



Stable Oxygen Isotope Records ($\delta^{18}\text{O}$) of a High-Andean Cushion Peatland in NW Argentina (24°S) Imply South American Summer Monsoon Related Moisture Changes During the Late Holocene

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OPEN ACCESS

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Specialty section:

This article was submitted to
Quaternary Science, Geomorphology
and Paleoenvironment,
a section of the journal
Frontiers in Earth Science

Received: 21 December 2018

Accepted: 27 February 2019

Published: 22 March 2019

Citation:

Kock ST, Schitteck K, Wissel H,
Vos H, Ohlendorf C, Schäbitz F,
Lupo LC, Kulemeyer JJ and Lücke A
(2019) Stable Oxygen Isotope
Records ($\delta^{18}\text{O}$) of a High-Andean
Cushion Peatland in NW Argentina
(24°S) Imply South American
Summer Monsoon Related Moisture
Changes During the Late Holocene.
Front. Earth Sci. 7:45.
doi: 10.3389/feart.2019.00045

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High-elevation cushion peatlands are promising archives for paleoenvironmental studies in their extreme habitat of the Central Andean highlands between ~ 4000 and 4800 m a.s.l. The Cerro Tuzgle cushion peatland (CTP, $24^\circ 09' \text{S}$, $66^\circ 24' \text{W}$), located in the NW Argentine Andes, is formed by the vascular cushion plants *Oxychloe andina* (*O. andina*) and *Zameioscirpus muticus* (*Z. muticus*). To extend the knowledge base on the modern ecology of these peatlands, we investigated the stable isotope composition of bulk material and cellulose ($\delta^{18}\text{O}$, $\delta^{13}\text{C}$, $\delta^{15}\text{N}$) of the dominant cushion-forming species *O. andina* (Juncaceae) and *Z. muticus* (Cyperaceae) as well as water samples ($\delta^{18}\text{O}$, $\delta^2\text{H}$) of several pools interspersed within the peatland. We further applied a multiproxy approach for a peat core from CTP spanning the last 2900 years with XRF scanning, bulk geochemistry and stable isotope analyses on bulk peat and cellulose size fractions. Modern samples of *O. andina* and *Z. muticus* expose significant differences in cellulose $\delta^{18}\text{O}$, e.g., between leaves and rhizomes of *O. andina* ($\Delta\delta^{18}\text{O}_{\text{L-R}} = 4.11\text{‰}$) and between leaves of *O. andina* and *Z. muticus* ($\Delta\delta^{18}\text{O}_{\text{L-L}} = 2.8\text{‰}$). Modern water samples exhibit strong isotopic differences between single water pools (max. $\Delta\delta^{18}\text{O} = 13.09\text{‰}$) due to local variable evaporative enrichment. Within the peat core, we observe considerable multi-centennial variations in $\delta^{18}\text{O}$ composition of cellulose confirmed by all size fractions. Based on the regional relation between decreasing $\delta^{18}\text{O}_{\text{prec}}$ values with increasing precipitation amounts and ^{18}O enrichment in the peatland waters due to evaporation, we suggest an interpretation of our $\delta^{18}\text{O}$ cellulose record as moisture proxy for CTP. This interpretation is corroborated by a high correlation between

oxygen isotopes, peat growth and geochemical data. Accordingly, CTP indicates dryer conditions between 2190 and 2120, 1750 and 1590, 1200 and 1080 and since 130 cal. yr BP, whereas periods with increased humidity prevailed from 2750 to 2250 and from 600 to 130 cal. yr BP. Temporal changes in the match to South American Summer Monsoon (SASM) reconstructions suggest impacts of other large-scale atmospheric variability modes or a different SASM expression at our southerly location.

Keywords: late Holocene, cushion peatlands, Central Andes, SASM, precipitation, cellulose, stable isotopes

INTRODUCTION

The Andes represent a prominent longitudinal climatic barrier for the South American continent. Two major atmospheric circulation systems influence the climate of the Andes, the South American summer monsoon (SASM) and the Southern Hemisphere westerly winds (SHW) (Garreaud et al., 2003). Interactions between these systems affect climate conditions of the entire Southern Hemisphere. Hence, interest in paleoclimate archives from the Central Andes has grown during the recent decades (e.g., Flantua et al., 2016). Between these systems, the Arid Diagonal is developed, characterized by low annual precipitation amounts and a limited number of paleoclimate archives. Due to this lack of continuous and high resolution records, knowledge on the climatic evolution of the Andean region during the Holocene is still limited and insufficient compared to the hemispheric and global significance of this region.

Suitable archives to fill the gap of knowledge are high-Andean cushion peatlands, as several studies could provide in the past (Schitteck et al., 2016; Kock et al., 2018, 2019). Continuous, high accumulation rates ensure highly resolved records for paleoenvironmental investigations with a multiplicity of proxy data (Schitteck, 2014). While stable isotope analyses are a well-established tool in paleoecological investigations of Northern Hemisphere peatlands (e.g., Ménot-Combes et al., 2002; Moschen et al., 2009), isotope analyses on cushion peatlands reside in an initial stage. Different to the Northern Hemisphere dominant peat-forming species *Sphagnum*, a non-vascular plant, the cushion-forming species are vascular plants. Vascular plants are able to regulate their gas exchange by stomata, affecting the isotope fractionation processes of both oxygen ($\delta^{18}\text{O}$) and carbon isotopes ($\delta^{13}\text{C}$) during metabolic processes (Ménot and Burns, 2001).

The stable isotope values of peatland plants underlie a strong dependency on the respective source values of atmospheric CO_2 and plant available H_2O . Overall, carbon isotope fractionation is dependent on the atmospheric source value, on the difference between intercellular and atmospheric partial pressure of CO_2 , which is controlled by the stomatal conductance in vascular plants, and on the biochemical fractionation (Farquhar et al., 1989). During more humid phases, an increasing stomatal conductance is leading to decreased $\delta^{13}\text{C}$ values in plant organic matter. During more arid phases, an increased leaf to air vapor pressure gradient is leading to a closure of the stomata to prevent passive water loss as a result of transpiration

(Amesbury et al., 2015a). One of the first responses to water deficit in vascular plants is the inhibition of growth, which allows plants to reduce their transpiration rates and to avoid metabolic and cellular damage caused by low water potentials (Cominelli et al., 2008). Besides carbon isotopes, oxygen isotopes are very often applied within paleoecological investigations (e.g., Ménot-Combes et al., 2002). The $\delta^{18}\text{O}$ value of plant cellulose is directly related to the respective leaf water ($\delta^{18}\text{O}_{\text{LW}}$) and enriched by about 27.0‰ due to biochemical fractionation during formation of primary photosynthates (glucose) (DeNiro and Epstein, 1981; Sternberg, 2009). The aperture of plant stomata has a strong influence on $\delta^{18}\text{O}_{\text{LW}}$. Stomatal closure, caused, e.g., by drought, lead to a decrease of stomatal conductance and increased enrichment of $\delta^{18}\text{O}_{\text{LW}}$ values during transpiration (Farquhar et al., 2007). Due to the dependence on the respective water source value ($\delta^{18}\text{O}_{\text{SW}}$) and the leaf water enrichment, reported $\delta^{18}\text{O}$ values for extracted cellulose of vascular plants in peatlands cover a considerable range (Brenninkmeijer et al., 1982; Hong et al., 2009; Amesbury et al., 2015b).

Stable isotope investigations on cushion peatlands in South America prior to this study have been executed in Peru (*Distichia* peatlands; Skrzypek et al., 2011; Engel et al., 2014) and Bolivia (Gouze et al., 1987). Further studies on vascular plants in peatlands only have been conducted in Switzerland (Ménot and Burns, 2001), China (e.g., Hong et al., 2000, 2009; Liu et al., 2018) and New Zealand (Amesbury et al., 2015a,b). For carbon isotopes, a relationship between $\delta^{13}\text{C}$ and temperature (Ménot and Burns, 2001; Skrzypek et al., 2011; Engel et al., 2014; Amesbury et al., 2015a) was described. Ménot and Burns (2001) also reported that on an altitudinal gradient the atmospheric CO_2 concentration is a stronger determinant for $\delta^{13}\text{C}$ values of peat plants than temperature. Hong et al. (2000, 2009) interpret $\delta^{18}\text{O}$ as a proxy for surface air temperature changes, while Amesbury et al. (2015b) used $\delta^{18}\text{O}$ for tracking precipitation moisture sources.

Here we investigated modern water and plant (leaves, rhizomes, fruits) samples of the Cerro Tuzgle cushion peatland (CTP) as well as a 2.6 m composite peat profile. In the peat profile, we analyzed the major and trace elements, ash content, humification tendency, organic carbon and total nitrogen isotope composition of bulk peat and carbon and oxygen isotope composition of cellulose. Prior to cellulose extraction, samples were separated into three different size fractions (<200 μm , 200–1000 μm , >1000 μm). Additional macrofossil analyses were used to identify changes between different plant species within the record. As a general hypothesis for the study we postulate that changes in moisture conditions at CTP are mirrored in

$\delta^{18}\text{O}_{\text{cell}}$ values of the peat profile and that disturbances in such a relation, e.g., because of decomposition, can be detected within the multi-proxy approach.

Overall, the study strives (1) to set-up a modern dataset as base for interpretation of the isotope records, (2) to establish the first continuous cellulose stable carbon and oxygen isotope record of cushion peatlands from the Central Andes, (3) to discuss moisture fluctuations at CTP in the context of available SASM reconstructions.

STUDY AREA

NW Argentine, High-Andean “Puna”

The CTP (24°09' S, 66°24' W, 4,350 m a.s.l., **Figure 1a**) is located in the northwest Argentine Andes close to the Cerro Tuzgle (24°09' S, 66°24' W) in the province of Jujuy. The area is characterized by recent tectonic processes and geothermal activity, including several seasonal and permanent springs in the so called Cerro Tuzgle – Tocomar geothermal volcanic area (Giordano et al., 2013). The occurrence of block glaciers or high-elevation cryogenic soils is not reported for the study area. The region is part of the internally drained Puna comprising the southern part of the Altiplano-Puna plateau (Allmendinger et al., 1997; Norini et al., 2014). Due to differences in precipitation amount, the Argentine Puna can be subdivided into a semi-humid part in the east, where the CTP is situated and an arid area in the west (Cabrera, 1968; Troll, 1968; Kuentz, 2009).

Its location is in the north to the Arid Diagonal at the crossroads between tropical and extratropical precipitation regimes (Vuille and Ammann, 1997; Garreaud et al., 2009). The weather station at the Concordia mine (4,144 m a.s.l.) shows the main climatic characteristics of the study area (**Figure 1b**).

Annual temperatures range between 2 and 10°C (mean annual temp.: 6.9°C) with temperature maxima prevailing in the austral summer months. Annual precipitation sums are 109 mm on average while more than 90% occur between November and March, indicating the main source for meteoric water reaching the CTP is the SASM. Cold front or cuts off events that lead to precipitation in the austral winter months are rare (**Figure 1b**).

Cushion Peatlands as Environmental Archives

In the recent decade, high-elevation cushion peatlands turned out to be high-potential archive for paleoenvironmental investigations (Squeo et al., 2006; Engel et al., 2014; Schitteck et al., 2015). These sensitive ecosystems are comparably less vulnerable to the availability of water due to their special textural adaptations like cushion-growth that increase the resistance to water fluctuations (Schitteck, 2014). Thus, they have the potential for continuous records from the harsh environment in the (semi-) arid region of the Central Andes. High-Andean cushion peatlands can accumulate several meters of thickness, which are composed of Juncaceae like *Oxychloe*, *Distichia* and *Patosia* and are frequently supplemented by Cyperaceae of the genera *Zameioscirus* and *Phylloscirus* (Ruthsatz, 2008). Furthermore, cushion peatlands are very sensitive to climatic changes due to their occurrence close to the limits of vegetation growth and contingent upon temperature, water economy and growing period (Ruthsatz, 1977).

A number of high-elevation peatlands occur in the study region (Werner, 1974). In contrast to most of the cushion peatlands in the area, CTP is situated between comparatively flat slopes of outcropping Ordovician volcanic rocks and sedimentary units as well as Cretaceous conglomerates and sands. Thus, it is protected against increased allochthonous sediment input

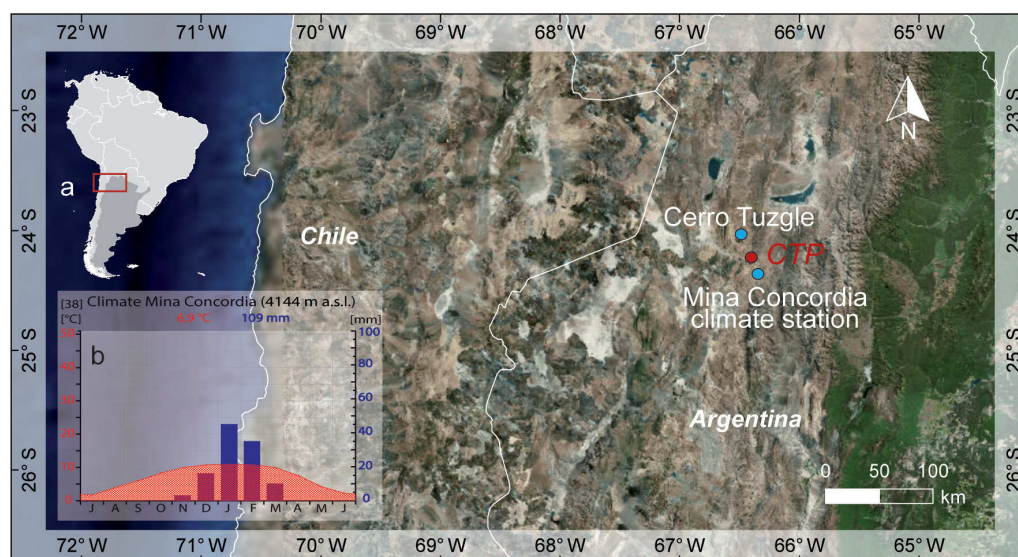


FIGURE 1 | (a) Location of the CTP study area, Cerro Tuzgle and Mina Concordia climate. **(b)** Climate diagram Mina Concordia [data source: Mina Concordia (1950 – 1990; 1966 and 1970 are missing) data based on Bianchi et al., 2005]. Map data source is GLCF World Data and DGM-GTOPO30.

from lateral stream channels or valleys during extreme rainfall events (Schitteck et al., 2016). A presently inactive alluvial fan at the southernmost section of the peatland developed a natural dam and led to continuous accumulation of peat in the past. Modern vegetation of the investigated section of the peatland is characterized by stable mats formed by *Oxychloe andina* (*O. andina*), partially supplemented by *Zameioscirpus muticus* (*Z. muticus*). *O. andina* is forming dense cushions by keep growing at their tops and form rhizomes at the bottom (Figure 2b). The leaves are lignified and have a prickly at their tops, potentially as protection against grazing of hoofed animals (Figure 2c) (Rauh, 1988; Ruthsatz, 2008; Schitteck et al., 2016). *Z. muticus* grows in small gaps and fissures of the *O. andina* cushions. Furthermore, Cyperaceous species like *Z. muticus* can overgrow open peat surfaces that were damaged by mechanical, frost or drought harms, and thus can protect the peatland against erosion (Ruthsatz, 2008). In contrast to *O. andina*, *Z. muticus* has a higher physiological and ecological tolerance, showing, e.g., greater resistance against frost or droughts. However, the species cannot compete with the growth of the cushion plant *Oxychloe*. The structure of their rhizomes is too insecure, the leaves are

neither lignified nor possess a prickly and the growth does not always continue at their tops (Ruthsatz, 2008).

As the CTP is situated in a valley of an ephemeral stream, the Juncaceous cushion plants are interspersed by shallow pools, which can be connected by superficial water flows, originating from springs and during rainfall events from run-off from the surrounding slopes, but also occur isolated without superficial connection (Figure 2a). Underground flow can additionally lead to hydrologic connectivity within cushion peatlands (Schitteck et al., 2012). However, for CTP, consisting of homogeneous peat with interspersed minerogenic matter of smaller grain sizes, the water flow velocity is likely to be low.

MATERIALS AND METHODS

Modern Water and Plant Samples

In October 2013, two crossing transects (from northwest to southeast and from southeast to northwest) were set to enable a systematic sampling of the peatlands surface waters. Furthermore, various modern plant samples of different species (*O. andina* and *Z. muticus*) were taken in a close radius around the coring location in 2012. In November 2014, further water samples were taken from selected positions within the peatlands surface. The peat water sample was extruded by hand from a respective peat sample close to the surface of the intact cushion.

Modern plant samples were separated into leaves, rhizomes, and fruits of *O. andina* and *Z. muticus* in the field. In the laboratory, the separated plant parts were washed several times with deionized water to remove all visible sedimentary remains. After freeze-drying, the plants were milled with a high-speed mill grinder (Retsch, MM 400) to generate homogeneous samples. Since dispersion was not necessary for these samples, cellulose extraction after Wissel et al. (2008) was conducted without a treatment of NaOH.

For $\delta^{18}\text{O}_{\text{cell}}$ analyses, about 275 μg of freeze-dried cellulose was weighed in silver capsules, crimped and stored for at least 24 h in a vacuum drier at 100°C before analyses. Samples were then pyrolysed at 1450°C by using a high temperature pyrolysis analyzer (HT-O, HEKAtech) and measured on-line with a coupled IRMS (Isoprime, GV Instruments) to determine the oxygen stable isotope ratio. Peak integration of m/z 28, 29, and 30 was used to determine the oxygen content (O). The $\delta^{13}\text{C}$ -values were determined using 200–300 μg of dried cellulose weighed into tin foil cups. Samples were combusted at 1080° using an element analyzer (EuroEA, Eurovector) interfaced on-line to an isotope IRMS (Isoprime, GV Instruments). Peak integration of m/z 44, 45 and 46 was used to determine the carbon content (C). The element contents of C and O respectively the C/O ratio can be used as a quality indicator for the extracted cellulose in comparison to the theoretical C/O ratio in cellulose of 0.90 (Wissel et al., 2008).

The overall precision of replicate analyses is estimated to be better than 5% (rel.) for carbon and oxygen content, $<0.1\text{‰}$ for $\delta^{15}\text{N}$ and $\delta^{13}\text{C}$ and $<0.25\text{‰}$ for $\delta^{18}\text{O}$.

Stable isotope analyses of the water samples were carried out using a Cavity Ringdown Spectrometer (L2130-I, Picarro).

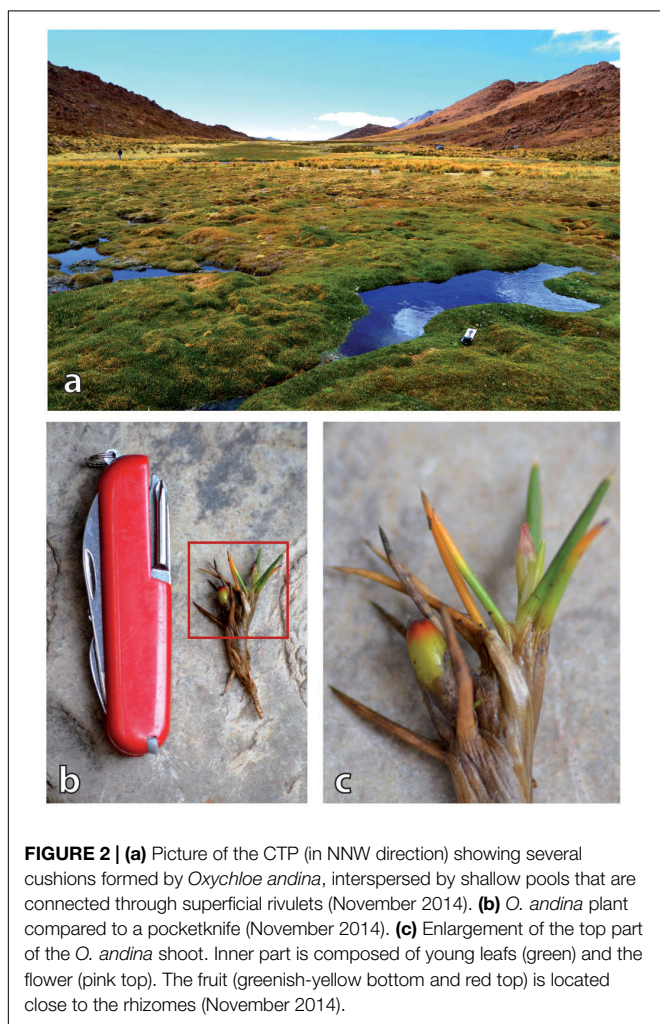


FIGURE 2 | (a) Picture of the CTP (in NNW direction) showing several cushions formed by *Oxychloe andina*, interspersed by shallow pools that are connected through superficial rivulets (November 2014). (b) *O. andina* plant compared to a pocketknife (November 2014). (c) Enlargement of the top part of the *O. andina* shoot. Inner part is composed of young leaves (green) and the flower (pink top). The fruit (greenish-yellow bottom and red top) is located close to the rhizomes (November 2014).

Results are reported as δ values relative to the VSMOW (Gonfiantini, 1978). Internal standards, calibrated against VSMOW, Standard Light Antarctic Precipitation, and Greenland Ice Sheet Precipitation were used to ensure long-term stability of analyses. The precision of the analytical system was $\leq 0.1\text{‰}$ for $\delta^{18}\text{O}$ and $\leq 1.0\text{‰}$ for $\delta^2\text{H}$.

All isotope results are presented as δ -values (‰) according to the equation:

$$\delta = (R_s/R_{st} - 1) \cdot 1000$$

where R_s is the isotope ratio of the sample ($^2\text{H}/^1\text{H}$, $^{13}\text{C}/^{12}\text{C}$, $^{15}\text{N}/^{14}\text{N}$, $^{18}\text{O}/^{16}\text{O}$) and R_{st} is the isotope ratio of the respective standard. Calibrated laboratory standards were used to control the quality of the analyses and to relate the raw values to the isotopic reference scales (VSMOW for hydrogen and oxygen, VPDB for carbon, AIR for nitrogen). Standards, including the International Atomic Energy Agency (IAEA) standard IAEA-CH6 cellulose powder ($\delta^{18}\text{O} = 37.09 \pm 0.09\text{‰}$), two further commercially available cellulose powders, Merck ($\delta^{18}\text{O} = 29.97 \pm 0.08\text{‰}$) and Fluka cellulose ($\delta^{18}\text{O} = 28.84 \pm 0.12\text{‰}$), and two in-house standards, Rice ($\delta^{18}\text{O} = 23.64 \pm 0.15\text{‰}$) and Peanut cellulose ($\delta^{18}\text{O} = 23.93\text{‰} \pm 0.11\text{‰}$), were calibrated against the reference standards IAEA-601 ($\delta^{18}\text{O} = 23.14\text{‰}$) and IAEA-602 ($\delta^{18}\text{O} = 71.28\text{‰}$) for oxygen isotopes. The standards USGS24 ($\delta^{13}\text{C} = -10.45 \pm 0.03\text{‰}$), IAEA-CH6 ($\delta^{13}\text{C} = -16.05 \pm 0.04\text{‰}$) and IAEA-CH7 ($\delta^{13}\text{C} = -32.15 \pm 0.05\text{‰}$) were used for calibration of carbon isotopes values. The calibration of nitrogen isotopes values was ensured by the use of the standards IAEA-N-1 ($\delta^{15}\text{N} = 0.4 \pm 0.2\text{‰}$), IAEA-N-2 ($\delta^{15}\text{N} = 20.3 \pm 0.2\text{‰}$), USGS25 ($\delta^{15}\text{N} = -30.4 \pm 0.4\text{‰}$), and USGS26 ($\delta^{15}\text{N} = 53.7 \pm 0.4\text{‰}$).

Core Sampling and Geochemical Measurements

In December 2012, the core Tuz-694 which reached a total depth of 8 m was extracted in 1 m segments by percussion coring. In this study, we present investigations on the upper 3 m of the core Tuz-694. The core was opened lengthwise with a modified core saw, cut into two halves, photographed and described lithologically.

For major and trace elements analyses, the first core half was measured with a resolution of 2 mm using an ITRAX X-ray fluorescence (XRF) core scanner (Cox Analytical Systems; Croudace et al., 2006). A molybdenum (Mo) tube at 30 kV and 10 mA was used with an exposure time of 10 s for every measurement respectively. The same core half was cut in one centimeter slices, from which boundary areas were removed to exclude possible contamination, which were used for cellulose extraction. The second half was accordingly prepared in 1 cm slices for stable isotope ($\delta^{13}\text{C}_{\text{OM}}$, $\delta^{15}\text{N}_{\text{bulk}}$) and element content analyses as well as loss on ignition (LOI_{550}) and humification (T_{535}) measurements on bulk material. From these samples, subsamples at 8 cm intervals were selected for macrofossil analyses.

Prior to the cellulose extraction after Wissel et al. (2008), bulk peat samples were treated with NaOH (5%) to ensure dispersion and enable separation into three size fractions by sieving at 200

and 1000 μm (established fractions: $<200 \mu\text{m}$, $200\text{--}1000 \mu\text{m}$, $>1000 \mu\text{m}$). Oxygen and carbon isotope measurements of the extracted cellulose followed methods described above.

For stable isotope and element content analyses as well as for LOI_{550} and T_{535} , bulk peat samples were freeze-dried and milled (Retsch, MM 400). For the analyses of the stable isotope ratios of carbon ($\delta^{13}\text{C}_{\text{OM}}$) and total carbon content (TC) as well as total nitrogen isotopes ($\delta^{15}\text{N}_{\text{bulk}}$) and nitrogen content (TN) one measurement was conducted using tin capsules and combusted at 1080°C in an elemental analyzer (EuroEA, Eurovector) with an automated sample supply linked to an isotope ratio mass spectrometer (Isoprime, Micromass). TN and TC contents were calculated according to the amounts of N_2 and CO_2 released after sample combustion (peak integration) and calibrated against element standards. Various tests with 5% HCl proved that decalcification was not necessary for the determination of organic carbon content and for $\delta^{13}\text{C}_{\text{OM}}$ measurements, thus, the TC content is an equivalent of the total organic carbon content (TOC). Loss on ignition (LOI_{550}) determination followed standard procedures with an initial milled material weight of 25–30 mg (see Heiri et al., 2001). Accordingly, the ash content was calculated by subtracting the LOI_{550} from 100%. Humification measurements followed the procedures recommended by Blackford and Chambers (1993) and Chambers et al. (2011), standardized to a target value of 20 mg organic matter in the sample.

Composite Profile, Radiocarbon Dating and Data Analyses

The upper sections of the second and third Tuz-694 core segment were determined as caving material because of loose bedding and deviations in XRF results and were thus discarded. A small hiatus occurs between 98 and 100 cm due to loss of material owing to the coring procedure. Moreover the retrieved core segments were compacted unequally during the coring due to the high water content. Within the first 1 m section, the peat material was compacted to 80 cm (20% compaction), the peat material of the second segment to 49 cm (51% compaction) and the third segment was compacted to 67 cm (33% compaction). To build a composite profile, the core segments were linearly decompacted by using a correction factor (core length/compacted length). The developed composite profile is the basis for the age-depth model. All depths mentioned in the text are decompacted composite core depths except for the depth given in the stratigraphical description below.

Nine samples were selected from characteristic stratigraphical layers for radiocarbon dating on bulk peat samples (0.5–1.0 cm^3) by Accelerator Mass Spectrometry at Poznań Radiocarbon Laboratory (Poznań, Poland) (Table 1). Radiocarbon ages were calibrated using CALIB 7.0.4 (Stuiver et al., 2013) and the SHCal13 data set for Southern Hemisphere calibration (Hogg et al., 2013).

In an earlier contribution (Schitteck et al., 2016), the calibration of radiocarbon ages for the Tuzgle composite record was based on the IntCal13 calibration data set (Reimer et al., 2013). Meanwhile, Marsh et al. (2018) proposed a mixed-curve model

TABLE 1 | Radiocarbon ages of core Tuz-694 from the CTP.

Lab-ID	Profil depth (cm)	Measured ^{14}C -age	Measured error ($\pm a$)	2 σ calibrated age (cal. yr BP)	MC age depth age (cal. yr BP)
Poz-56032	37–38	600	35	603 (541–654)	602 (541–654)
Poz-56034	77–78	1095	30	1001 (938–1060)	1000 (941–1061)
Poz-56035	103–104	1245	25	1211 (1082–1268)	1210 (1084–1264)
Poz-66440	144–145	1620	30	1514 (1412–1591)	1514 (1419–1588)
Poz-56036	160–161	1715	30	1620 (1557–1700)	1619 (1558–1700)
Poz-66442	187–188	1960	30	1911 (1830–1988)	1910 (1836–1984)
Poz-56037	234–235	2475	35	2576 (2380–2720)	2581 (2392–2710)
Poz-56038	246–247	2705	30	2804 (2757–2855)	2803 (2760–2860)
Poz-56039	258–259	2820	30	2921 (2849–3002)	2920 (2855–3009)

Calibrated ages (2 σ) were calculated using CALIB 7.0.4 (Stuiver et al., 2013) and the SHCal13 data set (Hogg et al., 2013). Given are the radiocarbon ages and their measurement error, the mean calibrated ages (2 σ error in brackets) and the modeled median ages (2 σ error in brackets).

between IntCal13 and SHCal13 for the regions affected by the SASM and receiving unknown amounts of carbon dioxide input from both hemispheres. Although atmospheric CO₂ from the Northern Hemisphere is brought to the eastern Andes during the austral summer, its potential influence declines with increasing southern latitude. At the latitude of CTP, the impact of northern hemispheric CO₂ should be negligible. Accordingly, radiocarbon ages for the Tuzgle composite record were recalibrated here with the SHCal data set, the age-depth model was recalculated (Tuz.v2) and geochemical proxies from Schitteck et al. (2016) are also shown on this updated age-depth scale. The modeled ages for the decompacted core depths are the medians of 800 Monte Carlo simulations also used to generate confidence intervals integrating the probabilistic nature of calibrated ^{14}C -dates (Higuera et al., 2009; MCAgeDepth software package). All discussed ages are calibrated ages BP (cal. yr BP), if not mentioned otherwise. Pearson correlation coefficients ($p > 0.05$) were determined by using the R software (version 3.2.2) and the package “rioja” (version 0.9-15; Juggins, 2017) to describe parameter relations.

RESULTS

Stratigraphy and Chronology

The stratigraphy of core Tuz-694 is dominated by homogeneous, faintly layered peat, partially interspersed with clastic material (Figure 3). Between 14 and 8 cm depth, sand is supplementing the highly organic peat. A short hiatus, occurring between 100 and 98 cm, is caused by core loss due to the coring procedure. At 134 cm, a fine gravel dominated layer occurs. Between 245 and 230 cm, an increased content of silt is observed.

The chronology is based on nine AMS radiocarbon dates and the drilling date as anchor point for the core top (Table 1). The age-depth model (Figure 3) shows a continuous peat accumulation of the CTP for the last ~2900 cal. years. The average time-resolution of our CTP record is about 11 years per centimeter with a range between 7 to 18 years.

Modern Data

The $\delta^{18}\text{O}$ values of modern water scatter between -11.1 and 2.0‰ , while the respective $\delta^2\text{H}$ values vary between -78.2‰ and -13.8‰ (Supplementary Table S1). Overall, the data reveal

a very good correlation between $\delta^{18}\text{O}$ and $\delta^2\text{H}$ ($r = 0.99$). The extruded water sample shows relatively low $\delta^{18}\text{O}$ (-8.7‰) and $\delta^2\text{H}$ (-65.1‰) values, however, only a small offset to isotopic of an adjacent pool (-9.5‰ $\delta^{18}\text{O}$ and -71.1‰ $\delta^2\text{H}$) could be revealed.

The analysis of plant samples from the CTP reveal distinct differences between *O. andina* and *Z. muticus* (Table 2). With regard to the cellulose yield, a large difference between leaves of *O. andina* (19.3%) and *Z. muticus* (10.9%) becomes apparent. The difference between the rhizomes is much smaller (*O. andina* 14.1% and *Z. muticus* 16.6%) and almost indistinguishable with respect to the small sample number. The isotope measurements of the extracted cellulose indicate that leaves of *Z. muticus* have lower $\delta^{18}\text{O}_{\text{cell}}$ values (28.9‰) than leaves of *O. andina* (31.68‰), but that *Z. muticus* has slightly higher $\delta^{18}\text{O}_{\text{cell}}$ values in rhizomes compared to *O. andina* (28.2 to 27.6‰). The $\delta^{13}\text{C}_{\text{cell}}$ dataset indicate values as expected for C3 plants (Pate, 2001). $\delta^{13}\text{C}_{\text{cell}}$ values are consistently enriched in leaves and rhizomes of *Z. muticus* compared to leaves and rhizomes of *O. andina* (leave $\Delta\delta^{13}\text{C}_{\text{cell}}$: Zam–Oxy = 0.6‰, rhizome $\Delta\delta^{13}\text{C}_{\text{cell}}$: Zam–Oxy = 0.8).

Whereas the intraplant difference between leaves and rhizomes in the carbon isotope composition ($\delta^{13}\text{C}_{\text{I-r}}$) is very low, major differences occur in the oxygen isotope composition of the investigated species ($\Delta\delta^{18}\text{O}_{\text{I-r}}$ = 0.73‰ for *Z. muticus*; $\Delta\delta^{18}\text{O}_{\text{I-r}}$ = 4.11‰ for *O. andina*). Comparing distinctive leave-rhizomes pairs within the same cushion of *O. andina* results in an even higher spread ($\Delta\delta^{18}\text{O}_{\text{I-r}}$ = 4.95‰).

Core Data

Bulk Isotope Composition, Element Contents, LOI₅₅₀ and Humification

Values of $\delta^{13}\text{C}_{\text{OM}}$ reveal low variability between -26 and -25‰ during most parts of the record (Figure 4). Highest values occur between 1700 and 1270 cal. yr BP (up to -24.5‰), before a decreasing trend toward lower $\delta^{13}\text{C}_{\text{OM}}$ values prevail until 230 cal. yr BP. The $\delta^{15}\text{N}_{\text{bulk}}$ values shift considerably between 4.0 (~ 2060 cal. yr BP) and 1.1‰ (~ 390 cal. yr BP) with phases of higher (1730 to 1340, 700 to 610, and around 110 cal. yr BP) and lower (2910 to 2300, 1240 to 1010, 500 to 350 and 50 to -30 cal. yr BP) values. The TOC content ranges between 42.8

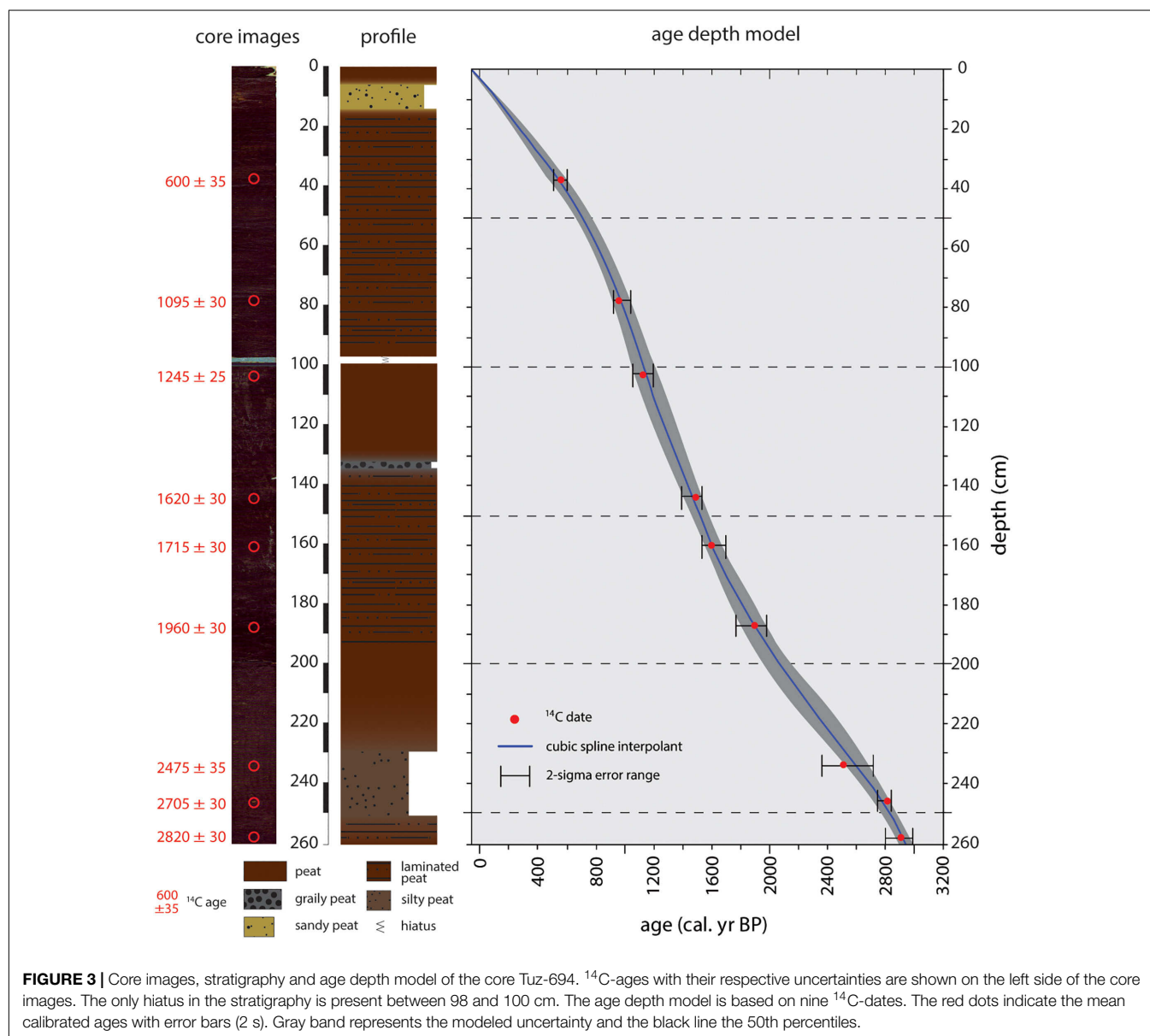


TABLE 2 | Cellulose isotope values of modern plant samples from the Cerro Tuzgle Peatland.

Plant samples	<i>n</i>	Cellulose yield (%)	<i>SD</i>	$\delta^{18}\text{O}$ (‰)	<i>SD</i>	$\delta^{13}\text{C}$ (‰)	<i>SD</i>	$\Delta \delta^{18}\text{O}$ l-r (‰)	$\Delta \delta^{13}\text{C}$ l-r (‰)
<i>Oxychloe andina</i>									
Leaves	6	19.3	3.06	31.68	3.88	-24.59	0.39	—	—
Rhizomes	4	14.1	1.54	27.57	1.84	-24.78	0.38	4.11*	0.19
Fruits	2	8.6	2.76	27.10	0.85	-25.51	0.72	—	—
<i>Zameioscirpus muticus</i>									
Leaves	4	10.9	1.39	28.88	2.15	-23.98	0.61	—	—
Rhizomes	4	16.6	4.39	28.15	2.20	-23.94	0.29	0.73	-0.04
Fruits	—	—	—	—	—	—	—	—	—

Number of samples (*n*), mean cellulose yields, mean $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ values of cellulose, mean isotopic difference between leaves and rhizomes ($\Delta \delta^{18}\text{O}_{l-r}$, $\Delta \delta^{13}\text{C}_{l-r}$), standard deviation (*SD*). *Comparison of *Oxychloe andina* leaves and rhizomes from the same plant cushion (*n* = 2) results in a $\Delta \delta^{18}\text{O}_{l-r}$ of 4.95 ‰.

(~2910 cal. yr BP) and 16.3% (~1390 cal. yr BP). On average, TOC contents remain between 25 and 40% even in sections with increased sedimentation rates, e.g., between 2740 and 2510 cal. yr BP. Only during short intervals where clastic input can be

observed, the TOC values are lower (e.g., 21.9% around 1370 cal. yr BP). The TN content shows low variations around 2%, but is well correlated with TOC ($r = 0.83$). A conspicuous peak occurs around 2060 cal. yr BP (3.3%). The TOC/TN ratio varies

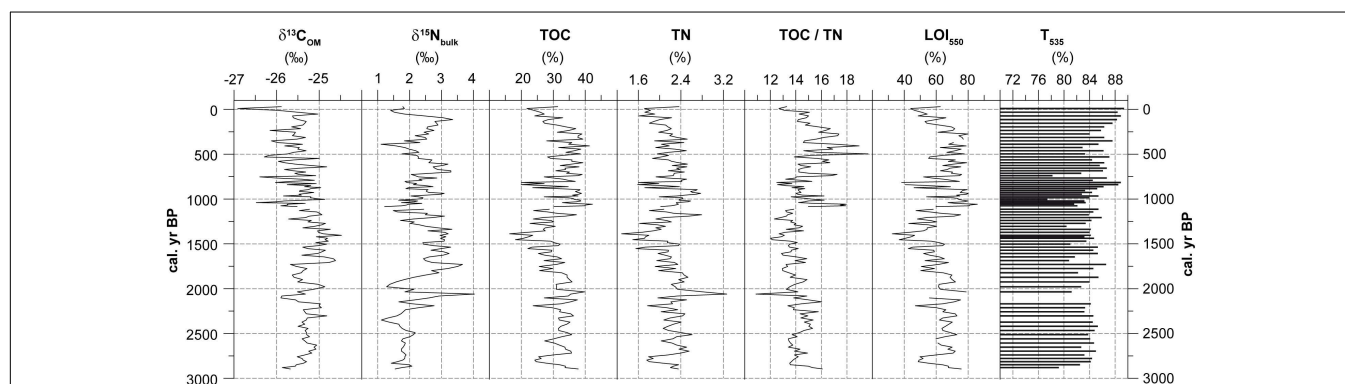


FIGURE 4 | Stable isotopes and element composition on bulk material. $\delta^{13}\text{C}_{\text{OM}}$ (‰ vs. VPDB), $\delta^{15}\text{N}_{\text{bulk}}$ (‰ vs. AIR), TOC (%), TN (%) values as well as the TOC/TN ratio, LOI₅₅₀ and T₅₃₅. Missing values are due to insufficient bulk material for analyses and the hiatus between 1080 and 1110 cal. yr BP.

between 10.9 (~2060 cal. yr BP) and 19.7% (~500 cal. yr BP). The transition from low TOC/TN values combined with low variability to higher values with an increased variability after 1080 cal. yr BP is prominent. LOI₅₅₀ ranges between 32.5 (~1390 cal. yr BP) and 85.7% (~1060 cal. yr BP). On average, LOI₅₅₀ values remain low between 2880 (67.7%) and 1080 (59.8%) cal. yr BP. Comparably high values occur between 1030 and 270 cal. yr BP. T₅₃₅ values shift between 77.4% (~1000 cal. yr BP) and 89.4% (~10 cal. yr BP). While moderate fluctuations dominate the record, a trend toward higher values is prominent between 270 and –10 cal. yr BP.

A slight negative correlation ($r = -0.17$) exists between TOC and $\delta^{13}\text{C}_{\text{OM}}$ while no correlation exists between TN and $\delta^{15}\text{N}_{\text{bulk}}$ (Supplementary Table S2). Similarly, only a weak correlation occurs between $\delta^{13}\text{C}_{\text{OM}}$ and $\delta^{15}\text{N}_{\text{bulk}}$ ($r = 0.16$). T₅₃₅ values reveal a negative correlation with $\delta^{13}\text{C}_{\text{OM}}$ ($r = -0.4$) and with TOC ($r = -0.39$).

Cellulose Oxygen and Carbon Isotopes

The C/O ratios of cellulose from all three investigated size fractions fall into a small range around 0.9 and, thus, indicate that the extraction provided cellulose free of inorganic and organic contaminants (Supplementary Figure S1, see Wissel et al., 2008 for details).

In general, the $\delta^{18}\text{O}_{\text{cell}}$ values of the different size fractions fluctuate between 28.0 and 31.0‰ and show comparable trends (Figure 5). Conspicuous peaks (e.g., ~1320 cal. yr BP) or phases with increased (e.g., 2210 to 2120, 1800 to 1600 or 1160 to 1080 cal. yr BP) or reduced values (e.g., 2530 to 2230 or 610 to 130 cal. yr BP) occur in all three time series within the same depths. The high correlation between the fractions < 200 μm and 200–1000 μm ($r = 0.93$) confirms a strong similarity between these size fractions. $\delta^{18}\text{O}_{\text{cell}}$ values of the fraction > 1000 μm are comparable to those of the two smaller fractions, however, with a reduced variability and slight differences in the temporal development. This is confirmed by weaker, but still remarkably high correlations ($r \geq 0.59$).

Comparing all size fractions, the $\delta^{13}\text{C}_{\text{cell}}$ values vary on average between –22.0 and –23.0‰. However, a transition from lower to higher variable $\delta^{13}\text{C}_{\text{cell}}$ values after 1080 cal.

yr BP is prominent. Further, a slight trend toward decreasing values after 1080 cal. yr BP can be distinguished that is mostly visible in the fraction > 1000 μm . The two smallest fractions reveal a high correlation ($r = 0.61$). Correlations between the smallest and the largest ($r = 0.35$), as well as between the middle and largest fraction ($r = 0.52$) are lower. Within fractions, correlations between $\delta^{18}\text{O}_{\text{cell}}$ and $\delta^{13}\text{C}_{\text{cell}}$ reach r values between 0.17 and 0.48.

In comparison to $\delta^{13}\text{C}_{\text{OM}}$ values, $\delta^{13}\text{C}_{\text{cell}}$ values are enriched by about ~3‰ on average. This difference is comparable with the results of Ménot and Burns (2001) for vascular plants and of Moschen et al. (2009) for *Sphagnum* peatlands respectively. The trends of $\delta^{13}\text{C}_{\text{cell}}$ and $\delta^{13}\text{C}_{\text{OM}}$ show similarities that are confirmed by correlation coefficients between $r = 0.34$ and $r = 0.42$. However, it is conspicuous that $\delta^{13}\text{C}_{\text{OM}}$ values are not decreasing within the upper 50 cm. No correlation occurs for $\delta^{13}\text{C}_{\text{cell}}$ and TOC, while slight negative correlations between $\delta^{18}\text{O}_{\text{cell}}$ and TOC values exist ($r \leq -0.26$). A slight negative correlation can be found between T₅₃₅ and $\delta^{13}\text{C}_{\text{cell}}$ ($r \leq -0.4$) while no correlation exists between T₅₃₅ and $\delta^{18}\text{O}_{\text{cell}}$ ($r \geq 0.1$).

Visual inspection of size fractions prepared for cellulose extraction revealed further details of sample compositions. According to nine inspected samples, the largest size fraction (> 1000 μm) is mainly comprised of rhizomes with contributions reaching almost 100% and minor admixtures of leaves. The medium size fraction (200–1000 μm) in comparison contains larger amounts of leaf fragments resulting roughly in a 40/60% mixture of rhizomes fragments and leaves. The content of the smallest size fraction (< 200 μm) was unidentifiable.

Geochemical Composition From XRF Measurements

Sulfur (S) and Calcium (Ca) values reveal a close similarity ($r = 0.99$) with very low intensities (total counts = cnts) between 2900 and 2070 cal. yr BP and again from 1060 to –60 cal. yr BP (Figure 6). Strongly increased values occur around 1900, 1650, 1520 and between 1310 and 1130 cal. yr BP. On average, Manganese (Mn) counts fluctuate between 50 and 200. Slightly increased values occur around 2780 and 870 cal. yr BP, while sharply increased values are evident between 260 and 10 cal. yr BP. The counts of Iron (Fe) vary between 5000 and 15000

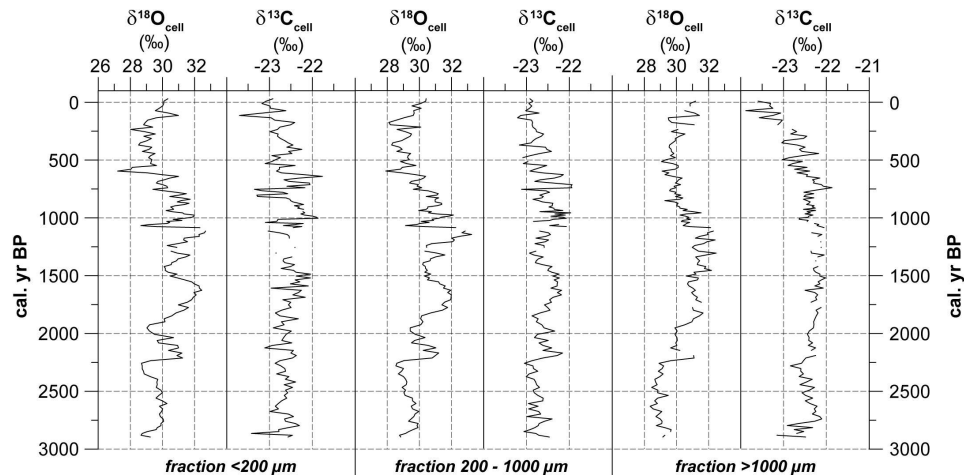


FIGURE 5 | Stable oxygen and carbon isotope composition of cellulose. Given are $\delta^{18}\text{O}_{\text{cell}}$ (‰ vs. VSMOW) and $\delta^{13}\text{C}_{\text{cell}}$ (‰ vs. VPDB) for the respective fractions < 200 μm , 200–1000 μm and >1000 μm . Missing values are due to insufficient cellulose material for analyses and the hiatus between 1080 and 1110 cal. yr BP.

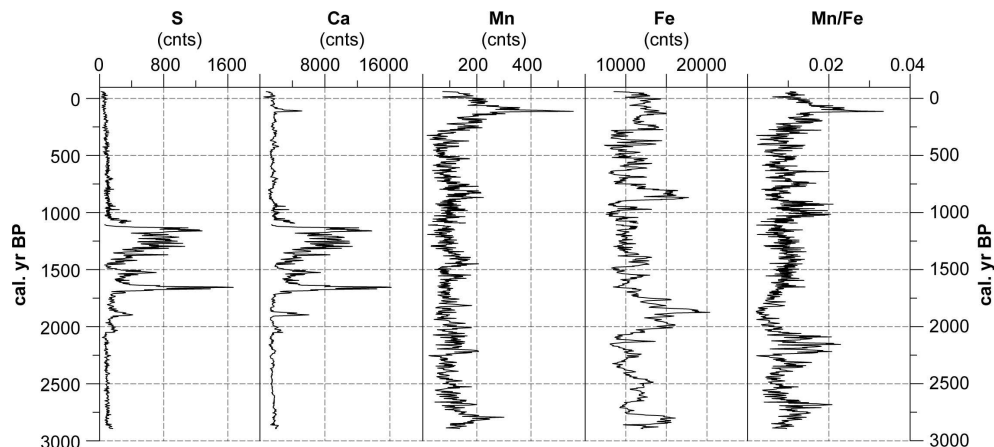


FIGURE 6 | Results of XRF analyses. Given are S, Ca, Mn and Fe values as well as the Mn/Fe-ratio. Missing values are due to the hiatus between 1080 and 1110 cal. yr BP.

cps during the most phases of the record. Highest values occur between 2850 and 2780, 2000 and 1750, and between 890 and 800 cal. yr BP. The Mn/Fe ratio fluctuates mostly around 0.01, however, with increased values around 2680, between 2220 and 2080, 1020 and 920 and between 210 and 10 cal. yr BP. Reduced values can be distinguished between 2000 and 1750 and between 400 and 320 cal. yr BP. Interestingly, the peaks of Ca and S occur in the phase with the highest $\delta^{18}\text{C}_{\text{cell}}$ values.

DISCUSSION

Implications From Modern Hydrology

The isotopic composition of modern water samples gives an insight into the present hydrological system of CTP. In the

headwater as well as the main accumulation section of the peatland (**Figure 7**), two springs occur (Schitteck et al., 2016). These springs are presumably seasonal cold-water springs, located in the Ordovician basement rocks that are characterized by secondary permeability (Giordano et al., 2013). Giordano et al. (2013) describe this part of the Cerro Tuzgle-Tocomar geothermal region as recharge area, thus, the CTP is affected by groundwater of recent precipitation periods and not by deeper geothermal groundwater sources. Following this concept, the isotopic source values of the peatlands' water are directly linked to the isotope composition of local precipitation, though local evaporation processes during infiltration may have additional influence. However, a multiyear smoothing of the input isotope signal has to be expected due to soil/groundwater residence time and the smoothing effect of the peat water reservoir.

The isotopic composition all CTP water samples are compared with the Global Meteoric Water Line [GMWL, $\delta^2\text{H} = 8.13\delta^{18}\text{O} + 10.8$, Rozanski et al. (1993)], as the existing database (e.g., Global Network of Isotopes in Precipitation - GNIP) is not sufficient to develop a Local Meteoric Water Line. Irrespective of the observed spatial heterogeneity, the samples reveal regularity as they fall on a single line in δ space, clearly describing the modern evaporation line (LEL) for CTP (Figure 7). This is also true for the extruded plant water sample. The slope of the observed relation ($\delta^2\text{H} = 4.85\delta^{18}\text{O} - 25.05$) is typical for evaporating surface waters (e.g., Mook, 2001) that have to be expected due to the local environmental conditions. We interpret this behavior as another evidence for a single, isotopic homogeneous water source for CTP, i.e., that meteoric water (precipitation, young groundwater) recharges the peatland. Thus, older and potentially isotopically different groundwater is not involved in the hydrology of the peatland which in turn is closely connected to the meteoric water cycle. This is supported by low electrical conductivity values of the water (Schitteck et al., 2016).

The surface water data set from October 2013 gives further insights into the spatial heterogeneity of water isotopes. Samples close to the two springs (Tuz-1, Tuz-2, Tuz-12, Tuz-13, Tuz-21, Tuz-22) are characterized by a low degree of ^{18}O enrichment compared to spring values, thus low evaporative enrichment (Figure 8). All other samples of the CTP suggest a higher degree of evaporative enrichment, e.g., in isolated pools (Tuz-5) that are surrounded by dense cushion vegetation without interspersed channels. The water samples of the lake section (Tuz-9 and Tuz-10) exhibit the strongest degree of evaporation due to their large open water surface prone to evaporation. On the one hand, these spatial heterogeneities indicate the strong effect of microtopographical differences for evaporative isotope enrichment of surface water in cushion peatlands. Similar effects have already been described for *Sphagnum* peatlands (Ménot-Combes et al., 2002). On the other hand, they point toward a limited movement respectively exchange of water within the peatland despite superficial connections through

visible rivulets (Figure 2). It seems likely that with increasing dryness or temporal drought, the number of isolated pools in the peatland will grow causing a respective response toward higher $\delta^{18}\text{O}$ and $\delta^2\text{H}$ values in these pools. The implication for a microtopographical habitat, i.e., a potentially temporal stable single cushion in the peat land is that variations in water availability through changes in e.g. evaporation, precipitation, connectivity or growth are systematically translated into a respective response of the local source water isotopic signature.

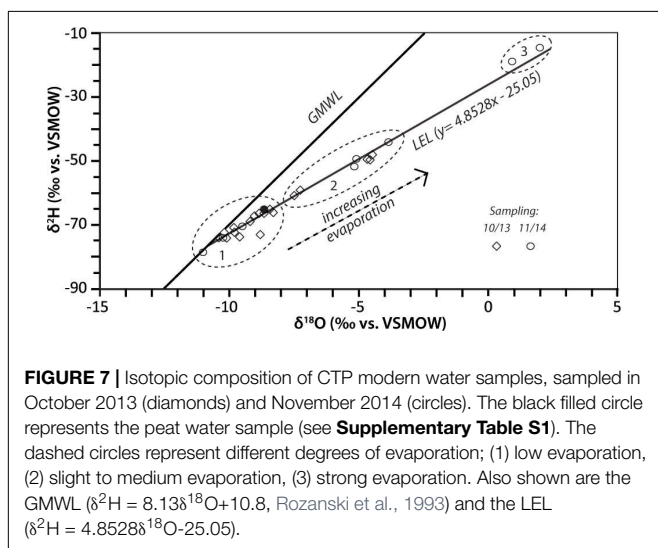
Implications From Modern Vegetation

The modern plant samples reveal distinct isotopic differences between *O. andina* and *Z. muticus* (Table 2). In contrast to *Z. muticus* ($\Delta\delta^{18}\text{O}_{\text{L-r}} = 0.7\text{‰}$) a huge offset in $\delta^{18}\text{O}$ between rhizomes and leaves of *O. andina* ($\Delta\delta^{18}\text{O}_{\text{L-r}} = 4.1\text{‰}$) is evident. Even larger differences between leaves and roots were reported for the vascular plant *Empodisma* ($\sim\Delta\delta^{18}\text{O}_{\text{L-r}} = 8.4\text{‰}$ to 11.8‰) (Amesbury et al., 2015b). We suggest that this offset arises from dissimilar leaf structures. Leaves of *O. andina* are described as lignified (Ruthsatz, 2008), thus, might possess a higher protection against transpiration, leading to increased $^{18}\text{O}/^{16}\text{O}$ ratios of leaf water and, finally, of leaf cellulose relative to *Z. muticus*. Considering these intraspecies $\delta^{18}\text{O}_{\text{cell}}$ ranges, vegetation changes should have a minor influence on the $\delta^{18}\text{O}$ signal in the size fractions, i.e., 200–1000 μm , since values of *Z. muticus* leaves and rhizomes are within the limits of *O. andina*. Given the small species difference for rhizomes (Table 2), such effects should be negligible for the largest size fraction ($>1000\text{ }\mu\text{m}$).

In comparison with other investigations on vascular (Amesbury et al., 2015a) and non-vascular peatlands (e.g., Loisel et al., 2009), the ranges of $\delta^{13}\text{C}$ cellulose values are small for both species (Table 2). However, previous studies reveal that interspecies variability can have significant influences on the $\delta^{13}\text{C}_{\text{cell}}$ signal (Ménot and Burns, 2001). In the present study, mean $\delta^{13}\text{C}_{\text{cell}}$ values of *Z. muticus* are enriched for leaves ($\delta^{13}\text{C}_{\text{Zam-Oxy}} = 0.61\text{‰}$) and rhizomes ($\delta^{13}\text{C}_{\text{Zam-Oxy}} = 0.84\text{‰}$) respectively in contrast to *O. andina*. Thus, the potential $\delta^{13}\text{C}_{\text{cell}}$ change to be expected for a complete (but unlikely) vegetation change from one species to the other would amount to 0.6–0.8‰. Further investigations on the range of $\delta^{18}\text{O}_{\text{cell}}$ and $\delta^{13}\text{C}_{\text{cell}}$ variations within a single cushion in the same microenvironment could help identify causes for the observed isotopic differences.

Peat Forming Vegetation and Decomposition

To use cushion peatlands as continuous high-resolution archives aspects like changes in plant composition need to be considered. Peatlands consisting of *Oxychloe* are poor in species diversity, in contrast to e.g., *Distichia* dominated peatlands, are especially suitable for paleoenvironmental studies (Ruthsatz, 2008) because species-specific effects are limited. This notion applies with regard to the CTP record, where only *O. andina* and *Z. muticus* dominate the peatland. Our measurements on modern plant samples indicate that changes in the occurrence between the species could cause a smoothing of $\delta^{18}\text{O}$ variations recorded



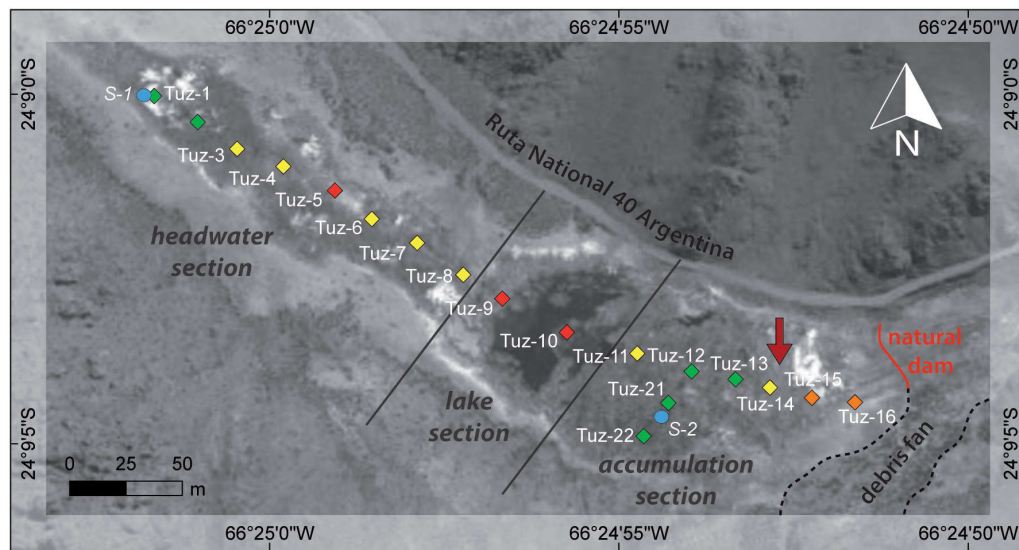


FIGURE 8 | Location of water samples from October 2013 on the CTP. Colors indicate relative $\delta^{18}\text{O}$ enrichment in the samples, i.e., the degree of evaporation (green: low or absent evaporation, yellow: slight evaporation, orange: medium evaporation, red: strong evaporation). Diamonds indicate surface peatland samples (Tuz), blue circles (S1 and S2) indicate the location of the springs, the red arrow the location of core Tuz-694. The debris fan (dashed line) presumably favors the development of the natural dam (red line). The main sections of the peatland are illustrated according to Schitteck et al. (2016) (data source: DGM-GTOPO30).

in the peat core and might lead to a slight shift in $\delta^{13}\text{C}_{\text{cell}}$ values (see above). However, macrofossil analyses confirm that *O. andina* is the dominant peat forming species in CTP within the whole record (Schitteck et al., 2016; Schitteck, unpublished data). Furthermore, macrofossil analyses indicate an increase of *Z. muticus* between 80 cm and 130 cm depth (Schitteck et al., 2016; Schitteck, unpublished data), without any visible response in our isotope proxies excluding any vegetational bias in the records.

Another question is, whether different size or compositional fractions of the peat possess different (climate-) signals. Correlation analyses reveal a strong respectively good accordance between our two smaller size fractions ($r = 0.93$ for $\delta^{18}\text{O}$, $r = 0.61$ for $\delta^{13}\text{C}$), indicating that the fractions $< 200 \mu\text{m}$ and $200\text{--}1000 \mu\text{m}$ comprise the same signal. Weaker correlations of $\delta^{18}\text{O}$ of these fractions with the fraction $> 1000 \mu\text{m}$ can be linked to the composition of the size fractions since the largest fraction mainly consist of rhizomes while the smaller fractions have varying contributions of rhizomes and leaves. According to our modern plant data (see above), *O. andina* dominated phases should possess an offset $\delta^{18}\text{O}$ between the smaller (enriched) fractions and the largest fraction dominated by (depleted) rhizomes. However, this expectation is not met by our data, even if some parts of the record exhibit more depleted $\delta^{18}\text{O}_{\text{cell}} > 1000$ values. Most likely a direct comparison between the isotopic compositions of living leaves with already decomposed leaves or other plant parts in the peat is not feasible. A way to overcome this problem could be to use already died off leaves age groups of preceding years of still living plants. Modern $\delta^{13}\text{C}_{\text{cell}}$ values of *O. andina* exhibit a small deviation between leaves and rhizomes ($\Delta\delta^{13}\text{C}_{\text{L-R}} = 0.2\text{‰}$), thus, the development in every fraction is expected to be very similar. This assumption is mainly confirmed despite some

deviations regarding the $\delta^{13}\text{C}_{\text{cell}} > 1000$ record. Reasons for these deviations could be a change in the contribution of the remaining plant parts (leaves, rhizomes) to the composition of the smaller fractions. Overall, the general isotopic trends in all fractions reveal a good match, wherefore we argue that the primordial environmental signal is imprinted and inherited in all three size fractions.

Decomposition can have an influence on the stable isotope composition of bulk peat or cellulose. Burial and humification of peat material removes at first the more labile organic fractions, thus may significantly change the $\delta^{13}\text{C}$ values of bulk organic matter (Ménot and Burns, 2001) toward enriched $\delta^{13}\text{C}$ values in the residual peat (e.g., Novak et al., 2010). A raised degree of humification can potentially be identified by low T_{535} values (e.g., Chambers et al., 2011). In the CTP record, the variation of T_{535} is very limited indicating that the influence of decomposition on isotope records of bulk peat is negligible, after the step of initial removal of mobile and soluble phases. Starting from fresh plant litter, cellulose is also decomposed to a comparable or even larger degree as lignin after initial removal of soluble components (e.g., Sjöberg et al., 2004; Fioretto et al., 2005). However, not much is known on isotopic changes through cellulose decomposition especially with respect to the oxygen isotopic composition of cellulose. During an artificial aging experiment, Schleser et al. (1999) observed a maximum deviation of 0.3‰ from the initial $\delta^{13}\text{C}$ value of wood cellulose, however, with increasing degree of cellulose decomposition the $\delta^{13}\text{C}$ values of the remaining cellulose again reached their initial values. Thus, we expect isotope effects of decomposition on cellulose, despite strong reduction of cellulose content, to be small. Identical (within analytical uncertainty) mean cellulose oxygen and carbon stable isotope values of all size fractions further support this concept.

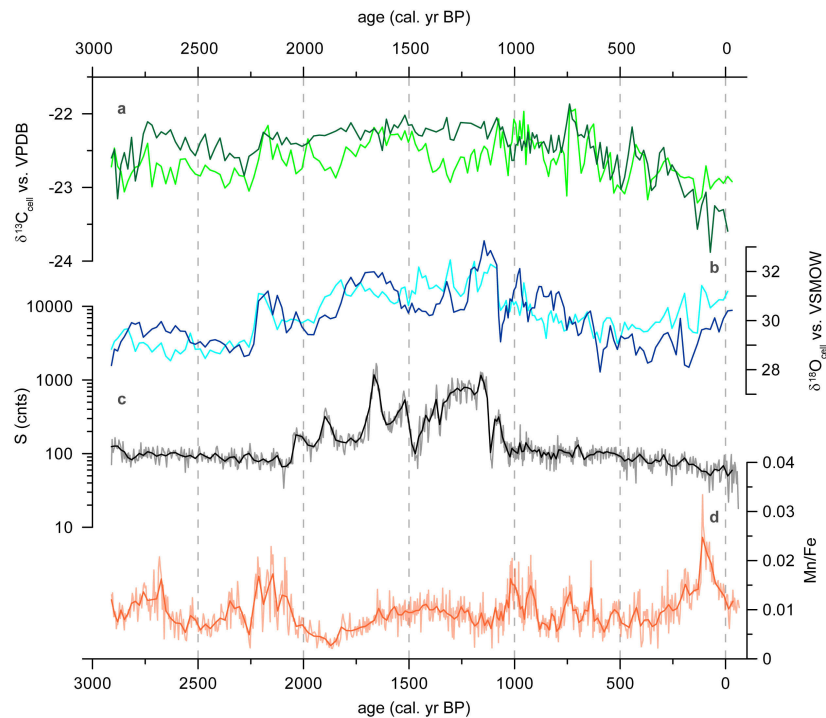


FIGURE 9 | Results of core Tuz-694. **(a)** $\delta^{13}\text{C}_{\text{cell}}$ (‰ vs. VPDB) for fraction 200–1000 μm (light green) and >1000 μm (dark green), **(b)** $\delta^{18}\text{O}_{\text{cell}}$ (‰ vs. VSMOW) for fraction 200–1000 μm (dark blue) and fraction > 1000 μm (light blue), **(c)** Sulfur content (black: 5 pt. average, log10-scale), **(d)** Mn/Fe ratio (orange: 5 pt. average).

Paleoclimatic Implications From the CTP Proxy Dataset

The $\delta^{18}\text{O}$ values recorded in plant cellulose are controlled by several (local) influences like evaporation degree of the surface water or changes in peat composition (see above). However, a major factor is the isotopic composition of the source water. Climate modeling studies (e.g., Vuille et al., 2003; Vuille and Werner, 2005) point out that $\delta^{18}\text{O}$ values of precipitation ($\delta^{18}\text{O}_{\text{prec}}$) in the Central Andes are affected by precipitation amount, moisture source, degree of rainout upstream and re-evaporation, condensation temperature and atmospheric circulation. Independent of the variety of factors, Global Network of Isotopes in Precipitation (GNIP) station data clearly indicate a significant negative correlation of $\delta^{18}\text{O}$ values with precipitation amount for different parts in South America (IAEA/WMO, 2004). While low gradients are observed from the coastal area of Brazil (e.g., Belém GNIP station, 24 m a.s.l.: $-0.113\text{‰}/10\text{ mm}$), higher gradients occur in the Andean Highlands (e.g., Gran Chocaya GNIP station, 4340 m a.s.l.: $-1.226\text{‰}/10\text{ mm}$) (**Supplementary Table S3**). The connection between precipitation amount and $\delta^{18}\text{O}$ is supported by Ballantyne et al. (2011) who infer decreased $\delta^{18}\text{O}_{\text{prec}}$ values in more humid periods in a study of tree ring cellulose in Brazil (3.18°S, 60.08°W), Peru (12.68°S, 69.28°W) and northern Argentina (22.08°S, 66.08°W).

Based on this fundamental relation, we argue that increased precipitation amounts at the study site are directly connected with more depleted source water for CTP cushion plants and vice

versa. Furthermore, we argue that recent precipitation forms the primary source of water for the cushion plants. Since water influx into the peatland from the north or from older groundwater can be excluded (see above), water solely reaches the CTP in form of (1) direct precipitation on the peatland and superficial run-off from the surrounding slopes as well as from the ephemeral stream and (2) recharge from two springs from shallow groundwater (see also Schitteck et al., 2016).

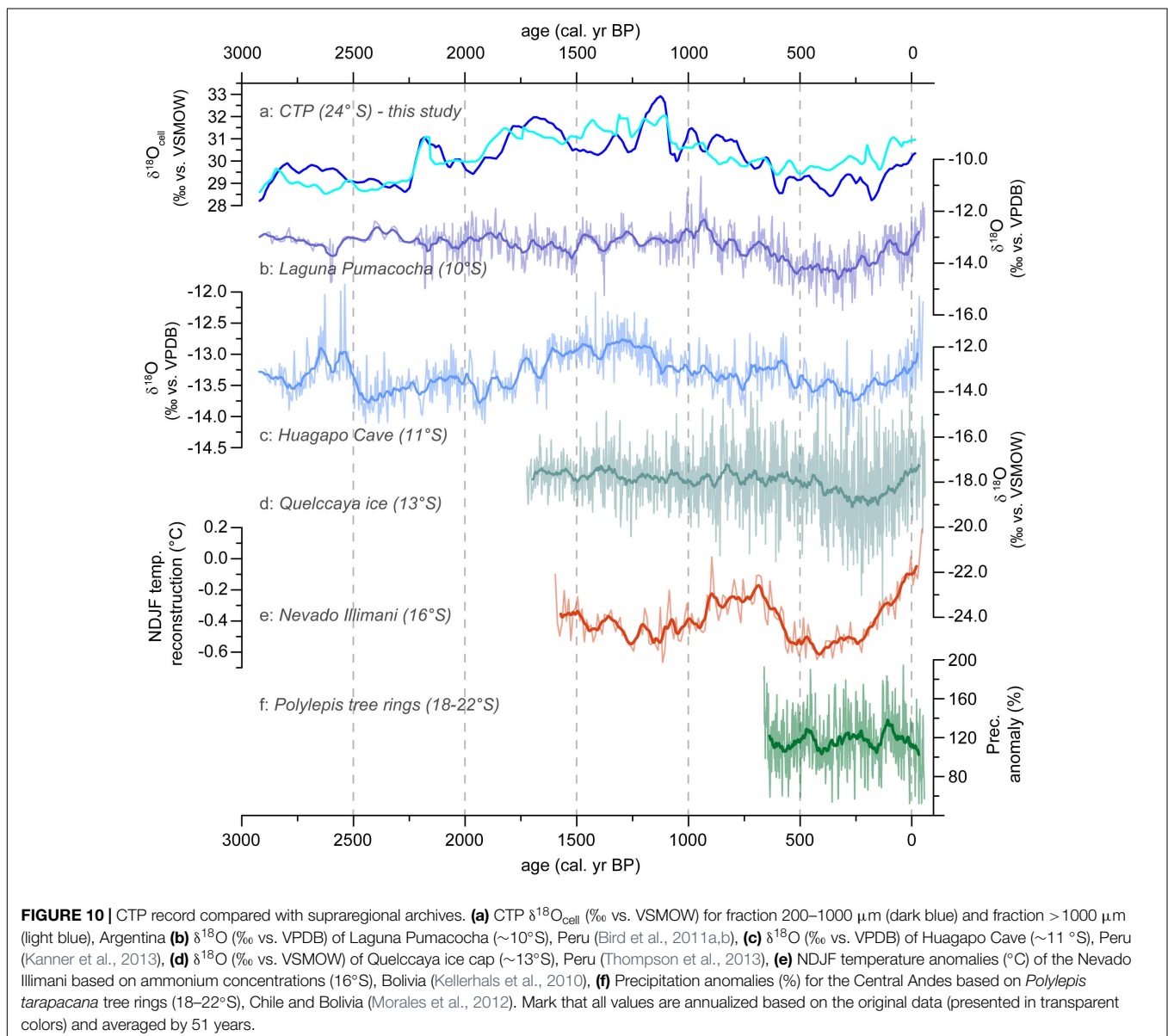
Accordingly, during phases of increased humidity, depleted precipitation reaches CTP. Due to a high water volume (but nearly constant surface area), evaporation has potentially lower influence on the $\delta^{18}\text{O}$ values of the water within the peatland. With sufficient water supply (considering site conditions), we expect the vascular plants *O. andina* and *Z. muticus* to have comparably open stomata, leading to comparably less ^{18}O enrichment in the leaf water (e.g., Roden et al., 2000). Overall, these factors operate additive and lead to relatively depleted $\delta^{18}\text{O}_{\text{cell}}$ values within the cushion plants. Vice versa, periods with increased aridity are characterized by low but enriched precipitation reaching the study area, stronger influence of local evaporation effects, reduced stomatal conductance and increasing fractionation with enriched $\delta^{18}\text{O}_{\text{LW}}$ values, and finally enriched $\delta^{18}\text{O}_{\text{cell}}$ values. Thus, we interpret $\delta^{18}\text{O}_{\text{cell}}$ values to be a proxy for moisture conditions at CTP (**Figure 9b**).

Beside oxygen isotopes, the stable carbon isotope compositions of bulk organic matter or cellulose also provide insights into local environmental and climatic conditions (**Figure 9a**). Through the stomatal aperture transpiration

and photosynthesis in leaves are coupled processes, which in turn have strong impacts on the stable carbon and oxygen isotope composition of new photosynthates. Due to this basic dependency of both variables on stomatal aperture, i.e., because gas exchange through the stomata influences both carbon and water cycle of the plants, a certain relation can be expected (Smedley et al., 1991; Barbour et al., 2002; Chaves et al., 2004; Gessler et al., 2009). Since photosynthetic carbon fixation solely influences the plant carbon cycle, such a relation might be very variable in strength. Nevertheless, further evidence for our interpretation of $\delta^{18}\text{O}_{\text{cell}}$ as moisture indicator for CTP may be derived from a correlation between $\delta^{13}\text{C}_{\text{cell}}$ and $\delta^{18}\text{O}_{\text{cell}}$. Indeed, positive significant correlations appear in our cellulose stable isotope records (Supplementary Table S2). Correlation coefficients between $\delta^{13}\text{C}_{\text{cell}}$ and $\delta^{18}\text{O}_{\text{cell}}$ reach values of

about $r = 0.5$ for the medium size fractions. Overall, these positive correlations between $\delta^{13}\text{C}_{\text{cell}}$ and $\delta^{18}\text{O}_{\text{cell}}$ support our interpretation that more enriched $\delta^{18}\text{O}_{\text{cell}}$ values are indicative for more severe moisture stress at CTP.

The ratio of Mn/Fe is used as an indicator of redox conditions in CTP (Schitteck et al., 2015, 2016). Accordingly, anoxic conditions in the peat water column and presumably stable water conditions are indicated by high Mn/Fe values (Figure 9d). Principally, the place of signal generation differs between these proxies (leave, “catotelm”) which might already cause temporal deviations. Also, Mn and Fe can be of lithogenic origin and transported to the CTP while redeposition processes can further hinder an unambiguous interpretation. $\delta^{18}\text{O}_{\text{cell}}$ values can be affected in detail by several factors like precipitation amount, moisture source, moisture transport history or the degree of



rainout upstream (Insel et al., 2013). Further investigations are needed to explain the differences between these two proxies connected to the peatland hydrology in more detail.

The striking parallel curves of S and Ca counts ($r = 0.99$) suggests the hypothesis that these element counts are linked to gypsum precipitates, developed under dry conditions without strong water-table fluctuations (e.g., Litaor et al., 2004). Although the values might indicate secondary gypsum precipitates after core opening, we argue that the ion concentration did not change in the plant pore water, and thus, reflect climatic conditions during synthesis. Interestingly, the phase of strongly increased counts of S and Ca appear together with highest values of $\delta^{18}\text{O}_{\text{cell}}$ (Figure 9c), which again strengthen our interpretation of $\delta^{18}\text{O}_{\text{cell}}$ as proxy for moisture conditions at CTP.

Regional Paleoclimate Comparison

Based on our $\delta^{18}\text{O}_{\text{cell}}$ record, we observe phases with enhanced and reduced moisture supply at the CTP (24°S, Figure 10a). Wetter conditions prevailed from 2750 to 2250 and from 600 to 130 cal. yr BP, while drier conditions can be identified between 2190 and 2120, 1750 and 1590, 1200 and 1080 and since 130 cal. yr BP. In order to interpret our findings in the context of changes in the large scale atmospheric circulation, we compare the CTP record with local and supraregional paleoclimate records of the high Andes between 10 and 24°S (Figure 10).

Regional atmospheric circulation is mainly driven by (1) the strength of the Atlantic Meridional Overturning Circulation, (2) latitudinal changes of the Intertropical Convergence Zone (ITCZ), (3) the dipole-like pattern of the South Atlantic Convergence Zone and the South American Low Level Jet (SALLJ) and (4) the position of the Bolivian High (Liebmann and Mechoso, 2011; Marengo et al., 2012; Vuille et al., 2012; Rojas et al., 2016; Novello et al., 2018). During austral spring, moisture transport from the North Atlantic to the Andean highlands starts with a weakening of the meridional temperature gradient between October and November, leading to a southward displacement of the SALLJ. Simultaneously, connected to this shift of the ITCZ over the tropical Atlantic and Pacific, deep convection over the central part of the continent starts to develop (Hastenrath, 1997; Marengo et al., 2012). Finally, this leads to the development of the SASM during the summer months, thus, to N-S moisture transport reaching the NW Argentine Andes (Garreaud, 2000; Vera et al., 2006; Vuille et al., 2012). In the northern Central Andes, numerous archives are described to have a close $\delta^{18}\text{O}$ dependence to the SASM development (e.g., lake sediments: Bird et al., 2011a,b; speleothems: Kanner et al., 2013; ice caps: Thompson et al., 2013; tree rings: Ballantyne et al., 2011). Comparing these sites (see Figure 10d) with the CTP $\delta^{18}\text{O}_{\text{cell}}$ record (Figure 10a) several phases with strong similarities and differences can be detected.

In many parts, comparable curve progressions of $\delta^{18}\text{O}$ records are visible for CTP, Laguna Pumacocha, Huagapo cave and Quelccaya ice cap. High similarity of all archives occurs between 600 cal. yr BP and recent times, covering the Little Ice Age, characterized by decreased $\delta^{18}\text{O}$ values (increased precipitation), and the Current Warm Period, dominated by increased $\delta^{18}\text{O}$ values (decreased precipitation). Comparing the entire CTP

record since 2900 cal. yr BP, highest conformity is visible for Huagapo cave. The good comparability of $\delta^{18}\text{O}$ records from Huagapo cave and CTP offers the potential to resolve latitudinal climate gradients due to transient changes in circulation systems (e.g., SASM) if uncertainties in age models can be accounted for. However, despite the spatial proximity to the Huagapo cave, $\delta^{18}\text{O}$ values of Laguna Pumacocha show stronger deviations compared to CTP.

If we assume that no shift of the seasonal precipitation regimes took place, thus, rainfall occurs only between November and March, other reasons linked to the overturning circulation systems need to be considered. Vuille and Keimig (2004) stated that due to easterly upper- and midtropospheric wind anomalies, three out of phase SASM modes can be differentiated. A northern mode is leading to enhanced convective activity and tropical precipitation in the Andes between ~10 and ~20°S, while a northeastern mode is affecting the eastern Cordillera of the high Andes between ~15 and ~22°S by increased precipitation amounts. Contrary, a southern mode leads to increased precipitation in the southern Central Andes between ~20 and ~30°S, due to a southern displacement of the Bolivian High and a Rossby wave dispersion of the Southern Hemisphere extratropics (Vuille and Keimig, 2004). A more frequent out-of-phase SASM in the past could be a possible explanation for temporal deviations between the proxy records, although extratropical precipitation events as cold air cut off and cold frontal snowfall are typical for the austral winter (Vuille and Ammann, 1997; Barrett et al., 2009). To disentangle the paleoclimatic features in the southern Central Andes, more isotope studies on cushion peatlands are needed.

Temperature reconstruction of the Amazon Basin as the source region for the Ammonium concentration in the Nevado Illimani (Kellerhals et al., 2010; Figure 10e) as well as local temperature reconstruction from the high-Andean Laguna Chepical (32°S, de Jong et al., 2013) excludes the influence of local temperatures on the $\delta^{18}\text{O}_{\text{cell}}$ values of the CTP, which could possibly extend/reduce a vegetation-active period. Previous works already showed that the amount effect and not temperature explains a significant fraction of isotopic variations on seasonal and interannual time scales (e.g., Bird et al., 2011b; Vuille et al., 2012).

As archives covering the last 2900 years are scarce, a local paleoclimate comparison is difficult. Nevertheless, taking small age depth model uncertainties into account, precipitation reconstruction for the Altiplano of Bolivia and Northern Chile based on tree-ring width series of *Polylepis tarapacana* also reveals a good progression with the $\delta^{18}\text{O}_{\text{cell}}$ values of the CTP (Morales et al., 2012; Figure 10f).

CONCLUSION

For the understanding of cushion peatlands as highly suitable paleoclimate archives our study could reveal several important findings. Modern plant samples expose significant differences in the isotope composition of the dominant plant species and clearly indicate the need for analyses of the modern vegetation

prior to the interpretation of any core sample results. Modern water samples reveal strong isotopic differences for single water pools of up to $\Delta\delta^{18}\text{O} = 13.09\text{‰}$ due to evaporation, implying limited movement respectively exchange of water within the peatland despite superficial rivulets. Overall, modern water samples describe a local evaporation line for CTP.

Macrofossil analyses of bulk samples showed that the dominating peat forming species of CTP is *O. andina*. Despite temporal changes in composition of the deposited peat between *O. andina* and *Z. muticus* no detectable influence on the isotopic composition of peat cellulose did occur. Microscopic inspection of peat size fractions revealed distinct differences in their composition (leaves, rhizomes). Irrespective of that, the respective isotope records of all three size fractions are overall consistent.

Increasing precipitation in the study area is connected with decreasing $\delta^{18}\text{O}_{\text{prec}}$ values while less precipitation is accompanied by enriched $\delta^{18}\text{O}_{\text{prec}}$ values (IAEA/WMO, 2004). Also, during more arid phases the effects of local evaporation on the isotopic composition of peat water is stronger than during humid phases. Additionally, the stomata of the cushion plants are constricted or closed under water stress, leading to respectively enriched $\delta^{18}\text{O}_{\text{LW}}$ values, and finally enriched $\delta^{18}\text{O}_{\text{cell}}$ values. Considering these factors together we argue that $\delta^{18}\text{O}_{\text{cell}}$ values are a suitable proxy for moisture conditions at the CTP reflecting local humidity changes during the last 2900 cal. yr BP that are closely connected to the SASM. Discrepancies in the temporal match with other SASM reconstructions suggest impacts of other large-scale atmospheric variability or a different SASM expression at the southerly location of the CTP. According to this finding, predominant dryer conditions dominated at the CTP from 2190 to 2120, 1750 to 1590 and are dominating since 130 cal. yr BP, while more humid conditions prevailed between 2750 and 2250 and between 600 and 130 cal. yr BP. The investigated size fractions show similar developments in their $\delta^{18}\text{O}_{\text{cell}}$ records, however, smaller deviations occur most likely indicating either different compositions or decomposition histories. The obvious differences between the CTP record and other existing climate records reveal the need for further investigations regarding the causes for those discrepancies and provide potential to reveal the history of the spatial (latitudinal) evolution of the SASM during the late Holocene.

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DATA AVAILABILITY

All datasets generated for this study are included in the manuscript and/or the **Supplementary Files**. Raw data can be found in the **Supplementary Table S1** and will additionally be made available via the National Oceanic and Atmospheric Administration (NOAA) Paleoclimate database.

AUTHOR CONTRIBUTIONS

All authors have made substantial contributions to the manuscript in an equal way and approved the submitted version. SK, KS, and AL contributed conception and design of the study. KS and FS provided approval for publication of the content. HV, CO, and HW performed the statistical analysis of the data and helped by the interpretation. SK, AL, and KS wrote sections of the manuscript. LL and JK revised the manuscript critically for important intellectual content.

FUNDING

This work was funded by the German Academic Exchange Service (DAAD) PROMOS – scholarship for Thesis preparation 2015 to SK, German Science Foundation to FS (SCHA 14-1/2), the German Federal Ministry for Education and Research to KS (ARG 06/009), and the National University of Jujuy to LL (SeCTER) which is greatly appreciated.

ACKNOWLEDGMENTS

We thank Gabriel Cortés, Fabio Flores, Gonzalo Torres, and Leandro Rojo for great help during fieldwork. Special thanks belong to Ulrike Langen, Heike Zimmermann, and Josip Sučić for their help in the laboratory. We also thank Sabine Stahl for support and help with the calibration of the XRF results.

SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: <https://www.frontiersin.org/articles/10.3389/feart.2019.00045/full#supplementary-material>

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Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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