

n-Alkane-based reconstructions of peat accumulations and depositional conditions at four locations around a shallow maar lake in the Changbai Mountains, northeastern China

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Key points:

- *n*-Alkane-based reconstruction of peat accumulation history around a maar lake.
- Hydrological conditions and plant communities related to lake basin caused by volcanism lead to unsynchronized peat depositional processes.
- Volcanism and anthropogenic have important impacts on local peat development.

Abstract

A high-resolution *n*-alkane biomarker study of peat cores from four locations around the Yuanchi maar lake in the Changbai Mountains of northeastern China, has revealed different histories of peat deposition in the closely located sequences, although they experienced the same paleoclimate changes. Comparisons of the *n*-alkane distributions of modern plants around the lake and those in the peat cores suggest that the disparate peat development patterns in the four sites are the consequence of different peat-forming communities growing around the lake. These floral differences were in turn controlled by different water depths associated closely with the volcanogenic lava- and tephra-shaped topography of the lake basin. Moreover, the *n*-alkane-inferred variations in peat development patterns and inferred recent climate changes around Yuanchi Lake relate closely to the histories of volcanic forcing in the tropical Pacific and local volcanic eruptions of the Changbai Mountains. These events evidently led to alterations of local climate that affected growth of land plants at the four locations. Finally, anthropogenic impacts since 1950 CE have had an additional effect on peat accumulation in the Changbai Mountains region.

Keywords: peat deposition, *n*-alkanes, local conditions, volcanic eruption, anthropogenic activities, Changbai Mountains

1. Introduction

Peat is the partially preserved remains of former plant communities that contains a high proportion of organic matter (OM). Peatlands therefore can be either sources or sinks of carbon depending on environmental conditions (Kayranli et al., 2010; Naafs, et al., 2019). Different types of peatlands are defined by their distinctive combinations of hydrology and chemistry that support characteristic plant communities (Andersson et al., 2012). Lipid biomarkers have special features of relative source specificity and resistance to decomposition. The composition and abundance of biomarkers in peat are mainly derived from waxes of plants that grew where the peat accumulated (e.g. Nichols et al., 2006; Zheng et al., 2007; Zhou et al., 2005; Andersson et al., 2011; Zhang et al., 2014, 2016, 2017). Lipid biomarkers in continuous peat cores consequently can provide information on variations in former plant communities that contributed to peat deposition and provide site-specific records of *in situ* environmental histories (e.g. Regnery et al., 2013; Zhang et al., 2018; Naafs et al., 2019).

Peatlands are widely distributed in northeast China. Many studies have focused on them to provide detailed paleoenvironmental reconstructions on centennial and millennial age-scales (e.g. Hong et al., 2001; Zhou et al., 2005; Zhang et al., 2015, 2016; Zhang et al., 2014, 2017). The Changbai Mountains in northeastern China are comprised of a group of volcanoes. Most peatlands in these mountains have developed in lava-dammed lakes or shallow crater lakes, and apparent differences in the initial time and development of peat deposition exist between the two types of volcanogenic landforms (Yang, 1995; Xing et al., 2019). Unlike the Jinchuan and Hani peatlands that

evolved from lava-dammed lakes and began to accumulate before 10 ka, more recent peatlands such as the Yuanchi and Chichi peatlands developed from maar lakes that formed in shallow and irregular depressions resulting from inactive volcanic craters. Therefore, both the accumulation time and the thickness of the peat deposited in those regions are smaller than those in the peatlands that evolved from lava-dammed lakes (Yang et al., 1995). To date, studies of the longer-lived peatlands have yielded valuable information about the long-term regional climate changes (e.g. Hong et al., 2001; Seki et al., 2009; Zhou et al., 2010). However, little is known about modern peat deposition process and its possible controlling factors in the peatlands that derived from the shallow crater lakes in the Changbai Mountains.

Even small volcanic eruptions commonly inject large quantities of aerosols and ash into the atmosphere that alter the radiative balance and chemical equilibrium of the stratosphere, thereby impacting regional climate (Xu et al., 2012; Sun et al., 2014, 2015). A Plinian eruption in the Changbai Mountains around 1000 years ago (the Millennium Eruption), which was accompanied by an earthquake with a magnitude of ~7, was one of the most violent eruptions in history (Sun et al., 2014, 2015). Hereafter, three small-scale volcanic eruptions, in 1668 CE, 1702 CE, and 1903 CE, also documented in the Mountains (Xu et al., 2012). In addition, population growth and economic development around the Changbai Mountains area in recent years also have had a significant impact on local conditions (Gao et al., 2016; Bao et al., 2019). These volcanism and anthropogenic impacts on local conditions consequently could have had important impacts on plant growth that directly affected peat accumulations. However,

scant information is available about details of the peat deposition process and its potential responses to volcanic or anthropogenic activities in this region.

We present here a well-resolved record of the amounts and distributions of *n*-alkane biomarkers in modern plant communities and associated peat cores from four different sites around a modern peatland that evolved from a shallow maar lake (Yuanchi Lake) in the Changbai Mountains. The main objectives of this study are (1) to reconstruct a decade-scale history of peat-forming communities that influenced peat deposition around the maar lake, (2) to understand the localized responses of peat depositional processes to variations in plant assemblages and local conditions caused by volcanic eruptions, and (3) to explore the influences of volcanism and modern anthropogenic impacts on peat deposition in this region.

2. Materials and methods

2.1 Study area

The Changbai Mountains are in the Jilin Province of northeastern China and extend along the boundary between China and North Korea. The area has a continental monsoon climate with long, cold winters and short, cool summers. More detailed descriptions on the regional character have been described by Bao et al. (2010) and Gao et al. (2016). Many lakes that formed from volcanic dams and in volcanic craters exist in this region. Both the climate conditions and the topographic features in the region result in extensive development of peatlands from lava-dammed lakes or shallow crater lakes (Chai, 1990).

Yuanchi Lake (42°01'52"~42°01'58" N, 128°26'03"~128°26'10" E; elevation ca. 1280

m asl) has a diameter of 180 m and a catchment area of 4.1 hm², located on the eastern slopes and about 30 km from the main peak of the Changbai Mountains (Figs.1 and 2). It partially fills a volcanic crater and has no significant inflow or outflow from surface water or groundwater (Yang, 1995; Bao et al., 2010). Peatlands are distributed around the perimeter of the lake and are populated by different peat-forming plant communities (Fig.2). The modern biomes in the surroundings of the lake are trees and shrubs, including *L. olgensis*, *B. fruticosa*, *V. uliginosum*, *R. parvifolium* and *L. palustre*. The plant community on the west side of YC lake (YC_W) is dominated by *C. lasiocarpa* and *Sphagnum* spp, and some *P. australis* and *S. triqueter* grow near the lake shore. On the south edge of lake (YC_S), the peat-forming plants are dominated by *C. lasiocarpa*, and extensive developments of emerged vascular plants (*T. orientalis* and *P. australis*) are in this location. On the east side of the lake (YC_E), the peat-forming plants are dominated by shrubs, including *V. uliginosum*, *R. parvifolium*, and *L. palustre*, and some *Sphagnum* mounds are distributed among the shrubs. The north side of the lake (YC_N) is dominated by *Sphagnum* spp. Large amounts of shrubs (*V. uliginosum* and *Rhododendron parvifolium*) mixed with *L. olgensis*, *B. fruticosa* and *Lonicera caerulea* also appear in this location.

2.2. Sample collection and stratigraphy

Four peat-mud cores were collected around the edges of Yuanchi Lake (YC_E, YC_W, YC_S and YC_N) with a Wardenaar peat sampler in September, 2006. The lithologies of each core show that the cores contain distinctive stratigraphic sequences based on texture, peat color, plant remains, and TOC contents (Fig.3). For YC_W, the top 8 cm

consists of brown peat having a small amount of partially decomposed *Carex* remains, then a layer from 8 cm to 29 cm containing dark brown peat with a high degree of decomposition, and finally black mud underlying the peat sequence. In the YCs core, the peat layer in the top 13 cm is composed of brown peat, the interval from 13 cm to 28 cm is dark brown peat, and black mud occupies the bottom of the core. For YC_E, the brown peat layer comprises the top 8 cm, dark brown peat with higher degree of decomposition occupies the interval from 7 to 35 cm, and black mud is present below 35 cm. In the YC_N core, a brown moss-peat layer dominated by *Sphagnum* spp. constitutes the top 14 cm, then a brown peat layer occupies the interval from 14 to 21 cm, the interval from 21 cm to 28 cm is dark brown peat with a high degree of decomposition, then the black mud occupies the bottom of the core. The cores were subsampled on-site by slicing into 1 cm intervals that were stored in polyethylene bags for transport to the laboratory for analysis.

2.3. Chronology of the peat-mud sequence

The peat cores were dated at 1 cm intervals by ²¹⁰Pb analysis. A collective total of 120 peat samples from the four cores were analyzed using a low-background γ -ray spectrometer with a high-purity Ge semiconductor (ORTEC Instruments Ltd., USA) at the State Key Laboratory of Lake Science and Environment, Nanjing Institute of Geography and Limnology, CAS. The radioisotope results and the age-depth model were described in more detail by Bao et al. (2010). Peat accumulation rates (PAR, cm/yr) were estimated from the 1 cm depth intervals and the corresponding ²¹⁰Pb ages.

2.4. Laboratory analysis

The peat samples taken at 1 cm intervals from the peat cores were dried at 105 °C and then combusted in a muffle furnace at 550 °C for 4h to determine their loss on ignition (LOI) values (Heiri et al., 2001). TOC concentration of each sample was estimated from OM content expressed as LOI values by multiplying the LOI by 0.5, and organic carbon accumulation rates (CAR, g C/m²/yr) were calculated from the TOC concentrations and the PAR values (Bao et al., 2010).

For *n*-alkane analyses, air-dried samples (1g) were extracted three times with MeOH/dichloromethane (1:1, v:v) for 15 min in an accelerated solvent extractor. The three extracts were combined after filtration and concentrated using a rotary evaporator. The *n*-alkanes were isolated using silica gel column chromatography by elution with 20 ml *n*-hexane. The eluate was concentrated with a N₂ stream and transferred to vials for gas chromatography-mass spectrometry (GC-MS) analysis. Details of the *n*-alkane analysis and determination are described by Zhang et al. (2014, 2016).

2.5 *n*-Alkane proxies

We employ a suit of *n*-alkane molecular ratios as proxies to identify changes in the origins of the biomarker molecules and in their relative amounts of preservation, two parameters that are commonly sensitive to climate changes. These ratios include the total *n*-alkane concentrations normalized to TOC (µg total *n*-alkanes/g TOC), carbon preference index (CPI), average chain length (ACL), and the proportion of aquatic components (P_{aq}). These derived ratios collectively summarize variations of *n*-alkane distributions in the peat sequences (e.g. Naafs et al., 2019). The *n*-alkane CPI values reflecting the odd-over-even predominance of the carbon-chain-length have been

interpreted as an indicator of extensive microbial degradation of *n*-alkanes in peat sequences (Zhou et al., 2010). The *n*-alkane ACL describes the weighted mean lipid chain length and reflects the dominant chain-lengths of the *n*-alkanes. We calculated the CPI and ACL values over the carbon atom range of 20-33. The P_{aq} was defined using lacustrine vegetation (Ficken et al., 2000), but it is also used to reconstruct the proportions of aquatic plants or mosses in peat, which is a parameter that indicates peat hydrological conditions (e.g, Zhou et al., 2005, Nichols et al., 2006; Zhang et al., 2016, 2017).

3. Results

3.1 The age-depth chronologies, TOC concentrations and PARs of the four peat sequences

The YC_w age-depth scale indicates that the 29-cm thick peat layer in this sequence was deposited during the past ca. 400 years, from 1600 to 2000 CE (Fig.3a). The TOC concentration ranges from 19.5 to 48.2%, with a mean value of $40.3 \pm 8.5\%$. Continuous increases in TOC concentrations occur in the bottom of the peat core, and notably high and stable values (mean $44.6 \pm 2.9\%$) appear in the upper 20-cm section that corresponds to peat deposition since 1800 CE (Fig.3b). PAR varies from 0.03 to 0.33 cm/yr in the YC_w sequence (Fig.3b). The rate is initially very slow (~ 0.04 cm/yr) and increases slightly to 0.1 cm/yr in the interval from 20 to 13 cm (from 1800 to 1900 CE). Then the rates increase markedly and peak at 8 cm (ca.1950 CE).

The peat in the upper 28 cm of the YCs sequence was deposited during the last 200 years, from 1800 to 2000 CE (Fig.3c). TOC concentrations range from 3.4 to 39.5 %,

with a mean value of $28.8 \pm 10.5\%$. Relatively higher TOC values (mean $31.6 \pm 7.0\%$) occur in the top 25 cm, corresponding to deposition since 1850 CE (Fig.3d). PAR ranges between 0.04 and 0.33 cm/yr in the YC_s peat section (Fig.3d). Lower rates less than 0.1 cm/yr occur below 25 cm, then the rate increases gradually to a maximum (0.33 cm/yr) at 8 cm (ca. 1950 CE) before decreasing sharply to ~ 0.1 cm/yr in the top 5 cm of the core.

The age-depth scale of the YC_E sequence indicates that the peat in the top 35 cm was deposited during the past ca. 400 years, from 1600 to 2000 CE (Fig.3e). The TOC concentrations in the peat vary between 10.7 and 51.1%, with a mean value of $33.6 \pm 8.3\%$. High values ($>30\%$) and a gradually increasing trend starting at 28 cm correspond to accumulation since 1800 CE (Fig.3f). The PAR in the YC_E peat section fluctuates between 0.04 and 0.59 cm/yr (Fig.3f). Markedly lower values occur below 28 cm (before 1800 CE), then the rates start to increase slightly but fluctuate strongly, and relatively higher values appear in the upper 8 cm (since 1950 CE).

The YC_N peat layer was deposited during the past ca. 250 years, from 1750 to 2000 CE (Fig.3g). The TOC concentrations range from 13.1 to 51.4% (mean $38.4 \pm 11.5\%$), with obviously high and ever-increasing values (mean $43.2 \pm 6.5\%$) in the upper 25 cm that corresponds to peat deposition since 1800 CE (Fig.3h). The PAR has lower values at the bottom of this peat section below 25 cm, and higher values occur from 22 to 19 cm. Markedly lower values (~ 0.08 cm/yr) appear again from 19 to 10 cm (1850 to 1950 CE) before increasing markedly in the top 10 cm (since 1950 CE) (Fig.3h).

3.2. *n*-Alkane distributions

The *n*-alkane distributions from the different layers in the four peat sequences range similarly from C₂₀ to C₃₃ and have an odd/even predominance with a dominance of middle and long chain components (Fig.4). The *n*-alkane distributions in the whole YC_w peat core have a strong dominance of the C₂₇ component, and *n*-C₂₃ also contributes high proportions in the upper and bottom peat units. The YC_s peat core is characterized by strong dominances of C₂₃, C₂₅ and C₂₇ *n*-alkanes in different peat layers. For the YC_E peat core, the distribution in the brown peat unit is dominated by the C₂₃ and C₃₁ *n*-alkanes and in the dark brown peat by the C₂₇ and C₃₁ *n*-alkanes. In the YC_N peat core, long-chain C₂₉ and C₃₁ *n*-alkanes are the major components in the upper unit and the bottom of the core, and the middle brown moss peat layer has abundant middle chain C₂₃ and C₂₅ in subequal proportions with the long chain *n*-alkanes.

3.3. *n*-Alkane concentrations and their molecular ratios

We focus on *n*-alkane-based reconstruction of the peat deposition history and paleoenvironmental information in the Yuanchi Lake region. Here we present the results of the changes in TOC concentrations, the PARs, the CARs, and the *n*-alkane concentrations and their derived molecular climate proxies (CPI, ACL and P_{aq}) for the four peat sequences that we collected around Yuanchi Lake (Figs.5-8).

The total *n*-alkane concentrations normalized to TOC in the YC_w core range between 42.2 and 595.7 µg/g TOC, with a mean value of 188.5±176.3 µg/g. Markedly higher values occur before 1800 CE and lower values appear from 1800 CE to present (Fig.5c). The *n*-alkane CPI varies between 2.1 and 4.2, with a mean value of 3.2±0.5 (Fig.5d). ACL and P_{aq} values range from 24.3 to 26.4 (mean 25.6±0.5) and 0.5 to 0.9 (mean

0.7±0.09), respectively (Fig.5e and f). Gradually increasing CPIs exist before 1800 CE, and lower ACL but higher P_{aq} values occur in this interval. Markedly decreasing CPIs, higher ACL but lower P_{aq} values appear from 1800 to 1900 CE, corresponding to the highest TOC concentrations (Fig.5a). The interval from 1940 to present is characterized by the lowest CPI and ACL values and the highest P_{aq} value, coinciding with the higher TOC concentrations and higher PAR and CAR values (Fig.5a and b).

In the YC_S sequence, the total *n*-alkane concentrations range between 56.6 and 529.4 µg/g TOC, with a mean value of 226.1±134.6 µg/g (Fig.6c). Relatively higher concentrations appear before 1850 CE, and the values decrease sharply to the minimum at ca. 1900 CE. Then higher values appear again in the interval between 1900 and 1950 CE, and stably lower values occur from 1950 CE to the present. This variation generally corresponds to the PAR changes in those time (Fig.6 and c). The *n*-alkane CPIs vary between 2.6 and 4.9 (mean 3.5±0.6), and ACLs vary between 24.5 and 26 (mean 25.4±0.5), and the P_{aq} values change inversely to the ACL values, ranging from 0.6 to 0.85 (mean 0.7±0.07) (Fig.6d-f). Markedly lower ACLs (<25) but higher P_{aq} values (>0.7) appear before 1850 CE and during the interval between 1900 and 1950 CE, corresponding to the relatively higher PAR, CAR and total *n*-alkane concentrations in these times (Fig.6b and c).

In the YC_E sequence, the total *n*-alkane concentrations range from 3.0 to 482.8 µg/g TOC, with a mean value of 83.8±103.4 µg/g (Fig.7c). The concentrations are generally higher (~120 µg/g) before 1900 CE, but the values are almost below 50 µg/g TOC from 1900 to the present. The *n*-alkane CPIs vary widely between 1.3 and 10.7 (mean

3.8±1.9), and most of them are lower than 4.0 before 1950 CE, except for the relatively higher values at ca. 1825 CE and 1900 CE, and in the interval from 1950 to the present (Fig.7d). ACLs and P_{aq} values fluctuate between 24.4 and 28.3 (mean 26.2±1.0), and 0.3 and 1.0 (mean 0.5±0.2), respectively (Fig.7e and f). Relatively higher ACLs (>26) but lower P_{aq} values (<0.4) appear before 1800 CE, and lower ACLs but higher P_{aq} values occur during the intervals from 1800 to 1950 CE. The ACLs then increase dramatically, in contrast to CPIs and P_{aq} values that decrease markedly during the recent decades (Fig.7d-f), coinciding with the highest TOC, PAR and CAR values but with the lowest *n*-alkane concentrations (Fig.5a-c).

The total *n*-alkane concentrations in the YC_N sequence range from 3.7 to 570 µg/g TOC, with a mean value of 172.4±182.8 µg/g (Fig.8c). Markedly higher concentrations (~400 µg/g TOC) appear before 1850 CE, accompanied with higher PARs but lower CARs (Fig.8b); then the values decrease rapidly and most of them below 100 µg/g TOC during the interval from 1850 CE to the present. The *n*-alkane CPIs vary between 1.8 and 5.5 (mean 4.0±0.9), with relatively high and stable values (mean 4.4±0.7) before 1900 CE (Fig.8d). ACL and P_{aq} values fluctuate between 26.3 and 29.7 (mean 27.8±1.0), and between 0.17 and 0.5 (mean 0.3±0.1), respectively (Fig.8e and f). The highest ACL and the lowest P_{aq} values appear before 1800 CE, and relatively lower ACLs (< 27) but higher P_{aq} values (> 0.4) occur during the intervals from 1800 to 1900 CE; Then markedly higher ACLs (>27.5) but lower P_{aq} values (<0.3) appear again from 1900 CE, which correspond with gradual increases in both CARs and PARs (Fig.8b).

4. Discussion

4.1 Peat TOC concentrations and peat accumulation processes around the Yuanchi Lake

Yuanchi Lake is in a circular shallow crater that was formed ca. 1060±100 a B.P (Yang et al., 1995) and was likely related to the Millennium Eruption. The peat layers were deposited in the basin of the maar lake over sediments that are represented by the basal black muds in the four cores (Fig.3). The peat sequences provide records of the succession of peat-forming vegetation that has partially filled the maar lake and evidence of conditions important to the production and preservation of the plant remains (Fig.3). Peat OM is mainly derived from the remains of local plants and provides information about the kinds and amounts of vegetation that has lived in a specific location and the processes that affect peat preservation (Chai, 1990; Meyers and Ishiwatari, 1993; Meyers, 2003; Zhang et al, 2016, 2017). Yuanchi Lake has been filled by ingrowth of peatlands around its edges that have been spreading gradually toward the center of the lake (Yang et al., 1995). The peat ²¹⁰Pb-based age-depth models of the four sequences around Yuanchi Lake indicate the initial contributions of TOC-rich vascular plants inputs in the four sites are asynchronous (Fig.3). We hypothesize that this process of transformation of a shallow lake into a peatland first started in the YC_w and YC_E locations ca. 1600 CE, followed by the YC_N ca.1750 CE, with the YC_s starting to infill last since ca.1800 CE (Fig.3).

Although the total histories of peat sequences differ at the four sites, their initiation times for abundant peat accumulations are similar as indicated by the collectively increasing PAR values at ca. 1800 CE (Fig.3). TOC concentrations and PARs in the four

peat sections exhibit increasing trends from 1800 to 1950 CE, suggesting a continuous
 and growing input of TOC-rich vascular plants promoted peat accumulation around
 Yuanchi Lake. However, we observe that the mean TOC concentration (~44.6%) in the
 YC_w peat sequence is higher than in the other three locations (mean 28.8% at YC_s, 36%
 at YC_E, and 39% at YC_N) since 1800 CE. Nevertheless, mean PAR in the YC_w peat
 sequence is slightly lower (~0.12 cm/yr) than those values in the other three cores (mean
 0.17 cm/yr in YC_s, 0.22 cm/yr in YC_E, and 0.17 cm/yr in YC_N peat cores) (Fig.3). In
 addition, differences in peat accumulation between the four locations is evident since
 1950 CE, when marked decreases in PAR values occurred in the YC_w and YC_s locations,
 and the rates in the YC_E and YC_N peat sequences showed rapidly increasing PARs during
 recent decades (Fig.3). We also observe that the four different locations around Yuanchi
 Lake are dominated by different vegetation communities (Fig.2), representing settings
 with different water depths that are likely responsible for the asynchronous peat
 accumulations. Except for *C. lasiocarpa*, the modern vegetation communities in the YC_w
 and YC_s sites are dominated by emerged aquatic vascular plants (*T. orientalis* and *P.*
australis) that could contribute to higher OM inputs but would be less likely to increase
 peat accumulate in recent decades as the water became shallower. Moreover, modern
 vegetation communities at the YC_E and YC_N sites are dominated by *Sphagnum* spp and
 shrubs growing in shallow water depths that are more likely to promote peat
 accumulation than *Carex* and the emerged aquatic vascular plants that dominate the YC_w
 and YC_s sites in deeper-water conditions (Figs.2 and 3).

4.2. Biomarker distributions in the Yuanchi peat sequences

4.2.1. *n*-Alkane biomarkers of modern plants around Yuanchi Lake

n-Alkane biomarkers in peat originate mainly from plant waxes and bacteria and are recognized as important proxies for indicating changes in the plant communities and reflecting any alterations from *in situ* processes that peat deposits may experience after their formation (Zhang et al., 2018). The *n*-alkane distribution patterns vary considerably among different peat-forming plant species, a feature that provides important evidence of the main sources of these biomarkers in the peat (e.g. Kirkels et al., 2013; Zhang et al., 2016, 2017). The dominant plants observed at the four sites around Yuanchi lake are clearly different (Fig.2). We compiled published information about the dominant *n*-alkane homologues found in modern plant species like those around the lake to interpret the main sources of the *n*-alkanes in the peat and thereby to provide a biomarker record for reconstructing the *in situ* conditions at the four sites (Table 1).

The modern plants in the YC_W site are dominated by *C. lasiocarpa*, *S. triqueter*, and *P. australis*, all of which have a predominance of *n*-C₂₉ in their *n*-alkane distributions (Cranwell et al., 1984; Street et al., 2013; Ronkainen et al., 2013). Some *Sphagnum* spp characterized by prominent abundances of C₂₃ and C₂₅ *n*-alkanes (Cranwell et al., 1984; Street et al., 2013) also develop in this site. The *C. lasiocarpa* and *E. fauriei*, which are respectively dominated by *n*-C₂₉ and *n*-C₂₇ (Tarasov et al., 2013; Ronkainen et al., 2013), are the major modern species growing at the YC_S site. In addition, abundant emerged aquatic vascular plants (*T. orientalis* and *P. australis*), both having a predominance of *n*-C₂₉, and some submerged aquatic plants with prominent abundances of C₂₃ and C₂₅ *n*-alkanes (Cranwell et al., 1984; Ficken et al., 2000) are also prevalent in this site under

deeper water conditions. The modern vegetation covering both the YC_E and YC_N sites is mainly dominated by *Sphagnum* spp and shrubs (*V. uliginosum* and *R. parvifolium*) under drier condition that are characterized by *n*-C₂₉ and *n*-C₃₁ (Salasoo, 1987; Tarasov et al., 2013; Street et al., 2013). Moreover, the two sites are surrounded by trees (*L. olgensis*, *B. fruticosa* and *L. caerulea*) that have a major predominance of the C₂₇ *n*-alkane (Tarasov et al., 2013).

4.2.2 Biomarker origins in each of the peat cores from Yuanchi Lake

The chain-length distributions of *n*-alkanes in the four peat sequences range from C₂₀ to C₃₃ with an odd/even predominance (Fig.3), indicating a major input from terrigenous plant waxes (Eglinton et al., 1967). The dominant *n*-alkane components have different distributions among the four sequences, which can be attributed to origins from different plant types associated with different water levels in their growth locations. Based on the dominant *n*-alkane homologues found in the modern plant covering on the four sites (Table 1), we can infer probable origins of the biomarkers in each of the peat cores (Fig.4).

The strong dominance of *n*-alkane C₂₃, C₂₅ and C₂₇ components in the bottom peat units in the YC_w peat core mainly originated from a combination of aquatic plants in a deeper water lake setting and trees from surrounding area. Shallower water along the western edge of the lake and larger amounts of terrestrial plant contributions resulted in the strong dominance of long-chain *n*-alkane in the middle section of the peat core (Fig.4). The abundances of C₂₃ and C₂₅ components that occur in the upper section is likely associated with increased amounts of *Sphagnum* spp input in recent years. A

strong dominance of C₂₃, C₂₅ and C₂₇ *n*-alkanes in the YC_s peat core relate to major contributions from both aquatic plants under high water level condition and terrestrial plants around the area. The *n*-alkane distributions in the YC_E and YC_N peat cores have the long-chain C₂₉ and C₃₁ *n*-alkanes that are diagnostic of shrub communities in these peat sites (Table 1). In addition, significant proportions of C₂₃ and C₂₅ *n*-alkanes in the middle and upper sections of the two peat cores suggest important contributions from *Sphagnum* spp, which is consistent with modern plant coverage on the two sites (Fig.2).

4.2.3 Changes in plant communities and site-specific conditions around Yuanchi Lake

The *n*-alkane-based reconstruction of peat deposition in the four settings reveals differences in the rates and composition of peat deposited around Yuanchi Lake (Figs. 5-8). In the YC_w site, inputs of vascular plant residues to the paleolake sediments beginning around 1600 CE initiated accumulations of TOC and elevated *n*-alkane concentrations (Fig.5 a and c). The plant communities that developed in this location before 1800 CE were probably dominated by abundant aquatic plants that are typified by short and middle chain lipids (Fig.5e and f). Terrigenous vascular plants characterized by TOC-rich and long-chain lipids began to dominate since 1800 CE, as indicated by the highest TOC concentrations, higher ACL and lower P_{aq} values from 1800 to 1900 CE (Fig.5 a, e and f). These features imply that the relative water depth at this site dropped quickly from 1600 CE as this lakeshore location filled in starting ca. 1800 CE, leading to greater expansion of subaerial TOC-rich vascular plants that promoted peat accumulation. The peat entered a rapid accumulation interval since 1940 CE as evident from the increases in the TOC concentrations and peaks in the PAR and CAR (Fig.3a

and b). Based on the *n*-alkane compositions of modern plants in this site, the lowest CPI and ACL values and the highest P_{aq} values (Fig.5 d-f) suggests a higher contribution of *Sphagnum* spp residues that are more resistant to decomposition and commonly accumulates in peat during shallow water conditions (Andersson et al., 2012; Loisel et al., 2014; Zhang et al., 2016, 2018).

At the YC_s site, the modern plants are characterized by emerged aquatic vascular plants (*T. orientalis* and *P. australis*) and other aquatic plants associated with relatively deeper water than at the other sites (Fig.2). Lower TOC concentrations, *n*-alkane CPI and ACL values and higher P_{aq} values, collectively imply a lacustrine condition with smaller subaerial plant inputs before 1850 CE (Fig.6 a, d-f). Higher *n*-alkane TOC-normalized concentrations in this time might have resulted from algae and bacteria that are characterized by short chain *n*-alkanes (Fig.6 c-f). Higher TOC, *n*-alkane concentrations, CPI and ACL values, and lower P_{aq} values indicate greater inputs of terrigenous vascular plants associated with continual lake infilling, promoted peat accumulation from 1850 to 1900 CE (Fig.6). An evident rise in water depth occurred from 1900 to 1950 CE when the aquatic plants became the major communities as indicated by the highest TOC-normalized *n*-alkane concentrations, lowest *n*-alkane ACL but the highest P_{aq} values (Fig.6 c, e and f). Obvious decreases in both TOC and TOC-normalized *n*-alkane concentrations (Fig.6 a and c) imply smaller inputs to the peats from terrigenous vascular plants (Eglinton and Hamilton, 1967). However, PAR reached a maximum and CAR also increased during this interval, although the terrigenous vascular plant contribution to the peat was less (Fig.6b). Peat preservation

and accumulation are mainly controlled by the combination of the kind of OM from the peat-forming plant inputs and microbial activity related to climate conditions (Zhou et al., 2005, 2010; Zhang et al., 2016). The variations in peat *n*-alkane CPIs are closely associated with peat-forming plants, temperature condition and microbial activity (e.g. Zhou et al., 2005; Andersson and Meyers, 2012; Naafs et al., 2019). We suggest that the weaker microbial activity that is indicated by relatively higher CPI values (Fig.6 d) in an inferred high water-level setting from 1900 to 1950 CE was a major factor for limiting peat decomposition. Since 1950 CE, the variations in TOC, PAR, TOC-normalized *n*-alkane concentrations and *n*-alkane proxies reflect local conditions, and the relative water level diminished to be like that in the period before 1900 CE (Fig.6).

Unlike YC_w and YC_s locations, the YC_E site was dominated by terrigenous vascular plants dominated by long-chain *n*-alkanes in low water depth conditions before 1800 CE (Fig.7). Major inputs to the peats from large amounts of terrigenous plants started at 1800 CE, when obvious increases in TOC concentrations appear in the YC_E location (Fig.7 a). Marked changes in *n*-alkane distributions of the peat core imply variations in vegetation communities in the YC_E locations since 1800 CE (Fig.7 c-f); Based on the *n*-alkane compositions of modern plants in this site, higher PARs and CARs from 1800 CE can be interpreted in terms of a greater contribution of *Sphagnum* spp. growing under wetter conditions that were the major contributors of middle chain carbon lipids, leading to higher *n*-alkane concentrations, elevated CPI and P_{aq} values, and the lowest ACL values in peat (Fig.7b-f).

Like the YC_E location, the peat in the YC_N site has clearly lower TOC

concentrations and accumulation rates during the initial time from 1750 to 1800 CE (Fig.8a and b), when fewer terrigenous vascular plants dominated by long-chain *n*-alkanes developed under drier *in situ* conditions (Fig.8d-f). Subsequently, the peat sequence received a high contribution of the middle-chain *n*-alkanes that are diagnostic of *Sphagnum* spp under inferred cold and wet conditions that stimulated peat TOC preservation and peat accumulation from 1800 to 1900 CE (e.g. Zhou et al., 2010; Andersson and Meyers, 2012; Naafs et al., 2019). After this period the *n*-alkane distributions are dominated by long chain carbon *n*-alkanes as evidenced from rapid decreases in CPI and P_{aq} values and an increase in ACL values (Fig.8 d-f), providing more evidence of inputs from predominantly terrigenous vascular plant such as *Carex* and shrubs during an interval of shallow water depths since 1950 CE.

Both the plant communities that contribute to the peat deposits and the degree of preservation can be the results of different hydrological conditions (Andersson et al., 2012). Compared to other three sites, the YC_N site evidently was deposited under a lower relative water level, given that the peat sequence has higher mean value of the *n*-alkane CPI (4.0 ± 0.9) and ACL (27.8 ± 1.0) but has lower P_{aq} values (0.3 ± 0.1) (Table 2), indicating that the peat originated mostly from terrigenous vascular plants. In contrast, the YC_w and YC_s peat cores have lower ACL (~ 25.4) but higher P_{aq} (~ 0.7) values (Table 2), implying the prevalence of aquatic plants typified by middle-chain *n*-alkanes under deeper water depth conditions at the two sites (e.g. Cranwell et al., 1984; Ficken et al., 2000; Andersson and Meyers, 2012; Naafs et al., 2019). The initial peat deposition in the YC_s site of the Yuanchi Lake was more than 200 years later than in the other three

sites (Fig.3). Therefore, we speculate that the Yuanchi Lake shallow basin that is composed of massive volcanic lava and tephra (Yang et al., 1995) is uneven and has a gradient with a higher north side and lower south and west sides. This topography results in different water depths around its edges and explains the contrast between the four peat sequences.

4.3. Controls on local climate and peat deposition around Yuanchi Lake

Because Yuanchi Lake is a closed system with no inflow or outflow (Bao et al., 2010), its hydrology is mainly controlled by evaporation and precipitation. Volcanic eruptions become important natural contributors to regional climate changes on many timescales, and large explosive volcanism is a leading forcing agent of natural climate changes on an interannual timescale (e.g. Robock, 2000; Liu et al., 2016). In general, air temperatures are reduced for several years after large or high frequency eruptions, because aerosols generated by volcanic eruptions could diminish solar radiation and thereby impact regional and sometimes global climates (Huang et al., 2013; Liu et al., 2015). *n*-Alkane-inferred climate reconstructions in the Great Hinggan Mountain, northeastern China, have revealed that cold climate intervals are associated with periods with a higher frequency of tropical Pacific volcanic eruptions (Mann et al., 2005; Zhang et al., 2014). In the Yuanchi peatland, the initial peat deposition occurred at the YC_w and YC_E locations around 1600 CE (Fig.3). Coincidentally, high frequency volcanic activities had occurred in the tropical Pacific in this period (Mann et al., 2005) and volcanic eruptions also occurred in the Changbai Mountains in 1668 and 1702 CE (Xu et al., 2012). We therefore postulate that lower OM accumulations accompanied with

higher ash contents in the sediments before 1700 CE in these two sites record tephra delivery and that cold climate conditions are consequences of the frequent volcanic activities in the Pacific and the local eruptions in the Changbai Mountains (Fig. 9 f and g).

The initial stage of transformation of lake edges into peatlands around 1800 CE corresponds to a time of warmer and drier climate in northeastern China (Hong et al., 2001; Lin et al., 2004; Zhang et al., 2014). Decreased volcanic activities during the interval from 1700 to 1800 CE led to warmer climate conditions that stimulated more terrigenous vascular plant growth and promoted peat accumulation along the shores of Yuanchi Lake (Fig.9). From 1800 to 1900 CE, cool climate condition in the Yuanchi region likely led to smaller OM inputs from terrigenous vascular plants as indicated by relatively lower TOC-normalized total *n*-alkane concentrations (except for the YCs sequence), lower CPI and ACL values, and higher P_{aq} values in the peat sequences (Fig.9 a, c-e). Moreover, the higher ash contents deposited in the cores is likely also a local response to an increase in volcanic forcing from the tropical Pacific (Fig.9f and g, Mann et al., 2005) and to volcanic eruptions in the Changbai Mountains in 1898 CE and 1903 CE (Huang et al., 2013). Subsequently, *n*-alkane proxies in the peat cores indicate that enhanced inputs of terrigenous vascular plant waxes under warm and dry climate conditions around Yuanchi Lake existed from 1900 to 1950 CE and promoted peat accumulation around the lake (Fig.9a-e). This multi-decadal warm/dry regional transition corresponds to the period characterized by diminished volcanic activities in both the Changbai Mountains and the tropical Pacific (Fig.9f).

After 1950 CE, total *n*-alkane concentrations in the four sequences drop to their minima, and obvious decreases in PAR values occurred in the high-water level locations (the YC_W and YC_S sites) (Fig.9a-b). These *n*-alkane proxies imply that the Yuanchi region experienced a significantly cooler climate accompanied with decreased terrigenous vascular plant inputs in recent years, as reflected in the higher CPI, lower ACL, and higher P_{aq} values in the peat sequences (Fig. 9c-e). This multidecadal period of cold climate might be attributed to a series of volcanic activities in the tropical Pacific (Fig.9f) that released abundant aerosols and thereby diminished solar radiation (Robock, 2000; Huang et al., 2013; Liu et al., 2016). In addition, the beginning of the Anthropocene in the Changbai Mountain region could be defined as 1950 CE due to the rapid industrial and urban development after the Reform and Opening-Up of China (Bao et al., 2019). As part of this development, greater combustion of fossil fuels and biomass from expanding anthropogenic sources led to increases in their soot emissions and more black carbon being deposited in peat (Gao et al., 2016). Substantial increases since 1950 CE in Pb accumulation rates in the Yuanchi peat further indicate significant deterioration of the mountain environment from magnifying human impacts (Fig.9g). Therefore, we postulate that anthropogenic impacts on the mountain region since 1950 CE likely suppressed growth of land plants, reduced OM inputs from subaerial vascular plants, and slowed peat accumulation.

5. Conclusion

n-Alkane biomarkers from the four peat cores around Yuanchi Lake in the Changbai Mountains, northeastern China, record peat depositional processes in a typical maar lake

on a decadal scale. Although different lengths of peat sequences were collected around Yuanchi Lake, the initiation for abundant peat accumulation (ca. 1800 CE) at the four locations was similar. Based on *n*-alkane compositions in modern plants at the four sites, *n*-alkane distributions in the four peat cores indicate that different vegetation communities under different hydrological conditions contributed to differences in peat accumulation rates and compositions of the four peat sequences. We postulate that the differences in peat deposition between the four sites is a consequence of different peat-forming communities growing in different water depths that are the results of the uneven basement rock topography of the lake basin caused by volcanogenic origin from lava and tephra.

n-Alkane-inferred regional climate changes have impacted peat accumulations, but in ways that are shared by the four locations. Lower peat accumulation rates record diminished OM inputs from terrigenous vascular plant during two long-term intervals from 1600 to 1700 CE and 1800 to 1900 CE, which both correspond to cooler climate associated with high-frequency volcanic eruptions in the tropical Pacific and in the Changbai Mountains. A lower rate of volcanic eruptions contributed to warm/dry regional conditions from 1700 to 1800 CE and 1900 to 1950 CE, when large terrigenous vascular plant input and promoted peat accumulation. Since 1950 CE, lower peat accumulation rates with diminished OM inputs from terrigenous vascular plants have existed during a significantly cooler climate in the Yuanchi region that coincides with both a series of volcanic eruptions and anthropogenic activities.

Data Availability Statement

Analytical data are available from Deep Blue Data () and from the cited references.

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Author Contributions

YZ, CG, SZ, PY, and GW conceived and designed the study. YZ, CG, SZ, and PY facilitated and conducted the field work. YZ did the lab analyses and analyzed the data. YZ and PAM wrote the paper. All authors contributed to the drafts of the paper and its final approval.

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Figure legends

Fig. 1. Location of the sampling site from the Yuanchi peatland in the Changbai Mountains, northeastern China.

Fig. 2. Panoramic views of Yuanchi Lake and of the four different sampling sites (west, south, east and north) in the peatlands surrounding the lake. (Photos by Guoping Wang)

Fig.3. Lithostratigraphy, ^{210}Pb -inferred age-depth model (a, c, e and g), the TOC concentrations (black lines, Bao et al., 2010) and peat accumulation rates (PAR, red lines) (b, d, f and h) of the four peat cores from the west, south, east, and north sides (YC_w, YC_s, YC_e, YC_n, respectively) around Yuanchi Lake.

Fig.4. Distributions relative to the major *n*-alkane in samples from different depths of the four peat cores around Yuanchi Lake.

Fig.5. Downcore variations of TOC concentrations (a), carbon accumulation rates (CAR) (b, black line) and PAR values (red line), Total *n*-alkane concentrations relative to TOC (c), *n*-alkane CPI (d), ACL (e) and P_{aq} (f) values from the western site of the Yuanchi peatland (YC_w). TOC and CAR values from Bao et al., 2010. The red dotted lines represent the intervals characterized by different variations in those proxies.

Fig.6. Downcore variations of TOC concentrations (a), carbon accumulation rates (CAR) (b, black line) and PAR values (red line), Total *n*-alkane concentrations relative to TOC (c), *n*-alkane CPI (d), ACL (e) and P_{aq} (f) values from the southern site of the Yuanchi peatland (YC_s). TOC and CAR values from Bao et al., 2010. The

red dotted lines represent the intervals characterized by different variations in those proxies.

Fig.7. Downcore variations of TOC concentrations (a), carbon accumulation rates (CAR) (b, black line) and PAR values (red line), Total *n*-alkane concentrations relative to TOC (c), *n*-alkane CPI (d), ACL (e) and P_{aq} (f) values from the eastern site of the Yuanchi peatland (YC_E). TOC and CAR values from Bao et al., 2010. The red dotted lines represent the intervals characterized by different variations in those proxies.

Fig.8. Downcore variations of TOC concentrations (a), carbon accumulation rates (CAR) (b, black line) and PAR values (red line), Total *n*-alkane concentrations relative to TOC (c), *n*-alkane CPI (d), ACL (e) and P_{aq} (f) values from the northern site of the Yuanchi peatland (YC_N) over the past 150 years. TOC and CAR values from Bao et al., 2010. The red dotted lines represent the intervals characterized by different variations in those proxies.

Fig.9. The colored dotted lines represent total *n*-alkane concentrations relative to TOC contents (a), PAR (b), *n*-alkane CPI (c), ACL (d), P_{aq} values (e) and ash contents (f, ASH%, Bao et al., 2010) of the four sites peat cores, respectively; Red dotted lines in (a-f) represent of those proxies in YC_W peat core; blue dotted lines represent of YC_S peat core; black dotted lines represent of YC_E peat core; green dotted lines represent of YC_N peat core. Volcanic forcing of the tropical Pacific from a model of the tropical Pacific coupled ocean-atmosphere system studied by Mann et al. (2005) (g); Red solid line in (g) represents Pb accumulation rate (Pb AR) in the peat core from Yuanchi

peatland studied by Bao et al., 2019. The gray shaded areas represent the intervals characterized by cold local conditions associated with high volcanic forcing of the tropical Pacific and anthropogenic activities in the Changbai Mountains region.

Table1.

Major modern plant species around Yuanchi Lake in Changbai Mountains with dominant *n*-alkane homologues in those plants.

Major modern species		Major homologue associated to plant
Trees	<i>Larix olgensis</i>	C ₂₇ , C ₂₉ ^a
	<i>Betula fruticosa</i>	C ₂₇ ^a
	<i>Vaccinium uliginosum</i>	C ₂₉ ^a
Shrubs	<i>Rhododendron parvifolium</i>	C ₂₉ , C ₃₁ ^{a,b}
	<i>Ledum palustre</i> L.	C ₂₉ , C ₃₁ ^{a,c}
	<i>Carex lasiocarpa</i>	C ₂₉ ^d
Sedges	<i>Scirpus triqueter</i>	C ₂₉ , C ₃₁ ^b
	<i>Eriophorum fauriei</i>	C ₂₇ ^a
	<i>Equisetum hiemale</i>	C ₂₉ ^e
Mosses	<i>Sphagnum</i> spp	C ₂₃ , C ₂₅ ^{d,f,g,h}
Emergent macrophytes	<i>Typha orientalis</i>	C ₂₉ ^{i,j}
	<i>Phragmites australis</i>	
Aquatic plants	Submerged and floating plants	C ₂₁ , C ₂₃ and C ₂₅ ^{i,j}

^a Tarasov et al. (2013); ^b Street et al. (2013); ^c Salasoo (1987); ^d Ronkainen et al. (2013); ^e Andersson et al. (2011); ^f Bass et al., 2000; ^g Nichols et al., 2006; ^h Bingham et al., 2010; ⁱ Cranwell et al. (1984); ^j Ficken et al. (2000);

Figure.

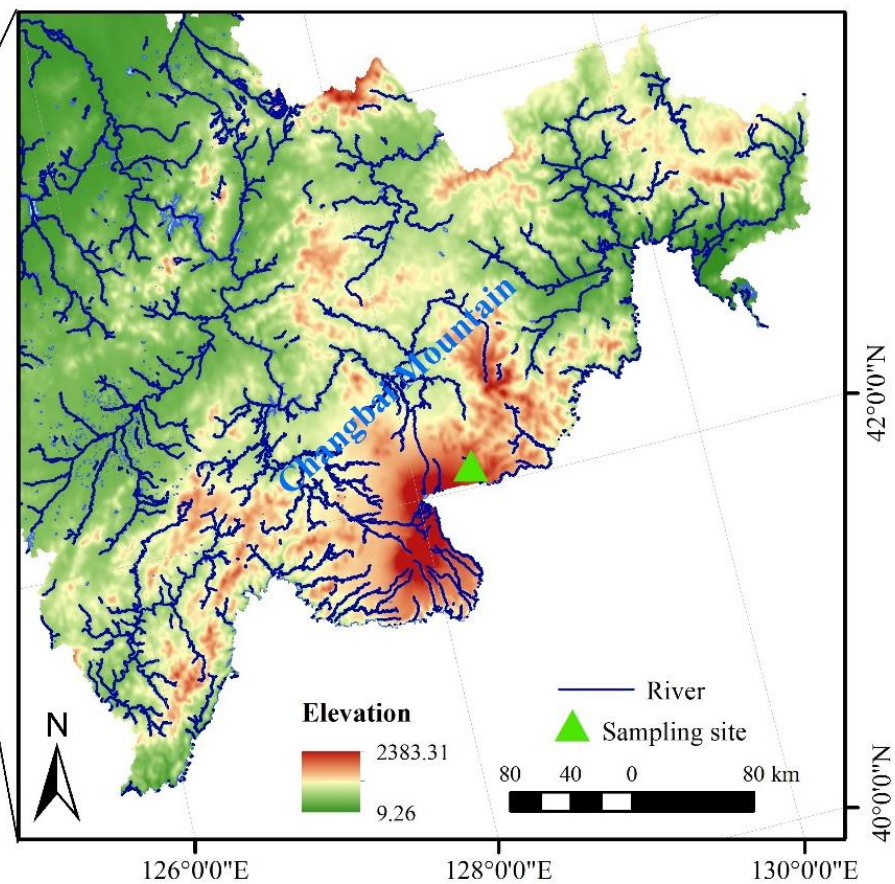
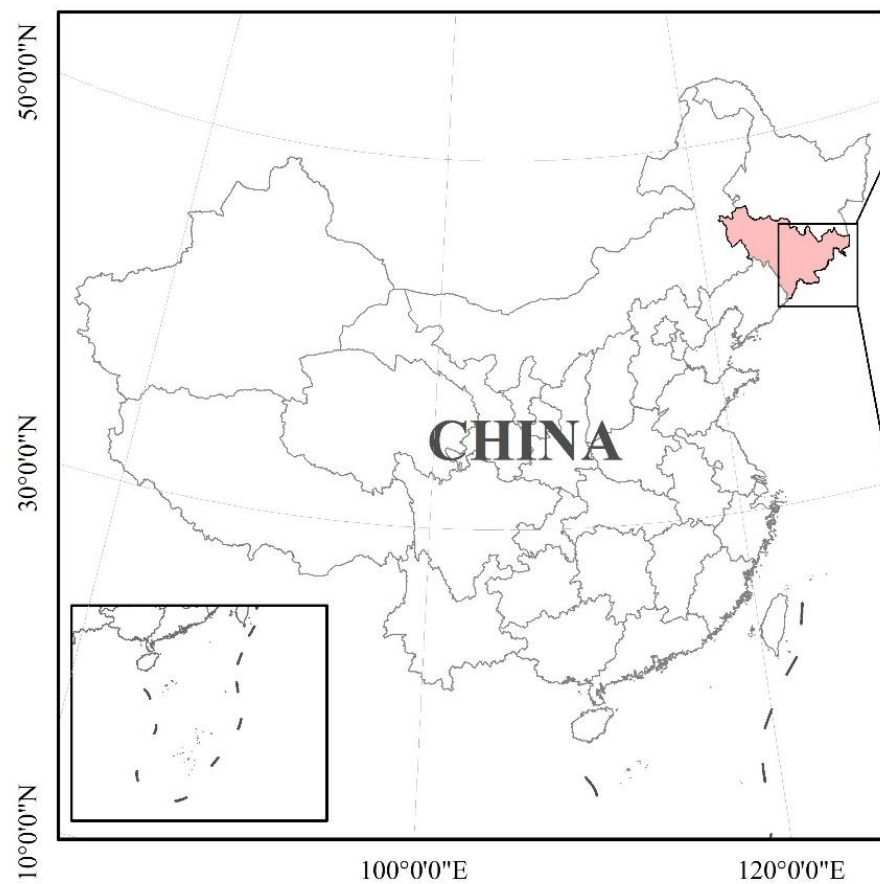


Figure.



Figure.

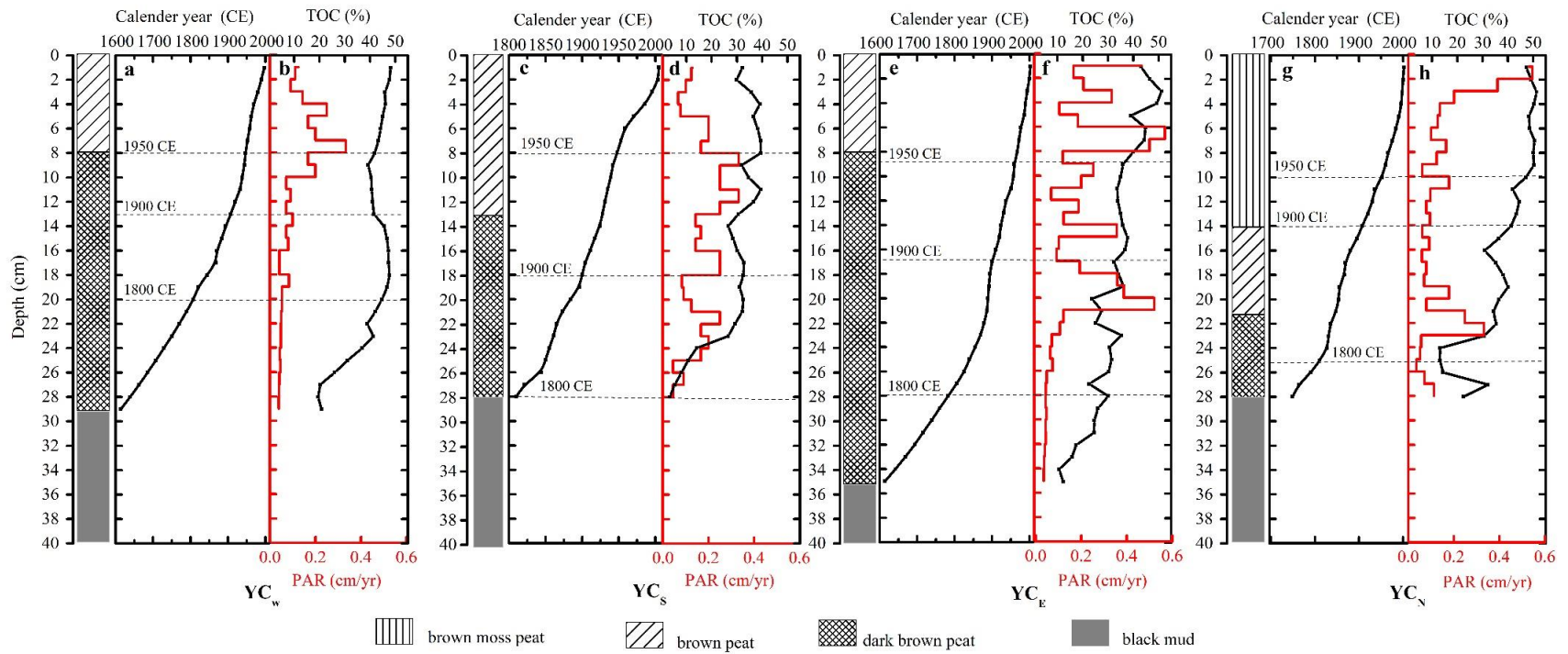


Figure.

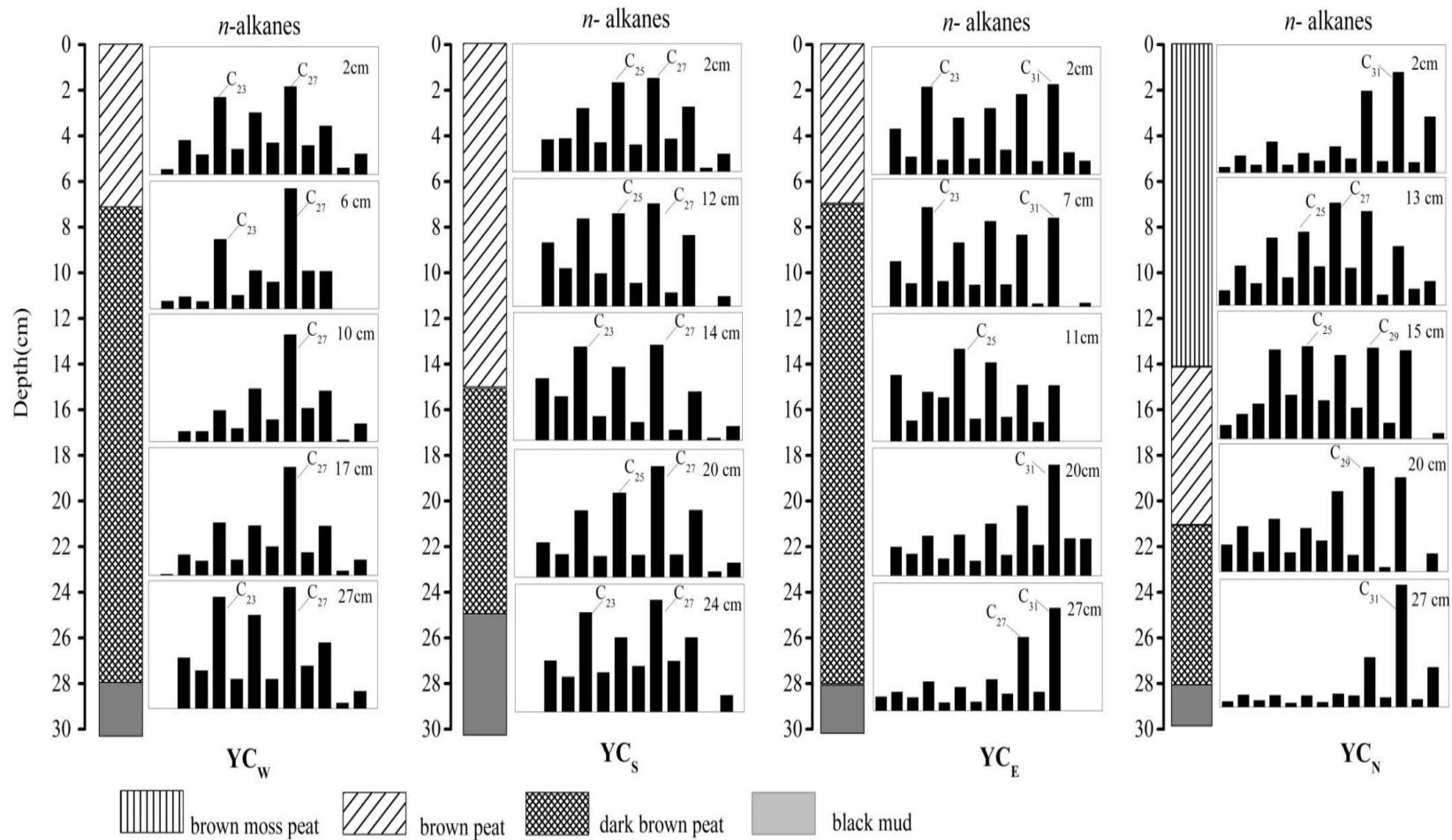


Figure.

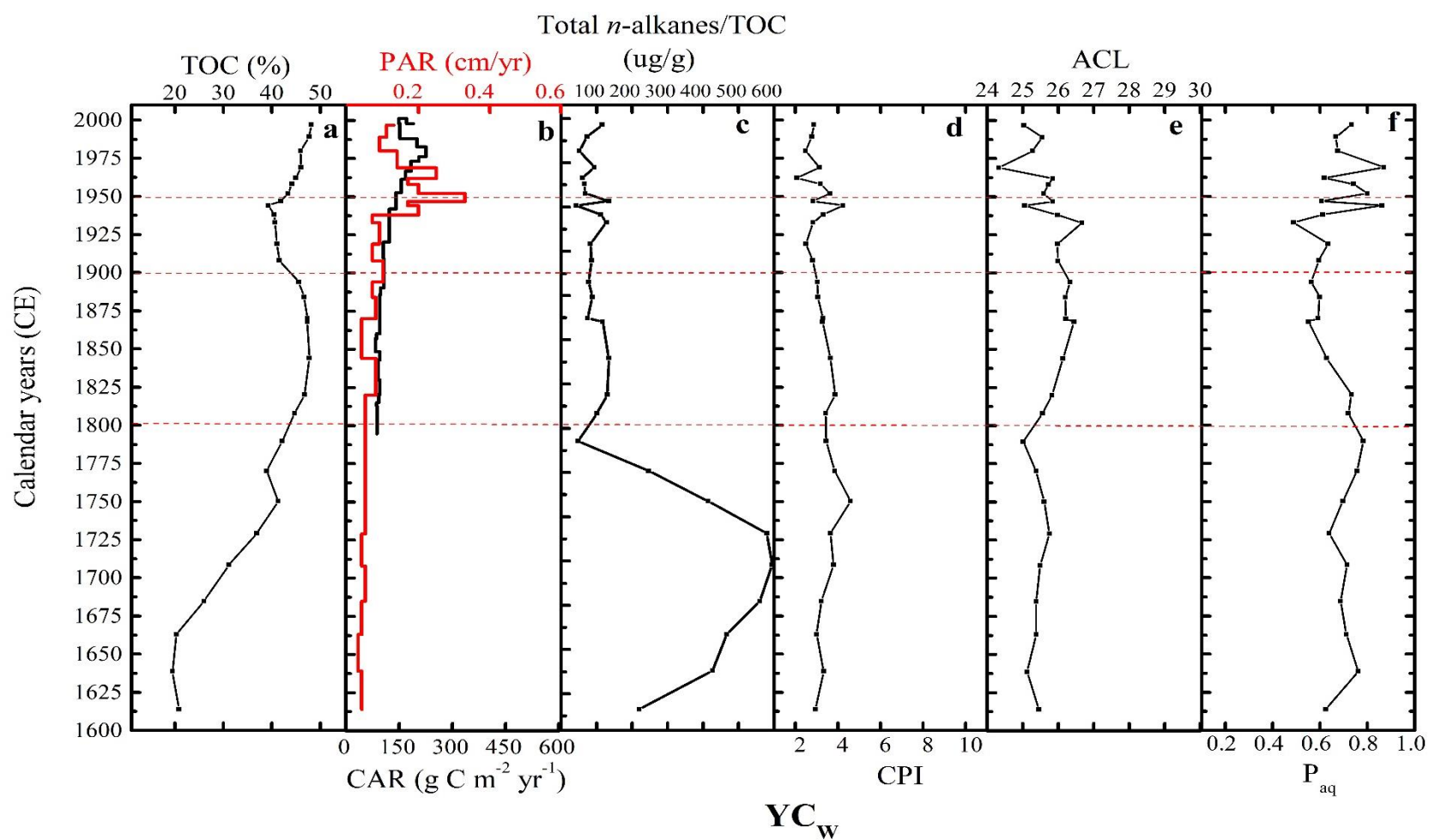


Figure.

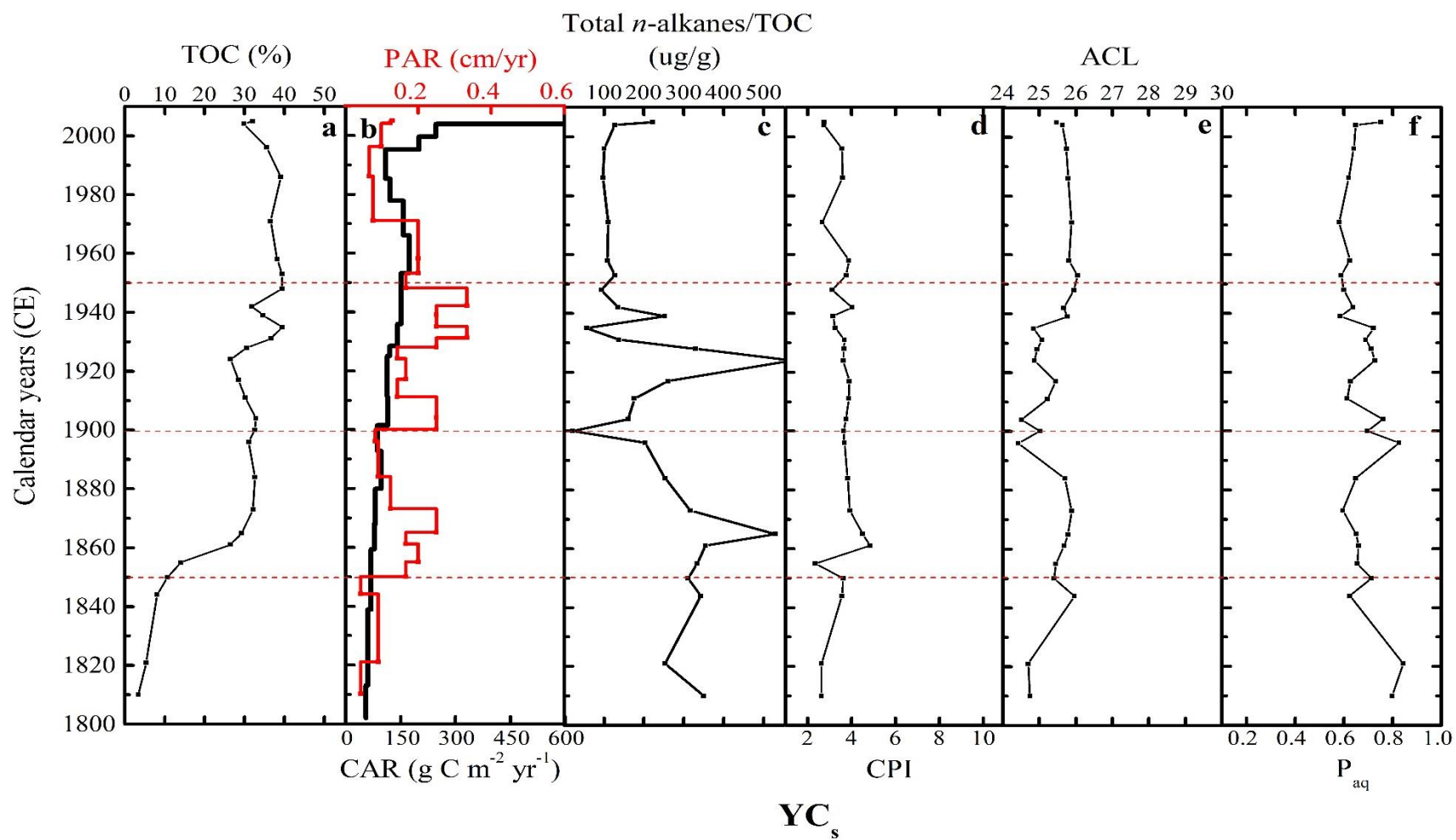


Figure.

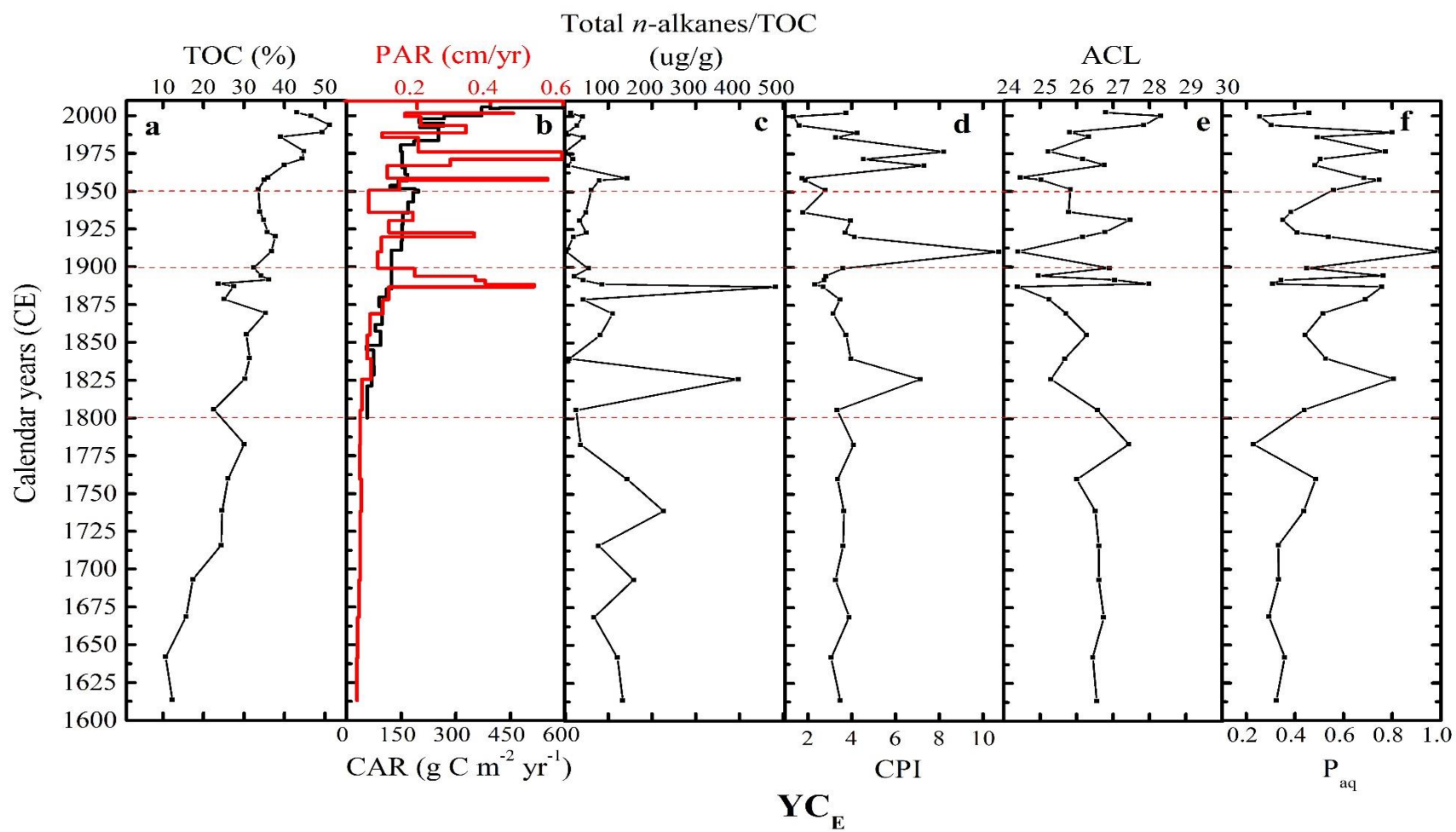


Figure.

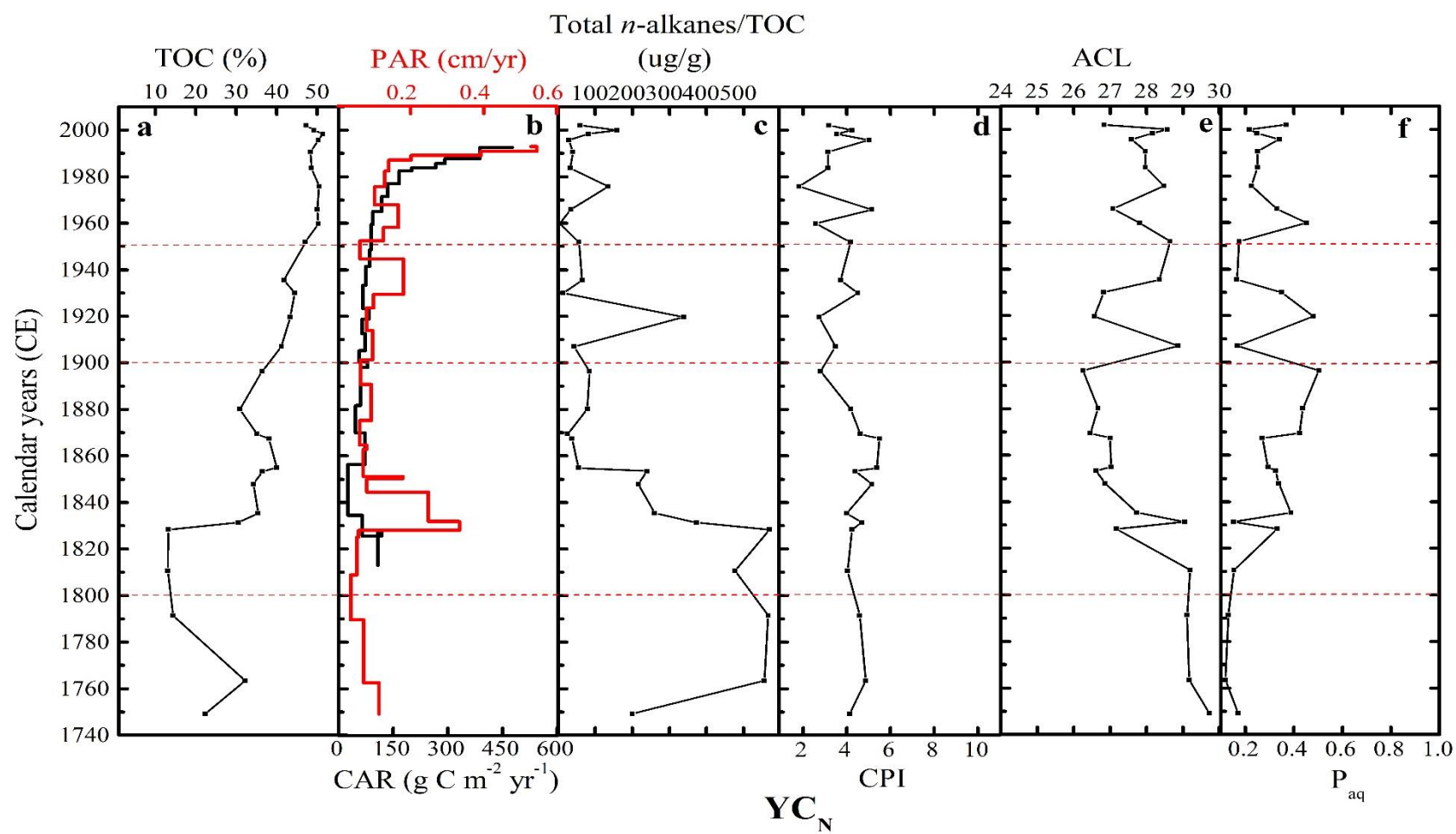


Figure.

