Co-evolution of the terrestrial and aquatic ecosystem in the Holocene Baltic Sea

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Abstract. The Baltic Sea experienced changes in marine input throughout the Holocene as substantial regional ice melt led to isostatic adjustment, eustatic sea level change, and periodic isolation from the North Sea. Here, we determine the distributions and isotopic signatures of organic compounds preserved in a sediment record spanning the last ~ 11 ka to reconstruct environmental change under these dynamic conditions. Carbon and hydrogen isotope ratios of short-, mid-, and long-chain *n*-alkanes, along with long-chain diol and glycerol dialkyl glycerol tetraether abundances, were analyzed in sediments sampled from the Arkona Basin in the western Baltic Sea. In the earliest part of the record (10 – 8.2 ka), hydrogen isotope values of higher plant-derived *n*-alkanes revealed a change in dominant water source from an ice melt-derived to a precipitation-dominated hydrological regime. Following this shift in water source, carbon isotope values of *n*-alkanes suggest diversification of vegetation. Shifts in hydrology and vegetation did not coincide with established phase boundaries, but instead occurred mid-phase or spanned phase transitions. For this reason, we suggest describing the Ancylus Lake in two phases: a meltwater phase (10.6 to 9.2 ka) and a precipitation phase (9.2 to 7.7 ka). Our results highlight the utility of using C and H isotope ratios in conjunction with other compound abundance proxies to better understand the complex environmental change recorded in paleo archives.

1 Introduction

The Baltic Sea, located in Northwest Europe, is a semi-enclosed marine basin characterized by restricted circulation, strong terrestrial influence, and relatively low oxygen concentrations that promote enhanced sedimentary organic matter contents and diverse phytoplankton communities (e.g., Wasmund and Uhlig, 2003; Golubkov et al., 2021). The Baltic Sea existed at least as far back as the Eemian interglacial (~130 ka, Andrén et al., 2015). Today, there is an open connection between the North Sea and the Baltic Sea via the Danish Straits in the west; however, this connection has not been permanent, causing the basin

to experience substantial environmental change throughout the Holocene. The Baltic Ice Lake – the manifestation of the Baltic Sea just prior to the start of the Holocene – emerged around 13 ka as large portions of the Scandinavian Ice Sheet (SIS) melted

35 and formed a meltwater lake filled with icebergs (Björck, 1995). The complex climate dynamics caused substantial shifts in the salinity of the Baltic Sea during the Holocene, reconstructed from geochemical and micropaleontological data (e.g., Alhonen, 1972; Ning et al., 2017; Ni et al., 2020; Weiss et al., 2020). Salinity changes in the Baltic Sea largely indicate periods of connection or isolation from the North Sea via the Danish Straits. The timing of different Holocene phases of the Baltic Sea is debated and appears to be divergent at different locations in the basin (Gustafsson and Westman, 2002; Moros et al., 2002).

40 The dates used hereafter for Baltic Sea phases are from two previously studied Arkona Basin cores (Warden et al., 2016, Fig. 1). As global and regional temperatures fluctuated at the start of the Holocene, eustatic sea level rise caused an inundation of saline North Sea water causing the transition of the basin from a freshwater to a brackish lake, a phase known as the Yoldia Sea (YS) (11.2 to 10.6 ka). Melting and subsequent calving stimulated retreat of the SIS, and in turn isostatic rebound of surrounding landmass led to changes in shoreline and water levels (e.g., Moros et al., 2002). The shoreline changes caused

45 isolation from the North Sea which led to freshening of the basin and a phase known as the Ancylus Lake (AL). The AL phase terminated at around 7.7 ka when a large marine transgression reconnected the Baltic Sea with the North Sea. The transgression reached a maximum at 7.2 ka and the prolonged inundation of saline North Sea water resulted in a second brackish phase called the Littorina Sea (LS), which lasted until 3 ka. After 3 ka, the Modern Baltic phase was established (Andrén et al., 2000, 2011), with further freshening due to continued continental uplift. The present day halocline in the Baltic proper sits around

50 60-80 m; surface salinity is around 7-8 g kg^{-1}, and salinity below the halocline is almost double at around 12 g kg^{-1} (Gustafsson and Westman, 2002). The halocline in the Kattegat is at the much shallower depth of 15 m, with salinities between 15-25 g kg^{-1} above and 32-35 g kg^{-1} below (Gustafsson and Westman, 2002). It is likely that the halocline depth changed throughout time as a result of various climatic conditions (e.g., Ni et al., 2020).

55 Regional pollen records from southern Sweden (Krageholmssjön, Färskesjön, and Trehörningen lakes, Antonsson and Seppä, 2007; Berglund et al., 2008a,b, Fig 1) and central Sweden (Lake Flarken, Seppä et al., 2005, Fig. 1) suggest an establishment of deciduous woody forests (first appearance of e.g., *Alnus* and *Quercus*) starting ~ 10 – 9.5 ka, while gymnosperm (especially *Pinus*) and *Betula* slowly retreated. At the same time, the northern region of Sweden and Finland showed the growth of gymnosperm forests (mainly *Pinus*, Lake Tsuolbmajavri, Seppä and Weckström 1999; Seppä and Birks, 2001), while

60 *Juniperus* decreased (maximum at ~ 9 ka). These major changes in vegetation coincide with an increase in mean annual air temperature in the region (e.g., Hammarlund et al., 2003; Antonsson et al., 2006). The mid-Holocene (9 – 3 ka) is characterized by a slow decrease in *Pinus* in the south, an overall low abundance of herbs, while in the north the vegetation remained stable (Seppä and Weckström 1999; Seppä and Birks 2001; Seppä et al., 2005; Berglund et al., 2008a,b; Antonsson and Seppä, 2007). Notable events during that period are short-lived cold-events (9.2 and 8.2 ka) that are recorded in the southern pollen records

65 indicated by a decrease in angiosperm taxa (e.g., *Alnus*) and increase in *Pinus* – although not all lakes record these events at the same scale or exact time. The last 3 ka are characterized by a decrease in deciduous tree taxa, an increase in herbs and

Betula, and the appearance of *Picea* in the south, in conjunction with a general cooling (Hammarlund et al., 2003; Antonsson et al., 2006). Overall, the species richness is the lowest at around 10.5 ka followed by a gradual increase that reaches a maximum at ~2.5 ka (Berglund and al., 2008b).

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We aim to complement the terrestrial pollen records with isotopic and lipid biomarker data from an Arkona Basin sediment record spanning the Holocene. Biomarkers, organic compounds indicative of a particular environmental process or produced exclusively by a particular group of organisms, are widely used organic geochemical tools that provide a wealth of paleoclimatic information. An overview of common proxies and the parameters they reflect can be found in Table 1.

75   Historically, geochemists have made use of isotope measurements of bulk sediments or biomass, which present a general overview of a given system (e.g., Leng and Lewis, 2017). Compound−specific carbon isotope ratios (δ^{13}C, the ratio of ^{13}C/^{12}C in a sample relative to a standard, Vienna Pee Dee Belemnite, aka VPDB) of organic molecules give insight into sources of organic matter (e.g., terrestrial versus marine; Meyers, 1994) and atmospheric CO_2 conditions (e.g., Freeman and Hayes, 1992). Compound−specific hydrogen isotope ratios (δ^2H, the ratio of ^{2}H/^{1}H in a sample relative to a standard, Vienna Standard Mean

80   Ocean Water, aka VSMOW) of lipids permit further understanding of compound biosynthesis and environmental conditions, such as precipitation and salinity (e.g., Sachse, Billault et al., 2012). These two elemental signatures are widely applied to study past environmental change across the planet and throughout the geologic record (e.g., Tipple and Pagani, 2010; Schefuß et al., 2011; Sinninghe Damsté et al., 2011). Straight-chain hydrocarbons, such as *n*-alkanes, are especially resistant to degradation and alteration over geological timescales and are found as far back as the Precambrian (McKirdy and Hahn, 1982;

85   Gordadze et al., 2017). In addition to being relatively recalcitrant, *n*-alkanes are ideal biomarkers for a C and H isotope approach because they contain non-exchangeable C and H, and are synthesized by a variety of organisms from bacteria to higher plants (e.g., Hemingway et al., 2016). Generally, long-chain *n*-alkanes are found in the protective wax layer of leaves from terrestrial higher plants (Eglinton and Hamilton, 1967), and the C isotopic compositions of *n*-alkanes from these plants can differ by up to 20‰ depending on metabolism (C_3, C_4, Crassulacean Acid Metabolism, aka CAM; see Diefendorf &

90   Freimuth, 2017 for review). C_4 plants produce more ^{13}C-enriched *n*-alkanes than C_3 plants, and *n*-alkanes synthesized by CAM plants show a wide range of isotopic compositions (Chikaraishi and Naraoka, 2003, 2005, 2007; Diefendorf and Freimuth, 2017). Mid-chain *n*-alkanes (C_{23} and C_{25}) are mainly produced by submerged macrophytes in freshwater and marine environments (Ficken et al., 2000), and have characteristic C and H signatures indicative of their aquatic origins (Aichner et al., 2010, 2017). H isotopes of *n*-alkanes longer than C_{21} reflect plant growth water at the time of leaf formation, making them

95   useful for paleohydrology studies (e.g., Sternberg, 1988; Sachse et al., 2012; McFarlin et al., 2019). Shorter-chain *n*-alkanes ($< C_{21}$) are principally produced by phytoplankton and bacteria (Meyers and Ishiwatari, 1993; Zhang and Sachs, 2007), and their H isotopic composition also correlates strongly with growth water (Sachse et al., 2004). Moreover, *n*-alkanes can be transported to sediments as aerosolized particles that are deposited directly in coastal waters and lakes or transported by rivers (Schefuss et al., 2005). The spatial coverage of this atmospheric transfer is dependent upon climate conditions and density of

vegetation, but *n*-alkanes are generally deposited within weeks after being aerosolized (e.g., Nelson et al., 2018), making them ideal for paleoenvironmental reconstructions.

In addition to *n*-alkanes, other lipid biomarkers – such as glycerol dialkyl glycerol tetraethers (GDGTs) and long-chain diols (LCDs) – can provide information about the environmental conditions under which they were produced. An abundance of branched GDGTs (brGDGTs) reflects bacterial communities – believed to be Acidobacteria – in soils (Weijers et al., 2007, 2009; Sinninghe Damsté et al., 2018), thus should be representative of terrestrial environments or terrestrial input at aquatic sites. However, aquatic in situ production complicates the strictly terrestrial interpretation of brGDGTs (De Jonge et al., 2014, 2015). Luckily, the process is characterized by a number of indices that distinguish between allochthonous and autochthonous production (Hopmans et al., 2004; Sinninghe Damsté, 2016; Xiao et al., 2016; Martin et al., 2020). LCDs are synthesized by both marine and freshwater phototrophic organisms (Volkman et al., 1992), and have been used to trace freshwater input into coastal environments using one isomer, the C_{32} 1,15-diol (Lattaud et al., 2017). LCDs are also used to reconstruct sea surface temperature via the long-chain diol index (LDI, Rampen et al., 2012; de Bar et al., 2020). In addition, the presence of 1,14- and 1,12-diols are characteristic of diatom abundance, more precisely of the presence of *Proboscia* spp. (de Bar et al., 2020; Rampen et al., 2007, 2014). The presence of *Proboscia* diatoms has been shown to hinder LDI temperature reconstructions due to their contribution to the pool of 1,13-diols used in the index (de Bar et al., 2020). At a minimum, pairing brGDGT with LCD data can contribute to understanding of complex riverine transport of organic matter and soil erosion in a given location (Lattaud et al., 2017), and can help reconstruct sea surface temperature in some environments.

Recently, attempts to understand the complex salinity history of the Baltic Sea using H isotope values of long-chain alkenones highlighted hydrological shifts that did not coincide with a change in the Haptophyte population in the Early Holocene (Weiss et al., 2020). Long-chain alkenones are a species-specific biomarker produced by haptophyte algae that have been widely used to discern changes in sea surface temperature and salinity (e.g., Brassell et al., 1986; Weiss et al., 2019). This observation, in conjunction with previous studies of Baltic Sea climate history, serves as the basis for the present investigation, which aims to compare information gathered from a variety of lipid biomarkers in order to delineate regional effects of large-scale hydrological changes. We compare the biomarker data generated here with pollen records from regional lakes in Sweden and Finland (Fig. 1).

2 Methods

Expedition 64PE410 aboard the RV *Pelagia* retrieved a 12 m-long piston core (10 cm in diameter) from the Arkona Basin (64PE410_S7, 54°55.208 N, 13°29.992 E, Fig. 1) in 2016 to study the climate history of the marginal marine environment of the Baltic Sea. The Arkona Basin (water depth ~45 m) is situated in the western part of the Baltic Sea, where saline North Sea water enters into the basin (Fig. 1). The geochemical composition of the core was analyzed by X-ray fluorescence (XRF) using an Avaatech XRF core scanner with a 100W rhodium X-ray tube and a Rayspec cubed SiriusSD silicon drift detector.

Continuous measurement of core sections was conducted at 1 cm resolution. The spectral data was processed using bAxil spectrum analysis software to determine the element intensities in counts.

The age model was created by combining ^{14}C-ages of mollusk shells and correlation of Ca/Ti and Br records with two nearby cores (318310: 54°50.34' N; 13°32.03' E, and 318340: 54°55.77' N; 13°41.44' E , Warden et al., 2016, Fig. 1). Bromine, a good indicator of marine organic carbon (Ziegler et al., 2008) was correlated with the total organic carbon content determined by loss on ignition (LOI) in core 318310 (Warden et al., 2016). Five shifts in Arkona Basin sediments are clearly distinguishable in the carbonate (Ca/Ti) and organic matter (Br and LOI) contents of the two sediment cores from Warden et al. (2016) and described in detail in Weiss et al. (2020). Three mollusk shells were radiocarbon dated by Beta Analytic to improve age control for the interval younger than 7 ka. Radiocarbon ages were corrected for the local reservoir effects (reservoir age = 376 years, Lougheed et al., 2013) and calibrated using CALIB (Stuiver et al., 2021). The final age model for all depths was created using the Bacon software package in R (Blaauw & Christen, 2011). Error estimates for ^{14}C ages were also used for the XRF tie-points (as these are based on correlation to the nearby ^{14}C dated cores). In Bacon, student's t-test values were set to 33 and 34 to allow for more narrow error estimates as suggested in the Bacon manual.

2.1 Bulk isotope analyses

Samples of bulk sediment were taken every 100 cm at the top of the core and every 10 cm after 930 cm and freeze-dried prior to analyses. An aliquot of 10 to 15 mg of freeze-dried sediment was sampled for total organic carbon (TOC), total nitrogen (TN), and bulk carbon (δ^{13}C) and nitrogen (δ^{15}N) isotope composition. Aliquots were acidified with 2 M hydrochloric acid and rinsed with deionized water to remove carbonates and salts. Isotopic composition and elemental weight percentage of bulk organic matter were measured with an elemental analyzer (EA, Thermo Scientific Flash 2000) coupled to an isotope ratio mass spectrometer (Delta V IRMS, Thermo Scientific).

2.2 Lipids

Samples for lipid analyses were taken at 100 cm intervals at the top of the core and every 10 cm after 930 cm to aim for a high resolution of the marine transgression and extracted as described in Weiss et al. (2020). Briefly, freeze-dried sediment samples (11 – 17 g) were extracted via an accelerated solvent extractor (ASE 350, Dionex, Thermo-Scientific, Sunnyvale, CA, USA) with dichloromethane (DCM) : methanol (MeOH) 9 : 1 (v / v). Internal standards – C_{46} GDGT (Huguet et al., 2006) C_{22} 7,16-diol, C_{19} ketone, and C_{36} n-alkane – were added to all total lipid extracts. The extracts were dried under a stream of N_2 and separated over an aluminum oxide column (activated for 3 h at 150 °C). An apolar fraction (containing n-alkanes) was eluted with four column volumes of hexane : DCM 9 : 1 (v / v), a ketone fraction with three column volumes of hexane : DCM 1 : 1 (v / v), and the polar fraction (containing glycerol dialkyl glycerol tetraethers, GDGTs, and long-chain diols, LCDs) with three column volumes of DCM : MeOH 1 : 1 (v / v).

2.2.1 *n*-Alkanes

The *n*-alkanes were quantified based on a known concentration of a C_{36} *n*-alkane standard on a HP 7890A gas chromatograph (GC) equipped with a flame ionization detector (FID) and a VF-1 ms capillary column (Agilent, 30 m × 0.25 mm × 0.25 µm). The temperature program started with a 1 min hold at 50 °C, followed by a 10 °C min^{-1} ramp to 320 °C, and held at 320 °C for 5 min. Reported concentrations are normalized to TOC content (ng g_{TOC}^{-1}).

2.2.2 Long-chain diols (LCDs)

Aliquots of the polar fraction were derivatized using N,O-Bis(trimethylsilyl)trifluoroacetamide (BSTFA) and pyridine (10 µL each) at 60 °C for 30 min, after which ethyl acetate was added. Long-chain diols (LCDs) were analyzed following Rampen et al. (2012) by GC (Agilent 7990B GC) coupled to a mass spectrometer (Agilent 5977A MSD; GC-MS) equipped with a fused silica capillary column (Agilent CP Sil-5, 25 m × 0.32 mm × 0.12 µm). The temperature program was as follows: started at 70 °C, increased to 130 °C at 20 °C min^{-1}, increased to 320 °C at 4 °C min^{-1}, held at 320 °C for 25 min. Flow was held constant at 2 mL min^{-1}. The LCDs were identified via single ion monitoring (SIM) of the $m/z = 299.3$ (C_{28} 1,14-diol), 313.3 (C_{28} 1,13-diol, C_{30} 1,15-diol), 327.3 (C_{28} 1,12-diol, C_{30} 1,14-diol), and 341.3 (C_{30} 1,13-diol, C_{32} 1,15-diol) ions (Versteegh et al., 1997; Rampen et al., 2012; de Bar et al., 2020). LCDs were quantified based on a known concentration of the C_{22} 7,16-diol.

2.2.4 Glycerol dialkyl glycerol tetraethers (GDGTs)

For GDGT analyses, a second aliquot of the polar fraction – at a concentration of 2 mg mL^{-1} – was dissolved in hexane : isopropanol (IPA) 99:1 (*v / v*) and analyzed using ultra high performance liquid chromatography mass spectrometry on an Agilent 1260 UHPLC coupled to a 6130 Agilent MSD following Hopmans et al. (2016). The Branched and Isoprenoid Tetraether (BIT) index (Hopmans et al., 2004) was calculated as follows:

$$BIT = \frac{GDGT-I + GDGT-II + GDGT-III}{GDGT-I + GDGT-II + GDGT-III + crenarchaeol} \tag{1}$$

where roman numerals refer to the number of methyl groups attached to the 5 or 5' positions of the alkyl chain (zero, one, and two, respectively). GDGT concentrations were calculated based on a known concentration of the C_{46} GDGT standard (Huguet et al., 2006).

The #ring$_{tetra,}$ used to assess the source of brGDGT in the marine water column, where values higher than 0.7 indicate sedimentary in situ production (Sinninghe Damsté, 2016) was also calculated. #ring$_{tetra}$ is defined as follows:

$$\#ring_{tetra} = \frac{GDGT-Ib + 2 \times GDGT-Ic}{GDGT-Ia + GDGT-Ib + GDGT-Ic} \tag{2}$$

The roman numeral I refers to the number of methyl groups, as indicated for the BIT index. The letters 'a', 'b', and 'c' refer to the number of cyclopentane moieties (zero, one, and two, respectively).

2.3 Compound-specific isotope analyses

190 Compound–specific carbon isotope analyses of n-alkanes ($\delta^{13}C_{alkane}$) were performed in duplicate on a Thermo Trace GC (1310) coupled with a Thermo Delta V plus IRMS at the Climate Geology Department of the ETH Zurich. The GC was equipped with a RTX-200 60 m capillary column (Restek, 60 m × 0.25 mm × 0.25 µm). The temperature program was as follows: ramp from 40 °C to 120 °C at 40 °C min^{-1}, followed by a 6 °C min^{-1} ramp to 320 °C, held at 320 °C for 12 min.

Hydrogen isotope ratios of n-alkanes (δ^2H_{alkane}) were measured at the Royal Netherland Institute for Sea Research in duplicate
195 by GC coupled to an IRMS via a high temperature conversion reactor. The GC was equipped with a CP Sil-5 column (Agilent, 30 m × 0.25 mm × 0.25 µm). The temperature program was as follows: started at 70 °C, increased to 130 °C at 20 °C min^{-1}, increased to 320 °C at 4 °C min^{-1}, held at 320 °C for 10 min. At the start of each day, the H_3^+ factor was measured and corrected for (Sessions et al., 2001). H_3^+ values ranged from 4.3 to 4.6. Prior to measurement of samples for both C and H isotopes, a n-alkane mixture (Mix B, supplied by A. Schimmelmann, Indiana University) was measured. Sample analyses were only
200 conducted when the average difference and standard deviation between online and offline values was less than 5‰ for H and 0.5‰ for C. For both C and H isotope measurements, the reported standard error is based on duplicate analyses.

3 Results

The core covers the last 10.6 ka with average sedimentation rates between 55 and 250 cm ky^{-1} (Weiss et al., 2020). The large
205 variations in sedimentation rates for core 64PE410-S7, and the nearby cores to which it has been correlated, are likely related to the shallow water depth in the Arkona Basin (~ 45 m) as well as changes in connection with the North Sea. Bulk isotopes ($\delta^{13}C_{bulk}$ and $\delta^{15}N_{bulk}$ values, as well as TOC and TN content), lipid concentrations, and compound-specific n-alkane isotope analyses were conducted on 22 samples.

3.1 Bulk isotopes

210 TOC and TN ranged from 0.5 to 7.2 and 0.1 to 0.8 %, respectively, with higher values in the most recent part of the core (Fig. S1). $\delta^{13}C_{bulk}$ values varied by ~3‰ in the Early Holocene (before 7.7 ka) followed by a small shift after the marine transgression (Fig. S1). $\delta^{15}N_{bulk}$ values were elevated at the transition from the YS to the AL (5.5‰) followed by a drop to 2.8‰ in the early AL phase (10.1 ka). Values reached 4‰ during the latter half of the AL, succeeded by a second drop at the marine transgression (2.8‰). Values slowly increased during the early LS phase. Values of 4 ± 1‰ occurred across the MB phase (Fig. S1).
215 Elemental C and N (C/N) ratios ranged from 5.6 to 13.4, spanning common values for both marine and terrestrial ecosystems (Thornton and McManus., 1994). C/N ratios are plotted against $\delta^{13}C_{bulk}$ values for all samples; a mix of freshwater, marine, and terrestrial C_3 plant-derived values are recorded, which group by established Baltic Sea phases (Fig. 2).

3.2 n-alkane distributions and isotopic composition

Organic carbon-normalized concentrations of n-alkanes (C_{21} to C_{35}) were highest in the older part of the record (2.8 μg g_{TOC}^{-1}

220 at 10.6 ka, Fig. 3), followed by a decrease until 9.4 ka (with a notably low concentration at 10.2 ka). The relative abundance of n-alkane homologues is comparatively unchanged across the remainder of the record (Fig. 3), except for a slight increase in the abundance of the C_{21} n-alkane following the marine transgression and absence or low abundance of the C_{17} n-alkane during the AL. The most abundant n-alkane is the C_{29} (21 ± 4 %, n=23), followed by the C_{27} (19 ± 2 %, n=23), C_{25} (16 ± 3 %, n=23) and C_{31} (16 ± 3 %, n=23), suggesting a primarily terrestrial signal. C and H isotope values are plotted as weighted averages of

225 short-chain (C_{21}), mid-chain (C_{23} to C_{25}), and long-chain (C_{27} to C_{31}). Both $\delta^{13}C_{alkane}$ and δ^2H_{alkane} values captured substantial fluctuations during the early part of the record (before ca. 7.7 ka), but both show relatively stable signals following the marine transgression (Tables 2 and 3; Figs. 4 and 5). XRF data for this portion of the core also suggests stable conditions. Weighted-average $\delta^{13}C$ values of the mid- and long-chain n-alkanes were similar from 10.6 to 8 ka, after which the mid-chain n-alkanes shifted toward higher values and the long-chain n-alkanes remained similar (Fig. 4). The LEaf-Wax Isotopic Spread (LEWIS)

230 index reflects changes in plant species diversity (Magill et al., 2019) and is calculated as follows:

$$LEWIS = \max \delta^{13}C_{23-31} - \min \delta^{13}C_{23-31} \qquad (3)$$

Values varied from 2.0 ± 0.7 ‰ between 11 – 7.7 ka to 6.7 ± 1.4 ‰ between 7.3 – 1.3 ka (Fig. 6a). H isotope values of short-, mid-, and long-chain n-alkanes tracked the same trends across the record. The δ^2H values of all n-alkanes were higher in the oldest part of the record (YS, prior to 10.6 ka), and recorded a rapid decrease of ~50‰ at the onset of the AL phase (Fig. 5).

235 Subsequently, a second, considerable increase of ~50‰ occurred between 10 ka and 9.2 ka, succeeded by a gradual decrease lasting until the marine transgression at 7.7 ka. δ^2H values showed higher values across the marine transgression. Data is limited for the LS phase, but no large isotopic shifts were noted in the samples analyzed here.

3.3 Abundance of LCDs and BIT index

Low concentrations of LCDs were detected in a majority of samples prior to the marine transgression. Furthermore, no 1,12-

240 diols were detected before the MB phase indicating limited input from *Proboscia* diatoms (de Bar et al., 2020). The C_{28} 1,12- and C_{28} 1,14-diols were present during the MB phase (in relatively high amounts for the C_{28} 1,14-diol). For all other LCDs, the main change in abundance (relative to all LCDs) occurred at 7.9 ka with a drop in the fractional abundance of the C_{32} 1,15-diol ($F_{C32\ 1,15\text{-diol}}$, from 20 to 1%) and an increase in $F_{C30\ 1,15\text{-diol}}$ (from 30 to 80%). LDI-temperature reconstructions (Table S1) were similar to those of Kotthoff et al. (2017) with relatively high temperatures during the YS (18.5 ± 0.2 °C) and a decrease

245 during the AL (11.5 ± 2.4 °C). Maximum temperatures occurred during the LS phase, in agreement with the Holocene thermal maximum in Europe (21.5 ± 1.5 °C, 24 °C at 7.2 ka). Temperatures during the MB were similar to modern summer temperatures (17.2 ± 0.1 °C). The coldest period (AL) is also the period with the highest proportion of C_{32} 1,15-diol. However, as noted by De Bar et al. (2020) and Kothoff et al. (2017), the absolute LDI-reconstructed temperatures are likely affected by the shifts between freshwater and marine environments that impact the composition of diol-producers (e.g., an increase in diol-

250 producing diatoms at higher salinities or freshwater diol-producers at lower salinities), and should be taken with caution. For

this reason, we use the LCDs primarily as indicators of continental runoff into the basin rather than for absolute temperature reconstructions.

The distribution of brGDGTs changed drastically across phases (Fig. S2) with a distribution similar to soils for all phases except the AL, which suggests mixing between lacustrine and terrestrial brGDGTs. BIT values were higher in the older part
255 of the record (0.8 before 9.9 ka), then continuously decreased to 0.1 by 6.5 ka. Similar BIT values were calculated for all LS samples (Fig. 6). #ring$_{tetra}$ was lower than 0.7 before the transgression (0.5 ± 0.1). A rapid increase occurred at the transgression (~7.7 ka) and values remained consistently high afterwards (0.7 ± 0.1, Table S2). Due to the high BIT values in the older part of the record, as well as the high #ring$_{tetra}$ after 7.7 ka, temperature reconstructions using the TEX$_{86}$ index are unreliable (Sinninghe Damsté, 2016).

260 **4 Discussion**

Compilation of biomarker distributions with corresponding *n*-alkane C and H isotope compositions provide new insights into changes in the regional hydrology and vegetation in the western Baltic region throughout the Holocene. Isotope and biomarker data are discussed below by Baltic Sea phases.

4.1 Yoldia Sea

265 The record covers only a brief snapshot of the end of the YS phase, a period in Baltic Sea history thought to have higher salinity because of the presence of marine phytoplankton species (Sohlenius et al., 1996). The appearance of short-chain C$_{17}$ – C$_{21}$ *n*-alkanes is suggestive of aquatic in situ production as the C$_{17}$ *n*-alkane in particular is produced by algae (Gelpi et al., 1970; Meyers and Ishiwatari, 1993). The only LCDs present are the C$_{30}$ 1,13-, C$_{30}$ 1,15-, and the C$_{32}$ 1,15-diols. The C$_{32}$ 1,15-diol, produced by eustigmatophyte species in rivers and stagnant freshwater pools (e.g., Lattaud et al., 2017; Balzano et al.,
270 2018), implies low salinity to freshwater conditions. LDI values suggest high SST (the warmest of the record, n = 2), similar to modern summer SST, which is in agreement with previous studies (Borzenkova et al., 2015; Kothoff et al., 2017). Corresponding δ^{13}C$_{bulk}$ values versus C/N ratios for this phase fall just on the edge of values common for freshwater ecosystems (Fig. 2), and along with the C$_{32}$ 1,15-diol, suggest that salinity of the basin may have already significantly declined by the end of the YS phase. This aligns with other records that divide the YS into two parts: an early marine phase followed by a freshwater
275 phase (e.g., Sohlenius et al., 1996). The highest *n*-alkane concentrations of the whole record, together with highest δ^{2}H$_{alkane}$ values, occur just at the phase boundary (Fig. 3 and 5). The increase in δ^{2}H$_{alkane}$ values varies from around 12‰ (C$_{21}$ and C$_{23}$ *n*-alkanes) to 47‰ (C$_{29}$ *n*-alkane), and may be linked to regional climate conditions – like regional warming that caused ice melt and continental uplift – which drove the shift from the YS phase into the low salinity AL phase.

4.2 Ancylus Lake
280 The AL phase, from 10.6 to 7.7 ka, was the most dynamic of the whole record and includes a number of large isotopic shifts. C isotope compositions of mid- and long-chain *n*-alkanes follow similar trends during the AL. A series of C and H isotope

fluctuations were recorded during this phase which can be linked to changes in regional hydrology and vegetation discussed in detail below.

4.2.1 Hydrology

285 At 10.2 ka, Moros et al. (2002) suggest that a large regression occurred, a likely consequence of continental uplift that resulted in shallowing and freshening of the basin. Low concentrations of n-alkanes (~150 ng g_{TOC}^{-1}, Fig. 3) occurred at the AL regression, implying less continental runoff into the Arkona Basin at this time. Simultaneously, there was a 15 – 20‰ increase in δ^2H_{alkane} values (Fig. 5). Increased δ^2H values are common when evaporation exceeds precipitation (Craig and Gordon, 1965). Such evaporative conditions may have occurred when the basin was more restricted as precipitation is thought to be

290 low at this time (Dahl and Nesje, 1996; Seppä and Birks, 2001; Hammarlund et al., 2003). Following the Ancylus Regression, δ^2H_{alkane} values become lower, coincident with a rapid, order of magnitude increase in concentration of n-alkanes (Fig. 3 and 5). The SIS was retreating at this time (Muschitiello et al., 2015; Cuzzone et al., 2016), thus it is plausible that a meltwater pulse transported a higher concentration of n-alkanes from the north into the basin just after 10.2 ka. Additionally, the availability of meltwater for plant growth during this period can explain the large change to lower δ^2H_{alkane} values. A coincident

295 increase in the BIT index from 0.5 to 0.8 (Fig. 6b) and #ring$_{tetra}$ values below 0.7 (suggesting a lack of in situ production, Sinninghe Damsté, 2016), strengthen the argument for increased input of allochthonous material into the basin. The higher proportion of GDGT-III (Table S2) is indicative of combined in situ brGDGT production and permafrost or soil erosion (Russell et al., 2018; Warden et al., 2018; Kusch et al., 2019), likely the latter in this case. No major changes appear to have occurred in the water column, but progressive melting of the SIS likely transported soil (terrestrial) brGDGTs resulting in the

300 increased BIT index values. Furthermore, a peak in $F_{C32\ 1,15\text{-}diol}$ to around 50% signals increased freshwater runoff (or a decrease in other diols) as the C_{32} 1,15-diol originates from the low flow area of freshwater systems (ponds and lakes; Lattaud et al., 2018; Häggi et al., 2019). The high abundance of C_{32} 1,15-diol hinders the LDI-temperature reconstruction which results in very low SST estimates (11.5 ± 2.4 °C, n = 25), while pollen data indicates a steep warming in air and water temperature (Borzenkova et al., 2015). The combined increase in n-alkane concentrations, BIT index, and $F_{C32\ 1,15\text{-}diol}$ points towards

305 increased runoff from the continent. Additional evidence for increased terrestrial input comes from a concurrent, rapid decrease in $\delta^{13}C_{bulk}$ values (1.5‰) in parallel with a large increase in C/N ratios (+6%) and high TOC values (+4%) just after the Ancylus Regression. Altogether, an increase in terrestrial-derived organic matter either from enhanced coastal erosion or riverine input, both resulting from the melting of the SIS, occurred at this time.

A significant increase in δ^2H_{alkane} values (~ 50‰) began after 10 ka and peaked at 9.2 ka (Fig. 5), precisely at the time when a

310 similar increase in δ^2H values of haptophyte-derived alkenones from the same core is noted (Weiss et al., 2020). The change in alkenone δ^2H values occurred when alkenone distributions were constant, suggesting that environmental factors rather than species related effects were the cause of the isotopic shift. The H isotope composition of terrestrial plants reflects local meteoric water signals (Sachse et al., 2012), and is thus subject to large fluctuations resulting from variations in the hydrological cycle. H isotope values of precipitation were reconstructed from C_{23} and C_{29} n-alkanes using linear models from McFarlin et al.

315   (2019), and range from -125‰ to -62‰ (residual standard error = 14) and -107‰ to -36‰ (residual standard error = 15), respectively, across the AL phase (Fig. 5). The lowest δ^2H_{alkane} values – and in turn the lowest δ^2H values of reconstructed precipitation – occur immediately preceding the 50‰ increase (Fig. 5). Reconstructed δ^2H values of precipitation using the C_{23} n-alkane (originating from an aquatic source) are lower than those for the C_{29} n-alkane (originating from terrestrial vegetation), suggesting an influence from an isotopically lighter (more negative) water source affected isotopic signatures of

320   aquatically-produced n-alkanes. Model data suggests that the H isotope composition of the SIS was around -310‰ (De Boer et al., 2014), and modern mean annual precipitation in the region is -63‰ (Bowen and Revenaugh, 2003), highlighting the marked isotopic offset between the two sources. Larger volumes of meltwater from the SIS would result in significantly more negative δ^2H values for aquatically-derived mid-chain n-alkanes, in line with what is observed here at around 10 ka (Fig. 5) and also recorded by haptophyte algal biomarkers from the same core (Weiss et al., 2020). The depleted n-alkane values imply

325   that terrestrial plants may have relied more heavily on glacial melt water as a water source during this period. After 10 ka the decreased influence of melt water and increase in precipitation in the region (Dahl and Nesje, 1996; Seppä and Birks, 2001; Hammarlund et al., 2003) likely caused the dominant water source for aquatic and terrestrial plants to become isotopically heavier, resulting in increased n-alkane H isotope values.

Based on these interpretations, we suggest division of the AL into two parts to highlight the dynamicity of the phase: an earlier

330   ice melt subphase followed by a precipitation-driven subphase. The ice-melt subphase is the older portion of the AL, from 10.6 to 9.2 ka, where the SIS provided glacial melt water as an isotopically light water source. After 9.2 ka, precipitation became the dominant, isotopically heavier water source resulting in higher H isotope values preserved in organic compounds.

4.2.2 Vegetation

The n-alkane distributions and C isotope signatures imply largely terrestrial, higher plant input during the AL, supported by

335   the absence of the aquatically produced C_{17} n-alkane during most of this phase (Fig. 3). The LEWIS index values are the lowest of the record (Fig. 6a), suggesting limited or low species diversity (Magill et al., 2019). This is supported by nearby pollen records that indicate an absence of higher plant diversification, likely due to the relatively cold temperatures in Northern Europe at this time (Seppä and Birks, 2001; Seppä et al., 2005; Antonssson et al., 2006, Berglund et al., 2008). In contrast, the $\delta^{13}C$ values of C_{21} n-alkane were higher compared with the longer-chain (> C_{27}) n-alkanes (-29 versus -21‰ before 7.8 ka),

340   suggestive of mixed terrestrial and aquatic production or increased aquatic productivity (Ficken et al., 2000).

C isotope values of all n-alkanes record a decrease at the time of the AL regression and LEWIS index values are low, indicating particularly low species diversity. C/N values of bulk sediments suggest a stronger C_3 plant signal at this time implying terrestrial plants were the dominant producers of n-alkanes. Following the Ancylus Regression, there were two shifts in mid- and long-chain n-alkane C isotope signatures: an increase (+2‰) around 9.2 ka, followed by a return to lower values, and a

345   second increase (again +2‰) at the onset of the marine transgression (Fig. 4). These shifts in $\delta^{13}C_{alkane}$ values during the AL phase might reflect the advance and retreat of coniferous cover because conifers, such as *Pinus* and *Juniperus*, have higher

$\delta^{13}C_{alkane}$ values compared to deciduous angiosperm trees (Diefendorf and Freimuth, 2017). Since Pinaceae produce low concentrations of n-alkanes compared to *Juniperus* (Diefendorf et al., 2011; Lane, 2017), the increases in C isotope values can be tentatively attributed to *Juniperus* shrub expansion. These isotopic shifts align with the main climatic changes recorded in Swedish and Finnish lakes and note similar vegetation changes (Seppä and Birks, 2001; Seppä et al., 2005; Antonsson and Seppä, 2007; Berglund et al., 2008), but do not always find an increase in gymnosperms at 9.2 ka, which might be the result of low sampling resolution around that time period (Seppä and Birks, 2001; Seppä et al., 2005; Antonssson et al., 2006). A large increase in the angiosperm *Alnus* (which tends to have lower $\delta^{13}C_{alkane}$ values) was observed at 8.7 ka at the time of a decrease in *Juniperus* cover north of the Baltic Sea (Antonsson and Seppä, 2007), paralleling the return to lower $\delta^{13}C$ values before the second C isotope increase.

The low variability in n-alkanes C isotope values suggest limited sources of vegetation until the Marine Transgression. However, the data presented here suggests that expansion of *Juniperus* began to occur at around 9.2 ka when we detect a shift to a precipitation-dominated hydrological regime. Thus, the meltwater dominated portion of the AL is characterized by low species diversity and the shift to a precipitation-controlled climate coincided with the start of a transformation in regional vegetation.

4.3 Marine Transgression

A large marine transgression re-established the connection between the Baltic Sea and the North Sea at around 7.7 ka. At the onset of the transgression (which lasted from 7.7 to 7.2 ka, Warden et al., 2016), long-chain n-alkane ($> C_{27}$) concentrations increased and shorter-chain n-alkanes are detected ($< C_{19}$, Fig. 3), indicating greater input from both terrestrial and algal sources. Regional warming that began prior to the Holocene and caused the SIS to melt (Seppä and Birks, 2001; Seppä et al., 2005; Antonsson and Seppä, 2007) promoted the growth and diversification of terrestrial and aquatic plants. At the start of the transgression, mid- and long-chain n-alkane C isotope values began to diverge (LEWIS index increases, Fig. 6b), with the mid-chain n-alkanes becoming more positive by about 4‰ (Fig. 4). The $\delta^{13}C_{alkane}$ values of long-chain n-alkanes showed a C_3 higher plant signal (-30.4 ± 0.7‰), while the mid-chain n-alkanes were higher (-27.7 ± 0.4‰), characteristic of submerged aquatic plants (Ficken et al., 2000). The C_{21} n-alkane, produced by submerged aquatic plants as well as freshwater and marine phytoplankton, was the highest prior to the transgression, and became lower by ~5‰ across the transgression, despite the fact that large volumes of saline water entered the basin and the $\delta^{13}C_{bulk}$ values showed increased values. Terrestrial vegetation and freshwater organisms generally have lower C isotope signatures than marine organisms (Fig. 2), so the more negative values noted for the C_{21} n-alkane might indicate an increase in the abundance of submerged aquatic plants and freshwater phytoplankton rather than the influx of saline water. The organisms producing the C_{21} n-alkane during this period may have been growing in freshwater pools in the catchment rather than in the Arkona Basin itself. Alternatively, salinity stratification may have occurred during the transgression allowing freshwater phytoplankton to thrive in a less saline surface water layer. An increase in δ^2H_{alkane} values also occurred across the transgression (except the δ^2H values of the C_{23} n-alkane which

decreased). The C$_{23}$ *n*-alkane is produced by *Sphagnum* species (Baas et al., 2000; Vonk and Gustafsson, 2009), and thus may

380 be reflecting a different water source than the rest of the *n*-alkanes. An alternative water source for *Sphagnum* may be stagnant meltwater ponds/peat bogs which would have very negative H isotope values. TOC and $\delta^{13}C_{bulk}$ values increased, while C/N ratios decreased, likely due to enhanced phytoplankton production (cyanobacterial blooms in particular, Bianchi et al., 2000), and amplified input of organic matter into the sediments from increased erosion and primary production (Sohlenius et al., 2001). The BIT index continued to decrease (Fig. 6b), while #ring$_{tetra}$ increased (> 0.7) suggesting a change in the main

385 brGDGT producers from (freshwater) lacustrine production towards (marine) sedimentary pore-water production. The decrease in BIT values at the transgression may be the result of a shift in freshwater versus marine producers rather than a decrease in soil-input. A rapid decrease in F$_{C32\ 1,15\text{-diol}}$ occurred from the end of the AL phase into the marine transgression (8.0 to 7.3 ka; Fig. 6b) and is likely linked to a change in the LCD-producing population from freshwater to mixed brackish and marine LCD-producing species – which synthesize the C$_{32}$ 1,15 diol in lower amounts – caused by increased salinity in the

390 basin.

4.4 Littorina Sea and Modern Baltic phases

The marine transgression is followed by the brackish Littorina Sea (LS) phase. The early part of the LS is characterized by variable δ^2H_{alkane} and $\delta^{13}C_{alkane}$ values (Figs. 4 and 5). Multiple studies (e.g., Hammarlund et al., 2003; Seppä et al., 2005; Antonssson et al., 2006; Antonsson and Seppä, 2007) indicate that the period between 8.0 and 4.2 ka was warm, dry, and

395 stable, followed by a period of rapid hydrological changes. This is supported by the LDI-reconstructed summer sea surface temperatures (Table S1), indicating an optimum between 7.3 and 5.3 ka (LS average of 21.5 ± 1.5 °C). During the LS, values are comparable with pollen records from southern Sweden (e.g., Seppä et al., 2005) and the LDI record of Kothoff et al. (2017). In addition, stable #ring$_{tetra}$, BIT, and F$_{C32\ 1,15\text{-diol}}$ suggest relatively unchanged bacterial communities and river inflow. However, the C and H isotope data indicate instability and diversification in the catchment from the marine transgression until

400 the middle of the LS phase. We note, however, that the rapid hydrological changes evidenced in other records and the variation suggested by isotopic data discussed here likely do not capture the nuances of the time period as a result of low sampling resolution. At the transition from the LS into the Modern Baltic phase, the C$_{28}$ 1,14- and C$_{28}$ 1,12-diols, characteristic of *Proboscia* diatoms (Rampen et al., 2007) are present. This aligns with data from Andrén et al. (2000), which indicated that freshwater diatoms were low throughout the LS phase, but were more prominent during the Modern Baltic phase. During the

405 MB, LDI-reconstructed SST is lower than during the LS (17.2 ± 0.1 °C, n = 2) following the regional trend in mean annual air temperature (Berglund et al., 2008a,b; Kothoff et al., 2017). However, diols from surface sediments analyzed in De Bar et al. (2020) and close by the core analysed here (54°53.50', 13°24.62') result in a reconstructed temperature of 12.87 °C, higher than the annual average and lower than the summer temperature (which is also seen for the other surface sediments in that study, n = 5), which indicates that reconstructing temperature using LDI in the MB may be unreliable, likely due to the

410 influence of diatoms and freshwater input.

5 Conclusion

The Holocene history of the Baltic Sea consisted of variations in regional vegetation and hydrology that were connected to global climate phenomena. The melting of the Scandinavian Ice Sheet caused the Ancylus Lake phase to be particularly dynamic. This time period was characterized by fluctuations in the extent of gymnosperm cover across the newly ice-free landscape and substantial shifts in source water from a significantly light water source primarily from ice melt to an isotopically heavier precipitation-dominated source once the SIS had melted. The change in source water was also observed by a previous study focusing on phytoplankton-derived biomarkers showing that the source water shift affected aquatic microscopic organisms and terrestrial higher plants alike. We suggest that future studies of the Ancylus Lake time period divide the phase into an early ice melt-derived source water subphase and a later precipitation-dominated subphase. The marine transgression that ended the low salinity Ancylus Lake phase was accompanied by increased heterogeneity of the terrestrial higher plant population as well as changes in coastal aquatic ecosystems. A large hydrological change and diversification of higher plants were offset from the previously established Baltic Sea phase boundaries based on salinity and eustatic changes. Use of multiple proxies is essential to better understand the complexity of paleoenvironments, and in this case isotopic evidence provided new insights into the hydrological change that occurred during a time period where the conditions in the Baltic Sea are otherwise believed to be stable.

Author contribution

GW and JL designed the study, extracted samples, conducted isotopic analyses, and contributed equally to writing of the manuscript with contributions from all co-authors.

Competing interests

The authors declare that they have no competing interests.

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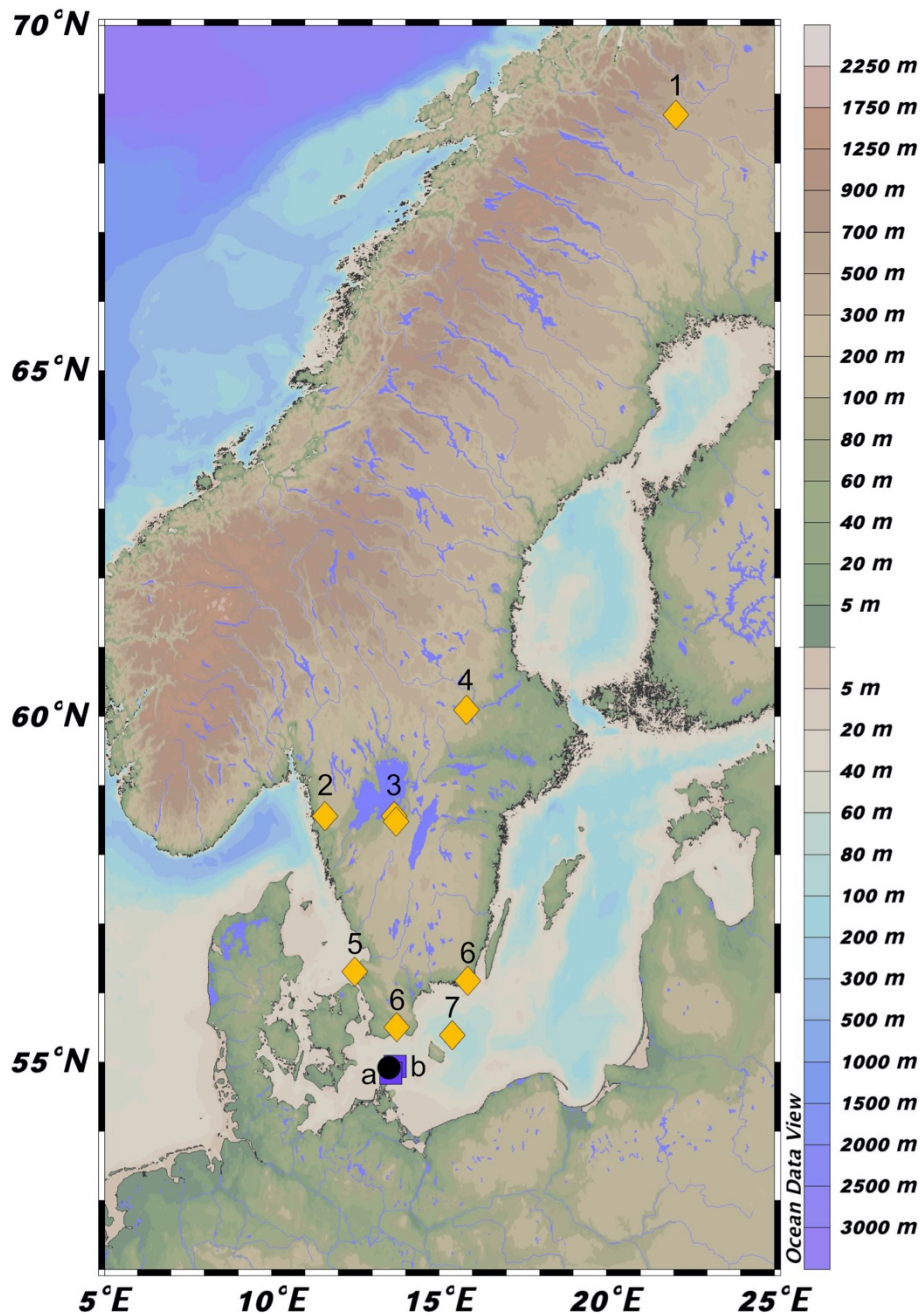
Figure 1: Location of 64PE410_S7 (black circle) in the Southwestern Baltic Sea near the connection with the North Sea via the Danish Straits. Map was created in Ocean Data View (Schlitzer, R., Ocean Data View, https://odv.awi.de, 2018). Squares a and b represent the location of cores from Warden et al. (2016) used to construct the age model. Yellow diamonds correspond to regional pollen records. 1 = Seppä and Weckström (1999), Seppä and Birks (2001); 2 = Antonsson and Seppä (2007); 3 = Hammarlund et al. (2003), Seppä et al. (2005); 4 = Antonsson et al. (2006); 5 = Berglund et al. (2008a); 6 = Berglund et al. (2008b); 7 = Andrén et al. (2000).
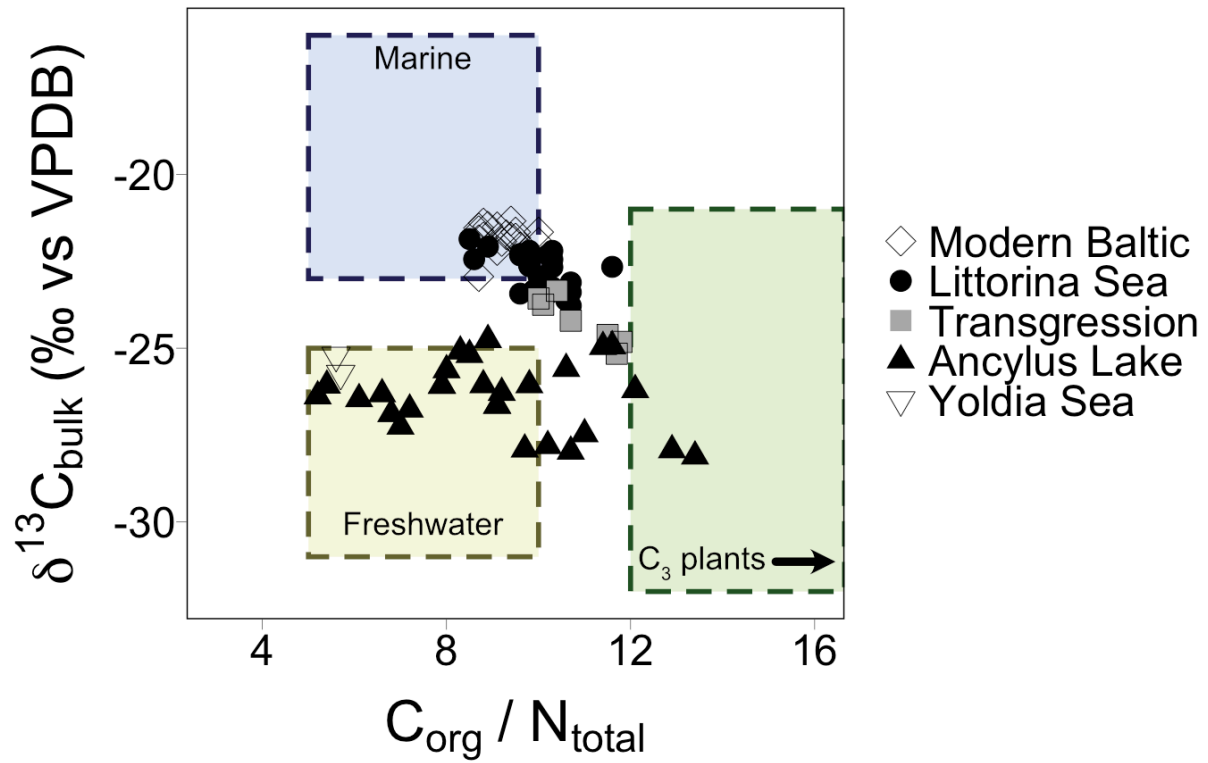
Figure 2: Elemental C/N ratios plotted against δ¹³C values of bulk sediment. Shapes correspond to different Baltic Sea phases and align with a mix of values corresponding to marine, freshwater, and C₃ plant signatures. Arrow in the C₃ plants box indicates that values for C₃ plants encompass a wider range of values which are excluded from this plot for clarity.

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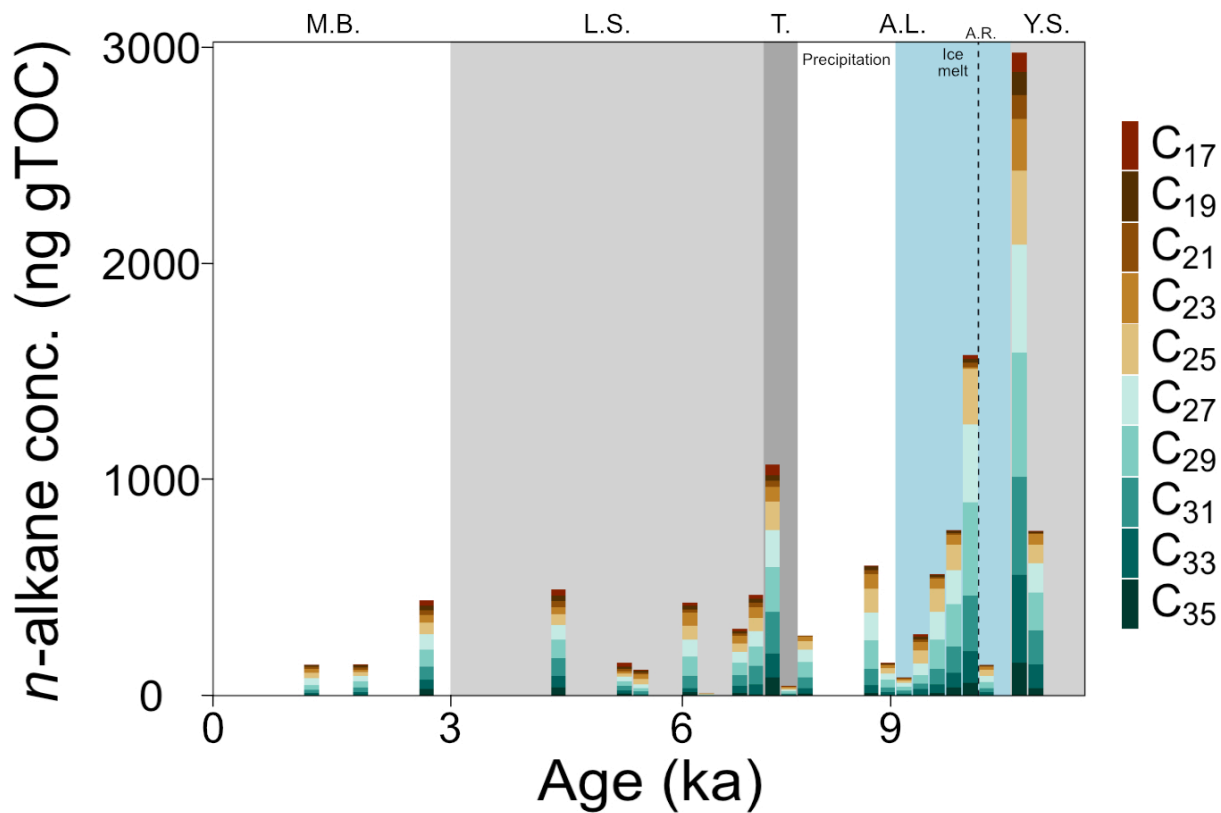
Figure 3: TOC weighted concentrations of $C_{17} - C_{35}$ *n*-alkanes in core 64PE410_S7 (n = 22). Concentrations were highest in the Yoldia Sea brackish phase and lowest in the Modern Baltic. Phases are abbreviated as follows: Y.S. = Yoldia Sea, A.L. = Ancylus Lake, T. = Marine Transgression, L.S. = Littorina Sea, M.B. = Modern Baltic. A. R. stands for Arkona Regression. New subdivisions for the AL phase showing the shift from ice melt-derived to precipitation dominated source water are shown in blue and white.

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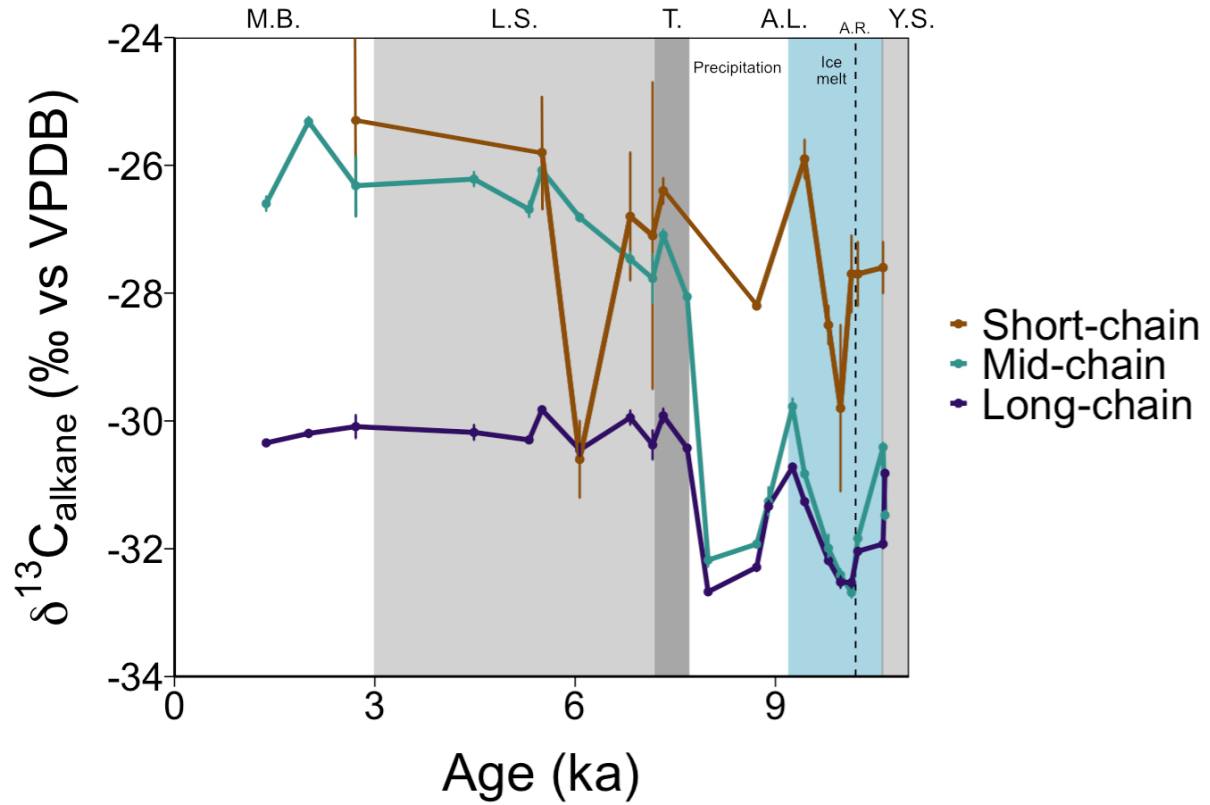
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Figure 4: Carbon isotope ratios of short-chain (C_{21}), mid-chain ($C_{23} - C_{25}$), and long-chain ($C_{27} - C_{31}$) *n*-alkanes from core 64PE410_S7. Phase boundaries are indicated as well as a regional climate event at 10.2 ka (A.R., = Ancylus Regression) and new subdivisions for the A. L. phase are shown in blue and white.
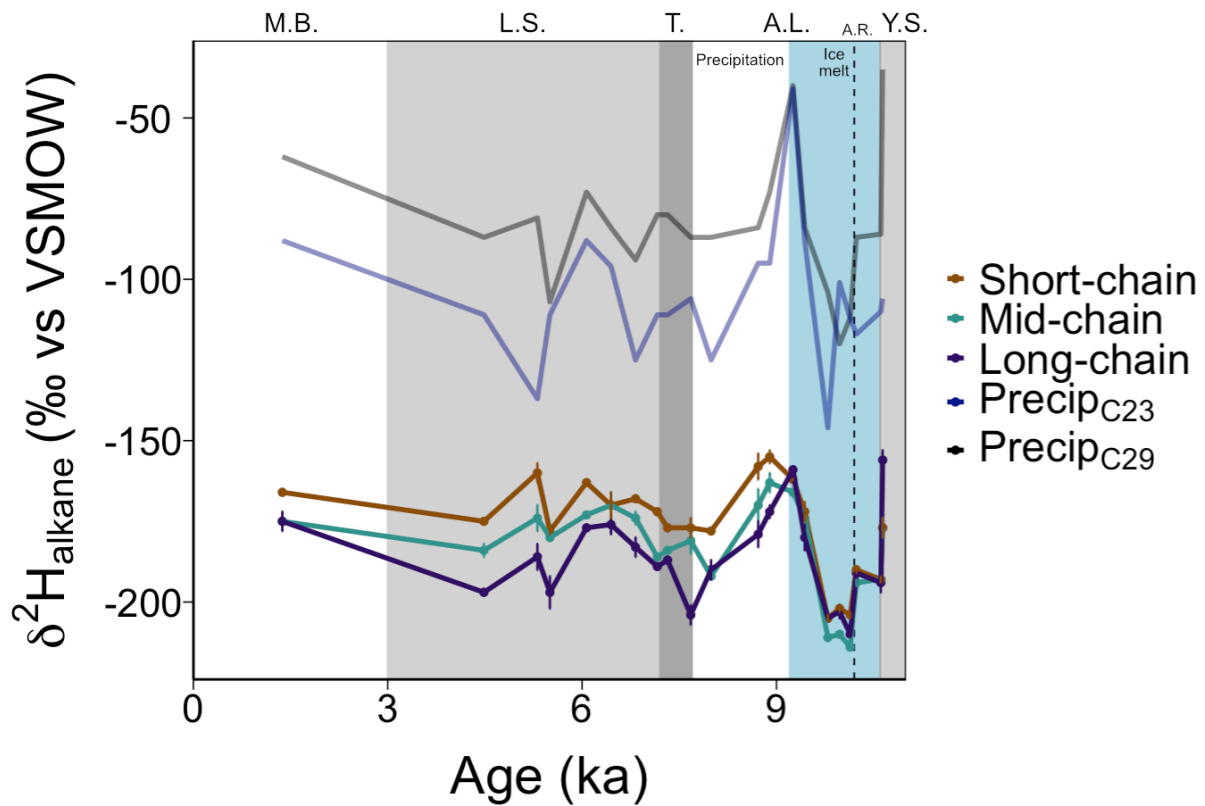
Figure 5: Hydrogen isotope ratios of short-chain, mid-chain, and long-chain $C_{21} - C_{33}$ *n*-alkanes from core 64PE410_S7. Phase boundaries are indicated as well as a regional climate event at 10.2 ka (A.R., = Ancylus Regression) and new subdivisions for the A. L. phase are shown in blue and white. Blue and black lines represent reconstructed δ^2H values of precipitation from linear models of McFarlin et al. (2019) using δ^2H values of C_{23} and C_{29} *n*-alkanes.
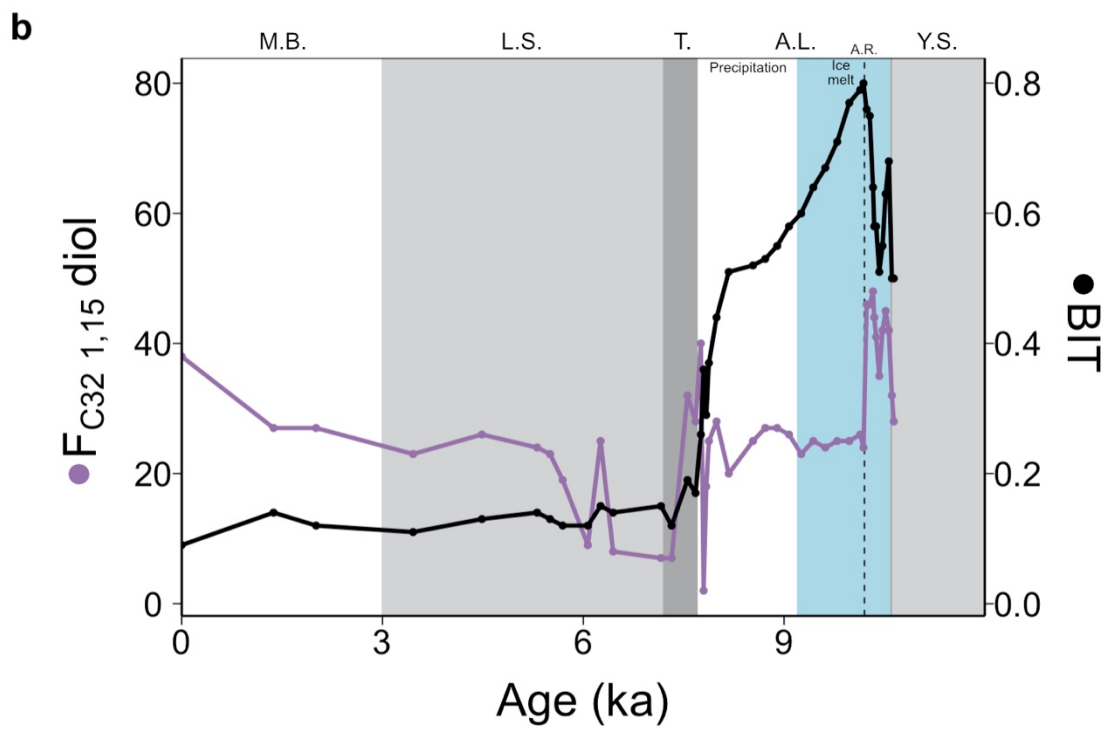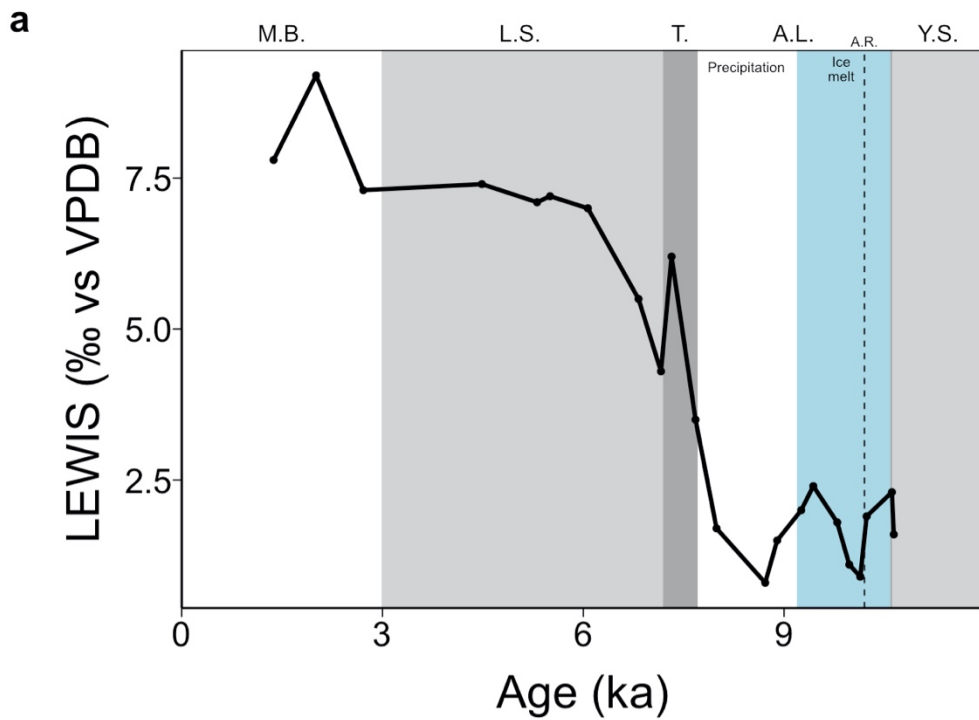
Figure 6: (a) The LEaf-Wax Isotopic Spread (LEWIS) index (Magill et al., 2019) shows a marked increase from the end of the Ancylus Lake phase into the Modern Baltic indicating increased species diversity in the younger half of the record. (b) Fractional abundance of C_{32} 1,15-diol ($F_{C32\ 1,15\text{-diol}}$, de Bar et al., 2016; Lattaud et al., 2017) and BIT index (Hopmans et al., 2004) for core 64PE410_S7, indicators of riverine and terrestrial runoff into the basin, suggest the highest terrestrial input occurred during the Ancylus Lake phase and decreased thereafter. Phase boundaries are indicated as well as a regional climate event at 10.2 ka (A.R., = Ancylus Regression) and new subdivisions for the A. L. phase are shown in blue and white.

Table 1: Summary of the proxies applied in this study, how they are used as paleoclimate indicators, and their source(s).

Organic compound	Uses	Source	References
Bulk C and N isotopes	Sources of organic matter	Organic matter	Meyers, 1994; Leng and Lewis, 2017
Short-chain *n*-alkanes (C_{17} - C_{21})	Reconstruction of vegetation (concentration and C isotopes); growth water, hydrology (H isotopes)	Phytoplankton, bacteria	Meyers and Ishiwatari, 1993; Zhang and Sachs, 2007; Sachse et al., 2012;
Mid-chain *n*-alkanes (C_{23} - C_{25})	Reconstruction of vegetation (concentration and C isotopes); growth water, precipitation (H isotopes)	Submerged macrophytes	Ficken et al., 2000; Sachse et al., 2012
Long-chain *n*-alkanes (> C_{27})	Reconstruction of vegetation (concentration and C isotopes); growth water, precipitation (H isotopes)	Terrestrial higher plants	Eglinton and Hamilton, 1967; Sachse et al., 2012; Freimuth et al., 2017
BIT index	Aquatic production and soil erosion	Heterotrophic bacteria	Hopmans et al., 2004; Weijers et al., 2007, 2009
C_{32} 1,15 diol	Riverine input	Marine and freshwater Eustigmatophytes	Volkman et al., 1992; Rampen et al., 2012; de Bar et al., 2016; Lattaud et al., 2017; Balzano et al., 2018;
LDI	Sea surface temperature	Marine and freshwater Eustigmatophytes	Rampen et al., 2012; de Bar et al., 2020
#ring$_{tetra}$	Sediment pore-water branched glycerol dialkyl glycerol tetraether (brGDGT) production	Heterotrophic bacteria	Sinninghe Damsté, 2016