Highly branched isoprenoids for Southern Ocean semi-quantitative sea ice reconstructions: a pilot study from the Western Antarctic Peninsula

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Abstract. Organic geochemical and micropaleontological analyses of surface sediments collected in the southern Drake Passage and the Bransfield Strait, Antarctic Peninsula, enable a proxy-based reconstruction of recent sea ice conditions in this climate sensitive area. The distribution of the sea ice biomarker $IPSO_{25}$ supports earlier suggestions that the source diatoms seem to be common in near-coastal environments characterized by an annually recurring sea ice cover. We here propose and evaluate the combination of $IPSO_{25}$ with a more unsaturated highly branched isoprenoid alkene and phytosterols and introduce the $PIPSO_{25}$ index as a potentially semi-quantitative sea ice proxy. This organic geochemical approach is complemented with diatom data. $PIPSO_{25}$ sea ice estimates are used to discriminate between areas characterized by permanently ice-free conditions, seasonal sea ice cover and extended sea ice cover. These trends are consistent with satellite sea ice data and winter sea ice concentrations estimated by diatom transfer functions. Minor offsets between proxy-based and satellite-based sea ice data are attributed to the different time intervals recorded within the sediments and the instrumental records from the study area, which experienced rapid environmental changes during the past 100 years.

Key Words: biomarker, $IPSO_{25}$, sea ice, Bransfield Strait, satellite observation

1   Introduction

In the last century, the Western Antarctic Peninsula (WAP) has undergone a rapid warming of the atmosphere of $3.7 \pm 1°$ C, which exceeds several times the average global warming (Vaughan et al., 2003). Simultaneously, a

1 reduction in sea ice coverage (Parkinson and Cavalieri, 2012), a shortening of the sea ice season (Parkinson, 2002)

2 and a decreasing sea ice extent of ~4-10 % per decade (Liu et al., 2004) are recorded in the adjacent Bellingshausen

3 Sea. The loss of seasonal sea ice and increased melt water fluxes impact the formation of deep and intermediate

4 waters, the ocean-atmosphere-exchange of gases and heat, the primary production and higher trophic levels

5 (Arrigo et al., 1997; Morrison et al., 2015; Orsi et al., 2002; Rintoul, 2007). Since the start of satellite-based sea

6 ice observations, however, a slight increase in total Antarctic sea ice extent has been documented, which contrasts

7 the significant decrease of sea ice in Western Antarctica, especially around the WAP (Hobbs et al., 2016).

8 For an improved understanding of the oceanic and atmospheric feedback mechanisms associated with the observed

9 changes in sea ice coverage, reconstructions of past sea ice conditions in climate sensitive areas such as the WAP

10 are of increasing importance. A common approach for sea ice reconstructions in the Southern Ocean is based on

11 the investigation of sea ice associated diatom assemblages preserved in marine sediments (Bárcena et al., 1998;

12 Gersonde and Zielinski, 2000; Heroy et al., 2008; Leventer, 1998; Minzoni et al., 2015). By means of transfer

13 functions, this approach can provide quantitative estimates of a paleo sea ice coverage (Crosta et al., 1998; Esper

14 and Gersonde, 2014a). The application of diatoms for paleoenvironmental studies, however, can be limited by the

15 selective dissolution of biogenic opal frustules (Burckle and Cooke, 1983; Esper and Gersonde, 2014b) in the

16 photic zone (Ragueneau et al., 2000) and in surface sediments (Leventer, 1998). As an alternative or additional

17 approach to diatom studies, Massé et al. (2011) proposed the use of a specific biomarker lipid – a diunsaturated

18 highly branched isoprenoid alkene (HBI $C_{25:2}$, Fig. 1) – for Southern Ocean sea ice reconstructions. The HBI diene

19 was first described by Nichols et al. (1988) from sea ice diatoms. ^{13}C isotopic analyses of the HBI diene suggest a

20 sea ice origin for this molecule (Sinninghe Damsté et al., 2007; Massé et al., 2011) and this is further corroborated

21 by the identification of the sea ice diatom *Berkeleya adeliensis* as a producer of this HBI diene (Belt et al., 2016).

22 In a survey of surface sediments collected from proximal sites around Antarctica, Belt et al. (2016) note a

23 widespread sedimentary occurrence of the HBI diene and – by analogy with the Arctic HBI monoene termed IP$_{25}$

24 (Belt et al., 2007) – proposed the term IPSO$_{25}$ (Ice Proxy for the Southern Ocean with 25 carbon atoms) as a new

25 name for this biomarker.

26 In previous studies, an HBI triene (HBI $C_{25:3}$; Fig. 1) found in polar and sub-polar phytoplankton samples (Massé

27 et al., 2011) has been considered alongside IPSO$_{25}$ and the ratio of IPSO$_{25}$ to this HBI triene hence has been

28 interpreted as a measure for the relative contribution of organic matter derived from sea ice algae versus open

29 water phytoplankton (Massé et al., 2011; Collins et al., 2013; Etourneau et al., 2013; Barbara et al., 2013, 2016).

30 Collins et al. (2013) further suggested that the HBI triene might reflect phytoplankton productivity in marginal ice

31 zones (MIZ) and, based on the observation of elevated HBI triene concentrations in East Antarctic MIZ surface

waters, this has been strengthened by Smik et al. (2016a). Known source organisms of $C_{25:3}$ HBI trienes (Fig. 1 shows molecular structures of both the E- and Z-isomer) are, for example, *Rhizosolenia* and *Pleurosigma* diatom species (Belt et al., 2000, 2017). In the subpolar North Atlantic, the HBI Z-triene has been used to further modify the so-called PIP_{25} index (Smik et al., 2016b) - an approach for semi-quantitative sea ice estimates. Initially, PIP_{25} was based on the employment of phytoplankton-derived sterols, such as brassicasterol and dinosterol (Kanazawa et al., 1971; Volkman, 2003), to serve as open-water counterparts, while IP_{25} reflects the occurrence of a former sea ice cover (Belt et al., 2007; Müller et al., 2009, 2011). Consideration of these different types of biomarkers helps to discriminate between ice-free and permanently ice-covered ocean conditions, both resulting in a lack of IP_{25} and $IPSO_{25}$, respectively (for further details see Belt, 2018; Belt and Müller, 2013a). Uncertainties in the source-specificity of brassicasterol (Volkman, 1986) and its identification in Arctic sea ice samples (Belt et al., 2013), however, require caution when pairing this sterol with a sea ice biomarker lipid for Arctic sea ice reconstructions. In this context, we note that Belt et al. (2018) reported that brassicasterol is not evident in the $IPSO_{25}$ producing sea ice diatom *Berkeleya adeliensis*. While the applicability of HBIs (and sterols) to reconstruct past sea ice conditions has been thoroughly investigated in the Arctic Ocean (Belt, 2018; Stein et al., 2012; Xiao et al., 2015), only two studies document the distribution of HBIs in Southern Ocean surface sediments (Belt et al., 2016; Massé et al., 2011). The circum-Antarctic data set published by Belt et al. (2016), however, does not report HBI triene abundances. Significantly more studies so far focused on the use of $IPSO_{25}$ and the HBI Z-triene for paleo sea ice reconstructions and these records are commonly compared to micropaleontologial diatom analyses (e.g., Barbara et al., 2013; Collins et al., 2013; Denis et al., 2010).

Here, we provide a first overview of the distribution of $IPSO_{25}$, HBI trienes, brassicasterol and dinosterol in surface sediments from the northern part of the WAP (southern Drake Passage and Bransfield Strait). These biomarker data are completed by diatom analyses and remote sensing sea ice data. We further introduce and discuss the so-called $PIPSO_{25}$ index (phytoplankton-$IPSO_{25}$ index), which, following the PIP_{25} approach in the Arctic Ocean (Müller et al., 2011), may serve as a semi-quantitative indicator of past Southern Ocean sea ice cover. These biomarker-based sea ice estimates are compared to sea ice concentrations derived from diatom transfer functions and satellite-derived data on the recent sea ice conditions in the study area.

2   Oceanographic setting

The study area includes the southern Drake Passage and the Bransfield Strait located between the South Shetland Islands and the northern tip of the WAP (Fig. 2a and b). While the oceanographic setting in the Drake Passage is dominated by the Antarctic Circumpolar Current (ACC), a complex current system prevails in the Bransfield Strait. According to Sangrá et al. (2011) a branch of the ACC enters the Bransfield Strait in the west as the Bransfield Current, carrying transitional waters under the influence of the Bellingshausen Sea (Transitional Bellingshausen Sea Water, TBW). The TBW is characterized by a well-stratified, fresh and warm water mass with summer sea surface temperatures (SST) above 0° C. Below the shallow TBW, a narrow tongue of circumpolar deep water (CDW) flows along the slope of the South Shetland Islands (Sangrà et al., 2011). In the eastern part, transitional water from the Weddell Sea (Transitional Weddell Sea Water, TWW) enters the Bransfield Strait through the Antarctic Sound and from the Antarctic Peninsula (AP). This water mass corresponds to the Antarctic Coastal Current (Collares et al., 2018; Thompson et al., 2009). The TWW is significantly colder (summer SST < 0° C) and saltier due to extended sea ice formation in the Weddell Sea Gyre. The two water masses are separated at the sea surface by the Peninsula Front characterized by a TBW anticyclonic eddy system (Sangrà et al., 2011). While the TWW occupies the deep water column of the Bransfield Strait (Sangrà et al., 2011), it joins the surface TBW in the southwestern Bransfield Strait (Collares et al., 2018).

Due to high concentrations of dissolved iron on the shelf (Klunder et al., 2014), the area around the WAP is characterized by a high primary production with high vertical export fluxes during early summer associated with the formation of fast sinking mineral aggregates and fecal pellets (Kim et al., 2004; Wefer et al., 1988). The Peninsula Front divides the Bransfield Strait into two biogeographic regimes of high chlorophyll and diatom abundance in the TBW and low chlorophyll values and a pre-dominance of nanoplankton in the TWW (Gonçalves-Araujo et al., 2015), which is also reflected in the geochemistry of surface sediments (Cárdenas et al., 2018).

1	**3 Materials and Methods**
2	**3.1 Sediment Samples and radiocarbon dating**
3	In total, 26 surface sediment samples obtained by multicorers and box corers during the RV *Polarstern* cruise
4	PS97 (Lamy, 2016) were analyzed (Fig. 2, Table 1). All samples were stored frozen and in glass vials. The
5	composition of the sediments ranges from foraminiferal mud in the Drake Passage to diatomaceous mud with
6	varying amounts of ice rafted debris in the Bransfield Strait (Lamy, 2016).
7	[14]C radiocarbon dating of two samples from the PS97 cruise and one from the *Polarstern* cruise ANT-VI/2
8	(Fütterer, 1988) was conducted using the mini carbon dating system (MICADAS) at the Alfred Wegener Institute
9	following the method of Wacker et al. (2010). The ^{14}C ages were calibrated to calendar years before present (cal
10	BP) using the Calib 7.1 software (Stuiver et al., 2019) with an estimated reservoir age of 1178 years, derived from
11	the six closest reference points listed in the Marine Reservoir Correction Database (www.calib.org).
12	
13	**3.2 Organic geochemical analyses**
14	For biomarker analyses, sediments were freeze-dried and homogenized using an agate mortar. Also after freeze-
15	drying, samples were stored frozen to avoid degradation. The extraction, purification and quantification of HBIs
16	and sterols follow the analytical protocol applied by the international community of researchers performing HBI
17	and sterol-based sea ice reconstructions (Belt et al., 2014; Stein et al., 2012). Prior to extraction, internal standards
18	7-hexylnonadecane (7-HND) and 5α-androstan-3β-ol were added to the sediments. For the ultrasonic extraction
19	(15 min), a mixture of CH_2Cl_2:MeOH (v/v 2:1; 6ml) was added to the sediment. After centrifugation (2500 rpm
20	for 1 min), the organic solvent layer was decanted. The ultrasonic extraction step was repeated twice. From the
21	combined total organic extract, apolar hydrocarbons were separated via open column chromatography (SiO_2) using
22	hexane (5 ml). Sterols were eluted with ethylacetate:hexane (v/v 20:80; 8 ml). HBIs were analyzed using an
23	Agilent 7890B gas chromatography (30 m DB 1MS column, 0.25 mm diameter, 0.250 μm film thickness, oven
24	temperature 60° C for 3 min, rise to 325° C within 23 min, holding 325° C for 16 min) coupled to an Agilent
25	5977B mass spectrometer (MSD, 70 eV constant ionization potential, ion source temperature 230° C). Sterols
26	were first silylated (200 μl BSTFA; 60° C; 2hours) and then analyzed on the same instrument using a different
27	oven temperature program (60° C for 2 min, rise to 150° C within 6 min, rise to 325° C within 56 min 40 sec). As
28	recommended by Belt (2018), the identification of $IPSO_{25}$ and HBI trienes is based on comparison of their mass
29	spectra with published mass spectra (Belt et al., 2000). Regarding the potential sulfurization of $IPSO_{25}$, we
30	examined the GC-MS chromatogram and mass spectra of each sample for the occurrence of the HBI C_{25} sulfide
31	(Sinninghe Damsté et al., 2007). The C_{25} HBI thiane was absent in all samples. For the quantification, manually

integrated peak areas of the molecular ions of the HBIs in relation to those of 7-HND were used. An external calibration for HBI diene and trienes was applied using a sample with known HBI concentrations from the Lancaster Sound, Canada, to account for the different response factors of the HBI molecular ions (m/z 346; m/z 348) and the fragment ion of 7-HND (m/z 266). The identification of sterols was based on comparison of their retention times and mass spectra with those of reference compounds run on the same instrument. Comparison of peak areas of individual analytes and the internal standard was used for sterol quantification. The error determined by duplicate GC-MS measurements was below 0.7%. The detection limit for HBIs and sterols was 0.5 ng/g sediment. Absolute concentrations of HBIs and sterols were normalized to total organic carbon contents (for TOC data see Cárdenas et al., 2018).

The herein presented phytoplankton-IPSO$_{25}$ index (PIPSO$_{25}$) is calculated using the same formula as for the PIP$_{25}$ index following Müller et al. (2011):

$$PIPSO_{25} = \frac{IPSO_{25}}{IPSO_{25}+(c \times phytoplankton\ marker)} \tag{1}$$

The balance factor c (c = mean IPSO$_{25}$ / mean phytoplankton biomarker) is applied to account for the high offsets in the magnitude of IPSO$_{25}$ and sterol concentrations.

Stable carbon isotope composition of IPSO$_{25}$, requiring a minimum of 50 ng carbon, was successfully determined on five samples using GC-irm-MS. The ThermoFisher Scientific Trace GC was equipped with a 30m Restek Rxi-5 ms column (0.25 mm diameter, 0.25 μm film thickness) and coupled to a Finnigan MAT 252 isotope ratio mass spectrometer via a modified GC/C interface. Combustion of compounds was done under continuous flow in ceramic tubes filled with Ni wires at 1000° C under an oxygen trickle flow. The same GC program as for the HBI identification was used. The calibration was done by comparison to a CO_2 reference gas. The values of $\delta^{13}C$ are expressed in per mill (‰) against Vienna PeeDee Belemnite (VPDB) and the mean standard deviation was <0.9 ‰. An external standard mixture was measured every six runs, achieving a long-term mean standard deviation of 0.2‰ and an average accuracy of <0.1 ‰.

3.3   Diatoms

Details of the standard technique of diatom sample preparation were developed in the micropaleontological laboratory at the Alfred Wegener Institute (AWI) in Bremerhaven. The preparation included a treatment of the sediment samples with hydrogen peroxide and concentrated hydrochloric acid to remove organic and calcareous remains. After washing the samples several times with purified water, the water was removed and the diatoms were embedded on permanent mounts for counting (see detailed description by Gersonde and Zielinski, 2000). The respective diatom counting was carried out according to Schrader and Gersonde (1978). On average, 400 to

600 diatom valves were counted in each slide using a Zeiss Axioplan 2 at x1000 magnification. In general preservation state of the diatom assemblages was moderate to good in the Bransfield Strait and decreased towards the Drake Passage where it is moderate to poor.

Diatoms were identified to species or species group level and if possible to forma or variety level. The taxonomy follows primarily Hasle and Syvertsen (1996), Zielinski and Gersonde (1997), and Armand and Zielinski (2001). Following Zielinski and Gersonde (1997) and Zielinski et al. (1998) we combined some taxa to groups: The *Thalassionema nitzschioides* group combines *T. nitzschioides* var. *lanceolata* and *T. nitzschioides* var. *capitulata*, two varieties with gradual transition of features between them and no significantly different ecological response. The species *Fragilariopsis curta* and *Fragilariopsis cylindrus* were combined as *F. curta* group taking into account their similar relationship to sea ice and temperature (Armand et al., 2005; Zielinski and Gersonde, 1997). Furthermore, the *Thalassiosira gracilis* group comprises *T. gracilis* var. *gracilis* and *T. gracilis* var. *expecta* because the characteristic patterns in these varieties are often transitional, which hampers distinct identification. Although the two varieties *Eucampia antarctica* var. *recta* and *E. antarctica* var. *antarctica* display different biogeographical distribution (Fryxell and Prasad, 1990), they were combined to the *E. antarctica* group. This group was not included in the transfer function (TF) as it shows no relationship to either sea ice or temperature variation (Esper and Gersonde, 2014a, b). Besides the *E. antarctica* group, we also discarded diatoms assembled as *Chaetoceros* spp. group from the TF-based re-constructions, following Zielinski et al. (1998) and Esper and Gersonde (2014a). This group combines mainly resting spores of a diatom genus with a ubiquitous distribution pattern that cannot be identified to species level due to the lack of morphological features during light microscopic inspection. Therefore, different ecological demands of individual taxa cannot be distinguished.

For estimating winter sea ice (WSI) concentrations we applied the marine diatom TF MAT-D274/28/4an, comprising 274 reference samples from surface sediments in the western Indian, the Atlantic and the Pacific sectors of the Southern Ocean, with 28 diatom taxa and taxa groups, and an average of 4 analogs (Esper and Gersonde, 2014a). The WSI estimates refer to September sea-ice concentrations averaged over a time period from 1981 to 2010 at each surface sediment site (National Oceanic and Atmospheric Administration, NOAA; Reynolds et al., 2002, 2007). The reference data set is suitable for our approach as it uses a 1° by 1° grid, representing a higher resolution than previously used and results in a root mean squared error of prediction (RMSEP) of 5.52% (Esper and Gersonde, 2014a). We defined 15% concentration as threshold for maximum sea-ice expansion following the approach of Zwally et al. (2002) for the presence or absence of sea ice, and 40% concentration representing the average sea-ice edge (Gersonde et al., 2005; Gloersen et al., 1993). MAT calculations were carried out with the statistical computing software R (R Core Team, 2012) using the additional packages Vegan (Oksanen

et al., 2012) and Analogue (Simpson and Oksanen, 2012). Further enhancement of the sea-ice reconstruction was obtained by consideration of the abundance pattern of the diatom sea-ice indicators allowing for qualitative estimate of sea-ice occurrence, as proposed by Gersonde and Zielinski (Gersonde and Zielinski, 2000).

3.4   Sea ice data

The mean monthly satellite sea ice concentration was derived from Nimbus-7 SMMR and DMSP SSM/I-SSMIS passive microwave data and downloaded from the National Snow and Ice Data Center (NSIDC; Cavalieri et al., 1996). An interval from 1980 to 2015 was used to generate an average sea ice distribution for each season, spring (SON), summer (DJF), autumn (MAM) and winter (JJA).

4   Results and Discussion

4.1   Distribution of IPSO$_{25}$, HBI trienes and sterols

The sea ice biomarker IPSO$_{25}$ was detected in 14 samples, with concentrations ranging between 0.37 and 17.81 µg g^{-1} TOC (Table 1). The HBI Z-triene was present in all 26 samples (0.33-26.86 µg g^{-1} TOC) and the HBI E-triene was found in 24 samples (0.15-13.87 µg g^{-1} TOC). Brassicasterol was present in all measured samples with concentrations ranging from 3.39 to 5017.44 µg g^{-1} TOC while dinosterol was detected in 22 samples (0.0002-1983.75 µg g^{-1} TOC).

The distribution of IPSO$_{25}$ in the study area shows a clear northwest-southeast gradient (Fig. 3a) with concentrations increasing from the continental slope and around the South Shetland Islands towards the continental shelf. Maximum IPSO$_{25}$ concentrations are observed in the Bransfield Strait. According to Belt et al. (2016), deposition of IPSO$_{25}$ is highest in area covered by landfast sea ice and platelet ice during early spring and summer. We suppose that core sites PS97/068 to PS97/073 in the central and eastern Bransfield Strait are located too distal to be covered by fast ice and suggest that peak IPSO$_{25}$ concentrations at these sites may refer to the frequent drift and melt of sea ice exported from the Weddell Sea into the Bransfield Strait. The vertical export of biogenic material from sea ice towards the seafloor may be accelerated significantly by the formation of organic-mineral aggregates, fecal pellets or by (cryogenic) gypsum ballasting, which promotes a rapid burial and sedimentation of organic matter in polar settings (De La Rocha and Passow, 2007; Wefer et al., 1988; Wollenburg et al., 2018). Lateral subsurface advection of organic matter (incl. biomarkers) through the TWW, however, may also contribute to elevated IPSO$_{25}$ concentrations at these sites.

IPSO$_{25}$ was not detected in sediments from the permanently ice-free areas in the Drake Passage. Highest concentrations of both HBI trienes are found in the eastern Drake Passage and along the continental slope, while their concentrations in the Bransfield Strait are rather low (Fig. 3b and c) suggesting unfavorable environmental conditions (ocean temperature, sea ice cover) for their source diatoms.

We observe higher concentrations of brassicasterol and dinosterol in the eastern part of the Drake Passage and, in contrast to the observation made for HBI trienes, also in the eastern and central Bransfield Strait (Fig. 3d and e). Dinosterol and, in particular, brassicasterol are known to have different source organisms including diatoms, dinoflagellates, prymnesiophycean algae and cyanobacteria (Volkman, 1986) and we assume that this diversity may account for the higher concentration of these lipids in Bransfield Strait sediments, while concentrations of HBI trienes, mainly derived from diatoms, are significantly lower. Sediments collected along the Hero Fracture Zone in the western Drake Passage (Fig. 2) contain only minor amounts of biomarkers except for elevated brassicasterol concentrations observed at stations PS97/048-1 and 049-2 (Fig. 3d). This part of the Drake Passage

1 is mainly barren of fine-grained sediments and dominated by sands (Lamy, 2016), which may point to intensive

2 winnowing by ocean currents impacting the deposition and burial of organic matter. We consider that also

3 degradation of HBIs and sterols may affect their distribution within surface sediments. Rontani et al. (2014) report

4 a higher sensitivity of tri-unsaturated HBIs to oxidation but also note that oxidation conditions in pelagic

5 environments (i.e. their source organisms' habitat) are not as significant as those within sea ice. A more recent

6 study by Rontani et al. (2019) on surface sediments shows that $IPSO_{25}$ may potentially be affected by autoxidative

7 and bacterial degradation but oxidation products are found in only minor proportions. In general, further

8 investigations into degradation processes affecting both HBIs and sterols within sediments would address an

9 important knowledge gap regarding in-situ biochemical modifications of the biomarker signal.

10 The $\delta^{13}C$ values of $IPSO_{25}$ are between -10.3‰ and -14.7‰ which is the commonly observed range for $IPSO_{25}$ in

11 surface sediments and sea ice derived organic matter (Massé et al., 2011, Belt et al., 2016), and contrasts the low

12 $\delta^{13}C$ values of marine phytoplankton lipids in Antarctic sediments (-38‰ to -41‰ after Massé et al., 2011).

13 Contrary to the finding of elevated Z-triene concentrations in surface waters along an ice-edge (Smik et al., 2016a)

14 and earlier suggestions that this biomarker may be used as a proxy for MIZ conditions (Belt et al., 2015; Collins

15 et al., 2013), we observe highest concentrations of the Z- and E-triene at the permanently ice-free northernmost

16 stations in the eastern Drake Passage. This is also apparent for brassicasterol and dinosterol supporting an open

17 marine (pelagic) source for these sterols. Moderate concentrations of HBI trienes at the continental slope along

18 the WAP and in the Bransfield Strait likely refer to primary production at the sea ice margin during spring and

19 summer indicating seasonally ice-free waters in high production coastal areas influenced by upwelling (Gonçalves-

20 Araujo et al., 2015). The similarity in the distribution of the Z- and the E-triene in our surface sediments – the

21 latter of which so far is not often considered for Southern Ocean paleoenvironmental studies – supports the

22 assumption of a common diatom source for these HBIs (Belt et al., 2000, 2017). Since brassicasterol and dinosterol

23 are highly abundant in both seasonally ice-covered Bransfield Strait sediments as well as in permanently ice-free

24 Drake Passage sediments, their use as an indicator of fully open-marine conditions is questionable. Elevated

25 concentrations of both sterols in the Bransfield Strait could either point to an additional input of these lipids from

26 melting sea ice (Belt et al., 2013) or a better adaptation of some of their source organisms to cooler and/or ice-

27 dominated ocean conditions. Production and accumulation of these lipids in (late) summer (i.e. after the sea ice

28 season) may be considered as well. This observation highlights the need for a better understanding of the source

29 organisms and the mechanisms involved in the synthesis of these sterols.

30

1 **4.2   A novel sea ice index for the Southern Ocean: PIPSO$_{25}$**

2 The main concept of combining the sea ice proxy with an indicator of an ice-free ocean environment (i.e. a

3 phytoplankton biomarker, Müller et al., 2011), aims at a semi-quantitative assessment of the sea ice conditions.

4 By reducing the light penetration through the ice, a thick and perennial sea ice cover limits the productivity of

5 bottom sea ice algae (Hancke et al., 2018), which results in the absence of both sea ice and pelagic phytoplankton

6 biomarker lipids in the underlying sediments. Vice versa, sediments from permanently ice-free ocean areas only

7 lack the sea ice biomarker but contain variable concentrations of phytoplankton biomarkers (Müller et al., 2011).

8 The co-occurrence of both biomarkers in a sediment sample suggests seasonal sea ice coverage promoting algal

9 production indicative of sea ice as well as open ocean environments (Müller et al., 2011).

10 Following the PIP$_{25}$-approach applied in the Arctic Ocean (Müller et al., 2011; Belt and Müller, 2013; Xiao et al.,

11 2015), we used IPSO$_{25}$, HBI triene and sterol data to calculate the PIPSO$_{25}$ index. Depending on the biomarker

12 reflecting pelagic (open ocean) conditions, we define P$_Z$IPSO$_{25}$ (using the Z-triene), P$_E$IPSO$_{25}$ (using the E-triene),

13 P$_B$IPSO$_{25}$ (using brassicasterol), and P$_D$IPSO$_{25}$ (using dinosterol). Since the concentrations of IPSO$_{25}$ and both HBI

14 trienes are in the same range, the application of the c-factor is not needed here. For the calculation of P$_B$IPSO$_{25}$

15 the c-factor is 0.0048, for P$_D$IPSO$_{25}$ it is 0.0137.

16 The PIPSO$_{25}$ values are 0 in the Drake Passage and increase to intermediate values at the South Shetland Islands

17 and the continental slope and reach highest values in the Bransfield Strait (Fig. 4a-d). Minimum PIPSO$_{25}$ values

18 are supposed to refer to a predominantly ice-free oceanic environment in the Drake Passage, while moderate

19 PIPSO$_{25}$ values mark the transition towards a marginal sea ice coverage at the continental slope and around the

20 South Shetland Islands. Elevated PIPSO$_{25}$ values in samples from the northeastern Bransfield Strait suggest an

21 increased sea ice cover (probably sustained through the drift of sea ice originating in the Weddell Sea). This pattern

22 reflects the oceanographic conditions of a permanently ice-free ocean north of the South Shetland Islands and a

23 seasonal sea ice zone at the WAP influenced by the Weddell Sea as described by Cárdenas et al. (2018). Both HBI

24 triene-based PIPSO$_{25}$ indices show constantly high values at the coast of the WAP of >0.7 (P$_Z$IPSO$_{25}$) and >0.8

25 (P$_E$IPSO$_{25}$), respectively, in the southern Bransfield Strait paralleling the southwest-northeast oriented Peninsula

26 Front described by Sangrà et al. (2011). This front is reported to act as a barrier for phytoplankton communities

27 (Gonçalves-Araujo et al., 2015) and is associated with the encounter between TWW carrying Weddell Sea sea ice

28 through the Antarctic Sound and the TBW. The high PIPSO$_{25}$ values suggesting an extended sea ice cover west of

29 the Peninsula Front (station PS97/054 and PS97/056) result from minimum concentrations of pelagic biomarkers

30 and slightly higher concentrations of IPSO$_{25}$.

1    $PIPSO_{25}$ values based on the E-triene are about 0.2 higher compared to $P_Z IPSO_{25}$, due to the generally lower

2   concentrations of the E-triene (Table 1). The sterol-based $PIPSO_{25}$ values display a generally similar pattern as

3    $P_Z IPSO_{25}$ and $P_E IPSO_{25}$, respectively, and we note a high comparability between the $P_E IPSO_{25}$ and $P_B IPSO_{25}$

4   values (r^2 = 0.73). Some differences, however, are observed in the southwestern part of the Bransfield Strait

5   (station PS97/056) where $P_B IPSO_{25}$ indicates a lower sea ice cover and in the central Bransfield Strait (stations

6   PS97/068 and PS97/069) where $P_B IPSO_{25}$ and $P_D IPSO_{25}$ reflect only MIZ conditions. The HBI-triene based

7    $PIPSO_{25}$ indices hence seem to reflect the oceanographic conditions within the Bransfield Strait more

8   satisfactorily. We note that the brassicasterol or dinosterol-based $PIPSO_{25}$ index links environmental information

9   derived from biomarker lipids belonging to different compound classes (i.e. HBIs and sterols), which have

10   fundamentally different chemical properties. This requires special attention as, for example, selective degradation

11   of one of the compounds may affect the sedimentary concentration of the respective lipids. Previous studies linking

12   HBI and sterol-based sea ice reconstructions with satellite-derived or, with respect to downcore paleo studies,

13   paleoclimatic data, however, demonstrate that the climatic/environmental conditions controlling the production of

14   HBIs and sterols seem to exceed the influence of a potential preferential degradation of these biomarkers within

15   the sediments (e.g., Berben et al., 2014; Cabedo-Sanz et al., 2013; Müller et al., 2009, 2012; Müller and Stein,

16   2014; Stein et al., 2017; Xiao et al., 2015). A comparison of PIP_{25} records determined using brassicasterol and the

17   HBI Z-triene for three sediment cores from the Arctic realm covering the past up to 14.000 years BP (Belt et al.,

18   2015) reveals very similar trends for both versions of the PIP_{25} index in each core, which may point to, at least, a

19   similar degree of degradation of HBI trienes and sterols through time. More such studies are needed to evaluate

20   the preservation potential of HBIs and sterols in Southern Ocean sediments.

21

22   **4.3   Comparison of satellite-derived modern sea ice conditions and biomarker data**

23   Satellite-derived sea ice data were averaged over the time period from 1980 to 2015 for all four seasons (Table 2)

24   and are considered to reflect the modern mean state of sea ice coverage around the WAP. The sea ice concentration

25   is expressed to range from 0 to 100 % and, although the error can be up to 15 %, concentrations below 15 % still

26   suggest the occurrence of sea ice. These low sea ice concentrations are usually neglected for the determination of

27   the sea ice extent, which is defined as the ocean area with a sea ice cover of at least 15 %. The spring and winter

28   sea ice concentrations are shown in Figure 4 e-f. Winter sea ice is estimated to not extend north of 61° S (Fig. 4 f)

29   and varies between 1 % and 50 % in the study area, while sea ice is reduced to less than 20 % in spring (Fig. 4e,

30   Table 2).

1   Sea ice concentrations of up to 50 % are common in winter between the South Shetland Islands and north of the

2   Antarctic Sound where the influence of TWW is highest. Permanent sea ice cover is uncommon in the Bransfield

3   Strait and around the WAP and this area is mainly characterized by a high sea ice seasonality and drift ice from

4   the Weddell Sea (Collares et al., 2018). Comparisons of individual biomarker concentrations with satellite sea ice

5   data reveal a weak and positive correlation between $IPSO_{25}$ concentrations and winter sea ice concentrations ($r^2 =$

6   0.5), while no correlation is found between sea ice and pelagic biomarker concentrations ($r^2 < 0.1$ for all relations).

7   Correlations of $PIPSO_{25}$ values with satellite-derived sea ice concentrations (for spring, summer, autumn and

8   winter) contrast earlier observations made for the PIP_{25} index in the Arctic Ocean, where the closest linear

9   relationship is found mainly with the spring sea ice coverage (i.e. the blooming season of sea ice algae; Müller et

10   al., 2011; Xiao et al., 2015).

11   Esper and Gersonde (2014a) studied the response of diatom species to changes in environmental conditions and

12   their response to the non-linear behaviour of sea ice dynamics (Zwally et al., 2002). In contrast to ice free areas or

13   areas of permanent sea ice cover, areas characterized by the transition from consolidated to unconsolidated sea ice

14   show rapid changes in satellite derived sea ice concentrations (ranging from 90 % to 15 %) and exhibit a large

15   variability in species composition. To reflect this curve in sea ice we hence chose a cubic polynomial regression

16   (polynomial of third degree) to determine the relation between $PIPSO_{25}$ values and satellite data depicting sea ice

17   concentrations of more than 20 %.

18   We observe a remarkably low correlation between $PIPSO_{25}$ values and spring sea ice concentrations of less than

19   20 % with a coefficient of determination $r^2 = 0.37$ for P_ZIPSO_{25}, $r^2 = 0.50$ for P_EIPSO_{25} (Fig. 5a), $r^2 = 0.31$ for

20   P_BIPSO_{25}, and $r^2 = 0.34$ for P_DIPSO_{25} (Fig. 5b). The highest correlation is observed between winter sea ice

21   concentrations and P_EIPSO_{25} ($r^2 = 0.72$), and P_ZIPSO_{25} ($r^2 = 0.65$, Fig. 5c). A weaker correlation is noted for the

22   sterol-based $PIPSO_{25}$ values (P_BIPSO_{25}: $r^2 = 0.52$; P_DIPSO_{25}: $r^2 = 0.44$, Fig. 5d). The slightly sigmoid-shaped

23   regression line of winter sea ice concentrations and $PIPSO_{25}$ values reflects the non-linearity of sea ice cover in

24   different sea ice regimes as mentioned above.

25   The contour lines in Figure 4 a-d show the observed extent of 15 %, 30 %, 40 % and 50 % winter sea ice compared

26   to the $PIPSO_{25}$ values. In the northeastern part of the study area, the HBI triene based $PIPSO_{25}$ indices align well

27   with the contour lines of winter sea ice concentrations and depict the gradient from the marginally ice-covered

28   southern Drake Passage towards the intensively ice-covered Weddell Sea. In the southwestern part of the

29   Bransfield Strait, all $PIPSO_{25}$ indices suggest a higher sea ice cover than it is reflected in the satellite data. This

30   may be explained by the transport (and melt) of drift ice through the TWW, joining the TBW at the southwestern

31   Peninsula Front. In contrast, also the absence of $IPSO_{25}$ at stations PS97/052 and PS97/053, off the continental

1 slope, is in conflict with the satellite data depicting an average winter sea ice cover of 23 %. Earlier documentations

2 that the IPSO$_{25}$ producing sea ice diatom *Berkeleya adeliensis* favors land-fast ice communities in East Antarctica

3 (Riaux-Gobin and Poulin, 2004) and platelet ice occurring mainly in near coastal areas (Belt et al., 2016) could

4 explain this mismatch between biomarker and satellite data, which further strengthens the hypothesis that the

5 application of IPSO$_{25}$ seems to be confined to continental shelf or near-coastal and meltwater affected

6 environments (Belt et al., 2016). Alternatively, strong ocean currents (i.e. the ACC) could impact the deposition

7 of IPSO$_{25}$ in this region.

8 As photosynthesis is not possible and a release of sea ice diatoms from melting sea ice is highly reduced during

9 the Antarctic winter, the observation of a stronger correlation between recent winter sea ice concentrations and

10 PIPSO$_{25}$ sea ice estimates is unexpected. We hence suggest that this offset may be related to the fact that the

11 sediment samples integrate a longer time interval than is covered by satellite observations. Radiocarbon dating of

12 selected samples that contain calcareous material reveals an age of 100 years BP in the vicinity of the South

13 Shetland Islands (station PS97/059-2) and 142 years BP at the Antarctic Sound (station PS1546-2, Table 3). A

14 significantly older age was determined for a sample of *N. pachyderma* from station PS97/044-1 (4830 years BP)

15 which likely denotes the winnowing and/or very low sedimentation rates in the Drake Passage. Bioturbation effects

16 and uncertainties in reservoir ages potentially mask the ages of the near-coastal samples.

17 Nevertheless, since also other published ages of surface sediments within the Bransfield Strait (Barbara et al.,

18 2013; Barnard et al., 2014; Etourneau et al., 2013; Heroy et al., 2008) are in the range of 0-270 years, we consider

19 that our surface samples likely reflect the paleoenvironmental conditions that prevailed during the last two

20 centuries (and not just the last 35 years covered by satellite observations). In the context of the rapid warming

21 during the last century (Vaughan et al., 2003) and the decrease of sea ice at the WAP (King, 2014; King and

22 Harangozo, 1998), we suggest that the biomarker data of the surface sediments relate to a spring sea ice cover,

23 which must have been enhanced compared to the recent (past 35 years) spring sea ice recorded via remote sensing.

24 Presumably, the average spring sea ice conditions over the past 200 years might have been similar to the modern

25 (past 35 years) winter conditions, which would explain the stronger correlation between PIPSO$_{25}$ sea ice estimates

26 and winter sea ice concentrations.

27

28 **4.4   Comparison of diatom-based sea ice estimates and biomarker data**

29 The diatoms preserved in sediments from the study area (Table 4) can be associated with open ocean and sea ice

30 conditions. North of the South Shetland Islands, the strong influence of the ACC is reflected in the high abundance

31 of open ocean diatom species such as *Fragilariopsis kerguelensis* and *Thalassiosira lentiginosa* (Esper et al.,

2010). The two diatom species *Fragilariopsis curta* and *Fragilariopsis cylindrus* – known to not produce HBI (Belt et al., 2016; Sinninghe Damsté et al., 2004) mark the vicinity to sea ice (Buffen et al., 2007; Pike et al., 2008) and indicate fast and melting ice, a stable sea ice margin and stratification due to melting processes and the occurrence of seasonal sea ice. The high abundance of these species in our samples is in good agreement with high and moderate IPSO$_{25}$ concentrations and PIPSO$_{25}$ values in the Bransfield Strait and around the South Shetland Islands, respectively. The only HBI source diatom identified is the HBI Z-triene producing *Rhizosolenia hebetata* (Belt et al., 2017), which is present in four samples in rather small amounts and does not show a relation to the measured Z-triene concentrations (Table 1 and 4). The source diatom of IPSO$_{25}$ *Berkeleya adeliensis* was not observed (or preserved) in the samples, and we assume that other, hitherto unknown, producers may exist.

We applied the transfer function of Esper and Gersonde (2014a) with four analogs (4an, Table 4) to our samples to compare the different estimates of sea ice cover based on biomarkers and diatoms. A positive correlation of the linear relationship is found between WSI concentrations derived from diatoms and the PIPSO$_{25}$ indices based on HBI trienes (P$_Z$IPSO$_{25}$ with $r^2 = 0.76$; P$_E$IPSO$_{25}$ with $r^2 = 0.77$, Fig. 6a). The correlations of sterol-based PIPSO$_{25}$ values with WSI are slightly lower but in the same range (P$_B$IPSO$_{25}$ with $r^2 = 0.74$; P$_D$IPSO$_{25}$ with $r^2 = 0.69$, Fig. 6b). A slightly weaker correlation is noted for diatom- and satellite-based winter sea ice concentrations ($r^2 = 0.63$; Fig. 6c). Overall, the diatom approach indicates higher sea ice concentrations than the satellite data with an offset of up to 65 %. This may be due to different sources of satellite reference data used for the transfer function or also due to the fact that the sediment samples integrate a longer time period with a higher sea ice cover than the satellite data (see discussion in section 4.3).

4.5   Application of PIPSO$_{25}$ as a semi-quantitative sea ice index

Precise and, in particular, quantitative reconstructions of past sea ice coverage are crucial for a robust assessment of feedback mechanisms in the ice-ocean-atmosphere system. While diatom transfer functions provide a valuable tool, additional information on sea ice conditions in coastal ice-shelf proximal areas, which are often affected by opal dissolution, are essential. The PIPSO$_{25}$ approach seems to be a promising step into this direction, though our data obtained for the WAP are not yet sufficient for a full calibration. PIPSO$_{25}$, diatom and satellite sea ice data, however, reveal positive correlations (Figs. 5 and 6) and depict similar gradients in sea ice cover. The observed offset between satellite data and biomarker- and diatom-based sea ice estimates likely relates to the fact that the instrumental records cover a significantly shorter or more recent time interval than the studied sediments. The recent rapid warming along the WAP (Vaughan et al., 2003) hence complicates attempts to calibrate these proxy data against observational data. Regarding the interpretation of PIPSO$_{25}$ in terms of sea ice coverage in the study

1   area, lower PIPSO$_{25}$ values (<0.15 for P$_Z$IPSO$_{25}$; <0.31 for P$_E$IPSO$_{25}$; <0.22 for P$_B$IPSO$_{25}$ and P$_D$IPSO$_{25}$) roughly

2   seem to reflect unconsolidated, drifting winter sea ice and a nearly ice-free spring season. Higher values (>0.71

3   for P$_Z$IPSO$_{25}$; >0.9 P$_E$IPSO$_{25}$; >0.6 for P$_B$IPSO$_{25}$ and P$_D$IPSO$_{25}$) would refer to an extended winter sea ice cover

4   (up to 91 % in some years) with ice floes remaining until summer.

5 Conclusion

The distribution of the sea ice biomarker $IPSO_{25}$ and related HBI trienes and sterols as well as diatoms in a suite of surface sediments from the southern Drake Passage and the WAP reflects recent sea ice conditions reasonably well. The herein proposed sea ice index $PIPSO_{25}$ indicates seasonal sea ice cover along the coast of the WAP and in the Bransfield Strait, whereas mainly ice-free conditions prevail in the Drake Passage. In general, this pattern is consistent with satellite-derived sea ice data and diatom-based sea ice estimates and we note that the $PIPSO_{25}$ index seems a promising approach towards semi-quantitative sea ice reconstructions in the Southern Ocean. The recent rapid warming in the study area, however, affects the comparability of proxy and satellite data. The fact that the surface sediments integrate a significantly longer time interval than the remote sensing data thwarts attempts to calibrate $PIPSO_{25}$ values against observed sea ice concentrations. Additional data from other circum-Antarctic coastal (and distal) environments and investigations into potential calibration methods are needed to further develop this index as a quantitative sea ice proxy.

1 **Data Availability**

2 All data can be found in this paper and will be available at the open access repository *www.pangaea.de*.

3

4 **Author contributions**

5 The study was conceived by MV and JM. Data collections and experimental investigations were done by MV

6 together with OE (diatoms), GM (radiocarbon dating), CH (satellite data), and ES (isotope data). MV wrote the

7 manuscript and did the visualizations. KF provided technical support. JM supervised the study. All authors

8 contributed to the interpretation and discussion of the results and the conclusion of this study.

9

10 **Competing interests**

11 None of the authors has a conflict of interest.

12

20

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23

1   **Figures**

a

b

c
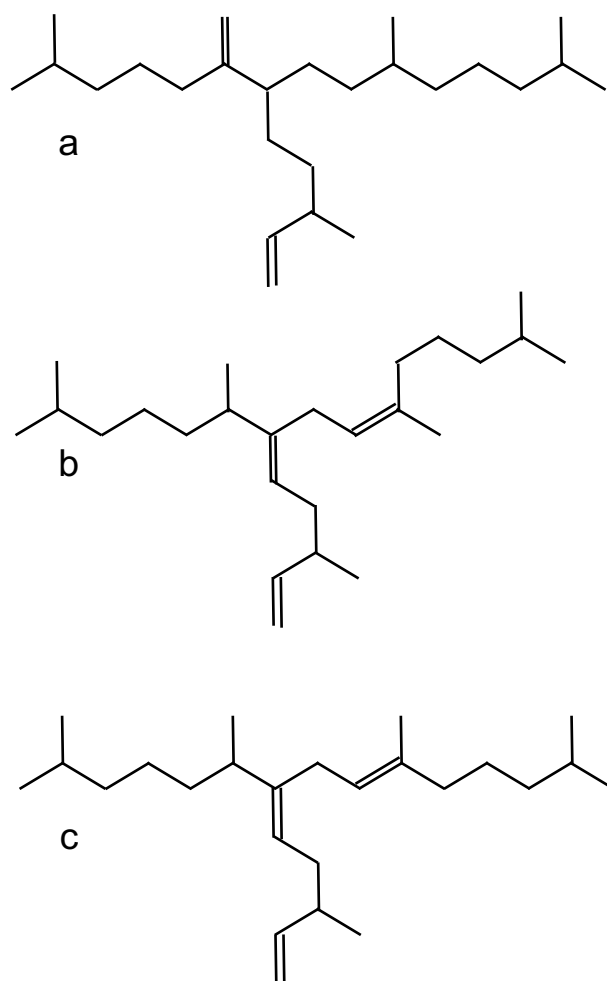
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3   **Figure 1: The molecular structures of a) IPSO$_{25}$, b) the HBI Z-triene, and c) the HBI E-triene.**

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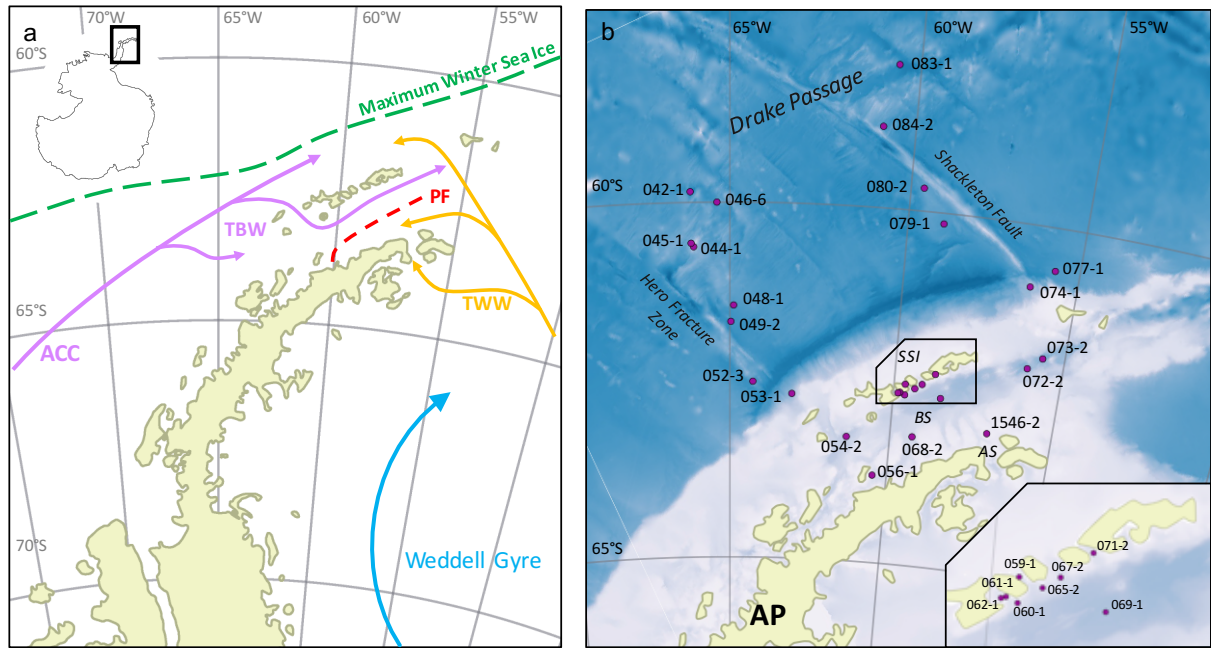
Figure 2: a) Oceanographic setting of the study area (modified after Hofmann et al., 1996; Sangrà et al., 2011) with

ACC = Antarctic Circumpolar Current, TBW = Transitional Bellingshausen Water, TWW = Transitional Weddell

Water, and PF = Peninsula Front, and the maximum winter sea ice extent (after Cárdenas et al., 2018). b) The

bathymetric map of the study area with locations of all stations; AP = Antarctic Peninsula, AS = Antarctic Sound, BS

= Bransfield Strait, and SSI = South Shetland Islands. A detailed station map at the South Shetland Islands is integrated.

The overview maps were done with QGIS 3.0 from 2018 and the bathymetry was taken from GEBCO_14 from 2015.

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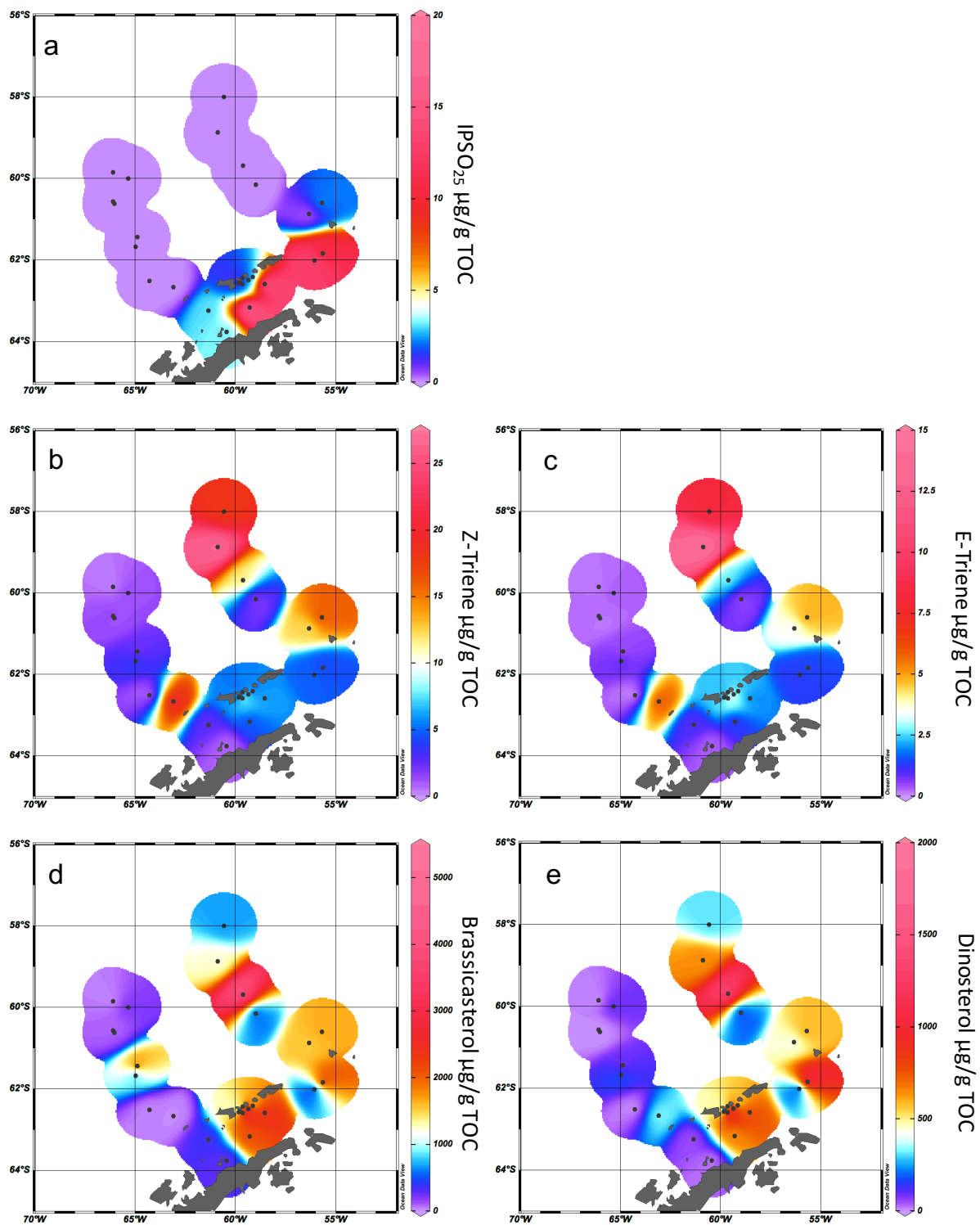
Figure 3: Distribution of a) IPSO$_{25}$, b) HBI Z-triene, c) HBI E-triene, d) brassicasterol, and e) dinosterol concentrations normalized to TOC. All distribution plots were made with Ocean Data View 4.7.10 (2017).

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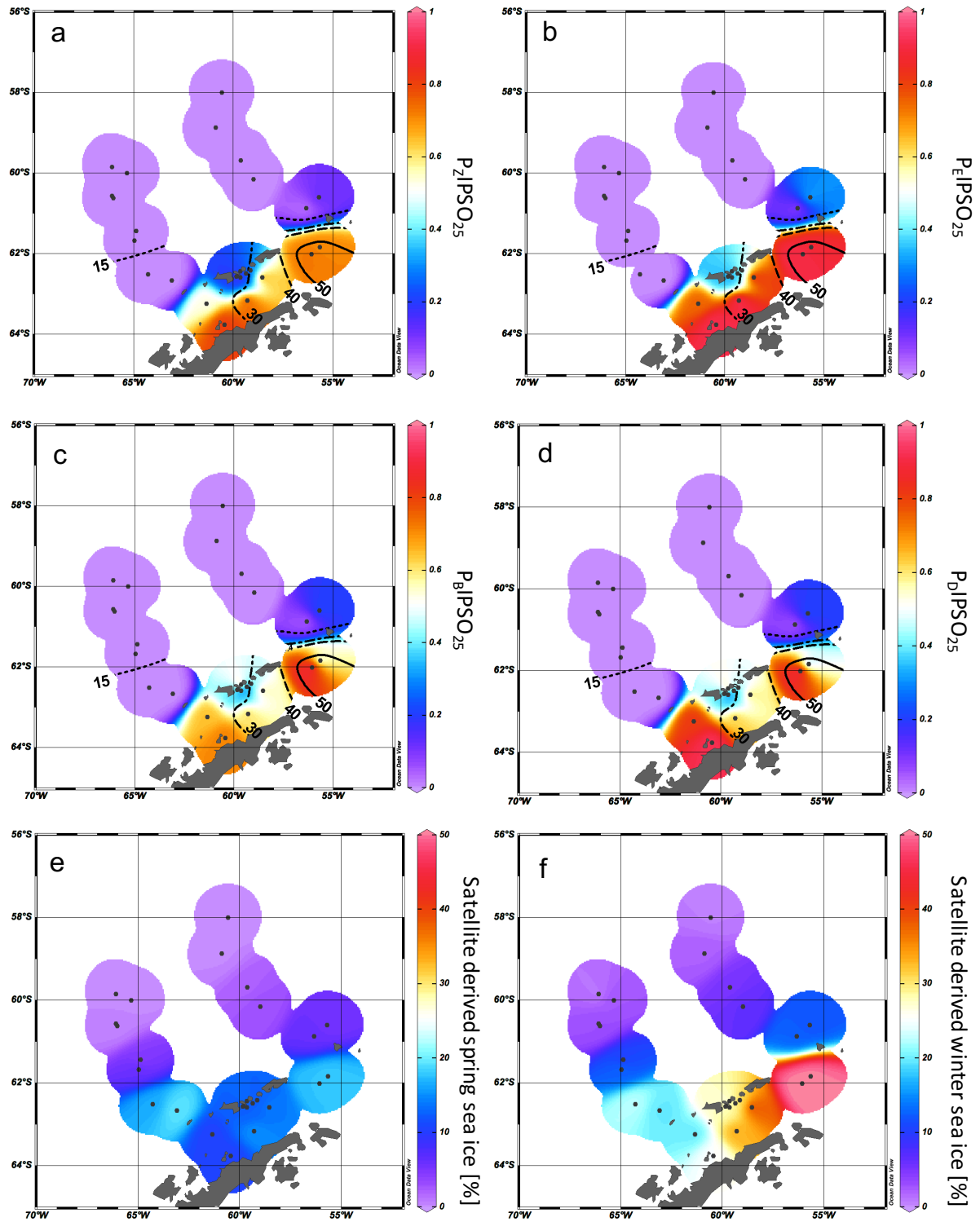
Figure 4: Distribution of a) $P_Z IPSO_{25}$, b) $P_E IPSO_{25}$, c) $P_B IPSO_{25}$, and d) $P_D IPSO_{25}$ values in the study area. The extent of 15 %, 30 %, 40 % and 50 % satellite sea ice concentrations during winter is added as contour lines. Satellite derived mean sea ice concentrations at each sampling station are shown for e) spring and f) winter.

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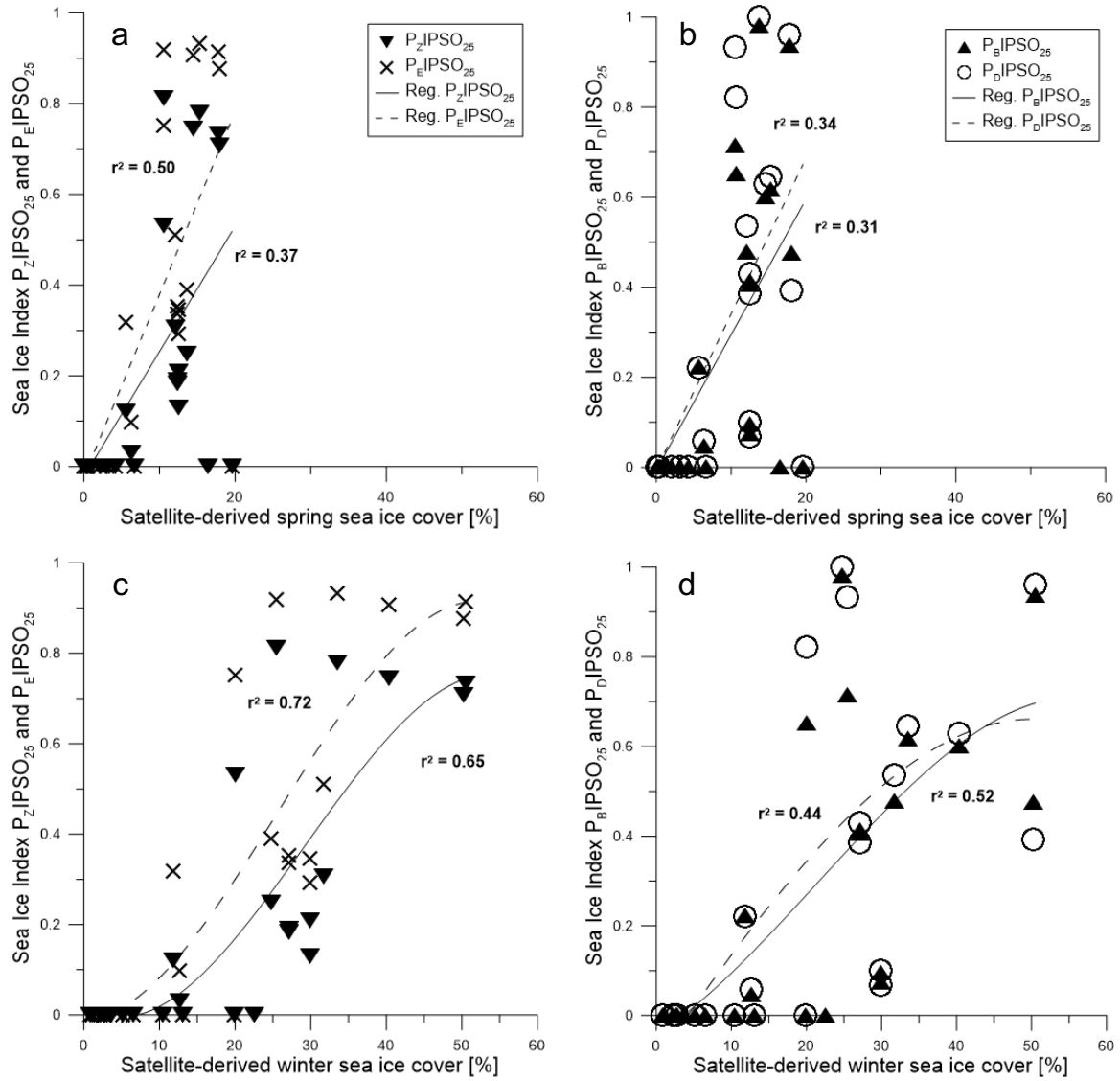
2

3   **Figure 5: Scatter plots of satellite spring sea ice concentrations and a) $P_Z IPSO_{25}$ (triangles, solid regression line) and**

4   **$P_E IPSO_{25}$ (crosses, dashed regression line) and b) $P_B IPSO_{25}$ (triangles, solid regression line) and $P_D IPSO_{25}$ (circles,**

5   **dashed regression line). Scatter plots of satellite winter sea ice concentrations and c) $P_Z IPSO_{25}$ (triangles, solid**

6   **regression line) and $P_E IPSO_{25}$ (crosses, dashed regression line) and d) $P_B IPSO_{25}$ (black triangles, solid regression line)**

7   **and $P_D IPSO_{25}$ (circles, dashed regression line). All scatter plots were done with Grapher^{TM} 13.**

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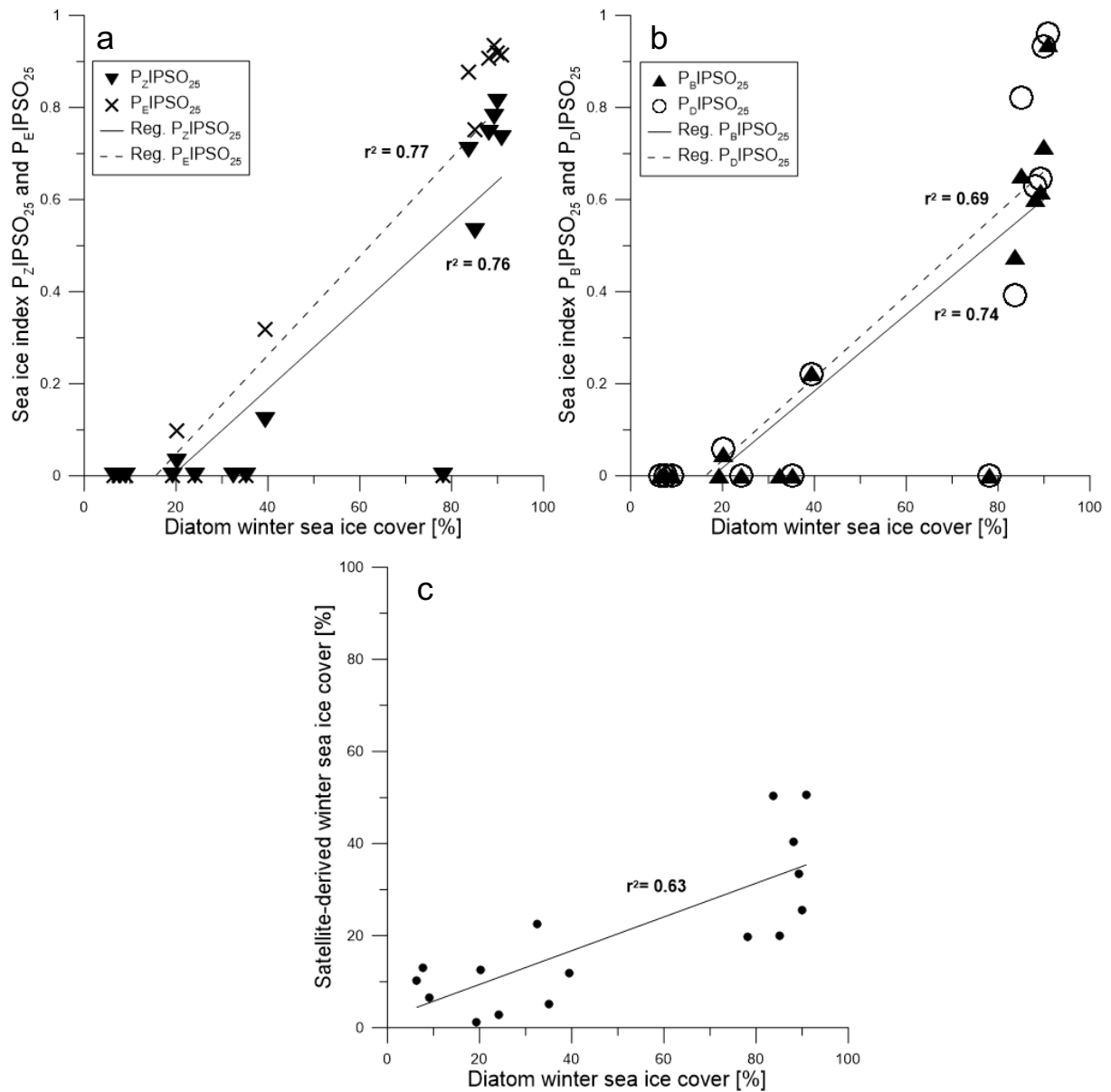
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3 **Figure 6: Scatter plots of a) P_ZIPSO_{25} (triangles, solid regression line) and P_EIPSO_{25} (crosses, dashed regression line)**

4 **and b) P_BIPSO_{25} (triangles, solid regression line) and P_DIPSO_{25} (circles, dashed regression line) against diatom derived**

5 **winter sea ice concentrations. c) Scatter plot of diatom transfer function winter sea ice concentrations and satellite**

6 **winter sea ice concentrations.**

7   **Tables**

8   **Table 1: Coordinates of sample stations with water depth, concentrations of IPSO$_{25}$, Z- and E-trienes, brassicasterol and dinosterol normalized to TOC, $\delta^{13}C$ values for IPSO$_{25}$, and values**

9   **of sea ice indices PIPSO$_{25}$ based on the Z- and E-triene, brassicasterol and dinosterol. Concentrations below the detection limit are expressed as 0. The PIPSO$_{25}$ could not be calculated**

10   **where IPSO$_{25}$ and the phytoplankton marker is absent (blank fields).**

Station	Lon [dE]	Lat [dN]	Water Depth [m]	IPSO$_{25}$/TOC [µg/g TOC]	HBI Z-Triene /TOC [µg/g TOC]	HBI E-Triene /TOC [µg/g TOC]	Brassi-casterol/TOC [µg/g TOC]	Dino-sterol/TOC [µg/g TOC]	δ^{13}C of IPSO$_{25}$ [‰]	P$_Z$IPSO$_{25}$	P$_E$IPSO$_{25}$	P$_B$IPSO$_{25}$	P$_D$IPSO$_{25}$
PS97/042-1	-66.10	-59.85	4172	0	0.333	0.152	12.997	0		0.000	0.000	0.000	
PS97/044-1	-66.03	-60.62	1203	0	1.080	0	143.688	0		0.000		0.000	
PS97/045-1	-66.10	-60.57	2292	0	1.531	0.386	36.902	0		0.000	0.000	0.000	
PS97/046-6	-65.36	-60.00	2803	0	1.359	0.291	214.634	101.809		0.000	0.000	0.000	0.000
PS97/048-1	-64.89	-61.44	3455	0	2.085	0.375	1859.609	73.532		0.000	0.000	0.000	0.000
PS97/049-2	-64.97	-61.67	3752	0	3.924	0.851	719.155	178.446		0.000	0.000	0.000	0.000
PS97/052-3	-64.30	-62.51	2890	0	0.679	0	26.554	0		0.000		0.000	
PS97/053-1	-63.10	-62.67	2021	0	19.350	5.948	13.356	332.868		0.000	0.000	0.000	0.000
PS97/054-2	-61.35	-63.24	1283	3.033	2.675	1.000	337.686	48.579	-14.741	0.531	0.752	0.652	0.820
PS97/056-1	-60.45	-63.76	633	3.232	0.752	0.290	268.190	17.158	-10.3 ± 0.9	0.811	0.918	0.716	0.932
PS97/059-1	-59.66	-62.44	354	0.835	2.523	1.305	3.386	0.0002		0.249	0.390	0.981	0.999
PS97/060-1	-59.65	-62.59	462	1.934	12.937	4.693	5017.437	1983.750		0.130	0.292	0.074	0.066
PS97/061-1	-59.80	-62.56	467	1.018	4.341	1.870	302.356	119.512		0.190	0.352	0.413	0.383
PS97/062-1	-59.86	-62.57	477	0.907	4.044	1.787	276.372	88.272		0.183	0.337	0.407	0.428
PS97/065-2	-59.36	-62.49	480	2.416	9.184	4.549	4788.292	1587.309		0.208	0.347	0.095	0.100
PS97/067-2	-59.15	-62.42	793	1.785	4.038	1.710	406.567	113.728		0.307	0.511	0.478	0.533
PS97/068-2	-59.30	-63.17	794	16.206	4.558	1.152	2096.690	653.977	-14.1 ± 0.6	0.780	0.934	0.617	0.643
PS97/069-1	-58.55	-62.59	1642	17.814	6.115	1.824	2472.025	774.345	-12.6 ± 0.4	0.744	0.907	0.601	0.626
PS97/072-2	-56.07	-62.01	1992	13.689	4.997	1.277	192.625	40.686	-13.6 ± 0.3	0.733	0.915	0.937	0.961
PS97/073-2	-55.66	-61.84	2624	10.369	4.283	1.451	2388.458	1180.752		0.708	0.877	0.476	0.390
PS97/074-1	-56.35	-60.87	1831	0.371	12.075	3.409	1539.629	438.073		0.030	0.098	0.048	0.058
PS97/077-1	-55.71	-60.60	3587	2.267	16.356	4.874	1647.616	589.731		0.122	0.317	0.223	0.219
PS97/079-1	-59.00	-60.15	3539	0	1.893	0.510	479.917	154.400		0.000	0.000	0.000	0.000
PS97/080-2	-59.64	-59.68	3113	0	12.021	2.705	4019.003	1329.129		0.000	0.000	0.000	0.000
PS97/083-1	-60.57	-58.00	3756	0	18.256	8.280	686.502	308.610		0.000	0.000	0.000	0.000
PS97/084-2	-60.88	-58.87	3617	0	26.857	13.871	1245.652	648.474		0.000	0.000	0.000	0.000

11

Table 2: Seasonal sea ice concentrations from satellite observations for spring, summer, autumn and winter with standard deviations.

Station	Sea Ice Spring [%]	Sea Ice Spring StDev [%]	Sea Ice Summer [%]	Sea Ice Summer StDev [%]	Sea Ice Autumn [%]	Sea Ice Autumn StDev [%]	Sea Ice Winter [%]	Sea Ice Winter StDev [%]
PS97/042-1	0.04	0.19	0.00	0.00	0.01	0.05	1.14	5.00
PS97/044-1	0.92	3.25	0.02	0.23	0.00	0.00	3.67	9.38
PS97/045-1	0.52	2.08	0.01	0.08	0.00	0.04	2.65	7.81
PS97/046-6	0.29	1.35	0.00	0.00	0.00	0.00	2.84	8.55
PS97/048-1	4.22	8.52	0.00	0.00	0.00	0.00	10.36	18.17
PS97/049-2	6.65	11.85	0.00	0.00	0.00	0.04	13.02	19.91
PS97/052-3	16.48	21.62	0.40	2.95	0.04	0.31	22.59	24.94
PS97/053-1	19.59	23.59	0.29	2.45	0.04	0.35	19.86	24.13
PS97/054-2	10.62	15.18	0.44	0.79	0.76	2.62	20.06	20.72
PS97/056-1	10.55	16.21	4.73	3.25	2.77	4.44	25.47	23.02
PS97/059-1	13.67	16.13	4.23	2.25	5.03	5.48	24.77	20.33
PS97/060-1	12.53	16.84	1.87	2.15	5.43	9.24	29.93	22.05
PS97/061-1	12.43	16.18	1.86	2.07	4.15	7.30	27.14	21.31
PS97/062-1	12.43	16.18	1.86	2.07	4.15	7.30	27.14	21.31
PS97/065-2	12.53	16.84	1.87	2.15	5.43	9.24	29.93	22.05
PS97/067-2	12.08	17.22	0.82	1.88	5.60	10.10	31.74	22.69
PS97/068-2	15.30	19.35	4.89	3.40	6.44	10.45	33.49	23.13
PS97/069-1	14.51	19.85	0.40	2.34	7.83	13.78	40.41	24.27
PS97/072-2	17.74	22.74	1.46	5.38	16.69	20.35	50.49	25.09
PS97/073-2	17.99	23.28	1.81	6.14	16.43	19.85	50.29	26.01
PS97/074-1	6.30	13.65	0.02	0.12	0.55	2.29	12.65	19.30
PS97/077-1	5.60	12.20	0.04	0.13	0.77	2.99	11.83	17.81
PS97/079-1	3.10	8.91	0.03	0.27	0.01	0.12	6.50	15.49
PS97/080-2	2.08	7.52	0.01	0.08	0.00	0.04	5.14	14.17
PS97/083-1	0.03	0.23	0.00	0.00	0.00	0.04	0.87	4.27
PS97/084-2	0.40	2.21	0.00	0.00	0.00	0.04	2.23	9.59

Table 3: Details of the radiocarbon dates and calibrated ages.

Sample Name	AWI-No.	Material	F[14]C ± error	Conventional ^{14}C age [a]	Calibrated age (cal BP) [a]
PS97/044-1	1657.1.1	N. pachyderma	0.5076	5447 ± 111	4830
PS97/059-2	1434.1.1	calcareous	0.8507	1299 ± 49	100
PS1546-2	1602.1.1	Moll.-Echinod	0.8456	1347 ± 64	142

Table 4: Estimations of winter sea ice (WSI) derived from diatom species and the distribution of main diatom species in each sample.

Station	Diatoms WSI (4an)	A.tabularis	E.antarctica	F.vanheurckii	F.kerguelensis	F.obliquecostata	F.sublinearis	F.curta	F.cylindrus	N.directa	O.weißflogii	P.lineola-turgid.-gr.	R.alata	R.hebetata fo. semispina	S.microtrias	T.lentiginosa	T.oliverana	Thalassiosira MT 3	P.pseudodenticulata	Stephanopyxis sp.
	[%]	[%]	[%]	[%]	[%]	[%]	[%]	[%]	[%]	[%]	[%]	[%]	[%]	[%]	[%]	[%]	[%]	[%]	[%]	[%]
PS97/042-1	19.2	0	0	0	0.8	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
PS97/046-6	24.2	0	0	0	0.8	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
PS97/048-1	6.4	0	0	0	0.8	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
PS97/049-2	7.7	0	0	0	0.7	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
PS97/052-3	32.4	1.4	4.3	0	60.3	0.2	0	0.9	0	0	0.5	0	0	0	0	16.2	0.5	0	0	0
PS97/053-1	78.1	0.4	1.2	0	48.5	0.1	0.1	7.4	0.3	0	0.4	0	0.3	0	0.1	6.5	0	0	0	0
PS97/054-2	85.2	0	0.4	0	6.2	0	0	6.9	0.4	0.2	2.7	0.2	0.2	0.2	0.4	0.9	0	0.2	0	0.4
PS97/056-1	89.9	0	0	0	0.5	0.3	0	1.9	0.5	0.3	0.5	0	0	0	0.5	0.5	0	0.0	0.5	0.3
PS97/068-2	89.2	0	0	0	0.7	0.3	0.6	4.9	2.4	0.1	0.5	0.1	0	0	0.4	0.4	0	0.4	0.2	0
PS97/069-1	88.2	0	0.2	0.4	2.1	0.2	0.4	4.3	1.3	0	0.6	0.2	0	0	0.2	0.4	0	0.2	0	0
PS97/072-2	90.9	0	0.2	1.1	1.7	0.6	0.9	8.1	0	0	5.7	0.2	0.2	0	0	1.7	0	0.9	0.9	0
PS97/073-2	83.7	0	0.2	0.2	1.8	0	0.4	7.4	1.0	0	1.0	0	1.6	0.8	0	0.4	0	2.1	0.6	0
PS97/074-1	20.1	0.4	0.6	0	63.1	0	0	2.3	0	0	0.2	0	0.0	0.2	0	7.4	0.4	0	0.2	0
PS97/077-1	39.4	0.6	3.3	0	49.1	0.8	0	3.1	0.2	0	0.2	0	0.0	1.3	0	10.0	0.2	0	0	0
PS97/079-1	9.1	0	0	0	0.7	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
PS97/080-2	35.1	0	0	0	0.7	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0

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