A late Holocene multi-proxy record from the northern Norwegian margin: temperature and salinity variability

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Abstract

This study presents a late Holocene multi-proxy analysis performed on a marine sediment core from the northern Norwegian margin. Palaeo sub-surface temperature and salinity estimates were reconstructed in order to elucidate the natural variability of Atlantic and Coastal water influences. Planktic foraminiferal fauna and their preservation indicators, stable isotopes (δ^{18} O, δ^{13} C) and trace elements were analysed. Paired Mg/Ca and δ^{18} O measurements of N. pachyderma resulted in temperature (SST $_{\mathrm{Mg/Ca}}$) and salinity reconstructions in addition to transfer function derived temperature (SST_{Transfer}) estimates. The overall poor preservation conditions at the core site might have influenced the Mg/Ca ratios and are therefore likely the primary cause of the recorded underestimated SST_{Mg/Ca} and SSS values. The high resolution record shows a general cooling trend with fluctuating palaeoceanographic conditions which are attributed to shifting NAO conditions. Period I (ca. 3500 – 2900 cal yr BP) is characterized by a stronger influence of Coastal water and stratified water masses most likely associated with negative NAO-like conditions. During period II (ca. 2900 – 1600 cal yr BP) warm Atlantic water linked with a positive NAO mode dominates the core site. A returned influence of Coastal water is observed throughout period III (ca. 1600 – 950 cal yr BP). The stable and colder sub-surface temperatures during this period correlate with the Dark Ages. Within period IV (ca. 950 – 550 cal yr BP) the core site experienced a stronger influence of Atlantic water likely due to positive NAO conditions correlating to the Medieval Warm Period.

1 Introduction

Throughout the late Holocene, a general cooling trend has been observed in the North Atlantic associated with a reduced influence of warm Atlantic water (e.g. Slubowska et al., 2005; Hald et al., 2007; Skirbekk et al., 2010). A similar cooling trend, recorded by lake and tree records from north-western Europe, has been ascribed to reduced insolation at high latitudes (e.g. Bjune et al., 2009; Kaufman et al., 2009). Contradictory, fluctuations of a strengthened Atlantic water inflow towards the Arctic Ocean have been observed at the Vøring plateau (e.g. Andersson et al., 2003; Risebrobakken et al., 2003; Andersson et al., 2010), the Barents Sea (e.g. Duplessy et al., 2001; Lubinski et al., 2001; Berben et al., 2014)

and the Svalbard margin (e.g. Slubowska-Woldengen et al., 2007; Jernas et al., 2013; Werner et al., 2013; Zamelczyk et al., 2013). Additionally, throughout the late Holocene, several observations of fluctuating climatic conditions have been found in the Nordic Seas (e.g. Giraudeau et al., 2004; Nyland et al., 2006; Solignac et al., 2006; Slubowska-Woldengen et al., 2007) as well as in north-western Europe (e.g. Lauritzen and Lundberg, 1999; Bjune and Birks, 2008). These include the 'Roman Warm Period' (RWP, ca. BCE 50 – CE 400), the 'Dark Ages' (DA, ca. CE 400 - 800), the 'Medieval Warm Period' (MWP, CE 900 – 1500) and the 'Little Ice Age' (LIA, ca. CE 1500 - 1900) (e.g. Lamb, 1977).

These fluctuating conditions have been ascribed to different causes such as solar forcing, volcanic eruptions (e.g. Bryson and Goodman, 1980; Lean, 2002; Jiang et al., 2005; Wanner et al., 2008) or changes in atmospheric forcing linked to the North Atlantic Oscillation (NAO) controlling the inflow of Atlantic water to the Arctic Ocean (e.g. Trouet et al., 2009; Olsen et al., 2012). Warm and salty Atlantic water is brought into the Nordic Seas by the North Atlantic Current (NAC) and flows parallel with colder and less saline Coastal water along the Norwegian margin. These two water masses possess opposite characteristics with respect to temperature and salinity. Further, they respond opposite to the strengthened or reduced westerlies attributed to positive and negative NAO modes, respectively (e.g. Sætre, 2007). A positive versus negative NAO mode affects the climatic conditions in north-western Europe by generating warmer and wetter versus colder and dryer conditions (e.g. Wanner et al., 2001). Along the Norwegian coast the impact of the variable NAO is seen in precipitation, temperature and wind intensity changes (Ottersen et al., 2001). And thus, the northern Norwegian margin is a key location to investigate the natural variability of Atlantic water inflow throughout the late Holocene linked to fluctuating NAO modes. With respect to the recent global warming, it is of great importance to understand the natural variability of the Atlantic water inflow towards the Arctic Ocean as it strongly influences north-western Europe climatic conditions.

For the Nordic Seas, several studies have reconstructed water mass properties based on planktic foraminiferal fauna. Transfer functions reflect sub-surface temperatures whereas stable oxygen isotopes reflect both the temperature and $\delta^{18}O$ of ambient sea water (e.g. Sarnthein et al., 2003; Rasmussen and Thomsen, 2010; Risebrobakken et al., 2010; Berben et al., 2014). However, in order to reconstruct palaeo sub-surface salinities, additional proxies are required. The Mg-uptake in foraminiferal calcite is primarily temperature depended (e.g.

Nürnberg, 1995). The positive correlation between the Mg-uptake in foraminiferal calcite and temperature of the ambient sea water during growth allows palaeo reconstructions of subsurface temperatures based on Mg/Ca ratio measurements (e.g. Mashiotta et al., 1999; Elderfield and Ganssen, 2000; Kozdon et al., 2009). Therefore, paired calcite $\delta^{18}O$ and Mg/Ca measurements enable the reconstruction of palaeo seawater $\delta^{18}O$ (e.g. Thornalley et al., 2009; Elderfield et al., 2010) and subsequently, via modern $\delta^{18}O_{seawater}$:salinity relationships, enable the reconstruction of palaeo salinity.

In order to investigate the fluctuating interplay of Atlantic and Coastal water related to variable NAO modes throughout the late Holocene, the variability of sub-surface temperature and salinity has been reconstructed. Paired Mg/Ca and δ^{18} O measurements of *Neogloboquadrina pachyderma* were analysed in addition to the distribution of planktic foraminiferal fauna and preservation conditions. Subsequently, a new record of both palaeo sub-surface temperature and salinity estimates from the northern Norwegian margin is presented.

2 Present day oceanography

The Norwegian Sea is dominated by relatively warm and saline Atlantic water (>2 °C, >35 %; Hopkins, 1991). Atlantic water is brought to the area by the two-branched Norwegian Atlantic Current (NwAC) (Orvik and Niiler, 2002) (Figure 1A). Both branches follow a topographically steered northwards pathway through the Nordic Seas and eventually reach the Arctic Ocean via the Fram Strait. The western branch crosses the Iceland-Faroe Ridge entering the Norwegian Sea as the Iceland-Faroe frontal jet (Perkins et al., 1998) (Figure 1A). The eastern branch passes through the Faroe-Shetland channel and follows a pathway along the Norwegian shelf edge towards the Arctic Ocean with a branch flowing into the Barents Sea (Orvik and Niiler, 2002) (Figure 1A).

The Norwegian Coastal Current (NCC) transports Coastal water (2-13 °C, 32-35 ‰; Hopkins, 1991) northwards originating from the North Sea, the Baltic and the Norwegian coast (Figure 1A). Coastal water is characterized by its low salinities due to the influence of freshwater run off from the Norwegian mainland. The NCC is density driven which is mainly influenced by its salinity distribution. Mixing with Atlantic water takes place and increases northwards, and

thus salinity increases whereas stratification reduces. In general, cold Coastal water can be found above warmer Atlantic water in the upper 50 to 100 m of the water column as a thinning wedge westwards (Ikeda et al., 1989) (Figure 1B). A boundary is formed as a well-defined front between the cold, low salinity Coastal water and the warmer, more saline Atlantic water (Ikeda et al., 1989). The overall properties and movements of the NCC are influenced by several factors such as freshwater, tides, wind conditions, bottom topography and Atlantic water (Sætre, 2007). In the study area, the topography causes the NCC to extend much further westwards and hence, closer to the influence of the NwAC (Figure 1C). Therefore, this is a key area to investigate the interplay between Atlantic and Coastal water fluctuations as a result of climatic variability.

3 Material and methods

For this study, a marine sediment core from the northern Norwegian margin (Vøring plateau in front of Trænadjupet south of the Lofoten) was investigated. The core (W00-SC3) (67.24° N, 08.31° E) was retrieved in 2000 by the SV *Geobay* at a water depth of 1184 m (Laberg et al., 2002) (Figure 1). Its recovery was 385 cm from which the top 19 cm was disturbed and therefore not used. The core consists of very soft clay sediments and was sampled for each 1.0 cm between 19 and 263 cm. CTD data nearby the core site (67.10° N, 08.26° E) indicate the presence of Atlantic water till a water depth of ca. 375 m for present day conditions (Blaume, 2002) (Figure 2).

3.1 Chronology

A depth-age model of W00-SC3 based on four AMS 14 C dates measured on *N. pachyderma* was developed. All four AMS 14 C dates were calibrated using Calib 7.0.0 software (Stuiver and Reimer, 1993), the Marine13 calibration curve (Reimer et al., 2013) and a local reservoir age (Δ R value) of 71 \pm 21 following recommendations by Mangerud et al. (2006) (Table 1). The calibration was constrained on a 2- σ range for both calendar years Before Present (cal yr BP) and calendar years Before Common Era (BCE)/Common Era (CE) (cal yr BCE/CE). For this study the cal yr BP depth-age model will be used, however, the cal yr BCE/CE scale was

added for all the plotted data in order to compare with different studies. The AMS ¹⁴C date at 23.5 cm was omitted from the final depth-age model as it appeared too old. The new depth-age model was constrained using linear interpolation between dated levels. Further, based on the homogeneous lithology throughout the core, the sedimentation rate from 120-40 cm was extrapolated towards the top, more specifically between 40 and 19 cm (Figure 3). The resulting depth-age model is constrained between 3485 and 550 cal yr BP (1536 BCE – 1395 CE) and showed sedimentation rates between 0.79 and 0.87 mm/yr and thus, enabled the sampling on a multi-decadal temporal resolution (Figure 3).

3.2 Planktic foraminifera and preservation indicators

All samples were freeze-dried, wet-sieved through three different size fractions (1000, 100 and 63 μm) and subsequently dried in an oven at 40 °C. Every 4 cm from the 100 - 1000 μm size fraction was analysed for its planktic foraminiferal content (>300 specimens) following Knudsen (1998). The identification of left and right coiling *N. pachyderma* was done following Darling et al. (2006), meaning that the right coiling form is identified as *Neogloboquadrina incompta* (Cifelli, 1961). Subsequently, relative abundances (%) of each species were calculated for 62 samples. The planktic foraminiferal concentration (#/g sediment) and fluxes (#/cm²/yr) were calculated. For the latter, a theoretical value for the dry bulk density of 0.76 g/cm³ was assumed based on marine sediment core T-88-2 (Aspeli, 1994) and subsequently calculated according Ehrmann and Thiede (1985).

As carbonate dissolution might affect the planktic foraminiferal assemblages, it is important to investigate the preservation conditions in order to assess the potential dissolution induced pre- and post-depositional alterations (e.g. Zamelczyk et al., 2013). Preservation indicators such as the mean shell weight (μg) of *N. pachyderma* (Broecker and Clark, 2001; Barker and Elderfield, 2002; Beer et al., 2010) and fragmentation (%) of planktic foraminiferal tests (Conan et al., 2002) were inspected. The mean shell weight of *N. pachyderma* was determined for 118 samples using a Mettler Toledo microbalance (0.1 μg sensitivity). In order to minimize problems of variations due to size difference and ontogeny, 50 square shaped and well preserved (visually) specimens were selected from a narrow size range (150 - 250 μm) (Barker et al., 2004). For 62 samples within the 100 - 1000 μm size fraction, the fragmentation was calculated using the equation of Pufhl and Shackleton (2004) (Equation 1).

Fragmentation (%) =
$$\frac{\# fragments/g}{\# fragments/g/2 + \# tests/g} * 100$$
 [Equation 1]

Within this equation, a divisor (2) is included as it is most likely that each test breaks into more than one fragment. The latter is in order to reduce possible misinterpretations of the dissolution sensitivity in changes and progress (Le and Shackleton, 1992; Pufhl and Shackleton, 2004).

3.3 Geochemical analysis

Weight percentages (wt. %) of total carbon (TC) and total organic carbon (TOC) were analysed at one cm resolution using a LECO SC-444 (ES-2) at the Laboratory of the Geological Survey of Norway (NGU). Prior to TOC analysis, the inorganic carbon (carbonate) was removed by treating the samples with 10 % hydrochloric acid (HCl). The calcium carbonate (CaCO₃) content was calculated according Espitalié et al. (1977) (Equation 2).

$$CaCO_3 = (TC - TOC) * 8,33$$
 [Equation 2]

3.4 Stable isotope analysis

Stable isotope analysis (δ^{18} O, δ^{13} C) was performed on 117 samples containing 50 specimens of *N. pachyderma*. In order to minimize size dependent effects on isotopic composition, all specimens were picked in a 150 - 250 µm size fraction (Keigwin and Boyle, 1989; Oppo and Fairbanks, 1989; Donner and Wefer, 1994; Aksu and Vilks, 1998; Bauch et al., 2000). The stable isotope measurements were performed with a Finnigan MAT 253 Mass Spectrometer coupled to an automated Kiel device at the Geological Mass Spectrometer (GMS) Laboratory of the University of Bergen. The analytical errors for δ^{18} O and δ^{13} C were \pm 0.06 ‰ and \pm 0.03 ‰ respectively. The data were reported in ‰ versus VPDB and calibrated with NBS-19. For δ^{18} O, a vital effect of 0.6 ‰ has been applied. This value has previously been suggested for the Nordic Seas (Simstich et al., 2003) as well as for the Norwegian Sea (Nyland et al., 2006).

3.5 Trace element analysis

Trace element analysis was performed on 101 samples containing an average of 75 visually well preserved specimens of the planktic foraminifer *N. pachyderma*. In order to minimize the possible size-dependent bias on the Mg/Ca measurements, the tests were picked from a narrow (as narrow as possible) size fraction (150 - 250 µm) (Elderfield et al., 2002). The tests were cracked open between glass plates in order to optimize the chemical cleaning procedure which consisted of a reduction step to remove metal oxides and an oxidation step to remove any organic matter (e.g. Boyle and Keigwin, 1985). Subsequently, the samples were analysed by ICP-MS method for foraminiferal analysis including simultaneous measurements of Mg, Mn, Al, and Fe, all of which are reported as ratios to Ca. The measurements were performed on a Finnigan Element2 at the Marine Science Institute, UC Santa Barbara with an analytical precision of 1.2 % for Mg/Ca. Fe, Al and Mn are tracers of contaminating phases and thus investigated in order to identify remnant post cleaning sample contamination which might bias the measured Mg/Ca ratios in foraminiferal calcite (Barker et al., 2003).

3.6 Temperature and salinity reconstructions

To reconstruct summer sub-surface temperatures (SST_{Transfer}) (July-August-September) for a water depth of 100 m, a transfer function and the modern training set of Husum and Hald (2012) were used. Although the core site lays at the geographical southern boundary of this modern training set, it was assumed to be the most appropriated as it uses material analysed at the >100 µm size fraction and thus include smaller specimens as *Turborotalita quinqueloba* and *Globigerinita uvula*. The transfer function derived SST_{Transfer} reconstructions were performed using C2 version 1.7.2. (Juggins, 2010) and a three component Weighted Average-Partial Least Square (WA-PLS) statistical model with a leave-one-out cross validation (Ter Braak and Juggins, 1993; Telford and Birks, 2005).

Mg/Ca derived sub-surface temperature estimates ($SST_{Mg/Ca}$) were reconstructed using the refined species-specific temperature:Mg/Ca equation of Kozdon et al. (2009) (Equation 3).

$$Mg/Ca \ (mmol/mol) = 0.13 \ (\pm 0.037) * T \ (^{\circ}C) + 0.35 \ (\pm 0.17)$$
 [Equation 3]

This equation is based on cross-calibrated $\delta^{44/40}$ Ca and Mg/Ca temperature estimates from Holocene core top samples of *N. pachyderma* from the Norwegian Sea and Arctic domain. Although the Mg-uptake into foraminiferal calcite is controlled by exponential thermodynamics (Mashiotta et al., 1999), in Equation 3, it is assumed that within the narrow temperature range occupied by *N. pachyderma*, a linear temperature function is appropriate and works well for reconstructed temperatures above ca. 2.5 °C (Kozdon et al., 2009). However, for temperatures below 2.5 °C, associated with salinities less than 34.5 ‰, the method loses its precision (Kozdon et al., 2009).

In order to reconstruct sub-surface salinities, the oxygen isotope ratios of the ambient seawater ($\delta^{18}O_w$) were first calculated using the temperature: $\delta^{18}O$ equation of Shackleton (1974) modified after O'Neil et al. (1969) (Equation 4).

$$T(^{\circ}C) = 16.90 - 4.38 * (\delta^{18}O_c - \delta^{18}O_w) + 0.10 * (\delta^{18}O_c - \delta^{18}O_w)^2$$
 [Equation 4]

Within this equation, T refers to the previously calculated $SST_{Mg/Ca}$, $\delta^{18}O_c$ to the measured oxygen isotope value of foraminiferal calcite (*N. pachyderma*) and $\delta^{18}O_w$ to the oxygen isotope composition of ambient seawater. Both $\delta^{18}O_c$ and $\delta^{18}O_w$ are expressed as % VPDB. In order to convert $\delta^{18}O_w$ VPDB values to the $\delta^{18}O_w$ VSMOW scale, the following equation was applied (Hut, 1987) (Equation 5).

$$\delta^{18}O_{WVPDB} = \delta^{18}O_{WVSMOW} - 0.27\%_0$$
 [Equation 5]

Subsequently, Equation 4 and 5 are rearranged in order to calculate $\delta^{18}O_w$ VSMOW which results in Equation 6 (Nyland et al., 2006).

$$\delta^{18}O_w = \delta^{18}O_c + 0.27 - \left[\frac{4.38 - \sqrt{4.38^2 - 0.4 * (16.9 - T_{Mg/Ca})}}{0.2}\right]$$
 [Equation 6]

Subsequently, sub-surface salinities (SSS) were reconstructed using the salinity (S) to $\delta^{18}O_w$ relation for the central and eastern Nordic Seas by Simstich et al. (2003) (Equation 7).

$$\delta^{18}O_w = -12.17 + 0.36 * S$$
 [Equation 7]

4 Results

4.1 Planktic foraminifera and preservation indicators

The planktic foraminiferal fauna consist of six species: *N. pachyderma*, *T. quinqueloba*, *N. incompta*, *Globigerinita glutinata*, *Globigerina bulloides* and *Globigerinita uvula* (Table 2) (Figure 4). Overall, the record is dominated by *N. incompta* and *T. quinqueloba* with a mean value of ca. 34 and 29 % respectively.

Between ca. 3500 and 2900 cal yr BP, the planktic foraminiferal concentration and flux show relative low values of ca. 750 #/g sediment and ca. 50 #/cm²/yr (Figure 4A; 5E). *N. pachyderma* and *N. incompta* show a small decrease from ca. 18 to 14 % and from ca. 40 to 22 % (Figure 4B; 4D). Simultaneously, *T. quinqueloba* and *G. uvula* show a gradual increase from ca. 28 to 35 % and from ca. 5 to 20 % (Figure 4C; 4G). *G. glutinata* and *G. bulloides* show relative abundances of ca. 6 and 3 % (Figure 4E; 4F).

The planktic foraminiferal concentration and flux show slightly higher values of ca. 1000 #/g sediment and ca. 65 #/cm²/yr between ca. 2900 and 2300 cal yr BP which are followed by a sharp increase reaching the highest recorded values (3765 #/g sediment, 226 #/cm²/yr) towards ca. 1600 cal yr BP (Figure 4A; 5E). Between ca. 2900 and 1600 cal yr BP, stable abundances of *N. pachyderma* (ca. 11 %) and relative high and stable values of *N. incompta* (ca. 33 %) are recorded (Figure 4B; 4D). *T. quinqueloba* shows high values (ca. 32 %) with a gentle decrease towards ca. 1600 cal yr BP (Figure 4C). *G. glutinata* and *G. bulloides* show, in particular between ca. 2900 and 2300 cal yr BP, the highest recorded values (ca. 7 and 5 %) followed by a small reduction between 2300 and 1600 cal yr BP (Figure 4E; 4F).

At ca. 1600 cal yr BP, the concentration and flux records show a rapid drop from 3765 to 865 # g/sediment and from 226 to 52 #/cm²/yr followed by stable values (ca. 1330 #/g sediment, ca. 80 #/cm²/yr) between ca. 1600 and 950 cal yr BP (Figure 4A; 5E). Gently increasing values of *G. uvula* (ca. 15 to 26 %) and *G. glutinata* (ca. 2 to 5 %) are recorded; whereas *T. quinqueloba* shows a slight decrease from ca. 25 to 16 % (Figure 4G; 4E; 4C). *N. pachyderma*, *N. incompta* and *G. bulloides* remain relatively constant throughout this period (Figure 4B; 4D; 4F).

Between ca. 950 and 550 cal yr BP, the concentration and flux record show slightly decreasing values towards 450 #/g sediment and 27 #/cm²/yr (Figure 4A; 5E). *N. pachyderma* and *N. incompta* both show an increase with values up to ca. 23 and 42 % (Figure 4B; 4D). Simultaneously, decreasing values are recorded for *G. glutinata* from ca. 5 to 2 % and for *G. uvula* from ca. 26 to 8 % (Figure 4E; 4G). *T. quinqueloba* and *G. bulloides* remain constant around values of ca. 21 and 2 % respectively (Figure 4C; 4F).

The preservation indicators show overall values between 47 and 94 % of planktic foraminiferal fragmentation and between 1.6 and 4.8 µg for the mean shell weight of *N. pachyderma* (Figure 5A-B). Between ca. 3500 and 2900 cal yr BP, the fragmentation shows generally high values around ca. 75 %, whereas the mean shell weight decreases slightly from 3.7 to 2.6 µg. Between ca. 2900 and 1600 cal yr BP, the fragmentation remains relatively stable around slightly reduced values of ca. 69 %. The mean shell weight of *N. pachyderma* shows decreasing values towards 1.7 µg at ca. 2300 cal yr BP followed by an increase reaching ca. 3 µg at ca. 1600 cal yr BP. Between ca. 1600 and 950 cal yr BP, the fragmentation increases from 49 to 79 % and the mean shell weight remains stable around ca. 2.7 µg. Between ca. 950 and 550 cal yr BP, both preservation indicators show a pronounced increase from ca. 47 to 90 % for the planktic foraminiferal fragmentation and from 2.0 to 3.2 µg for the mean shell weight of *N. pachyderma*.

4.2 Geochemical analysis

The TOC and CaCO₃ results show an overall increasing trend with values between 0.7 and 1.3 wt. % and 17.0 and 30.6 wt. %, respectively (Figure 5C-D). Between ca. 3500 and 2900 cal yr BP, both records show an increase of ca. 0.7 to 0.9 wt. % for TOC and ca. 17 to 22 wt. % for CaCO₃. Towards ca. 2300 cal yr BP, TOC values increase reaching 1.1 wt. % followed by a quick drop to 0.8 wt. % around ca. 2200 cal yr BP and remain further stable around this value towards ca. 1600 cal yr BP. Between ca. 2900 and 1600 cal yr BP, the CaCO₃ continues its increasing trend from ca. 22 to 30 wt. %. Between ca. 1600 and 950 cal yr BP, the TOC values increase gradually from 1.0 to 1.1 wt. %, whereas the CaCO₃ data decreases from ca. 30 to 24 wt. %. Between ca. 950 and 550 cal yr BP, TOC values remain relatively stable around ca. 1.0 wt. %, whereas the CaCO₃ record increases slightly towards ca. 26 wt. %.

4.3 Stable isotope analysis

The stable isotope analysis results in values between 2.3 and 3.1 % for $\delta^{18}O$ and within the range -0.4 to 0.6 % for $\delta^{13}C$ (Figure 6). Between ca. 3500 and 2900 cal yr BP, $\delta^{18}O$ values decrease gently from ca. 2.9 to 2.7 % whereas the $\delta^{13}C$ record shows simultaneously increasing values from ca. 0.2 to 0.4 %. Between ca. 2900 and 1600 cal yr BP, the $\delta^{18}O$ record continues its decreasing trend towards values of 2.5 % at ca. 1600 cal yr BP. The $\delta^{13}C$ values decrease from ca. 0.4 % at ca. 2900 cal yr BP towards ca. 0.1 % at ca. 2300 cal yr BP followed by relative stable values around ca. 0.3 % towards ca. 1600 cal yr BP. Between ca. 1600 and 950 cal yr BP, $\delta^{18}O$ values initially continue their decreasing trend towards 2.4 % at ca. 1300 cal yr BP before increasing to values of ca. 2.7 % between ca. 1200 and 950 cal yr BP. The $\delta^{13}C$ record shows increasing values from ca. 0.1 to 0.4 % during this time interval. Between ca. 950 and 550 cal yr BP both records increase simultaneously towards ca. 2.8 % for $\delta^{18}O$ and ca. 0.5 % for $\delta^{13}C$.

4.4 Trace element analysis

Tracers of contamination are plotted against Mg/Ca ratios (Figure 7). Fe and Al are tracers of detrital material contamination such as silicate minerals, whereas Mn is a tracer of secondary diagenetic Mn-rich carbonate (Boyle, 1983). The results show high correlations between Mg/Ca and Fe and Al tracers, with $R^2 = 0.57$ for Fe/Ca and $R^2 = 0.47$ for Al/Ca (Figure 7A-B). The correlation between Mg/Ca and Mn/Ca is low ($R^2 = 0.06$) and thus further not relevant (Figure 7C). When deleting samples with ratios higher than 100 µmol/mol of the contamination tracers following the contamination threshold used by Barker et al. (2003), the correlations are strongly reduced to non-existing with $R^2 = 0.05$ for Fe/Ca and $R^2 = 0.09$ for Al/Ca (Figure 7D-E). This change suggests that contamination by detrital material (i.e. high values of Fe/Ca and/or Al/Ca) impact the results by elevating the Mg/Ca values. Therefore, all samples (n=32) with ratios >100 µmol/mol of Fe and Al are omitted from the further study.

The resulting Mg/Ca record consists of 69 data points with ratios in the range 0.70 to 0.96 mmol/mol (Figure 8A). Between ca. 3500 and 2900 cal yr BP, the Mg/Ca record shows relative stable values around ca. 0.76 mmol/mol. Between ca. 2900 and 2500 cal yr BP, the ratios increase towards 0.89 mmol/mol. Due to the omitted data as a result of possible

contamination, there is no reliable data between ca. 2500 and 2100 cal yr BP. Between ca. 2100 and 1600 cal yr BP, the record remains relatively stable around ca. 0.81 mmol/mol. Around ca. 1600 cal yr BP the Mg/Ca ratios drop quickly towards 0.70 mmol/mol, thereafter they remain stable around ca. 0.78 mmol/mol towards the top of the record (i.e. due to the omitted values after ca. 800 cal yr BP).

4.5 Temperature and salinity reconstructions

The transfer function derived temperature (SST_{Transfer}) reconstructions (n=62) range between 6.3 and 7.7 °C (Figure 8C; 4H). Between ca. 3500 and 2900 cal yr BP, the SST_{Transfer} record decreases from 7.2 to 6.7 °C. Thereafter, the SST_{Transfer} values increase towards ca. 7.7 °C at ca. 2500 cal yr BP. Hereafter the SST_{Transfer} record decreases towards ca. 1600 cal yr BP reaching ca. 6.6 °C. Between ca. 1600 and 950 cal yr BP, the SST_{Transfer} estimates remain relatively stable around ca. 6.6 °C. Eventually, between ca. 950 and 550 cal yr BP, the SST_{Transfer} record shows a decrease of ca. 0.5 °C (from ca. 6.8 to 6.3 °C).

The Mg/Ca derived sub-surface temperature reconstructions (SST_{Mg/Ca}) (n=69) range between 2.7 and 4.7 °C (Figure 8B). Between ca. 3500 and 3300 cal yr BP, the SST_{Mg/Ca} values show a small decrease from 3.1 to 2.7 °C followed by relatively stable values of ca. 3.2 °C towards ca. 2900 cal yr BP. Between ca. 2900 and 2500 cal yr BP, the SST_{Mg/Ca} values show a gradual increase from ca. 3.2 to 4.2 °C and between ca. 2100 and 1600 cal yr BP, SST_{Mg/Ca} values are relatively stable around 3.5 °C. Between ca. 1600 and 950 cal yr BP, the SST_{Mg/Ca} estimates are slightly lower around ca. 3.2 °C with one exception around ca. 1300 cal yr BP where SST_{Mg/Ca} estimates peak to 4.7 °C.

Reconstructed sub-surface salinities (SSS) (n=65) range between 31.6 and 34.0 ‰ (Figure 8D). Between ca. 3500 and 2900 cal yr BP, the reconstructed SSS estimates show a gradual decrease from ca. 33 to 32 ‰. After ca. 2900 cal yr BP, the SSS data increases to 33.6 ‰ at ca. 2800 cal yr BP followed by a decrease to ca. 2500 cal yr BP. Between ca. 2100 and 1600 cal yr BP, palaeo SSS estimates are stable around ca. 33 ‰, showing the highest values of the record (i.e. 34.0 ‰). Between ca. 1600 and 950 cal yr BP, the SSS values show stable values of ca. 32 ‰.

5 Discussions

5.1 Palaeo SST and SSS assessment

Modern sea temperature measured nearby the core site is ca. 7 °C within the upper 300 m of the water column (Blaume, 2002) (Figure 2), and thus corresponds to the reconstructed palaeo SST_{Transfer} estimates (Figure 4H; 8C). However, the SST_{Mg/Ca} estimates, which range 3.0 to 3.6 °C lower than the SST_{Transfer}, do not correspond with the modern sea temperature data (Figure 8B). Further, the SSS estimates (ca. 33 %) show (consequently) lower values than the modern salinity values (>35 %) (Blaume, 2002) (Figure 2). Furthermore, the SST_{Mg/Ca} estimates also seem to be rather low compared to palaeo reconstructions of the last 1200 years from the Vøring plateau (Nyland et al., 2006). This dissimilarity is in the first place attributed to the use of a different temperature:Mg/Ca equation by Nyland et al. (2006), more specifically the equation from Elderfield and Ganssen (2000). The temperature:Mg/Ca equations from Elderfield and Ganssen (2000) and Kozdon et al. (2009) are calibrated differently resulting in two different potential temperature ranges of the equations, 0 to 7 °C and 3 to 6 °C respectively. In this study, applying the temperature:Mg/Ca equation of Elderfield and Ganssen (2000) would have resulted in 0.7 to 1.8 $^{\circ}\text{C}$ warmer $SST_{Mg/Ca}$ and 0.71 to 0.88 %higher SSS estimates. Nonetheless, a different equation would still have resulted in lower $SST_{Mg/Ca}$ estimates in comparison with the $SST_{Transfer}$ and/or present day instrumental data. This discrepancy might be caused by variability of the species-specific habitat depth preference and/or main calcification period (i.e. season) of N. pachyderma, in addition to initial low Mg/Ca values (Meland et al., 2006; Kozdon et al., 2009).

Whereas the SST_{Transfer} record is reconstructed for a water depth of 100 m, the SST_{Mg/Ca} record reflects the average calcification depth of *N. pachyderma*. Studies have shown a poor correlation between modern Mg/Ca inferred temperatures and their corresponding hydrographic data and thus, contest the previously made assumption that the SST_{Mg/Ca} reflects a constant average calcification depth (Meland et al., 2006; Nyland et al., 2006). The water depth of chamber formation and encrustation of *N. pachyderma* is linked to the stratification of the water column and can therefore be highly variable (e.g. Kohfeld et al., 1996; Stangeew, 2001; Simstich et al., 2003). Furthermore, Kozdon et al. (2009) argued that the main factor controlling the habitat depth of *N. pachyderma* in the Nordic Seas is the density-driven water mass stratification. Therefore, it seems that the by Mg/Ca recorded temperature signal is

bound to the isopycnal layer and hence, only reflects a narrow temperature range. Hemleben et al. (1989) indicated that *N. pachyderma* is probably a passive dweller implying that the water mass stratification is preconditioning the habitat and thus, causing a lower temperature response of the Mg/Ca signal. This effect might even be amplified in high latitudes due to the stronger water mass stratification. It is thus most likely that *N. pachyderma* does not possess a constant calcification depth and that the species therefore records a dampened amplitude of the total sub-surface water temperature range. Due to the distinctive seasonality and strong stratification in high latitude regions, the assignment of a true calcification depth and hence temperature remains one of the major issues (Barker et al., 2003). The apparent calcification depth of *N. pachyderma* can be found around ca. 250 m water depth off Norway (Simstich et al., 2003). Palaeo water mass properties at ca. 250 m might have differed from those at ca. 100 m and thereby possibly partly contributed to the observed discrepancy between SST_{Mg/Ca} and SST_{Transfer} estimates.

On an annual time scale, the biological production of *N. pachyderma* peaks in spring and a second time during maximum stratification in late summer (Jonkers et al., 2010). Therefore, the Mg-uptake by *N. pachyderma* reflects a combined spring and summer temperature signal resulting in lower $SST_{Mg/Ca}$ estimates compared to the $SST_{Transfer}$ values which are reconstructed for summer. However, Jonkers et al. (2013) measured Mg/Ca values of *N. pachyderma* from sediment traps in the central Irminger Sea which only show a weak seasonal trend. Jonkers et al. (2013) speculate that the Mg-uptake is controlled by other factors than temperature such as sensitivity of *N. pachyderma* to $[CO_3^{2-}]$ changes which could partly mask the summertime increases due to a parallel increase in $[CO_3^{2-}]$.

In addition to temperature, several secondary factors might influence the absolute Mg/Ca ratios including variability in salinity (Nürnberg et al., 1996; Lea et al., 1999), calcite dissolution (Dekens et al., 2002), test size (e.g. Elderfield et al., 2002) and the applied cleaning protocol (e.g. Barker et al., 2003), which may provide an explanation to the initially low Mg/Ca values. The presented Mg/Ca ratios are relatively low and are ca. 0.16 mmol/mol lower than previous observations from the Vøring plateau performed within a comparable size fraction using a similar cleaning method (Nyland et al., 2006). In particular, the "Cd cleaning method" (Boyle and Keigwin, 1985) comprising a reductive cleaning step could result in absolute lowering of Mg/Ca ratios of up to ~15 % (Barker et al., 2003; Meland et al., 2006). Artificially correcting the present Mg/Ca data for this potential reduction would translate into

0.8 to 1.1 °C higher $SST_{Mg/Ca}$ and 0.6 to 0.8 % higher SSS estimates, which are still lower than modern day values and the $SST_{Transfer}$ record. Hence, this cleaning step does not seem to explain the differences observed between the two temperature proxies. This was also concluded by Aagaard-Sørensen et al. (2013) after comparisons between transfer function and Mg/Ca derived temperature estimates from a core retrieved in the Polar North Atlantic.

The observed preservation indicators show generally high planktic foraminiferal fragmentation and low mean shell weights of N. pachyderma and thereby, argue for overall poor preservation conditions (Figure 5A-B). In addition, the generally low CaCO₃ record (<30 wt. %) corresponds to earlier findings of low carbonate contents of surface sediments on the Norwegian continental margin associated with poor preservation conditions (Huber et al., 2000) (Figure 5D). At the continental margins, dilution by terrigenous material might lead to low percentages of sedimentary carbonate (Hebbeln et al., 1998), whereas high fertility results in a high input of organic matter and thereby, to enhanced carbonate dissolution (Berger, 1971; Emerson and Bender, 1981; Archer, 1991). Previous studies highlighted that Mg-rich parts might have been removed from the foraminiferal test due to dissolution and thereby, bias the Mg/Ca ratios resulting in underestimated SST_{Mg/Ca} values (Brown and Elderfield, 1996; Rosenthal et al., 2000; Johnstone et al., 2011). Although the risk of measuring biased material was minimized by picking the visually most preserved tests, it remains impossible to quantify the degree of preservation influence on the measured Mg/Ca ratios and hence, the reconstructed SST_{Mg/Ca} and SSS estimates. Due to the observed overall poor preservation conditions, it is likely that the Mg/Ca measurements were influenced by carbonate dissolution.

5.2 Palaeoceanographic evolution of the late Holocene

Throughout the late Holocene, the current proxy records show an overall cooling, with SST_{Transfer} values decreasing from 7.7 to 6.3 °C (Figure 4H; 8C), associated with an overall decreased inflow of Atlantic water. This corresponds well to previous observations of decreased Atlantic water inflow in the Nordic Seas (e.g. Slubowska et al., 2005; Hald et al., 2007; Skirbekk et al., 2010) as well as to lake and tree records from north-western Europe arguing for a late Holocene trend towards colder and dryer conditions (e.g. Bjune et al., 2009; Kaufman et al., 2009). Contrary, at the Vøring plateau south of the study site, planktic

foraminiferal data show an overall increased inflow of Atlantic water throughout this time period (Andersson et al., 2003; Risebrobakken et al., 2003; Andersson et al., 2010). In addition to the overall cooling, the current study also shows smaller scale fluctuations of the sub-surface water masses which will be discussed further for four separate time periods. The interpretation of the fluctuating influence of sub-surface water masses and their possible link with NAO conditions is presented as schematic profiles across the northern Norwegian margin (Figure 9).

5.2.1 Period I: ca. 3500 – 2900 cal yr BP

Between ca. 3500 and 2900 cal yr BP, the relative faunal abundances show increased values of G. uvula (ca. 5 to 20 %) and high values (ca. 30 %) of T. quinqueloba (Figure 4G; 4C). The latter has been associated with sub-polar conditions and Atlantic water (Bé and Tolderlund, 1971; Volkmann, 2000) and additionally considered to respond rapidly to changes in nutrient supply (Reynolds and Thunnel, 1985; Johannessen et al., 1994). G. uvula has been associated with Coastal water and reduced salinities (Husum and Hald, 2012) as well as with high food supplies and cold productive surface waters (e.g. Saito et al., 1981; Boltovskoy et al., 1996; Bergami et al., 2009). Hence, the high relative abundances of both T. quinqueloba and G. uvula argue for enhanced productivity most likely due to an increased influence of colder, less saline and more productive Coastal water. Additionally, the enriching δ^{13} C values might also argue for increased primary production corresponding to an increased influence of more productive Coastal water (Figure 6B).

However, the relatively low planktic foraminiferal concentrations and fluxes seem to argue for a rather low primary production (Figure 4A; 5E). These relatively low values might result from relatively poor preservation conditions as indicated by generally high planktic foraminiferal fragmentation, a decreasing mean shell weight of *N. pachyderma* and low CaCO₃ values (<22 wt. %) (Figure 5A; 5B; 5D). The latter might reflect the dilution by terrigenous material enhancing carbonate dissolution at the continental margin off Norway (Huber et al., 2000). In addition, the solubility of CaCO₃ increases with decreasing temperatures (Edmond and Gieskes, 1970), and thus, the enhanced dissolution conditions are most likely due to a combination of the core location at the continental margin and an increased influence of colder Coastal water.

The depleting $\delta^{18}O$ trend might indicate an increased temperature and/or decreased salinity signal (Figure 6A). The planktic foraminiferal fauna data and decreasing SST_{Transfer} estimates (7.2 to 6.7 °C) with an average value of ca. 6.9 °C argue strongly that the $\delta^{18}O$ record reflects less saline water masses associated with an increased influence of Coastal water (Figure 4H; 8C). Correspondingly, the reconstructed SSS estimates confirm the overall trend towards less saline conditions (Figure 8D). Nonetheless, contrary to decreasing SST_{Transfer} estimations, the overall lower SST_{Mg/Ca} values remain relatively stable throughout this period which might illustrate the different water depths and/or season that the two proxies represent (see section 5.1.3) possibly arguing for more stratified water masses (Figure 8B-C).

Overall, the multi-proxy data argue for an increased influence of Coastal water and possibly more stratified water masses at the core site. This is likely the result of a more westwards located thinning wedge of Coastal water above Atlantic water that might have been the result of a dominating negative NAO-like mode throughout this time interval (Figure 9A). Previously, a 5200 year NAO-index has been reconstructed using a multi-proxy geochemical record from a lake in south-west Greenland (Olsen et al., 2012). Although this NAO-index shows mainly positive values, Olsen et al. (2012) observed a stronger influence of negative NAO conditions between ca. 4500 and 2500 cal yr BP which might correspond to the negative NAO-like conditions suggested here (Figure 8E). Negative NAO conditions are generally associated with a reduced inflow of Atlantic water and a stronger influence of Coastal water (e.g. Sætre, 2007; Hurrell et al., 2013). Correspondingly, between ca. 3500 and 2500 cal yr BP, reconstructed SST estimates from the Vøring plateau (66.58° N, 07.38° E) show decreasing values arguing for a reduced influence of Atlantic water (Andersson et al., 2003; Risebrobakken et al., 2003; Andersson et al., 2010). Additionally, negative NAO conditions result in a colder and dryer climate in north-western Europe (e.g. Wanner et al., 2001). A decreasing temperature and precipitation trend throughout this time interval was observed by pollen and plant macrofossil analyses from a lake record in northern Norway (Svanåvatnet; 66.25° N, 14.03° E) (Bjune and Birks, 2008). Furthermore, based on the mean ablation-season temperature and winter snow accumulation, a decreased winter precipitation was also observed in western Norway (Nesje et al., 2001) and between 3500 and 3200 cal yr BP, a speleothem record from northern Norway indicates decreased surface ground temperatures (Lauritzen and Lundberg, 1999).

Between ca. 2900 and 1600 cal yr BP, the planktic foraminiferal fauna is characterized by high relative abundances of *T. quinqueloba*, *N. incompta*, *G. glutinata* and *G. bulloides* (Figure 4C-F). These species have all been associated with subpolar conditions and warm Atlantic surface water masses (e.g. Bé and Tolderlund, 1971; Johannessen et al., 1994; Carstens et al., 1997; Simstich et al., 2003) and thus, argue for a pronounced influence of Atlantic water brought to the study area by the NwAC.

The total planktic foraminiferal concentration and flux values show higher and increasing values, especially between ca. 2300 and 1600 cal yr BP, possibly indicating for increased primary production (Figure 4A; 5E). However, the δ^{13} C record shows an initial depletion towards ca. 2300 cal yr BP followed by relative stable values (ca. 0.3 ‰) arguing for less primary production (Figure 6B). The seemingly increased primary production, reflected by higher flux and concentration, likely results from the generally improved preservation conditions. The observed overall reduced fragmentation and increasing trend of CaCO₃ (up to ca. 30 wt. %) argue for a gradual trend towards reduced dissolution conditions throughout this period (Figure 5A; 5D). More favourable preservation conditions have previously been associated with increased influence of Atlantic surface water where pore waters are supersaturated with respect to calcium due to the lower organic matter productivity and a higher rain of CaCO₃ (e.g. Huber et al., 2000; Henrich et al., 2002).

The $\delta^{18}O$ record continues its depleting trend from Period I possibly reflecting an increased temperature and/or reduced salinity signal (Figure 6A). The planktic foraminiferal fauna data and the therefrom derived high $SST_{Transfer}$ estimates (ca. 7.3 °C) argue for increasing temperatures which are most likely related to an increased inflow of Atlantic water (Figure 4H; 8C). Correspondingly, the palaeo SSS record shows overall higher values up to ca. 34 % and thus, correlates to an increased influence of more saline Atlantic water (Figure 8D). Additionally, the $SST_{Mg/Ca}$ estimates show, compared to Period I, ca. 0.5 - 1.0 °C higher temperatures associated with an increased influence of Atlantic water (Figure 8B).

The increased influence of Atlantic water inflow suggested here might result from positive NAO conditions. The latter are associated with stronger westerlies across the North Atlantic (e.g. Hurrell et al., 2013). This resulted in an enhanced inflow of Atlantic water which possibly reduced the influence of Coastal water leading to a more eastwards located thinning

wedge of Coastal water above Atlantic water (Figure 9B). Throughout this time period, Olsen et al. (2012) observed a general increasing trend in their NAO-index which might correspond to the influence of a positive NAO mode suggested here (**Figure 8E**). Similarly high subsurface temperatures associated with a strengthened inflow of Atlantic water have been observed at the Vøring plateau between ca. 2500 and 1600 cal yr BP (Andersson et al., 2003; Risebrobakken et al., 2003; Andersson et al., 2010). Throughout the time interval of period II, increased temperatures in northern Norway have been observed from a lake record (Bjune and Birks, 2008) and a speleothem record (Lauritzen and Lundberg, 1999). In addition, between ca. 2700 and 1900 cal yr BP increased winter precipitation has been observed in western Norway (Nesje et al., 2001). This correlates well to the warmer and wetter climate scenarios attributed to positive NAO conditions in north-western Europe (e.g. Wanner et al., 2001). Further, the last part of this period might be corresponding to the earlier observed Roman Warm Period between ca. 2000 and 1550 cal yr BP (ca. BCE 50 – CE 400) (Lamb, 1977) and additionally correlate to the findings of warmer temperatures in northern Norway also linked to the Roman Warm Period by Lauritzen and Lundberg (1999).

5.2.3 Period III: ca. 1600 – 950 cal yr BP

Between ca. 1600 and 950 cal yr BP, the planktic foraminiferal fauna shows stable values for all species with particularly high values of *G. uvula* (ca. 25 %) (Figure 4G). This likely reflects a stable period characterized by a strongly increased influence of Coastal water (e.g. Husum and Hald, 2012). At ca. 1600 cal yr BP, both the planktic foraminiferal concentration and flux drop rapidly where after the record remains stable until ca. 950 cal yr BP (Figure 4A; 5E). The planktic foraminiferal fragmentation increases, whereas the CaCO₃ record shows an overall decreasing trend (Figure 5A; 5D). This might indicate slightly reduced preservation conditions, possibly related to an enhanced influence of Coastal water and dilution of terrigenous material (e.g. Huber et al., 2000). Correspondingly, the enriched δ^{13} C record indicates enhanced primary production conditions arguing for a returned influence of Coastal water (Figure 6B).

Although $\delta^{18}O$ values initially continue their decreasing trend towards ca. 1300 cal yr BP, they eventually increase and thereby argue for a possible reduction in temperature (Figure 6A). Compared to Period II, the SST_{Transfer} (ca. 6.6 °C) and SST_{Mg/Ca} records both reflect Paper III - 19

stable conditions with ca. 0.5 - 1.0 °C lower estimates indicative of a reduced influence of relatively warm Atlantic water (Figure 8B-C). Additionally, the SSS estimates show a rapid decrease followed by stable, and compared to period II, ca. 0.5 - 1.0 % lower values and thus, also arguing for a reduced influence of Atlantic water or enhanced influence of Coastal water (Figure 8D).

The multi-proxy data argues for a strong influence of Coastal water which is interpreted as a westwards migrated thinning wedge of Coastal water (Figure 9C) which would suggest a shift towards negative NAO conditions (e.g. Sætre, 2007). However, the negative NAO conditions suggested here do not correlate with the overall positive NAO mode reconstructed by Olsen et al. (2012) during this time interval (Figure 8E). The current multi-proxy record corresponds well with other marine and terrestrial observations from nearby the study area within this time interval. Palaeo SST estimates from the Vøring plateau showed a sharp decrease at ca. 1600 cal yr BP where after they remained stable around lower temperatures (compared to values before ca. 1600 cal yr BP) (Andersson et al., 2003; Risebrobakken et al., 2003; Andersson et al., 2010). This also suggests a reduced influence of Atlantic water towards the core site. Further, a colder and dryer climate associated with the Dark Ages was suggested by low surface ground temperatures between ca. 1500 and 900 cal yr BP in northern Norway (e.g. Lauritzen and Lundberg, 1999). Additionally, Bjune and Birks (2008) also observed decreasing air temperatures between ca. 1800 and 800 cal yr BP in northern Norway.

5.2.4 Period IV: ca. 950 – 550 cal yr BP

The increased relative abundance of *N. incompta* indicates a stronger influence of subpolar conditions and thus, a stronger influence of Atlantic water (e.g. Bé and Tolderlund, 1971) (Figure 4D). The concomitant decreasing values of *G. glutinata* and *G. uvula* correspondingly argue for a possible decreased influence of Coastal water (Figure 4E; 4G). The increased abundance of *N. pachyderma* argues for increased polar conditions (e.g. Bé and Tolderlund, 1971; Volkmann, 2000) (Figure 4B). A similar increased abundance of *N. pachyderma* was also found throughout the last ca. 1000 cal yr BP at the Vøring plateau (Risebrobakken et al., 2003). The planktic foraminiferal concentrations and fluxes slightly decrease which might indicate a reduction in primary production (Figure 4A; 5E). Fragmentation and mean shell weight records show a pronounced increase arguing for deteriorated preservation conditions Paper III - 20

in the sediment, i.e. increased fragmentation as well as to reduced dissolution of calcite, i.e. increased shell weight (Figure 5A-B). In addition, the $CaCO_3$ record shows a slight increase indicating somewhat reduced dissolution conditions associated with an increased influence of Atlantic water (Huber et al., 2000) (Figure 5D). The increasing $\delta^{13}C$ record might indicate a reduced stratification at the core site resulting from an increased influence of Atlantic water (Figure 6B).

Based on the planktic foraminiferal fauna and the therefrom derived decreasing $SST_{Transfer}$ record, the enriching $\delta^{18}O$ values might reflect a reduced temperature signal (Figure 6A). The $SST_{Transfer}$ record reaches its lowest estimate (ca. 6.3 °C) by the end of the record around ca. 550 cal yr BP (Figure 4H; 8C). The latter contradicts with the previously suggested returned influence of Atlantic water, however, the generally colder temperature estimates are most likely due to the overall cooling trend observed throughout the record and linked to decreasing solar insolation values throughout the late Holocene (e.g. Hald et al., 2007; Kaufman et al., 2009). Nonetheless, contemporaneous reconstructed SST estimates from the Vøring plateau have shown increasing values (Andersson et al., 2003; Risebrobakken et al., 2003; Andersson et al., 2010).

Despite the overall cooling trend, the increased influence of Atlantic water seems to be the result of generally positive NAO conditions (Figure 9D). The reconstructed NAO-index from south-west Greenland also indicate positive NAO conditions during this time interval which are associated with the Medieval Warm Period (Olsen et al., 2012) (Figure 8E). Furthermore, this also corresponds to the results of a NAO reconstruction based on tree rings and speleothems that indicate a dominating positive NAO mode associated with an intensified AMOC (Trouet et al., 2009). These conditions are expressed by a reduced influence of Coastal water and a stronger moisture and heat transport to Norway by the NwAC resulting in warmer and wetter climate conditions (e.g. Wanner et al., 2001; Hurrell et al., 2013). This correlates well to terrestrial records from northern Norway which show slightly increasing air temperatures for this time interval (Bjune and Birks, 2008) and high surface ground temperature estimates between ca. 800 and 500 cal yr BP (Lauritzen and Lundberg, 1999). This time interval most likely reflects the warmer conditions associated with the Medieval Warm Period correlating with observation at the Vøring plateau between ca. 1150 and 650 cal yr BP (ca. 800 – 1300 AD) (Nyland et al., 2006) and in northern Norway between ca. 800 and 500 cal yr BP (ca. 1150 – 1450 AD) (Lauritzen and Lundberg, 1999).

6 Conclusions

A marine core from the northern Norwegian margin was investigated to elucidate the natural variability of water mass properties throughout the late Holocene. In this study, paired Mg/Ca and δ^{18} O measurements of *N. pachyderma* were presented in order to reconstruct palaeo subsurface temperature and salinity estimates. Additionally, preservation conditions were analysed and the reconstructed palaeo SST_{Mg/Ca} and SSS estimates were assessed. The palaeoceanographic evolution of the late Holocene was discussed in terms of changing influences of Atlantic and Coastal water and linked to fluctuating modes of the NAO.

The recorded discrepancy between the reconstructed $SST_{Mg/Ca}$ and $SST_{Transfer}$ estimations is likely caused by a combination of several factors such as the restriction of Mg/Ca recordings to a narrow temperature signal, inaccurate estimations of the preferred depth habitat and calcification season of *N. pachyderma* and the initially low Mg/Ca measurements. The overall low measured Mg/Ca ratios might have been influenced by the cleaning method including a reductive step. However, they are likely a direct result of the overall poor preservation conditions associated with the continental margin, resulting in potential underestimations of true $SST_{Mg/Ca}$ and SSS values.

Overall, the proxy results indicate a general cooling trend (7.7 to 6.3 °C, SST_{Transfer}) throughout the late Holocene. In addition, fluctuating conditions of Atlantic and Coastal water intensity are observed. Period I (ca. 3500 – 2900 cal yr BP) is influenced by relatively cold (ca. 6.9 °C, SST_{Transfer}), less saline and more productive Coastal waters and a stronger vertical stratification of the water column. These conditions are attributed to dominating negative NAO-like conditions. Throughout Period II (ca. 2900 – 1600 cal yr BP), the core site experiences a stronger influence of warm Atlantic water (ca. 7.3 °C, SST_{Transfer}) with more favourable preservation conditions associated with positive NAO conditions. The last part of this period likely corresponds to the Roman Warm Period. Within period III (ca. 1600 – 950 cal yr BP), stable conditions and cold sub-surface temperatures (ca. 6.6 °C, SST_{Transfer}) are observed indicating a returned and stronger influence of Coastal water. This period correspond very well with the colder and dryer Dark Ages. Period IV (ca. 950 – 550 cal yr BP) experienced a returned stronger influence of Atlantic water attributed to dominating positive NAO conditions and correlates well with the Medieval Warm Period. Nonetheless,

Atlantic water reaching the core site was less warm (ca. 6.3 $^{\circ}$ C, SST_{Transfer}) compared to period II due to the overall late Holocene cooling.

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Figure captions

Table 1: AMS 14 C dates and calibrated radiocarbon ages of W00-SC3. The calibration is performed using Calib 7.0.0 software (Stuiver and Reimer, 1993), the Marine13 calibration curve (Reimer et al., 2013) and a local reservoir age (ΔR value) of 71 \pm 21 following Mangerud et al. (2006). The AMS 14 C date highlighted in grey is omitted from the final depth-age model.

Table 2: Planktic foraminiferal species list.

Figure 1: Study area of W00-SC3 (67.24° N, 08.31° E) (green) with the main surface currents: Norwegian Atlantic Current (NwAC) (red) and Norwegian Coastal Current (NCC) (black). A) Surface currents presented on a bathymetric map. B) Generalized schematic profile across the northern Norwegian margin modified after Rørvik et al. (2010). Coastal water (CW), Atlantic water (AW), Atlantic intermediate water (AIW) and Deep water (DW). C) Detail map of the surface currents nearby the core site modified after Sætre (1983). Red and black arrows indicate Atlantic and Coastal water respectively.

Figure 2: CTD data from PO181_311 (67.10° N, 08.26° E) (Blaume, 2002). Temperature (black) and salinity (grey) versus water depth. Atlantic water mass is highlighted in grey.

Figure 3: Depth-age model of W00-SC3 based on three calibrated AMS 14 C dates. The 2- σ range of the calibrated radiocarbon ages is indicated by an error bar. The exact values are noted in black for the used dates and grey for one omitted AMS 14 C date. A) Calibrated calendar years BP versus core depth. B) Sedimentation rates versus core depth.

Figure 4: Planktic foraminiferal fauna versus cal yr BP (left y-axis) and cal yr BCE/CE (right y-axis). The black diamonds on the Y-axis indicate the AMS ¹⁴C converted to calibrated radiocarbon ages. The horizontal lines (grey) in the background indicate the boundaries between the different periods. A) Planktic foraminiferal concentration (black) and flux (grey) versus age. B-G) Species-specific relative abundances (black) and fluxes (grey) of planktic foraminifera versus age. H) Transfer function derived SST_{Transfer} estimates using a modern foraminiferal dataset by Husum and Hald (2012) versus age.

Figure 5: Preservation and geochemical analysis versus cal yr BP (left y-axis) and cal yr BCE/CE (right y-axis). The black diamonds on the Y-axis indicate the AMS ¹⁴C converted to calibrated radiocarbon ages. The horizontal lines (grey) in the background indicate the boundaries between the different periods. A) Planktic foraminiferal fragmentation versus age. B) Mean shell weight of *N. pachyderma* versus age. C) Total organic carbon versus age. D) Calcium carbonate versus age. E) Planktic foraminiferal concentration (black) and flux (grey) versus age.

Figure 6: Stable isotopes analysis performed on *N. pachyderma* versus cal yr BP (left y-axis) and cal yr BCE/CE (right y-axis). The black diamonds on the Y-axis indicate the AMS ^{14}C converted to calibrated radiocarbon ages. The horizontal lines (grey) in the background indicate the boundaries between the different periods. A) $\delta^{18}O$ measurements corrected for a vital effect of 0.6 % according Simstich et al. (2003) and Nyland et al. (2006) versus age. B) $\delta^{13}C$ measurements versus age.

Figure 7: Trace element analysis plotted as correlations of contamination tracers versus Mg/Ca ratios. A-C) Correlations are plotted for all measurements. D-F) Samples with ratios higher than 100 μ mol/mol are omitted following the contamination threshold of Barker et al. (2003). A; D) Fe/Ca versus Mg/Ca. B; E) Al/Ca versus Mg/Ca. C;F) Mn/Ca versus Mg/Ca.

Figure 8: Water mass properties versus cal yr BP (left y-axis) and cal yr BCE/CE (right y-axis). The black diamonds on the Y-axis indicate the AMS 14 C converted to calibrated radiocarbon ages. The horizontal lines (grey) in the background indicate the boundaries between the different periods. A) Mg/Ca ratios of *N. pachyderma* versus age. B) SST_{Mg/Ca} estimates obtained by the temperature:Mg/Ca equation of Kozdon et al. (2009) versus age. C) SST_{Transfer} estimates using the modern foraminiferal dataset by Husum and Hald (2012) versus age. D) SSS estimates derived by a combined $\delta^{18}O_c$ and SST_{Mg/Ca} approach using the salinity to $\delta^{18}O_w$ relation for the central and eastern Nordic Seas by Simstich et al. (2003) versus age. E) Reconstructed NAO-index from a lake record in south-west Greenland (Olsen et al., 2012) versus age.

Figure 9: The interpretation of fluctuating influence of sub-surface water masses based on multi-proxy data from the study area is presented as a schematic profile across the northern Norwegian margin for four separate time periods. Coastal water (CW), Atlantic water (AW), Atlantic intermediate water (AIW) and Deep water (DW).

Table 1

Lab ID	Core depth	Material	Uncorrected	cal yr	2-σ range	cal yr	2-σ range	Reference
	(cm)		¹⁴ C age	BP	BP	BCE/CE	BCE/CE	
Beta-334422	23.5	N. pachyderma	1590 ± 30	1069	968 - 1170	881 CE	780 - 982 CE	This study
TUa-2931	40.0	N. pachyderma	1360 ± 65	821.5	684 - 959	1128.5 CE	991 - 1266 CE	Laberg et al., 2002
TUa-2930	120.0	N. pachyderma	2305 ± 55	1836	1688 - 1984	114.5 CE	35 BC - 262 CE	Laberg et al., 2002
TUa-2929	263.0	N. pachyderma	3660 ± 95	3484.5	3237 - 3732	1535.5 BCE	1783 - 1288 BCE	Laberg et al., 2002

Table 2

Planktic foraminiferal species

Globigerina bulloides (d'Orbigny), 1826

Globigerinita glutinata (Egger), 1893

Globigerinita uvula (Ehrenberg), 1861

 $Neogloboquadrina\ incompta\ (Cifelli),\ 1961$

Neogloboquadrina pachyderma (sinistral) (Ehrenberg), 1861

Turborotalita quinqueloba (Natland), 1838

Figure 1

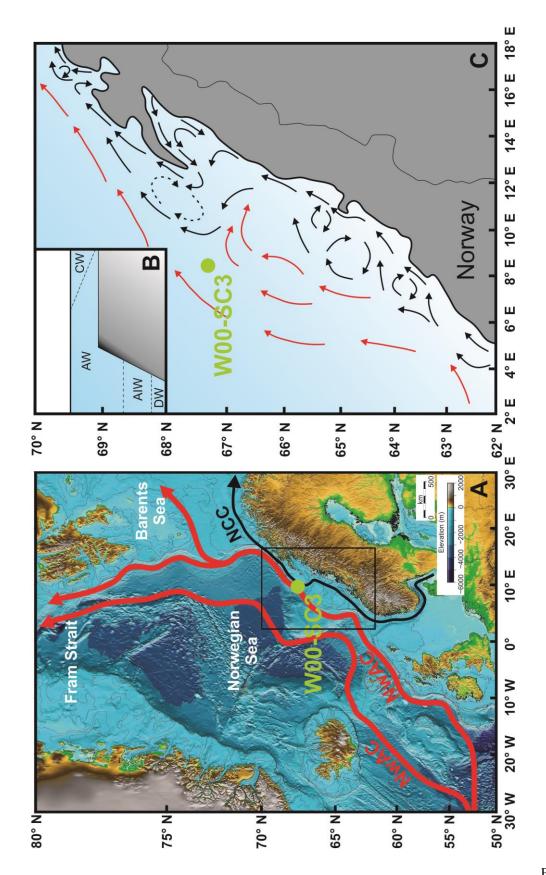


Figure 2

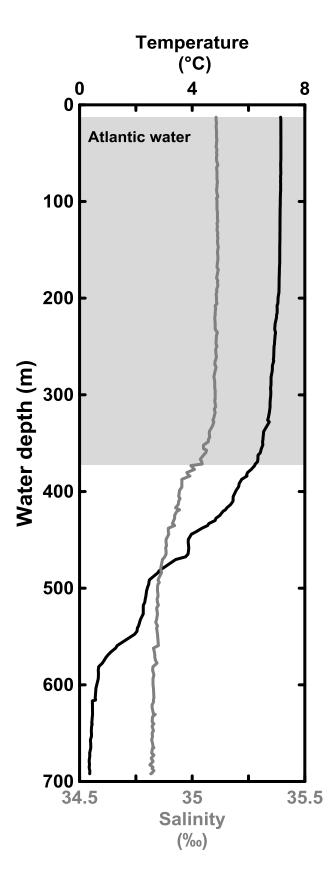
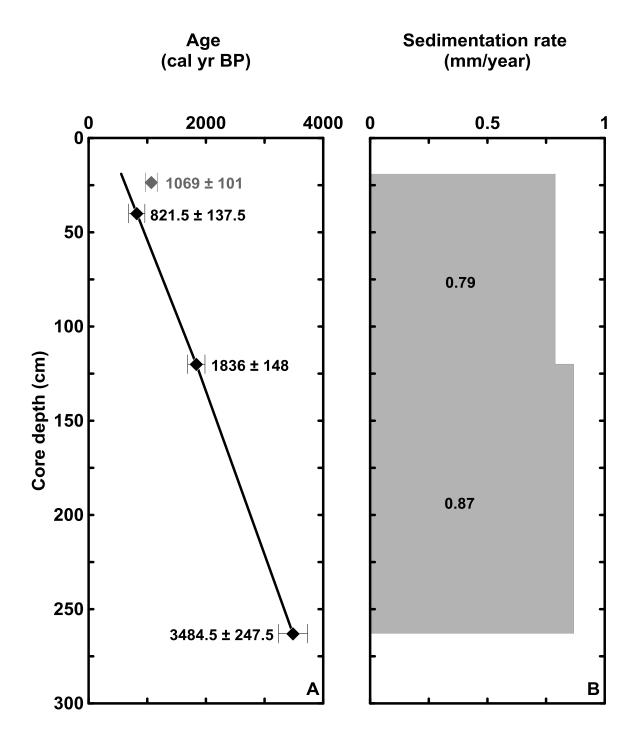


Figure 3



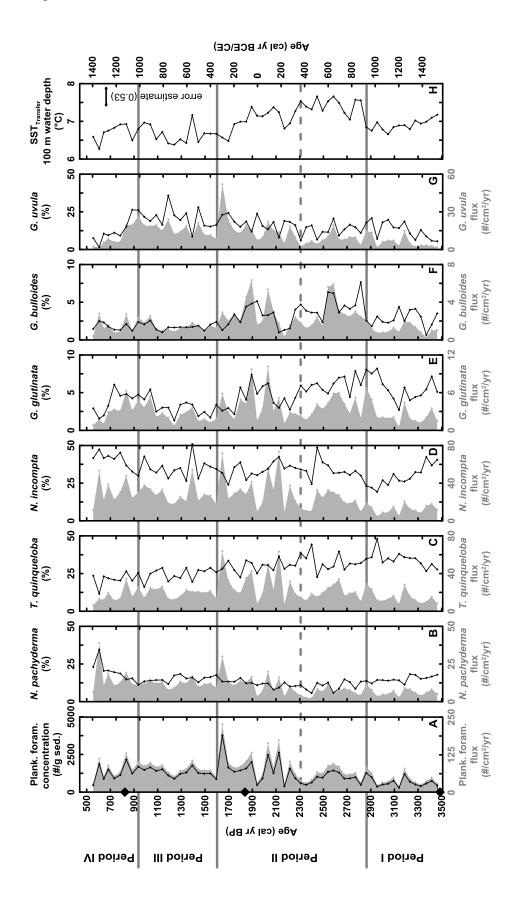


Figure 5

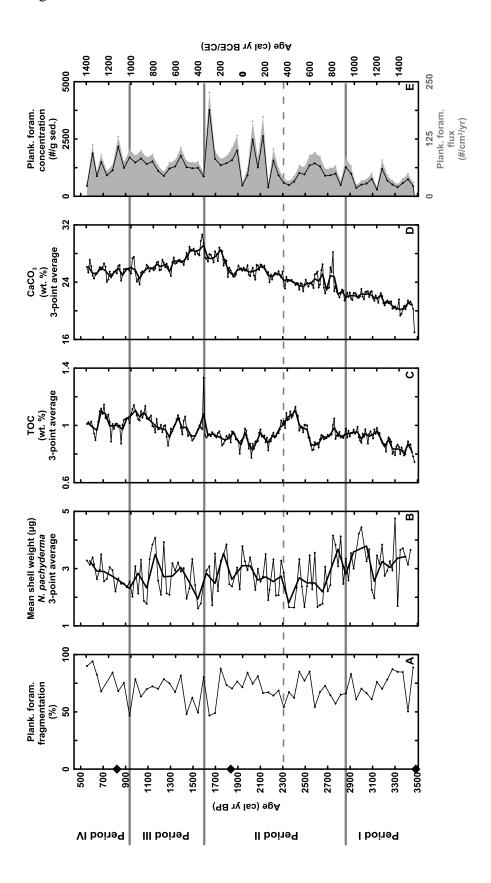


Figure 6

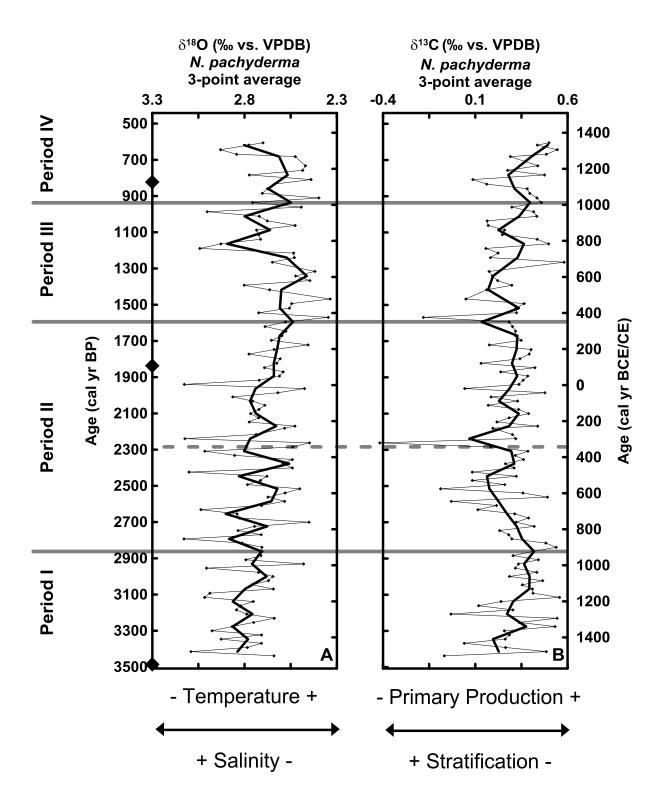
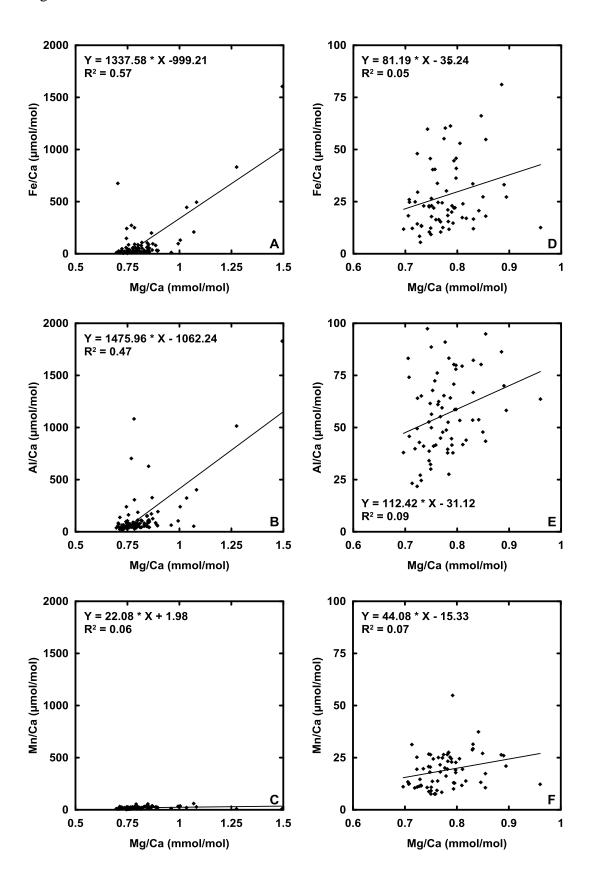


Figure 7



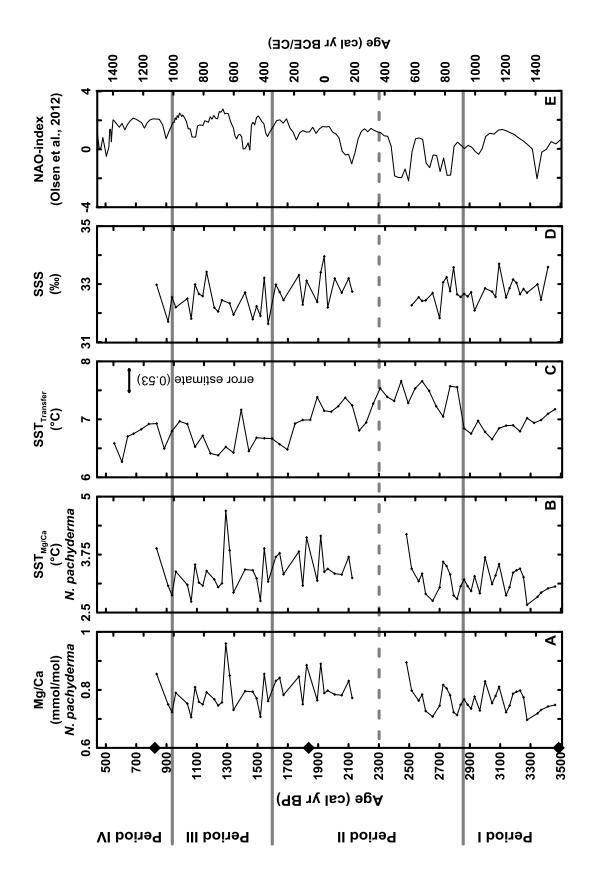


Figure 9

