The history of subaquatic volcanism recorded in the sediments of Lake Kivu; East Africa

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Abstract Subaquatic volcanic activity has been ongoing in Lake Kivu since the early Holocene and has a dynamic effect on the biological productivity in the surface water, and the preservation of carbonate in the deep anoxic water. Groundwater discharge into the lake's deepwater propels the upward advection of the water column that ultimately supplies nutrients to the surface water for biological production. The amount of nutrients supplied from the deepwater can be increased suddenly by (1) a cold meteorological event that drives deep seasonal mixing resulting in increased nutrients from below and oxygen from above, or (2) subaquatic volcanic activity that induces a buoyant hydrothermal plume, which entrains nutrients from the

deepwater and results in anoxia or suboxic conditions in the surface water. Previous sedimentological studies in Lake Kivu have hypothesized that regional climatic changes are responsible for sudden changes in the preservation of carbonates in the Main Basin. Here we reveal that sublacustrine volcanic events most likely induce the abrupt changes to the geochemistry in the sediment in Lake Kivu. An unprecedented look into the sediment stratigraphy and geochemistry from high-resolution seismic-reflection, and ^{15}N -isotope analyses was conducted in the Main Basin. The results reveal that buoyant hydrothermal plumes caused by subaquatic volcanic activity are a possible trigger for increased biological productivity and organic matter preservation, and that ongoing hydrothermal activity increases the alkalinity in the deepwater, leading to carbonate preservation. The onset of carbonate preservation since the 1970s that is currently observed in the

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sediment could indicate that hydrothermal discharge has recently increased in the lake.

Keywords XRF · ^{15}N · Lake sediment · Subaquatic volcanism · Limnic eruption · High-resolution seismic

Introduction

Lake Kivu is a dynamic system that has been heavily influenced by volcanic activity since its development in the Miocene (Ross et al. 2014). Phreatomagmatic cones border the northern shoreline, and are reminiscent of lavas of the Virunga Volcanic Province (VVP) that have dammed the northern outflow of the lake sometime in the late Pleistocene ~ 10 – 14 ka (Haberyan and Hecky 1987). Furthermore, the sublacustrine basin is marred by submerged volcanic features, which include volcanic domes, edifices, lava flows, and tuff cones (Ross et al. 2014). The lake-damming resulted in an ~ 400 -m lake-level rise that eventually drained south to Lake Tanganyika via the Ruzizi River (Degens et al. 1973; Ross et al. 2014). It was following this blockage that Lake Kivu's thermohaline stratification was initiated as a result of ongoing hydrothermal activity in its deepwater. Since then, at least two periods are described where there has been a possible breakdown of the stratification linked to intense hydrothermal activity and volcanism within the last ~ 10 kyr (Haberyan and Hecky 1987), likely causing a degassing of the lake's deepwater. A similar eruption from the present-day lake would cause significant and detrimental damage to the riparian population, including the two million people living along the shorelines of Lake Kivu.

Presently the lake is meromictic with seasonal mixing in the top ~ 65 m, referred to as the mixolimnion. The monimolimnion is the thermohaline-stratified anoxic water that persists below this depth. It is separated from the mixolimnion by a steep salinity gradient. The stable stratification in the monimolimnion is maintained by warm and salty subaquatic groundwater discharge (SGD), or simply hydrothermal springs (Ross et al. 2015). Major density gradients are observed within the monimolimnion that are maintained at their depths by relatively cold SGD into the lake above the respective gradients (Ross et al. 2015). Observations reveal only small changes

in temperature and conductivity in the monimolimnion since the 1970s, indicating that these properties are currently near steady state. Nevertheless, they also indicate that ongoing warm SGD has been lifting the lower boundaries of the density gradients (Schmid et al. 2012). The cumulative SGD displaces the lake water upwards at vertical advection rates of 0.15 – 0.9 m year^{-1} without disturbing the permanent stratification (Schmid et al. 2005).

It is estimated from the calculated advection rates that the monimolimnion has persisted without complete mixing for ~ 1000 years (Schmid et al. 2005), and has been accumulating dissolved CO_2 and CH_4 during this period. SGD is the main source of CO_2 , whereas CH_4 is a result of ongoing methanogenesis (Schoell et al. 1988; Pasche et al. 2011). CO_2 and CH_4 would erupt from the monimolimnion, and consequently mix the lake, if the combined partial pressure of these gases were to exceed the hydrostatic pressure of the lake at all depths (presently at a maximum of 55 % saturation). The gases could also hypothetically erupt from the lake if subaquatic volcanic activity were significant enough to create a large thermal plume and induce gas oversaturation (Schmid et al. 2003). Degens and Kulbicki (1973) suggested that the lake was mixed at the onset of intense hydrothermal activity ~ 5 ka, and then possibly again at ~ 1 ka. The youngest lake-mixing event correlates well with the estimated minimum age of the monimolimnion calculated by Schmid et al. (2005). However, there is still a paucity of data that indicates a lake eruption has occurred in the past. Furthermore, there is no information that correlates a lake eruption to either gas oversaturation or subaquatic volcanic activity.

The mixolimnion as its own entity behaves similarly to other tropical lakes; a thermal stratification develops during the rainy season and complete mixing occurs during the dry season. A stable thermal stratification persists in the mixolimnion of Lake Kivu from September to June that separates it into an epilimnion where most of the primary production occurs, and a hypolimnion that becomes increasingly anoxic. The thermal stratification is obliterated in the dry season, which typically begins in June and ends in August. During the dry season the surface waters are cooled down by a low relative humidity, an increase in evaporation, strong winds, and a lower intensity of downward long-wave radiation as a result of reduced

cloud coverage (Schmid and Wüest 2012). These factors culminate in convective mixing down to a maximum depth of 65 m. The strong, permanent thermohaline stratification at the top of the monimolimnion prevents deeper seasonal mixing. The stratification in the deepwater of e.g. Lakes Malawi and Tanganyika is much weaker. Vertical exchange of solutes by turbulent transport in these lakes is therefore more dominant compared to Lake Kivu (Verburg et al. 2003), where upwelling by advection is the dominant process of solute exchange.

Nutrients have been accumulating for thousands of years in the monimolimnion of Lake Kivu as a result of its persistent thermohaline stratification. Most of the settling organic matter (OM) from biological production in the mixolimnion is mineralized at the sediment-water interface. Therefore, nutrients continue to accumulate in the anoxic water, and are only supplied to the mixolimnion by the upward advection of the water column and the downward seasonal mixing. The contribution of dissolved inorganic phosphorous (DIP) and NH_4^+ from the monimolimnion to the mixolimnion in Lake Kivu presently accounts for 80 and 85 %, respectively, of the total nutrient inputs used for biological production (Pasche et al. 2009). The high C:P and N:P ratios measured from the seston in the mixolimnion indicate that the supply of DIP is limiting primary production (Sarmiento et al. 2006), while the C:N ratios also indicate moderate N limitation (Sarmiento et al. 2012). Primary production is dominantly cyanobacteria in the rainy season, and pennate diatoms and cryptophytes in the dry season (Sarmiento et al. 2012). Seasonal and interannual variations in biological production, including phytoplankton growth, are dependent on changes to the seasonal mixing depth. This is because the SGD is near steady-state, and therefore upwelling yields a constant rate of supply of nutrients to the bottom of the mixolimnion. High productivity occurs in the dry season when deeper mixing supplies nutrients from the monimolimnion. A low productivity persists throughout the rainy season with a stable thermal stratification in the mixolimnion and therefore a reduced nutrient supply (Darchambeau et al. 2014).

Exceptional mixing events will increase the supply of nutrients from the monimolimnion and fuel a higher productivity. These changes in nutrient dynamics are recorded in the form of anomalous sediment layers

that consequently yield information on past climate conditions that have altered the seasonal mixing depth, or past lake eruptions and subaquatic volcanic activity that might have eliminated the thermohaline stratification and entrained nutrients from below. The present study investigates the sediment stratigraphy, lithology, and geochemistry in the Main Basin in order to disentangle changes in nutrient dynamics as a consequence of climate change, from past lake eruptions and subaquatic volcanic activity.

Study site

Lake Kivu is situated at an elevation of 1463 m above sea level within the western branch of the East African Rift system. It has a 1:2 area, ratio of lake-water (2385 km²) to watershed (4940 km²). The lake volume is 549 km³, and a total of 300 km³ at standard temperature and pressure of CO₂, and 60 km³ at standard temperature and pressure of CH₄, are dissolved in the monimolimnion (Schmid et al. 2005). The presence of subaquatic volcanic features, where active hydrothermal springs are observed, has been outlined by a previous study (Ross et al. 2014), and their location is delineated in Fig. 1. In the Holocene, a large lake-level rise followed the increased volcanic activity of the Nyiragongo chain to the north and the onset of wetter climatic conditions (Ross et al. 2014).

The hydroclimate in tropical East Africa varies considerably on all time-scales (Russell and Johnson 2005, 2007). Zonal shifts in the Intertropical Convergence Zone are often accountable for the observed spatial and interannual variability of rainfall. When considering longer time-scales, variations in the sea surface temperature gradient across the Indian Ocean are suggested to be the principal controller of multi-decadal variability (Tierney et al. 2013). Recent hydroclimate data indicates drought-like conditions within the last two decades in this region that is also recorded in the Kivu Basin, albeit with considerable interannual fluctuations (Muvundja et al. 2014). Major drought events of decadal, centennial, and millennial-scale duration have been recorded in the lacustrine records in Lake Edward, Tanganyika, Malawi and Victoria (Tierney et al. 2010; Stager et al. 2005, 2009; Brown and Johnson 2005; Powers

Fig. 1 Overview map of Lake Kivu. The *inset* is the location of the Kivu Basin within Africa, and reveals its proximity to other lakes within the Western Rift. Gray scale bathymetric map is overlain by contour lines with a 50 m vertical distance. Cores used for analysis are marked directly onto the map

et al. 2011; Johnson et al. 2002; Russell and Johnson 2005, 2007).

Materials and methods

Geophysical surveys

Seismic stratigraphic analysis of the shallow subsurface sediment stratigraphy down to a depth of ~ 10 m was achieved using 3.5 kHz pinger single-channel data acquired in October, 2010. The system was mounted to the boat RV Gloria and the survey was conducted at an average speed of 2 m s^{-1} using a GPS navigation system. Processing included a flat gain and a bandpass filter (1600–6000 Hz). Seismic data

interpretation was performed with the SMT Kingdom Suite software.

Sediment parameters

The locations of gravity cores given in Fig. 1 were chosen based on the seismic survey results, and the geomorphology presented in Ross et al. (2014). As a consequence of the high gas concentration in the monimolimnion leading to degassing upon pressure relief, complete undisturbed cores were only retrieved above 200 m water depth. The top sections of cores KV10-20 and -21 retrieved from 459 and 458 m depth, respectively, were lost from gas bubbling. The cores were scanned for petrophysical properties with a GEOTEKTM multisensor core logger (MSCL)

calibrated with an aluminium-water standard. The cores were then opened into half-sections and immediately photographed. The core-photographs were overexposed for easy visualization of the colour variations. Lithological successions were macroscopically described and smear-slide analysis were conducted on each section using a petrographic microscope with alternating plane and cross-polarized light.

The intensities of Ca, Fe, S, Mn, Ti, and Sr were determined on half-cores using an X-ray fluorescence (XRF) core scanner (ITRAX, Cox Ltd., Sweden) with a Mo-tube at 30 kV and 30 mA for cores KV10-04 and -21. Cores KV10-19 and -20 were scanned with an Avaatech XRF core scanner using a measurement duration of 20 s at 10 kV and a 2000 μ A current for light elements, whereas 30 kV and 1000 μ A were applied for the heavier elements. The scanning was conducted with an integration time of 10 s per measurement at a 2-mm resolution for KV10-04 and -21, and at 5-mm for cores KV10-19 and -20, and the results are reported as counts.

Cores KV10-04 and -21 were sampled at 1-cm intervals, and samples were taken from representative lithological sections in each core including KV10-19 and -20. The samples were then subsequently frozen, lyophilized, and ground until homogeneous. Total carbon (TC), total nitrogen (TN) and the stable nitrogen isotope composition (^{15}N) of the homogenized samples were analyzed with an elemental analyzer (EuroVector, Euro EA) linked by continuous flow to an isotope-ratio mass spectrometer (Micro-mass, IsoPrime). The ^{15}N values were calibrated using international reference materials (IAEA-N1) with an assigned $\delta^{15}\text{N}$ value of 4‰ (versus AIR). The analytical reproducibility for ^{15}N measurements was ± 0.37 ‰. ^{15}N is reported as (δ) in units ‰ according to Eq. 1:

$$\delta^{15}\text{N} = \frac{R_{\text{sample}} - R_{\text{AIR}}}{R_{\text{AIR}}} 1000 \text{‰} \quad (1)$$

Total inorganic carbon (TIC) was analyzed as CO_2 on a CM 5012 Coulometer (UIC) following acidification with HCl (3 M). The precision for TIC was $\pm 0.4\%$. Total organic carbon (TOC) was calculated from the difference of TC and TIC. Biogenic silica (BSi) was measured according to the methods described by Pasche et al. (2010) on two separate cores taken at the

same location, where the cores were aligned based on their TIC profiles, and the duplicate core was sampled onsite.

Chronology

Cores KV10-04, -19, -20, and -21 taken from the Main Basin were sampled for terrestrial macrofossils in order to determine an age model for the system. The cores were aligned based on their lithology and patterns in XRF-derived Ca and Zr counts. The depth down to 9.5 cm in KV10-04 was dated using ^{210}Pb and ^{226}Ra concentrations provided by Pasche et al. (2010) from Kivu-06 by applying the Constant Initial Concentration (CIC) model (Appleby and Oldfield 1978), and assuming the same depositional environment leading to similar sedimentation rates between the cores. The bottom 58 cm of KV10-04, and the portion of cores KV10-19, -20, and -21 that were not disturbed by gas, were dated using ^{14}C ages of terrestrial organic material calibrated with the OxCal 4.2 program (Ramsey 2009). The metadata for radiocarbon analyses are found in ESM1.

Results

Sedimentology

Core KV10-04 was used in this study to describe the key sedimentary successions. Its location of origin and alignment to cores Kivu-06, -07 (Fig. 2), and core #10 (Fig. 4) from preceding studies, make it an ideal core for analysing paleo-environmental changes. The sediment from KV10-04 is composed mainly of finely laminated carbonate- and organic-rich mud. The mm-sized laminations consist of aragonite and diatoms, or OM and diatom frustules, in lithofacies A and B, respectively (Fig. 4). These laminated sections are punctuated with the visible lithofacies; C, D, E, and F (Fig. 3). In addition to their different colours, these layers are also recognized by geochemical changes in Fig. 3, and the lithological changes observed in the smear slides in Fig. 4. They are characterized by a high diatom content for lithofacies C, a high OC content for lithofacies D, a mixture of diatoms and carbonates with traces of tephra for lithofacies E, and dominantly tephra material for lithofacies F.

Fig. 2 Two age models are hypothesized following analyses of dates from all sediment cores: Age model 1 applies the radiocarbon ages from KV10-04, -20, and -21, and the ^{210}Pb calculated age, Age model 2 applies the latter ^{210}Pb age and three radiocarbon ages at 22 and 24 cm depth from KV10-04, and the age for the tephra layer at 49.5 cm depth taken from the sediment-mixed tephra in KV10-19 at 58 cm depth. *Dagger symbol* (†) is the age calculated from unsupported Pb, and all

other dates are in cal yrBP calculated using OxCal 4.2 (Ramsey 2009). *Double dagger symbol* (‡) is taken from the corresponding depth in Kivu-07 (Pasche et al. 2010). The *red dates* are three anomalously old dates >10 kyr BP obtained in cores KV10-04, KV10-21 and Kivu-07, which were assumed to be reworked terrestrial plant macrofossils and therefore not considered in the calculation of the age models

KV10-04 was taken on a ridge at 187 m water depth southeast of the area of subaquatic volcanic cones (Fig. 1). Based on its location, this core should be exempt from mass-movement deposits and sediment mobilization from a gas eruption in the Main Basin. The lithology in core KV10-04 is directly correlated to the lithology in cores Kivu-06 and -07 taken by Pasche et al. (2010), and core #10 taken by Hecky and Degens (1973) (Fig. 1). Core KV10-21 is correlated to core KV10-04 by the layers C1 and D1 (Fig. 3), observed within the gas-disturbed portion of this core (Fig. 2). The sediment directly below this depth is not

correlatable to core KV10-04 due to a mass-movement deposit that is observed in Fig. 2. Cores KV10-20 and -21 can be correlated to core KV10-19 taken near the location of subaquatic volcanic features 15 km to the northwest (Fig. 1). The three cores each contain a mixed-sediment layer that consists of diatoms, carbonates, and traces of tephra, which are labeled ‘tephra layer’ in Fig. 2. The prominent tephra layer F1 is used to align KV10-04 with core #10 taken by Hecky and Degens (1973) in 1972 (Fig. 4).

The sedimentation rate in the top 9.5 cm of core KV10-04 is high relative to the lower core-section as a

Fig. 3 Geochemical data from KV10-04. Geochemical parameters TIC, TOC, TN, and BSi in % weight. Isotope data for ^{15}N . XRF data for Fe, S, Mn, Ti, and Sr in counts. Geochemical

proxy data for SGD versus allochthonous input (Fe:Ti), abrupt changes to seasonal mixing depth (Mn:Fe), and authigenic pyrite formation (Fe:S). The lithofacies legend is given in Fig. 4

direct effect of carbonate precipitation (Pasche et al. 2010). A linear sedimentation rate (LSR) of $2.75 \text{ mm year}^{-1}$ yields a calendar age of 1976 at 9.5 cm core depth (Fig. 2). This age was calculated to be at least 15 years younger than in previous age models that did not always include ^{226}Ra concentrations (Pasche et al. 2010). Age model 1 applies the 1976 age, in addition to four radiocarbon calibrated dates, to produce an age versus depth LSR of $0.36 \text{ mm year}^{-1}$. The 1976 age and the three radiocarbon dates used in Age model 2 yield a LSR of 0.5 mm year^{-1} (Fig. 2). Age model 2 applies the radiocarbon age obtained within the tephra layer of core KV10-19. The younger age here implies that the calibrated ages from KV10-20 and -21 at 58.5 and 43 cm depth, respectively, were reworked to yield older ages. This scenario seems likely since the macrofossil samples were taken from what appears to be a distal

mass-movement deposit, with visible coarse-grained mica flakes and shell fragments (Fig. 2). The rates presented in Fig. 2 are comparable to the ^{14}C derived LSR of 0.3 mm year^{-1} provided by Hecky and Degens (1973) for the overlapping section of their core #10 (Fig. 3).

Geochemistry

XRF data

The XRF counts for Ca, Fe, S, Mn, Ti, and Sr in Fig. 3 represent most of the variation observed in the data of core KV10-04. Ca intensities correlate with TIC concentrations ($r = 0.86$), implying that Ca in the sediment is mostly bound to carbonates. Carbonate precipitation and preservation in Lake Kivu is controlled by (1) aragonite oversaturation in the

Fig. 4 Core #10 aligned with KV10-04 by layer R3/F1. The *colours* in core #10 are from the descriptions given in Hecky and Degens (1973), Degens et al. (1973), Stoffers and Hecky (1978), Degens and Kulbicki (1973), and Haberyan and Hecky (1987) that correspond to the lithology in KV10-04. *Lithological colours and letters* are represented by their corresponding smear slides with a *scale bar* of 50 μm in each image

mixolimnion and (2) Ca^{2+} and CO_3^{2-} concentrations in the water column, which are further controlled by the pH and alkalinity (Ross et al. 2015). A similar trend to Ca is observed for Sr, which is commonly exchanged for Ca in aragonite and therefore also primarily controlled by carbonate precipitation and dissolution. Sr in Lake Kivu is enriched in the hydrothermal SGD within the monimolimnion (Ross et al. 2015), and therefore large peaks in Sr could indicate increased hydrothermal discharge; e.g. the onset of layer A1 in Fig. 3.

The dissolved Fe, S, and Mn flux into Lake Kivu is primarily controlled by SGD (Ross et al. 2015). Fe correlates with S in KV10-04 ($r = 0.85$) outside of the tephra layer F1, which suggests that Fe in Lake Kivu sediments is primarily present in the form of authigenic Fe-sulfides. Mn does not correlate with S to the same degree as Fe ($r = 0.6$), likely due to the higher solubility of Mn^{2+} compared to Fe^{2+} in the reducing waters of the monimolimnion (Ross et al. 2015). There is an overall increase in Fe and S relative to Mn beginning at layer D2, which lowers the general Mn:Fe ratio above this

layer (Fig. 3). A peak in Mn^{2+} is observed just below the oxycline in the water column that is formed by the partial redissolution of MnOOH and MnCO_3 precipitates from mixing in the dry season (Pasche et al. 2009). Under these concurrent conditions large peaks in Mn:Fe in KV10-04 could represent peaks in MnOOH and MnCO_3 precipitates, which are positively influenced by the depth of mixing in the dry season.

Ti is not susceptible to biogeochemical alterations in the water column or during diagenesis, and represents the influx of terrestrial and/or volcanic material into the lake (Vogel et al. 2013). The large peaks in Ti in KV10-04 are attributed to tephra layers, e.g. layer F1 and E1 in Fig. 3. Elevated Fe relative to Ti corresponds to lower inputs of allochthonous material or higher inputs of Fe from SGD (Ross et al. 2015). Therefore elevated Fe:Ti ratios, including that observed in layer B3 in Fig. 2, indicate the importance of authigenic Fe precipitation in Lake Kivu at certain periods. K revealed a nearly identical trend to Ti ($r = 0.98$; data not shown), which is likely an effect of the K-rich volcanics in the VVP (Smets et al. 2010).

TOC/TIC/TN/BSi

The aragonite-OM laminations in Fig. 3 are observed between 0 and 10 cm, and again between 50 and

67 cm depth in lithofacies A of KV10-04. The dilution of TOC by aragonite precipitation (Pasche et al. 2010), corresponds to the negative correlation in Fig. 5i ($r = -0.8$). A high %TOC found within lithofacies D observed in Fig. 3 indicate a high OC production in the mixolimnion at the time of their deposition. These layers contain on average 1.5x more OC content relative to the composition in lithofacies B.

Figure 5ii in core KV10-04 ($r = 0.93$) yields an average C:N ratio of 15 (σ 2.4), suggesting OM originates primarily from autochthonous biological production, which is also supported by the small catchment to lake ratio and the trophic status of the lake (Sarmento et al. 2012). Comparatively, the D lithofacies have a C:N ratio of 12 (σ 0.8) that could indicate decreased N-limitation in these layers. Troughs in the C:N ratio are observed at the depths of D1, D2 and D3 in Fig. 3.

The BSi measurements in KV10-04 reconstruct the phytoplankton abundance that is enriched in diatoms in the dry season. Nutrient fluxes into the mixolimnion from seasonal mixing is the most important parameter controlling diatom abundance (Sarmento et al. 2012). A large peak in BSi observed in layer C1 of Fig. 3 implies an increased mixing depth at this time.

Fig. 5 Scatter plots of TIC, TOC, TN and ^{15}N data. The data are coloured according to their lithofacies origin

Isotopes

The ^{15}N does not correlate with the fluctuations observed in C:N in Fig. 5iii ($r = -0.02$). This indicates that the observed ^{15}N ratios in Fig. 3 do not represent changes in allochthonous versus autochthonous OM. Therefore, the variations in ^{15}N are likely caused by lake-internal processes. Within lakes, denitrification and ammonification become increasingly dominant processes with an increasing N-nutrient supply. Ultimately, the preferential elimination of isotopically light ^{14}N by these metabolic pathways enriches ^{15}N in the remaining DIN available for primary production (Struck et al. 2001). Alternatively, nitrogen fixation by cyanobacteria ($\sim 0\%$) during periods of reduced N nutrients will lower the ^{15}N ratio of the OM (Olsen et al. 2013). These metabolic effects of ^{15}N enrichment are underlined in KV10-04 by the positive correlation observed in Fig. 5iv ($r = 0.7$), where deviations from the trend are the highest within the D lithofacies. Within the D lithofacies, the ^{15}N begins to increase near the top of the deposited layer at depths where the %TOC begins to decrease (Fig. 3).

Seismic stratigraphy

The penetration, and therefore the achievable seismic stratigraphy, at the location of KV10-04 was poor

Fig. 6 Seismic profiles S1 and S2 in Fig. 1. The vertical and horizontal scales are located in the lower corner of each frame. Annotations are explained within the text

owing to the gas content and steep slopes of the ridge observed here. However, the stratigraphy along seismic lines S1 and S2 in Fig. 6 provides a lateral sedimentologic dimension to the sediment cores KV10-20, 21, and core #10 (Fig. 1). As a consequence of the similar depth and nearby location of core #10 to our seismic profiles (Fig. 1), we can extrapolate the lithofacies observed in this core in Fig. 4 to the seismic stratigraphy in Fig. 6.

The prominent reflections R1–R4 traced in the seismic data in Fig. 6 refer to the lithology of core #10 that are also marked R1–R4 in Fig. 4. The reflection R4 in Fig. 6 is the sediment-water interface represented by the top of the core KV10-04. The tephra layer F1 in KV10-04 aligns to the tephra layer in KV10-20 and -21, and R3 in core #10. The seismic-unit between reflections R4 and R3 is semi-transparent, and its lateral extension changes thickness and evens out the morphology in the deep basin (Fig. 6).

The high-amplitude reflection R2 in Fig. 6 corresponds to tephra layer R2 observed in core #10 in Fig. 4. The tephra layers R3 and R2 were deposited at ~ 1 and 5 ka, respectively, according to Hecky and Degens (1973). Either a gas-eruption or lake-mixing event is postulated to have occurred at the time of their deposition (Haberyan and Hecky 1987). Both tephra layers mark the abrupt termination of laminated aragonite-OM alternations (lithofacies A in Fig. 4). Furthermore, these layers coincide with the bottom of a transparent seismic facies, with varying thickness, observed in Fig. 6.

The reflection R1 in Fig. 6 corresponds to the tephra layer R1 that forms the base of core #10 in Fig. 4. This tephra layer was postulated to have been deposited during the increased volcanic activity in the Kivu Basin, which dammed the lake to the north ~ 10 ka (Ross et al. 2014).

Discussion

Significance of aragonite/OM laminations

The mixolimnion in Lake Kivu is oversaturated with respect to CO_2 (Borges et al. 2012), and mixing in the dry season results in aragonite precipitation from CO_2 degassing and a concomitant increase in pH. This is manifested in the sediment-trap data collected by Pasche et al. (2010), where the highest amount of

carbonate deposition is observed in June at the onset of the dry season. Within the sediment, there have been three shifts within the last ~ 10 ka from high OC to aragonite/OM laminations (from lithofacies B to A; Fig. 4). The most recent onset of carbonate preservation in the sediment was initiated in 1976 at 9.5 cm core depth in KV10-04 (A1; Fig. 3), and persists today. Layer A1 in conjunction with the additional carbonate layers observed below this depth in KV10-04 and core #10 beginning at 108 and 280 cm depth (Fig. 4) does not support the conclusions that carbonate preservation is a consequence of climate aridity causing the lake to mix, or of a monimolimnion that preexisted corrosive hydrothermal activity (Degens et al. 1973; Hecky and Degens 1973), or of anthropogenic changes to the catchment (Pasche et al. 2010). Although the current drought-like conditions in the Kivu Basin may be a factor influencing carbonate preservation, the observed stratification in the lake indicates that preservation is not a consequence of lake mixing. Additionally, the present-day monimolimnion is corrosive from ongoing hydrothermal activity, and the observations of prehistoric carbonate preservation cycling eliminate the anthropogenic-hypothesis.

We posit that carbonate preservation in the sediment is the product of an increase in alkalinity and CO_2 in the monimolimnion from hydrothermal SGD culminating in a high dissolved inorganic carbon (DIC), and therefore a CO_3^{2-} concentration that eventually favours aragonite preservation (Ross et al. 2015). Since the onset of measurements conducted by Damas (1937), a ~ 5 -mM increase in alkalinity below 300 m depth has been observed. Such an increase in alkalinity alongside reducing reactions that occur in the monimolimnion, e.g. SO_4^{2-} reduction to H_2S , will shift the DIC equilibrium to the saturation threshold of CaCO_3 over time. This shift could hypothetically produce the sudden onset of carbonate preservation that we observe at lithofacies A2 and A1 in Fig. 3, if the monimolimnion is near the equivalence point of either HCO_3^- or CO_3^{2-} . The idea that the CaCO_3 saturation varies with depth and time in Lake Kivu is synonymous to the changes in the lysocline and the aragonite compensation depth in the deep ocean (Zeebe and Westbroek 2003). The present lysocline in Lake Kivu appears within the first density gradient beneath the mixolimnion (Pasche et al. 2009).

Intense hydrothermal activity or subaquatic volcanism will increase the flux of CO_2 and other acidic gases such as SO_2 into the lake, and increase the thermal energy in the monimolimnion (Carey et al. 2013). As a consequence CO_2 could become saturated and be expelled from the monimolimnion, thereby decreasing the total DIC in the lake and lowering the overall alkalinity as an effect of complete mixing. These gases, in addition to oxidation reactions that are likely associated with a lake eruption, will lower the pH with the net effect being carbonate dissolution. This scenario could explain the abrupt termination of the aragonite laminae at the tephra layers R2 and R3/F1 (Figs. 3, 4). The abrupt termination of the carbonate layer corresponding to a tephra layer suggests the interplay between subaquatic volcanic activity and carbonate preservation in the sediment. Following the partial to complete obliteration of the stratification in Lake Kivu that might have occurred at these tephra layers, the ongoing hydrothermal SGD into the monimolimnion, and the cold SGD above the main pycnocline at 260 m depth, would reinitiate the stratification in the lake almost immediately (Ross et al. 2015; Schmid et al. 2012). The subsequent phases of carbonate preservation would be delayed compared to the increased SGD input, until the carbonate concentration in the water column reaches its saturation threshold. The preservation of carbonates in lithofacies A2 may have required more hydrothermal contribution than that resulting in A1, as a consequence of the already high alkalinity in the lake following a mixing event.

Origin of event layers

The lithofacies C, D, E, and F in KV10-04 represent geochemical changes in the mixolimnion at the time of their deposition (Fig. 3). We assume that the rate of sediment deposition in KV10-04 likely changed within an order of magnitude at the depth of these lithofacies. These layers are also observed by Pasche et al. (2010) in cores Kivu-06 and Kivu-07 taken from the Main Basin (Fig. 1). We suggest that punctuations in high productivity culminate in the C and D lithofacies, which are an effect of top-down versus bottom-up mixing events, respectively.

Layer C1 at 13 cm core depth in Fig. 3 is characterized by a high diatom content and coincides

with a peak in BSi and Mn:Fe. This is indicative of a diatom bloom due to increased mixing-depth in the dry season resulting in increased production, but would not have reached a depth more significant than the peak in Mn^{2+} observed just below the oxycline in the water column. The high Mn:Fe suggests oxygen diffusion and precipitation of MnO minerals, indicating top-down mixing. This contrasts with D layers, which lack clear indications of metal oxides. The D lithofacies are the result of enhanced OM production and contain up to 31 % TOC (ESM2). The present-day OM preservation in Lake Kivu is 10–20 % with most of the degradation occurring during early diagenesis (Pasche et al. 2010). We postulate that the D lithofacies in KV10-04 were produced by single events that resulted in a high flux of nutrients into the mixolimnion from below, consequently yielding a high productivity and an increase in OM flux and preservation. Such an event could be caused by a buoyant hydrothermal plume (Woods 2010), which would entrain nutrients from below and could penetrate the mixolimnion.

Since the residence time of the nutrients in the monimolimnion of the lake is on the order of thousands of years (Schmid et al. 2005), the isotopic composition of the nutrients supplied to the surface layer is likely constant over shorter time scales. Therefore changes to the internal nutrient recycling, such as the preferential assimilation of isotopically light ^{14}N during phytoplankton growth, will result in the greatest fractionation between the DIN and the OM when nitrogen is not limiting (Olsen et al. 2012). We propose that the observed peaks in ^{15}N ratios at depths corresponding to the top of the D lithofacies in KV10-04 (Fig. 3), Kivu-06, and Kivu-07 reflect a strong increase of N-nutrient input into the mixolimnion. Increased OM production is concomitant with increased denitrification and ammonification, which could continuously deplete the nutrient pool in the mixolimnion while enriching it in ^{15}N . This results in a delayed peak in ^{15}N in Fig. 3 relative to the peak in TN within the D lithofacies. The peaks in ^{15}N observed in layers D1, D2, and D3 are enriched by a factor (ϵ) of 1.7, 1.5, and 2.4, respectively (ESM2). We hypothesize that the D layers are formed as a result of single events, which are of volcanic origin based on that layers D3 and D1 in KV10-04 overlay tephra deposits F1, and E1, respectively. Significant volcanic

events would have resulted in oversaturation of the gases in the lake and degassing to the atmosphere as suggested by our biogeochemical data. An induced oversaturation of gases is consequent of the high DIC concentrations that are inferred during periods of carbonate preservation discerned from the presence of the A lithofacies. These eruptions result in an abrupt change in the geochemical depositional regime initiating the B lithofacies (Fig. 4).

To further investigate the hypothesis that a bottom-up mixing event occurred at the D lithofacies, rather than a top-down mixing event, we compared the regional climate-proxy records that have revealed periods of intense aridity or temperature cooling in Lake Edward (Russell and Johnson 2005, 2007), Tanganyika (Tierney et al. 2010), and Malawi (Johnson et al. 2002) with the record observed in Lake Kivu in Fig. 3 and age models in Fig. 2. We do not observe any correlation in these records to the D lithofacies observed in KV10-04, however, the temporal resolution may not be refined enough to define a mixing-event in Lake Kivu. Furthermore, we calculated the mixing depth, and corresponding temperatures and heat dissipation required to achieve this mixing depth in Lake Kivu and Lake Tanganyika, in order to entrain sufficient nutrients from below to produce the D lithofacies observed (ESM3). These calculations resulted in heat dissipation and lake surface temperature constraints that would completely overturn Lake Tanganyika, where the density stratification is much weaker (Verburg et al. 2003). Such an indication of mixing has not been observed in the relatively refined sediment record of Lake Tanganyika, however this possibility has not been investigated in detail. Additional wind-induced mixing models also indicated the necessity of continuous, strong winds of $>15 \text{ m s}^{-1}$ for an extended period of time in order to mix to a depth below 80 meters in Lake Kivu (Aaberg 2013).

Corresponding D lithofacies were originally observed by Hecky and Degens (1973) in core #10 (Fig. 4), and considered to be indicators of hydrothermal discharge. Hecky and Degens (1973) predicted that these layers were composed almost entirely of algal biomass, which was later confirmed by Al-Mutlaq et al. (2008). Degens et al. (1973) describe the D lithofacies as synonymous to sapropels observed in the anoxic ocean (Calvert et al. 1992). They conclude that an increased frequency of the

D lithofacies is associated with wetter climatic conditions and hydrothermal activity, which resulted in the thermohaline stratification present today. Here we confirm that the D lithofacies have a high OM content from autochthonous origin, are low in diatoms, and enriched in ^{15}N (Fig. 3). We also suggest that these layers are analogous to the sapropels observed in the anoxic ocean, where the enriched ^{15}N and high OC content are a result of denitrification (Struck et al. 2001). Furthermore, we hypothesize that these layers are a consequence of episodic intense hydrothermal activity.

Paleoenvironmental reconstruction

Given the cumulative data collected from the cores taken below the mixolimnion (Fig. 1), the past sedimentological history of Lake Kivu since the lake-level rise in the late Pleistocene can be reconstructed. Core #10 terminated in tephra layer R1 hypothesized to be deposited during the lake-level rise ~ 10 ka. The age model constructed by Degens et al. (1973) yielded a sedimentation rate of 0.3 mm year^{-1} for this core, where this rate was calculated from only one sample younger than 10 cal kyr BP (6200 cal year BP; Fig. 6). Furthermore, the radiocarbon analysis was conducted on bulk-OM that could result in anomalously old ages from the reservoir effect of old carbon. From Age model 1 and 2 (Fig. 2) an age of ~ 9.3 – 6.6 ka for layer R1 is calculated. This age range correlates to the first evidence of overflow via the Ruzizi River into Lake Tanganyika at ~ 8 ka (Felton et al. 2007).

The high preservation of carbonate beginning in core #10 at 280 cm (Fig. 4) is dated to 7.6–5.4 ka, and was abruptly terminated at tephra layer R2, at 4.2–3 ka (160 cm; Fig. 4). This period of carbonate preservation corresponds to the increase in the Ca/Al observed by Felton et al. (2007) in the Lake Tanganyika sediment record at $< \sim 4$ ka. We suggest that this carbonate was preserved in the sediment as a consequence of increasing DIC in the monimolimnion, and not due to climatic changes that might have led to a lake-level low stand and a closed basin, as suggested by Haberyan and Hecky (1987) and Hecky and Degens (1973). However, there were periods of intense aridity that are recorded in other lakes in East Africa during the mid-Holocene, including Lake Edward from

5400–2000 ka (Russell et al. 2003), Lake Turkana at 5270 cal year BP (Garcin et al. 2012), and Lake Chala from 5.9 to 4.7 cal kyr BP and again from 3.6 to 3.0 cal kyr BP (Verschuren et al. 2009). In general, the time period after ~ 5 ka was drier than the early Holocene in East Africa, termed the ‘African Humid Period’. The 2nd carbonate layer was initiated at 110 cm corresponding to 2.8–2 ka (A2; Fig. 4). This date coincides with a peak in aridity documented in Lake Edward at 2 cal kyr BP (Russell et al. 2003). However, layer A2 spans the onset of a wetter climate beginning ~ 1.7 ka documented in both Lake Edward and Tanganyika (Stager et al. 2009). This carbonate layer was also abruptly terminated, at 1.2 ka (Fig. 2, Age model 1) or at no later than 765 cal year BP (Fig. 2, Age model 2). Finally, the onset of the carbonate layer that is currently preserved in the sediments of the monimolimnion (A1; Fig. 3) was not initiated until 1976 AD, and corresponds to a large peak in Sr and Zr (Figs. 2, 3, respectively). This change in depositional regime might indicate a controlling effect of hydrothermal activity corresponding to the 1977 Nyiragongo eruption (Tazieff 1977).

The D lithofacies observed in core #10 (above layer R2; Fig. 4) and KV10-04 (D3 and D1 above layer R3/F1 and E1, respectively; Fig. 4) overlay tephra layers, and could therefore correspond to large inputs of nutrients into the mixolimnion as a consequence of subaquatic volcanic activity in the monimolimnion. Furthermore, these subaquatic volcanic pulses likely resulted in a gas eruption, and subsequently a lake-mixing event, if they materialized with high DIC concentrations in the monimolimnion. The last mixing event occurred at 1.2 ka or no later than 765 cal yr BP where a large peak in Ti and Fe is observed in KV10-04 at 49.5 cm depth. Fe and Mn would be oxidized upon a lake-mixing event, and a concave peak in Mn:Fe observed at F1 is likely a consequence of the higher solubility of Mn at a low pH. The large influx of nutrients into the monimolimnion upon lake mixing results in the deposition of the high OC layer D3. The ensuing layer B4 ends abruptly with the high OC layer D2, which likely also corresponds to subaquatic volcanic activity. The high authigenic Fe recorded in the following layer B3 and B2 could be due to wetter climatic conditions that would result in increased SGD into the monimolimnion (Pasche et al. 2009). Layers B3 and B2 are separated by layer D1 at ~ 400 cal yr BP (Fig. 2), which marks the last indication of intense

hydrothermal activity. This date corresponds to an increase in Ti into the lake up until the present. A Ti increase was also recorded in core Kivu-06 by Pasche et al. (2010) and attributed to changes in the catchment including deforestation and heightened population. This explanation seems likely given that the most significant input of Ti recorded outside of the tephra layers in KV10-04 has been within the last 30 years (down to 8 cm depth; Fig. 3).

Two anomalously old radiocarbon samples dated 10 and 11 cal kyr BP were acquired from layer B4 in core KV10-04 and Kivu-07 (Fig. 1). The correlating B4-lithofacies in Kivu-06 was determined to have an unusually low abundance of plankton (Pasche et al. 2010). Additionally, a third radiocarbon sample with a date of 11 cal kyr BP was acquired from the mass-movement deposit in KV10-21. The appearance of three samples dated ~ 10 cal kyr BP (Fig. 2) is attributed to the high influx of terrestrial material with the lake-level rise in the late Holocene. However, massive sediment mobilization is the only reasonable explanation for the presence of terrestrial macrofossils that are > 10 cal kyr BP at less than 50 cm depth in three cores from three locations. We suggest that these anomalously old dates are from terrestrial macrofossils that remain buoyant in the water column following sediment mobilization concomitant with seismic events or a lake eruption. Such reoccurring events could account for the observation of these macrofossils in the upper layers of the sediment cores.

The sediment-mixed tephra layer observed in KV10-20, and -21 correlates to the reflection R3 in the seismic profile in Fig. 6, across the basin to KV10-19 (52 cm core depth; Fig. 2), and to the tephra layer F1 in KV10-04 (Fig. 3). The ensuing transparent seismic facies observed in Fig. 6 above this layer is postulated to be mobilized sediment as an effect of either seismic events corresponding to basin volcanic activity, or a gas eruption from the deepwater. The resulting basin-wide transparent seismic facies is mapped in ESM4. A total of two transparent seismic facies are observed in Fig. 6 overlying what appears to correspond to layers R3 and R2 in Fig. 4. The variation in thickness observed within this facies in ESM4 could be formed by a gravitational sediment focusing of mass movements upon a lake-wide event. This phenomenon was also observed in Lake Nyos by Freeth and Rex (2000) following its eruption as an effect of gas oversaturation in 1986.

Conclusions

From the analysis of the sediment stratigraphy, lithology, and geochemistry of sediment cores taken in the Main Basin (Fig. 1), we are able to deduce the major environmental changes that have occurred since the onset of increased volcanic activity and lake-level rise at ~ 10 ka. From these results, we hypothesize that the evolution from OC to aragonite/OM-laminae in the sediment, which is then abruptly terminated at a tephra layer, is coeval with SGD and subaquatic volcanic activity within the monimolimnion. This sequence of events has occurred twice following ~ 10 ka, presumably due to gas eruptions as a consequence of volcanic activity at ~ 5 and at ~ 1 ka. Furthermore, pulses in subaquatic volcanism that resulted in buoyant hydrothermal plumes have occurred no later than 400 cal year BP. The present-day carbonate saturation in the monimolimnion since the 1970s implies that there has been a recent increase in hydrothermal activity. Given that indications of a gas eruption from the lake have only occurred at periods of carbonate saturation, these results highlight the importance of monitoring subaquatic volcanic activity in Lake Kivu.

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