

# 1 **Characterization of particulate organic matter in the Lena** 2 **River Delta and adjacent nearshore zone, NE Siberia. Part** 3 **II: Lignin-derived phenol compositions**

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## 17 18 **Abstract**

19 The Lena River in central Siberia is one of the major pathways translocating terrestrial  
20 organic matter (OM) from its vast catchment area to the coastal zone of the Laptev Sea and  
21 the Arctic Ocean. The permafrost soils of its far south stretching catchment, which store huge  
22 amounts of OM, will most likely respond differently to climate warming and remobilize  
23 previously frozen OM with distinct properties specific for the source vegetation and soil. To  
24 characterize the material discharged by the Lena River, we analyzed the lignin phenol  
25 composition in total suspended matter (TSM) from surface water collected in spring and  
26 summer, surface sediments from the Buor Khaya Bay along with soils from the Lena Delta's  
27 first (Holocene) and third terraces (Pleistocene ice complex), and plant samples. Our results  
28 show that lignin-derived cinnamyl:vanillyl (C/V) and syringyl:vanillyl (S/V) ratios are >0.14  
29 and 0.25, respectively, in TSM and surface sediments, whereas in delta soils they are >0.16  
30 and >0.51, respectively. These lignin compositions are consistent with significant inputs of

1 organic matter from non-woody angiosperm sources mixed with organic matter derived from  
2 woody gymnosperm sources. We applied a simple linear mixing model based on the C/V and  
3 S/V ratios and the results indicate the organic matter in delta TSM samples and Buor Khaya  
4 Bay surface sediments contain comparable contributions from gymnosperm material, which is  
5 primarily derived from the taiga forests south of the delta, and angiosperm material typical for  
6 tundra vegetation. Considering the small catchment area covered by tundra (~12%), the input  
7 is substantial and tundra-derived OM input is likely to increase in a warming Arctic. The  
8 similar and high acid to aldehyde ratios of vanillyl and syringyl ( $Ad/Al_{V,S}$ ) in Lena Delta  
9 summer TSM ( $>0.7$  and  $>0.5$ , respectively) and Buor Khaya Bay surface sediments ( $>1.0$  and  
10  $>0.9$ , respectively) suggest that the OM is highly degraded and Lena River summer TSM  
11 could be a possible source for the surface sediments. The  $Ad/Al_{V,S}$  ratios of the first and third  
12 delta terraces were generally lower (mean ratios  $>0.4$  and  $>0.4$ , respectively) than summer  
13 TSM and surface sediments. This implies that TSM contains additional contributions from a  
14 more degraded OM source (southern catchment and/or finer more degraded particle size).  
15 Alternatively, OM degradation on land after permafrost thawing and subaqueously during  
16 transport and sedimentation could be considerable. Despite the high natural heterogeneity of  
17 OM stored in delta soils and exported by the Lena River, the catchment characteristic  
18 vegetation is reflected by the lignin biomarker composition. Climate warming related changes  
19 in the Lena River catchment may be detectable in changing lignin biomarker composition and  
20 diagenetic alteration.

21

## 22 **1 Introduction**

23 Within the permafrost affected soils of the high northern latitudes lies a huge organic carbon  
24 (OC) reservoir, estimated to be as big as 1400-1850 Pg carbon representing about 50% of the  
25 global soil OC (Tarnocai et al., 2009). Currently most of this OC pool remains frozen and is  
26 therefore excluded from biogeochemical cycles. Over the last decades mean annual air  
27 temperatures in the Arctic increased more strongly than the global mean and this trend is  
28 projected to continue (IPCC, 2013). As a result annual permafrost thaw depths and arctic river  
29 runoff increase (McClelland et al., 2012; Peterson et al., 2002) likely leading to enhanced  
30 mobilization and export of old, previously frozen soil-derived OC (e.g. Guo et al., 2004;  
31 Schuur et al., 2008; Vonk et al., 2010). Consequently, the great arctic rivers play an important  
32 role in global biogeochemical cycles by connecting the large permafrost carbon pool of their  
33 hinterlands with the arctic shelf seas and the Arctic Ocean.

1 Terrigenous sediments reaching the nearshore zone and shelves serve as archives recording  
2 changes in material derived from river catchments and from erosion of permafrost coasts. The  
3 particulate organic matter associated with these sediments consists of a complex mixture of  
4 compounds from different aquatic and terrigenous sources with different chemical/physical  
5 recalcitrance towards decomposition and mineralization. Determining the sources (e.g.  
6 phytoplankton, vegetation, surface soil, mineral-associated soil, peat, etc.) and quality of OC  
7 transported by arctic rivers is therefore important to understand the effects of climate change  
8 on the river watersheds as well as on the arctic coastal zone.

9 Recent studies using molecular organic compounds and their carbon isotopes have shown that  
10 there are great differences in the age, quality, and source of OM exported by individual rivers  
11 (Dickens et al., 2011; Drenzek et al., 2007; Feng et al., 2013; Goñi et al., 2013; 2000;  
12 Gustafsson et al., 2011; Karlsson et al., 2011; Kuzyk et al., 2008; Unger et al., 2005; Vonk et  
13 al., 2010). The catchments of the great arctic rivers in North America and Siberia cover  
14 several climate zones. Their response to climate change will most likely vary strongly  
15 between the temperate and high latitude regions affecting river biogeochemical carbon  
16 cycling in different ways. Knowing where the OM derives from (southern vs. northern part of  
17 the catchment), if and how the relative contributions of climatic zones to riverine POC may  
18 change with climate warming, is important to understand and evaluate different permafrost  
19 thawing scenarios and their projected effect on the global climate.

20 Research efforts on studying arctic rivers increased in the last decades and the spatial and  
21 temporal data resolution on dissolved and particulate organic matter has improved.  
22 Nonetheless, the resolution is still relatively low, especially for riverine POC. The main  
23 reasons for that are the great logistical difficulties of conducting fieldwork in these remote  
24 arctic regions under mainly severe climate conditions, especially for winter and spring  
25 campaigns.

26 This is the first of two papers (see same issue) dealing with particulate organic matter from  
27 the Lena River Delta and adjacent Buor Khaya Bay. The Lena River is one of the biggest  
28 Siberian rivers in terms of water and sediment discharge and an important source of sediment  
29 as well as dissolved and particulate organic matter to the Laptev Sea and Arctic Ocean  
30 (Holmes et al., 2002; 2012; Rachold, 1999). In recent years, several studies have investigated  
31 the input, composition, and transport mechanisms of sediments delivered by the Lena River  
32 and by erosion of permafrost coasts (e.g. Charkin et al., 2011; Günther et al., 2013; Karlsson  
33 et al., 2011; Rachold and Hubberten, 1999; Semiletov et al., 2011). However, it is still under

1 debate how OM from the two main sources (riverine vs. coastal erosion) affects the total  
2 carbon budget and cycling in the Laptev Sea. Our samples were taken during field campaigns  
3 in the summers of 2009 and 2010 as well as in spring 2011. Here, we present new data on  
4 particulate OC composition and quality from riverbank soil profiles of the eastern Holocene  
5 first delta terrace and the Pleistocene third terrace of Kurungnakh Island (e.g. Schwamborn et  
6 al., 2002), surface water particulate matter along the main delta channels, and surface  
7 sediments from the Buor Khaya Bay. We used the lignin phenol composition to distinguish  
8 the sources of OM transported by the river, namely the taiga forest in the southern catchment  
9 versus the tundra covering the northernmost part of the watershed including the delta. The  
10 alkaline cupric oxide (CuO) oxidation products are also used to characterize the degree of  
11 aerobic degradation of lignin in these samples.

12 Lignin is a biopolymer produced almost exclusively by terrestrial vascular plants. Through  
13 CuO oxidation it is possible to break up the polymer structure and analyze the main building  
14 blocks, the lignin-derived phenols, as well as other CuO oxidation products by gas  
15 chromatography mass spectrometry (GC-MS). This method has been successfully applied in  
16 numerous studies to a variety of environments including the Arctic to trace soil-derived OM  
17 and differentiate between gymnosperm and angiosperm plants as well as between woody and  
18 non-woody tissues as sources (see Bianchi et al., 2007; Goñi et al., 2000; Hedges and Mann,  
19 1979; Kuzyk et al., 2008; Onstad et al., 2000; Opsahl et al., 1999; Prahl et al., 1994; Tesi et  
20 al., 2011). Furthermore, lignin is believed to be a rather recalcitrant fraction of soil organic  
21 matter, although this model is currently under debate (Feng et al., 2008).

22 Considering that, our study in the Lena Delta can serve as possible benchmark against which  
23 future changes in OM composition and quality associated with a warming Siberian Arctic  
24 could be assessed. Because of our sampling location in the delta covered by tundra vegetation  
25 we provide lignin compositional information from the Lena River including the whole  
26 catchment and compare these results with data from more southern Lena River sampling  
27 locations (e.g. Amon et al., 2012). Further, characterizing the riverine particulate organic  
28 matter can improve our understanding of organic matter delivery cycling in the near coastal  
29 zone of the Buor Khaya Bay and Laptev Sea.

30

## 1 **2 Material and Methods**

### 2 **2.1 Study Area**

3 The Lena River is one of the largest Russian Arctic rivers draining an area of  $\sim 2.46 \times 10^6$  km<sup>2</sup>  
4 in central Siberia (Fig. 1A). Its watershed stretches from 53°N near Lake Baikal to 71°N  
5 where the river discharges into the Laptev Sea and Arctic Ocean. Because of its huge  
6 extension, the Lena River basin comprises a diverse flora and fauna. In general, the basin can  
7 be divided into two major vegetation zones transitioning from south to north: 1) the boreal  
8 forest or taiga which covers about 72% of the watershed and 2) a small tundra zone in the  
9 north representing 12% of the basin area (the remaining area is categorized as water bodies,  
10 cropland, etc.; see Amon et al., 2012) consisting mainly of wet and dry dwarf-shrub tundra  
11 and sedge/grass wetland tundra (CAVM Team, 2003). About 90% of the Lena River  
12 catchment are characterized by continuous and discontinuous permafrost (72-80% and 6-10%  
13 of basin area, respectively; (Amon et al., 2012; Zhang et al., 2005). The permafrost table  
14 beneath the seasonally thawed layer (active layer) acts as water-impermeable layer and thus  
15 its distribution has a large impact on regional hydrology and hydrochemistry. Because of the  
16 extreme continental climate of central Siberia with average temperatures around -45°C in  
17 January and up to +35°C in August, the Lena River water discharge varies strongly  
18 throughout the seasons (e.g. Holmes et al., 2012). The river is covered by a thick ice layer  
19 (~2m) from October to late May/June and runoff is comparatively low during this time of the  
20 year (Yang et al., 2002). It reaches its maximum during the spring ice-breakup and snowmelt  
21 in late May to June when more than 50% of the annual freshwater, sediment, and dissolved  
22 and particulate organic matter discharge into the Laptev Sea take place (Rachold et al., 2004).  
23 With a mean annual water discharge of  $\sim 588$  km<sup>3</sup> between 1999 and 2008 (Holmes et al.,  
24 2012) the Lena ranks second largest of the Russian rivers after the Yenisey. Corresponding  
25 annual sediment, dissolved organic carbon (DOC) and particulate organic carbon (POC)  
26 fluxes are 20.7 Tg/yr (Holmes et al., 2002), 5.7 Tg/yr (Holmes et al., 2012), and 1.2 Tg/yr  
27 respectively (Rachold and Hubberten, 1999). A second major source for terrestrial organic  
28 matter delivered to the Laptev Sea is the sediment input by thermal erosion of the ice-rich  
29 Pleistocene ice complex or Yedoma deposits along the coast (see Gustafsson et al., 2011;  
30 Mueller-Lupp et al., 2000; see Rachold and Hubberten, 1999). Annual supply of sedimentary  
31 material and total organic carbon to the Laptev Sea by coastal erosion is estimated to be  $\sim 58.4$   
32 Tg/yr and 1.8 Tg/yr, respectively (Stein and Fahl, 2004).

1 The Lena River Delta is the largest arctic delta with an area of ~32,000 km<sup>2</sup>. It can be divided  
2 into three geomorphological terraces (Grigoriev, 1993; Schwamborn et al., 2002). The first  
3 terrace includes the active floodplains that were formed during the Holocene and makes up  
4 about 55% of the total delta area (Morgenstern et al., 2008) covering the central and eastern  
5 part. Within the first delta terrace remains of a Pleistocene accumulation plain, also called ice  
6 complex or Yedoma deposits, form the third terrace. Covering about 6% of the total delta area  
7 (Morgenstern et al., 2008). Sandy islands forming the second terrace cover the rest of the  
8 delta area in the west. The first and third terraces formed under completely different  
9 conditions. Whereas, fluvial high energy depositional regime characterize the Holocene (e.g.  
10 Schwamborn et al., 2002), the Pleistocene terraces were formed under a comparatively low  
11 energy alluvial and proluvial depositional regime (e.g. Schirrmeister et al., 2011). These  
12 contrasts result in distinct differences in OC content and quality, extent of soil formation,  
13 composition of the soil matrix, and ice content. Erosion of exposed surfaces means that both  
14 terraces contribute to the suspended particulate matter in the Lena Delta surface water  
15 sampled for this study, as well as suspended matter transported by the river from the southern  
16 catchment area.

17 Lena River water and sediment discharge is not equally distributed through the different delta  
18 channels (Fig. 1B). Approximately 80-90% of the total water and up to 85% of the sediment  
19 discharge are delivered through the three main eastern channels to the Buor Khaya Bay east  
20 of the delta, i.e. through the Sardakhsko-Trofimovskaya channel system (60-75% water, 70%  
21 sediment) and the Bykovskaya channel (20-25% water, 15% sediment). Only a minor portion  
22 is discharged to the north and west through the Tumatskaya and Olenyokskaya channels (5-  
23 10% water, 10% sediment; (Ivanov and Piskun, 1999).

24 All riverbank bluffs sampled here belong to the first terrace, which is elevated (5 to 16 m)  
25 over the active floodplains. The bluff profiles vary strongly in sediment composition and  
26 organic matter content. Within the profiles sandy layers derived from extreme flooding events  
27 (Schwamborn et al., 2002) and aeolian input (Kutzbach et al., 2004; Sanders, 2011) alternate  
28 with buried surface soil layers and peat layers rich in fibrous plant and root detritus in  
29 different stages of decomposition. The peat layers are either of autochthonous or of  
30 allochthonous origin. Allochthonous material is eroded from river banks further upstream and  
31 re-deposited in the delta.

32 The first terrace is characterized by wet polygonal tundra with depressed polygon centers and  
33 elevated polygon rims. Phytologically, the polygon centers are dominated by hydrophilic

1 sedges like *Carex aquatilis*, *Carex chordorrhiza*, *Carex rariflora*, and mosses (e.g.  
2 *Drepanocladus revolvens*, *Aulacomnium turgidum*) and the rims by mesophilic dwarf shrubs  
3 (e.g. *salix glauca*) and mosses (e.g. *Hylocomnium splendens*, *Timmia austriaca*) (Boike et al.,  
4 2013; Kutzbach et al., 2004; Sachs et al., 2010).

## 5 **2.2 Sampling**

6 The sampling sites presented in this study are located in the eastern part of the Lena Delta and  
7 adjacent Buor Khaya Bay (Fig. 1B). Permafrost soil samples, total suspended matter (TSM)  
8 from surface waters, and surface sediments were collected during two expeditions in August  
9 2009 and July/August 2010. Additional TSM samples were collected during the Lena River  
10 freshet in late May 2011. Four Holocene permafrost peat bluffs of different heights (3 to 8 m  
11 above river level in August 2009 and July/August 2010) were sampled along the main  
12 channels of the first delta terrace (all sampling sites in Fig. 1B and Table 1). In order to obtain  
13 samples that reflect the original state of the frozen permafrost soils, thawed material was  
14 removed with a spade for the total height of each bluff. Frozen pieces of peat were excavated  
15 at different depths using hatchet and hammer.

16 Suspended particulate matter of Lena River surface water was sampled at different stations in  
17 the main river channels of the delta on the Russian vessel Puteyski 405 (Fig. 1B, Table 1).  
18 Between 1 and 30 L of water were filtered on pre-combusted (4.5h at 450°C) and pre-  
19 weighed glass fiber filters (GF/F Whatman, 0.7µm membrane, Ø142mm) for biomarker  
20 analysis. Additionally, water samples of 15 and 20 L from the spring freshet in 2011 were  
21 stored cooled in opaque canisters for several days to allow for the suspended matter to settle.  
22 Before decanting the supernatant water it was filtered on pre-combusted and pre-weighed  
23 GF/F filters to check for the TSM remaining in suspension. For the sample presented here  
24 (sample ID 37) the TSM of the supernatant water represented 0.1% of the settled material on  
25 a dry weight basis and therefore the loss of material in suspension can be neglected.

26 Surface sediment samples from the Lena riverbed and off Muostakh Island were taken 2009  
27 using a grab sampler on board the Russian vessel Puteyski 405. Surface sediments from the  
28 Buor Khaya Bay were taken in 2010 with the Russian vessel PTS using a steel tube (Ø 5cm)  
29 connected to a rope. Penetration depths into sediment were between 3 and 6cm.

30 The peat and sediment samples were stored in pre-combusted glass jars (4.5h at 450°C) and  
31 GF/F filters were wrapped in pre-combusted aluminum foil. All samples were kept frozen at –  
32 20°C during storage and transport until analysis.

1 Additionally to the samples taken for this study, we analyzed 5 samples (2 from the early  
2 Holocene, 3 from the Pleistocene) from a profile on Kurungnakh Island, which were taken in  
3 2002 and provided by Lutz Schirrmeister from the AWI Potsdam, Germany. A detailed  
4 description of the study site and the paleoenvironmental interpretation was published by  
5 Wetterich et al. (2008). Furthermore, vegetation samples collected further south along the  
6 Lena River were provided by Ulrike Herzschuh and Juliane Klemm from the AWI Potsdam,  
7 Germany (for more information on the sampling sites see: Herzschuh et al. 2009; Klemm and  
8 Zubrzycki, 2009; Zubrzycki et al., 2012). Plant species analyzed here were *Aulacomnium*  
9 *turgidum* (moss), *Ledum palustre* (wild rosemary), *Carex spp.* (sedges), *Betula nana* (dwarf  
10 birch), *Salix* (willow), and *Larix* (larch).

## 11 **2.3 Laboratory analyses**

12 Peat and sediment samples were freeze-dried, homogenized, and subsampled for elemental  
13 and biomarker analysis.

14 All filters were oven-dried at 40°C for 24h. Due to expected problems with alkaline CuO  
15 oxidation of glass fiber filters in the microwave (dissolution of glass fiber), the particulate  
16 matter from samples selected for CuO oxidation was carefully scraped off the filter with a  
17 scalpel. During the filtering process a large portion of the particulate matter settles within the  
18 membrane structure. Therefore it was only possible to scrape off the material sitting directly  
19 on the filter surface. This material made up between 23-72 % (mean: 50%) of the total TSM  
20 on the filters. Because of this treatment the measured CuO oxidation products cannot  
21 accurately be related to the original water volume filtered and are rather treated like sediment  
22 samples normalized to the sample weight and weight of organic carbon.

### 23 **2.3.1 Elemental analyses**

24 Weight percent organic carbon (OC) and total nitrogen (TN) content of soil and sediment  
25 samples were determined by high temperature combustion after removal of carbonates as  
26 described by Goñi et al. (2003). The particulate organic carbon (POC) and particulate  
27 nitrogen (PN) content of TSM were analyzed on Ø25mm and Ø47mm GF/F obtained from  
28 the same water sample as the respective Ø142mm filters, which were scraped for lignin  
29 phenol analysis.

### 1 **2.3.2 CuO oxidation products**

2 Alkaline CuO oxidation was performed at Oregon State University based on the method  
3 described by Goñi and Montgomery (2000). Alkaline oxidations were carried out with  
4 nitrogen-purged 2N NaOH at 150°C for 1.5h using a microwave digestion system. After the  
5 oxidation, recovery standards (ethyl vanillin, *trans*-cinnamic acid) were added and the  
6 solution was acidified to pH 1 with concentrated HCl. Subsequently, samples were extracted  
7 with ethyl acetate. Extracts were evaporated to dryness under a stream of nitrogen. CuO  
8 reaction products were re-dissolved in pyridine and derivatized with bis-trimethylsilyl  
9 trifluoroacetamide (BSTFA)+1% trimethylchlorosilane (TCMS) to silylate exchangeable  
10 hydrogens prior to analysis by gas chromatography-mass spectrometry (GC-MS). Compounds  
11 were separated chromatographically in a 30m x 250µm DB1 (0.25µm film thickness)  
12 capillary GC column, using an initial temperature of 100°C, a temperature ramp 4°C/min and  
13 a final temperature of 300°C. Gas chromatography-mass spectrometry was used to identify  
14 and quantify individual biomarkers (e.g., Goñi et al., 2009). The GC-MS was set to scan from  
15 50 to 650 amu and used to acquire full spectra of compounds of interest that were compared  
16 to those of standards to confirm identities. Individual compounds were quantified based on  
17 the intensities of selected ions using multi-level calibrations run routinely during the analysis  
18 period. Yields of non-lignin products were quantified using the detector response of *t*-  
19 cinnamic acid. External calibration standards were determined for individual compounds  
20 using ions specific to each chemical structure. The calibrations, which were performed on a  
21 weekly basis to test the response of the GC-MS, were highly linear ( $r^2 > 0.99$ ) over the  
22 concentration ranges measured in the samples. A more detailed method description can be  
23 found in Goñi et al. (2009) and Hatten et al. (2012).

24 Quantified reaction products included eight lignin-derived compounds: vanillyl phenols (V =  
25 vanillin, acetovanillone, vanillic acid), syringyl phenols (S = syringaldehyde,  
26 acetosyringone, syringic acid), and cinnamyl phenols (C = *p*-coumaric acid, ferulic acid).

27 In addition, also non-lignin-derived phenols were quantified including *p*-hydroxybenzenes (P  
28 = *p*-hydroxybenzaldehyde, *p*-hydroxybenzophenone, *p*-hydroxybenzoic acid).

### 29 **2.4 End-member unmixing**

30 The concentration of different lignin phenol groups of marine sediment samples and riverine  
31 suspended matter samples was used to infer the contribution of gymnosperms and  
32 angiosperms to the total lignin derived OM. The end-member (EM) properties from the

1 literature (as shown in Amon et al., 2012) in the form of C/V and S/V ratios were transformed  
2 into relative concentrations of the respective lignin compounds (see Table 7). The linear  
3 mixing system of lignin concentrations in the samples can be written in matrix notation as:

$$4 \quad \mathbf{X} = \mathbf{AS} + \mathbf{R}$$

6  
7  $\mathbf{X}$  represents a  $n$ -by- $m$  matrix of  $n$  samples and  $m$  of lignin compounds.  $\mathbf{A}$  ( $n$ -by- $l$ ) denotes the  
8 mixing coefficients of  $l$  EMs for the  $n$  samples. The  $m$  EM properties (lignin concentrations)  
9 for the  $l$  EMs are represented by matrix  $\mathbf{S}$  ( $l$ -by- $m$ ).  $\mathbf{R}$  ( $n$ -by- $m$ ) denotes the residual matrix.  
10 This linear problem can be solved using non-negative least-squares fitting (NNLSQ,  
11 (Löfberg, 2004). Since the mixing coefficients must be positive and the abundances must add  
12 up to unity, a non-negativity-constraint ( $\mathbf{A} \geq 0$ ) and sum-to-one constraint for the rows in  $\mathbf{A}$   
13 was defined. Because the relative abundances of lignin represent a closed data set, we  
14 performed the centered-log-ratio transformation (Aitchison, 1982) to bring the data  $\mathbf{X}$  into  
15 real space. We implemented a Monte-Carlo simulation with 500 iterations, each with  
16 randomized first guess within the constraints formulated above. The resulting probability  
17 density function of possible solutions for each sample and EM contribution characterized by  
18 its median and interval containing 90 % of the possible solutions.

## 19 20 **3 Results**

### 21 **3.1 General characteristics and elemental composition**

22 The surface water TSM concentrations showed a strong spatial (within the delta) and  
23 temporal (seasonal/annual) variability (Table 2). The concentrations varied from 3.1 mg/L to  
24 174.9 mg/L in 2009 and from 3.5 mg/L to 32.2 mg/L in 2010. The maximum value of  
25 174.9 mg/L in 2009 of sample 17 (Fig. 1B, Table S2) was determined offshore Bykovsky  
26 Peninsula close to shore in shallow water depth. The particulate organic carbon (POC)  
27 concentrations and POC to particulate nitrogen (PN) ratios are from the companion paper  
28 (Winterfeld et al., 2015; submitted as companion paper) and additionally given in Table 2.  
29 The sample taken in 2011 shortly after the ice-breakup off Samoylov Island (sample ID 37)  
30 showed with 494 mg/L the highest TSM loads determined during this study.

31 OC and TN contents of first terrace soil samples varied strongly within individual riverbank  
32 bluffs and between the bluffs. The OC contents ranged from 1.02 to 17.14 wt% and the TN

1 contents from 0.03 to 0.45 wt% (Table 3, Fig. S6). The highest values (>10 wt% OC) were  
2 not necessarily found in the topsoil layers, but also within bluff profiles associated with layers  
3 containing plant remains like twigs and leaves. Lower OC and TN contents (<2 wt% and  
4 <0.1 wt%, respectively) were found in layers with high sand contents. The atomic OC to TN  
5 ratios (OC:TN) of these samples show a similar distribution pattern. The ratios varied from  
6 21.7 to 68 with the highest values (>40) in samples rich in plant remains.

7 Buor Khaya Bay surface sediments showed generally lower OC and TN contents than  
8 observed for the first and third delta terraces (Table 3) ranging from 1.67 to 2.47 wt% and  
9 from 0.09 to 0.18 wt%, respectively. The highest OC and TN contents (2.47 wt% OC and  
10 0.18 wt% TN) were analyzed for sample L09-34 off Muostakh Island (see Fig. 1B). The  
11 island is mainly composed of Pleistocene Yedoma deposits and highly affected by coastal  
12 erosion providing a lot of particulate matter throughout the open water season. The highest  
13 OC:TN ratio of 20.9 was determined off the Sardakh-Trofimovskaya channel system (sample  
14 L10-36, see Fig. 1B, Table 3), where the majority of the Lena River water and sediment  
15 discharge occurs.

## 16 **3.2 CuO oxidation products**

17 Table 4 and 5 summarize the sediment- and OC-normalized CuO product yields of samples  
18 presented in this study. Yields of individual samples can be found in the supplementary  
19 material (Table S4 and S5).

### 20 **3.2.1 Sediment- and carbon-normalized CuO oxidation yields**

21 On average the plant samples exhibit the highest V, S, C, and P phenol yields per gram dried  
22 sediment/plant tissue (dws), i.e.,  $\Sigma 8$  (sediment-normalized sum of V, S, and C phenols)  
23 ranging from 4.64 to 17.08 mg/g dws. Only a few soil samples of the first terrace reach  
24 similar yields. Generally first terrace  $\Sigma 8$  contents vary from 0.04 to 7.10 mg/g dws (mean  $\Sigma 8$   
25 1.93 mg/g dws). Contents from the third terrace on Kurungnakh Island are generally lower  
26 (<2.0 mg/g dws) except for the two Pleistocene samples from unit III ( $\Sigma 8$  is 1.81 mg/g dws  
27 for both samples). Suspended matter from 2009 to 2011 and surface sediment samples have  
28 CuO product yields in a similar range from 0.04 to 0.47 mg/g dws over all phenol groups. In  
29 the Buor Khaya Bay the yields decrease with distance from the delta. Highest values were  
30 determined in front of the Sardakh-Trofimovskaya channel and offshore Muostakh Island. As  
31 already shown for the OC and TN contents above, also the V, S, C, and P phenol yields vary

1 strongly within the first delta terrace soils samples and TSM samples. In general, the P and V  
2 phenol groups were most abundant followed by the S and C phenol groups.

3 An overview of the CuO yield per 100 mg OC ( $\Lambda 8$ ) for the different locations and sample  
4 types is presented in Figure 2. The overall patterns described for the sediment-normalized  
5 yields are also true for the carbon-normalized yields. The highest  $\Lambda 8$  were analyzed in  
6 samples from the first and third delta terraces varying between 0.78 and 8.81 mg/100mg OC  
7 over all phenol groups (Table 5). The  $\Lambda 8$  were lower in TSM from the summers 2009 and  
8 2010 (<1.5 mg/100mg OC) and notably higher for the spring flood sample from 2011  
9 (5.16 mg/100 mg OC) as well as for the surface sediments of the Buor Khaya Bay (mean  
10 value 1.96 mg/100 mg OC). Also the amounts of individual phenol groups are different  
11 between the delta soil samples, the TSM, and the surface sediments samples. Generally the P  
12 and V phenols were most abundant followed by S and C phenols (Fig. 2). Again, the two  
13 samples from the third terrace from unit III were slightly different. Here, the S phenols were  
14 most abundant followed by the P, V, and C phenols. The distribution of V, S, C, and P  
15 phenols in the summer TSM samples of 2009 and 2010 were similar with the V and P phenols  
16 being most abundant. The spring flood sample from 2011 and the surface sediment samples  
17 were comparable with V and S phenols having the highest yields (Fig. 2).

### 18 **3.2.2 Vegetation source parameters**

19 The bulk samples of the first delta terrace show a broad range of C/V and S/V ratios (0.16 to  
20 1.16 for C/V and 0.58 to 1.58 for S/V, Table 5 and S5 and Fig. S7). As shown in Fig. 4B the  
21 values fall on a mixing line between woody gymnosperm and non-woody angiosperm tissues.  
22 The P/V ratios show a similar range of variation. The samples from the third terrace have  
23 comparable ratios as those from the first terrace, with the highest C/V and S/V ratios  
24 determined for the two Pleistocene samples from unit III. The values of the TSM samples  
25 taken in summer and spring and in three consecutive years are within the same range. Mean  
26 C/V ratios were 0.21, 0.18, 0.25 and mean S/V ratios were 0.44, 0.38, 0.5 for the years 2009,  
27 2010, and 2011, respectively. The P/V ratios were higher in the summers of 2009 and 2010  
28 (0.65 to 1.25 and 0.62 to 0.89, respectively) than in spring 2011 (0.44). The C/V, S/V, and  
29 P/V ratios vary only slightly in the Buor Khaya surface sediments and are generally in the  
30 range of the TSM samples and lower than the mean of the first delta terrace and the third  
31 terrace soil samples. Except for the sample of larch needles, the C/V, S/V, and P/V ratios of  
32 the vegetation samples reflect their tissue and plant origin closely (Fig. 4B and 5).

### 1 **3.2.3 Degradation indicators**

2 The acid to aldehyde ratios of vanillyl and syringyl phenols ( $Ad/Al_{V,S}$ ) of the first delta  
3 terrace vary strongly from moderately degraded (0.5 to 0.6) to highly degraded ( $>0.6$ ) (Figs.  
4 4A and S7, Table 5). Ratios of the third terrace on Kurungnakh Island are generally lower  
5 ( $<0.6$ ) than ratios from the first terrace. Notably, the lowest ratios were analyzed for the oldest  
6 sample S45 ( $<0.4$ , Table 5).  $Ad/Al_{V,S}$  ratios of the summer TSM are in the range of the first  
7 delta terrace or higher, e.g. varying between 0.68 and 3.97 for  $Ad/Al_V$  in 2009 and between  
8 0.69 and 2.02 in 2010. The spring flood sample from 2011 is characterized by one of the  
9 lowest ratios of all samples presented here (0.32 for both,  $Ad/Al_V$  and  $Ad/Al_S$ ). Buor Khaya  
10 Bay surface sediments showed ratios  $>0.6$ , which are in the range of the first delta terrace and  
11 summer TSM samples (0.98-1.75 for  $Ad/Al_V$  and 0.77-1.37 for  $Ad/Al_S$ ). The highest ratios  
12 were analyzed off Muostakh Island (sample L09-34). The vegetation samples have low  
13  $Ad/Al_{V,S}$  ratios ( $<0.4$ ) except for the larix needles, the herb sample (*Ledum palustre*) and the  
14 moss sample (*Aulacomnium turgidum*), which have ratios  $>0.4$ . It is known that different  
15 plant species or even different parts of a plant (e.g. mosses and needles, respectively) can  
16 have naturally higher acid concentrations resulting in higher  $Ad/Al_{V,S}$  (=more degraded) even  
17 when they are fresh (see Benner et al., 1990). This is most likely also the case for the  
18 vegetation samples presented here.

### 19 **3.3 End-member unmixing**

20 The EM unmixing was performed for the TSM and surface sediment samples. The EM  
21 properties of moss and peat contribution in this model do not represent the range of values  
22 observed in our samples. Fig. 5 shows our  $Pn/P$  and  $P/V$  ratios in relation to several published  
23 values, amongst others the end-members used for moss, soil, and peat.

24 Therefore, we applied an unmixing model distinguishing between the four major vegetation  
25 sources for OM: woody and non-woody gymnosperm and angiosperm tissues. We used  $C/V$   
26 and  $S/V$  ratios and took the EMs (Table 6) from Amon et al. (2012) and references therein,  
27 which covered the complete range measured in TSM and surface sediment samples (Fig. 4B).

28 The median values of the unmixing solutions (obtained by Monte-Carlo simulation) of  
29 angiosperms (woody + non-woody) and gymnosperms (woody + non-woody) are shown in  
30 Table 7. The relative contributions show a broad range for the summer TSM samples of 2009  
31 and 2010, i.e. gymnosperm contribution varies from 0.24 to 0.69 and from 0.49 to 0.63,  
32 respectively. The low gymnosperm contribution of 0.24 is inferred for sample 17, located off

1 Bykovsky Peninsula (Fig. 1B). The contributions to the Buor Khaya Bay surface sediments  
2 vary to a lesser extent from 0.49 to 0.56 for gymnosperms.

3

## 4 **4 Discussion**

### 5 **4.1 Spatial and temporal patterns of Lena Delta total suspended matter**

#### 6 **4.1.1 Suspended sediment distribution and particulate lignin biomarker** 7 **abundances**

8 Surface water suspended particulate matter sampled in highly dynamic systems like a river  
9 delta can only provide very local snapshots of the suspended matter properties. The Lena  
10 Delta is characterized by a dynamic hydrology and fast changes of local conditions of erosion  
11 and accumulation, which are related to changes in water velocity and turbidity leading to  
12 channel migration and branching (Fedorova et al., 2013). Longer time series covering several  
13 years and seasons are needed to observe catchment related changes in these properties  
14 independent of the natural variability. Further, it is important to consider the season of TSM  
15 sampling: In the summer season in July and August the active layer depth is deepest,  
16 riverbank erosion along the delta channels is very pronounced, and streams draining ice  
17 complex deposits and thermokarst lakes transport more sediment providing local delta-  
18 derived sediment to the river surface water. During the ice break-up and associated spring  
19 flood in late May to early June the soils in the delta and northern catchment are still frozen.  
20 Riverbanks and bluffs are eroded by ice jamming against the riverbank and by thermal  
21 abrasion by relatively warmer Lena River water. The eroded material mixes with sediment  
22 transported from the south and is exported with the flood to the Laptev Sea coastal zone.

23 Our TSM concentrations from July/Aug 2009 and 2010 (mean values are 28.5 and  
24 19.85 mg/L, respectively) showed a high spatial and inter-annual variability (Fig. 3A-C).  
25 Surface water TSM from the Lena Delta has been sampled during several expeditions in the  
26 past, mainly during the summer season, and by the Federal Service of Hydrometeorology and  
27 Environmental Monitoring of Russia (Roshydromet) at several stations throughout the delta  
28 (see Fedorova et al., 2013). All concentrations measured in this study were well within the  
29 range of published values of samples taken in July to early September between 1989 and 2003  
30 (16.5 to >30 mg/L) (Cauwet and Sidorov, 1996; Rachold and Hubberten, 1999). Our single  
31 measurement from the spring flood in late May 2011 taken offshore Samoylov Island  
32 (72.37°N, 126.47°E) was more than 10 times higher (494 mg/L) than the summer values. It

1 clearly reflects the distinct seasonality of the hydrograph of the Lena River, where more than  
2 50% of the annual TSM export happen during the spring freshet (Cauwet and Sidorov, 1996;  
3 Rachold et al., 2004). The only additional spring flood values from the Lena River we are  
4 aware of are provided by the Arctic Great Rivers Observatory Project (A-GRO, [www.wwww.arcticgreativers.org](http://www.wwww.arcticgreativers.org)) and are taken at Zhigansk gauging station (66.77°N, 123.37°E)  
5 approximately 900 km south of the Lena Delta. The TSM concentrations reported by Arctic  
6 GRO for late May/early June 2004 to 2010 are lower than our measurement varying from  
7 28.8 to 221 mg/L. The higher Lena Delta value from 2011 could be a result of the flood wave  
8 eroding and entraining more sedimentary material on its way to the north. Despite the low  
9 sample resolution for the time of the spring flood, our single delta sample and the few  
10 samples from Zhigansk (A-GRO) highlight the strong seasonal differences in river supply  
11 suspended sediments and the need for more sampling campaigns during the spring freshet to  
12 improve flux estimates of terrestrial OM to Laptev Sea.

14 We chose to only discuss the carbon-normalized ( $\Lambda 8$ ) yields instead of sediment-normalized  
15 ( $\Sigma 8$ ) results of our TSM samples, because during sample preparation some glass fiber filter  
16 material was included in the analyzed sample, thus biasing the sediment-normalized  
17 calculation (described above in Sect. 2.3). Like the TSM concentrations discussed above, the  
18  $\Lambda 8$  concentrations reflect the strong seasonality of the Lena River hydrograph.  $\Lambda 8$   
19 concentrations were similar in the summers 2009 and 2010 (mean  $\Lambda 8$  1.03 and  
20 1.09 mg/100mg OC, respectively; Table 5) and about five times higher in spring 2011 ( $\Lambda 8$  of  
21 5.16 mg/100 mg OC). The normalization to the total organic carbon measured in our TSM  
22 samples, which is a mixture of terrestrial- and plankton-derived organic matter, most likely  
23 alters the ratio of  $\Lambda 8$  to organic carbon in the sample. The presence of aquatic plankton-  
24 derived OM dilutes carbon-normalized lignin concentrations lowering  $\Lambda 8$  values relative to  
25 those from terrestrial source material. The particulate organic carbon to nitrogen ratios  
26 (POC:PN) from 2009 to 2011 (Table 2, from Winterfeld et al., 2015 companion paper)  
27 suggest a considerable amount of nitrogen-rich plankton-derived OM is present in our TSM  
28 samples. This is further supported by lower  $\delta^{13}\text{C}$  values determined for the bulk POM taken in  
29 summer 2009 and 2010 (mean  $\delta^{13}\text{C}$  of  $-30.6\text{‰}$  and  $-29.2\text{‰}$ , respectively; Winterfeld et al.,  
30 companion paper), which suggests a considerable fraction of freshwater plankton utilizing  
31 dissolved inorganic carbon depleted in  $\delta^{13}\text{C}$  is present in the surface water as has been shown  
32 for the Lena and Yenisey Rivers (e.g. Alling et al., 2012; Galimov et al., 2006; Rachold and  
33 Hubberten, 1999). Furthermore, Winterfeld et al. (2015, companion paper) estimated fractions  
34 of phytoplankton OM contributing to the bulk POM of the same water samples analyzed here

1 for lignin phenol composition using two simple binary mixing models between  
2 phytoplankton- and soil-derived POM based on the POC:PN ratios and based on the  $\delta^{13}\text{C}$   
3 values. For the lignin phenol samples presented here the phytoplankton fractions ranged from  
4 47% to 77% and 23% to 53% based on POC:PN ratios and  $\delta^{13}\text{C}$  values, respectively  
5 (Winterfeld et al, companion paper). However, low POC:PN ratios in combination with  
6 elevated  $\Delta 8$  values were also found in a river-dominated estuary in the US (Goñi et al., 2003).  
7 Here, the authors suggested that contributions of bacterial nitrogen-rich exudation products  
8 generated during vascular plant decomposition (Rice and Hanson, 1984), could be responsible  
9 for the lower POC:PN ratios (Goñi et al., 2003). Yet, OC:TN ratios from Lena Delta soils  
10 presented in this study are  $>20$  (Table 3) and from soils along a North-South transect in the  
11 Lena watershed ( $73.5^\circ - 69.5^\circ\text{N}$ ) are  $>13.5$  (Zubrzycki et al., 2012), which points to less  
12 degraded soil OM and together with the low POM  $\delta^{13}\text{C}$  values (Winterfeld et al., 2015,  
13 companion paper) supports the idea of higher phytoplankton OM contributions to our TSM  
14 samples instead of highly degraded mineral soil.

#### 15 **4.1.2 Tracers of vegetation sources from the Lena River catchment**

16 The C/V and S/V ratios allow to distinguish different vegetation sources, such as woody and  
17 non-woody tissues as well as gymnosperm and angiosperm tissues, respectively, (e.g. Hedges  
18 and Mann, 1979; Hedges et al., 1982; Kuzyk et al., 2008). As shown in Fig. 4B, the TSM  
19 values of 2009-2011 reflect a mixture of woody gymnosperm and non-woody angiosperm  
20 vegetation sources. However, cinnamyl phenols are known to degrade relatively fast during  
21 early diagenesis resulting in decreased C/V ratios, while S/V ratios seem to be only  
22 moderately altered (Benner et al., 1990; Opsahl and Benner, 1995). That implies our low C/V  
23 ratios do not unambiguously reflect high woody gymnosperm contribution. As a result, any  
24 estimate of woody gymnosperm contribution based on C/V ratios alone must be considered a  
25 maximum value. But this process seems to be less important here, because our data do not  
26 deviate from a linear trend in the C/V vs. S/V diagram. Furthermore, note that the sample  
27 preparation of filters was done by scraping off supernatant material (section 2.3) resulting in  
28 some kind of sediment fractionation, which could affect the lignin phenol compositions. Yet,  
29 our TSM samples lie within the range of soils and sediments presented here and we suggest  
30 this effect to be negligible for the OM source and degradation interpretation below.

31 Our C/V and S/V values are slightly higher than values measured for particulate and dissolved  
32 lignin sampled in the Lena Delta in 1994 (Lobbess et al., 2000) and dissolved lignin sampled  
33 from the Lena River at Zhigansk (Amon et al., 2012). This could either be because the

1 contribution of non-woody angiosperm sources, most likely tundra vegetation to Lena Delta  
2 TSM increased since 1994 due to active layer deepening and increased riverbank abrasion.  
3 Alternatively, a detectable difference in contribution of non-woody angiosperm OM to delta  
4 samples compared to the southern sample location at Zhigansk is simply due to the fact that  
5 the latter location lies in the taiga-tundra transition zone, where higher woody gymnosperm  
6 contributions would be expected. Thirdly, a large fraction of the particulate river load might  
7 be trapped in floodplains and/or the lower reaches of the Lena River. Particularly material  
8 from the distal parts of the watershed carrying the predominantly woody gymnosperm signal  
9 would not be transported efficiently to the delta. This inefficient transport mechanism of  
10 riverine particulate load is characteristic for large river system and has for instance been  
11 reported for the Amazon River and the Fly River, Papua New Guinea (e.g. Alin et al., 2008;  
12 Aufdenkampe et al., 2007; 2011; Blair and Aller, 2012; Goñi et al., 2014; Moreira-Turcq et  
13 al., 2013; Zocatelli et al., 2013). The slightly lower C/V and S/V values from Lobbes et al.  
14 (2000) and Amon et al. (2012) could also be well within the range of the natural variability of  
15 Lena River TSM composition, which was not covered by samples from 2009 and 2010 in this  
16 study. However, it should be noted that dissolved and particulate lignin might be derived from  
17 different terrigenous sources, i.e. from a modern OM pool and a pre-aged OM pool,  
18 respectively. In general, dissolved organic matter in Arctic rivers is of modern age while  
19 POM is much older ( $10^2$ - $10^3$  years; e.g. Guo and Macdonald 2006; Guo et al., 2007;  
20 Raymond et al., 2007), which could result in differences of the lignin phenol composition and  
21 hence be an additional explanation for the differences of C/V and S/V values between our  
22 particulate lignin values and the dissolved lignin of Amon et al. (2012). Nonetheless, our C/V  
23 and S/V ratios clearly depict the catchment vegetation characteristics of the Lena River being  
24 a mixture of taiga forest in the south and tundra in the north. They therefore distinguish the  
25 Lena River catchment from other arctic river catchments like the Ob' River (Dickens et al.,  
26 2011) or Mackenzie River (Goñi et al., 2000).

27 Although *p*-hydroxybenzenes (P) have multiple sources, the CuO oxidation of fresh  
28 Sphagnum and other mosses, which do not produce the typical lignin phenols, release  
29 considerable amounts of *p*-hydroxybenzenes and the P/V and *p*-hydroxyacetophenone to P  
30 ratios (Pn/P) have been used as tracer for Sphagnum-derived OC in peats (Dickens et al.,  
31 2011; Tsutsuki and Kondo, 1995; Williams et al., 1998). The higher P/V ratios of summer  
32 TSM from 2009 and 2010 (mean ratios 0.9 and 0.8, respectively) compared to spring 2011  
33 (0.4; Fig. 5, Table 5) might indicate a higher contribution of mosses to Lena Delta TSM in the  
34 summer season, presumably derived from local tundra vegetation. Alternatively, the rather

1 high fraction of phytoplankton-derived OM contributing to our summer samples (see  
2 Winterfeld et al., 2015 companion paper) could also result in higher P/V ratios as  
3 phytoplankton has been shown to contain elevated amounts of P (e.g. Goñi and Hedges,  
4 1995). Overall, the summer and spring P/V ratios in TSM were lower than the mean bulk P/V  
5 ratio of the first and third terrace (1.1 and 1.0, respectively), the moss sample (*Aulacomnium*  
6 *turgidum*) analyzed here (P/V ratio of 2.3) and other values from the literature for Sphagnum  
7 moss and peat (see Fig. 5). The relatively lower P/V ratios in TSM samples thus indicate that  
8 moss contribution is minor compared to a dominant non-woody angiosperm source,  
9 particularly when considering possible P contributions from phytoplankton to TSM.

#### 10 **4.1.3 State of diagenetic alteration of suspended particulate lignin biomarkers**

11 Lignin phenol composition has been widely used to identify sources of terrigenous OM in  
12 aquatic and soil systems and characterize the degree of aerobic degradation (e.g. Benner et al.,  
13 1990; Goñi and Hedges, 1992; Hedges and Mann, 1979; Hernes and Benner, 2002; Tesi et al.,  
14 2007). The acid to aldehyde ratios of vanillyl and syringyl ( $Ad/Al_{V,S}$ ) usually increase with  
15 increasing OM oxidation. In general, values  $<0.4$  for both ratios are considered fresh and  
16 samples with values  $>0.4$  have undergone some degree of degradation (Goñi et al., 1993;  
17 Hedges et al., 1988).

18 The TSM  $Ad/Al_{V,S}$  ratios vary annually and with the hydrograph. The spring flood value from  
19 2011 appears to be derived mainly from fresh plant litter and/or surface soils (Fig. 4A) in  
20 agreement with the dissolved organic matter (DOM) exported during the flood, which was  
21 also found to be younger than summer DOM (Amon et al., 2012). In contrast, the  $Ad/Al_{V,S}$   
22 ratios of our TSM collected in summer, indicate a more degraded OM source presumably  
23 from deeper soil horizons that thawed during the summer months. The deeper soil OM could  
24 partly originate from the first and third delta terraces. However, most of the  $Ad/Al_{V,S}$  ratios  
25 we determined for the TSM were higher than the bulk soil  $Ad/Al_{V,S}$  ratios of the first and third  
26 terraces. Such finding points to either an additional more degraded source, most likely from  
27 south of the Lena Delta, or a more degraded fraction of soil present in suspended matter, most  
28 likely the fine fraction. The fine grain-size fraction of soils and riverine suspended matter are  
29 generally associated with higher  $Ad/Al_{V,S}$  ratios (Carrington et al., 2012; Guggenberger et al.,  
30 1994; Hedges et al., 1986) and the fine fraction is also most likely to be held in suspension  
31 during lower summer flows compared to coarser grain sizes. Similarly degraded terrigenous  
32 OM was found in the surface waters of the Mackenzie River Delta also draining a permafrost  
33 affected watershed (Goñi et al., 2000).

1 Additionally, sorption of dissolved lignin to mineral surfaces could have an effect on the  
2  $Ad/Al_{V,S}$  ratios. Dissolved lignin in the Lena River has high  $Ad/Al_{V,S}$  ratios of  $\sim 0.9$  to 1.6  
3 during the peak flow and  $\sim 0.6$  to 1.3 during mid and base flow (Amon et al., 2012). Higher  
4  $Ad/Al_{V,S}$  ratios of dissolved lignin are not necessarily associated with highly degraded lignin,  
5 but are also observed when dissolved lignin is derived from leaching of litter or soil (Hernes  
6 et al., 2007). The Lena Delta summer TSM ratios from 2009 and 2010 were higher than any  
7 other values reported for particulate lignin in the Lena Delta or other arctic rivers (Dickens et  
8 al., 2011; Goñi et al., 2000; Lobbes et al., 2000), and to a large fraction also higher than  
9 values in dissolved lignin. Thus, they cannot be explained by sorption of dissolved lignin, but  
10 potentially reflect input from a highly degraded source, e.g., from greater soils depths of the  
11 southern catchment.

## 12 **4.2 Spatial patterns in Buor Khaya Bay surface sediments**

### 13 **4.2.1 Lignin biomarker abundances**

14 In contrast to the surface water TSM snapshots, the surface sediments from the Buor Khaya  
15 Bay integrate the sedimentary OM and associated lignin phenol signal over a certain period of  
16 time depending on the local accumulation rates and the sediment re-working by waves and  
17 land-fast ice affecting the shallow coastal zone. The surface sediments therefore reflect an  
18 average of the OM transported to the coastal zone and smooth the seasonal and interannual  
19 differences in OM properties as well as the differences between OM sources. Buor Khaya  
20 Bay sedimentary OM is mainly derived from three sources, i.e. terrigenous OM transported  
21 by the Lena River, terrigenous OM derived from coastal erosion of the Buor Khaya coast  
22 predominantly consisting of Pleistocene ice complex deposits, and aquatic (riverine and  
23 marine) primary production. The latter source is negligible when discussing lignin phenols.

24 The sediment-normalized ( $\Sigma 8$ ) and carbon-normalized ( $\Lambda 8$ ) lignin phenol concentrations of  
25 Buor Khaya Bay surface sediments are high in front of the two main delta outlets, the  
26 Sardakh-Trofimovskaya channel and the Bykovskaya channel (Fig. 3G, Table 4 and 5), and  
27 decrease offshore. That points to the Lena River as the dominant source of lignin phenols with  
28 decreasing influence offshore, presumably as a result of hydrodynamic sorting where a less  
29 lignin phenol-rich finer sediment fraction is transported further offshore. Highest  $\Sigma 8$   
30 contributions from coastal erosion are evident at the site off Muostakh Island (L09-34), while  
31 the carbon-normalized yield ( $\Lambda 8$ ) at this location was slightly smaller than at the river outlets  
32 (samples L10-23 and L10-36).

1 High lignin phenol concentrations are generally associated with the coarse particulate OM  
2 fraction in soils and suspended material and they decrease with decreasing grain size  
3 (Carrington et al., 2012; Guggenberger et al., 1994; Hedges et al., 1986). An offshore gradient  
4 of decreasing grain size off the delta coast and towards greater water depths has been reported  
5 for the Buor Khaya Bay (Charkin et al., 2011). The spring flood could play a major role in  
6 transporting coarser lignin bearing OM to the coastal zone, which is in agreement with the  
7 high spring flood  $\Sigma 8$  and  $\Lambda 8$  concentrations from 2011. In contrast, slower current velocities  
8 during summer would transport fine particulate material to the delta edge or further offshore,  
9 carrying a lower  $\Lambda 8$  signature. Alternatively, the increased sedimentation of particulate and  
10 dissolved material through flocculation in the mixing zone of fresh and salt water in the  
11 prodelta area (marginal filter; cf. Lisitsyn, 1995) could be an additional reason for increased  
12 lignin phenol concentrations at these sample locations. Unfortunately, there is not much  
13 known about transport of sediment along the delta channels to the coastal zone.

#### 14 **4.2.2 Vegetation sources contributing to sedimentary organic matter**

15 We observed a generally high contribution of terrestrial organic matter to the Buor Khaya Bay  
16 sediments based on the OC:TN ratios (Table 3). An offshore trend of decreasing OC:TN  
17 ratios likely reflects the increasing marine contributions by plankton as well as decreasing  
18 amounts of terrigenous material reaching offshore locations, which supports similar findings  
19 on OM sources in Buor Khaya Bay surface sediments (e.g. Karlsson et al., 2011; Tesi et al.  
20 2014).

21 The contribution of woody and non-woody gymnosperm and angiosperm tissues based on  
22 C/V and S/V ratios as well as the contribution from mosses based on the P/V ratios was rather  
23 similar for all surface sediment samples. The C/V and S/V ratios were notably lower than in  
24 the delta soil samples in this study (Fig. 3L-S and 4B) and the surface soil and ice complex  
25 samples taken in the Lena watershed and along the Buor Khaya coast by Tesi et al. (2014).  
26 This suggests a considerable contribution of woody gymnosperm tissues from the southern  
27 Lena catchment to Buor Khaya Bay sediments. Additional angiosperm-derived OM  
28 contributed by coastal erosion of Pleistocene ice complex deposits (Tesi et al., 2014) seems to  
29 dilute the gymnosperm signal from the Lena River with distance to the delta (slight increase  
30 in C/V and S/V ratios offshore, Table 5). The angiosperm-derived OM originating from the  
31 Lena catchment and from coastal erosion of ice complex, respectively, contribute to the Buor  
32 Khaya Bay sediments and these two sources cannot unambiguously be distinguished based on  
33 their C/V and S/V ratios. Here, the use of bulk as well as lignin-specific  $^{14}\text{C}$

1 concentrations might be helpful as the Pleistocene ice complex deposits are more depleted in  
2  $^{14}\text{C}$  than the soil-derived OM from the Lena catchment (e.g. Karlsson et al., 2011; Vonk et al.,  
3 2012; Winterfeld et al., 2015 companion paper).

4 Further, sedimentary P/V ratios (0.4 to 0.5) suggest the fraction of moss-derived OM in the  
5 sediments is smaller than in the summer TSM samples (0.62 to 1.25) and bulk soils from the  
6 first and third delta terrace (Fig. 5, Table 5). Instead, the sedimentary P/V ratios were similar  
7 to the spring TSM sample from 2011 (value of 0.4) pointing to a considerable contribution of  
8 spring flood TSM to Buor Khaya Bay sediments. However, we cannot exclude that lower P/V  
9 ratios could be partially the result of selective degradation of more labile P phenols compared  
10 to V phenols resulting in lower P/V ratios than carried by the original OM source (Hedges and  
11 Weliky, 1989; Williams et al., 1998). And as discussed above for the TSM samples we also  
12 cannot exclude possible contributions from phytoplankton containing elevated P  
13 concentrations to sedimentary OM. The slight increase in P/V ratios further offshore (samples  
14 L10-23 to L10-25, Table 5) might point to higher phytoplankton OM contributions with  
15 larger distance to the coast. The estimated relative contributions of the different vegetation  
16 zones in the Lena catchment will be discussed in section 4.3.1.

#### 17 **4.2.3 Degradation of terrigenous organic matter**

18 The relatively high  $\text{Ad}/\text{Al}_{\text{V,S}}$  ratios (0.77 to 1.75, Table 5, Fig. 3K) imply a rather strong  
19 degradation of lignin phenols in the surface sediments. There is a small gradient towards less  
20 degraded material along the offshore transect (Fig. 3K) of progressively finer sediments  
21 (Charkin et al., 2011). This in contrast to analysis of soils and Amazon River suspended  
22 material, where  $\text{Ad}/\text{Al}_{\text{V,S}}$  ratios increased with decreasing particle size (Amelung et al., 1999;  
23 Carrington et al., 2012; Hedges et al., 1986). As summer TSM in our samples is  
24 predominantly strongly degraded (see section 4.1), and as discussed above inferred to be more  
25 fine grained than the less degraded spring flood material, this observation argues against a  
26 dominant control of hydrodynamic sorting on the lignin monomer distribution. An additional  
27 OM source with a less degraded  $\text{Ad}/\text{Al}_{\text{V,S}}$  signature contributing to Buor Khaya Bay  
28 sediments, e.g. Lena River spring flood TSM or material derived from erosion of ice complex  
29 deposits along the Buor Khaya coast, could explain the offset. Efficient sediment  
30 redistributing processes such as bottom erosion and nepheloid layer bottom transport of  
31 sediments in the Buor Khaya Bay have been identified by Charkin et al. (2011).

32 It is difficult to assess where the lignin degradation occurred. Oxidative degradation of the  
33 lignin macromolecule in soils by fungi is known to increase  $\text{Ad}/\text{Al}_{\text{V,S}}$  ratios severely (e.g.

1 Goñi et al., 1993). Subaqueous decay of lignin has also been shown to increase the Ad/Al  
2 ratios (e.g. Opsahl and Benner, 1995). We favor the explanation of aerobic degradation on  
3 land, because Ad/Al<sub>v,s</sub> ratios of the sediment samples are in the upper range of values found  
4 in the bulk first terrace soils and well within the range of summer TSM samples from the  
5 delta, suggesting that river-transported material is the dominant source of OM deposited in  
6 surface sediments. The one spring flood sample from 2011 appears to be relatively fresh. If  
7 the majority of particle discharge to the Laptev Sea occurs during spring, we would expect  
8 that similarly low Ad/Al<sub>v,s</sub> ratios would be observed in the sediments. The fact that we do not  
9 observe such a signal might be either due to insufficient information about the heterogeneity  
10 of material transported during the spring flood, or due to efficient degradation of lignin during  
11 transport and/or early diagenesis. Karlsson et al. (2011) studied bulk parameters and lipid  
12 biomarker contents of surface sediments in the Buor Khaya Bay. Using a three EM and dual  
13 isotope ( $\delta^{13}\text{C}$  and  $\Delta^{14}\text{C}$ ) Monte Carlo simulation, these authors suggested that about 60% of  
14 the sedimentary OM is derived from ice complex deposits and roughly 20% each from surface  
15 soils and primary production. Further, using lipid biomarker indicators for OM degradation  
16 and  $^{14}\text{C}$  dating, they found marked differences between river-derived POC that was younger  
17 and more degraded and sedimentary OM that was older but less degraded. These lipid-based  
18 findings are in contrast to our lignin data, which suggest similar states of degradation for both,  
19 river-derived OM and surface sediments.

20 Notably, the highest Ad/Al<sub>v</sub> ratio of 1.75 was measured offshore Muostakh Island (Fig. 3K,  
21 Table 5), which consists of Pleistocene ice complex deposits. In contrast, the Ad/Al<sub>v</sub> ratios of  
22 ice complex deposits on Muostakh Island were much lower varying between 0.16 and 0.95  
23 (see Tesi et al., 2014). Additionally, other ice complex deposits from the Buor Khaya Cape  
24 and along the Lena channel (Tesi et al., 2014) as well as from Kurungnakh (this study)  
25 showed similar low Ad/Al<sub>v</sub> ratios supporting the idea of relatively fresh ice complex OM  
26 (e.g. Karlsson et al., 2011; Vonk et al., 2012). Therefore, the lignin fraction of the OM must  
27 have been strongly degraded between erosion and deposition in surface sediments. This could  
28 have happened after thawing on land and/or post-depositional under subaqueous conditions.  
29 Vonk et al. (2012) suggested that substantial degradation of Muostakh Island ice complex  
30 material occurs rapidly and immediately after thawing. Sanchez-García et al. (2011)  
31 determined rapid particulate OM degradation rates to occur subaqueously in shallow Laptev  
32 Sea waters.

33

## 4.3 Terrigenous organic matter sources of Lena Delta suspended matter and Buor Khaya Bay surface sediments

### 4.3.1 Unmixing of taiga and tundra vegetation contributions

As in a first approximation, gymnosperm vegetation is restricted exclusively to the taiga part of the Lena River catchment, we use the gymnosperm to angiosperm ratio as estimate for the relative contributions of taiga and tundra. Therefore, we combined the model solutions for woody and non-woody contributions of gymnosperms and angiosperms, respectively. According to the model presented here, the fractions of gymnosperm and angiosperm-derived OM varied strongly in the summer TSM samples. However, the mean gymnosperm contributions for spring 2011 and the summers 2009 and 2010 were very similar (Table 7), i.e. 0.4 (n=1), 0.5 (n=7), and 0.6 (n=8), respectively. The Buor Khaya Bay surface sediment total gymnosperm fraction was in the range of the TSM values (mean fraction = 0.5, n=5). In summary the model suggests roughly equal contributions of total gymnosperm and total angiosperm-derived OM in suspended particulate OM and sediments based on the lignin monomer distribution. This implies that a large fraction of the gymnosperm POM derived from the taiga gets trapped in floodplains along the course of the Lena River and hence the contribution of angiosperm vegetation, mainly present in the tundra, to surface water TSM is relatively big. The Buor Khaya Bay surface sediments receive additional angiosperm-derived OM through coastal erosion of ice complex deposits (Tesi et al., 2014), which adds to the angiosperm-derived OM from the Lena catchment and dilutes the gymnosperm signal with distance to the delta. Moreover, we cannot distinguish and therefore exclude small angiosperm contributions from the taiga zone itself being a heterogeneous landscape with some amount of angiosperm vegetation and contributions from higher elevated areas, where bushes and grasslands are favored over trees. Thus, taiga-derived material might indeed account for more than 50% of the total lignin in our samples, which in turn implies that based on our model the maximum contribution by the tundra zone (angiosperm OM) is 50%. Also the C/V and S/V values we chose for the non-woody angiosperm source (“a” in Fig. 4B, see Hedges and Parker, 1976; Hedges and Mann, 1979) add some uncertainty to the model results that has to be considered when interpreting the data. As shown in Fig. 4B the end-member values lie well within most of the soil samples, but there are also several soil and vegetation samples with higher and lower C/V and S/V ratios. Using more representative end-member values including the natural variability of the different permafrost soil sources could reduce

1 the model uncertainty here. Unfortunately, this kind of data is not available for Lena  
2 watershed, yet.

3 The comparison of C/V and S/V ratios between dissolved and particulate lignin can be  
4 complicated by fractionation processes occurring during leaching of lignin phenols from plant  
5 tissues and soils as well as sorption of dissolved lignin to minerals in soils or sediments  
6 (Hernes et al., 2007). However, Lobbes et al. (2000) and Amon et al. (2012) have shown that  
7 C/V and S/V ratios of dissolved lignin in arctic rivers, including the Lena River, reflect the  
8 vegetation signal of the individual catchments and are not significantly altered by degradation  
9 or fractionation. Therefore, we feel confident to compare our Lena Delta data with C/V and  
10 S/V ratios generated by Amon et al. (2012).

11 We compared the gymnosperm fractions in our samples with the results from Amon et al.  
12 (2012), who estimated a total gymnosperm contribution of 70 % to Lena River dissolved  
13 lignin. Despite a broad range of ratios in our summer TSM and surface sediments, we infer a  
14 substantially lower gymnosperm contribution to particulate OM in the delta surface water and  
15 Buor Khaya Bay surface sediments than further upstream at Zhigansk. This finding clearly  
16 indicates the overprint of TSM signatures by higher contributions of angiosperm OM  
17 contributing to the total TSM load between Zhigansk, located at the taiga-tundra transition  
18 zone, and our sampling sites in the delta. As mentioned above this might be due to the  
19 inefficient transport of POM from distal catchment areas to the delta and its intermediate  
20 storage on floodplains (e.g. Aufdenkampe et al., 2007; 2011; Moreira-Turcq et al., 2013;  
21 Zocatelli et al., 2013). In contrast, dissolved organic matter including dissolved lignin is  
22 transported with the flow of the water, which might lead to a more efficient transport of taiga-  
23 derived DOM to the delta, thus explaining the difference of modeled gymnosperm  
24 contributions by Amon et al. (2012) and our study. The resulting considerable impact of the  
25 northern part of the catchment area to the POM composition is disproportional to its small  
26 spatial extent within the Lena River drainage area. It further implies environmental changes  
27 associated with above average climate warming expected for the high northern latitudes will  
28 most likely increase the disproportional OM input by enhanced permafrost thawing in the  
29 north compared to the southern catchment.

30

## 31 **5 Conclusions**

32 Despite the annual, seasonal, and spatial variability, the distribution of lignin phenols in our  
33 Lena delta surface water TSM samples clearly reflects the main vegetation characteristics of

1 the Lena River catchment. The gymnosperm fraction derived from the taiga covering most of  
2 the catchment and the angiosperm fraction derived predominantly from the northern tundra  
3 zone contribute about equally to the spring and summer TSM samples. However, because of  
4 possible contributions of angiosperm OM from the taiga zone, for example from elevated  
5 treeless areas, the 50% angiosperm vegetation have to be interpreted as the maximum  
6 contribution from tundra zone. Considering the relatively small area covered by tundra  
7 (~12%; e.g. Amon et al., 2012) this still relatively high angiosperm contribution emphasizes  
8 the importance of this small area as organic matter source to the Lena Delta surface water  
9 TSM and Laptev Sea coastal zone, where it adds to the angiosperm OM contributed by  
10 coastal erosion.

11 Based on the low acid to aldehyde ratios of vanillyl and syringyl phenols ( $Ad/Al_{V,S}$ ), the  
12 spring flood sample seems to have organic matter that has undergone a relatively low extent  
13 of degradation and most likely originates from surface soils and fresh vegetation. This could  
14 be due to the fact that particularly in the northern part of the catchment the soils are still  
15 frozen at the time of the spring freshet, which may favors surface erosion by the flood wave.  
16 On the other hand, less degraded OM with low  $Ad/Al_{V,S}$  ratios is also found in ice complex  
17 deposits of Pleistocene age (e.g. Tesi et al., 2014) and does not necessarily have to be derived  
18 from surface soils. The summer TSM samples displayed compositions ( $Ad/Al_{V,S}$ ) consistent  
19 with higher degrees of degradation, and presumably originated from greater soil depths  
20 thawed during the summer months. As the first and third delta terrace bulk soil samples  
21 analyzed here had generally lower  $Ad/Al_{V,S}$  ratios than the summer TSM, we speculate that  
22 there must be an additional more degraded organic matter source. This source could be  
23 organic matter derived from the southern catchment, where annual permafrost thaw depths are  
24 greater than in the Lena Delta. Because these materials are transported to the delta during  
25 lower flow conditions, it is likely they are predominantly composed of finer particles, which  
26 usually contain more highly altered lignin and may have been affected by sorptive processes  
27 with DOM, all of which can contribute to the higher  $Ad/Al_{V,S}$  ratios.

28 The marginal filter leading to flocculation of dissolved and particulate organic matter and  
29 rapid sedimentation seems to be the dominant reason for high lignin contents off the major  
30 delta outlets. Similar to the TSM samples, the lignin distribution within the surface sediments  
31 of the Buor Khaya Bay points to a mixed gymnosperm and angiosperm vegetation source for  
32 organic matter and the modeled contributions are as well about equal for both sources. The  
33 additional contribution of angiosperm-derived OM to Buor Khaya Bay sediments through  
34 coastal erosion makes it difficult to unambiguously distinguish between angiosperm-derived

1 OM from the Lena watershed and from coastal erosion of ice complex deposits. However, as  
2 gymnosperm vegetation is not present in the Lena Delta and along the Buor Khaya coast  
3 today and their respective Holocene and Pleistocene deposits but covers the southern part of  
4 the Lena River catchment, the fact that we find gymnosperm-derived OM in surface  
5 sediments suggests that a substantial amount of sedimentary organic matter in the Buor Khaya  
6 Bay originates from Lena River catchment. The additional source of angiosperm OM  
7 contributed by coastal erosion results in dilution of the gymnosperm signal with distance to  
8 the delta.

9 The surface sediments were strongly degraded resembling the Lena Delta summer samples  
10 and implying at least some summer TSM is transported from the delta to the coastal zone.  
11 However, the strong degradation of sedimentary organic matter close to Muostakh Island  
12 consisting of Pleistocene ice complex and being affected by coastal erosion, which most  
13 likely happened after thawing on land, makes it complicated to distinguish between degraded  
14 ice complex and degraded summer TSM derived organic matter.

15 In the future more severe warming is expected for the high northern latitudes (IPCC, 2013),  
16 which will presumably influence the northernmost part of the Lena River catchment, i.e. the  
17 tundra zone with the delta, stronger than the southern part. On the basis of our data it should  
18 be possible to trace changes in OM contribution and quality from different parts of the Lena  
19 River catchment area. Additionally, more research is needed to investigate the fate of Lena  
20 River and ice complex organic matter, particularly their degradability on land, in the water  
21 column, and post-depositionally to understand their potential for possible increase in  
22 greenhouse gas release from the Arctic.

23

24 Additional data on individual CuO oxidation products for the samples presented here can be  
25 found in PANGAEA ([www.pangaea.de](http://www.pangaea.de)).

26

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12

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9

1 Table 1. Samples presented in this study and analyzed for lignin phenol composition. Bluff  
 2 height is given in meters above river level [m a.r.l.] measured in Aug 2009 and Jul/Aug 2010.  
 3 All total suspended matter samples 2009-2011 were taken from the surface water layer with a  
 4 sampling depth of ca. 0.5m. Additional surface water samples used for total suspended matter  
 5 determination can be found in table S1 in the supplement. Not applicable denoted by n.a.

Sample code	Sample & site description	Date of sampling	Latitude N [dec]	Longitude E [dec]	Bluff height [m a.r.l.]	Water depth [m]
<i>Lena Delta first terrace bluff profiles</i>						
L09-08	Gorgolevsky Island, 3 depths sampled	17-Aug-2009	72.6158	127.2627	3.4	n.a.
L09-12	Samoylov Island, 5 depths sampled	18-Aug-2009	72.3775	126.4954	7.5	n.a.
L09-28-2	Bykovskaya Channel, 2 depths sampled	21-Aug-2009	72.0586	128.6309	1.7	n.a.
L10-04	Baron Belkey Island, 6 depths sampled	31-Jul-2010	72.5378	126.8608	6.5	n.a.
<i>Kurungnakh Island third terrace<sup>a</sup></i>						
S29	unit V, middle Holocene	Aug-2002	72.3447	126.3092	37.0	n.a.
S17	unit IVb, early Holocene	Aug-2002	72.3447	126.3092	37.0	n.a.
S13	unit IVa, Pleistocene ice complex	Aug-2002	72.3447	126.3092	37.0	n.a.
S22D	unit III, Pleistocene ice complex	Aug-2002	72.3447	126.3092	37.0	n.a.
S45	unit III, Pleistocene ice complex	Aug-2002	72.3431	126.3056	37.0	n.a.
<i>Lena River total suspended matter</i>						
4	Lena River main channel south of Tit Ari Island	16-Aug-2009	71.9040	127.2544	n.a.	n.a.
10	Lena River main channel	19-Aug-2009	72.2760	126.9041	n.a.	n.a.
11	Lena River main channel	19-Aug-2009	72.5159	126.7142	n.a.	n.a.
13	Lena River Bykovskaya Channel	20-Aug-2009	72.2352	127.9619	n.a.	n.a.
14	Lena River Bykovskaya Channel	20-Aug-2009	72.0341	128.5232	n.a.	n.a.
16	Lena River Bykovskaya Channel	21-Aug-2009	72.0586	128.6309	n.a.	n.a.
17	offshore Bykovsky Peninsula	22-Aug-2009	71.7889	129.4189	n.a.	n.a.
25	Lena River Trofimovskaya Channel	31-Jul-2010	72.4764	126.6250	n.a.	n.a.
26	Lena River Trofimovskaya Channel	31-Jul-2010	72.4764	126.8588	n.a.	n.a.
27	Lena River main channel south of Samoylov	1-Aug-2010	72.3776	126.7478	n.a.	n.a.
28	Lena River main channel north of Tit Ari Island	1-Aug-2010	72.2102	126.9423	n.a.	n.a.
29	Lena River main channel south of Tit Ari Island	1-Aug-2010	71.9514	127.2582	n.a.	n.a.
30	Lena River main channel off Kurungnakh	2-Aug-2010	72.2808	126.2091	n.a.	n.a.
31	Lena River main channel	2-Aug-2010	72.3567	126.3521	n.a.	n.a.
32	Lena River Bykovskaya Channel	3-Aug-2010	72.3604	127.6761	n.a.	n.a.
37	Lena River main channel off Samoylov Island	29-May-2011	72.3651	126.4757	n.a.	n.a.
<i>Buor Khaya Bay surface sediments</i>						
L09-34	surface sediment (grab sampler) off Muostakh Island	23-Aug-2009	71.5750	129.8200	n.a.	10.5
L10-23	surface sediment (steel tube)	4-Aug-2010	71.7778	130.0872	n.a.	11.5
L10-24	surface sediment (steel tube)	4-Aug-2010	71.9250	130.8227	n.a.	17.0
L10-25	surface sediment (steel tube)	4-Aug-2010	72.0725	131.5896	n.a.	17.0
L10-36	surface sediment (steel tube)	6-Aug-2010	72.7411	130.1324	n.a.	5.8

*Vegetation samples<sup>b</sup>*

09-TIK-04	<i>Aulacomnium turgidum</i>	Jul/Aug 2009	72.8087	124.9121	n.a.	n.a.
09-TIK-01	<i>Carex spp.</i>	Jul/Aug 2009	73.1731	124.5757	n.a.	n.a.
09-TIK-13	<i>Ledum palustre</i>	Jul/Aug 2009	69.3991	123.8261	n.a.	n.a.
09-TIK-13	<i>Betula nana</i>	Jul/Aug 2009	69.3991	123.8261	n.a.	n.a.
09-TIK-13	<i>Salix spp.</i>	Jul/Aug 2009	69.3991	123.8261	n.a.	n.a.
09-TIK-13	<i>Larix</i> (mostly needles)	Jul/Aug 2009	69.3991	123.8261	n.a.	n.a.

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1 <sup>a</sup> samples provided by L. Schirrmeister (Alfred Wegener Institute Potsdam, Germany), data  
2 from Wetterich et al. (2008)

3 <sup>b</sup> samples provided by U. Herzsuh (Alfred Wegener Institute Potsdam, Germany),  
4 expedition field reports by Herzsuh et al. (2009) and Klemm and Zubrzycki (2009)

5

1 Table 2. Total suspended matter (TSM) concentrations in Lena Delta surface waters (2009 to  
 2 2011) and atomic particulate organic carbon (POC) to particulate total nitrogen (PN) ratios.  
 3 Note that there is one sample less for the TSM [mg/L] calculation (n=20) of the TSM August  
 4 2009 data than for POC and POC:PN (both n=21), because one filter weight was missing  
 5 while the volume filtered was known and thus POC:PN could be calculated.

	TSM [mg/L]	POC <sup>a</sup> [mg/L]	POC <sup>a</sup> [wt%]	atomic POC:PN <sup>a</sup>
<i>TSM Aug 2009</i>	<i>n=20</i>	<i>n=21</i>	<i>n=20</i>	<i>n=21</i>
mean	28.50	1.21	7.2	9.6
median	14.94	0.83	4.7	9.2
min	3.10	0.35	1.9	6.8
max	174.92	7.24	37.7	19.3
<i>TSM July/Aug 2010</i>	<i>n=15</i>	<i>n=13</i>	<i>n=13</i>	<i>n=13</i>
mean	19.85	0.57	3.05	7.6
median	19.88	0.47	3.05	7.8
min	3.52	0.15	1.42	3.7
max	32.23	1.30	4.74	10.3
<i>TSM late May 2011</i>				
sample 37	494.00	8.20	1.66	7.5

6 <sup>a</sup> from Winterfeld et al. (2015, submitted as companion paper)

7

8

1 Table 3. Organic carbon (OC), total nitrogen (TN), and atomic OC:TN ratios of the Lena  
 2 Delta soil samples (first and third terrace) and Buor Khaya Bay surface sediments

Sample code	OC [wt%]	TN [wt%]	atomic OC:TN
<i>Lena Delta first terrace bulk, n=19</i>			
mean	7.48	0.21	38.5
median	7.61	0.24	35.1
min	1.02	0.03	21.7
max	17.14	0.45	68.0
<i>Lena Delta third terrace (Kurungnakh Island)<sup>a</sup></i>			
S29 (unit V)	3.76	0.19	19.4
S17 (unit IVb)	1.97	0.38	5.2
S13 (unit IVa)	1.69	0.19	9.1
S22D (unit III)	6.91	0.54	12.8
S45 (unit III)	3.72	0.31	12.1
<i>Buor Khaya Bay surface sediments</i>			
L09-34	2.47	0.18	15.7
L10-23	2.33	0.17	16.4
L10-24	1.88	0.15	14.7
L10-25	1.93	0.16	11.7
L10-36	1.67	0.09	20.9

3 <sup>a</sup> from Wetterich et al. (2008)

4

1 Table 4. Sediment-normalized yields of CuO oxidations products of Lena Delta soils, total  
 2 suspended matter (TSM), surface sediments, and vegetation samples in milligram per gram  
 3 dry weight sediment (mg/g dws). Trivial names of analyzed plant species in brackets. V =  
 4 vanillyl phenols (sum of vanillin, acetovanillone, vanillic acid), S = syringyl phenols (sum of  
 5 syringaldehyde, acetosyringone, syringic acid), C = cinnamyl phenols (sum of *p*-coumaric  
 6 acid, ferulic acid),  $\Sigma 8$  = sum of V, S, and C phenols, P = *p*-hydroxybenzenes (sum of *p*-  
 7 hydroxybenzaldehyde, *p*-hydroxyacetophenone, *p*-hydroxybenzoic acid), Pn = *p*-  
 8 hydroxyacetophenone.

	V	S	C	$\Sigma 8$	P	Pn
	[mg/g dws]					
<i>Lena Delta first terrace bulk, n=19</i>						
mean	0.75	0.74	0.43	1.93	0.84	0.13
median	0.73	0.66	0.31	1.60	0.69	0.10
min	0.04	0.04	0.02	0.09	0.05	0.00
max	2.41	2.82	1.87	7.10	3.68	0.42
<i>Lena Delta third terrace (Kurungnakh Island)</i>						
S29 (unit V)	0.24	0.17	0.21	0.63	0.13	0.04
S17 (unit IVb)	0.17	0.14	0.06	0.37	0.15	0.02
S13 (unit IVa)	0.13	0.11	0.06	0.29	0.12	0.02
S22D (unit III)	0.53	0.69	0.59	1.81	0.97	0.13
S45 (unit III)	0.54	0.73	0.54	1.81	0.62	0.09
<i>TSM Aug 2009, n=7</i>						
mean	0.16	0.07	0.04	0.27	0.14	0.05
median	0.17	0.07	0.03	0.27	0.15	0.05
min	0.10	0.04	0.02	0.17	0.07	0.04
max	0.22	0.17	0.08	0.47	0.21	0.07
<i>TSM July/Aug 2010, n=8</i>						
mean	0.21	0.08	0.04	0.32	0.16	0.04
median	0.20	0.08	0.03	0.31	0.15	0.05
min	0.08	0.03	0.01	0.12	0.07	0.02
max	0.34	0.14	0.06	0.53	0.30	0.06
<i>TSM late May 2011, n=1</i>						
	0.47	0.24	0.12	0.83	0.21	0.24
<i>Buor Khaya Bay surface sediments</i>						

L09-34	0.33	0.14	0.05	0.52	0.13	0.07
L10-23	0.41	0.18	0.06	0.64	0.15	0.08
L10-24	0.16	0.07	0.03	0.25	0.07	0.05
L10-25	0.11	0.05	0.02	0.18	0.05	0.05
L10-36	0.28	0.12	0.04	0.45	0.12	0.07

*Vegetation samples*

<i>Aulacomnium turgidum</i> (moss)	1.57	1.63	1.44	4.64	3.64	1.74
<i>Carex spp.</i> (sedge)	4.13	6.24	6.71	17.08	3.58	0.70
<i>Ledum palustre</i> (wild rosemary)	2.76	2.59	3.62	8.97	3.51	0.82
<i>Betula nana</i> (dwarf birch)	5.78	7.43	3.19	16.40	1.27	0.34
<i>Salix</i> (willow)	6.22	4.21	2.17	12.59	2.19	0.75
<i>Larix</i> needles (larch)	7.93	1.41	7.32	16.66	5.48	1.46

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1 Table 5. Carbon-normalized yields of CuO oxidation products of Lena Delta soils, surface  
 2 water total suspended matter (TSM), and surface sediments in milligram per 100 milligram  
 3 organic carbon [mg/100mg OC] and related lignin parameters. Abbreviations for phenol  
 4 groups are the same as in table 4. Ad/Al<sub>v</sub> = acid to aldehyde ratio of vanillyl phenols, Ad/Al<sub>s</sub>  
 5 = acid to aldehyde ratios of syringyl phenols, C/V = cinnamyl to vanillyl phenols, S/V =  
 6 syringyl to vanillyl phenols, P/V = *p*-hydroxybenzenes to vanillyl phenols, Pn/P = *p*-  
 7 hydroxyacetophenone to *p*-hydroxybenzenes. Not determined denoted by n.d.

	V	S	C	Λ8	P	Pn	Ad/Al <sub>v</sub>	Ad/Al <sub>s</sub>	C/V	S/V	P/V	Pn/P
	[mg/100 mg OC]											
<i>Lena Delta first terrace bulk, n=19</i>												
mean	1.08	1.02	0.54	2.64	0.99	0.14	0.76	0.64	0.53	0.96	1.05	0.14
median	0.91	0.88	0.47	2.18	0.95	0.13	0.77	0.62	0.48	0.92	1.08	0.14
min	0.30	0.31	0.14	0.78	0.33	0.03	0.41	0.37	0.16	0.58	0.24	0.07
max	3.50	3.62	1.69	8.81	2.38	0.27	1.19	1.01	1.16	1.58	1.53	0.22
<i>Lena Delta third terrace (Kurungnakh Island)</i>												
S29 (unit V)	0.73	0.37	0.57	1.66	0.35	0.10	0.43	0.58	0.79	0.51	0.49	0.28
S17 (unit IVb)	1.05	0.55	0.28	1.89	0.75	0.11	0.46	0.51	0.27	0.53	0.71	0.15
S13 (unit IVa)	0.87	0.55	0.33	1.74	0.74	0.12	0.47	0.49	0.38	0.63	0.85	0.16
S22D (unit III)	0.80	0.96	0.86	2.62	1.40	0.18	0.48	0.38	1.07	1.20	1.75	0.13
S45 (unit III)	1.52	1.88	1.46	4.87	1.68	0.24	0.35	0.31	0.96	1.24	1.10	0.14
<i>TSM Aug 2009, n=7</i>												
mean	0.63	0.27	0.13	1.03	0.54	0.05	1.71	0.99	0.21	0.44	0.85	0.10
median	0.62	0.26	0.11	0.99	0.58	0.05	1.36	0.98	0.19	0.41	0.81	0.09
min	0.43	0.17	0.09	0.73	0.29	0.04	0.68	0.52	0.14	0.25	0.65	0.07
max	0.80	0.48	0.24	1.35	0.71	0.07	3.97	1.51	0.39	0.77	1.25	0.13
<i>TSM July/Aug 2010, n=8</i>												
mean	0.70	0.27	0.12	1.09	0.53	0.05	1.36	0.81	0.18	0.38	0.77	0.09
median	0.74	0.30	0.14	1.19	0.51	0.05	1.44	0.87	0.18	0.39	0.81	0.09
min	0.28	0.09	0.05	0.42	0.25	0.02	0.69	0.48	0.15	0.32	0.62	0.07
max	0.93	0.37	0.16	1.44	0.81	0.06	2.02	1.11	0.20	0.45	0.89	0.11
<i>TSM late May 2011, n=1</i>												
	2.94	1.47	0.74	5.16	1.29	0.24	0.32	0.32	0.25	0.50	0.44	0.19
<i>Buor Khaya Bay surface sediments</i>												
L09-34	1.34	0.55	0.21	2.09	0.52	0.07	1.75	1.37	0.15	0.41	0.39	0.13
L10-23	1.74	0.76	0.26	2.76	0.65	0.08	1.36	1.16	0.15	0.44	0.37	0.12
L10-24	0.84	0.36	0.14	1.33	0.37	0.05	1.19	1.06	0.16	0.43	0.44	0.14
L10-25	0.57	0.25	0.12	0.94	0.28	0.05	0.98	0.85	0.21	0.45	0.50	0.16
L10-36	1.68	0.74	0.27	2.68	0.72	0.07	1.10	0.77	0.16	0.44	0.43	0.10

*Vegetation samples*

<i>Aulacomnium turgidum</i> (moss)	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	0.67	0.80	0.92	1.04	2.32	0.48
<i>Carex spp.</i> (sedges)	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	0.23	0.22	1.63	1.51	0.87	0.20
<i>Ledum palustre</i> (wild rosemary)	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	0.41	0.49	1.31	0.94	1.27	0.23
<i>Betula nana</i> (dwarf birch)	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	0.18	0.13	0.55	1.29	0.22	0.27
<i>Salix</i> (willow)	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	0.20	0.24	0.35	0.68	0.35	0.34
<i>Larix</i> needles (larch)	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	0.14	0.87	0.92	0.18	0.69	0.27

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1

1 Table 6. Endmember ratios taken from the literature used for the unmixing model here and  
 2 our calculated relative amounts of V, S, and C phenols. For abbreviations see description in  
 3 table 4 and 5.

Endmember	C/V	S/V	V	S	C
			[%]		
woody gymnosperm	0.04*	0.03*	0.93	0.03	0.04
non-woody gymnosperm (needles)	0.17*	0.04*	0.83	0.03	0.14
woody angiosperm	0.05*	2.42*	0.29	0.70	0.01
non-woody angiosperm (leaves, grasses)	0.7*	0.98*	0.37	0.37	0.26

4 \*endmember ratios from table 4 in Amon et al. (2011) including Hedges and Mann (1979),  
 5 Hedges and Parker (1976), Prokushkin et al. (in preparation), Williams et al. (1998)

6

1 Table 7. Results of unmixing model including relative abundances of V, S, and C phenols,  
 2 median mixing coefficients and gymnosperm to angiosperm ratio. See tables 4 and 5 for  
 3 abbreviations.

sample code	rel. V [%]	rel. S [%]	rel. C [%]	median (500 iterations) mixing coefficients				median total gymnosperm	median total angiosperm	proportion of median gymnosperm/angiosperm
				woody gymnosperm	non-woody gymnosperm	woody angiosperm	non-woody angiosperm			
<i>TSM Aug 2009</i>										
4	0.67	0.23	0.10	0.34	0.27	0.21	0.17	0.61	0.38	1.6
10	0.65	0.24	0.10	0.31	0.27	0.23	0.18	0.58	0.41	1.4
11	0.72	0.18	0.10	0.37	0.31	0.15	0.14	0.69	0.29	2.4
13	0.59	0.28	0.13	0.22	0.26	0.21	0.31	0.47	0.53	0.9
14	0.60	0.27	0.13	0.22	0.26	0.20	0.32	0.48	0.51	0.9
16	0.63	0.25	0.12	0.26	0.28	0.21	0.26	0.53	0.46	1.2
17	0.46	0.36	0.18	0.02	0.22	0.21	0.57	0.24	0.78	0.3
<i>TSM July/Aug 2010</i>										
25	0.64	0.25	0.12	0.27	0.28	0.20	0.25	0.55	0.44	1.2
26	0.64	0.25	0.11	0.29	0.27	0.22	0.20	0.56	0.43	1.3
27	0.62	0.26	0.12	0.24	0.27	0.20	0.28	0.52	0.48	1.1
28	0.65	0.24	0.11	0.30	0.27	0.20	0.22	0.57	0.42	1.4
29	0.68	0.22	0.10	0.33	0.30	0.20	0.16	0.63	0.36	1.8
30	0.64	0.26	0.10	0.30	0.26	0.24	0.19	0.56	0.43	1.3
31	0.60	0.27	0.12	0.24	0.25	0.22	0.29	0.49	0.51	1.0
32	0.66	0.22	0.12	0.28	0.30	0.16	0.25	0.58	0.41	1.4
<i>TSM late May 2011</i>										
37	0.57	0.29	0.14	0.17	0.25	0.19	0.37	0.43	0.57	0.8
<i>Buor Khaya Bay surface sediments</i>										
L09-34	0.64	0.26	0.10	0.31	0.25	0.25	0.18	0.56	0.44	1.3
L10-23	0.63	0.28	0.09	0.30	0.25	0.28	0.17	0.55	0.45	1.2
L10-24	0.63	0.27	0.10	0.28	0.26	0.26	0.20	0.54	0.46	1.2
L10-25	0.61	0.27	0.13	0.23	0.26	0.21	0.29	0.49	0.50	1.0
L10-36	0.63	0.27	0.10	0.29	0.25	0.27	0.19	0.54	0.46	1.2

4

5

## 1 **Figure captions**

2 Figure 1. A) Lena River catchment area with approximate tundra and taiga forest distribution,  
3 B) Lena Delta and Buor Khaya Bay sampling sites from 2009 to 2011 and associated sample  
4 codes.

5 Figure 2. Carbon-normalized yields of phenols groups shown as Whisker plots when the  
6 number of samples was large enough and as individual samples for smaller numbers.

7 Figure 3. Spatial distribution of carbon-normalized lignin concentrations ( $\Lambda_8$ ) and lignin  
8 parameters of Lena Delta total suspended matter (TSM) and Buor Khaya Bay surface  
9 sediments.  $Ad/Al_{vanillyl}$  = acid to aldehyde ratio of the vanillyl phenols,  $C/V$  = ratio of  
10 cinnamyl to vanillyl phenols, and  $S/V$  = ratio of syringyl to vanillyl phenols.

11 Figure 4. A) Lignin degradation indices ( $Ad/Al_V$  vs.  $Ad/Al_S$ ) and B) vegetation source  
12 parameters ( $C/V$  vs.  $S/V$ ) including compositional ranges of major vascular plant types (Goñi  
13 et al., 1998; Goñi and Hedges, 1992; Hedges and Mann, 1979; Hu et al., 1999). For  
14 abbreviations see Fig. 3. Note the different scales. Literature values: <sup>a</sup> end-member values  
15 used in this study taken from table 4 in Amon et al. (2012) and references therein, <sup>b</sup>Lobbés et  
16 al. (2000), <sup>c</sup>Amon et al. (2012). Note in A) the two values on the  $Ad/Al_V$  axis where  $Ad/Al_S$  is  
17 zero, because there were no values given in Lobbés et al. (2000).

18 Figure 5. The ratios of *p*-hydroxyacetophenone to *p*-hydroxybenzenes ( $Pn/P$ ) versus *p*-  
19 hydroxybenzenes to vanillyl phenols ( $P/V$ ) of samples analyzed in this study and values from  
20 the literature used as indicator for moss contributions. <sup>a</sup>Amon et al. (2012), <sup>b</sup>table 4 in Amon  
21 et al. (2012) and references therein, <sup>c</sup>Williams et al. (1998).

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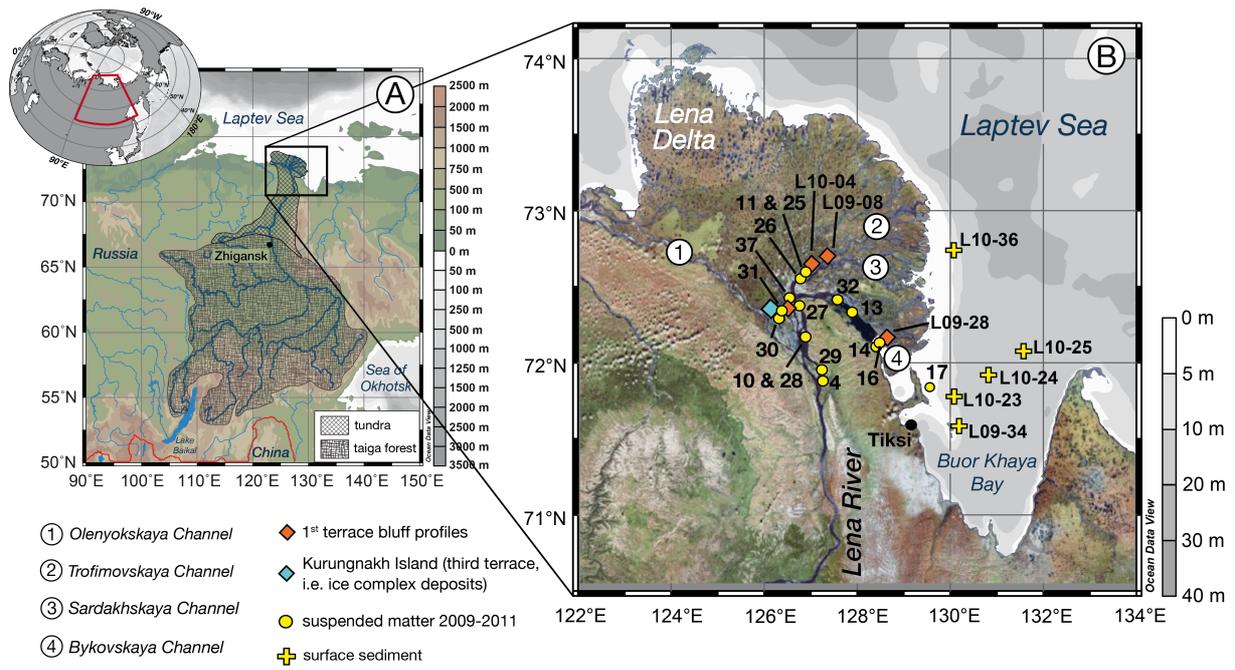
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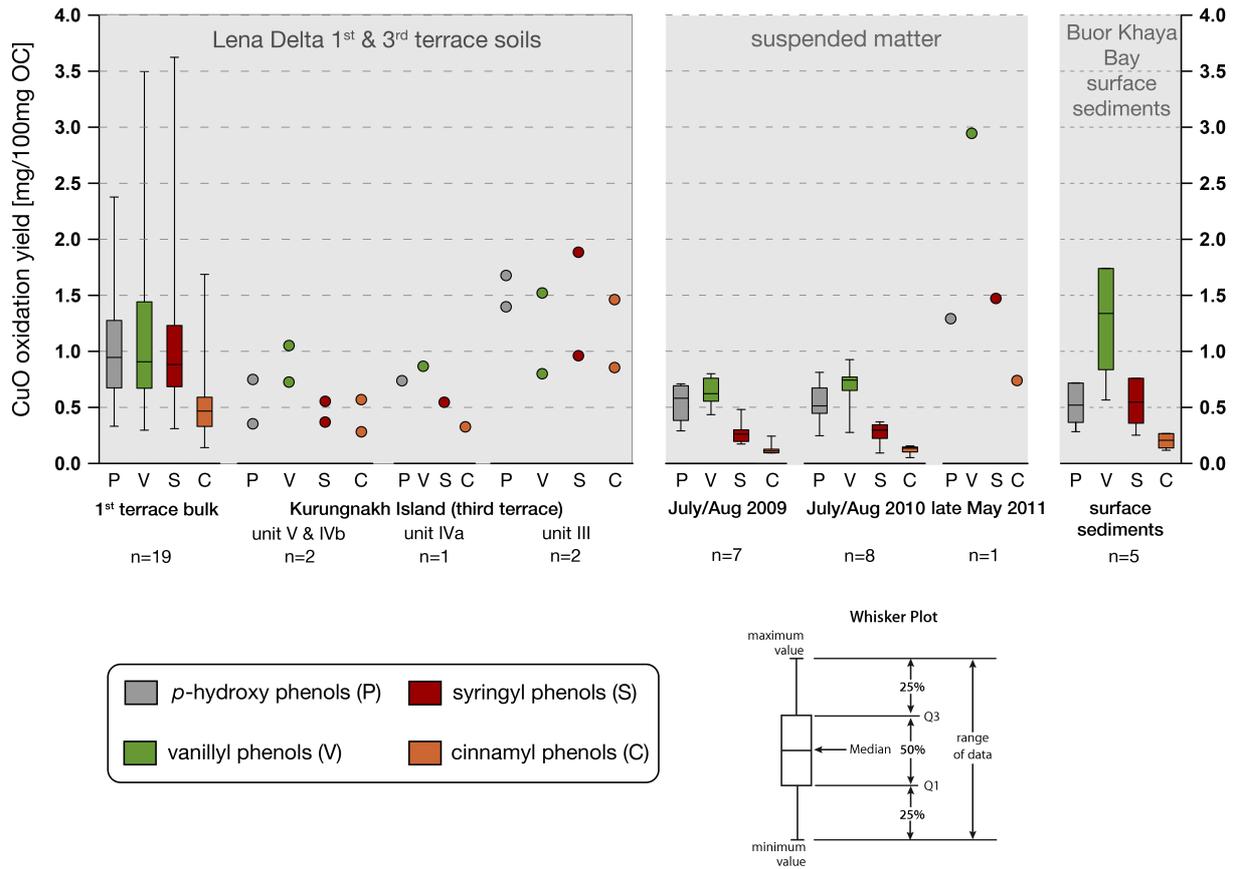
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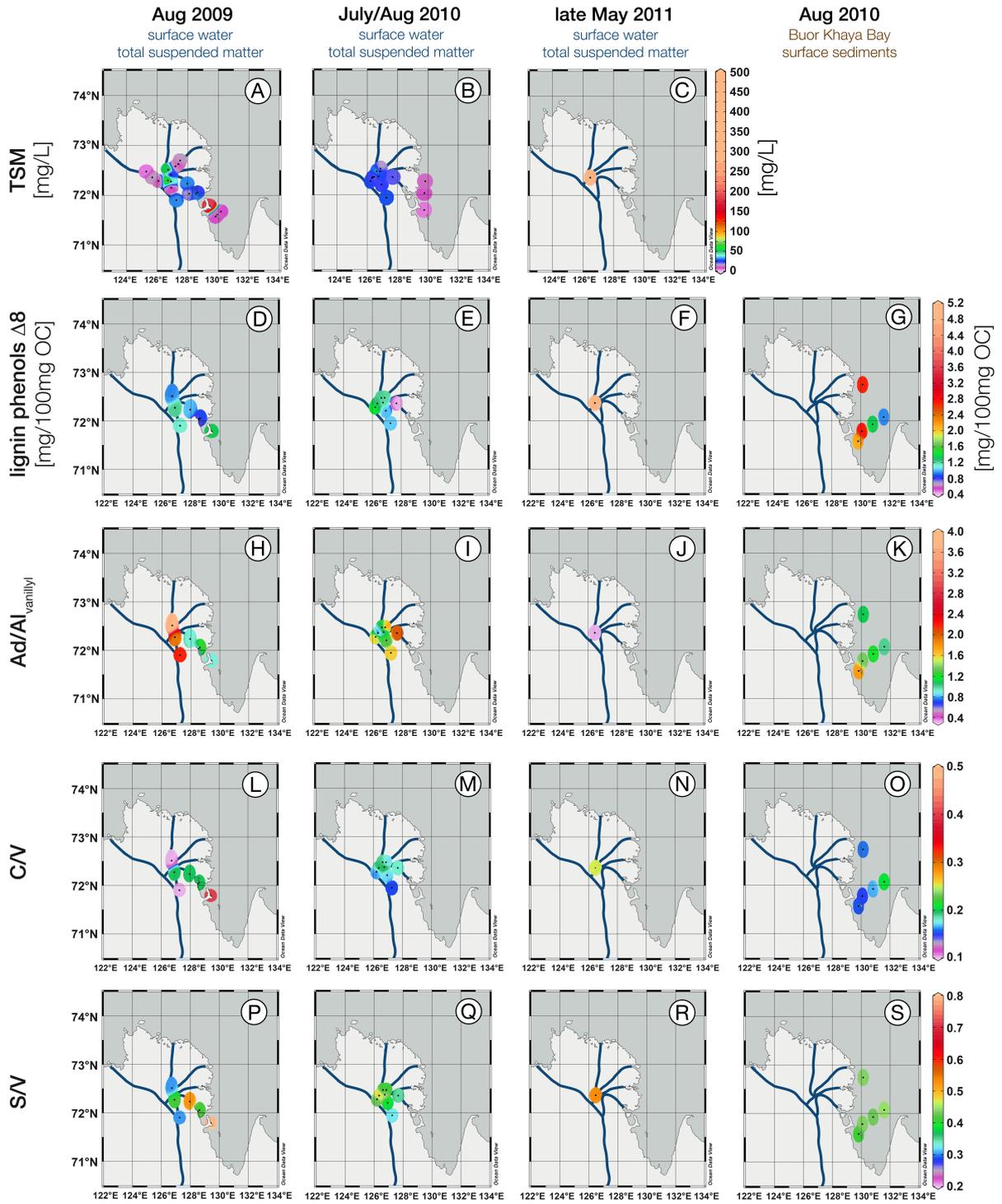
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2 Fig. 1



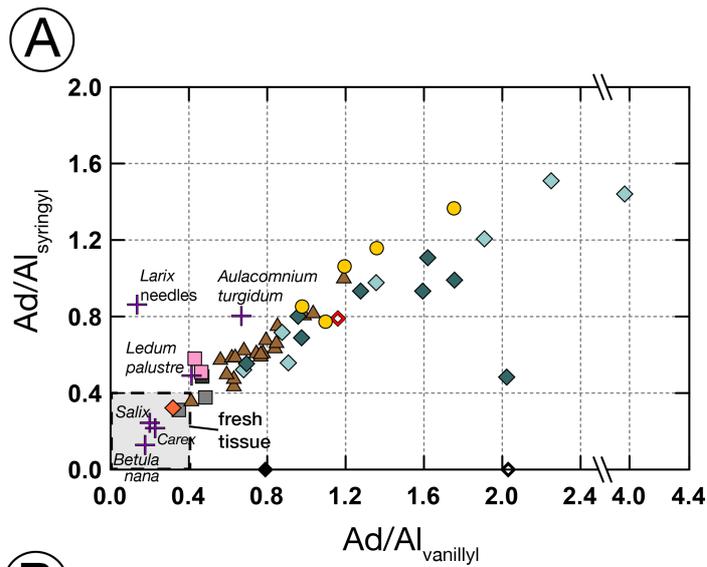
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2 Fig. 2



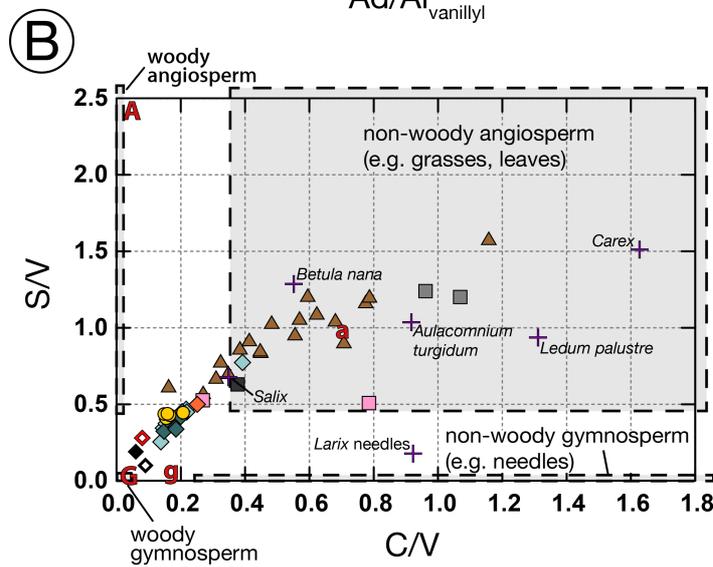
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2 Fig. 3



this study:

- ▲ Lena Delta 1<sup>st</sup> terrace bulk
- Kurungnakh Island unit IVb & V
- Kurungnakh Island unit IVa
- Kurungnakh Island unit III
- ◆ suspended matter Aug 2009
- ◆ suspended matter July/Aug 2010
- ◆ suspended matter late May 2011
- Buor Khaya Bay surface sediments
- + vegetation samples



literature values:

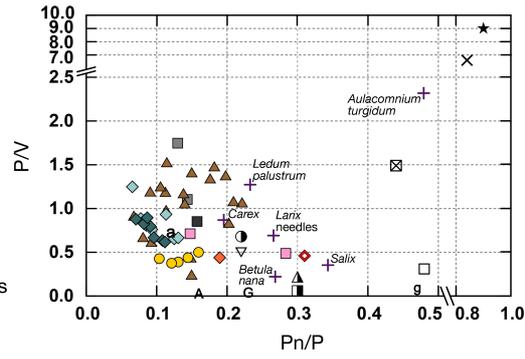
- G woody gymnosperm<sup>a</sup>
- g non-woody gymnosperm<sup>a</sup>
- A woody angiosperm<sup>a</sup>
- a non-woody angiosperm<sup>a</sup>
- ◆ Lena Delta DOC<sup>b</sup>
- ◆ Lena Delta POC<sup>b</sup>
- ◆ Lena River DOC (at Zhigansk)<sup>c</sup>

1

2 Fi.g 4

this study:

- ▲ Lena Delta 1<sup>st</sup> terrace bulk
- Kurungnakh Island unit IVb & V
- Kurungnakh Island unit IVa
- Kurungnakh Island unit III
- ◇ suspended matter Aug 2009
- ◆ suspended matter July/Aug 2010
- ◇ suspended matter late May 2011
- Buor Khaya Bay surface sediments
- + vegetation samples



literature values:

- ◇ Lena River DOC (at Zhigansk)<sup>a</sup>
- G woody gymnosperm<sup>b</sup>
- g non-woody gymnosperm<sup>b</sup>
- A woody angiosperm<sup>b</sup>
- a non-woody angiosperm<sup>b</sup>
- grasses<sup>b</sup>
- ★ moss<sup>b</sup>
- ▽ wetland plants<sup>b</sup>
- X peat (Sphagnum sp.)<sup>b</sup>
- ⊠ peat<sup>c</sup>
- ▲ boreal forest soil org. h.<sup>b</sup>
- DOM soil<sup>b</sup>
- DOM boreal lakes<sup>b</sup>

1

2 Fig. 5