Millennial-scale variability of marine productivity and terrigenous matter supply in the western Bering Sea over the past 180 kyr

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Abstract. We used piston cores recovered in the western Bering Sea to reconstruct millennial-scale changes in marine productivity and terrigenous matter supply over the past \(\sim 180 \text{kry}\). Based on a geochemical multi-proxy approach, our results indicate closely interacting processes controlling marine productivity and terrigenous matter supply comparable to the situation in the Okhotsk Sea. Overall, terrigenous inputs were high, whereas export production was low. Minor increases in marine productivity occurred during intervals of Marine Isotope Stage 5 and interstadials, but pronounced maxima were recorded during interglacials and Termination I. The terrigenous material is suggested to be derived from continental sources on the eastern Bering Sea shelf and to be subsequently transported via sea ice, which is likely to drive changes in surface productivity, terrigenous inputs, and upper-ocean stratification. From our results we propose glacial, deglacial, and interglacial scenarios for environmental change in the Bering Sea. These changes seem to be primarily controlled by insolation and sea-level forcing which affect the strength of atmospheric pressure systems and sea-ice growth. The opening history of the Bering Strait is considered to have had an additional impact. High-resolution core logging data (color b*, XRF scans) strongly correspond to the Dansgaard–Oeschger climate variability registered in the NGRIP ice core and support an atmospheric coupling mechanism of Northern Hemisphere climates.

1 Introduction

The subarctic North Pacific (N Pacific) is a high-nitrate, low-chlorophyll (HNLC) region (e.g. Kienast et al., 2004; Tyrrell et al., 2005), characterized by salinity driven stratification (permanent halocline), which is suggested to control glacial/interglacial variations in atmospheric carbon dioxide (CO\(_2\)) concentrations (Haug et al., 1999, 2005; Sigman and Boyle, 2000; Sigman et al., 2004, 2010; Jaccard et al., 2005). The halocline prevents the formation of deep water (Warren, 1983; Emile-Geay et al., 2003) and modulates the supply of nutrient-rich deep water into the euphotic zone, thereby influencing the extent of marine productivity and nutrient utilization. Since the halocline also acts as a barrier for atmospheric-oceanic gas exchange, the modern N Pacific, with its high carbon export efficiency (Honda et al., 2002), is considered a net sink of atmospheric CO\(_2\) (Takahashi et al., 2002a).

Several studies have reported low export production in the N Pacific during glacial times (Narita et al., 2002; Kienast et al., 2004; Jaccard et al., 2005, 2009, 2010; Brunelle et al., 2007; Shigemitsu et al., 2007; Galbraith et al., 2008; Gebhardt et al., 2008). However, it remains unclear whether this was caused by stronger stratification or by enhanced sea-ice cover. Both processes would result in a less efficient export of organic carbon to the deep ocean and its subsequent degradation to CO\(_2\) (“biological pump”), and hamper
the release of deep-sequestered CO$_2$ to the atmosphere. Inter-
glacial maxima in export production at ODP Site 882 were
related to reduced stratification rather than to sea-ice influ-
ence (Jaccard et al., 2005). Since the modern Bering Sea
is marked by high marine productivity and seasonal sea-
ic formation (e.g. Springer et al., 1996; Niebauer et al.,
1999), it might have had a different influence on past ocean–
climate–atmosphere CO$_2$ exchange. Paleoceanographic recon-
structions in the Bering Sea also revealed reduced export pro-
duction during the last glacial period, which increased dur-
ing Termination I and remained high in the Holocene (Gor-
barenko, 1996; Cook et al., 2005; Gorbarenko et al., 2005;
Okada et al., 2005; Okazaki et al., 2005; Brunelle et al.,
2007, 2010; Itaki et al., 2009; Khim et al., 2010; Kim et al.,
2011). This variability was explained by a complex interplay
of changes in sea surface temperatures (SST), sea-ice cover-
age, inflow of Pacific surface waters, and upper-ocean strat-
tification (e.g. Katsuki and Takahashi, 2005; Brunelle et al.,
2007, 2010; Kim et al., 2011).

Knowledge of past sea-ice variability in the Bering Sea
comes from diatom and radiolarian assemblages, and IP$_{25}$
biomarker studies (Cook et al., 2005; Katsuki and Takahashi,
2005; Tanaka and Takahashi, 2005; Max et al., 2012). Al-
though sea ice is considered an important transport agent
for terrigenous material, geochemical or sedimentological
studies assessing past terrigenous matter supply in the N
Pacific are rare. Existing studies focused on the Okhotsk
Sea (Sato et al., 2002; Nürnberg and Tiedemann, 2004;
Nürnberg et al., 2011), the NW Pacific (Shigemitsu et
al., 2007; VanLaningham et al., 2009), and the Southern
Ocean (Latimer and Filippelli, 2001). For the Okhotsk Sea,
Nürnberg and Tiedemann (2004) suggested that nearly syn-
chronous glacial/interglacial changes in biological and ter-
rigenous fluxes were modulated by sea-ice processes driven
by variations in the strength of the Siberian High. For the
Bering Sea some provenance studies involving sedimento-
logical and geochemical characteristics of surface sediments
are available, but were mainly conducted on the eastern
Bering Sea shelf (Gardner et al., 1980; Lisitzin, 2002; Asa-
harma et al., 2012; Nagashima et al., 2012). These studies and
results from the Meiji Drift in the NW Pacific indicate that
a large fraction of the supplied terrigenous material was de-
livered from Yukon–Bering Sea sources (VanLaningham et
al., 2009). However, downcore records reflecting the compo-
sitional variability of terrigenous matter in the Bering Sea are
missing.

Recent progress has been made in detecting millennial-
scale climate variability in Bering Sea sediments (Cook et al.,
2005; Gorbarenko et al., 2005, 2010; Okazaki et al., 2005;
Brunelle et al., 2010; Khim et al., 2010; Kim et al., 2011;
Max et al., 2012; Rella et al., 2012). Most of these studies are
restricted to the last ~ 70 kyr or focus on deglacial changes
in the northern, southern, and southeastern Bering Sea. To-
gether with studies from the NE Pacific (e.g. Hendy and Ken-
nett, 2000) they imply that short episodes of increased ma-
rine productivity are connected with interstadials recorded in
Greenland ice cores. However, the existing reconstructions
within the N Pacific realm are constrained by the shallow
lysocline, which limits the use of carbonate-based proxies
and causes stratigraphic uncertainties. In the Bering Sea ac-
cording reconstructions are thus restricted to shallow shelf
areas or topographic highs. Here, we present millennial-scale
reconstructions of export production and terrigenous matter
supply for the hitherto only poorly studied western Bering
Sea. Results were derived from a suite of geochemical prox-
ies and high-resolution core logging data covering the last
~ 180 kyr.

2 Regional setting

The Bering Sea links the Pacific Ocean with the Arctic Ocean
via the only shallow (~ 50 m) Bering Strait. In its eastern and
northern part it is characterized by a wide and shallow (0–
200 m) continental shelf area. Pacific surface waters, trans-
ported by the westward-flowing Alaskan Stream, enter the
Bering Sea through several passes within the Aleutian Island
Arc (e.g. Takahashi, 2005). Inside the Bering Sea, a large-
scale cyclonic surface circulation pattern develops with the
Bering Slope Current (BSC) and the East Kamchatka Current
acting as eastern and western boundary currents, respectively
(Fig. 1). Outflow occurs through the Bering Strait into the
Arctic Ocean and through the Aleutian passes, mainly Kam-
chatka Strait, into the N Pacific (Stabeno et al., 1999). Deep
waters flow northward and eastward from Kamchatka Strait
with return outflow above 3000 m water depth (Reed et al.,
1993; Stabeno et al., 1999). The oceanographic and climatic
conditions are characterized by a strong seasonal variability
of SST and sea-ice coverage that result from the interaction
of the Siberian High and the Aleutian Low. The Arctic Osc-
cillation, Pacific Decadal Oscillation, and the Pacific–North
American pattern are reported to be related with decadal vari-
ations of both atmospheric pressure cells (Niebauer, 1988,
1998; Mantua et al., 1997; Overland et al., 1999, 2002). Dur-
ing winter, a strong Siberian High results in advection of
cold Arctic air masses and mainly northerly wind directions
(Stabeno et al., 1999). This causes a significant cooling of
the sea surface, sea-ice formation, as well as enhanced vertical
mixing in the mixed layer, thereby returning nutrients from
the subsurface. In contrast, during summer, the reduced
strength of the Aleutian Low and enhanced insolation result
in a stratified mixed layer and increased marine productivity.

Primary productivity is dominated by diatoms mainly
blooming during spring, whereas increased biological
CaCO$_3$ fluxes (coccolithophores, planktonic foraminifera)
occur during spring and late summer/early fall (e.g. Taka-
hashi et al., 2002b). Highest annual production rates are as-
sociated with shelf areas and vary regionally between > 200
and > 800 g C m$^{-2}$ (Arzhanova et al., 1995; Springer et al.,
1996; Stabeno et al., 1999). Available nutrients are reported


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to be often fully consumed during seasonal blooms (Niebauer et al., 1995).

Sea ice begins to form during October/November on the northern Bering Sea continental shelf (Anadyr Bay, Bering Strait), reaches its maximum distribution in March/April, and subsequently declines until July (Tomczak and Godfrey, 1994; Niebauer et al., 1999; Lisitzin, 2002). Sea-ice formation takes place in shallow shelf areas, bays, and coastal areas. As in the Arctic, coastal polynyas play an important role for the build-up of sea ice, and consequently for water mass ventilation due to brine rejection (Niebauer et al., 1999; Stabeno et al., 1999). Processes entraining sediment into newly formed ice involve tidal sea-level oscillations, wind mixing, resuspension of sediments from the seafloor (suspension freezing), beach-ice formation, nearshore anchor ice formation, and seabed freezing (e.g. Nürnberg et al., 1994, 2011, and references therein; Stein, 2008, and references therein). Especially during fall and winter, storms affect reworking and resuspension processes by sea-ice crushing, mixing of the water column, and detachment of the sediment-laden ice from the coast. The sediment freight is released by
sea-ice melting, especially during spring/summer, and then contributes to (hemi-) pelagic sedimentation.

3 Material and methods

This study is based on piston cores SO201-2-77KL, -85KL, and -101KL from Shirshov Ridge, western Bering Sea (Fig. 1, Table 1). The cores were recovered along a ~280 km-long north-south transect from intermediate to deep water levels (630–2135 m) during R/V Sonne cruise SO201-KALMAR Leg 2 in 2009 (Dullo et al., 2009). The sedimentary succession is characterized by monotonous sequences of mainly clay- and silt-sized siliciclastic material, which are repeatedly interrupted by layers of diatomaceous ooze. Except for the younger part of the Holocene, which mainly consists of diatom-rich sediment, our sediment cores contained low but sufficient concentrations of CaCO3 to also allow for high-resolution foraminifera-based reconstructions. However, for cores 85KL and 101KL sediments younger than 7.5 and 12.5 ka BP, respectively, were not recovered.

3.1 Stratigraphic approach

3.1.1 Core logging

Color reflectance measurements were carried out using a Minolta CM 508d hand-held spectrophotometer at 1 cm-spaced intervals (Dullo et al., 2009). Reflectance data were automatically converted by Spectramagic software into CIE L*, a* and b* color space (CIELAB). The Avaatech X-ray fluorescence (XRF) core scanner at Alfred Wegener Institute for Polar and Marine Research, Bremerhaven, was used to determine relative changes in the sedimentary elemental composition. Core scanning was performed on the split core surface covered with SPEX CertiPrep 3525 Ultralene foil (4 µm thick). Each core segment was triple-scanned for analysis of elements Al through to Ba at 1 mA, but at different tube voltages and count times (10 kV, 10 s; 30 kV, 15 s; 50 kV, 30 s), using a sampling resolution of 1 cm. Results are considered semiquantitative (Richter et al., 2006; Tjallingii et al., 2007) and are given as count rates (in cps) or as log-ratios (natural logarithm) of element count rates.

3.1.2 Stable oxygen isotopes

For stable oxygen isotope (δ18O) stratigraphy we used endobenthic foraminiferal species Uvigerina peregrina and Uvigerina auberiana, since Uvigerina δ18O values are reported to be in equilibrium with seawater (Shackleton and Hall, 1984). δ18O was measured every 5 cm on 2–3 tests of U. peregrina collected from the 315–355 µm size fraction. For core 77KL we used tests of U. auberiana when U. peregrina was not present. The measurements were performed at GEOMAR, Kiel, using a MAT253 mass spectrometer (Thermo Scientific, Germany) coupled with a Kiel IV Carbonate device (Thermo Scientific, Germany). Results were referenced to the NBS19 standard and calibrated to the VPDB scale. Long-term precision (n > 1000 samples) for δ18O of the used carbonate standard (Solnhofen limestone) was ±0.06‰. In core 77KL benthic foraminifera were only preserved until 865 cm core depth.

3.1.3 Paleomagnetics

Sedimentary natural remanent magnetization (NRM) was determined in core 85KL based on saturation magnetization in magnetic-hysteresis parameters, and by differential thermomagnetic analyses using a Faraday magnetic balance and a coercive spectrometer (Burov et al., 1986; Yasonov et al., 1998). The NRM module and direction were measured with a AGICO JR-5A spinner magnetometer after the stepwise demagnetization of a reference sample. Magnetic cleaning was performed using an alternating magnetic field with an amplitude of 10 mT to recognize the characteristic component of NRM (ChRM). Anhysteretic remanent magnetization (ARM) was generated in the preliminarily demagnetized samples using a AGICO AMU-1A anhysteretic magnetizer under a constant field of 0.05 mT and a maximum alternating field of 100 mT. Relative paleointensity of the magnetic field (RPI) was then calculated by normalization of ChRM to ARM. Scalar petromagnetic properties (SPP) were additionally determined (Enkin et al., 2007; Malakhov et al., 2009). All measurements were conducted at NEISRI, Magadan.

Takahashi et al. (2011) reported on the transformation of oxide magnetic minerals into paramagnetic FeS2 in Bering Sea sediments. They associated the formation of FeS2 with sulfate reduction processes under anaerobic methane oxidation. We performed quality control of the magnetic hysteresis parameters (Day-plot, orthogonal projections of AF demagnetization of NRM, et al.) and the thermomagnetic analyses of the magnetic fraction, which also indicate an influence by sediment diagenesis in core 85KL. Specifically, the thermomagnetic analyses suggest the occurrence of paramagnetic FeS2 in the major ferrimagnetic phase from 880 to 1760 cm core depth. We assume that the dissolution of very finely dispersed, almost single-domain magnetic particles and the preservation of coarse pseudo-single-domain and multi-domain particles just led to decreasing RPI values within that depth interval. However, this most probably had no effect on the relative variability of the reconstructed geomagnetic field.

3.1.4 Age models

The chronostratigraphic approach included high-resolution core logging data (color b*, XRF scanning), benthic δ18O stratigraphy, magnetostratigraphy, and accelerator mass spectrometry (AMS) radiocarbon dating of planktonic foraminifera for absolute age control. A detailed presentation
of the stratigraphic framework for the last 20 kyr, including the AMS-14C dating results, is provided in Max et al. (2012). The pre-deglacial (> 20 ka BP) stratigraphic framework of our cores is primarily based on the graphic correlation between color b* recorded in core 85KL and the Dansgaard–Oeschger (D-O; e.g. Dansgaard et al., 1993) climate variability registered in the NGRIP δ18O record (NGRIP members, 2004; GICC05 timescale, Rasmussen et al., 2006) (Fig. 2a). For ages > 122 ka BP and for identification of the Marine Isotope Stage (MIS) 5.5 climate optimum (∼ 125 ka BP) color b* and XRF Ca/Ti log-ratios were correlated to the Sanbao stalagmite δ18O record (Wang et al., 2008). The Laschamp (∼ 42 ka BP), Norwegian–Greenland Sea (∼ 65 ka BP), and Blake (∼ 117 ka BP) paleomagnetic excursions were identified in the RPI record and correlated with the PISO-1500 geomagnetic paleointensity stack (Channell et al., 2009). Further ages were derived from comparison of benthic δ18O values and SPP (not shown) with the global reference stack LR04 (Lisiecki and Raymo, 2005), which was also used to identify boundaries of MIS 1 to 6. The stratigraphy of core 85KL was transferred via intercore correlations (color b*, XRF Ca/Ti log-ratios) to cores 77KL and 101KL (Fig. 2b), for which similar stratigraphic approaches were carried out. All age-depth points are provided in Appendix A.

Figure 3 shows the age models for our cores by direct comparison with the used reference records and a respective age versus depth diagram. Age models were tested via spectral analysis of the color b* and benthic δ18O records in the time domain to detect orbital frequencies. Spectral analysis was performed using the AnalySeries 2.0 software (Paillard et al., 1996). Despite the shortness of our records with respect to orbital-scale changes, we found dominant cyclicities of ∼ 23 and ∼ 39 kyr, which within appropriate bandwidths match frequencies of orbital precession (0.047 ± 0.005 kyr−1) and obliquity cycles (0.025 ± 0.0015 kyr−1) (Fig. 3).

### 3.2 Sedimentation and accumulation rates

Linear sedimentation rates (LSR, in cm kyr−1) were calculated between age control points. Bulk accumulation rates (ARBulk, in g cm−2 kyr−1) were calculated as the product of LSR and the dry bulk density (DBD, in g cm−3, determined each 5 cm). Records of LSR and ARBulk are shown in Fig. 4. Cores from Shirshov Ridge have average LSR (ARBulk) of 11–16 cm kyr−1 (7–15 g cm−2 kyr−1), and hence allow for centennial- to millennial-scale reconstructions. Variability and average values of LSR and ARBulk increase toward the northernmost site. In general, LSR and ARBulk are higher during cold intervals (MIS 6, 5.4, 5.2, and 4) than during warm intervals (MIS 5.5, 5.3, 5.1, 3, and 1), but highest during Termination I (20–10 ka BP) (Fig. 4). Holocene sediments are either not fully recovered or subject to low LSR.

### Table 1. Site information.

<table>
<thead>
<tr>
<th>Core</th>
<th>Latitude</th>
<th>Longitude</th>
<th>Depth (m b.s.l.)</th>
<th>Recovery (m)</th>
</tr>
</thead>
<tbody>
<tr>
<td>SO201-2-77KL</td>
<td>56°19.83′N</td>
<td>170°41.98′E</td>
<td>2135</td>
<td>11.78</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>57°30.30′N</td>
<td>170°24.77′E</td>
<td>968</td>
<td>18.13</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>58°52.52′N</td>
<td>170°41.45′E</td>
<td>630</td>
<td>18.32</td>
</tr>
</tbody>
</table>

**Fig. 2.** (A) Correlation of sediment core SO201-2-85KL with the PISO-1500 (thick grey line) paleomagnetic reference record (Channell et al., 2009) based on relative paleointensity (RPI, smoothed by a 5-point-running average), as well as with the Sanbao (grey lines) and Hulu (orange lines) stalagmite δ18O records (Wang et al., 2001, 2008) and the NGRIP δ18O record (NGRIP members, 2004; GICC05 timescale, Rasmussen et al., 2006) based on color b*. Black lines mark correlation lines. The Laschamp (L), Norwegian–Greenland Sea (N), and Blake (B) paleomagnetic events are indicated. Bottom numbers mark D-O events. (B) Intercore correlation of sediment cores SO201-2-77KL (green) and -101KL (blue) with core -85KL (red) based on color b* and XRF Ca/Ti log-ratio records.
Fig. 3. Left: comparison of proxy records from sediment cores SO201-2-77KL (green lines), -85KL (red lines), and -101KL (blue lines) with published reference records (grey lines). Age models are primarily based on the graphic correlation between color b* records and the NGRIP $\delta^{18}$O record (NGRIP members, 2004; GICC05 timescale, Rasmussen et al., 2006). Benthic $\delta^{18}$O values from *U. peregrina* (plus symbols) and *U. auberiana* (open triangles) are in agreement with the global reference stack LR04 (Lisiecki and Raymo, 2005). Relative paleointensity (RPI, smoothed by a 5-point-running average) recorded in core 85KL compares with the paleomagnetic reference record PISO-1500 (Channell et al., 2009). L, N, and B mark the Laschamp, Norwegian–Greenland Sea, and Blake paleomagnetic events. Absolute age control is provided by AMS-$^{14}$C-dating (coloured triangles; see Max et al., 2012). Top numbers indicate Marine Isotope Stages (boundaries after Lisiecki and Raymo, 2005) and D-O events. Upper right: age versus depth diagram showing the age-depth points and their underlying stratigraphic approach (see also Appendix A). Lower right: spectral analysis of benthic $\delta^{18}$O and color b* records performed in the time domain revealed dominant cyclicities that lie within the frequency bands of Earth’s obliquity and precession cycles.

3.3 Assessment of marine productivity

Past changes in marine productivity (paleo-export production) were approximated from total organic carbon (TOC), CaCO$_3$, biogenic opal, biogenic barium (Ba$_{bio}$), and XRF logging data. This multi-proxy approach was necessary due to specific restrictions of the respective proxies. TOC preservation in sediments is highly debated (e.g. Hartnett et al., 1998; Ganeshram et al., 1999; Thunell et al., 2000; Hedges et al., 2001) as it is, e.g. influenced by oxidation processes and bottom water ventilation (De La Rocha, 2007). Also, its source can be of marine and terrestrial origin. In the N Pacific the preservation of CaCO$_3$ is limited by the shallow lysocline. Accordingly, CaCO$_3$ concentrations rather reflect changes in the bottom water calcite saturation state (e.g. Jacard et al., 2005; Gebhardt et al., 2008), although changes in
biological CaCO$_3$ production can not be fully excluded to explain changes in CaCO$_3$. Opal dissolves during settling to the seafloor, but its preservation is independent from bottom water oxygenation. Since opal-rich sediments are linked to biogenic silica production (e.g. Honjo, 1990; Nelson et al., 1995; Ragueneau et al., 2000; Pondaven et al., 2000) biogenic opal is most often used in N Pacific reconstructions of marine productivity. Barium, present as sedimentary barite is used in several studies to reconstruct paleoproductivity (e.g. Dymond et al., 1992; Francois et al., 1995; Dymond and Collier, 1996; Gingele et al., 1999). Barite particles occur in areas of high new production (Dehairs et al., 1991) and laboratory experiments showed that the decay of phytoplankton in undersaturated seawater results in barite formation (Ganeshram et al., 2003).

3.3.1 CN-analyses and biogenic opal

Total carbon (TC), TOC, and total nitrogen (TN) were determined on freeze-dried bulk sediment samples of 20 mg using a Carlo Erba NA-1500 CNS analyzer. TOC was measured on previously decalcified samples and CaCO$_3$ concentrations were calculated by multiplication of the difference between TC and TOC with a factor of 8.333. Reproducibility of the TOC and TN measurements was ±0.03 wt % and ±0.01 wt %, respectively.

The molar ratio of TOC to TN, corrected for inorganic nitrogen compounds (referred to as [C/N]$_a$ hereafter), was used to distinguish between marine and terrestrial sources of TOC. The Redfield ratio (Redfield et al., 1963) translates the C/N ratio of marine organic matter to ~ 7. Typical terrigenous values lie between 20–200 (Hedges et al., 1986). We applied a correction for total inorganic nitrogen (TIN), usually clay-bound inorganic ammonium (Müller, 1977), based on a linear relationship between TOC and TN (after Göth et al., 1998). Results showed TIN concentrations of 0.01–0.02 wt % that were assumed constant downcore.

Biogenic opal was measured following Müller and Schneider (1993) using molybdate-blue spectrophotometry. Silica was extracted from 20 mg of freeze-dried bulk sediment samples. Results were evaluated applying the procedure of DeMaster (1981). Replicate measurements showed a reproducibility of 1–2 wt %.

3.3.2 Biogenic barium (Ba$_{bio}$)

Concentrations of major (Al, Ti, Fe, K) and trace (Ba) elements were determined on discrete samples of 0.6 g of freeze-dried bulk sediment using a Philips PW1480 XRF spectrometer without determination of loss on ignition following standard procedures. Analytical precision, determined for the BHVO standard, was < 2 % RSD for the major elements and ± 30 ppm for Ba (N = 15). Results for barium (Ba$_{total}$) are the sum of biogenic (Ba$_{bio}$) and nonbiogenic Ba-portions. Ba$_{bio}$ was therefore calculated via concentrations of Al by estimating the aluminosilicate contribution of Ba considering the global average Ba/Al ratio for pelitic rocks of 6.5 mg g$^{-1}$ (Wedepohl, 1971):

$$\text{Ba}_{bio} = \text{Ba}_{total} - \text{Al} \times 0.0065$$

(1)

Ba$_{bio}$ was subsequently used to assess new production ($P_{New}$) by applying the relationship of Nürnberg (1995) (see also Fig. 7). Annual primary production (PP) was calculated after Eppley and Peterson (1979) using the $P_{New}$ estimates.

3.3.3 XRF logging data (Br, Ca/Ti, Si/Al)

Several logging data and bulk geochemical analyses showed a similar temporal evolution. In this respect, XRF count rates of Br correlated with TOC concentrations (0.35 < $R^2$ < 0.74), which supports a relationship between TOC and biophilic halogen bromine (Ziegler et al., 2008). XRF Ca/Ti log-ratios correlated with CaCO$_3$ concentrations (0.07 < $R^2$ < 0.65), which is explained by assuming a detrital origin of Ti. Normalization of XRF-derived Ca to Ti and/or Al abundances has been applied before (Jaccard et al., 2005) and is thought to reflect biogenic CaCO$_3$ contents within the sediment. The XRF signals for Al were better than for Ti and were subsequently used for normalization. However, we favored Ca/Ti over Ca/Al log-ratios due to better correlation to CaCO$_3$. Although opal concentrations...
were close to reproducibility for most samples, they correlated with color \( b^* \) (0.20 < \( R^2 < 0.60 \)) and XRF Si/Al log-ratios (0.04 < \( R^2 < 0.73 \)). This is in accordance with other studies considering a connection between Si/Al ratios and biogenic opal (McDonald et al., 1999) and between color \( b^* \) and organic matter/opal (Debret et al., 2006).

All linear relationships found are strong for cores 77KL and 85KL (0.57 < \( R^2 < 0.74 \)), but weak for core 101KL (0.04 < \( R^2 < 0.35 \)). Since the linear correlation coefficients strongly varied between both, the respective sites and the respective proxies, the logging data were not used to apply calibration functions, but are shown together with the quantitative results.

### 3.4 Assessment of terrigenous matter supply

#### 3.4.1 Coarse material, siliciclastics, and terrigenous matter

Changes in terrigenous fluxes were approximated from a set of sedimentological and geochemical proxies together with XRF logging data. The proportions of coarse (> 63 µm) and fine (< 63 µm) material (in wt %) were determined every 5 cm by weighing of the dried sediment before and after wet-sieving through a 63 µm mesh. Magnetic susceptibility (not shown) was logged using a GEOTEK Multi-Sensor Core Logger in combination with a Bartington MS2C sensor loop each 1 cm on the unopened core segments directly after recovery (Dullo et al., 2009). Records of > 63 µm showed only weak relationships with magnetic susceptibility records (\( R^2 < 0.13 \)), which were characterized by low values (< 15 SI units). Hence, magnetizable minerals are most probably mainly bound to the fine fractions.

Relative amounts of siliciclastics (%Siliciclastics) were calculated by considering the bulk sediment to be composed of siliciclastics, CaCO\(_3\), TOC, and biogenic opal. TN concentrations were generally < 0.3 wt % and not included in the calculation:

\[
\text{%Siliciclastics} = 100\% - (\text{CaCO}_3\% + \text{TOC}\% + \text{Opal}\%).
\] (2)

Respective accumulation rates (AR\(_{\text{Siliciclastics}}\)) were calculated by dividing %Siliciclastic by 100 and subsequent multiplication with AR\(_{\text{Bulk}}\). The accumulation rate of the biogenic components (AR\(_{\text{Biogenic}}\)) was calculated by subtracting AR\(_{\text{Siliciclastics}}\) from AR\(_{\text{Bulk}}\).

In a second approach, relative amounts of terrigenous matter (%Terrigen) were calculated by normalizing bulk sedimentary Al and Ti concentrations to their concentration in average continental crust (Al = 3117 µmol g\(^{-1}\), Ti = 112.8 µmol g\(^{-1}\); Taylor and McLennan, 1995). Both normalizations resulted in a similar temporal evolution of the records, but Al-normalized results, which were on average 4–6 % higher than Ti-normalized results, better compared to the records of %Siliciclastics.

#### 3.4.2 Lithogenous elements

The geochemistry of lithogenous elements can be used to approximate continental input (e.g. Duce and Tindale, 1991; Bareille et al., 1994), dust supply (e.g. Boyle, 1983; Calvert and Fortugue, 2001), terrestrial runoff (e.g. Schmitz, 1987; Jansen et al., 1998), and mineralogical variations (e.g. Schneider et al., 1997; Yarincik et al., 2000). As described above, concentrations of Al, Ti, Fe, and K were determined via discrete XRF bulk analyses. The elements and their respective ratios were used to identify sources of terrigenous matter and to reconstruct variations in terrigenous fluxes. In cores 85KL and 101KL XRF logging data of K/Ti log-ratios correlated well with molar K/Ti ratios (\( R^2 > 0.64 \)) and are shown for comparison.

### 4 Results and discussion

#### 4.1 Marine productivity

##### 4.1.1 Proxy data

Our records for export production show a similar temporal evolution in all cores. Proxy concentrations increase toward the southernmost site, whereas the ranges of AR\(_{\text{Biogenic}}\) are comparable and low for all sites (< 3 g cm\(^{-2}\) kyr\(^{-1}\); Fig. 4). Results for TOC, opal, CaCO\(_3\), as well as their approximating logging data are shown in Fig. 5. In general, concentrations and their approximating XRF data remained low during cold intervals (MIS 6, 5.4, 5.2, 4 to 2), but high during warm intervals (MIS 5.5, 5.3, 5.1, and 1), with maximum values recorded during interglacials in core 77KL. Core 101KL exhibits lowest proxy concentrations and amplitude variations (Table 2). Overall concentrations hardly exceed ~1 wt % for TOC, ~3 wt % for opal, and ~2 wt % for CaCO\(_3\). MIS 5.3 and 5.1, as well as interstadials are characterized by ~1 to 3-times higher values at most. Notably, cores 85KL and 101KL recorded interstadial-like events during MIS 6 (at ~173, 164, 148, 137, and 133 ka BP), characterized by increased proxy concentrations or their approximating XRF data. Deglacial and interglacial maxima in TOC, opal, and CaCO\(_3\) reach values of ~2, ~50, and ~30 wt %, respectively. Maxima recorded during Termination I reflect the warm phases of the Bølling–Allerød (B/A; 14.7–12.9 ka BP, Blockley et al., 2012) and Preboreal (PB; ~11.7–11.0 ka BP), whereas deglacial minima are considered to correspond to the Heinrich Stadial 1 (HS1; 18.0–14.7 ka BP, Sarnthein et al., 2001) and Younger Dryas cold phases (YD; 12.9–11.7 ka BP, Blockley et al., 2012).

During Termination I, TOC appears to lead the deglacial increase of the other productivity proxies by ~2 kyr at our sites, which is in agreement with previous results from the whole Bering Sea realm (Gorbarenko, 1996; Okazaki et al., 2005; Kim et al., 2011). Although records of TOC and XRF count rates of Br correspond well in our cores (Fig. 5), Br
does not follow this deglacial TOC increase, suggesting a changing source of TOC. In contrast to TOC and CaCO$_3$, we observed only minor increases in opal during the B/A, but a subsequent gradual increase into the Holocene. These results from the western Bering Sea are comparable to opal records from the northern slope (Itaki et al., 2009; Kim et al., 2011). Despite showing a similar temporal evolution, opal records from Bowers Ridge (Okada et al., 2005; Okazaki et al., 2005; Brunelle et al., 2007, 2010), Umnak Plateau (Okada et al., 2005; Okazaki et al., 2005), and the southern Okhotsk Sea (Gorbarenko, 1996; Gorbarenko et al., 2002a, b; Narita et al., 2002; Brunelle et al., 2010) have 1.5 to 3-times higher values.

CaCO$_3$ concentrations at our sites are generally low and related to the abundance of foraminifera and nannoplanktonic remains (coccoliths). High XRF Ca/Ti log-ratios during MIS 5.5 are the result of decreased XRF counts of Ti and not of increased Ca counts. Short-lived increases of up to $\sim$3 wt% were recorded during MIS 6, interstadials, and Termination I (Fig. 5). Similar observations, especially for Termination I, were also reported for cores from the northern slope, Bowers Ridge, and Umnak Plateau (Cook et al., 2005;
Okazaki et al., 2005; Brunelle et al., 2007, 2010; Itaki et al., 2009; Khim et al., 2010; Kim et al., 2011). At ODP Site 882 in the N Pacific interglacial maxima in CaCO$_3$ are accompanied by maxima in Ba$_{bio}$ (Jaccard et al., 2005). Since enhanced preservation of CaCO$_3$ is explained by a release of deep-sequestered CO$_2$ from the deep-ocean basin (Broecker and Peng, 1987; Marchitto et al., 2005), these CaCO$_3$ maxima were suggested to be the result of a higher bottom water calcite saturation state in response to the weakening of the N Pacific halocline (Jaccard et al., 2005). Deglacial maxima of CaCO$_3$ in Bering Sea sediments were also explained by denitrification on continental shelves, which might have resulted in an increase in alkalinity and, thus, in enhanced carbonate preservation (Chen, 2002; Okazaki et al., 2005). Today, the calcite saturation horizon in the Bering Sea is reported to lie above 500 m water depth (Feely et al., 2002) and at our sites lies above 200 m water depth (Riethdorf et al., 2013). Accordingly, we consider CaCO$_3$ maxima in our cores to mainly reflect a higher bottom water calcite saturation state, but enhanced biological CaCO$_3$ production can not be ruled out.

4.1.2 Organic carbon source

Average concentrations of bulk sedimentary TN varied at ~0.1 wt %. Hence, sediments from Shirshov Ridge contain a considerable amount of TIN, which results in [C/N]a ratios that are by up to 4 units higher than (uncorrected) C/N ratios (Fig. 6a). [C/N]a ratios in our cores mainly vary between 10 and 15 (Fig. 6a, Table 2), indicating that TOC input contains considerable amounts of terrestrial organic material. These values compare to those found at the northern slope (Khim et al., 2010) and in the central Okhotsk Sea (Nürnberg and Tiedemann, 2004). For the eastern and southern part of the Aleutian Basin (Nakatsuka et al., 1995) and for the southern Okhotsk Sea (Ternois et al., 2001) lower C/N ratios of 6–9 were reported. All our records show a rise in [C/N]a ratios during Termination I. At Site 77KL, the deglacial increase in [C/N]a ratios starts at ~17 ka BP, with maximum values during the YD, and a subsequent decrease into the Holocene. The same deglacial evolution was observed at the northern slope (Khim et al., 2010) and in the southern Okhotsk Sea (Ternois et al., 2001; Seki et al., 2003). It was related to the discharge of terrestrial material from the flooded shelf due to sea-level rise. In contrast, cores from the eastern and southern Bering Sea show a gradual decrease of C/N ratios since the last glacial maximum (LGM) (Nakatsuka et al., 1995).

4.1.3 $\text{Ba}_{bio}$ and export production

Our records of $\text{Ba}_{bio}$ are shown in Fig. 7 and overall match the TOC and opal records, which argues against a preservation effect due to sulfate reduction and associated barite dissolution. Accordingly, we consider changes in $\text{Ba}_{bio}$ concentrations to mainly reflect variations in $\text{Ba}_{bio}$ accumulation. A potential source of error in the calculation of $\text{Ba}_{bio}$ comes from estimating the aluminosilicate contribution of Ba via Al. Ba/Al ratios range between 5–10 mg g$^{-1}$ in crustal rocks (Taylor, 1964; Röslér and Lange, 1972) with a crustal average of 7.5 mg g$^{-1}$ (Dymond et al., 1992). We estimated the regional detrital Ba/Al ratio from surface sediment samples following Klump et al. (2000), which resulted in a value of 7 mg g$^{-1}$ (unpublished data). This value is close to the global average of pelitic rocks of 6.5 mg g$^{-1}$ (Wedepohl, 1971), which was used for reconstructing $\text{Ba}_{bio}$ in the central Okhotsk Sea (Nürnberg and Tiedemann, 2004). $\text{Ba}_{bio}$ variability is low during most of the time covered by the cores, with average concentrations increasing from Site 101KL (~300 ppm), via Site 85KL (~400 ppm), to Site 77KL (~500 ppm) (Fig. 7, Table 2). Significant increases only occurred at Site 77KL during MIS 5.5 and the Holocene with concentrations of ~1000 ppm and ~1700 ppm, respectively. Also at Site 77KL, XRF Ba/Al log-ratios covary with $\text{Ba}_{bio}$ concentrations (not shown; $R^2 = 0.74$), thereby revealing minor increases during MIS 5.3, 5.1, and short-lived maxima during interstadials. A similar range and variability as found for core 77KL was observed at Bowers Ridge (Brunelle et al., 2007), ODP Site 882 (Jaccard et al., 2005),

Table 2. Statistics of parameters approximating marine productivity.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>SO201-2-77KL</th>
<th>SO201-2-85KL</th>
<th>SO201-2-101KL</th>
</tr>
</thead>
<tbody>
<tr>
<td>TOC (wt %)</td>
<td>Avg. 0.85</td>
<td>StDev. 0.30</td>
<td>Avg. 0.96</td>
</tr>
<tr>
<td>[C/N]a</td>
<td>Opal 13.0</td>
<td>2.9</td>
<td>11.9</td>
</tr>
<tr>
<td>CaCO$_3$ (wt %)</td>
<td>1.9</td>
<td>3.2</td>
<td>1.9</td>
</tr>
<tr>
<td>$\text{Ba}_{bio}$ (ppm)</td>
<td>733</td>
<td>330</td>
<td>436</td>
</tr>
<tr>
<td>$P_{\text{N}_\text{Tot}}$ (gC m$^{-2}$ yr$^{-1}$)</td>
<td>50.6</td>
<td>47.5</td>
<td>34.9</td>
</tr>
<tr>
<td>$PP$ (gC m$^{-2}$ yr$^{-1}$)</td>
<td>131.9</td>
<td>54.2</td>
<td>113.1</td>
</tr>
</tbody>
</table>

as well as in the sea-ice-free Antarctic Zone of the Southern Ocean (Nürnberg et al., 1997). Cores from the southern (Brunelle et al., 2010) and central Okhotsk Sea (Sato et al., 2002; Nürnberg and Tiedemann, 2004), as well as from the sea-ice-influenced Antarctic Zone of the Southern Ocean (Nürnberg et al., 1997) exhibit generally lower glacial (~200–400 ppm) and peak interglacial (~800–1000 ppm) contents of Ba\textsubscript{bio}, being more comparable to sites 85KL and 101KL. Notably, Sato et al. (2002) and Brunelle et al. (2010) for the Okhtosk Sea report a deglacial lead in the rise of Ba\textsubscript{bio} prior to that observed for opal contents, which we can not verify from our records.

We calculated \( P_{\text{New}} \), i.e. primary production that results from allochthonous nutrient inputs to the euphotic zone, from Ba\textsubscript{bio} following Nürnberg (1995) rather than from TOC, which at our sites is affected by terrestrial carbon (Fig. 7). This approach was also followed by Nürnberg and Tiedemann (2004) for a respective reconstruction in the Okhtosk Sea. Surface sediment samples from Shirshov Ridge reveal a north–south gradient in modern \( P_{\text{New}} \) values within \( \sim 3–54 \text{ gC m}^{-2} \text{ yr}^{-1} \), translating into PP values of \( \sim 35–145 \text{ gC m}^{-2} \text{ yr}^{-1} \) (estimated from \( P_{\text{New}} \), see Appendix B). Considering the uncertainties for AR\textsubscript{Bulk} and Ba\textsubscript{bio}, this result is in agreement with Springer et al. (1996), who reported a modern PP range of 50–100 gC m\textsuperscript{-2} yr\textsuperscript{-1} (average of 61 gC m\textsuperscript{-2} yr\textsuperscript{-1}) for the “oceanic domain” of the Bering Sea, in which our sites are located. On the eastern Bering Sea shelf edge, modern PP is reported to lie between 175–275 gC m\textsuperscript{-2} yr\textsuperscript{-1} (Springer et al., 1996). Stabeno et al. (1999) reported modern PP values of \( \sim 200 \text{ gC m}^{-2} \text{ yr}^{-1} \) over the southeastern shelf and \( > 800 \text{ gC m}^{-2} \text{ yr}^{-1} \) north of St. Lawrence Island, whereas Arzhanova et al. (1995) found \( > 400 \text{ gC m}^{-2} \text{ yr}^{-1} \) over the western shelf.

Our downcore results for \( P_{\text{New}} \) show that primary production was commonly low (< 50 gC m\textsuperscript{-2} yr\textsuperscript{-1}) at our sites during most of the last 180 kyr (Fig. 7, Table 2). Only core 77KL is characterized by two maxima > 150 gC m\textsuperscript{-2} yr\textsuperscript{-1} during Termination I. Although these maxima correspond to higher concentrations of Ba\textsubscript{bio}, they are likely overestimated due to the use of AR\textsubscript{Bulk} in the calculation of \( P_{\text{New}} \). Nevertheless, our downcore results are comparable to those reported for the central Okhotsk Sea (Nürnberg and Tiedemann, 2004) and the Antarctic continental margin (Gingele et al., 1999), albeit these show lower glacial values of \(< 10 \text{ gC m}^{-2} \text{ yr}^{-1} \). PP mainly remained below 150 gC m\textsuperscript{-2} yr\textsuperscript{-1} at our sites (not shown). Only the deglacial \( P_{\text{New}} \) maxima in core 77KL would translate into significantly higher PP values of \( \sim 250–300 \text{ gC m}^{-2} \text{ yr}^{-1} \) (Table 2). Consequently, PP most probably remained permanently low at our sites and never reached values as observed today on the eastern Bering Sea shelf.

4.2 Terrigenous matter supply

4.2.1 Proxy data

Sediments from Shirshov Ridge contain on average > 85 % siliciclastics and only < 10 % coarse material (Table 3), thereby tying terrigenous matter mainly to the fine fractions. Light microscope observations reveal silt to fine sand-sized, angular-shaped quartz grains as major components, supplemented by minor portions of feldspar and mica. Dropstones (1–2 cm in diameter) appear as well-rounded pebbles, but they are commonly rare although occurring over the entire length of the cores. The records approximating terrigenous matter supply exhibit a similar temporal evolution at our sites (Fig. 8), but in contrast to the productivity proxies decreasing concentrations toward the southernmost site are observed. This and the observation that proxy records reflecting export production and terrigenous matter supply are negatively correlated is mainly attributed to sedimentary dilution by the biogenic components. However, changes in AR\textsubscript{Bulk} (and LSR) are obviously largely determined by changes in...
Fig. 7. Left: concentrations of lithogenous elements Ti, Fe, and Al (angular brackets), as well as of biogenic barium (Ba\textsubscript{bio}) and new production (P\textsubscript{New}) for the last 150 kyr. P\textsubscript{New} was calculated from Ba\textsubscript{bio} using the equation of Nürnberg (1995): 
\[ P_{\text{New}} = 3.56 \times F_{\text{Ba}_{\text{bio}}}^{1.504} \times z^{-0.0937}, \]
where F\textsubscript{Ba\textsubscript{bio}} = AR\textsubscript{Ba\textsubscript{bio}} / \left[ 0.209 \times \log_{10}(AR_{\text{Bulk}} \times 1000) - 0.213 \right] (Dymond et al., 1992), and z = water depth (in m). F\textsubscript{Ba\textsubscript{bio}} is the flux of biogenic Ba to the seafloor (in µg cm\textsuperscript{-2} yr\textsuperscript{-1}), AR\textsubscript{Ba\textsubscript{bio}} is the accumulation rate of biogenic Ba (in mg cm\textsuperscript{-2} kyr\textsuperscript{-1}). Note that for records of [Ti], [Fe], and [Al] y-axes are inverted. Right: correlation between lithogenous elements and respective linear correlation coefficients (R\textsuperscript{2}) indicate a shared terrigenous source of these elements.

Table 3. Statistics of parameters approximating terrigenous matter supply.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>SO201-2-77KL</th>
<th>SO201-2-85KL</th>
<th>SO201-2-101KL</th>
</tr>
</thead>
<tbody>
<tr>
<td>&gt; 63 µm (wt %)</td>
<td>5.6</td>
<td>3.1</td>
<td>7.5</td>
</tr>
<tr>
<td>%Siliciclastics</td>
<td>88.9</td>
<td>8.8</td>
<td>94.1</td>
</tr>
<tr>
<td>%Terrigen (Al-norm.)\textsuperscript{(1)}</td>
<td>82.7</td>
<td>10.2</td>
<td>88.2</td>
</tr>
<tr>
<td>%Terrigen (Ti-norm.)\textsuperscript{(2)}</td>
<td>76.6</td>
<td>9.6</td>
<td>84.4</td>
</tr>
<tr>
<td>[Ti] (µmol g\textsuperscript{-1})</td>
<td>86.4</td>
<td>10.9</td>
<td>95.2</td>
</tr>
<tr>
<td>[Fe] (µmol g\textsuperscript{-1})</td>
<td>762</td>
<td>112</td>
<td>792</td>
</tr>
<tr>
<td>[Al] (µmol g\textsuperscript{-1})</td>
<td>2578</td>
<td>316</td>
<td>2751</td>
</tr>
</tbody>
</table>

\textsuperscript{(1)} Using [Al] = 3117 µmol g\textsuperscript{-1} of average continental crust (Taylor and McLennan, 1995). \textsuperscript{(2)} Using [Ti] = 112.8 µmol g\textsuperscript{-1} of average continental crust (Taylor and McLennan, 1995).
Fig. 8. Records approximating changes in terrigenous matter supply over the past 180 kyr. Relative contents of coarse material (> 63 µm), terrigenous matter (normalized to Al concentrations of continental crust; Taylor and McLennan, 1995), and molar K/Ti ratios (colored lines), in comparison to relative amounts of siliciclastics, and XRF K/Ti log-ratios (underlying grey lines) for cores SO201-2-77KL (green lines), -85KL (red lines), and -101KL (blue lines). Note that all y-axes are inverted, except for > 63 µm.

the accumulation of siliciclastic material (AR\textsubscript{Siliciclastics}), and therefore by changes in terrigenous matter supply (Fig. 4). Accordingly, cold intervals (MIS 6, 5.4, 5.2, 4 to 2) are characterized by high terrigenous matter supply, whereas warm intervals (MIS 5.5, 5.3, 5.1, and 1) show reduced proxy concentrations or ratios. Notably, a meridional gradient in AR\textsubscript{Siliciclastics} is observed along the core transect with lower values and therefore potentially decreasing terrigenous matter supply toward the southernmost site (Fig. 4). The most pronounced minima in the proxy records are found during interglacials. In core 77KL higher amounts of > 63 µm and of CaCO\textsubscript{3} concentrations (XRF Ca/Ti log-ratios) occur synchronously, indicating that changes in CaCO\textsubscript{3} result from higher foraminiferal abundances. Although independently derived, covariation is also found between %Terrigen and %Siliciclastics, which are characterized by almost identical ranges (Fig. 8, Table 3).

K/Ti ratios show a range of 5–7 mol mol\textsuperscript{-1} in all cores (Fig. 8, Table 3). We consider this ratio to be indicative of changes in the geochemical composition of the terrigenous matter itself. During MIS 5.5 and the Holocene, %Siliciclastics are reduced to ~50% at Site 77KL, ~60% at
Fig. 9. Records of XRF Si/Al log-ratios reflecting changes in biogenic opal concentrations (marine productivity), XRF records of Fe (in cps) reflecting relative changes in terrigenous matter supply, and linear sedimentation rates (LSR) at sites SO201-2-77KL (green lines), -85KL (red lines), and -101KL (blue lines) for the last 180 kyr. Logging data are smoothed by 5-point-running averages. Relative sea level (RSL) is after Waelbroeck et al. (2002). The dashed horizontal line indicates the sill depth of the Bering Strait (∼50 m). Summer insolation at 65° N (July–August) was calculated after Laskar et al. (2004). Records of diatom-bound nitrogen isotope ratios (δ\(^{15}\)N\(_{db}\)), assumed to reflect changes in nitrate utilization, are shown for Bowers Ridge core 17JPC (Brunelle et al., 2007). Comparison of these records suggests that marine productivity and terrigenous matter supply are subject to external forcing by summer insolation and sea-level changes.

Site 85KL, and ∼70% at Site 101KL. At the same time, K/Ti ratios decrease to ∼5 mol mol\(^{-1}\). Minor drops were recorded during MIS 5.3 and 5.1 (∼15% in siliciclastics; ∼0.5 mol mol\(^{-1}\) in K/Ti), whereas K/Ti ratios increased by ∼1 mol mol\(^{-1}\) during MIS 4 and subsequently decreased during MIS 3. In core 85KL MIS 6 is characterized by short-lived maxima in XRF K/Ti log-ratios at ∼173 and ∼163 ka BP. This core and core 101KL also reveal a minor peak in >63 μm found at ∼133 ka BP, which occurs simultaneously with minima in %Siliciclastics but with maxima in CaCO\(_3\) and TOC. Notably, highest amounts of >63 μm were recorded at sites 85KL and 101KL at the end of Termination II (∼127–125 ka BP) with maxima of ∼30% and ∼60%, respectively. These maxima are concurrent with high [C/N]\(_a\) ratios and lead maxima in TOC and opal concentrations by ∼3 kyr. During Termination I, the B/A is characterized by reductions in %Siliciclastics and %Terrigen of up to ∼25% and of up to ∼0.5 mol mol\(^{-1}\) in K/Ti. Maxima
in > 63 µm were recorded during both, the B/A and the PB. During HS1 maxima in K/Ti are observed, and during the YD all proxies briefly return to glacial values, followed by a gradual decrease into the Holocene. The described proxy ranges and their temporal variability almost compare to those reported for the central Okhotsk Sea (Nürnberg and Tiedemann, 2004), which showed lower interglacial minima in %Siliciclastics of ∼35%.

In all cores from Shirshov Ridge linear relationships between concentrations of lithogenous elements Al, Fe, and Ti were found (Fig. 7). This indicates that these elements originate from the same geochemical source and/or share the same transport mechanism. [Al] correlated with %Siliciclastics in cores 77KL ($R^2 = 0.42$) and 85KL ($R^2 = 0.74$), but not in core 101KL ($R^2 = 0.08$). We consider an influence of scavenging by organic material and/or bottom sediment resuspension on [Al] (Orlans and Bruuland, 1986; Nameroff et al., 2004) insignificant, which is supported by low interglacial concentrations and Al/Ti ratios that are close to crustal values. Records for [Al], [Fe], and [Ti] follow the same temporal evolution described for the terrigenous proxies with general high values during most of the last 180 kyr ($∼2600–2800 \mu\text{mol g}^{-1}$ for [Al], $∼700–900 \mu\text{mol g}^{-1}$ for [Fe], $∼90–100 \mu\text{mol g}^{-1}$ for [Ti]) (Table 3). During MIS 5.5 and the Holocene [Al], [Fe], and [Ti] decreased by up to $∼800 \mu\text{mol g}^{-1}$, $∼300 \mu\text{mol g}^{-1}$, and $∼30 \mu\text{mol g}^{-1}$, respectively. Notably, from MIS 3 to 1 [Fe] became slightly higher in core 77KL than in the other cores (Fig. 7). The described temporal evolution overall compares to that recorded in sediment cores from the central Okhotsk Sea (Sato et al., 2002) and the Antarctic Zone of the Southern Ocean (Lattimer and Filippelli, 2001), except that the respective studies reported on glacial [Al] and [Ti] that were by ∼30% lower than those recorded at Shirshov Ridge. Moreover, these studies described very low interglacial values (∼100 µmol g⁻¹ for [Al], ∼100 µmol g⁻¹ for [Fe], ∼10–20 µmol g⁻¹ for [Ti]) not found at our sites. This argues that the interglacial reduction in terrigenous fluxes at Shirshov Ridge was on the order of ∼30–40%.

4.2.2 Geochemical signature
Molar Al/Ti and Fe/Al ratios were used to characterize the geochemical signature of the sediments from Shirshov Ridge. In general, Al/Ti and Fe/Al ratios remained fairly constant during the last 180 kyr with similar ranges at all sites. Only at the end of both glacial terminations minor increases in Al/Ti and minor decreases in Fe/Al are found. Distinct glacial-interglacial differences are not observed, indicating that the source of the terrigenous matter or its underlying transport mechanism did not change. This is an interesting result, since records of %Siliciclastics, %Terrigen, and lithogenous element concentrations are characterized by strong variations on the glacial-interglacial level. Al/Ti ratios varied between 22–34 (mol mol⁻¹), with short-lived increases (∼3) at the end of MIS 4, and during Terminations I and II. Average downcore Al/Ti values (∼29.5 ± 3; Table 4), as well as surface sediment values from Shirshov Ridge (29.4 ± 4) and from the eastern Kamchatka continental margin (32.1 ± 3) (unpublished data) compare well, which indicates that past and modern sources of terrigenous matter are identical. These values are close to Al/Ti ratios reported for Paleozoic (∼29) and Mesozoic/Cenozoic shales (∼32) from the Russian Platform (Ronov and Migidisov, 1971), but also compare with average values for sediment and continental crust (∼28; McLennan, 1995; Taylor and McLennan, 1995), river particulate and mud (∼30; McLennan, 1995), pelagic clay (∼32; McLennan, 1995), as well as the range reported for loess deposits (∼26–31; Taylor et al., 1983; Pye, 1987; McLennan, 1995). Values are clearly lower than that of the North American shale composite (∼38; Gromet et al., 1984), but higher than Al/Ti ratios of oceanic tholeiitic basalt (∼17; Engel et al., 1965) and surface samples from St. George Basin, SE Bering Sea (∼24; Gardner et al., 1980). Nürnberg and Tiedemann (2004) reported a different range for sediments from the central Okhotsk Sea (∼24–45).

The downcore Fe/Al ratios varied between 0.17–0.36 (mol mol⁻¹) (Table 4). This range is also found in surface sediment samples from Shirshov Ridge (0.22–0.39; unpublished data), and it is in agreement with downcore results from the central Okhotsk Sea (0.28 ± 0.03; Nürnberg and Tiedemann, 2004). Our Fe/Al ratios compare to those reported for the North American shale composite and average mud (∼0.24; Gromet et al., 1984; McLennan, 1995), Mesozoic/Cenozoic (∼0.26) and Paleozoic (∼0.28) Russian Platform shales (Ronov and Migidisov, 1971), average values for river particulate (∼0.25) and sediment (∼0.27) (McLennan, 1995). Oceanic tholeiitic basalt (∼0.28; Engel et al., 1965) and surface samples from St. George Basin (∼0.28; Gardner et al., 1980) also apply to the overall range. The value for bulk continental crust is higher (∼0.41; Taylor and McLennan, 1995), while loess deposits show a lower range between ∼0.17–0.25 (Taylor et al., 1983; Pye, 1987; McLennan, 1995). From these results we conclude that sediments from Shirshov Ridge represent a mixture of geochemical signatures from aeolian sediments (loess) and continental sources that are supposedly not influenced by North American shales.

| Table 4. Ranges, averages and variability of molar elemental ratios in cores SO201-2-77KL, -85KL, and -101KL. |
|------------------|------------------|------------------|------------------|
|                  | SO201-2-77KL     | SO201-2-85KL     | SO201-2-101KL    |
| Al/Ti            | Average          | 29.9             | 30.0             | 29.5             |
|                  | StDev.           | 1.5              | 0.03             | 0.26             |
|                  | Maximum          | 34.1             | 32.3             | 32.4             |
|                  | Minimum          | 24.9             | 22.1             | 27.8             |
| Fe/Al            |                  |                  |                  |
|                  |                  |                  |                  |
4.2.3 Transport mechanism and source area

Possible mechanisms of terrigenous matter transport encompass aeolian and fluvial supply, as well as sea-ice rafting. Wind transported aerosols present in N Pacific sediments are restricted to the vicinity of their respective source areas, which in our case are most likely situated in NE Siberia. Model results of global desert dust deposition show fluxes of 0.5–1.0 g m⁻² yr⁻¹ for the Bering Sea (Mahowald et al., 2005), which implies that today aeolian input is negligible. This seems to have been different during past cold intervals as indicated by our higher lithogenous element concentration and generally enhanced glacial dust fluxes (e.g. Ruth et al., 2003). Major rivers entering the Bering Sea are the Yukon and Anadyr rivers, which are situated at rather long distances from Shirshov Ridge (cf. Fig. 1). Today the Yukon River provides ~63% of the total sediment load to the Bering Sea and it is suggested that glacial sediments from the Meiji Drift, NW Pacific, contain a larger fraction of terrigenous material delivered from Yukon–Bering Sea sources (VanLaningham et al., 2009). Surface sediments from the eastern Bering Sea shelf consist of Yukon River-derived detrital material (Asahara et al., 2012; Nagashima et al., 2012). Yet it remains unclear how the terrigenous material is transported from the eastern Bering Sea shelf to the Meiji Drift in the NW Pacific. We suggest that sea ice plays a considerable role in transporting terrigenous matter toward Shirshov Ridge, although we can not exclude a significant contribution from suspension load carried by the BSC. For the Arctic, Nürnberg et al. (1994) reported on sediments entrained in sea ice to be generally fine grained (clayey silts, silty clays) and mainly composed of quartz, clay minerals and diatom flora. Sediments from the central Okhotsk Sea, which are also assumed to originate from sea-ice rafting are described as mainly clay and silt-sized siliciclastics (> 65% siliciclastics) featuring dropstones (3–5 cm), and various lithogenic components (mainly quartz, rock fragments, mica, and dark minerals), with regionally different ice-rafted debris (IRD) composition (Nürnberg and Tiedemann, 2004; Nürnberg et al., 2011). This sedimentary composition is similar to that found in sediments recovered from Shirshov Ridge, although IRD records are needed for verification. Since dropstones are rather uncommon in our cores, which, if present, appear as well-rounded pebbles, and due to the dominance of silt- and clay-sized terrigenous material we suggest a beach deposit origin and tidal pumping, suspension freezing and beach-ice formation to be responsible for the entrainment of terrigenous matter into newly formed sea ice.

The influence of ice in the Bering Sea realm during glacials, either in the form of icebergs or sea ice is still debated. Kaufman et al. (1996) suggested that the most recent major ice advance in SW Alaska occurred between ~90–75 ka BP. For MIS 3 Bigg et al. (2008) proposed a Kamchatka–Koryak Ice Sheet with marine-terminating ice margins. Our cores lack clear evidence for iceberg discharge at ~40 kaBP (Bigg et al., 2008), therefore not supporting their suggested iceberg migration paths. For the LGM, the presence of the Beringian Ice Sheet (e.g. Grosswald and Hughes, 2002) has been disproved (e.g. Brigham-Grette et al., 2001, 2003; Karhu et al., 2001), and Kamchatkan climate seems to have been too arid for the development of large continental ice sheets during that time (Barr and Clark, 2011). We therefore favor sea ice over iceberg transport.

Lisitzin (2002) identified mineralogical provinces for western Bering Sea surface sediments together with their possible migration paths and proposed that coarse silts and the larger grain size fractions (pebbles, gravel, boulder) are controlled by sea ice, sharing the same provinces and transport pathways. Accordingly, the Koryak Coast, Olyutorskii Bay, and northern Kamchatka provinces (cf. Fig. 1) are potential source areas for the ice-rafted material. Anadyr Bay, where modern seasonal sea-ice formation begins during fall, is as well taken into consideration. The K/Ti ratio is assumed to reflect temporal changes in sediment sources weathered from acidic (more K) and basaltic (more Ti) source rocks (Richter et al., 2006). Relatively increased K/Ti ratios are observed at times of enhanced terrigenous matter supply. If sea-ice rafting is the main driver of these changes, the geochemical source of the terrigenous matter (and consequently the ice-rafted material) would be characterized as being relatively increased in K. Of the considered provinces only Anadyr Bay is reported to contain acidic index rocks (Lisitzin, 2002), although geochemical data to clearly identify it as the proposed source area of Shirshov Ridge sediments do not exist. That the source area supposedly lies in the vicinity of Anadyr Bay is supported by the finding that interannual variability in Bering Sea ice cover is controlled by wind-driven ice mass advection, which is clearly south-westward from there (Zhang et al., 2010). However, although stronger ocean/bottom currents can not be excluded as potential transport mechanism for the detrital fractions, the available data seem to congruently argue for a source area that is located on the eastern Bering Sea shelf (VanLaningham et al., 2009; Asahara et al., 2012; Nagashima et al., 2012).

4.3 Environmental changes during the last 180 kyr

4.3.1 Glacial situation

Cold intervals of the last 180 kyr seem to have been subject to strongly reduced, but maintained export production at our sites. At the same time, our records indicate high input of terrigenous matter, which is in agreement with records from the Okhotsk Sea (Nürnberg and Tiedemann, 2004; Nürnberg et al., 2011). Although regionally different, low glacial export production characterized the whole N Pacific during the last 800 kyr (for a review see Kienast et al., 2004). Nürnberg et al. (2011) argued for extreme glacial ice conditions in the Okhotsk Sea during late MIS 6 with a potentially perennial ice coverage. From our results, we can not
unambiguously argue for similar conditions during late MIS 6, although the coarse fraction maxima found during Termination II might indicate the sudden melt of an expanded sea-ice cover that was perennial during the final phase of MIS 6. Glacial conditions in the Okhotsk Sea are supposed to be similar to the modern winter situation of a strong Siberian High and a weak Aleutian Low, thereby resulting in strong offshore, northerly winds (Nürnberg and Tiedemann, 2004; Nürnberg et al., 2011). This situation might as well apply to the Bering Sea and would have resulted in enhanced sea-ice formation during a generally colder climate, thereby explaining the higher LSR and terrigenous inputs. At Bowers Ridge, at least for MIS 3 and 2, colder subsurface temperatures are supported by low total numbers of planktonic foraminifera, dominated by polar species Neogloboquadrina pachyderma (sin.) and reduced/absent numbers of subpolar species Globigerina bulloides (Gorbarenko et al., 2005). A spatially extended sea-ice coverage fostered by a shortened summer season might have resulted in enhanced upper-ocean stratification and reduced nutrient supply to the euphotic zone. Since primary productivity is limited by temperature and nutrient availability, the prolonged sea-ice season during colder intervals in hand with stronger stratification would have consequently led to the reduction in export production observed at our sites. This should be reflected by more efficient nitrate utilization, and indeed, bulk sediment and diatom-bound nitrogen isotope ratios (δ^{15}N) recorded at Bowers Ridge (Brunelle et al., 2007) show higher glacial than interglacial values (Fig. 9), thereby confirming earlier assumptions of enhanced stratification in the glacial Bering Sea (Nakatsuka et al., 1995; Brunelle et al., 2007; Kim et al., 2011).

It has been speculated that the net inflow of Alaskan Stream waters into the Bering Sea was reduced at times when the Bering Strait and/or other Aleutian passes, like Unimak Pass, were closed due to lower glacial sea level, thereby affecting Beringian climate (Pushkar et al., 1999; Tanaka and Takahashi, 2005). During MIS 2, a strengthened Subarctic Front could have additionally led to a reduced inflow of warmer and nutrient-enriched Pacific surface waters (Gorbarenko et al., 2005). In consequence, nutrient supply to the Bering Sea should have been further reduced and have resulted in stronger nitrate utilization. The relative sea level (RSL) reconstruction of Waebbroeck et al. (2002) implies that the Bering Strait was closed during MIS 6, as well as in between MIS 4 and 2 (Fig. 9), while its last major re-opening occurred at 12–11 ka BP (Keigwin et al., 2006). RSL changes in the Bering Strait are predominantly controlled by eustatic changes and suggested to considerably influence deep convection in the N Atlantic (Hu et al., 2010). At Bowers Ridge δ^{15}N values are highest during MIS 3 and 2 (Fig. 9), supporting an influence of RSL on upper-ocean stratification and, hence, nitrate utilization in the Bering Sea. A closed Bering Strait has previously been suggested to have resulted in a low-salinity surface layer that reinforced vertical stratification in the Bering Sea (Sancetta, 1983).

The low glacial CaCO3 concentrations and abundances of oxic benthic foraminifera species point toward the presence of corrosive bottom waters as a result of organic matter degradation under oxic bottom water conditions (Kim et al., 2011). This implies that either O2-rich intermediate water masses were formed in the Bering Sea, or flew in from the Pacific side. Sea-ice formation in the Bering Sea due to brine rejection results in denser, O2-rich surface waters (e.g. Niebauer et al., 1999), and thus might have maintained the production and ventilation of intermediate water. Rella et al. (2012) reported on benthic δ^{18}O excursions in sediments from the northern slope, which implies that the Bering Sea was a source of intermediate water during past stadial episodes, which is supported by microfossil assemblages (Okushi et al., 2003). However, as the modern origin of NPIW lies in the Okhotsk Sea (Yasuda, 1997) and during glacial stages a closed Bering Strait prevented inflow of surface waters from the Arctic Ocean (Takahashi, 1998, 1999), inflow from the N Pacific can not be ruled out.

4.3.2 Deglacial situation

During Termination I high export production but low terrigenous input is observed during the B/A and PB warm phases, whereas the opposite situation occurred during the HS1 and YD cold phases. Remarkably, SST records from our sites mirror the deglacial SST evolution recorded in the N Atlantic, supporting quasi-synchronicity of Northern Hemisphere climate changes (Max et al., 2012). Our records suggest that the early deglacial phase was characterized by higher input of terrestrial-derived organic matter from flooded shelf areas as previously suggested for the Okhotsk and Bering seas (Ternois et al., 2001; Seki et al., 2003; Khim et al., 2010). Alternatively, organic matter preservation became better due to poorly oxygenated conditions at the sediment–water interface. Notably, at the northern slope similar changes were reported to have occurred ~ 2 kyr earlier (Khim et al., 2010). This time lag for increased terrestrial organic carbon input might indicate that the northern slope and Shirshov Ridge were supplied from different sources or by different mechanisms.

Our records indicate enhanced sea-ice influence during HS1 and the YD, which for our sites is supported by qualitative reconstructions of the sea-ice-related IP25 biomarker (Max et al., 2012) and by increasing abundances of the sea-ice-related diatom genus Nitzschia at Unmak Plateau (Cook et al., 2005). Like during previous cold intervals, marine productivity was restricted by lowered temperature. A sudden decrease in bulk sedimentary and diatom-bound δ^{15}N (Fig. 9) point to decreased nitrate utilization (Brunelle et al., 2007, 2010). However, this drop in δ^{15}N is not fully understood since higher nutrient supply due to stronger vertical mixing should have resulted in enhanced export
production, which is not observed. Deglacial changes in ventilation/overturning in the N Pacific realm have been inferred from ventilation ages (Adkins and Boyle, 1997; Ahagon et al., 2003; Ohkushi et al., 2004; Galbraith et al., 2007; Sarnthein et al., 2007; Sagawa and Ikehara, 2008; Okazaki et al., 2010; Lund et al., 2011; Jaccard and Galbraith, 2013), foraminiferal isotope data (e.g. Matsumoto et al., 2002; Rella et al., 2012), as well as modeling studies (e.g. Okazaki et al., 2010; Chikamoto et al., 2012; Menivel et al., 2012). Notably, enhanced ventilation in the N Pacific during HS1 seems to have been restricted to ∼1400–2400 m depth (Jaccard and Galbraith, 2013), whereas the deep abyssal Pacific was better ventilated during the subsequent B/A (Galbraith et al., 2007).

Hence, it was suggested that NPIW formation was more enhanced, that it reached deeper, and that its source might have shifted to the Bering Sea during HS1 and during other cold periods (e.g. Matsumoto et al., 2002; Ohkushi et al., 2003; Rella et al., 2012; Jaccard and Galbraith, 2013).

For the onset of the B/A, and to a lesser degree also the PB phase, our records imply enhanced marine productivity and the sudden release of IRD from melting sea ice. Similar observations regarding changes in marine productivity have previously been reported for the whole Bering Sea realm (Gorbarenko, 1996; Cook et al., 2005; Gorbarenko et al., 2005; Okazaki et al., 2005; Khim et al., 2010) and other parts of the N Pacific (Keigwin and Jones, 1990; Keigwin et al., 1992; Gorbarenko, 1996; Keigwin, 1998; Crusius et al., 2004; McKay et al., 2004; Okazaki et al., 2005; Gebhardt et al., 2008). Increasing insolation should have amplified the surface ocean warming and led to more dynamic ice conditions, northward propagating ice margins, and a prolonged sea-ice-free summer season. Indeed, recent alkenone- and Mg/Ca-based studies indicate rising SST, strengthened thermal mixed layer stratification (stronger seasonal contrasts), and the onset of coccolithophorid production during the B/A (Caisse et al., 2010; Max et al., 2012; Riethdorf et al., 2013).

Enhanced surface freshening due to melting sea ice is supported by higher abundances of radiolarian species *Paralia sulcata* (Gorbarenko et al., 2005). A maximum in δ15N values (Fig. 9) implies increased nitrate utilization or even denitrification (Brunelle et al., 2007, 2010; Khim et al., 2010). It might further be speculated, that the increase in marine productivity associated with reduced sea-ice formation resulted in organic matter supply exceeding its degradation at the seafloor (Kim et al., 2011). In consequence, bottom water conditions might have become dysoxic or anoxic, impeding benthonic life and favoring laminated formation. This notion would be in agreement with the formation of dysoxic or laminated sediments observed at oxygen minimum zone (OMZ) depths in the N Pacific and Bering Sea during the B/A and PB (van Geen et al., 2003; Cook, 2006, and references therein) and most probably was related to an intensification of the OMZ (Zheng et al., 2000). At the same time the respired carbon pool was obviously removed from the deep basin, which resulted in enhanced carbonate preservation due to a deepened lysocline, and to an increase in atmospheric CO2 (e.g. Galbraith et al., 2007).

Our records show that at sites 85KL and 101KL the deglacial development during Termination II does not resemble that of Termination I. Most notably, extremely high amounts of coarse material are found at the end of Termination II showing values that are not reached previously or later at any site. At the same time minor excursions toward lower Fe/Al but higher Al/Ti ratios were recorded, indicating a slightly different geochemical composition of the sediments. We attribute these results to the sudden release of IRD in response to strong melting of a potentially perennial ice coverage present during the final phase of MIS 6 in agreement with conditions proposed for the Okhotsk Sea (Nürnberg et al., 2011). Also, records of benthic δ18O do not follow the LR04 stack, which is indicative of a strong regional effect on the δ18O signal, for example by melting of continental ice sheets in the Bering Sea realm. This might, of course, also be the result of constraints in our age models during that time frame.

### 4.3.3 Interglacial situation

In contrast to the situation during cold intervals, interglacials, as well as warm MIS 5.3 and 5.1 were characterized by increased marine productivity and decreased terrigenous matter supply. Our records for Site 77KL better compare to those reported for Bowers Ridge, whereas records from sites 85KL and 101KL resemble those from the northern slope and the central Okhotsk Sea. Marine productivity as reflected by opal concentrations seems to have been higher during MIS 5.5 than during the Holocene. Interglacial CaCO3 concentrations remained at or close to glacial values, implying a similar bottom water calcite saturation state with limited carbonate preservation. The interglacial increase in marine productivity most probably resulted from warmer temperatures and reduced sea-ice formation during a prolonged summer season (stronger seasonal contrasts). However, our records imply a reduction of only ∼30–40% in terrigenous matter supply. Overall, records of XRF Si/Al log-ratios are in phase with mean summer insolation at 65° N (July–September; calculated after Laskar et al., 2004; Fig. 9), showing maxima during the warm intervals (MIS 5.5, 5.3, 5.1, and 1). This implies a dominant insolation forcing for environmental changes in the Bering Sea. At Bowers Ridge, δ15N values are lowest during interglacials (Brunelle et al., 2007, 2010) and they also remain low during most of MIS 5 (Fig. 9). This indicates reduced nitrate utilization as a consequence of enhanced vertical mixing, which supplies sufficient amounts of nitrate from the subsurface nitrate pool into the euphotic zone. This situation might have been additionally influenced by an open Bering Strait and Unimak Pass, allowing for enhanced inflow of water masses from the Alaskan Stream and enhanced outflow of relatively fresh surface waters into...
the Arctic. Reduced sea-ice formation, but enhanced nutrient supply by stronger vertical mixing as a result of a strengthened Alaskan Stream and BSC during higher sea level has been proposed before (Gorbarenko et al., 2005; Okazaki et al., 2005; Kim et al., 2011). Accordingly, sea-level changes must be considered as additional forcing mechanism to explain the recorded environmental changes.

4.3.4 Interstadial situation

High-resolution core logging resulted in the detection of short-lived maxima in our color b* and XRF Ca/Ti log-ratio records during MIS 6 to 3. Notably, these maxima and maxima in >63 μm occur simultaneously. Most peaks were detected in cores 85KL and 101KL within which they are characterized by only 1–3 cm sediment thickness corresponding to a duration of ~100–300 yr. These changes in sediment composition seem to be related to D-O events (interstadials) registered in the NGRIP ice core. The most prominent peaks correspond to D-O events 7, 8, 12, 17–21, and 24 (Fig. 5), but events off the NGRIP record are found. We consider them as brief intervals of enhanced marine productivity, sudden sea-ice melt associated with the subsequent release of IRD, and a higher bottom water calcite saturation state. However, due to our stratigraphic approach, which relies on the NGRIP record, we can neither argue for nor against an in-phase evolution between abrupt climate changes recorded in Greenland ice and western Bering Sea sediments. Sediment records from Bowers Ridge (Gorbarenko et al., 2005, 2010), the northern slope (Kim et al., 2011; Rella et al., 2012), as well as the NE Pacific (Behl and Kennett, 1996; Hendy and Kennett, 2000; Ortiz et al., 2004) also implied millennial-scale climate changes connected to N Atlantic D-O events, especially during MIS 3. Similar features were detected in stalagmites from China (Wang et al., 2001, 2008), which indicates a Northern Hemisphere-wide acting atmospheric coupling that is related to the intensity of the East Asian monsoon. Kennett and Ingram (1995) proposed that such an atmospheric coupling mechanism directly affected the ventilation of NPIW. Accordingly, interstadials seem to have been characterized by weak ventilation of NPIW in combination with increased marine productivity (Behl and Kennett, 1996; Hendy and Kennett, 2000; Kim et al., 2011) and warmer SSTs (Gorbarenko et al., 2005). Recently, Kim et al. (2011) reported on D-O event-related brief episodes of high bulk δ15N values at the northern slope, implying increased nitrate utilization/denitrification. These results to a lesser degree reflect conditions inferred for the B/A warm phase and thus might point to the release of deep-sequestered CO2 during interstadials.

5 Summary and conclusions

From our results we propose scenarios for environmental change in the Bering Sea during glacial, deglacial, and interglacial times which compare to those previously suggested for the Okhotsk Sea. During the last 180 kyr, the Bering Sea paleoenvironment was characterized by closely interacting processes controlling marine productivity and terrigenous matter supply. External forcing is attributed to Northern Hemisphere summer insolation and sea-level changes controlling atmospheric circulation patterns, sea-ice dynamics and upper-ocean stratification. Marine productivity, dominated by diatoms, remained low during cold intervals (MIS 6, 5.4, 5.2, 4 to 2) when the Bering Strait was closed and summer insolation was weak. Increases occurred during warm intervals (MIS 5.5, 5.3, 5.1, and 1), when insolation was high and the Bering Strait was open. Sediments from Shirshov Ridge are dominated by terrigenous, siliciclastic material mainly bound to the fine fractions. Terrigenous matter supply was generally high, with reductions of up to ~30–40% during interglacial. Changes in terrigenous matter supply and marine productivity occurred synchronously with negatively correlated proxy behavior due to the sedimentary dilution by the biogenic components. Terrigenous input was supposedly stronger toward the northernmost site, and sea ice is considered as the transport agent for terrigenous matter, limiting marine productivity during cold intervals. Sedimentary geochemical signatures are a mixture of aeolian and continental sources, indicating that Shirshov Ridge sediments most likely originate from material entrained on the eastern Bering Sea shelf. Especially for the last glacial termination our records support the notion of an atmospheric, Northern Hemisphere-wide acting climate coupling. The situation during the HSI and YD cold phases compared to that of cold intervals with enhanced sea-ice rafting limiting marine productivity during a shortened summer season (weak seasonal contrasts). In contrast, the B/A and PB warm phases were characterized by enhanced marine productivity as a result of a prolonged summer season and reduced sea-ice influence (strong seasonal contrasts). At the same time, reduced ventilation of intermediate waters is in accordance with stronger nitrate utilization and better CaCO3 preservation indicative of a release of deep-sequestered CO2 to the atmosphere. Moreover, we found supporting evidence for the occurrence of abrupt environmental changes that are related to interstadials recorded in the NGRIP ice core.
Table A1. Age-depth points for core SO201-2-77KL. Ages < 20 ka BP and AMS-¹⁴C ages have been derived from Max et al. (2012) and calendar ages are given with 1σ-ranges (in ka BP).

<table>
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<tr>
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<th>Depth (cm)</th>
<th>Cal. Age (ka BP)</th>
<th>Approach</th>
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</thead>
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<td>2.0⁽¹⁾</td>
<td>color b* vs. color b* (SO201-2-12KL)</td>
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⁽¹⁾ Uncertain age; ⁽²⁾ Used to define carbonate spike 1 (see Max et al., 2012); ⁽³⁾ Used to define carbonate spike 2 (see Max et al., 2012).

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Appendix A

Age models

Age-depth points for cores SO201-2-77KL, -85KL, and -101KL are given in Tables A1, A2, and A3, respectively.

Appendix B

Surface sediment samples

Calculated results of Ba₉₀bio, P₉₀New, and PP from discrete XRF measurements conducted on surface sediment samples recovered in direct vicinity of the studied sediment cores using a MultiCorer (MUC) are given in Table B1.
Table A2. Age-depth points for core SO201-2-85KL. Ages < 20 ka BP and AMS-^{14}C ages have been derived from Max et al. (2012) and calendar ages are given with 1σ-ranges (in ka BP).

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<th>Approach</th>
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<td>45.5</td>
<td>–</td>
<td>AMS-^{14}C dating (10.79–10.97)(^{(3)})</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>52</td>
<td>11.6</td>
<td>color b* vs. NGRIP</td>
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<td>13.10</td>
<td>AMS-^{14}C dating (13.10–13.22)</td>
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<td>SO201-2-85KL</td>
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<td>13.87</td>
<td>AMS-^{14}C dating (13.67–13.89)</td>
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<td>SO201-2-85KL</td>
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<td>14.6</td>
<td>color b* vs. NGRIP</td>
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<td>SO201-2-85KL</td>
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<td>14.9</td>
<td>color b* vs. NGRIP</td>
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<td>SO201-2-85KL</td>
<td>95.5</td>
<td>15.84</td>
<td>AMS-^{14}C dating (15.80–15.82)</td>
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<tr>
<td>SO201-2-85KL</td>
<td>135.5</td>
<td>19.90</td>
<td>AMS-^{14}C dating (19.58–19.90)</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>135.5</td>
<td>23.78</td>
<td>AMS-^{14}C dating (23.71–24.19)</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>209</td>
<td>26.2</td>
<td>Transfer of AMS-^{14}C age from core 101KL, but not well constrained</td>
</tr>
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<td>SO201-2-85KL</td>
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<td>SO201-2-85KL</td>
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<td>color b* vs. NGRIP</td>
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<td>305</td>
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<td>color b* vs. NGRIP</td>
</tr>
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<td>SO201-2-85KL</td>
<td>350</td>
<td>41.9</td>
<td>RPI vs. PISO-1500 (Laschamp)</td>
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<td>SO201-2-85KL</td>
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<td>43.4</td>
<td>color b* vs. NGRIP</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>398</td>
<td>46.9</td>
<td>color b* vs. NGRIP</td>
</tr>
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<td>SO201-2-85KL</td>
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<td>48.4</td>
<td>color b* vs. NGRIP</td>
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<td>SO201-2-85KL</td>
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<td>51.7</td>
<td>color b* vs. NGRIP</td>
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<tr>
<td>SO201-2-85KL</td>
<td>490</td>
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<td>color b* vs. NGRIP</td>
</tr>
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<td>SO201-2-85KL</td>
<td>567</td>
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<td>color b* vs. NGRIP</td>
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<td>SO201-2-85KL</td>
<td>657</td>
<td>64.0</td>
<td>color b* vs. NGRIP</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>675</td>
<td>65.0</td>
<td>RPI vs. PISO-1500 (Norwegian–Greenland Sea)</td>
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<tr>
<td>SO201-2-85KL</td>
<td>796</td>
<td>72.3</td>
<td>color b* vs. NGRIP</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>853</td>
<td>76.4</td>
<td>color b* vs. NGRIP</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>876</td>
<td>77.8</td>
<td>color b* vs. NGRIP</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>937</td>
<td>84.7</td>
<td>color b* vs. NGRIP</td>
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<td>SO201-2-85KL</td>
<td>976</td>
<td>87.7</td>
<td>color b* vs. NGRIP</td>
</tr>
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<td>SO201-2-85KL</td>
<td>1006</td>
<td>91.7</td>
<td>color b* vs. NGRIP</td>
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<td>SO201-2-85KL</td>
<td>1100</td>
<td>104.1</td>
<td>color b* vs. NGRIP</td>
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<td>SO201-2-85KL</td>
<td>1149</td>
<td>108.0</td>
<td>color b* vs. NGRIP</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>1180</td>
<td>110.2</td>
<td>color b* vs. NGRIP</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>1210</td>
<td>113.3</td>
<td>color b* vs. NGRIP</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>1224</td>
<td>114.8</td>
<td>color b* vs. NGRIP</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>1240</td>
<td>117.0</td>
<td>RPI vs. PISO-1500 (Blake)</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>1280</td>
<td>125.0</td>
<td>color b* vs. Sanbao δ^{18}O (MIS 5.5 climate optimum)</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>1300</td>
<td>130.0(^{(4)})</td>
<td>SMP vs. LR04 (MIS 5/6 Boundary)</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>1320</td>
<td>132.0</td>
<td>color b* vs. Sanbao δ^{18}O (HE11)</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>1365</td>
<td>135.0</td>
<td>color b*, Ca/Ti vs. Sanbao δ^{18}O</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>1530</td>
<td>151.0</td>
<td>color b*, Ca/Ti vs. Sanbao δ^{18}O</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>1600</td>
<td>159.0</td>
<td>RPI vs. PISO-1500</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>1735</td>
<td>174.0(^{(4)})</td>
<td>SMP vs. LR04 (MIS 6 Minimum)</td>
</tr>
<tr>
<td>SO201-2-85KL</td>
<td>1813</td>
<td>181.8(^{(5)})</td>
<td>Extrapolation</td>
</tr>
</tbody>
</table>

\(^{(1)}\) Uncertain age; \(^{(2)}\) Used to define carbonate spike 1 (see Max et al., 2012); \(^{(3)}\) Used to define carbonate spike 2 (see Max et al., 2012); \(^{(4)}\) Scalar magnetic properties (SMP) correlated with MIS boundaries of LR04; \(^{(5)}\) By extrapolation using linear sedimentation rate (LSR).
Table A3. Age-depth points for core SO201-2-101KL. Ages < 20 ka BP and AMS-\(^{14}\)C ages have been derived from Max et al. (2012) and calendar ages are given with 1\(\sigma\) -ranges (in ka BP).

<table>
<thead>
<tr>
<th>Core Depth (cm)</th>
<th>Cal. Age (ka BP)</th>
<th>Approach</th>
</tr>
</thead>
<tbody>
<tr>
<td>SO201-2-101KL</td>
<td>4</td>
<td>12.9 Ca/Ti vs. Ca/Ti (85KL)</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>10.5</td>
<td>13.56 AMS-(^{14})C dating (13.69–13.84)</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>41</td>
<td>14.6 color b* vs. NGRIP</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>67</td>
<td>15.4 color b* vs. NGRIP</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>90.5</td>
<td>17.25 AMS-(^{14})C dating (17.17–17.51)</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>110.5</td>
<td>18.95 AMS-(^{14})C dating (19.54–19.92)</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>140</td>
<td>23.78 Transfer of AMS-(^{14})C Age from Core 85KL</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>190.5</td>
<td>25.74 AMS-(^{14})C dating (25.88–26.35)</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>234</td>
<td>28.6 color b* vs. NGRIP</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>249</td>
<td>30.3 color b* vs. NGRIP</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>260.5</td>
<td>32.0 AMS-(^{14})C dating (32.12–33.54)</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>274</td>
<td>33.5 Ca/Ti vs. Ca/Ti (85KL)</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>280</td>
<td>35.1 Ca/Ti vs. Ca/Ti (85KL)</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>284</td>
<td>35.7 Ca/Ti vs. Ca/Ti (85KL)</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>302</td>
<td>36.9 Ca/Ti vs. Ca/Ti (85KL)</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>349</td>
<td>51.6 Ca/Ti vs. Ca/Ti (85KL)</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>387</td>
<td>56.6 Ca/Ti vs. Ca/Ti (85KL)</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>454</td>
<td>71.3 Color b* vs. Color b* (85KL)</td>
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<tr>
<td>SO201-2-101KL</td>
<td>514</td>
<td>79.7 Color b* vs. Color b* (85KL)</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>650</td>
<td>86.8 Color b* vs. Color b* (85KL)</td>
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<td>SO201-2-101KL</td>
<td>685</td>
<td>92.4 Color b* vs. Color b* (85KL)</td>
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<td>SO201-2-101KL</td>
<td>723</td>
<td>97.7 Color b* vs. Color b* (85KL)</td>
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<td>SO201-2-101KL</td>
<td>799</td>
<td>100.5 Color b* vs. Color b* (85KL)</td>
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<tr>
<td>SO201-2-101KL</td>
<td>923</td>
<td>103.5 Color b* vs. Color b* (85KL)</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>1526</td>
<td>116.0 Benthic (^{18})O vs. LR04</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>1585</td>
<td>125.0 Color b* vs. Sanbao (^{18})O (MIS 5.5 climate optimum)</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>1625</td>
<td>130.0 Ca/Ti vs. Ca/Ti (85KL)</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>1635</td>
<td>132.0 Color b* vs. Sanbao (^{18})O (HE11)</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>1657</td>
<td>137.0 Ca/Ti vs. Ca/Ti (85KL)</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>1738</td>
<td>141.5 Ca/Ti vs. Ca/Ti (85KL)</td>
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<td>SO201-2-101KL</td>
<td>1765</td>
<td>145.1 Color b* vs. Color b* (85KL)</td>
</tr>
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<td>SO201-2-101KL</td>
<td>1804</td>
<td>147.8 Ca/Ti vs. Ca/Ti (85KL)</td>
</tr>
<tr>
<td>SO201-2-101KL</td>
<td>1832</td>
<td>149.7 Extrapulation</td>
</tr>
</tbody>
</table>

\(^{(1)}\) By extrapolation using linear sedimentation rate (LSR).

Table B1. Site information, concentrations of \(\text{Ba}_{\text{bio}}\), and calculated values for \(P_{\text{New}}\) and PP in surface sediment samples.

<table>
<thead>
<tr>
<th>Station</th>
<th>Latitude</th>
<th>Longitude</th>
<th>Depth (mbsl)</th>
<th>Tube Depth (cm)</th>
<th>(\text{Ba}_{\text{bio}}) (^{(1)}) (ppm)</th>
<th>(P_{\text{New}}) (^{(2)}) (gC m(^{-2}) yr(^{-1}))</th>
<th>PP (^{(3)}) (gC m(^{-2}) yr(^{-1}))</th>
</tr>
</thead>
<tbody>
<tr>
<td>SO201-2-76MUC</td>
<td>56(^{\circ}) 19.80' N</td>
<td>170(^{\circ}) 41.96' E</td>
<td>2137</td>
<td>1–2</td>
<td>932</td>
<td>31.8</td>
<td>113</td>
</tr>
<tr>
<td>SO201-2-76MUC</td>
<td>56(^{\circ}) 19.80' N</td>
<td>170(^{\circ}) 41.96' E</td>
<td>2137</td>
<td>2–3</td>
<td>1327</td>
<td>54.0</td>
<td>147</td>
</tr>
<tr>
<td>SO201-2-76MUC</td>
<td>56(^{\circ}) 19.80' N</td>
<td>170(^{\circ}) 41.96' E</td>
<td>2137</td>
<td>3–4</td>
<td>1319</td>
<td>53.5</td>
<td>146</td>
</tr>
<tr>
<td>SO201-2-83MUC</td>
<td>57(^{\circ}) 30.28' N</td>
<td>170(^{\circ}) 24.82' E</td>
<td>970</td>
<td>0–1</td>
<td>636</td>
<td>19.2</td>
<td>88</td>
</tr>
<tr>
<td>SO201-2-83MUC</td>
<td>57(^{\circ}) 30.28' N</td>
<td>170(^{\circ}) 24.82' E</td>
<td>970</td>
<td>1–2</td>
<td>681</td>
<td>21.3</td>
<td>92</td>
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<tr>
<td>SO201-2-83MUC</td>
<td>58(^{\circ}) 52.53' N</td>
<td>170(^{\circ}) 41.48' E</td>
<td>643</td>
<td>2–3</td>
<td>177</td>
<td>2.9</td>
<td>34</td>
</tr>
</tbody>
</table>

\(^{(1)}\) Via Al using the global average Ba/Al of pelitic rocks of 6.5 mg g\(^{-1}\) (Wedepohl, 1971). \(^{(2)}\) After Nürnberg (1995) using an average AR\(_{\text{Bulk}}\) of 4 g cm\(^{-2}\) kyr\(^{-1}\). \(^{(3)}\) After Eppley and Peterson (1979).
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