5 W	GPS	437.52	4	17	CTD,LADCP,O2,SiO3,PO4
2854	1	ROS	091812	1450	EN	66 01.0 N	026 47.3 W	GPS	437.52	4	17	CTD,LADCP,O2,SiO3,PO4
2855	1	ROS	091812	1552	BE	66 05.0 N	027 02.7 W	GPS	655.56	3	18	CTD,LADCP,O2,SiO3,PO4
2855	1	ROS	091812	1609	BO	66 04.9 N	027 02.6 W	GPS	655.56	3	18	CTD,LADCP,O2,SiO3,PO4
2855	1	ROS	091812	1637	EN	66 04.5 N	027 02.7 W	GPS	655.56	3	18	CTD,LADCP,O2,SiO3,PO4
2856	1	ROS	091812	1729	BE	66 09.1 N	027 14.7 W	GPS	517.64	6	17	CTD,LADCP,O2,SiO3,PO4
2856	1	ROS	091812	1746	BO	66 09.0 N	027 15.0 W	GPS	517.64	6	17	CTD,LADCP,O2,SiO3,PO4
2856	1	ROS	091812	1811	EN	66 08.8 N	027 14.7 W	GPS	517.64	6	17	CTD,LADCP,O2,SiO3,PO4
2857	1	ROS	091812	1948	BE	66 14.9 N	027 45.4 W	GPS	474.15	6	18	CTD,LADCP,O2,SiO3,PO4
2857	1	ROS	091812	1959	BO	66 14.8 N	027 45.6 W	GPS	474.15	6	18	CTD,LADCP,O2,SiO3,PO4
2857	1	ROS	091812	2023	EN	66 14.8 N	027 46.2 W	GPS	474.15	6	18	CTD,LADCP,O2,SiO3,PO4
2858	1	ROS	091812	2137	BE	66 19.9 N	028 08.1 W	GPS	340.76	5	15	CTD,LADCP,O2,SiO3,PO4

2858	1	ROS	091812	2148	BO	66 19.9 N	028 08.3 W	GPS	340.76	5	15	CTD,LADCP,O2,SiO3,PO4
2858	1	ROS	091812	2206	EN	66 19.8 N	028 09.0 W	GPS	340.76	5	15	CTD,LADCP,O2,SiO3,PO4
2859	1	ROS	091812	2323	BE	66 25.0 N	028 31.2 W	GPS	297.10	4	14	CTD,LADCP,O2,SiO3,PO4
2859	1	ROS	091812	2334	BO	66 25.2 N	028 31.1 W	GPS	297.10	4	14	CTD,LADCP,O2,SiO3,PO4
2859	1	ROS	091812	2351	EN	66 25.3 N	028 31.1 W	GPS	297.10	4	14	CTD,LADCP,O2,SiO3,PO4
2860	1	ROS	091912	0102	BE	66 29.9 N	028 54.0 W	GPS	319.15	5	12	CTD,LADCP,O2,SiO3,PO4
2860	1	ROS	091912	0113	BO	66 30.1 N	028 54.0 W	GPS	319.15	5	12	CTD,LADCP,O2,SiO3,PO4
2860	1	ROS	091912	0129	EN	66 30.1 N	028 53.7 W	GPS	319.15	5	12	CTD,LADCP,O2,SiO3,PO4
2861	1	ROS	091912	0246	BE	66 34.9 N	029 15.9 W	GPS	305.42	5	12	CTD,LADCP,O2,SiO3,PO4
2861	1	ROS	091912	0257	BO	66 34.9 N	029 15.8 W	GPS	305.42	5	12	CTD,LADCP,O2,SiO3,PO4
2861	1	ROS	091912	0312	EN	66 34.9 N	029 15.6 W	GPS	305.42	5	12	CTD,LADCP,O2,SiO3,PO4
2862	1	ROS	091912	0425	BE	66 39.9 N	029 38.7 W	GPS	296.45	6	13	CTD,LADCP,O2,SiO3,PO4
2862	1	ROS	091912	0434	BO	66 39.8 N	029 38.8 W	GPS	296.45	6	13	CTD,LADCP,O2,SiO3,PO4
2862	1	ROS	091912	0455	EN	66 39.6 N	029 38.8 W	GPS	296.45	6	13	CTD,LADCP,O2,SiO3,PO4
2863	1	ROS	091912	0608	BE	66 45.1 N	030 02.8 W	GPS	329.76	5	13	CTD,LADCP,O2,SiO3,PO4
2863	1	ROS	091912	0620	BO	66 45.1 N	030 03.0 W	GPS	329.76	5	13	CTD,LADCP,O2,SiO3,PO4
2863	1	ROS	091912	0637	EN	66 45.0 N	030 03.0 W	GPS	329.76	5	13	CTD,LADCP,O2,SiO3,PO4
2864	1	ROS	091912	0747	BE	66 49.9 N	030 26.6 W	GPS	404.76	5	16	CTD,LADCP,O2,SiO3,PO4
2864	1	ROS	091912	0801	BO	66 49.8 N	030 27.0 W	GPS	404.76	5	16	CTD,LADCP,O2,SiO3,PO4
2864	1	ROS	091912	0822	EN	66 49.7 N	030 27.0 W	GPS	404.76	5	16	CTD,LADCP,O2,SiO3,PO4
2865	1	ROS	091912	0938	BE	66 55.0 N	030 49.6 W	GPS	574.37	0	19	CTD,LADCP,O2,SiO3,PO4
2865	1	ROS	091912	0953	BO	66 55.0 N	030 49.6 W	GPS	574.37	0	19	CTD,LADCP,O2,SiO3,PO4
2865	1	ROS	091912	1023	EN	66 55.0 N	030 49.7 W	GPS	574.37	0	19	CTD,LADCP,O2,SiO3,PO4
2866	1	ROS	091912	1137	BE	66 59.9 N	031 12.4 W	GPS	353.70	4	15	CTD,LADCP,O2,SiO3,PO4
2866	1	ROS	091912	1150	BO	66 59.9 N	031 12.4 W	GPS	353.70	4	15	CTD,LADCP,O2,SiO3,PO4
2866	1	ROS	091912	1208	EN	67 00.0 N	031 12.4 W	GPS	353.70	4	15	CTD,LADCP,O2,SiO3,PO4
2867	1	ROS	091912	1323	BE	67 04.8 N	031 35.6 W	GPS	290.89	7	14	CTD,LADCP,O2,SiO3,PO4
2867	1	ROS	091912	1334	BO	67 04.9 N	031 35.7 W	GPS	290.89	7	14	CTD,LADCP,O2,SiO3,PO4
2867	1	ROS	091912	1351	EN	67 04.9 N	031 35.8 W	GPS	290.89	7	14	CTD,LADCP,O2,SiO3,PO4
2868	1	ROS	091912	1502	BE	67 09.8 N	031 58.7 W	GPS	301.49	2	14	CTD,LADCP,O2,SiO3,PO4
2868	1	ROS	091912	1513	BO	67 09.7 N	031 59.0 W	GPS	301.49	2	14	CTD,LADCP,O2,SiO3,PO4
2868	1	ROS	091912	1529	EN	67 09.6 N	031 59.3 W	GPS	301.49	2	14	CTD,LADCP,O2,SiO3,PO4
2869	1	ROS	091912	1638	BE	67 15.2 N	032 22.1 W	GPS	364.06	8	15	CTD,LADCP,O2,SiO3,PO4

2869	1	ROS	091912	1651	BO	67 15.1 N	032 22.4 W	GPS	364.06	8	15	CTD,LADCP,O2,SiO3,PO4
2869	1	ROS	091912	1711	EN	67 14.9 N	032 22.5 W	GPS	364.06	8	15	CTD,LADCP,O2,SiO3,PO4
2870	1	ROS	091912	1818	BE	67 19.8 N	032 44.0 W	GPS	455.84	8	14	CTD,LADCP,O2,SiO3,PO4
2870	1	ROS	091912	1830	BO	67 19.8 N	032 44.0 W	GPS	455.84	8	14	CTD,LADCP,O2,SiO3,PO4
2870	1	ROS	091912	1852	EN	67 19.8 N	032 43.9 W	GPS	455.84	8	14	CTD,LADCP,O2,SiO3,PO4
2871	1	ROS	091912	1954	BE	67 23.0 N	033 01.1 W	GPS	250.74	7	13	CTD,LADCP,O2,SiO3,PO4
2871	1	ROS	091912	2002	BO	67 22.9 N	033 01.2 W	GPS	250.74	7	13	CTD,LADCP,O2,SiO3,PO4
2871	1	ROS	091912	2017	EN	67 22.8 N	033 01.3 W	GPS	250.74	7	13	CTD,LADCP,O2,SiO3,PO4
2872	1	ROS	091912	2153	BE	67 19.8 N	032 44.0 W	GPS	428.74	7	14	CTD,LADCP,O2,SiO3,PO4
2872	1	ROS	091912	2204	BO	67 19.8 N	032 44.1 W	GPS	428.74	7	14	CTD,LADCP,O2,SiO3,PO4
2872	1	ROS	091912	2224	EN	67 19.8 N	032 44.2 W	GPS	428.74	7	14	CTD,LADCP,O2,SiO3,PO4
2873	1	ROS	091912	2335	BE	67 15.2 N	032 22.5 W	GPS	360.57	6	14	CTD,LADCP,O2,SiO3,PO4
2873	1	ROS	091912	2348	BO	67 15.2 N	032 22.3 W	GPS	360.57	6	14	CTD,LADCP,O2,SiO3,PO4
2873	1	ROS	092012	0006	EN	67 15.1 N	032 22.3 W	GPS	360.57	6	14	CTD,LADCP,O2,SiO3,PO4
2874	1	ROS	092012	0122	BE	67 09.9 N	031 59.0 W	GPS	301.47	7	14	CTD,LADCP,O2,SiO3,PO4
2874	1	ROS	092012	0133	BO	67 09.9 N	031 58.8 W	GPS	301.47	7	14	CTD,LADCP,O2,SiO3,PO4
2874	1	ROS	092012	0149	EN	67 10.0 N	031 58.6 W	GPS	301.47	7	14	CTD,LADCP,O2,SiO3,PO4
2875	1	ROS	092012	0300	BE	67 04.9 N	031 35.8 W	GPS	292.13	5	14	CTD,LADCP,O2,SiO3,PO4
2875	1	ROS	092012	0309	BO	67 05.0 N	031 35.5 W	GPS	292.13	5	14	CTD,LADCP,O2,SiO3,PO4
2875	1	ROS	092012	0326	EN	67 04.9 N	031 35.2 W	GPS	292.13	5	14	CTD,LADCP,O2,SiO3,PO4
2876	1	ROS	092012	0431	BE	67 00.0 N	031 12.5 W	GPS	358.74	7	12	CTD,LADCP,O2,SiO3,PO4
2876	1	ROS	092012	0451	BO	66 59.8 N	031 12.6 W	GPS	358.74	7	12	CTD,LADCP,O2,SiO3,PO4
2876	1	ROS	092012	0508	EN	66 59.8 N	031 12.5 W	GPS	358.74	7	12	CTD,LADCP,O2,SiO3,PO4
2877	1	ROS	092012	0617	BE	66 55.1 N	030 49.9 W	GPS	576.25	5	19	CTD,LADCP,O2,SiO3,PO4
2877	1	ROS	092012	0635	BO	66 55.1 N	030 49.6 W	GPS	576.25	5	19	CTD,LADCP,O2,SiO3,PO4
2877	1	ROS	092012	0702	EN	66 55.0 N	030 49.3 W	GPS	576.25	5	19	CTD,LADCP,O2,SiO3,PO4
2878	1	ROS	092012	0814	BE	66 50.1 N	030 26.8 W	GPS	408.11	7	15	CTD,LADCP,O2,SiO3,PO4
2878	1	ROS	092012	0828	BO	66 50.0 N	030 26.6 W	GPS	408.11	7	15	CTD,LADCP,O2,SiO3,PO4
2878	1	ROS	092012	0849	EN	66 50.0 N	030 26.5 W	GPS	408.11	7	15	CTD,LADCP,O2,SiO3,PO4
2879	1	ROS	092012	1007	BE	66 45.1 N	030 02.9 W	GPS	327.57	6	15	CTD,LADCP,O2,SiO3,PO4
2879	1	ROS	092012	1019	BO	66 45.1 N	030 02.6 W	GPS	327.57	6	15	CTD,LADCP,O2,SiO3,PO4
2879	1	ROS	092012	1037	EN	66 45.1 N	030 02.2 W	GPS	327.57	6	15	CTD,LADCP,O2,SiO3,PO4
2880	1	ROS	092012	1154	BE	66 39.9 N	029 38.8 W	GPS	300.11	7	15	CTD,LADCP,O2,SiO3,PO4

2880	1	ROS	092012	1206	BO	66 39.9 N	029 38.5 W	GPS	300.11	7	15	CTD,LADCP,O2,SiO3,PO4
2880	1	ROS	092012	1223	EN	66 39.7 N	029 38.2 W	GPS	300.11	7	15	CTD,LADCP,O2,SiO3,PO4
2881	1	ROS	092012	1332	BE	66 35.0 N	029 16.3 W	GPS	309.20	5	14	CTD,LADCP,O2,SiO3,PO4
2881	1	ROS	092012	1346	BO	66 35.0 N	029 16.2 W	GPS	309.20	5	14	CTD,LADCP,O2,SiO3,PO4
2881	1	ROS	092012	1402	EN	66 35.0 N	029 16.3 W	GPS	309.20	5	14	CTD,LADCP,O2,SiO3,PO4
2882	1	ROS	092012	1519	BE	66 30.0 N	028 54.3 W	GPS	314.40	4	15	CTD,LADCP,O2,SiO3,PO4
2882	1	ROS	092012	1531	BO	66 30.0 N	028 54.1 W	GPS	314.40	4	15	CTD,LADCP,O2,SiO3,PO4
2882	1	ROS	092012	1547	EN	66 29.9 N	028 54.1 W	GPS	314.40	4	15	CTD,LADCP,O2,SiO3,PO4
2883	1	ROS	092012	1659	BE	66 25.1 N	028 31.5 W	GPS	297.15	6	14	CTD,LADCP,O2,SiO3,PO4
2883	1	ROS	092012	1709	BO	66 25.1 N	028 31.4 W	GPS	297.15	6	14	CTD,LADCP,O2,SiO3,PO4
2883	1	ROS	092012	1725	EN	66 25.1 N	028 31.2 W	GPS	297.15	6	14	CTD,LADCP,O2,SiO3,PO4
2884	1	ROS	092012	1840	BE	66 20.0 N	028 08.4 W	GPS	341.13	5	10	CTD,LADCP,O2,SiO3,PO4
2884	1	ROS	092012	1850	BO	66 20.0 N	028 08.5 W	GPS	341.13	5	10	CTD,LADCP,O2,SiO3,PO4
2884	1	ROS	092012	1904	EN	66 19.9 N	028 08.5 W	GPS	341.13	5	10	CTD,LADCP,O2,SiO3,PO4
2885	1	ROS	092012	2028	BE	66 14.9 N	027 45.3 W	GPS	474.42	5	12	CTD,LADCP,O2,SiO3,PO4
2885	1	ROS	092012	2044	BO	66 14.9 N	027 45.4 W	GPS	474.42	5	12	CTD,LADCP,O2,SiO3,PO4
2885	1	ROS	092012	2101	EN	66 14.8 N	027 45.6 W	GPS	474.42	5	12	CTD,LADCP,O2,SiO3,PO4
2886	1	ROS	092012	2241	BE	66 09.1 N	027 14.9 W	GPS	517.71	6	16	CTD,LADCP,O2,SiO3,PO4
2886	1	ROS	092012	2253	BO	66 09.1 N	027 14.5 W	GPS	517.71	6	16	CTD,LADCP,O2,SiO3,PO4
2886	1	ROS	092012	2317	EN	66 09.1 N	027 14.3 W	GPS	517.71	6	16	CTD,LADCP,O2,SiO3,PO4
2887	1	ROS	092112	0010	BE	66 05.2 N	027 03.0 W	GPS	653.96	7	20	CTD,LADCP,O2,SiO3,PO4
2887	1	ROS	092112	0026	BO	66 05.1 N	027 01.9 W	GPS	653.96	7	20	CTD,LADCP,O2,SiO3,PO4
2887	1	ROS	092112	0059	EN	66 05.3 N	027 01.0 W	GPS	653.96	7	20	CTD,LADCP,O2,SiO3,PO4
2888	1	ROS	092112	0154	BE	66 01.0 N	026 47.9 W	GPS	439.15	3	14	CTD,LADCP,O2,SiO3,PO4
2888	1	ROS	092112	0208	BO	66 01.0 N	026 47.3 W	GPS	439.15	3	14	CTD,LADCP,O2,SiO3,PO4
2888	1	ROS	092112	0227	EN	66 01.4 N	026 45.7 W	GPS	439.15	3	14	CTD,LADCP,O2,SiO3,PO4
2889	1	ROS	092112	0338	BE	65 56.0 N	026 29.1 W	GPS	285.47	4	13	CTD,LADCP,O2,SiO3,PO4
2889	1	ROS	092112	0348	BO	65 56.0 N	026 28.6 W	GPS	285.47	4	13	CTD,LADCP,O2,SiO3,PO4
2889	1	ROS	092112	0404	EN	65 56.1 N	026 27.9 W	GPS	285.47	4	13	CTD,LADCP,O2,SiO3,PO4
2890	1	ROS	092112	0544	BE	65 49.9 N	026 00.0 W	GPS	220.22	3	12	CTD,LADCP,O2,SiO3,PO4
2890	1	ROS	092112	0553	BO	65 50.0 N	025 59.7 W	GPS	220.22	3	12	CTD,LADCP,O2,SiO3,PO4
2890	1	ROS	092112	0607	EN	65 49.9 N	025 59.5 W	GPS	220.22	3	12	CTD,LADCP,O2,SiO3,PO4
2891	1	ROS	092112	0736	BE	65 45.0 N	025 38.7 W	GPS	269.83	2	13	CTD,LADCP,O2,SiO3,PO4

2891	1	ROS	092112	0746	BO	65 45.1 N	025 38.3 W	GPS	269.83	2	13	CTD,LADCP,O2,SiO3,PO4
2891	1	ROS	092112	0802	EN	65 45.1 N	025 38.1 W	GPS	269.83	2	13	CTD,LADCP,O2,SiO3,PO4
2892	1	ROS	092112	1100	BE	65 40.0 N	025 15.9 W	GPS	96.740	4	7	CTD,LADCP,O2,SiO3,PO4
2892	1	ROS	092112	1109	BO	65 40.1 N	025 15.8 W	GPS	96.740	4	7	CTD,LADCP,O2,SiO3,PO4
2892	1	ROS	092112	1118	EN	65 40.2 N	025 15.6 W	GPS	96.740	4	7	CTD,LADCP,O2,SiO3,PO4
2893	1	ROS	092112	1352	BE	65 34.9 N	024 54.9 W	GPS	65.270	5	6	CTD,LADCP,O2,SiO3,PO4
2893	1	ROS	092112	1411	BO	65 35.0 N	024 54.7 W	GPS	65.270	5	6	CTD,LADCP,O2,SiO3,PO4
2893	1	ROS	092112	1417	EN	65 35.0 N	024 54.5 W	GPS	65.270	5	6	CTD,LADCP,O2,SiO3,PO4
2894	1	ROS	092412	0151	BE	64 24.6 N	014 03.1 W	GPS	103.37	6	6	CTD,LADCP,O2,SiO3,PO4
2894	1	ROS	092412	0158	BO	64 24.7 N	014 03.3 W	GPS	103.37	6	6	CTD,LADCP,O2,SiO3,PO4
2894	1	ROS	092412	0205	EN	64 24.8 N	014 03.3 W	GPS	103.37	6	6	CTD,LADCP,O2,SiO3,PO4
2895	1	ROS	092412	0408	BE	64 17.0 N	013 35.9 W	GPS	166.53	6	9	CTD,LADCP,O2,SiO3,PO4
2895	1	ROS	092412	0415	BO	64 17.1 N	013 35.9 W	GPS	166.53	6	9	CTD,LADCP,O2,SiO3,PO4
2895	1	ROS	092412	0427	EN	64 17.2 N	013 35.9 W	GPS	166.53	6	9	CTD,LADCP,O2,SiO3,PO4
2896	1	ROS	092412	0530	BE	64 13.9 N	013 21.0 W	GPS	175.71	6	11	CTD,LADCP,O2,SiO3,PO4
2896	1	ROS	092412	0537	BO	64 14.0 N	013 21.0 W	GPS	175.71	6	11	CTD,LADCP,O2,SiO3,PO4
2896	1	ROS	092412	0550	EN	64 14.1 N	013 21.0 W	GPS	175.71	6	11	CTD,LADCP,O2,SiO3,PO4
2897	1	ROS	092412	0714	BE	64 07.9 N	013 03.0 W	GPS	166.74	4	10	CTD,LADCP,O2,SiO3,PO4
2897	1	ROS	092412	0721	BO	64 08.0 N	013 03.0 W	GPS	166.74	4	10	CTD,LADCP,O2,SiO3,PO4
2897	1	ROS	092412	0733	EN	64 08.0 N	013 02.9 W	GPS	166.74	4	10	CTD,LADCP,O2,SiO3,PO4
2898	1	ROS	092412	0827	BE	64 05.0 N	012 52.1 W	GPS	575.71	6	18	CTD,LADCP,O2,SiO3,PO4
2898	1	ROS	092412	0842	BO	64 05.1 N	012 51.8 W	GPS	575.71	6	18	CTD,LADCP,O2,SiO3,PO4
2898	1	ROS	092412	0908	EN	64 05.2 N	012 51.6 W	GPS	575.71	6	18	CTD,LADCP,O2,SiO3,PO4
2899	1	ROS	092412	1013	BE	64 01.0 N	012 38.2 W	GPS	565.37	6	18	CTD,LADCP,O2,SiO3,PO4
2899	1	ROS	092412	1029	BO	64 01.0 N	012 37.9 W	GPS	565.37	6	18	CTD,LADCP,O2,SiO3,PO4
2899	1	ROS	092412	1055	EN	64 01.1 N	012 37.7 W	GPS	565.37	6	18	CTD,LADCP,O2,SiO3,PO4
2900	1	ROS	092412	1211	BE	63 57.0 N	012 20.2 W	GPS	459.86	3	16	CTD,LADCP,O2,SiO3,PO4
2900	1	ROS	092412	1228	BO	63 56.9 N	012 20.2 W	GPS	459.86	3	16	CTD,LADCP,O2,SiO3,PO4
2900	1	ROS	092412	1250	EN	63 57.0 N	012 20.7 W	GPS	459.86	3	16	CTD,LADCP,O2,SiO3,PO4
2901	1	ROS	092412	1431	BE	63 50.0 N	012 00.1 W	GPS	388.67	7	15	CTD,LADCP,O2,SiO3,PO4
2901	1	ROS	092412	1443	BO	63 50.0 N	012 00.1 W	GPS	388.67	7	15	CTD,LADCP,O2,SiO3,PO4
2901	1	ROS	092412	1502	EN	63 50.1 N	012 00.1 W	GPS	388.67	7	15	CTD,LADCP,O2,SiO3,PO4
2902	1	ROS	092412	1630	BE	63 44.0 N	011 40.2 W	GPS	346.71	6	14	CTD,LADCP,O2,SiO3,PO4

2902	1	ROS	092412	1643	BO	63 44.0 N	011 40.0 W	GPS	346.71	6	14	CTD,LADCP,O2,SiO3,PO4
2902	1	ROS	092412	1700	EN	63 44.1 N	011 39.9 W	GPS	346.71	6	14	CTD,LADCP,O2,SiO3,PO4
2903	1	ROS	092412	1901	BE	63 36.0 N	011 15.0 W	GPS	302.44	3	13	CTD,LADCP,O2,SiO3,PO4
2903	1	ROS	092412	1911	BO	63 36.0 N	011 15.0 W	GPS	302.44	3	13	CTD,LADCP,O2,SiO3,PO4
2903	1	ROS	092412	1925	EN	63 36.1 N	011 14.8 W	GPS	302.44	3	13	CTD,LADCP,O2,SiO3,PO4
2904	1	ROS	092412	2112	BE	63 28.9 N	010 49.1 W	GPS	440.64	6	16	CTD,LADCP,O2,SiO3,PO4
2904	1	ROS	092412	2126	BO	63 28.9 N	010 49.3 W	GPS	440.64	6	16	CTD,LADCP,O2,SiO3,PO4
2904	1	ROS	092412	2146	EN	63 28.9 N	010 50.0 W	GPS	440.64	6	16	CTD,LADCP,O2,SiO3,PO4
2905	1	ROS	092412	2346	BE	63 19.9 N	010 25.1 W	GPS	331.52	10	14	CTD,LADCP,O2,SiO3,PO4
2905	1	ROS	092412	2357	BO	63 19.9 N	010 25.4 W	GPS	331.52	10	14	CTD,LADCP,O2,SiO3,PO4
2905	1	ROS	092512	0013	EN	63 20.0 N	010 25.9 W	GPS	331.52	10	14	CTD,LADCP,O2,SiO3,PO4
2906	1	ROS	092512	0202	BE	63 14.0 N	009 58.1 W	GPS	482.18	4	18	CTD,LADCP,O2,SiO3,PO4
2906	1	ROS	092512	0217	BO	63 13.9 N	009 58.0 W	GPS	482.18	4	18	CTD,LADCP,O2,SiO3,PO4
2906	1	ROS	092512	0239	EN	63 14.1 N	009 58.2 W	GPS	482.18	4	18	CTD,LADCP,O2,SiO3,PO4
2907	1	ROS	092512	0417	BE	63 05.9 N	009 38.0 W	GPS	492.98	5	16	CTD,LADCP,O2,SiO3,PO4
2907	1	ROS	092512	0430	BO	63 06.1 N	009 38.2 W	GPS	492.98	5	16	CTD,LADCP,O2,SiO3,PO4
2907	1	ROS	092512	0452	EN	63 06.3 N	009 38.5 W	GPS	492.98	5	16	CTD,LADCP,O2,SiO3,PO4
2908	1	ROS	092512	0649	BE	62 57.9 N	009 09.0 W	GPS	447.78	3	17	CTD,LADCP,O2,SiO3,PO4
2908	1	ROS	092512	0703	BO	62 58.0 N	009 09.0 W	GPS	447.78	3	17	CTD,LADCP,O2,SiO3,PO4
2908	1	ROS	092512	0725	EN	62 58.1 N	009 09.1 W	GPS	447.78	3	17	CTD,LADCP,O2,SiO3,PO4
2909	1	ROS	092512	0854	BE	62 50.9 N	008 50.0 W	GPS	462.03	7	17	CTD,LADCP,O2,SiO3,PO4
2909	1	ROS	092512	0907	BO	62 51.0 N	008 50.1 W	GPS	462.03	7	17	CTD,LADCP,O2,SiO3,PO4
2909	1	ROS	092512	0931	EN	62 50.9 N	008 50.7 W	GPS	462.03	7	17	CTD,LADCP,O2,SiO3,PO4
2910	1	ROS	092512	1115	BE	62 43.9 N	008 27.1 W	GPS	482.88	4	17	CTD,LADCP,O2,SiO3,PO4
2910	1	ROS	092512	1130	BO	62 44.0 N	008 27.3 W	GPS	482.88	4	17	CTD,LADCP,O2,SiO3,PO4
2910	1	ROS	092512	1152	EN	62 43.9 N	008 28.1 W	GPS	482.88	4	17	CTD,LADCP,O2,SiO3,PO4
2911	1	ROS	092512	1352	BE	62 35.0 N	008 02.0 W	GPS	400.88	1	15	CTD,LADCP,O2,SiO3,PO4
2911	1	ROS	092512	1418	BO	62 34.6 N	008 02.4 W	GPS	400.88	1	15	CTD,LADCP,O2,SiO3,PO4
2911	1	ROS	092512	1441	EN	62 34.9 N	008 02.6 W	GPS	400.88	1	15	CTD,LADCP,O2,SiO3,PO4
2912	1	ROS	092512	1629	BE	62 26.0 N	007 39.1 W	GPS	103.47	4	6	CTD,LADCP,O2,SiO3,PO4
2912	1	ROS	092512	1640	BO	62 26.0 N	007 38.8 W	GPS	103.47	4	6	CTD,LADCP,O2,SiO3,PO4
2912	1	ROS	092512	1647	EN	62 26.0 N	007 38.8 W	GPS	103.47	4	6	CTD,LADCP,O2,SiO3,PO4
2913	1	ROS	092512	1818	BE	62 20.0 N	007 15.0 W	GPS	091.42	5	6	CTD,LADCP,O2,SiO3,PO4

2913	1	ROS	092512	1828	BO	62 20.0 N	007 14.7 W	GPS	091.42	5	6	CTD,LADCP,O2,SiO3,PO4
2913	1	ROS	092512	1835	EN	62 20.1 N	007 14.6 W	GPS	091.42	5	6	CTD,LADCP,O2,SiO3,PO4
2914	1	ROS	092612	0221	BE	61 50.0 N	006 22.1 W	GPS	076.93	6	5	CTD,LADCP,O2,SiO3,PO4
2914	1	ROS	092612	0231	BO	61 49.9 N	006 22.0 W	GPS	076.93	6	5	CTD,LADCP,O2,SiO3,PO4
2914	1	ROS	092612	0236	EN	61 50.0 N	006 22.2 W	GPS	076.93	6	5	CTD,LADCP,O2,SiO3,PO4
2915	1	ROS	092612	0421	BE	61 42.6 N	005 57.5 W	GPS	299.98	5	14	CTD,LADCP,O2,SiO3,PO4
2915	1	ROS	092612	0430	BO	61 42.6 N	005 57.5 W	GPS	299.98	5	14	CTD,LADCP,O2,SiO3,PO4
2915	1	ROS	092612	0446	EN	61 42.8 N	005 57.5 W	GPS	299.98	5	14	CTD,LADCP,O2,SiO3,PO4
2916	1	ROS	092612	0615	BE	61 36.9 N	005 35.9 W	GPS	314.89	7	12	CTD,LADCP,O2,SiO3,PO4
2916	1	ROS	092612	0626	BO	61 37.0 N	005 35.8 W	GPS	314.89	7	12	CTD,LADCP,O2,SiO3,PO4
2916	1	ROS	092612	0642	EN	61 37.1 N	005 35.9 W	GPS	314.89	7	12	CTD,LADCP,O2,SiO3,PO4
2917	1	ROS	092612	0756	BE	61 30.9 N	005 19.1 W	GPS	286.30	6	13	CTD,LADCP,O2,SiO3,PO4
2917	1	ROS	092612	0808	BO	61 30.9 N	005 18.9 W	GPS	286.30	6	13	CTD,LADCP,O2,SiO3,PO4
2917	1	ROS	092612	0825	EN	61 31.0 N	005 19.0 W	GPS	286.30	6	13	CTD,LADCP,O2,SiO3,PO4
2918	1	ROS	092612	0945	BE	61 25.0 N	005 01.1 W	GPS	223.57	6	13	CTD,LADCP,O2,SiO3,PO4
2918	1	ROS	092612	0953	BO	61 24.9 N	005 01.3 W	GPS	223.57	6	13	CTD,LADCP,O2,SiO3,PO4
2918	1	ROS	092612	1008	EN	61 24.9 N	005 01.6 W	GPS	223.57	6	13	CTD,LADCP,O2,SiO3,PO4
2919	1	ROS	092612	1128	BE	61 19.9 N	004 43.0 W	GPS	722.76	5	20	CTD,LADCP,O2,SiO3,PO4
2919	1	ROS	092612	1146	BO	61 19.9 N	004 43.4 W	GPS	722.76	5	20	CTD,LADCP,O2,SiO3,PO4
2919	1	ROS	092612	1217	EN	61 19.7 N	004 44.2 W	GPS	722.76	5	20	CTD,LADCP,O2,SiO3,PO4
2920	1	ROS	092612	1333	BE	61 16.0 N	004 28.0 W	GPS	1049.13	5	21	CTD,LADCP,O2,SiO3,PO4
2920	1	ROS	092612	1400	BO	61 16.1 N	004 28.6 W	GPS	1049.13	5	21	CTD,LADCP,O2,SiO3,PO4
2920	1	ROS	092612	1437	EN	61 15.9 N	004 28.7 W	GPS	1049.13	5	21	CTD,LADCP,O2,SiO3,PO4
2921	1	ROS	092612	1537	BE	61 12.0 N	004 18.0 W	GPS	1079.30	6	21	CTD,LADCP,O2,SiO3,PO4
2921	1	ROS	092612	1603	BO	61 12.1 N	004 17.9 W	GPS	1079.30	6	21	CTD,LADCP,O2,SiO3,PO4
2921	1	ROS	092612	1640	EN	61 12.5 N	004 17.6 W	GPS	1079.30	6	21	CTD,LADCP,O2,SiO3,PO4
2922	1	ROS	092612	1748	BE	61 08.0 N	004 03.8 W	GPS	1119.52	7	21	CTD,LADCP,O2,SiO3,PO4
2922	1	ROS	092612	1811	BO	61 08.2 N	004 03.8 W	GPS	1119.52	7	21	CTD,LADCP,O2,SiO3,PO4
2922	1	ROS	092612	1848	EN	61 08.5 N	004 04.2 W	GPS	1119.52	7	21	CTD,LADCP,O2,SiO3,PO4
2923	1	ROS	092612	2017	BE	61 01.4 N	003 44.9 W	GPS	1099.23	6	21	CTD,LADCP,O2,SiO3,PO4
2923	1	ROS	092612	2040	BO	61 01.3 N	003 44.6 W	GPS	1099.23	6	21	CTD,LADCP,O2,SiO3,PO4
2923	1	ROS	092612	2117	EN	61 01.2 N	003 44.6 W	GPS	1099.23	6	21	CTD,LADCP,O2,SiO3,PO4
2924	1	ROS	092612	2157	BE	60 58.9 N	003 38.0 W	GPS	1100.88	4	21	CTD,LADCP,O2,SiO3,PO4

2924	1	ROS	092612	2222	BO	60 58.8 N	003 37.9 W	GPS	1100.88	4	21	CTD,LADCP,O2,SiO3,PO4
2924	1	ROS	092612	2300	EN	60 58.5 N	003 38.0 W	GPS	1100.88	4	21	CTD,LADCP,O2,SiO3,PO4
2925	1	ROS	092712	0023	BE	60 51.9 N	003 18.0 W	GPS	676.64	6	21	CTD,LADCP,O2,SiO3,PO4
2925	1	ROS	092712	0041	BO	60 51.8 N	003 18.0 W	GPS	676.64	6	21	CTD,LADCP,O2,SiO3,PO4
2925	1	ROS	092712	0109	EN	60 51.6 N	003 18.1 W	GPS	676.64	6	21	CTD,LADCP,O2,SiO3,PO4
2926	1	ROS	092712	0243	BE	60 45.0 N	002 55.0 W	GPS	352.57	6	15	CTD,LADCP,O2,SiO3,PO4
2926	1	ROS	092712	0254	BO	60 44.9 N	002 55.0 W	GPS	352.57	6	15	CTD,LADCP,O2,SiO3,PO4
2926	1	ROS	092712	0312	EN	60 44.8 N	002 54.8 W	GPS	352.57	6	15	CTD,LADCP,O2,SiO3,PO4
2927	1	ROS	092712	0435	BE	60 37.9 N	002 35.9 W	GPS	130.69	5	8	CTD,LADCP,O2,SiO3,PO4
2927	1	ROS	092712	0444	BO	60 37.9 N	002 35.9 W	GPS	130.69	5	8	CTD,LADCP,O2,SiO3,PO4
2927	1	ROS	092712	0453	EN	60 37.9 N	002 36.0 W	GPS	130.69	5	8	CTD,LADCP,O2,SiO3,PO4
2928	1	ROS	092712	0616	BE	60 30.9 N	002 15.9 W	GPS	141.42	5	8	CTD,LADCP,O2,SiO3,PO4
2928	1	ROS	092712	0623	BO	60 30.9 N	002 16.0 W	GPS	141.42	5	8	CTD,LADCP,O2,SiO3,PO4
2928	1	ROS	092712	0633	EN	60 30.9 N	002 16.0 W	GPS	141.42	5	8	CTD,LADCP,O2,SiO3,PO4
2929	1	ROS	092712	0801	BE	60 25.1 N	001 55.1 W	GPS	112.79	6	9	CTD,LADCP,O2,SiO3,PO4
2929	1	ROS	092712	0812	BO	60 25.0 N	001 54.9 W	GPS	112.79	6	9	CTD,LADCP,O2,SiO3,PO4
2929	1	ROS	092712	0821	EN	60 24.9 N	001 55.0 W	GPS	112.79	6	9	CTD,LADCP,O2,SiO3,PO4

Table 2

	O ₂ , ml/l	Silicate, μMol kg ⁻¹	Phosphate, μMol kg ⁻¹
Total amount of analyzed samples	1436	1498	1500
Number of duplicates	107	166	168
Mean difference of duplicates	0,007	0,210	0,010
Min difference of duplicates	0,000	0,000	0,000
Max difference of duplicates	0,021	0,926	0,047
Median difference of duplicates	0,005	0,134	0,008
Standard deviation of difference	0,005	0,208	0,009
Accuracy	0,081%	3,854%	1,302%

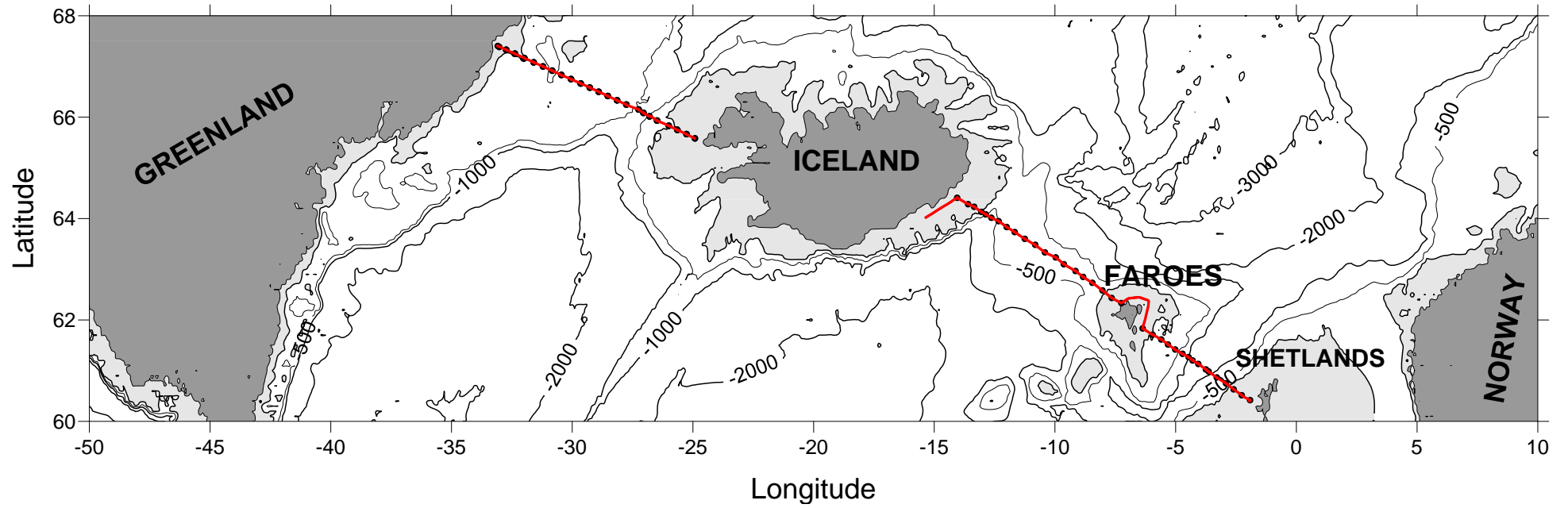


Fig. 1 Station locations (black circles) and ship track (in red). The shelf area with depth less than 200 m is shaded.

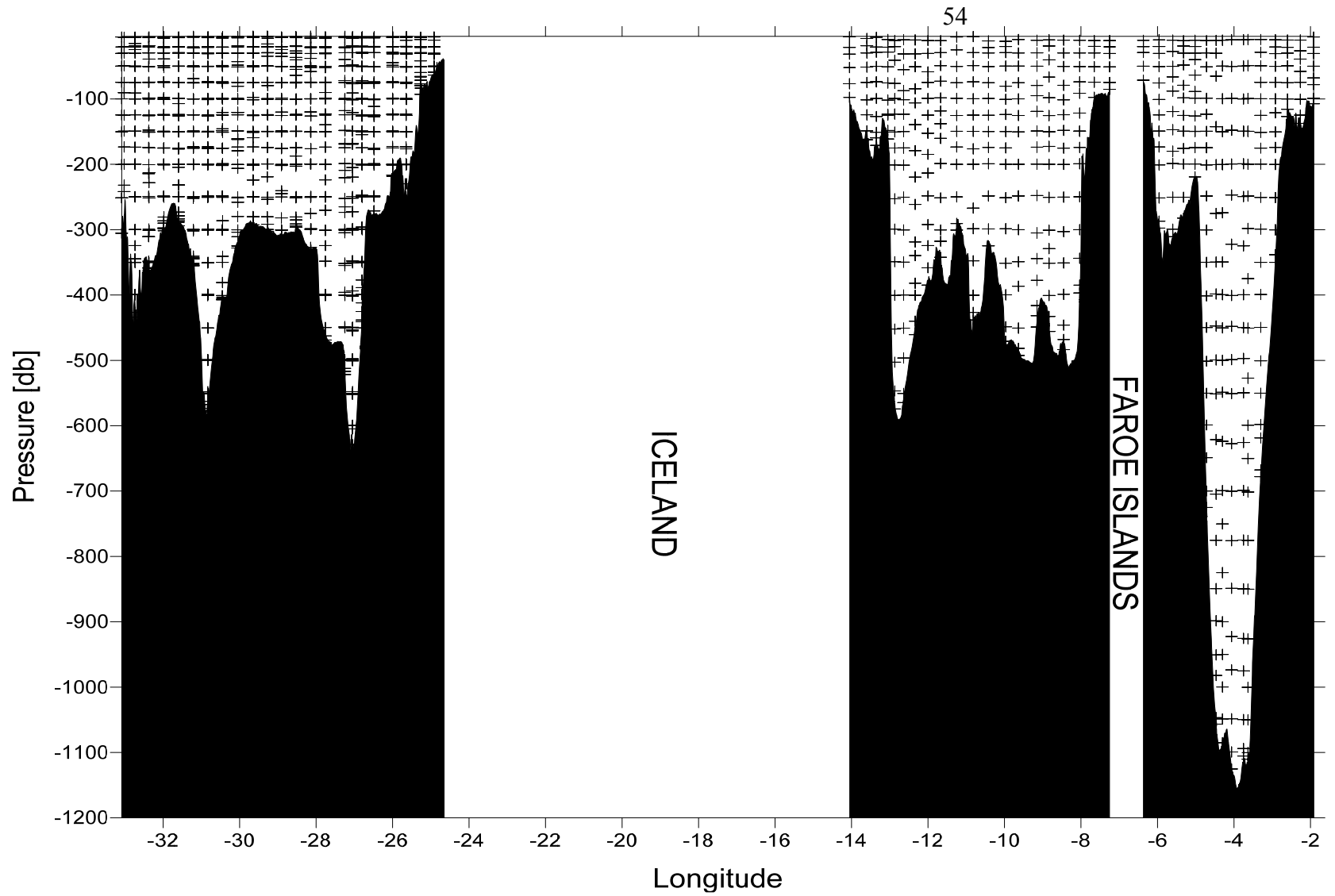


Figure 2. Vertical distribution of samples along the sill sections.

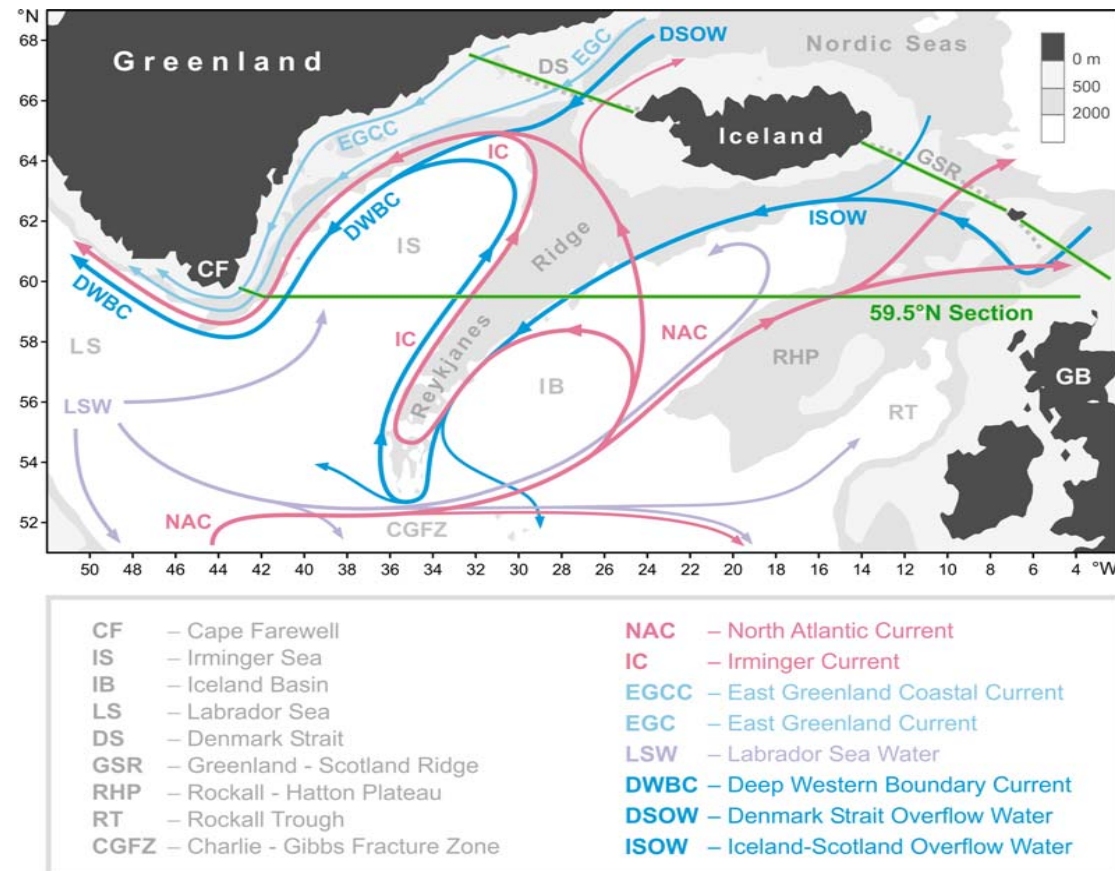


Figure 3. Schematic diagram of the large-scale circulation in the northern North Atlantic compiled from [Schmitz and McCartney, 1993; Schott and Brandt, 2007; Sutherland and Pickart, 2008; Lherminier et al., 2010]. Abbreviations for the main topographic features, currents and water masses are explained in the legend. The nominal locations of the 59.5°N hydrographic section (1997 – present) and sections across the straits between Greenland, Iceland, Faeroe and Shetland Islands (2011 – present) are shown with the solid green lines.

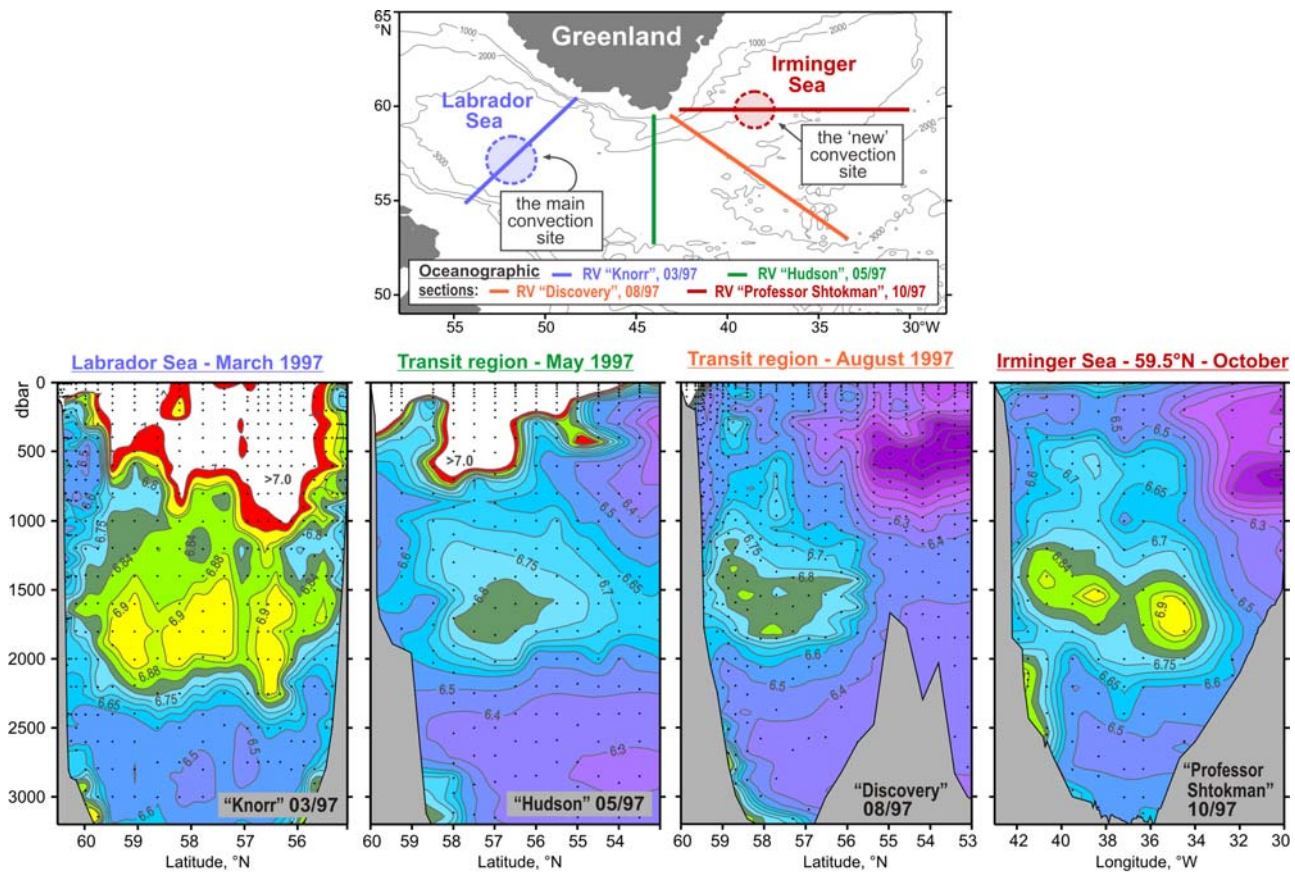


Figure 4. Oxygen concentrations (ml/l) in the water column (lower panel) as observed in March–October 1997 in four hydrographic sections (upper panel) ending nearby the southern tip of Greenland. A separate oxygen maximum in the LSW layer (1000–2000 m) in the Irminger Sea at 59.5°N strongly implies local convective renewal of LSW before 1997. Adapted from [Falina et al., 2007].

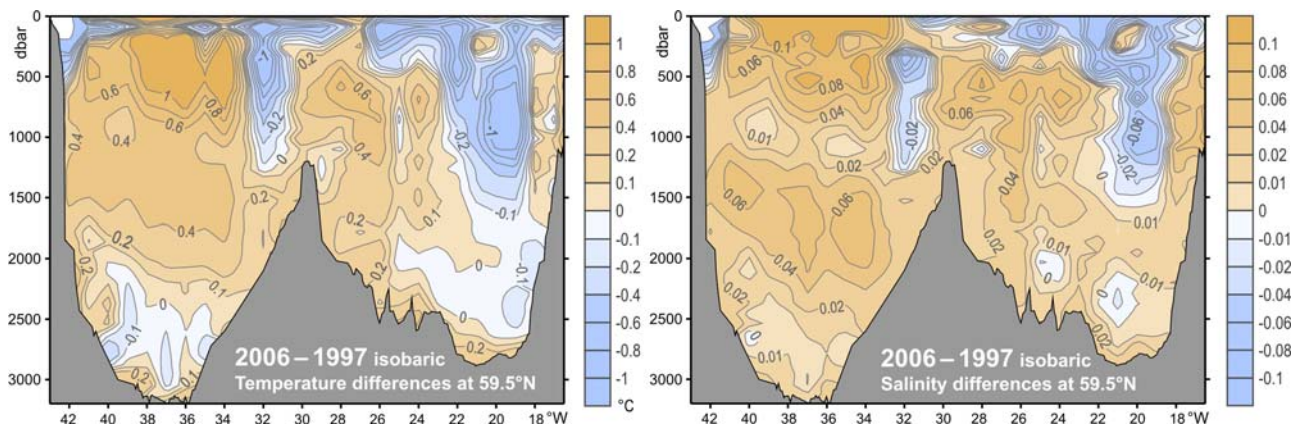


Figure 5. Warming and salinification in the northern North Atlantic between the mid-1990s and mid-2000s, as observed at 59.5°N. The figure shows the 2006–1997 temperature (°C, left) and salinity (right) differences on isobaric surfaces in the Irminger Sea and Iceland Basin. Adapted from [Sarafanov et al., 2007].

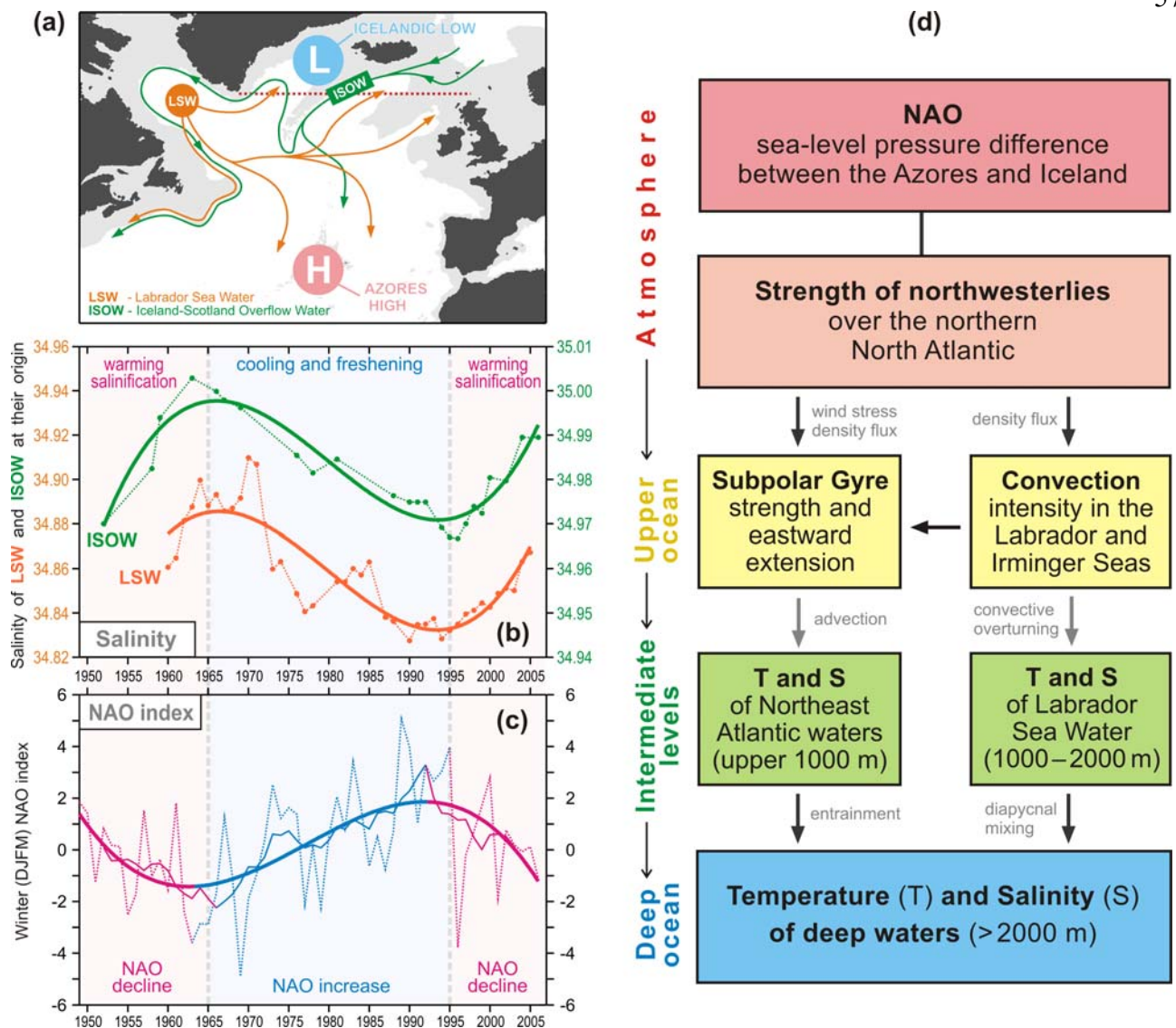


Figure 6. Coherence of the decadal salinity changes (1950s – 2000s) of the intermediate (LSW) and deep (ISOW) waters in the northern North Atlantic and their link to the North Atlantic Oscillation (NAO) index. **(a)** Schematic representation of the LSW and ISOW pathways and locations of the Icelandic Low (L) and Azores High (H) centers constituting the NAO dipole pattern. The red dotted line indicates the 59.5°N transatlantic section. **(b)** Salinity time series for LSW in the Labrador Sea [Yashayaev, 2007] and ISOW in the Iceland basin [Boessenkool et al., 2007; Sarafanov et al., 2007] overlaid by the third order polynomial fits. **(c)** Time series of the winter NAO index, after [Hurrell, 1995], overlaid by 7-year running mean and third order polynomial fit. **(d)** Mechanism of the NAO effect on the decadal changes in temperature (T) and salinity (S) of the northern North Atlantic intermediate and deep waters. Positive / negative links shown with the dark / light grey arrows mean that changes in ‘causative’ and ‘consequential’ characteristics have the same / opposite sign(s). The overall effect of the NAO on T and S of the in the water column is negative: persistent NAO decline leads to warming and salinification of the water masses and vice versa, as shown in (b) and (c). Adapted from [Sarafanov, 2009].

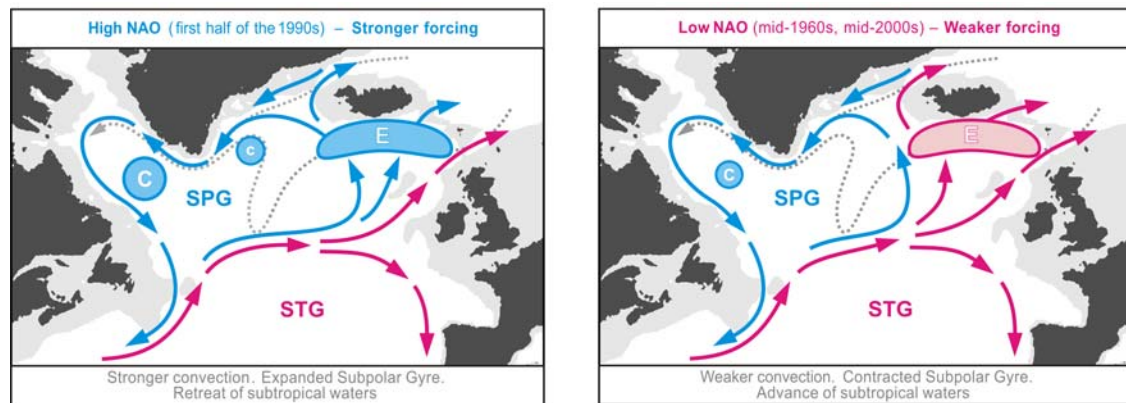


Figure 7. Schematic representation of the upper-ocean circulation and convection intensity in the northern North Atlantic under high (left) and low (right) NAO conditions. Blue (magenta) solid arrows indicate the upper-ocean flows with higher fraction of colder fresher subpolar (warmer saltier subtropical) waters. The main pathways of the Nordic overflow-derived deep waters are shown with the dotted curves. “C” and “E” symbols are used to denote, respectively, the deep convection sites and the domain, where the Atlantic waters are entrained into ISOW. Larger (smaller) circles indicate stronger (weaker) convection. SPG and STG – the subpolar and subtropical gyres, respectively. Adapted from [Sarafanov, 2009].

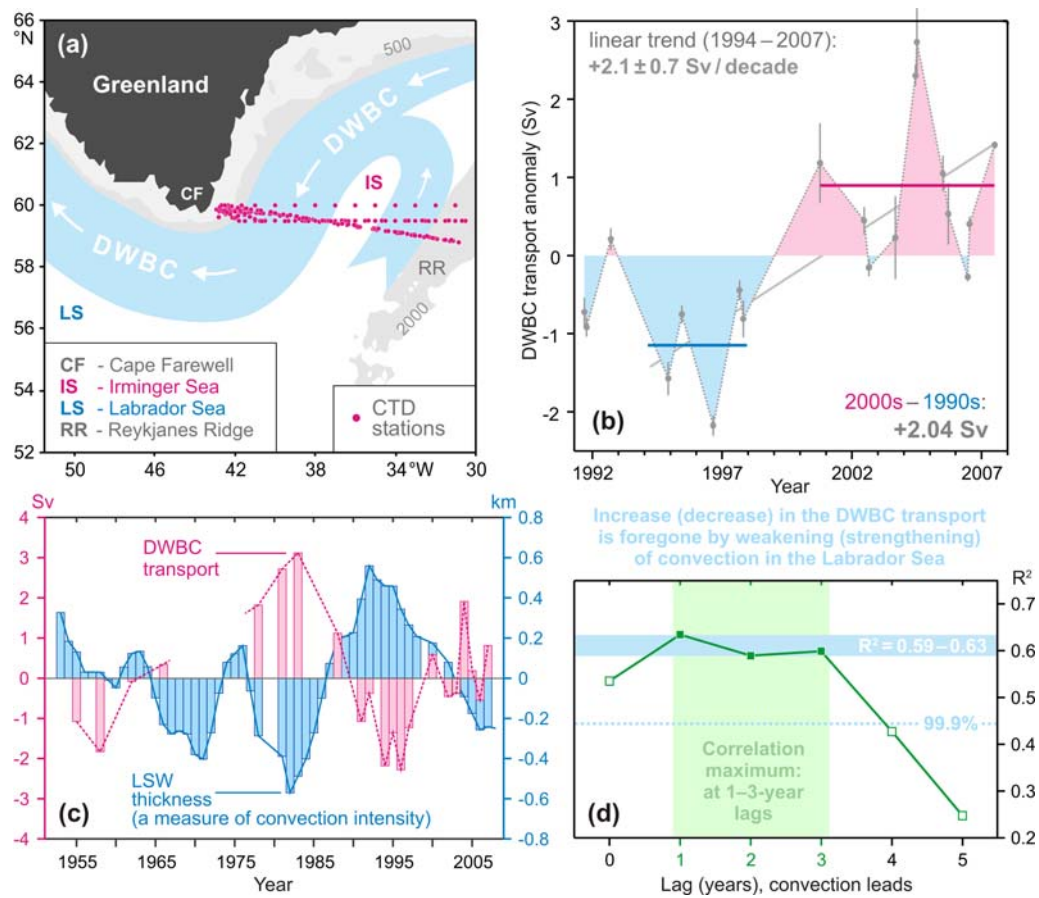


Figure 8. The Deep Western Boundary Current (DWBC) transport variability and its link to the convection intensity in the Labrador Sea. **(a)** Locations of the hydrographic sections (1991–2007) and schematic of the deep water circulation in the Irminger Sea. **(b)** The DWBC transport anomalies at Cape Farewell in 1991–2007, $1 \text{ Sv} = 10^6 \text{ m}^3 \text{ s}^{-1}$. The 1994–1997 and 2000–2007 mean anomalies and the 1994–2007 linear trend are shown. **(c)** Anomalies of the DWBC transport at Cape Farewell and the Labrador Sea Water (LSW) thickness in the Labrador Sea in the 1950s–2000s. **(d)** Correlation coefficient (R^2) for the two time series shown in (c) at the 0–5-year lag, the LSW thickness leads. The correlation maximum is achieved at the 1–3-year lag. The DWBC transport anomalies in the southern Irminger Sea are foregone by the convection intensity anomalies in the Labrador Sea. Adapted from [Sarafanov et al., 2009].

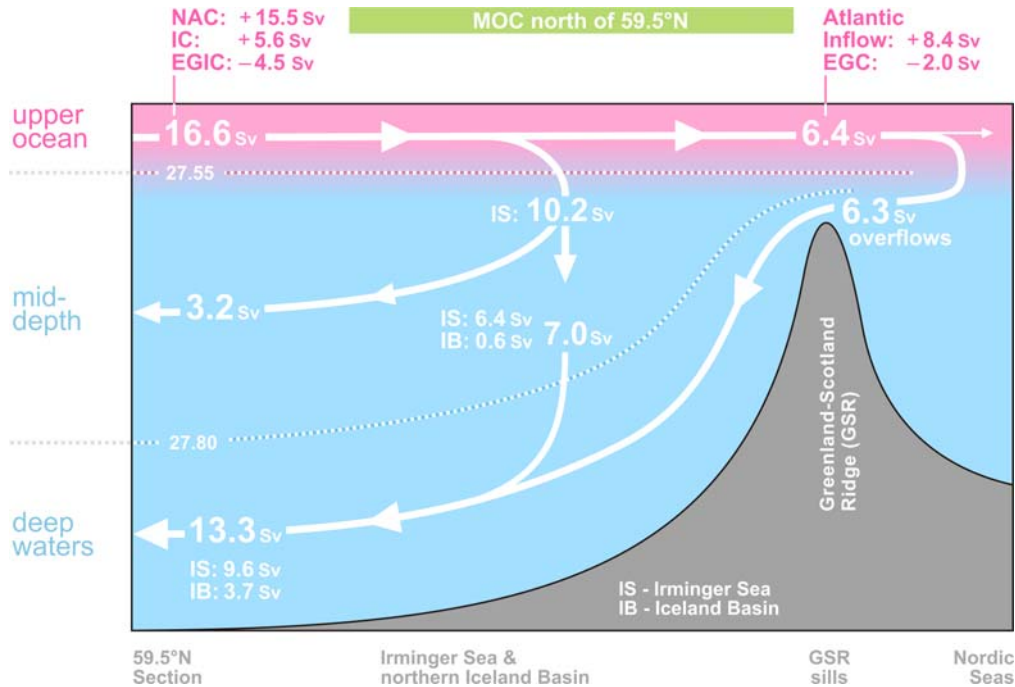


Figure 9. Schematic diagram of the Meridional Overturning Circulation (MOC) at the northern periphery of the Atlantic Ocean, northeast of Cape Farewell. The dotted lines refer to the σ_0 isopycnals 27.55 and 27.80. The arrows denote the integral meridional and diapycnal volume fluxes. Where the signs are specified, the positive (negative) transports are northward (southward). The NAC and EGIC transports in the upper layer ($\sigma_0 < 27.55$) at 59.5°N are the throughputs accounting for the recirculations. EGIC – the East Greenland / Irminger Current – refers to the upper part of the Western Boundary Current. Other abbreviations are explained in the legend to **Figure 3**. Adapted from [Sarafanov et al., 2012].

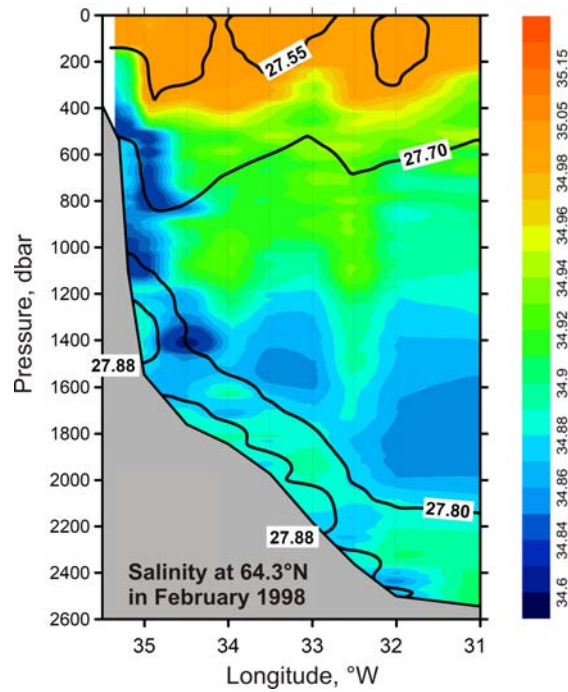


Figure 10. Salinity observed in the northwestern Irminger Sea at 64.3°N in February 1998. The σ_0 isopycnals 27.55, 27.70, 27.80 and 27.88 are plotted as the thick black lines; the station locations are marked with the ticks on the top axis. The plot shows fresh dense waters descending (cascading) down the continental slope of Greenland down to the LSW layer ($27.70 < \sigma_0 < 27.80$) and the layer of the Nordic Seas overflow-derived deep waters ($\sigma_0 > 27.80$). Adapted from [Falina et al., 2012].

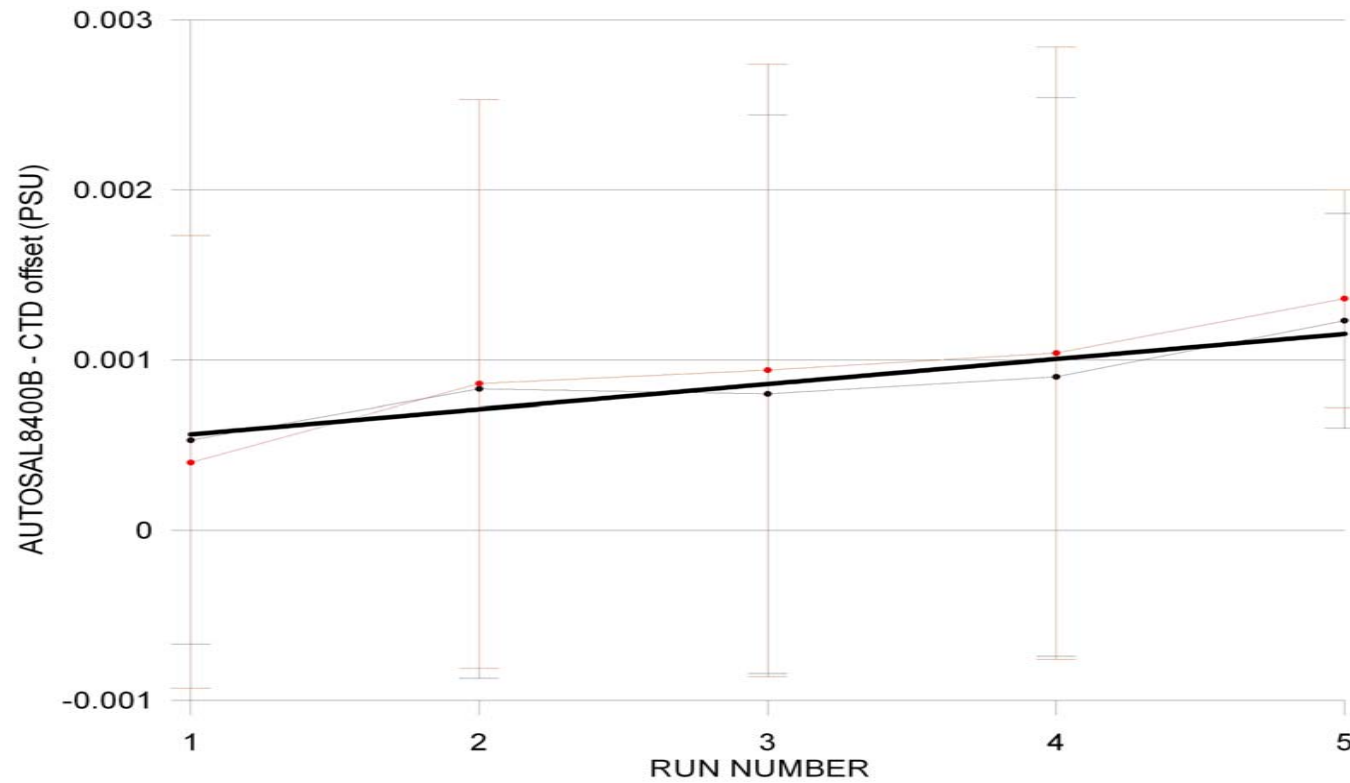


Figure 11 Autosal 8400B CTD salinity offset for the primary (in black) and the secondary (in red) conductivity sensors based on 327 salinity samples measurements (5 runs) during 39 cruise of R/V *Akademik Ioffe*. Vertical bars show standard deviation of each run. Thick black line denotes linear fit.

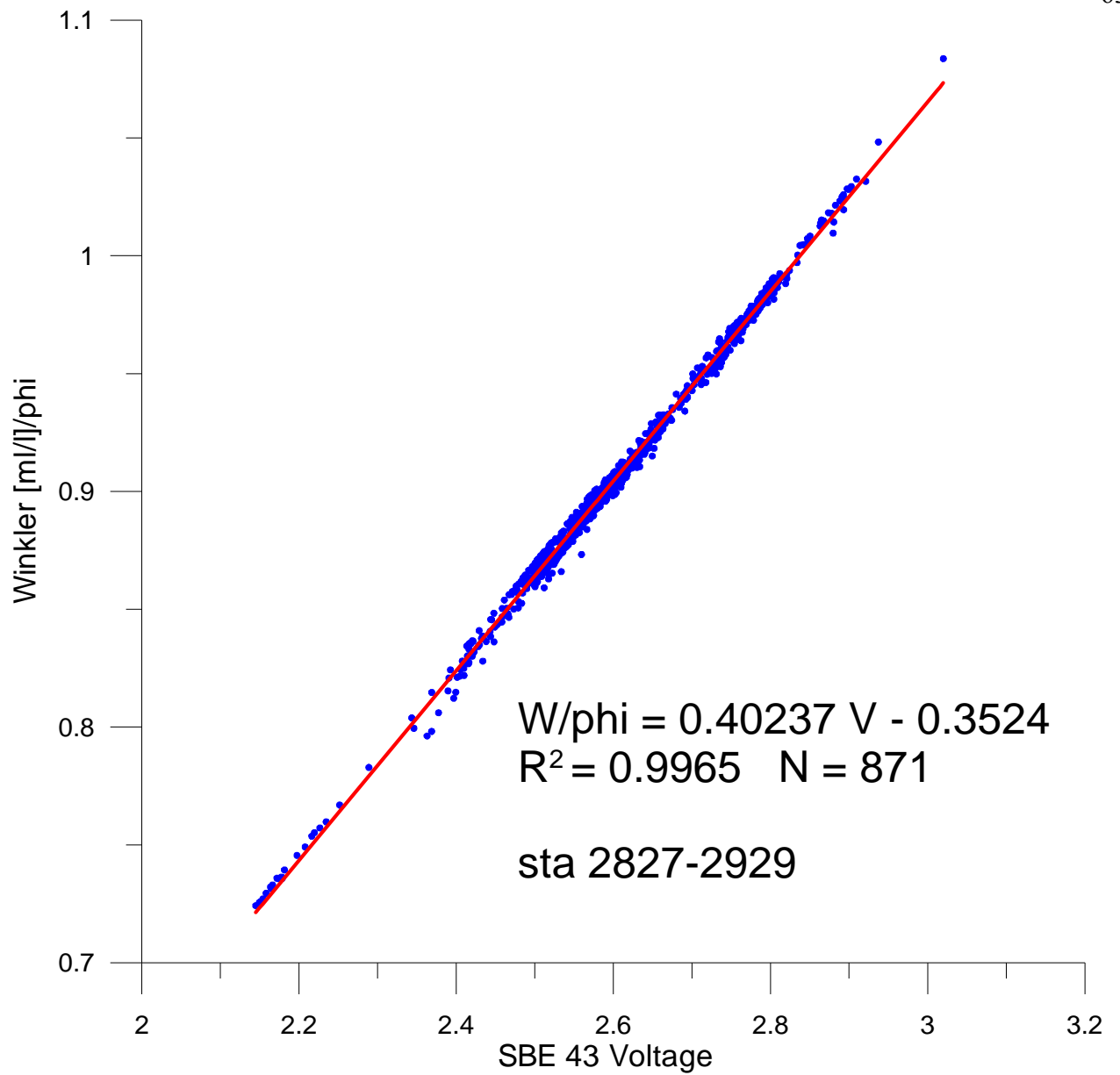


Figure 12 Regression line for Winkler oxygen divided by ϕ versus SBE 43 output voltage the sill sections.

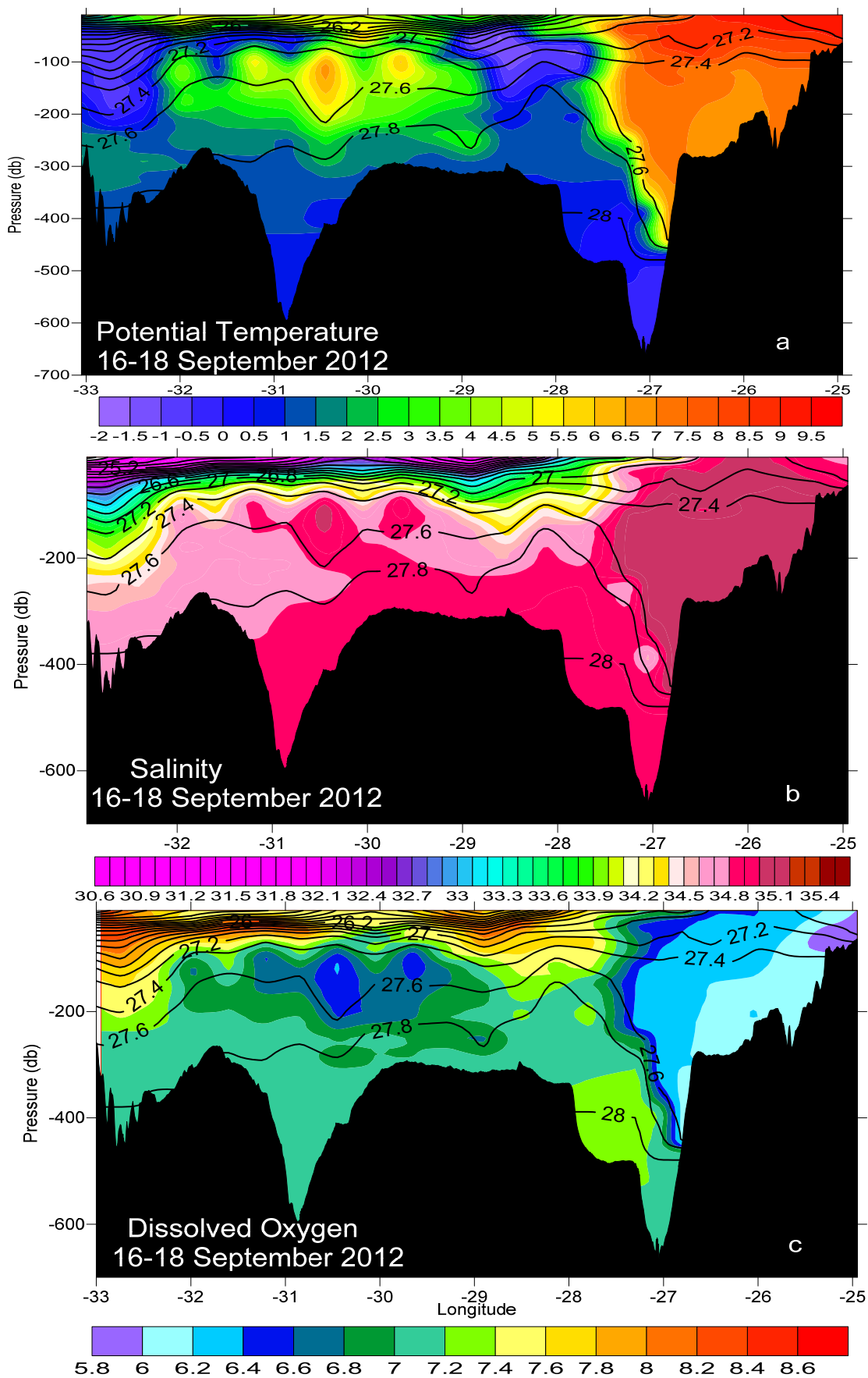


Figure 13 The vertical distribution of (a) potential temperature and (b) salinity and (c) dissolved oxygen between Greenland and Iceland 16-18 September 2012. Density is shown in black.

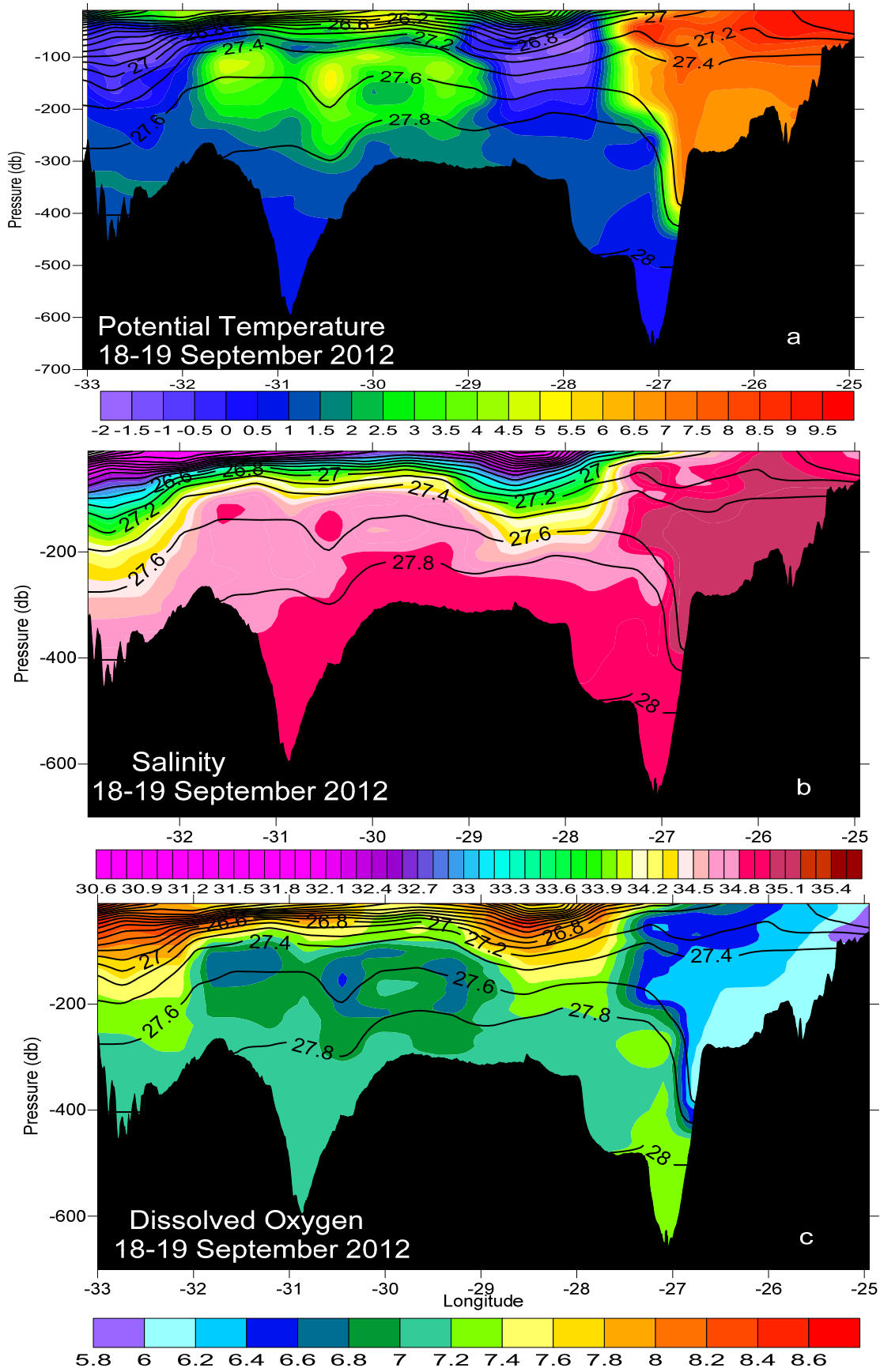


Figure 14 The vertical distribution of (a) potential temperature and (b) salinity and (c) dissolved oxygen between Greenland and Iceland 18-19 September 2012. Density is shown in black.

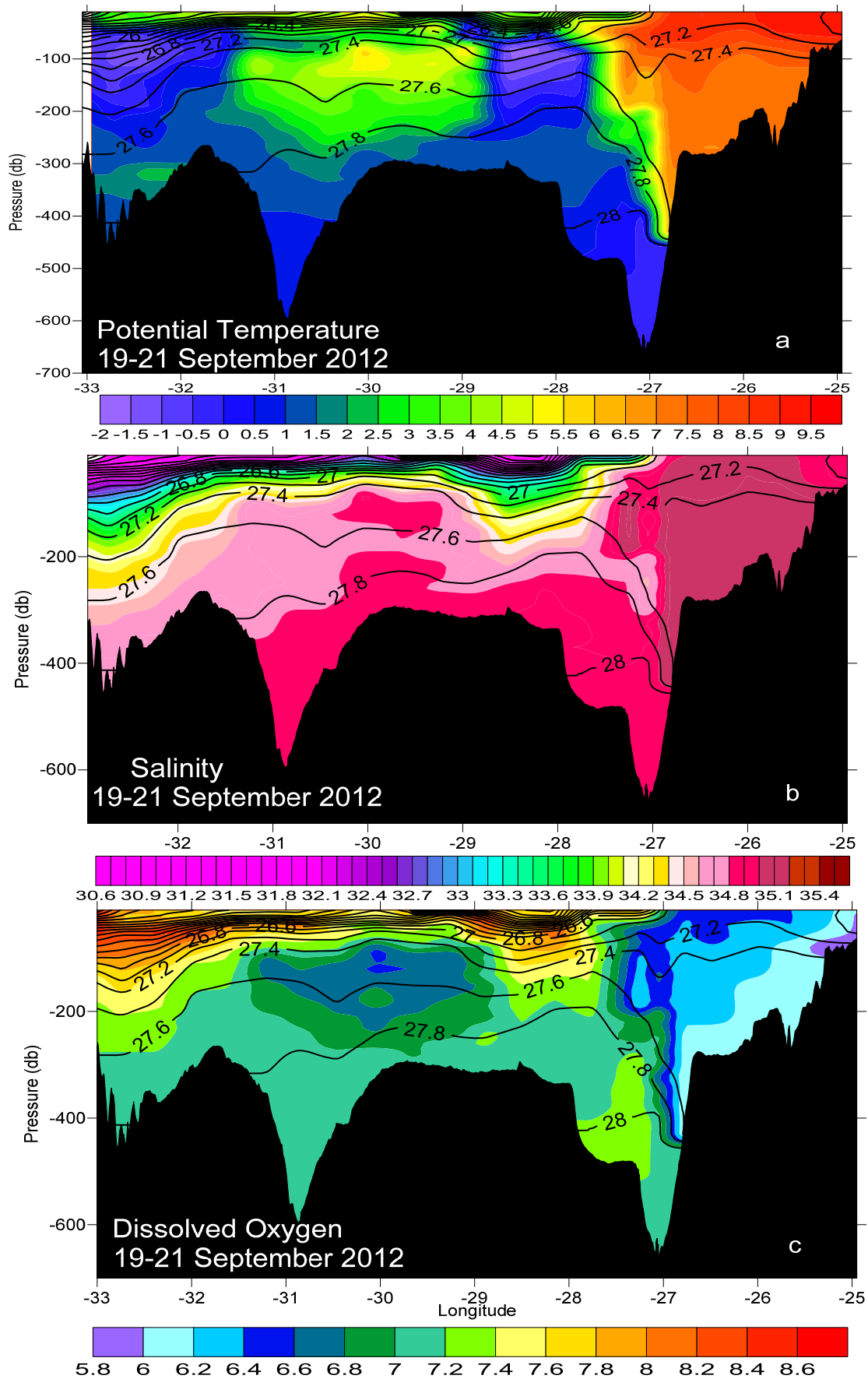


Figure 15 The vertical distribution of (a) potential temperature and (b) salinity and (c) dissolved oxygen between Greenland and Iceland 19-21 September 2012. Density is shown in black.

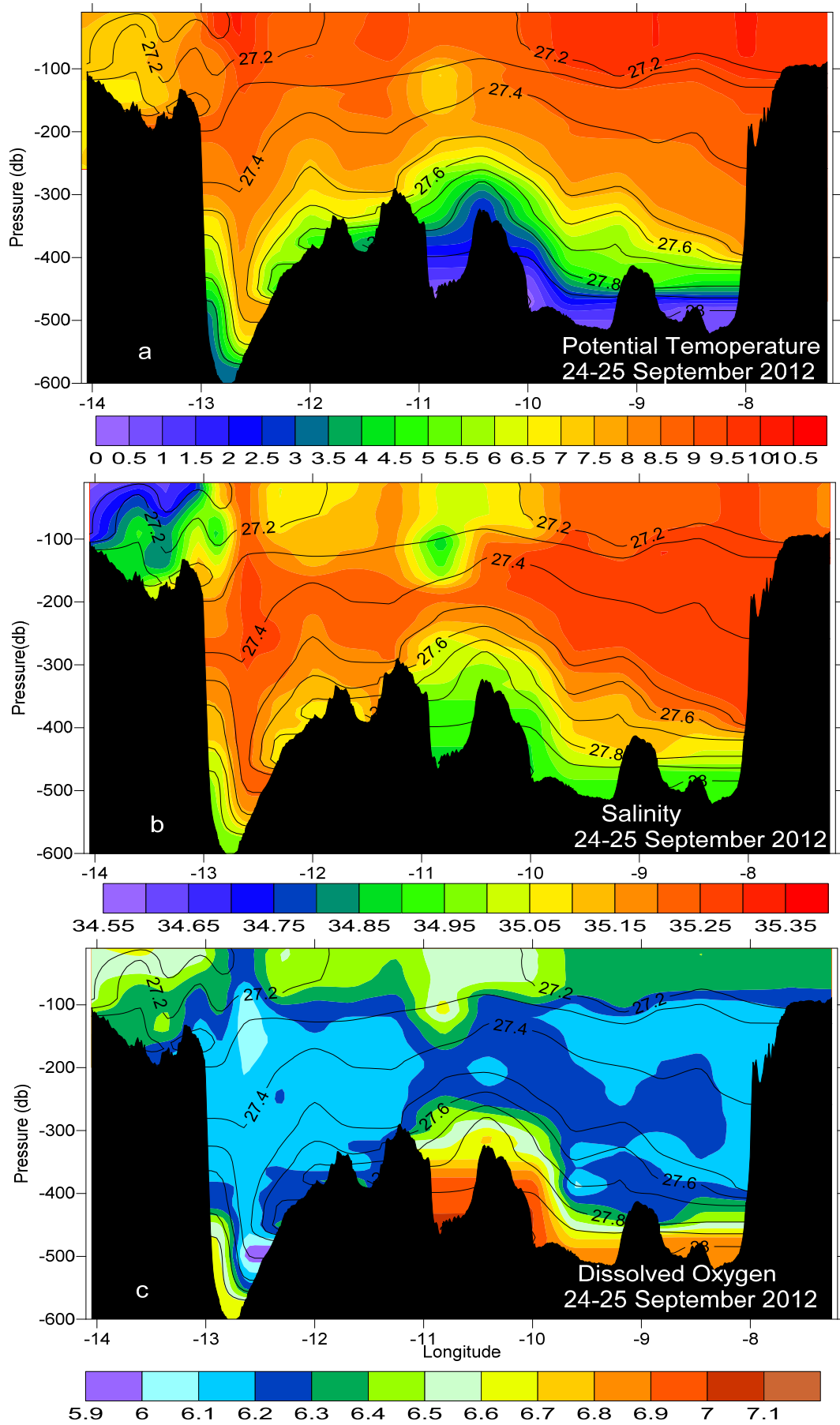


Figure 16 The vertical distribution of (a) potential temperature and (b) salinity and (c) dissolved oxygen between Iceland and Faroe Islands 24-25 September 2012. Density is shown in black.

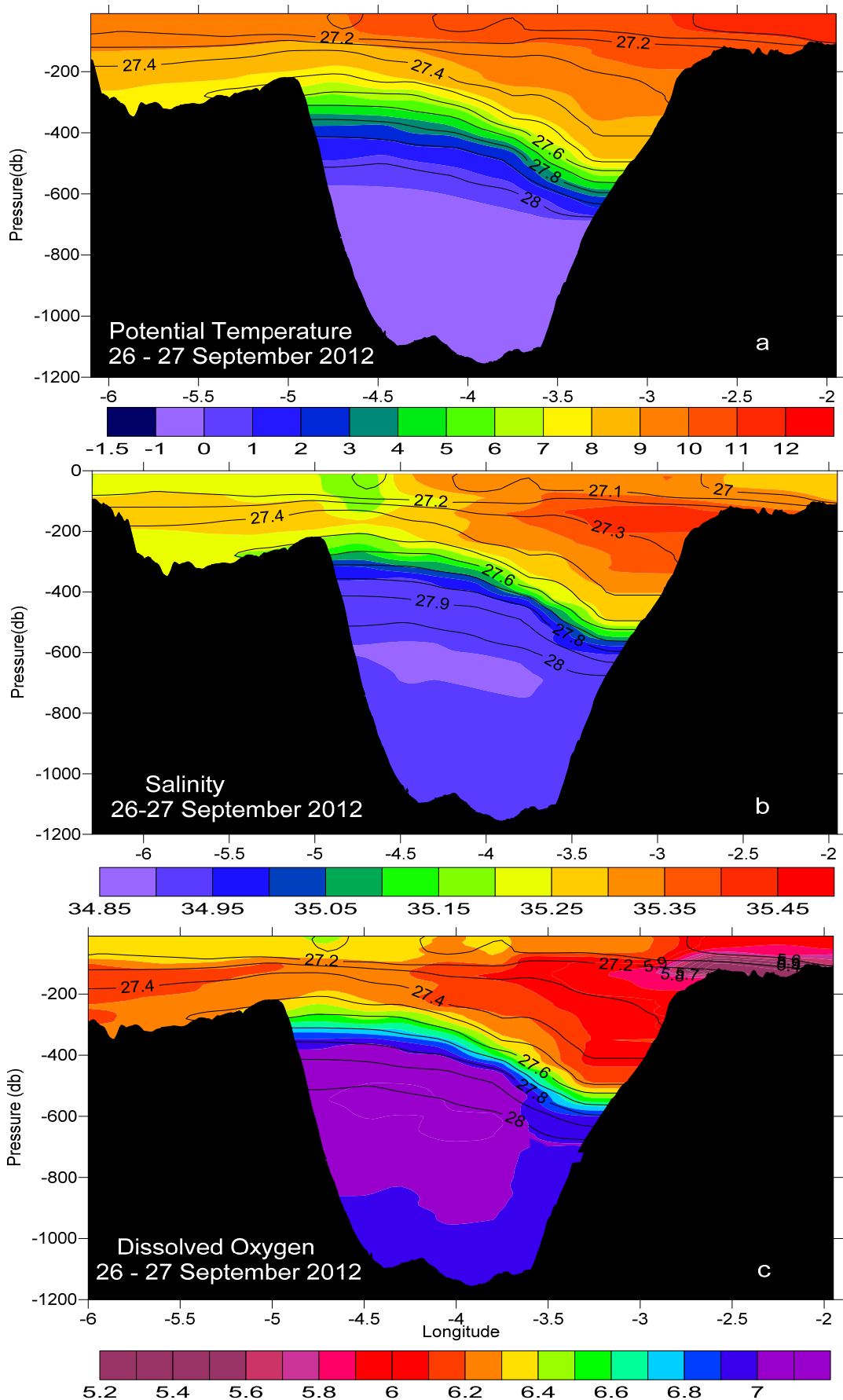


Figure 17 The vertical distribution of (a) potential temperature and (b) salinity and (c) dissolved oxygen between Shetlands and Faroe Islands 26-27 September 2012. Density is shown in black.

