- 1 Controls on the Provenance of late Eocene to Quaternary Mozambique Channel
- **2** shales (DSDP **25** Site **242**)
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13 Abstract

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Provenance determination of late Eocene to Quaternary sediments deposited at the DSDP 25 Site 242 14 may help to evaluate the mechanisms that have controlled the sediment deposition in the Mozambique 15 Channel . To determine the provenance of the clay fraction, we measure major and trace element 16 concentrations as well as strontium (87Sr/86Sr) and neodymium isotopic compositions (expressed as 17 18 $\mathcal{E}_{Nd(0)}$) of thirty-six sedimentary samples from DSDP 25 Site 242 on the eastern flank of the Davie Fracture Zone. Light Rare Earth Element (LREE) enrichment and Heavy REE (HREE) depletions 19 against Post Archean Australian Shales (PAAS) associated with Ce anomalies higher than or equal to 20 0.90 for all the analyzed samples suggest very little influence of authigenic smectite incorporation. The 21 absence of correlation of Al/Si and CIA with REE content, Eu/Eu*, Cr/Th, Th/Sc, and 87Sr/86Sr and 22 $\mathcal{E}_{Nd(0)}$ suggests that chemical weathering has not modified these provenance proxies. Slight increase of 23 Eu/Eu*, Cr/Th ratios, $\mathcal{E}_{Nd(0)}$ values associated with slight decrease in 87 Sr/ 86 Sr and Th/Sc ratios from 24 25 the late Eocene to the Quaternary point to input of less differentiated and younger detritus to the DSDP

25 Site 242 with age. Based on variations in $\mathcal{E}_{Nd(0)}$ and Nd concentrations, we estimate an overall decrease of the Southeast African rivers (Congo/Zambezi) contribution from ~ 65.7% (Standard Deviation (S.D.) ~ 13.2) in the late Eocene to ~28.5% (S.D. ~ 10.0) in the early Pliocene. The causes for the low $\mathcal{E}_{Nd(0)}$ values and high Southeast African contribution between the late Eocene and the early Oligocene remain to be determined. It could record the onset of modern sedimentation in the Zambezi delta or the discharge of the Paleo Congo river in the Indian Ocean. The decrease of Zambezi contribution between the late Oligocene and early Pliocene is attributed to the cumulative effect of increasing tectonic activity of the Davie Fracture Zone and intensification of the Mozambique Current. The late Pliocene to Quaternary glacial-interglacial cycles promoted higher Zambezi contribution during glacial sea-level lowstands and in turn lower $\mathcal{E}_{Nd(0)}$ detrital values, whereas low Zambezi contributions are favored during interglacial highstand causing higher $\mathcal{E}_{Nd(0)}$ values. Hence, the modern dispersal pattern of fine-grained sediments within the Mozambique Channel is no younger than late Pliocene.

Introduction

Provenance changes of clastic sediments deposited in deep offshore ocean settings may record the denudation history of adjacent continents and consequently changes in paleorelief, drainage configuration, as well as sediment transport associated with large-scale tectonic processes, climatic change, and variation in paleo-oceanic current pathways. The provenance of deep-sea sediments has been used to infer climate-driven changes in drainage sources during glacial-interglacial cycles (Bonneau et al., 2017; van der Lubbe et al., 2016), to evaluate the effect of Quaternary glacial and interglacial cycle on oceanic current intensities and paths (Fagel et al., 1999; Franzese et al., 2006; Innocent et al., 1997; Revel et al., 1996), to evaluate the respective role of tectonics and climate in sediment supply at geological timescale (Fagel et al., 1994), to reconstruct the evolution of continental paleodrainage basins (Liu et al., 2017; Mourlot et al., 2018) or even to unravel details about past deep ocean circulation patterns during the Cretaceous (Mourlot et al., 2018). With a few exceptions (e.g. Mourlot et al., 2018), most studies dealing with the provenance of deep-sea sediments concern present-

day to Pleistocene in age sediments. To date there is no provenance study of pre-Pleistocene Mozambique Channel or basin sediments.

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The Cenozoic sedimentary infill of the Mozambique Channel and Basin is controlled by the variations in sediment supply, subsidence and oceanic near-bottom current intensities. Onshore, the southern part of the African plateau was tectonically-reactivated during late Eocene to early Miocene times (40-15 Ma; Baby et al., 2018), which may have resulted in an increase of Zambezi river detritus exported to the Indian Ocean during Oligocene times (34-24 Ma, Walford et al., 2005). Also, this might have cut off the hydrological connection of the Congo drainage basin with Indian Ocean in the Oligocene or Eocene (30-40 Ma; Stankiewicz and de Wit, 2006). From late Miocene and onward, drainage capture and doubling of the size of the Zambezi catchment during the Pliocene would have also considerably increased the contribution of Zambezi river sediments to the Indian Ocean (Walford et al., 2005). Meanwhile an offshore middle Miocene graben that was formed along the Davie Ridge (Droz and Mougenot, 1987) as a result of the southerly extension of the East African Rifting System since the Oligocene (Mahanjane, 2014) may have controlled the subsidence in the Mozambique Channel. These may have influenced the variation in intensities of the Mozambique Current and its effect on sediment deposition during the Cenozoic. One way to evaluate the influence of any of these processes is to determine the provenance of well-dated siliciclastic sediments deposited in the Mozambique Channel, at DSDP 25 Site 242. In detail, Nd isotopes may be particularly useful as they have been extensively used as tracer of deep-sea sediment source (e.g. Goldstein and O'Nions; 1981; Dia et al., 1992; Grousset et al., 1990, 1988; Mourlot et al., 2018; Revel et al., 1996a,b). Furthermore, present-day sediments deposited in the Mozambique Channel and basin have contrasting Nd isotopic compositions (Fig.1). Indeed, sediments entering the Mozambique Channel from the North are characterized by input of geologically-young material, which results in $\mathcal{E}_{Nd(0)}$ values close to -11.2 (van der Lubbe et al., 2016), whereas sediments deposited southward in the Mozambique Basin by the Mozambique Current are characterized by lower $\mathcal{E}_{Nd(0)}$ values (-15.5, Franzese et al., 2006) due to contributions from relatively older source rock areas by the Zambezi river (Fig. 1).

The goal of this study is to determine the provenance of siliciclastic sediments deposited in the Mozambique Channel, since the late Eocene based on the major and trace element concentrations and Sr-Nd isotopic data of thirty-six clay samples of clay-rich sediments from the DSDP 25 Site 242 (Fig. 1). Determining the provenance of these sediments will provide new constraints on the geographical source of the sediments deposited in the Mozambique Channel and how long-term geological processes (drainage reorganization, tectonics of the East African Rift) may have controlled the sediments deposited by the Mozambique Current.

Geological settings and potential sediment sources

2.1 Geological settings

The Deep Sea Drilling Project (DSDP) 25 Site 242 was drilled in 1972 and is located on the eastern flank of the Davie Ridge (Simpson et al., 1974) in the Mozambique Channel at 2275 m water depth. Site 242 is intermittently cored to 676 m depth providing late Eocene to Holocene sediments. The Mozambique Channel is a broad trough bounded to the West by the Mozambique continental slope and to the East by the Madagascar continental slope (Fig. 1). It hosts the inner part of the Mozambique deep-sea fan and associated feeding canyons, the west-east Zambezi Valley, and the north-south Serpa Pinto Valley (Droz and Mougenot, 1987; Kolla et al., 1980). The Mozambique Channel is structurally partitioned by the asymmetric N-S trending Davie Ridge (Fig. 1). The Davie Ridge often referred to as the Davie Fracture Zone (DFZ) is a shear zone that developed during the southerly movement of East Gondwana between the middle Jurassic drift and the early Cretaceous (Franke et al., 2015; Mahanjane, 2014; Mougenot et al., 1986). During its active phase, the DFZ experienced pure strike-slip movement (Mahanjane, 2014). Since the Oligocene, southerly extension of the DFZ has been related to the East African Rifting System (Deville et al., 2018; Franke et al., 2015; Mahanjane, 2014).

The inner Mozambique Fan (also termed upper fan) is characterized by two main feeding paths that have been successively dominant (Droz and Mougenot, 1987). From the Oligocene to early Miocene, the inner fan was supplied by north Mozambique terrigenous sediments through the north-

south Serpa Pinto Valley. In contrast since the middle Miocene, the inner fan is mainly fed by sediments transported by the Zambezi river through the west-east Zambezi Valley (Droz and Mougenot, 1987). This change in sedimentary source was a consequence of a graben formation along the Davie Ridge that may have led to the abandonment of the Serpa Pinto Valley (Droz and Mougenot, 1987). Estimates of the Zambezi sediment flux to the Indian Ocean indicate that there were two periods of elevated sedimentary flux in the Cenozoic: i) during Oligocene times (34–24 Ma) as a response of rapid regional uplift of Southern Africa and ii) from late Miocene times (~10 Ma) to present-day due to a doubling of the size of the Zambezi catchment during the Pliocene, respectively (Walford et al., 2005).

2.2. Modern and past oceanic circulation along the Mozambique Channel

The Mozambique Current (MC, Fig. 1) is the southern extent of the Northeast Madagascar Current (NEMC, Fig.1), which corresponds to the northern branch of the South Equatorial Current (SEC, Fig.1). The Mozambique Current is responsible for the strong, net southward flow along the Mozambique Channel (Ridderinkhof et al., 2010). The flow through the Mozambique Channel occurs through the passage of large anticyclonic eddies (Ridderinkhof et al., 2010). Northward flowing bottom currents transporting North Atlantic Deep Water and Antarctic Intermediate Water masses are found at depths below 2000 m (Collins et al., 2016). Annual mean velocities of the southward flow decrease progressively in the upper 1500 m of the water column within the Mozambique Channel (de Ruijter Wilhelmus. et al., 2002; Ullgren et al., 2012).

Based on the fact that the gemorphology of the Zambezi deep-sea fan evolved from lobated to elongated during the Miocene, Walford et al. (2005) proposed that the strong southward Mozambique Current was established in the early Miocene (~24 Ma). Before the early Miocene, there was no "strong" influence of a north-south current on sediment distribution (Walford et al., 2005). Based on detailed seismic stratigraphic analyses, a similar early Miocene onset age for the intensification of the Mozambique Current is proposed by Preu et al. (2011). In contrast, several late Cretaceous current – controlled deposits influenced by north – south current in the Mozambique Basin were associated with

the identification of late Cretaceous contourite deposits in the Somali Basin thus suggesting that a southward coast-parallel current originating from or North of the Somali Basin flowed into and through the Mozambique Channel in late Cretaceous times (Castelino et al., 2016, 2015).

2.3. Potential sources of sediments deposited in the Mozambique Channel

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The Cenozoic fine-grained sediments deposited at DSDP 25 Site 242 are likely deposited by southward and northward paleo-oceanic currents and originated from by rivers draining the African continent and Madagascar. Southward paleo-oceanic current may have similar direction as the modern Mozambique Current and hence transported sediments may have similar composition. As the NEMC approaches the African coast entering the Mozambique Channel to feed into the Mozambique Current, it may receive sediments from: 1) African rivers draining the Neoproterozoic Mozambique Belt and younger volcanic and sedimentary deposits, 2) rivers of NW Madagascar that drain Paleozoic to Quaternary sedimentary rocks and Cenozoic volcanic rocks and 3) volcanic rocks from the Comoros and Mayotte island (van der Lubbe et al., 2016). A present-day surface sediment sampled close to the DSDP 25 Site 242 leg yields an $\mathcal{E}_{Nd(0)}$ value of -11.2 (van der Lubbe et al., 2016, Fig. 1), and may be considered as representative of sediments transported by the NEMC and deposited in the Mozambique Channel by the Mozambique Current. Northward paleo-undercurrent may follow a reverse trajectory to that of the presentday Agulhas Current (AC in Fig.1). Hence, sediments potentially transported by this near-bottom current may receive detrital input from southeastern African rivers draining the Archean Zimbabwe craton, the Mesoproterozoic Southern Irumide belt, Phanerozoic sedimentary rocks and volcanic deposits of the Karoo supergroup. These sedimentary end-members might be best represented by the Limpopo and Zambezi rivers (see Fig. 1A for localization), which are the largest single sediment contributors to the southwestern Indian Ocean. In addition to these rivers, one possible endmember could be the Congo river as it might have discharged into the Indian Ocean via the Rufiji Delta, located in Tanzania (northward the study area, see Fig. 1A for location), prior to the uplift of the East African rift system during the Eocene-Oligocene (Stankiewicz and de Wit, 2006). To constrain the Congo endmember, we calculated the mean $\mathcal{E}_{Nd(0)}$ values and Nd concentrations of the KZR-23 core sediments reported in Bayon et al. (2019). This gives a $\mathcal{E}_{Nd(0)}$ value of -15.6 and Nd concentration of 23.5 ppm. To characterize the Limpopo and Zambezi river sediments, volcanic rocks from the Comoros and Mayotte islands and sediments deposited on the

Mozambique margin, we have compiled 142 published Sr and Nd isotope data (supporting information Table S1 (Deniel, 1998; Garzanti et al., 2014; Pelleter et al., 2014; Tucker et al., 1999; van der Lubbe et al., 2016). Based on this compilation, there is a quite distinct average $\mathcal{E}_{Nd(0)}$ values on modern sedimentary input if considered samples are from the Limpopo (-26.2 Standard Deviation (S.D.4.3) and upstream Zambezi (-12.4, S.D.: 4.0) river sediments, as well as from the Comoros (2.3, S.D.:1.5) and Mayotte (4.1, S.D.:0.8) volcanic rocks. In comparison, modern Mozambique Channel sediments yield $\mathcal{E}_{Nd(0)}$ values range between -17.7 and -11.2 (mean value: -15.5, S.D.:1.5), which encompasses the average εNd(0) value for modern Zambezi shelf surface clay (-16.7; $1\sigma \pm 0.6$, n = 5) and, hence, modern Zambezi suspension load (van der Lubbe et al., 2016). This average value is lower than the $\varepsilon Nd(0)$ values of upstream Zambezi river sediments measured by Garzanti et al (2014), thus indicating that the Zambezi suspended load also receives contributions from old Zimbabwe and Mozambique belt cratonic terrains. Sediments from the Limpopo (0.737850, S.D.: 0.024686) and Zambezi (0.7182750, S.D.: 0.010343) river sediments, and Comoros (0.703538, S.D.: 0.000272) and Mayotte (0.703351, S.D.: 0.000130) volcanics also present distinct 87 Sr/ 86 Sr isotopic compositions. The Archean Madagascar terranes have a mean $\mathcal{E}_{Nd(0)}$ value and 87 Sr/ 86 Sr isotopic composition of -22.6 (S.D.:8.7) and 0.716081, (S.D.:0.014705), respectively (Table S1). Holocene sediments deposited in a deep-sea interfluve proximal to the mouths of major northwestern Madagascan rivers yield a mean $\mathcal{E}_{Nd(0)}$ of -19.5, S.D.: 0.3 (Table S1, Fontanier et al., 2018).

Materials and Methods

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3.1 Sampling

Thirty-six pelagic and hemipelagic sediments were sampled from DSDP leg 25 Site 242 (Table 1). The age of the samples has been determined based on nanoplankton zonation and planktonic foraminifera determination (Simpson et al., 1974). The samples range from late Eocene to Quaternary (Table 1). Details regarding the lithology, mineralogy and biostratigraphy of the analyzed samples can be found in the Shipboard Scientific Party Site report (Simpson et al., 1974).

3.2 Major and trace elements and Sr-Nd isotopes

3.3.1 Sample preparation

Each bulk sample was gently crushed in an agate mortar. About 3 g of the crushed sediment were placed into 50-ml centrifuge tubes for sequential leaching to remove the non-terrigenous sedimentary components following the method of Bayon et al. (2002). Three different solutions composed of 5% acetic acid, 15% acetic acid and hydroxylamine hydrochloride (0.05 M) and hydrogen peroxide (H₂O₂, 5%) were then added to remove the carbonates, Fe-Mn oxides and organic carbon, respectively. After the removal of non-terrigenous components, the residual fractions were cleaned with ultrapure water (MQ-H₂O, 18 MΩ). Clayey (<2 μm) and silty (2-63 μm) fractions were then separated by centrifugation in two steps. First, 25 ml of MQ-H₂O were added to the detrital residues in the tubes, shaken vigorously, and then centrifuged for 2 min at 1000 rpm (133 g). The clayrich supernatants were immediately transferred into new 50 ml centrifuge tubes. Another 25 ml of MQ-H₂O was added to silt-rich detrital residues, mixed thoroughly again, centrifuged for 2.5 min at 800 rpm (85 g), and transferred into corresponding centrifuge tubes. Finally, clay-size fractions were collected after decantation (48 hours) and centrifugation at 3500 rpm (1630 g) from the residual detrital material using low-speed centrifugation (Bayon et al., 2015). These clay-size fractions were then analyzed for their major and trace element concentrations and their Sr-Nd isotopic compositions.

3.3.2 Major and trace element analyses

The major and trace element concentrations of the clay-size fractions were measured at the Service d'Analyse des Roches et Minéraux (SARM, INSU facility, Vandoeuvre-Les-Nancy, France) by ICP-OES (Na, Mg, K, Ca, Sc, Ti, Mn, Fe, Al, Si and P) and ICP-MS (Rb, Cs, Ba, Sr, Th, U, Y, Zr, Nb, Hf, Cr, V, Co, Cu, Ni, Zn, La, Ce, Pr, Nd, Sm, Eu, Gd, Tb, Dy, Ho, Er, Tm, Yb, Lu and Ta) after alkali fusion. Analytical details are available at http://helium.crpg.cnrs-nancy.fr/SARM/ and in Carignan et al. (2001). Uncertainties are lower than 5% for elements measured by ICP-OES, and lower than 10% for elements measured by ICP-MS (see supplementary material Table S2).

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Among the samples analyzed for Sr and Nd isotopes, twenty-one were analyzed at the Geochronology Laboratory of the University of Brasília (Geochronos) and nine at the Geosciences Environment Toulouse (GET) in France. At GET, about 100 mg of the leached detrital residue fractions were accurately weighed into cleaned Savillex PFA vials prior to digestion at elevated temperatures in the GET clean room. Organic matter was first oxidized by hydrogen peroxide (H₂O₂) for 24 hours at ambient temperature, and then digested in HNO₃ for 24 hours at 80°C followed by HF-HNO₃ for 24 hours at 80°C, and, finally digested in HCl+HNO₃ for 24 hours at 115°C. Blank tests were performed to estimate the level of contamination induced by the acid digestion, but it was found to be negligible. Aliquots containing about 1000 ng of Sr and Nd were loaded into the ion exchange columns. Sr and Nd were separated using the Sr-SPEC, TRU-SPEC and LN-SPEC resins Eichrom. Nd and Sr isotopic ratios were measured using a Finnigan Mat 261 thermal ionization mass spectrometer and a Triton Thermal Ionization Mass Spectrometer (TIMS) in dynamic mode. During the Nd runs, a 146Nd/144Nd ratio of 0.7219 was used to correct the signal for mass fractionation. For each sample, checks were made for the absence of samarium (Sm). The accuracy of the measurements was estimated on the La Jolla standard. Repeated analyses of the La Jolla standard gave a 143 Nd/ 144 Nd ratio of 0.511867 \pm 0.000018 (2 S.D, n = 12) in agreement with the recommended value of 0.511858 (Lugmair et al., 1983). During the Sr runs, ${}^{86}\text{Sr}/{}^{88}\text{Sr} = 0.1194$ was used to correct the signal for mass fractionation. The accuracy of the ⁸⁷Sr/⁸⁶Sr measurements was checked against the NBS 987 standard (= 0.710240) Hodell et al., 2007). Repeated analyses of the NBS 987 standard gave an 87 Sr/ 86 Sr ratio of 0.710231 \pm 0.000032 (2SD, n = 15) in agreement with the recommended value of Hodell et al. (2007) therefore no instrumental bias needs to be taken into account. Total blanks (acid digestion plus column chemistry) for Nd and Sr were checked by ICP-MS and found to be negligible (<1 per mil) compared to the Nd and Sr amounts loaded onto the columns.

At the Geochronos Lab of the Geosciences Institute of the Universidade de Brasília, 20–40 mg aliquots of the sediments were digested under clean laboratory conditions in two stages using concentrated HF-HNO₃ and 6 N HCl on a hot plate. The Nd were separated using chromatographic columns, including AG 50W-X8 cation exchange resin for Sr and Rare Earth Elements (REEs). Neodymium was further isolated from the REE fraction using a second AG 50-X2 cation-exchange resin. Neodymium was loaded on double rhenium filaments with H₃PO₄ (0.1 M) and measured as a metal (Nd+) (Gioia and Pimentel, 2000). All of these analyses were performed on a TIMS (ThermoFisher, Triton).

The measured 143 Nd/ 144 Nd_{sample} ratios are expressed in epsilon notation as the fractional deviation in parts per 10^4 (units) from 143 Nd/ 144 Nd value of in the Chondritic Uniform Reservoir (CHUR). This notation is defined as: ε Nd(t) = [(143Nd/144Nd)sample(t) /(143Nd/144Nd)CHUR(t) - 11×104

where t indicates the time at which ε_{Nd} is calculated. Here, no time correction is applied (t = 0) and the ¹⁴³Nd/¹⁴⁴Nd_{CHUR(0)} = 0.512638 (Jacobsen and Wasserburg, 1980).

Results

Major and trace element concentrations and Sr-Nd isotopic compositions of samples analyzed in this study are reported in Table 2 and Table 3, respectively. Table 4 reports the mean values and associated standard deviation of elemental ratios and isotopic composition for each age period.

4.1 Major elements, Large-Ion Lithophile Elements (LILE), High Field Strength Elements (HFSE) and Trace Transition Elements (TTE)

The elemental concentrations of major elements, Large-Ion Lithophile Elements (LILE), High Field Strength Elements (HFSE) and Trace Transition Elements (TTE) of samples normalized to the Post Archean Australian Shales (PAAS) concentration are shown in Fig. 2. As a whole, the elemental composition of the analyzed DSDP samples is rather uniform. They show depletions in SiO₂, MnO₃

K₂O, Cs, Sr, U, Zr and Hf against PAAS, values which are representative of the upper continental crust composition (Taylor and McLennan, 1985). Other elemental concentrations are close to PAAS values. Four samples (3R3W, 6R3W, 14R3W, 17R1W) show slight enrichment in CaO and Sr against PAAS that might be due to incomplete removal of marine carbonates. No apparent trend of major depletion or enrichment in elemental concentration is recorded through time.

It might be possible to assess the degree of source-area weathering and subsequently paleoweathering conditions of ancient sediments using the Chemical Index of Alteration (CIA) (Nesbitt and Young, 1982; Fedo et al., 1995) (Table 2). This index measures the weathering intensity of feldspars relative to unaltered protoliths, and is defined as: $CIA = [Al_2O_3/(Al_2O_3+CaO^*+Na_2O+K_2O)]\times 100$ (in molar proportions), where CaO^* represents the CaO content in the silicate fraction. CIA values for unaltered plagioclase and K-feldspars typical of unaltered upper crustal rocks are approximately equal to 50, whereas higher CIA values represent higher degrees of weathering. For comparison, the CIA value of the PAAS standard is equal to 67 (Taylor and McLennan, 1985).

As aforementioned, four samples enriched in CaO, probably due to incomplete removal of marine carbonates, have anomalous low CIA values (between 44 and 64, Table 2). The remaining DSDP samples yield higher CIA values (72 to 84). There is a slight increase in CIA values from the late Eocene (mean \sim 72 \pm 4) to the late Oligocene (mean \sim 78 \pm 1). The CIA values from sample younger than late Oligocene are comparable. The Al/Si atomic ratio of analyzed DSDP samples range from 0.38 to 0.54 (Table 2). The analyzed DSDP site samples have slightly higher Cr/Th ratios (7.7 to 13.7) and lower Th/Sc ratios (0.50 to 0.74) than PAAS (Table 2). There is no apparent variation of the Cr/Th ratios as a function of the age of the sample (Table 2). There is a slight decrease of the mean Th/Sc ratio with the age.

4.2 Rare Earth Elements (REE)

The DSDP samples have comparable REE patterns with flat Light and Middle REE and depletion in Heavy REE (HREE) against PAAS (Fig. 3). To facilitate comparisons with sediments derived from basic or felsic sources (Cullers, 2000) and the PAAS (Taylor and McLennan, 1985), the Eu anomaly is calculated with respect to chondrites: $Eu/Eu^* = Eu_N/(Sm_N \times Gd_N)^{1/2}$ where N refers to

the chondrite-normalized concentration value. The Eu/Eu* ratios of DSDP samples are slightly higher (0.63-0.72) to that of the PAAS standard (*i.e.*, Eu/Eu* = 0.57 ± 0.02 , Pourmand et al., 2012). There is a slight increase of the mean Eu/Eu* ratio in DSDP samples from the late Eocene to the late Miocene (Fig. 3). The Cerium anomaly (Ω Ce) normalized to PAAS is calculated following De Baar et al. (1985): Ω Ce = 2 × (Cesample/CePAAS) / [(Lasample/LaPAAS) + (Prsample/PrPAAS)], where Xsample is the concentration of samples and X_{PAAS} refers to the concentrations of the PAAS standard (Pourmand et al., 2012). The analyzed Ce anomalies display a narrow range of values (0.90-1.02) close to the PAAS standard (*i.e.*, Ω Ce = 1).

4.3 Sr-Nd isotopes

Overall, the ${}^{87}\text{Sr}/{}^{86}\text{Sr}$ ratios of DSDP samples show a slight decrease from the late Eocene (mean ~ 0.719626 ± 0.001305 , n=5) and the early Oligocene (mean ~ 0.72065 ± 0.004375 , n=6) to the Quaternary (mean ~ 0.715576 ± 0.000685 , n=2). With the exception of one Quaternary sample (1R4W) and one late Pliocene sample (2R3W), all DSDP samples show an overall decrease in $\mathcal{E}_{Nd(0)}$ values from the late Eocene to the Quaternary (from ~-14.1 to -11.8).

Discussion

5.1 Provenance of Cenozoic sediments

The REE contents, Eu anomalies, elemental ratios (Cr/Th and Th/Sc) and Sr-Nd isotopic compositions of sedimentary rocks have been proven to be useful tools for determining the provenance of sediments (McLennan et al., 1993). For instance, high Eu anomalies and Th/Sc and low Cr/Th are characteristic of felsic and more highly differentiated source rocks, whereas high Cr/Th and low Eu anomalies and Th/Sc suggest more mafic and less differentiated source rocks (Cullers, 2000; McLennan et al., 1993). However, before drawing conclusions on provenance, the effects of chemical weathering and marine authigenesis should be carefully assessed as these processes may control the REE content and radiogenic isotope composition of marine sediments. The absence of any correlation (i.e. r>= 0.75) of CIA and Al/Si with REE content, Eu/Eu*, Cr/Th, Th/Sc, ⁸⁷Sr/⁸⁶Sr isotopic compositions and ENd(0) suggests that chemical weathering has not modified these provenance

proxies. Any influence from carbonates, Fe-Mn oxides or organic matter formed under marine conditions should have been removed by the sequential carbonate dissolution and leaching steps. Furthermore, all the analyzed DSDP samples have Ce anomalies higher than or equal to 0.90 and show LREE enrichment and HREE depletions against PAAS. As the REE patterns of authigenic smectites formed in deep-water environments are normally characterized by LREE depletion, HREE enrichment and strong negative Ce anomalies (Ω Ce < 0.50, De Baar et al., 1985) typical of seawater (Piper, 1974), this suggests that, even if authigenic smectites were incorporated during early diagenetic processes, its proportion was much less than the detrital fraction.

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Slight increase of Eu/Eu*, Cr/Th ratios, $\mathcal{E}_{Nd(0)}$ values associated with slight decrease in Sr isotopic composition and Th/Sc ratios from late Eocene to Quaternary point to slight input of more "basic" sensu Cullers (2000), less differentiated and younger detritus to the DSDP 25 Site 242 and DFZ through time (Figs 4 and 5). The late Eocene samples have the highest $\mathcal{E}_{Nd(0)}$ values of the analyzed samples (-14.1 \pm 0.5, n=4). They have $\mathcal{E}_{Nd(0)}$ values significantly higher than those of Madagascar Archean terranes (-22.6 \pm 4.4, n=15, Table S1, Tucker et al., 1999), Holocene sediments deposited by major northwestern Madagascan rivers (-19.5 \pm 0.2, n=14, Table S1, (Fontanier et al., 2018) and Limpopo River sediments (-26.2 ± 3.0, n=8, Table S1, Garzanti et al., 2014), hence ruling out Zimbabwe cratonic or Madagascar provenance as the main source of these sediments. In addition, the late Eocene DSDP sediments have mean $\mathcal{E}_{Nd(0)}$ values close to those which have been deposited at 1329 m water depth to the North of the Zambezi river during the past ~45,000 years ($\mathcal{E}_{Nd(0)}$ -14.1 ± 0.2, n = 52, Table S1, van der Lubbe et al. 2016) which may signify that they are sourced by paleo-rivers that drained similar terranes. This may indicate they were drained by the paleo-Zambezi river assuming that the late Eocene paleodrainage drained proportionally similar terranes as today. Alternatively, the late Eocene sediments could have been sourced by the Paleo Congo river as it might have discharged into the Indian Ocean prior to the uplift of the East African rift system during the Eocene-Oligocene (Stankiewicz and de Wit, 2006). When compared with $\mathcal{E}_{Nd(0)}$ values of the late Eocene DSDP 25 site 242, Holocene muds of the Congo fan have $\mathcal{E}_{Nd(0)}$ values slightly lower (-15.6 ± 0.1 1 S.D.; (Bayon et al., 2019)) than those of the late Eocene-Oligocene. Three late Pliocene to

Quaternary samples out of five have $\mathcal{E}_{Nd(0)}$ values similar to that of the northernmost present-day surface sediments deposited on the Davie Ridge ($\mathcal{E}_{Nd(0)} = -11.2$, van der Lubbe et al., 2016). Furthermore, the late Eocene to early Oligocene samples have $\mathcal{E}_{Nd(0)}$ values similar to those of the Zambezi and present-day surface sediments situated west to the DFZ and DSDP 25 Site 242 (i.e. $\mathcal{E}_{Nd(0)}$ = -14.4, van der Lubbe et al. 2016, Fig. 1). The high radiogenic $\mathcal{E}_{Nd(0)}$ value of the lower part of the DSDP core is interpreted to reflect mixing between low Nd radiogenic detritus originated from the Paleo Zambezi or the Paleo Congo rivers and high radiogenic detritus coming from the north and transported by the NEMC (van der Lubbe et al., 2016). Potential sources that could explain the contribution of high radiogenic Nd detritus to the NEMC include 1) the Lurio and Rovuma Rivers that drain the Neoproterozoic Mozambique Belt and geologically younger volcanic and sedimentary deposits, and 2) Cenozoic volcanic rocks of the Comoros and Mayotte islands. At present, it is impossible to further distinguish between these two potential sources, because there is no published Nd isotopic compositions of sediments from Lurio and Rovuma to the best of our knowledge. Whatever these contributors may be, we interpret the high radiogenic $\mathcal{E}_{Nd(0)}$ values of the youngest Quaternary DSDP sample (1R2W) analyzed in this study and the northernmost sediment deposited in the Mozambique Channel analyzed by van der Lubbe et al. (2016) as representative of sediments deposited by the NEMC before being mixed with Paleo Zambezi or Paleo Congo river detritus. Together with the $\mathcal{E}_{Nd(0)}$ values, Eu/Eu* and Cr/Th ratios also increase, whereas the 87 Sr/ 86 Sr and Th/Sc ratios decrease from the late Eocene to the Quaternary (Figs 4 and 5). These trends may be interpreted to reflect an increasing contribution of NEMC transported suspended material and/or lowering contributions of Zambezi or Congo suspended load trough time. In this scenario, using a simple mixing equation (1), we can calculate the relative contribution of the Zambezi/Paleo Congo and NEMC material in each sediment DSDP 25 site 242 sample.

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Unfortunately, there is no Nd concentration available for the study of van der Lubbe et al. (2016) to better constrain the NEMC and Paleo Zambezi/Paleo Congo (termed hereafter Southeast Africa) endmembers. We assume that the NEMC Nd isotopic composition and concentration endmember is represented by the youngest Quaternary sample (1R2W) analyzed in this study. To

constrain the Southeast Africa endmember, we used the Nd isotopic composition and concentration of the VM19-214 sample analyzed by Franzese et al. (2006), which represent the geochemical composition of sediments transported by the Mozambique Current after receiving the maximum input of Zambezi sediment. These end-members provide $\mathcal{E}_{Nd(0)}$ values of -11.8 and -15.5 and Nd concentrations of 24.8 and 33.3 ppm for the NEMC and South Africa (SeA) endmembers, respectively. The relative contribution of South Africa (Zambezi/Congo) and NEMC endmembers can be calculated using the following equations:

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$$\left(\varepsilon Nd_{(0)} \right)_{DSDP} = \left(\varepsilon Nd_{(0)} \right)_{SeA} * X + (1 - X) \times \left(\varepsilon Nd_{(0)} \right)_{NEMC}$$
 (1)

where X is the relative proportion of Nd delivered by the Southeast Africa (SeA) endmember in the sediment core sample.

$$X = M_{SeA} \times C_{SeA} / M_{DSDP} \times C_{DSDP}$$
 (2)

With M_{SeA} the sediment mass delivered by the Zambezi or Congo River, C_{SeA} the Nd concentration in the sediment of the Zambezi or Congo, M_{DSDP} the total sediment mass of the core sample and C_{DSDP} the Nd concentration in the core sample. Furthermore, we defined F_{SeA} as the mass fraction of the sediment in the core derived from the Zambezi or Congo rivers which is equal to the M_{SeA}/M_{DSDP} ratio. By combining the equations (1) and (2) we obtain the following equation:

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$$F_{SeA} = \frac{M_{SeA}}{M_{Core}} = \left(\left(\varepsilon N d_{(0)} \right)_{DSDP} - \left(\varepsilon N d_{(0)} \right)_{NEMC} \right) \div \left(\left(\varepsilon N d_{(0)} \right)_{SeA} - \left(\varepsilon N d_{(0)} \right)_{NEMC} \right)] \times C_{DSDP}$$
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$$\div C_{SeA}$$

Using the Nd concentrations of analyzed sediments (Table 2), we can calculate the proportion of sediment originating from the Zambezi or Paleo Congo river in the Eocene-Oligocene DSDP sediments and originating from the Zambezi river in younger DSDP sediments. According to this calculation, there is an overall decrease of the Southeast African contribution from ~ 65.7% (Standard Deviation (S.D.) ~ 13.2) in the late Eocene to ~28.5% (S.D. ~ 10.0) in the early Pliocene. Additionally, there is a very strong variability in Southeast African contributions during the late Pliocene to Quaternary between virtually no contribution and up to 71.9% (Table 3 and Fig.5). These variations in the Southeast African contribution are not linear and roughly follow four steps. The oldest period

between the late Eocene and early Oligocene is characterized by Southeast African contribution ≥to 50%. The late Oligocene to early Miocene period shows general decrease in the Southeast African contribution from ~ 37.4% to 51.2%. The third period between the middle Miocene and early Pliocene has Southeast African contributions between ~23.4% and 28.5%. Finally, the youngest Pliocene-Quaternary period shows strong variations between minor Southeast African contribution (<5 %) to strong Southeast African contributions (52.8% and 71.9%).

5.2 Controls on provenance variations and implications for past oceanic circulation in the Mozambique Channel

At present, the fine-grained sediments deposited at the location of DSDP 25 Site 242 are expected to have very minor Zambezi contribution due to the combined presence of the Davie Ridge and the strong southward Mozambique Current that may prevent Zambezi detritus reaching the DSDP 25 Site 242 location. In addition to variation in the Mozambique Current and tectonic activity of the Davie Ridge, the variation in provenance and decrease in Zambezi contribution since the late Eocene may be also caused by drainage change and variation in riverine supply in response to tectonic and climatic change that may have affected Southeast Africa. Hence, the late Eocene to Quaternary geochemical composition of DSDP 25 Site 242 sediments can be used as an archive of how the variation of in riverine supply, drainage changes, tectonic activity of the Davie Ridge and variation in the NEMC may have controlled their provenance (Fig.5).

The oldest period between the late Eocene and early Oligocene is characterized by the lowest $\mathcal{E}_{Nd(0)}$ values and strongest Southeast African contribution in the DSDP 25 site 242 samples. The onset of the modern deltaic sedimentation in the Zambezi delta is thought to occur around the Eocene-Oligocene boundary and is marked by Zambezi elevated sediment fluxes (Walford et al., 2005) and the presence of a major tectonic controlled unconformity related to the second uplift phase of the South African Plateau (Ponte et al., 2019). Before the late Eocene -early Oligocene (latest Bartonian to early Priabonian, 38.2–36.2 Ma, Ponte et al. 2019), the Zambezi delta sediments recorded a stable tectonic setting as emphasized by the presence of Eocene marine carbonates (Ponte et al., 2019). Given the

available stratigraphic resolution, it is unclear if the oldest samples analyzed here postdate or predate this unconformity. If predating, the elevated Southeast African contribution and low $\mathcal{E}_{Nd(0)}$ values recorded by the DSDP 25 site 242 samples cannot be explained by the onset of modern sedimentation in the Zambezi delta.

Alternatively, the late Eocene-early Oligocene sediments could have been sourced by a paleoriver draining similar terranes as the Zambezi. One possible candidate could be the Paleo Congo river which might have discharged into the Indian Ocean via the Rufiji Delta, located in Tanzania (see Fig. 1A), prior to the uplift of the East African rift system during the Eocene-Oligocene (Stankiewicz and de Wit, 2006). In this scenario, the thick early Eocene-early Miocene slope and basin fan sedimentary rocks (termed Mega-sequence 4) deposited over much of the Tanzania and Mozambique deepwater offshore (McDonough et al., 2013) could represent the distal fan of the Paleo Congo river. This scenario cannot be excluded on the basis of the $\mathcal{E}_{Nd(0)}$ values of Congo river sediments which are slightly lower (-15.6 \pm 0.1 (1 S.D.); Bayon et al., (2019)) than those of the late Eocene-early Oligocene. In this scenario, 100% of the sediment deposited at DSDP 25 Site 242 would be transported by the NEMC and in turn this mean that the $\mathcal{E}_{Nd(0)}$ values of sediment transported by the NEMC would have changed between the early and late Oligocene. Then, the second uplift phase of the South African Plateau (Baby et al., 2018) which coincides with the onset of modern sedimentation in the Zambezi delta (Ponte et al., 2019) and elevated Zambezi sediment flux (Walford et al., 2005) might have stopped the connection between the Paleo Congo and the Indian Ocean if it did exist.

Hence concerning the potential sources for the late Eocene-early Oligocene DSDP 25 site 242 samples, both a Zambezi and a Paleo Congo source are possible. If the Paleo Congo connection with the Indian Ocean is stopped before the deposition of late Eocene-early Oligocene sediments, the low $\mathcal{E}_{Nd(0)}$ values of the DSDP 25 site 242 samples associated with low sedimentation rates at DSDP 25 Site 242 between the late Eocene and early Oligocene (3-9 m/m.y., (Simpson et al., 1974), Fig. 5) suggest very minor contribution from a paleo southward NEMC. This could be due to the absence of tectonic activity of the DFZ during the late Eocene –early Oligocene (Deville et al., 2018; Franke et al., 2015; Mahanjane, 2014; Mougenot et al., 1986), which may not have acted as a barrier to sediments coming

from the South African coast to the DSDP 25 Site 242 or absence or low intensity of the NEMC which may have not been capable of transporting detritus with higher $\mathcal{E}_{Nd(0)}$ values.

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The DFZ is thought to have experienced a southerly extension related to East African Rifting System since the Oligocene (Deville et al., 2018; Franke et al., 2015; Mahanjane, 2014; Mougenot et al., 1986). This extension coincides with a period of an elevated sediment flux during Oligocene times (34-24 Ma) in the Zambezi deep-sea fan as a response of rapid regional uplift of Southern Africa (Walford et al. 2005; Baby et al., 2018) that may explain the strong contribution of Southeast African detritus to DSDP 25 Site 242 sediments. This is in agreement with the increase in mass accumulation rate at DSDP 25 site 242 between the late Oligocene and middle Miocene (14 to 23 m/m.y., (Simpson et al., 1974) when compared with those of the late Eocene-early Oligocene period (Fig. 5). In the middle Miocene, graben formation in the DFZ possibly led to the abandonment of the NS Serpa Pinto Valley and in turn to the export of Zambezi sediments to the Mozambique Channel through west-east Zambezi Valley (Droz and Mougenot, 1987). This renewed tectonic activity in the DFZ may have promoted the formation of a barrier preventing the arrival of Zambezi sediments to DSDP 25 Site 242 location, which would explain the decrease in Southeast African contribution between the early and middle Miocene (from ~51 to ~23%). Alternatively, this reduction can also be explained by strengthening of the southward flowing Mozambique Current. The initiation of the southward Mozambique Current and its effect on sediment dispersion in the Mozambique Channel are controverted. Recognition of late Cretaceous contourites deposits in the Somali Basin (Coffin et al., 1986) associated with identification of late Cretaceous north to south current–controlled deposits in the Mozambique Basin (Castelino et al., 2015) have been interpreted to document the existence of southward coast-parallel current that was flowing into and through the Mozambique Channel in late Cretaceous times (Castelino et al., 2015). In contrast, the morphology of the Zambezi deep-sea fan changed from lobate to elongate during the Miocene, which suggests that, there is no "strong" influence of a southward flowing current on sediment distribution in the Mozambique Channel before the early Miocene (Walford et al. 2005). Rather, these authors proposed that the southward Mozambique Current was established in the early Miocene (Walford et al., 2005). This early Miocene age for the presence of the Mozambique Current is also confirmed by the identification of plastered drift sequences south of the Mozambique Channel along the continental margin of Mozambique off the Limpopo River (Preu et al., 2011). As there is no published evidence for change in drainage configuration of the Zambezi catchment, we argue that the drop in Southeast African contribution between the early and middle Miocene could be explained by a combination of uplift of the DFZ and/or strengthening of the Mozambique Current.

The late Pliocene-Quaternary period is marked by a large spread in $\mathcal{E}_{Nd(0)}$ values and Southeast African contribution in sediments deposited at DSDP 25 Site 242. Calculated sedimentary fluxes exported by the Zambezi river to the Mozambique basin show a period of elevated flux from late Miocene times (10 Ma) to present day due to a doubling of the size of the Zambezi catchment during the Pliocene (Walford et al., 2005). This could explain the strong Southeast African contribution for late Pliocene 2R3W and Quaternary 1R4W samples. However, it cannot explain the low Southeast African contributions for other analyzed samples (Table 3 and Fig. 5). Alternatively, the low Southeast African contribution for other late Pliocene-Quaternary sediments (Table 3) could be explained by increasing tectonic activity of DFZ or increasing intensification of Mozambique Current. However, these mechanisms cannot account for the strong Southeast African contribution in samples 2R3W and 1R4W. We propose therefore that the strong $\mathcal{E}_{Nd(0)}$ variations and in turn inferred Zambezi (Southeast Africa) contributions during the late Pliocene-Quaternary are linked to the enhanced glacial-interglacial cycles.

During sea-level highstand interglacial conditions, the accommodation space is increased in the delta region which increased Zambezi sediment deposition in the delta and limited the amount of Zambezi sediment delivered to the Mozambique Basin and Channel (Wiles et al., 2017). This also result in higher Nd radiogenic isotopic composition of sediments (van der Lubbe et al., 2016) deposited in the Mozambique Basin and Channel by the NEMC. The reverse is thought to occur during glacial sea-level lowstand regression which promote bypassing of the shelf area and activation of deepsea canyons resulting in increasing delivery of Zambezi sediments to the Mozambique Basin and Channel (Wiles et al., 2017) and lowering strength of the NEMC. These induced lower radiogenic Nd

isotopic composition of sediments (van der Lubbe et al., 2016) deposited in the Mozambique basin and Channel. In this scenario, the late Pliocene 2R3W and Quaternary 1R4W would have been deposited during glacial lowstands and other analyzed samples during interglacial highstand.

Our Nd isotopic data on DSDP 25 Site 242 supports an intensification of the Mozambique Current from the late Pliocene and onwards, which prevented the arrival of Zambezi fine grained sediments at DSDP 25 Site 242 during interglacial highstand and a combined effect of increasing Zambezi sediment export to Mozambique Channel and Basin and lowering in the intensity of Mozambique Current strength during glacial lowstand. Hence, these suggest that the dispersion of fine-grained sediments within the Mozambique Channel is mainly controlled by strong glacial-interglacial cycles and that it is no younger than late Pliocene.

Conclusion

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The provenance study of late Eocene to Quaternary sediments deposited at the Davie Ridge (Mozambique Channel) at DSDP 25 Site 242 shows that they contain a mixture of detritus originating from the Zambezi or Congo river and transported from the North by the Mozambique Current. Based on variations in Nd isotopic compositions and Nd concentrations, this study infers an overall decrease of the Southeast African (Congo or Zambezi rivers) contribution from ~ 65.7% (S.D. ~ 13.2) in the late Eocene to ~1.8% (S.D.~2.5) in the late Pliocene-Quaternary. This decrease in Southeast African contributions roughly follows four steps. The oldest period between the late Eocene and early Oligocene is characterized by Southeast African contribution (Congo or Zambezi rivers) close to or superior to 50%. The late Oligocene- early Miocene period shows a first lowering of the Zambezi contribution between ~ 37.4% and 51.2%. The third period between the middle Miocene and early Pliocene has Zambezi contributions between ~23.4% and 28.5%. And finally the late Pliocene-Quaternary period shows strong variations in Zambezi contribution (0 to 4.6% and 52.8-71.9%). The causes for the low $\mathcal{E}_{Nd(0)}$ values and high Southeast African contribution between the late Eocene and the early Oligocene remain to be determined. It could record the onset of modern sedimentation in the Zambezi delta or the discharge of the Paleo Congo river in the Indian Ocean. The overall lowering in $\mathcal{E}_{Nd(0)}$ values and Zambezi contribution between the late Oligocene and the early Pliocene is thought to be due to the combined effect of increasing bathometry of the Davie Ridge and onset and intensification of the Mozambique Current. The strong variation in Zambezi contribution and $\mathcal{E}_{Nd(0)}$ values in the late Pliocene-Quaternary is interpreted to be caused by glacial-interglacial cycles with glacial lowstand favoring higher Zambezi contribution and lower $\mathcal{E}_{Nd(0)}$ values and interglacial highstand promoting lower Zambezi contribution and higher $\mathcal{E}_{Nd(0)}$ values. Hence, these suggest that the modern dispersion pattern of fine-grained sediments within the Mozambique Channel is no younger than the late Pliocene.

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References

Baby, G., Guillocheau, F., Boulogne, C., Robin, C., Dall'Asta, M., 2018. Uplift history of a 545 transform continental margin revealed by the stratigraphic record: The case of the Agulhas transform 546 547 margin along the Southern African Plateau. **Tectonophysics** 731–732, 104-130. https://doi.org/10.1016/j.tecto.2018.03.014 548

- Bayon, G., German, C.R., Boella, R.M., Milton, J.A., Taylor, R.N., Nesbitt, R.W., 2002. An
- 550 improved method for extracting marine sediment fractions and its application to Sr and Nd isotopic
- analysis. Chem. Geol. 187, 179–199. https://doi.org/10.1016/S0009-541(01)00416-8
- Bayon, G., Schefuß, E., Dupont, L., Borges, A.V., Dennielou, B., Lambert, T., Mollenhauer,
- 553 G., Monin, L., Ponzevera, E., Skonieczny, C., André, L., 2019. The roles of climate and human land-
- use in the late Holocene rainforest crisis of Central Africa. Earth Planet. Sci. Lett. 505, 30-41.
- 555 https://doi.org/10.1016/j.epsl.2018.10.016
- Bayon, G., Toucanne, S., Skonieczny, C., André, L., Bermell, S., Cheron, S., Dennielou, B.,
- Etoubleau, J., Freslon, N., Gauchery, T., Germain, Y., Jorry, S.J., Ménot, G., Monin, L., Ponzevera, E.,
- Rouget, M.-L., Tachikawa, K., Barrat, J.A., 2015. Rare earth elements and neodymium isotopes in
- 559 world river sediments revisited. Geochim. Cosmochim. Acta 170, 17–38.
- 560 https://doi.org/10.1016/j.gca.2015.08.001
- Bonneau, L., Toucanne, S., Bayon, G., Jorry, S.J., Emmanuel, L., Silva Jacinto, R., 2017.
- Glacial erosion dynamics in a small mountainous watershed (Southern French Alps): A source-to-
- sink approach. Earth Planet. Sci. Lett. 458, 366–379.
- 564 https://doi.org/10.1016/j.epsl.2016.11.004
- Castelino, J.A., Reichert, C., Klingelhoefer, F., Aslanian, D., Jokat, W., 2015. Mesozoic and
- Early Cenozoic sediment influx and morphology of the Mozambique Basin. Mar. Pet. Geol. 66, 890–
- 567 905. https://doi.org/10.1016/j.marpetgeo.2015.07.028
- Castelino, J.A., Eagles, G., Jokat, W., 2016. Anomalous bathymetry and palaeobathymetric
- models of the Mozambique Basin and Riiser Larsen Sea. Earth Planet. Sci. Lett. 455, 25-37.
- 570 https://doi.org/10.1016/j.epsl.2016.09.018
- Coffin, M.F., Rabinowitz, P.D., Houtz, R.E., 1986. Crustal structure in the Western Somali
- 572 Basin. Geophys. J. Int. 86, 331–369. https://doi.org/10.1111/j.1365-246X.1986.tb03832.x
- 573 Collins, C., Hermes, J.C., Roman, R.E., Reason, C.J.C., 2016. First dedicated hydrographic
- 574 survey of the Comoros Basin. J. Geophys. Res. Oceans 121, 1291–1305.
- 575 https://doi.org/10.1002/2015JC011418

- 576 Cullers, R.L., 2000. The geochemistry of shales, siltstones and sandstones of Pennsylvanian
- 577 Permian age, Colorado, USA: implications for provenance and metamorphic studies. Lithos 51, 181–
- 578 203. https://doi.org/10.1016/S0024-4937(99)00063-8
- De Baar, H.J.W., Bacon, M.P., Brewer, P.G., Bruland, K.W., 1985. Rare earth elements in the
- 580 Pacific and Atlantic Oceans. Geochim. Cosmochim. Acta 49, 1943–1959.
- 581 https://doi.org/10.1016/0016-7037(85)90089-4
- de Ruijter Wilhelmus P. M., Ridderinkhof Herman, Lutjeharms Johann R. E., Schouten Mathijs
- 583 W., Veth Cornelis, 2002. Observations of the flow in the Mozambique Channel. Geophys. Res. Lett.
- 584 29, 140–1. https://doi.org/10.1029/2001GL013714
- Deniel, C., 1998. Geochemical and isotopic (Sr, Nd, Pb) evidence for plume-lithosphere
- interactions in the genesis of Grande Comore magmas (Indian Ocean). Chem. Geol. 144, 281–303.
- 587 https://doi.org/10.1016/S0009-2541(97)00139-3
- Deville, E., Marsset, T., Courgeon, S., Jatiault, R., Ponte, J.-P., Thereau, E., Jouet, G., Jorry,
- 589 S.J., Droz, L., 2018. Active fault system across the oceanic lithosphere of the Mozambique Channel:
- Implications for the Nubia–Somalia southern plate boundary. Earth Planet. Sci. Lett. 502, 210–220.
- 591 https://doi.org/10.1016/j.epsl.2018.08.052
- Dia, A., Dupré, B., Allègre, C.J., 1992. Nd isotopes in Indian Ocean sediments used as a tracer
- of supply to the ocean and circulation paths. Mar. Geol. 103, 349–359. https://doi.org/10.1016/0025-
- 594 3227(92)90025-D
- Droz, L., Mougenot, D., 1987. Mozambique Upper Fan: Origin of Depositional Units. AAPG
- 596 Bull. 71, 1355–1365.
- Fagel, N., Debrabant, P., André, L., 1994. Clay supplies in the Central Indian Basin since the
- 598 Late Miocene: climatic or tectonic control? Mar. Geol. 122, 151–172.
- 599 https://doi.org/10.1016/0025-3227(94)90209-7
- Fagel, N., Innocent, C., Stevenson, R.K., Hillaire-Marcel, C., 1999. Deep circulation changes
- in the Labrador Sea since the Last Glacial Maximum: New constraints from Sm-Nd data on sediments.
- 602 Paleoceanography 14, 777–788. https://doi.org/10.1029/1999PA900041

- Fontanier, C., Mamo, B., Toucanne, S., Bayon, G., Schmidt, S., Deflandre, B., Dennielou, B.,
- Jouet, G., Garnier, E., Sakai, S., Lamas, R.M., Duros, P., Toyofuku, T., Salé, A., Belleney, D., Bichon,
- 605 S., Boissier, A., Chéron, S., Pitel, M., Roubi, A., Rovere, M., Grémare, A., Dupré, S., Jorry, S.J., 2018.
- Are deep-sea ecosystems surrounding Madagascar threatened by land-use or climate change? Deep
- 607 Sea Res. Part Oceanogr. Res. Pap. 131, 93–100. https://doi.org/10.1016/j.dsr.2017.11.011
- Franke, D., Jokat, W., Ladage, S., Stollhofen, H., Klimke, J., Lutz, R., Mahanjane, E.S.,
- 609 Ehrhardt, A., Schreckenberger, B., 2015. The offshore East African Rift System: Structural framework
- at the toe of a juvenile rift. Tectonics 34, 2086–2104. https://doi.org/10.1002/2015TC003922
- Franzese, A.M., Hemming, S.R., Goldstein, S.L., Anderson, R.F., 2006. Reduced Agulhas
- 612 Leakage during the Last Glacial Maximum inferred from an integrated provenance and flux study.
- Earth Planet. Sci. Lett. 250, 72–88. https://doi.org/10.1016/j.epsl.2006.07.002
- Garzanti, E., Padoan, M., Setti, M., López-Galindo, A., Villa, I.M., 2014. Provenance versus
- weathering control on the composition of tropical river mud (southern Africa). Chem. Geol. 366, 61–
- 74. https://doi.org/10.1016/j.chemgeo.2013.12.016
- Gioia, S.M.C.L., Pimentel, M.M., 2000. The Sm-Nd isotopic method in the geochronology
- 618 laboratory of the University of Brasília. An. Acad. Bras. Ciênc. 72, 219–245.
- 619 https://doi.org/10.1590/S0001-37652000000200009
- Goldstein, S.L., O'Nions, R.K., 1981. Nd and Sr isotopic relationships in pelagic clays and
- 621 ferromanganese deposits. Nature 292, 324–327. https://doi.org/10.1038/292324a0
- Grousset, F.E., Biscaye, P.E., Zindler, A., Prospero, J., Chester, R., 1988. Neodymium isotopes
- as tracers in marine sediments and aerosols: North Atlantic. Earth Planet. Sci. Lett. 87, 367–378.
- 624 https://doi.org/10.1016/0012-821X(88)90001-5
- Grousset, F.E., Henry, F., Minster, J.F., Monaco, A., 1990. Nd isotopes as tracers in water
- 626 column particles: the western Mediterranean Sea. Mar. Chem. 30, 389–407.
- 627 https://doi.org/10.1016/0304-4203(90)90083-O
- Hodell, D.A., Kamenov, G.D., Hathorne, E.C., Zachos, J.C., Röhl, U., Westerhold, T., 2007.
- Variations in the strontium isotope composition of seawater during the Paleocene and early Eocene

- 630 from ODP Leg 208 (Walvis Ridge). Geochem. Geophys. Geosystems 8.
- 631 https://doi.org/10.1029/2007GC001607
- Innocent, C., Fagel, N., Stevenson, R.K., Hillaire-Marcel, C., 1997. Sm Nd signature of
- 633 modern and late Quaternary sediments from the northwest North Atlantic: Implications for deep
- 634 current changes since the Last Glacial Maximum. Earth Planet. Sci. Lett. 146, 607-625.
- 635 https://doi.org/10.1016/S0012-821X(96)00251-8
- Jacobsen, S.B., Wasserburg, G.J., 1980. Sm-Nd isotopic evolution of chondrites. Earth Planet.
- 637 Sci. Lett. 50, 139–155. https://doi.org/10.1016/0012-821X(80)90125-9
- Kolla, V., Kostecki, J.A., Henderson, L., Hess, L., 1980. Morphology and Quaternary
- sedimentation of the Mozambique Fan and environs, southwestern Indian Oceans*. Sedimentology 27,
- 640 357–378. https://doi.org/10.1111/j.1365-3091.1980.tb01188.x
- Liu, C., Clift, P.D., Carter, A., Böning, P., Hu, Z., Sun, Z., Pahnke, K., 2017. Controls on
- modern erosion and the development of the Pearl River drainage in the late Paleogene. Mar. Geol.,
- 643 Evolution of the Deep South China Sea: Integrated IODP Expedition 349 Results 394, 52–68.
- 644 https://doi.org/10.1016/j.margeo.2017.07.011
- Lugmair, G.W., Shimamura, T., Lewis, R.S., Anders, E., 1983. Samarium-146 in the Early
- 646 Solar System: Evidence from Neodymium in the Allende Meteorite. Science 222, 1015–1018.
- 647 https://doi.org/10.1126/science.222.4627.1015
- Mahanjane, E.S., 2014. The Davie Fracture Zone and adjacent basins in the offshore
- Mozambique Margin A new insights for the hydrocarbon potential. Mar. Pet. Geol. 57, 561–571.
- 650 https://doi.org/10.1016/j.marpetgeo.2014.06.015
- McDonough, K.-J., Bouanga, E., Pierard, C., Horn, B., Emmet, P., Gross, J., Danforth, A.,
- Sterne, N., Granath, J., 2013. Wheeler-transformed 2D seismic data yield fan chronostratigraphy of
- offshore Tanzania. Lead. Edge 32, 162–170. https://doi.org/10.1190/tle32020162.1
- McLennan, S.M., Hemming, S., McDaniel, D.K., Hanson, G.N., 1993. Geochemical
- approaches to sedimentation, provenance, and tectonics, in: Geological Society of America Special
- Papers. Geological Society of America, pp. 21–40. https://doi.org/10.1130/SPE284-p21

- Mougenot, D., Recq, M., Virlogeux, P., Lepvrier, C., 1986. Seaward Extension of the East-
- 658 African Rift. Nature 321, 599–603. https://doi.org/10.1038/321599a0
- Mourlot, Y., Roddaz, M., Dera, G., Calvès, G., Kim, J., Chaboureau, A., Mounic, S., Raisson,
- 660 F., 2018. Geochemical Evidence for Large-Scale Drainage Reorganization in Northwest Africa During
- the Cretaceous. Geochem. Geophys. Geosystems 0. https://doi.org/10.1029/2018GC007448
- Pelleter, A.-A., Caroff, M., Cordier, C., Bachelery, P., Nehlig, P., Debeuf, D., Arnaud, N.,
- 663 2014. Melilite-bearing lavas in Mayotte (France): An insight into the mantle source below the
- 664 Comores. Lithos 208–209, 281–297. https://doi.org/10.1016/j.lithos.2014.09.012
- Piper, D.Z., 1974. Rare earth elements in ferromanganese nodules and other marine
- 666 phases.Geochim. Cosmochim. Acta 38, 1007–1022. https://doi.org/10.1016/0016-
- 667 7037(74)90002-7
- Ponte, J.-P., Robin, C., Guillocheau, F., Popescu, S., Suc, J.-P., Dall'Asta, M., Melinte-
- Dobrinescu, M.C., Bubik, M., Dupont, G., Gaillot, J., 2019. The Zambezi delta (Mozambique channel,
- East Africa): High resolution dating combining bio- orbital and seismic stratigraphies to determine
- climate (palaeoprecipitation) and tectonic controls on a passive margin. Mar. Pet. Geol. 105, 293–312.
- 672 <u>https://doi.org/10.1016/j.marpetgeo.2018.07.017</u>
- Pourmand, A., Dauphas, N., Ireland, T.J., 2012. A novel extraction chromatography and MC-
- 674 ICP-MS technique for rapid analysis of REE, Sc and Y: Revising CI-chondrite and Post-Archean
- 675 Australian Shale (PAAS) abundances. Chem. Geol. 291, 38–54.
- 676 https://doi.org/10.1016/j.chemgeo.2011.08.011
- Preu, B., Spieß, V., Schwenk, T., Schneider, R., 2011. Evidence for current-controlled
- sedimentation along the southern Mozambique continental margin since Early Miocene times. Geo-
- 679 Mar. Lett. 31, 427–435. https://doi.org/10.1007/s00367-011-0238-y
- Revel, M., Sinko, J.A., Grousset, F.E., Biscaye, P.E., 1996. Sr and Nd isotopes as tracers of
- North Atlantic lithic particles: Paleoclimatic implications. Paleoceanography 11, 95–113.
- 682 https://doi.org/10.1029/95PA03199

- Revel, M., Cremer, M., Grousset, F.E., Labeyrie, L., 1996. Grain-size and Sr-Nd isotopes as
- 684 tracer of paleo-bottom current strength, Northeast Atlantic Ocean. Mar. Geol. 131, 233-249.
- 685 https://doi.org/10.1016/0025-3227(96)00005-9
- Ridderinkhof, H., van der Werf, P.M., Ullgren, J.E., van Aken, H.M., van Leeuwen, P.J., de
- Ruijter, W.P.M., 2010. Seasonal and interannual variability in the Mozambique Channel from moored
- 688 current observations. J. Geophys. Res. 115. https://doi.org/10.1029/2009JC005619
- Simpson, E.S.W., Schlich, R., et al., 1974. Initial Reports of the Deep Sea Drilling Project, 25,
- 690 Initial Reports of the Deep Sea Drilling Project. U.S. Government Printing Office.
- 691 https://doi.org/10.2973/dsdp.proc.25.1974
- Stankiewicz, J., de Wit, M.J., 2006. A proposed drainage evolution model for Central Africa—
- 693 Did the Congo flow east? J. Afr. Earth Sci. 44, 75–84. https://doi.org/10.1016/j.jafrearsci.2005.11.008
- Taylor, S.R., McLennan, S.M., 1985. The continental crust: Its composition and evolution.
- Tucker, R.D., Ashwal, L.D., Handke, M.J., Hamilton, M.A., Le Grange, M., Rambeloson,
- R.A., 1999. U-Pb Geochronology and Isotope Geochemistry of the Archean and Proterozoic Rocks of
- 697 North-Central Madagascar. J. Geol. 107, 135–153. https://doi.org/10.1086/314337
- Ullgren, J.E., van Aken, H.M., Ridderinkhof, H., de Ruijter, W.P.M., 2012. The hydrography
- of the Mozambique Channel from six years of continuous temperature, salinity, and velocity
- 700 observations. Deep Sea Res. Part Oceanogr. Res. Pap. 69, 36–50.
- 701 https://doi.org/10.1016/j.dsr.2012.07.003
- van der Lubbe, J. (H.)J. L., Frank, M., Tjallingii, R., Schneider, R.R., 2016. Neodymium
- isotope constraints on provenance, dispersal, and climate-driven supply of Zambezi sediments along
- the Mozambique Margin during the past ~45,000 years. Geochem. Geophys. Geosystems 17, 181–
- 705 198. https://doi.org/10.1002/2015GC006080
- van der Lubbe, J. (H.)J. L., Tjallingii, R., Prins, M.A., Brummer, G.-J.A., Jung, S.J.A., Kroon,
- D., Schneider, R.R., 2014. Sedimentation patterns off the Zambezi River over the last 20,000 years.
- 708 Mar. Geol. 355, 189–201. https://doi.org/10.1016/j.margeo.2014.05.012

Walford, H.L., White, N.J., Sydow, J.C., 2005. Solid sediment load history of the Zambezi

Delta. Earth Planet. Sci. Lett. 238, 49–63. https://doi.org/10.1016/j.epsl.2005.07.014

Figure captions

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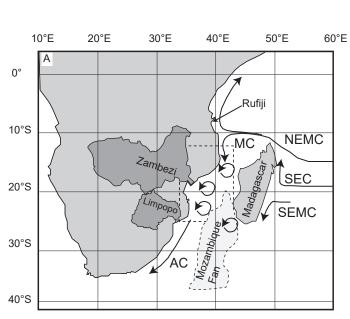
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- Figure 1. A) Map of southeast Africa showing the main surface currents (solid thin arrows) in the 712 southwest Indian Ocean (adapted from van der Lubbe et al., 2016, 2014). The Zambezi and Limpopo 713 drainage basins are indicated by the grey shaded area. The location of the present-day delta of the Rufiji 714 River is also indicated because it might represent the embouchure of the paleo-Congo River (see text for 715 details). The South Equatorial Current (SEC) is the main source of water in the Madagascar region. It 716 717 bifurcates into the North East Madagascar Current (NEMC) and the South East Madagascar Current 718 (SEMC) where approaching Madagascar coast. Part of the NEMC waters enter the Mozambique Channel (Ridderinkhof et al., 2010) and are responsible for the net southward flow through the Mozambique 719 Channel (MC). The flow through the Mozambique Channel occurs through the passage of large 720 anticyclonic eddies (Ridderinkhof et al., 2010). Sediments are mainly transported in southward direction 721 and deposited onto the large Mozambique deep-sea fan indicated in light gray (Kolla et al., 1980). The flow 722 through the Mozambique Channel is a source of the Agulhas Current (AC). The dashed rectangle indicates 723 the location of Fig. 1B. B) Detailed map of study area which shows the location of DSDP 25 Site 242 and 724 the surface distribution of selected previous published $\mathcal{E}_{Nd(0)}$ values in clay-sized lithogenic surface 725 sediments in the Mozambique Channel (Franzese et al., 2006; van der Lubbe et al., 2016). The Zambezi 726 Catchment is indicated by the dark shaded area. Depth contours are drawn at 500 m. S.P.V.: Serpa Pinto 727 Valley 728
- 729 Figure 2. Major and selected trace element contents normalized to PAAS (Taylor and McLennan, 1985;
- Pourmand et al., 2012). From left to right on the diagram, the major elements (SiO₂, TiO₂, Al₂O₃, Fe₂O₃,
- 731 MnO, MgO, CaO, Na₂O, K₂O and P₂O₅), LILE (Rb, Cs, Ba, Sr, Th and U), HFSE (Y, Zr, Nb, Hf), and
- 732 TTE (Sc, Cr, Co, V and Ni) are plotted for DSDP 25 Site 242 samples.
- 733 Figure 3. Rare earth element (REE) contents normalized to PAAS (Taylor and McLennan, 1985;
- Pourmand et al., 2012) for DSDP 25 Site 242 samples.

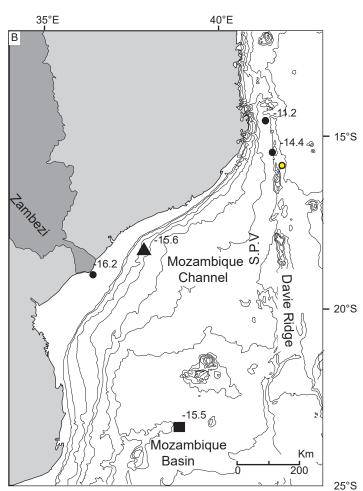
Figure 4. Variations of selected trace element ratios (Cr/Th, Th/Sc and Eu/Eu*, see text for details) since the Eocene. The lengths of the horizontal bar for the Cr/Th, Th/Sc, and Eu/Eu*, correspond to 15% (the sum of the Cr and Th concentration analytical errors), 25% (the sum of the Th and Sc concentration analytical errors) and 5% (the Eu concentration analytical error, see supplementary material Table S2). The main events with their duration affecting the studied area are also indicated. References are as follow: (1) Walford et al. (2005), (2) Droz and Mougenot (1987), (3) Mahanjane (2014), (4) Baby et al. (2018).

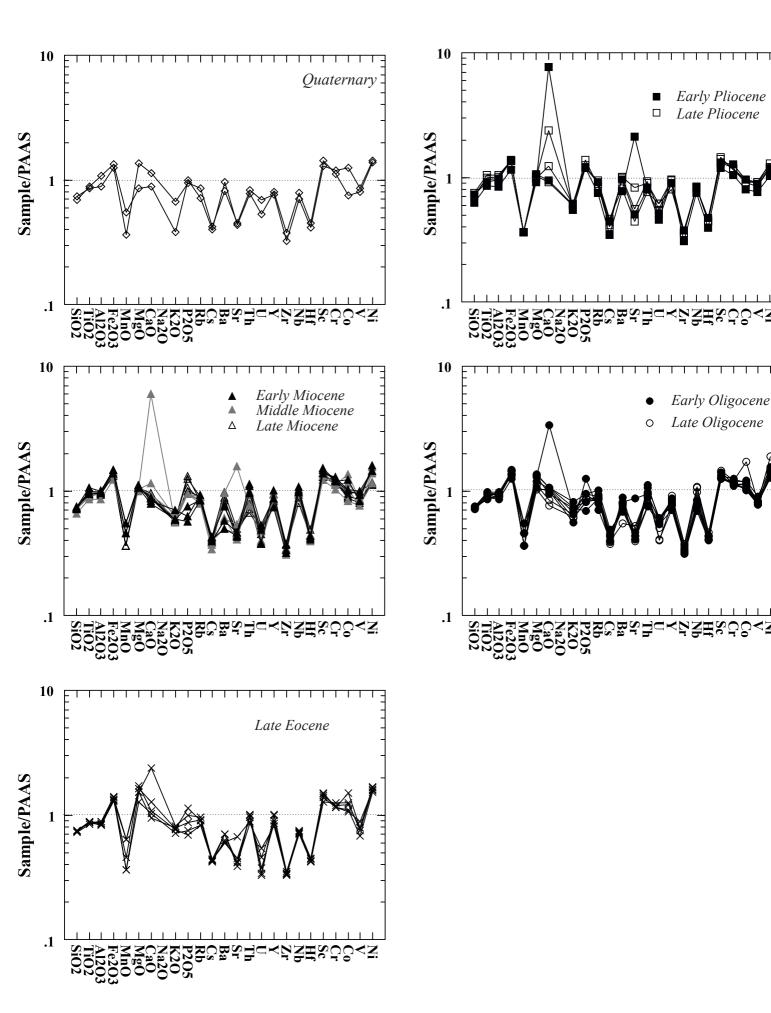
Figure 5. Variations of ⁸⁷Sr/⁸⁶Sr isotopic compositions, Nd isotopic compositions expressed as ε_{Nd(0)} values and of Southeast African contributions in % (see text for details) since the Eocene. The lengths of the horizontal bar for the Sr-Nd isotopic compositions correspond to 2σ analytical errors (see Table 3). The lengths of the horizontal bar for the Southeast contribution correspond to the sum of the Nd concentration analytical error (i.e 15%, see supplementary material Table S2) plus the % of the Nd isotopic composition 2σ analytical errors with respect to the measured Nd isotopic composition (see Table 3). The main events with their duration affecting the studied area are also indicated. References are as follow: (1) Walford et al. (2005), (2) Droz and Mougenot (1987), (3) Mahanjane (2014), (4) Baby et al. (2018). The sedimentation rates are taken from Simpson et al. (1974) and expressed in m/m.y.

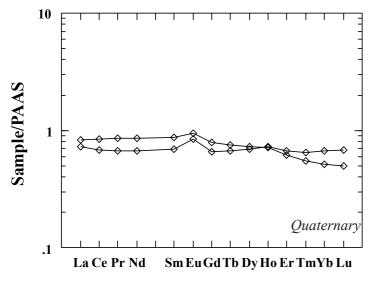


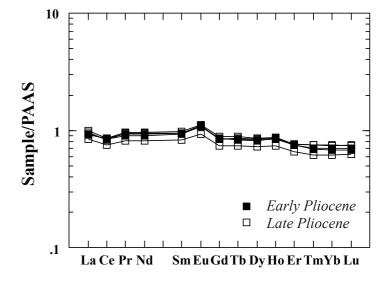


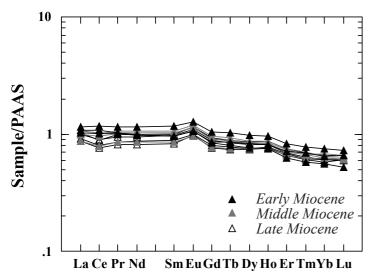
- Core 64PE304-80/GIK16160-3 (van der Lubbe et al., 2016)
- VM 19-214 (Franzese et al. (2006)
- O DSDP 25 Site 242 (this study)

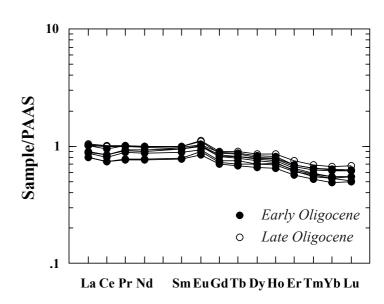


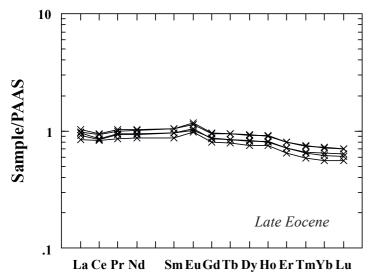


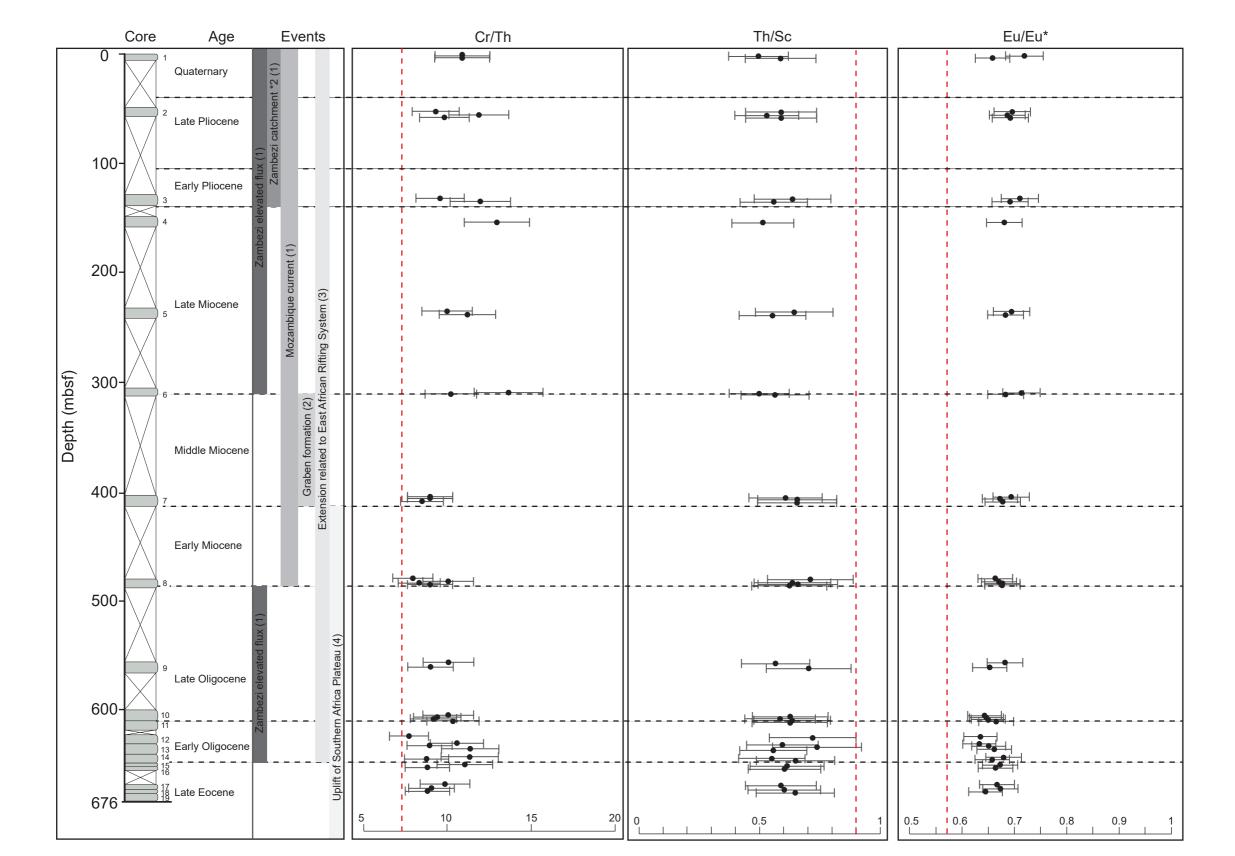


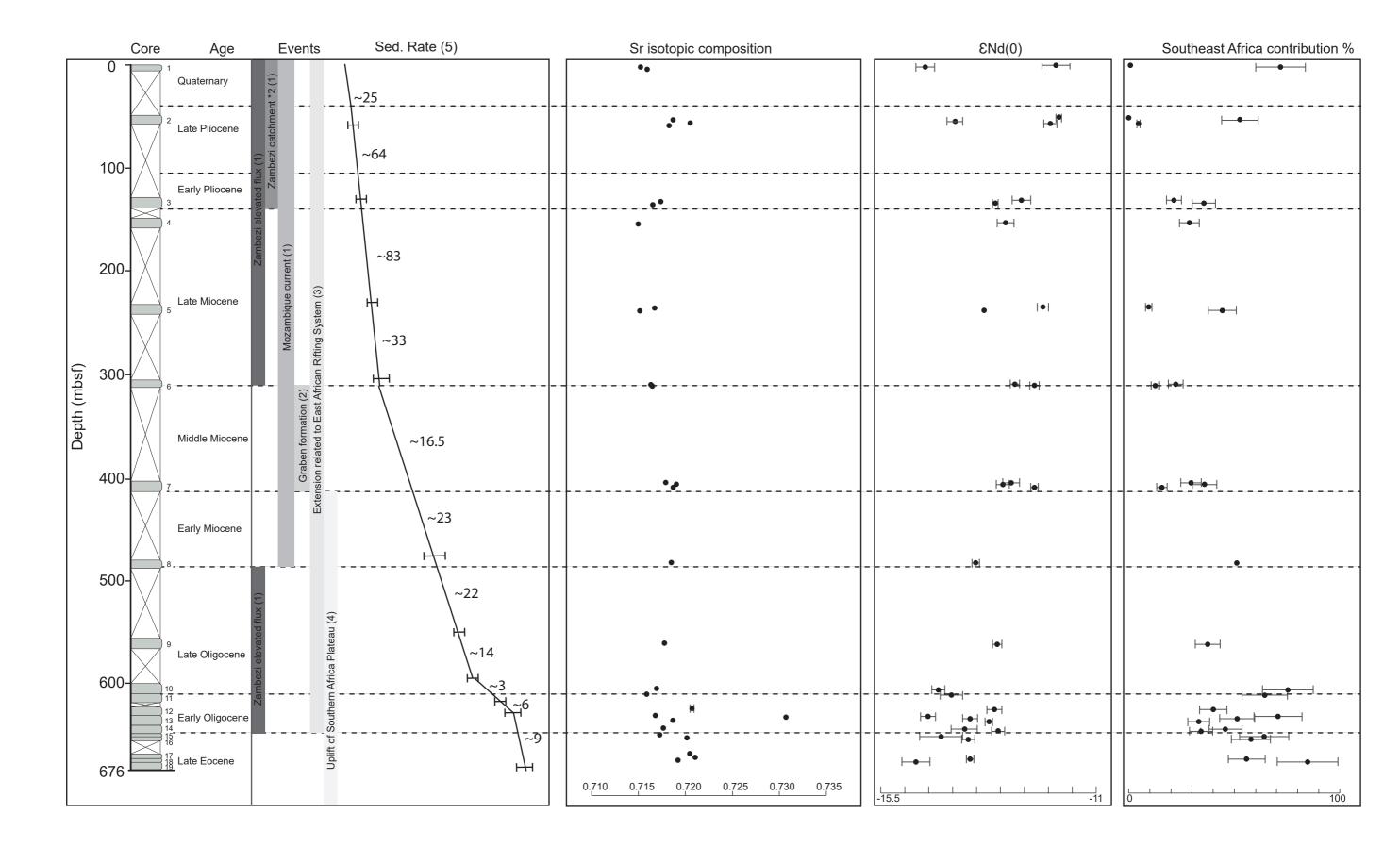












Sample	Core	Age	top CSF-B [m]	bottom CSF-B [m]
1R2W	1	Quaternary	1.3667	1.38
1R4W	1	Quaternary	3.3467	3.36
2R1W	2	Late Pliocene	52.555	52.58
2R3W	2	Late Pliocene	55.56	55.58
2RCC	2	Late Pliocene	58	58
3R3W	3	Early Pliocene	132.06	132.08
3R5W	3	Early Pliocene	135.025	135.045
4R5W	4	Late Miocene	154.06	154.08
5R2W	5	Late Miocene	235.565	235.59
5R4W	5	Late Miocene	238.55	238.57
6R2W	6	Late Miocene	310.04	310.06
6R3W	6	Middle Miocene	311.5	311.52
7R2W	7	Middle Miocene	405.51	405.53
7R3W	7	Middle Miocene	407.02	407.04
7R5W	7	Middle Miocene	409.98	410
8R1W	8	Early Miocene	480.025	480.05
8R3W	8	Early Miocene	483.21	483.23
8R4W	8	Early Miocene	484.5	484.52
8R5W	8	Late Oligocene	486.05	486.07
9R2W	9	Late Oligocene	557.055	557.075
9R5W	9	Late Oligocene	561.51	561.53
10R3W	10	Late Oligocene	605.58	605.6
10R4W	10	Late Oligocene	607.5	607.52
10R5W	10	Late Oligocene	609	609.03
11RCCW	11	Early Oligocene	611	611
12R2W	12	Early Oligocene	625.01	625.035
13R1W	13	Early Oligocene	631.515	631.535
13R2W	13	Early Oligocene	633.48	633.5
13R4W	13	Early Oligocene	636.57	636.59
14R3W	14	Early Oligocene	644.01	644.03
14R4W	14	Early Oligocene	645.93	645.95
15R2W	15	Late Eocene	650.8044	650.8222
16R1W	16	Late Eocene	653.535	653.56
17R1W	17	Late Eocene	668.7067	668.72
18R1W	18	Late Eocene	672.5567	672.5667
19R3W	19	Late Eocene	675.34	675.3533

Table 1: Names, cores, ages and depths of sampled DSDP sediments; top CSF-B [m]: top core depth below sea floor in m; bottom CSF-B [m]: bottom core depth below sea floor in m;

Sample		1R2W 1F	R4W 2	2R1W 2F	R3W 2	RCC 3R	3R5V	V 4F	R5W 5I	R2W 5R4	1W 6	R2W 6F	R3W* 7R2V	/ 7R3W	7R5V	V 8	8R1W 8	R3W 8	8R4W
Age													liddle Miocene Midd						
<u> </u>		, , ,	,				,,										,	,	
SiO2	%	43.07	46.14	47.04	45.62	45	39.13	45.61	44.79	46.4	45.55	46.88	41.16	46.92	45.17	46.86	46.03	44.91	46.2
AI2O3	%	20.51	16.76	19.69	18.58	18.99	16.00	18.14	18.23	18.92	17.57	17.98	16.10	18.77	18.25	18.73	18.27	18.42	18.8
Fe2O3	%	9.62	9.00	9.74	9.79	9.79	8.32	9.92	9.70	10.02	9.83	9.47	8.94	9.63	10.25	10.25	10.63	9.89	10.5
MnO	%	0.06	0.04	0.04	0.04	0.04	0.04	0.04	0.04	0.04	0.04	0.04	0.05	0.05	0.06	0.05	0.06	0.05	0.0
MgO	%	1.89	3.00	2.16	2.26	2.18	2.00	2.34	2.29	2.34	2.42	2.32	2.17	2.25	2.25	2.36	2.47	2.32	2.3
CaO	%	1.15	1.47	1.59	1.19	3.10	9.98	1.23	1.18	1.13	1.13	1.24	7.73	1.49	1.15	1.20	1.07	1.15	1.0
Na2O	%	0.11	0.14	0.21	0.17	0.18	0.16	0.13	0.11	0.11	0.11	0.15	0.14	0.14	0.09	0.13	0.13	0.09	0.1
K2O	%	1.41	2.48	2.27	2.20	2.17	2.02	2.23	2.13	2.16	2.11	2.18	2.05	2.26	2.27	2.45	2.59	2.15	2.2
TiO2	%	0.88	0.85	1.04	0.96	1.00	0.86	0.91	0.90	0.93	0.96	0.97	0.86	0.95	0.90	0.95	0.94	1.00	1.0
P2O5	%	0.16	0.15	0.20	0.19	0.22	0.19	0.19	0.20	0.16	0.17	0.21	0.15	0.15	0.15	0.15	0.10	0.12	0.0
LOI	%	20.18	19.24	17.22	18.41	17.43	21.10	18.45	19.59	18.15	19.30	19.03	20.42	16.86	18.70	16.55	16.82	18.72	16.9
Total	%	99.03	99.28	101.20	99.39	100.09	99.78	99.19	99.17	100.36	99.21	100.47	99.77	99.47	99.25	99.68	99.10	98.80	99.5
CIA		84	75	78	79	70	44	79	80	81	80	79	49	78	80	79	79	80	8
Al/Si		0.54	0.41	0.47	0.46	0.48	0.46	0.45	0.46	0.46	0.44	0.44	0.44	0.45	0.46	0.45	0.45	0.47	0.40
Sc	ppm	22.66	20.44	23.08	20.88	22.26	18.83	20.95	20.44	22.1	20.94	19.53	19.64	22.52	20.79	22.18	23.33	22	24.23
Rb	ppm	113.66	137.11	140.60	150.81	136.90	119.70	145.15	143.40	142.90	137.36	132.73	126.60	136.70	133.39	143.30	147.80	135.45	133.60
Cs	ppm	5.99	6.35	6.21	6.99	6.04	5.19	6.67	6.45	6.12	5.83	6.29	5.11	5.68	5.51	5.94	6.07	6.46	5.91
Ba	ppm	623.32	533.14	640.70	529.20	656.90	501.50	626.35	489.90	543.50	577.99	635.47	622.70	388.30	392.77	323.90	325.40	490.17	377.50
Sr	ppm	86.40	90.32	110.80	88.55	166.50	425.90	100.58	91.53	92.31	88.18	98.68	317.10	108.30	81.78	90.69	86.85	85.45	92.94
Th	ppm	11.23	12.00	13.60	11.07	13.13	11.99	11.72	10.50	14.23	11.60	9.73	11.08	13.69	13.65	14.54	16.58	14.00	15.96
U	ppm	1.66	2.16	1.69	1.92	1.90	1.42	1.60	1.57	1.39	1.17	1.48	1.23	1.47	1.22	1.25	1.18	1.51	1.63
Υ	ppm	22.04	20.95	26.04	21.85	25.82	24.72	24.68	22.48	25.83	23.28	23.70	21.59	23.73	23.08	25.21	27.36	19.99	23.90
Zr	ppm	78.91	67.84	75.21	75.36	75.06	64.25	79.10	72.82	79.31	71.64	78.40	65.03	72.85	71.15	70.27	70.14	66.79	76.93
Nb	ppm	14.94	13.42	15.77	15.95	16.07	14.20	15.86	15.18	19.38	17.51	16.90	17.76	18.53	17.83	18.29	18.99	18.51	20.51
Hf	ppm	2.24	2.06	2.24	2.23	2.24	1.97	2.34	2.19	2.45	2.03	2.21	1.99	2.21	2.10	2.17	2.12	2.05	2.18
Cr	ppm	122.48	130.72	126.70	131.52	129.10	114.80	140.26	136.05	142.20	129.92	132.84	113.20	123.00	122.51	123.60	131.90	140.87	133.00
V	ppm	119.93	127.45	137.20	127.20	137.00	114.30	131.59	126.71	150.00	120.43	127.78	114.10	127.50	113.74	130.40	124.50	149.69	139.60
Co	ppm	17.23	28.75	20.90	20.91	22.25	18.55	22.09	21.13	21.66	20.17	20.60	19.12	31.08	23.51	26.57	28.51	23.72	23.14
Cu	ppm	95.59	45.51	94.61	72.67	85.32	74.68	85.97	87.00	82.45	56.76	66.88	57.90	72.87	65.42	44.22	65.55	45.31	160.80
Ni -	ppm	76.27	78.14	66.29	65.31	70.88	56.78	64.60	63.88	63.66	62.57	61.48	63.59	84.71	75.69	78.44	87.66	79.90	79.63
Zn La	ppm	162.78	173.16 37.05	193.40 42.88	188.87	454.80	130.40 40.47	147.38 41.44	145.18	153.90	143.87	180.17	139.80	200.70 44.44	143.39 47.24	137.80 47.64	166.30	163.92	154.40 47.65
Ce	ppm	32.39 59.82	73.78	73.42	37.79 66.53	44.50 75.69	75.68	73.92	40.50 71.04	45.41 78.39	44.17 79.89	38.88 67.05	38.70 69.57	85.05	90.26	88.82	51.71 103.80	45.38 88.54	95.05
Pr	ppm ppm	6.75	8.62	9.55	8.29	9.81	9.46	9.07	8.72	10.10	9.59	8.28	8.73	10.18	10.49	10.73	11.73	10.16	10.36
Nd	ppm	24.82	31.97	35.14	30.32	36.07	34.66	33.51	32.36	37.24	35.40	30.52	31.77	37.22	38.41	39.60	42.84	36.70	36.92
Sm	ppm	4.77	5.98	6.48	5.69	6.69	6.46	6.40	6.04	6.82	6.67	5.66	5.87	6.95	7.04	7.26	8.04	6.67	6.62
Eu	ppm	1.03	1.15	1.31	1.14	1.35	1.34	1.30	1.20	1.41	1.34	1.19	1.17	1.40	1.37	1.44	1.54	1.28	1.30
Gd	ppm	3.99	4.75	5.11	4.49	5.31	5.12	5.16	4.81	5.64	5.36	4.61	4.67	5.48	5.49	5.81	6.27	5.13	5.19
Tb	ppm	0.60	0.67	0.76	0.66	0.78	0.75	0.74	0.69	0.79	0.75	0.66	0.67	0.79	0.78	0.82	0.91	0.72	0.7
Dy	ppm	3.68	3.84	4.51	3.86	4.53	4.41	4.34	3.98	4.58	4.30	3.94	4.00	4.50	4.51	4.64	5.19	4.09	4.39
Ho	ppm	0.77	0.75	0.91	0.78	0.92	0.88	0.89	0.81	0.92	0.86	0.80	0.79	0.89	0.88	0.92	1.01	0.79	0.8
Er	ppm	2.07	1.88	2.36	2.02	2.35	2.32	2.30	2.10	2.32	2.14	2.02	2.02	2.26	2.18	2.33	2.57	1.94	2.2
Tm	ppm	0.29	0.25	0.34	0.28	0.34	0.32	0.31	0.28	0.31	0.29	0.27	0.28	0.30	0.30	0.32	0.35	0.26	0.3
Yb	ppm	2.01	1.55	2.25	1.87	2.21	2.13	2.05	1.82	1.97	1.82	1.73	1.88	2.04	1.88	1.95	2.25	1.67	1.9
Lu	ppm	0.30	0.22	0.33	0.27	0.33	0.31	0.30	0.27	0.29	0.26	0.27	0.28	0.29	0.27	0.29	0.32	0.23	0.2
Та	ppm	1.16	1.10	1.31	1.29	1.32	1.10	1.25	1.18	1.47	1.36	1.28	1.28	1.40	1.39	1.38	1.43	1.46	1.5
		0.75	0.55	0.77	0.55	0.66	0.74	0.50	0.55	0.50	0.55	0.7:	0.50	0.50	0.67	0.55	0	0.5-	
Eu/Eu*		0.72	0.66	0.70	0.69	0.69	0.71	0.69	0.68	0.69	0.68	0.71	0.68	0.69	0.67	0.68	0.66	0.67	0.6
ΩCe		0.97	0.99	0.87	0.91	0.87	0.93	0.92	0.91	0.88	0.94	0.90	0.91	0.96	0.98	0.95	1.02	0.99	1.0
Cr/Th		10.91	10.89	9.32	11.89	9.83	9.57	11.97	12.95	9.99	11.20	13.65	10.22	8.98	8.98	8.50	7.96	10.06	8.3
Th/Sc		0.50	0.59	0.59	0.53	0.59	0.64	0.56	0.51	0.64	0.55	0.50	0.56	0.61	0.66	0.66	0.71	0.64	0.6

Table 2: Major and trace element concentrations of analyzed sediments. * denotes the samples that may have been affected by incomplete carbonate leaching leading to lower their CIA values

Mag	Sample		8R5W	9R2W	9R5W	10R3W	10R4W	10R5W	11RCCW	12R2W	13R1W	13R2W	13R4W	14R4W	14R3W	15RO2W	16R1W	17R1W	18R1W	19R3W
Single War	-																			
Part										, ,	, ,	, ,	, ,	, ,						
Page	SiO2	%	45.75	46.5	52 45	.16	46 45.81	45.8	1 45.57	46.	22 46.82	47.05	46.37	7 45.58	45.07	45.9	9 46.78	8 46.33	3 47.14	47.1
Mor	AI2O3		18.41	18.3	30 17	.97 17	.24 17.45	17.4	1 16.89	17.	37 17.08	17.11	17.58	18.44	16.11	16.66	6 16.61	1 15.69	16.19	9 15.8
MgO																				
Caco May 1.33																				
NACO W	-																			
No.																				
TOC 10																				
Page																				
Total %																				
Total % 98.59 98.74 99.14 99.34 98.68 98.75 98.59 98.82 100.86 99.47 100.71 100.17 99.41 99.23 99.17 99.47 A/Si CiA 78 79 70 70 70 70 70 70 70 70 70 70 70 70 70																				
CA																				
A/Sc pm 2.2.7 23.12 20.36 2.4.2 24.06 2.3.5 20.97 22.46 20.99 20.84 19.78 20.25 21.22 20.20 23.8 Rb ppm 138.70 137.60 137.60 141.76 140.00 143.00 134.88 110.10 120.90 20.84 19.78 20.25 21.27 20.22 22.85 22.87 Rb ppm 1585 5.79 5.67 6.84 6.22 6.63 6.57 6.47 6.42 6.84 6.62 5.98 6.84 6.50 8.75 9.75 5.53 6.64 6.27 8.90 31.10 11.11 1.17 1.14		/0																		
Sc ppm 22.27 23.12 20.36 21.62 24.06 23.5 20.97 22.46 20.99 20.84 19.78 20.25 21.22 20.22 23.86 21.30 Cs ppm 138.70 137.60 137.64 137.83 140.20 144.30 134.18 140.00 120.64 148.40 126.12 112.33 143.00 133.46 146.70 153.00 Ba ppm 358.60 499.50 516.45 456.47 380.90 391.40 912.63 456.70 402.00 488.20 514.33 779.55 533.50 453.66 407.50 393.90 Th ppm 13.88 13.10 14.33 13.56 14.08 14.93 13.12 16.18 11.02 11.15 13.78 11.24 14.40 11.87 V ppm 21.99 25.04 19.56 23.63 20.26 22.87 22.12 23.90 19.80 22.88 19.30 21.44 23.10																				
Rb ppm 138,70 137,60 137,60 137,60 140,20 144,30 134,18 140,10 120,69 148,40 151,21 112,33 143,00 133,46 146,70 53,00 63 6.64 6.63 6.57 6.47 6.42 6.84 6.62 5.98 6.68 6.33 6.48 6.50 98 Sr ppm 138,60 499,50 756,64 456,47 380,90 391,40 512,63 465,70 402,00 468,20 514,33 579,55 533,50 43,36 407,50 393,90 Th ppm 11.88 13.10 11.43 11.56 14.08 14.43 11.51 11.70 11.71 1.66 1.07 11.74 1.66 1.07 1.42 Y ppm 21.99 25.04 119.56 2.267 2.212 2.290 11.80 1.28 1.11 1.71 1.67 1.66 1.05 1.42 Y ppm 2.213 2.04 2.228	7.1.70.		0.10		.5				02			0.1.2	0.12	0.10	0.12	0	2 0	0.50	0.0.	0.5
Rb ppm 137.09 137.04 137.03 140.20 144.30 134.18 140.10 120.69 148.40 161.22 123.33 143.00 133.46 146.70 153.06 Ba ppm 335.60 499.50 51.645 456.47 380.90 391.40 512.63 456.70 402.00 468.20 514.33 579.55 533.50 453.66 407.50 393.90 Th ppm 11.88 13.10 11.43 11.56 14.08 14.08 11.71 11.71 11.75 11.66 1.00 11.24 11.55 1.77 1.59 1.33 1.51 1.90 1.77 1.18 1.10 11.11 1.14	Sc	ppm	22.27	23.1	12 20	.36 21	.62 24.06	23.	5 20.97	22.	46 20.99	20.84	19.78	3 20.25	21.22	20.22	2 23.86	6 21.87	7 23.37	7 22.7
Ba ppm 338,80 499,50 516,45 456,47 380,90 391,40 512,63 456,70 402,00 468,20 514,33 579,55 333,50 433,66 407,50 393,90 75 75 333,50 438,66 407,50 393,90 75 75 75 333,50 436,66 407,50 393,90 75 75 72 72 74 11 11 190 179 171 187 166 118,71 174 116 11,05 11,42 11,60 11,22 11,05 11,23 11,12 11,18 11,66 118,00 12,64 11,15 11,78 166 18,00 12,67 17,79 10,16 10,55 14,60 11,80 12,28 11,18 11,40 11,80 12,40 13,10 12,28 12,45 14,52 14,32 14,32 14,32 14,32 14,32 14,32 14,32 14,32 14,32 14,32 14,32 14,32 14,32 14,32 </th <th>Rb</th> <th>ppm</th> <th>138.70</th> <th>137.6</th> <th>50 137</th> <th>.04 137</th> <th>.63 140.20</th> <th>144.30</th> <th>134.18</th> <th>140.</th> <th>10 120.69</th> <th>148.40</th> <th>126.12</th> <th>112.33</th> <th>143.00</th> <th>133.46</th> <th>6 146.70</th> <th>0 153.00</th> <th>144.10</th> <th>131.7</th>	Rb	ppm	138.70	137.6	50 137	.04 137	.63 140.20	144.30	134.18	140.	10 120.69	148.40	126.12	112.33	143.00	133.46	6 146.70	0 153.00	144.10	131.7
Sr ppm 104.00 99.63 78.67 91.66 93.88 96.21 96.85 81.88 81.28 86.05 81.01 87.41 174.50 82.95 84.60 134.60 Th ppm 1.28 1.13.0 14.33 13.35 1.42 1.15 1.27 1.19 1.23 1.51 1.90 1.79 1.71 1.87 1.66 1.80 1.76 1.66 1.65 1.66 1.66 1.00 1.62 1.42 Zr ppm 2.19 2.50.4 1.95.6 2.63.8 2.22.87 22.12 2.39.0 1.98.0 2.288 1.93.0 21.44 23.10 22.22.8 27.66 24.62 2.77 7.92.8 70.46 80.09 56.36 58.51 64.01 67.01 62.24 68.30 65.44 72.80 65.31 71.03 66.47 71.39 Nb ppm 20.13 20.35 1.84 1.42.21 1.43.13 1.43.1 1.43.3 13.20	Cs	ppm	5.95	5.7	79 5	.67 6	.84 6.22	6.63	3 6.57	7 6.	47 6.42	6.84	6.62	5.98	6.48	6.33	3 6.43	1 6.50	6.48	6.6
Th	Ва	ppm	358.60	499.5	50 516	.45 456	.47 380.90	391.40	512.63	456.	70 402.00	468.20	514.33	579.55	533.50	453.66	6 407.50	393.90	390.10	453.6
U ppm 1.24 1.56 1.27 1.59 1.71 1.87 1.66 1.80 1.76 1.66 1.05 1.42 Y ppm 1.99 2.504 1.95.6 2.63 20.26 22.87 22.12 2.30 19.80 22.88 1.30 21.44 23.10 22.28 27.66 24.62 2.64 2.24 68.30 65.44 72.80 65.31 71.05 69.47 71.39 Nb ppm 2.18 2.33 2.04 1.85 1.78 1.90 2.00 2.17 1.92 2.18 1.99 2.04 2.04 2.16 2.20 2.14 Cr ppm 124.60 132.00 135.61 137.25 130.50 137.25 130.50 137.25 130.50 137.25 130.50 137.25 125.50 126.43 120.90 137.25 127.00 127.00 127.00 127.00 127.00 127.00 127.00 127.00 127.00 127.00		ppm																		
Y pm 21.99																				
Zr ppm 75.77 79.28 70.46 80.09 56.36 58.51 64.01 67.01 67.02 68.30 65.44 77.80 65.31 71.05 69.47 71.39 Nb ppm 20.13 20.35 18.44 14.21 14.32 14.11 14.69 14.31 14.33 13.20 14.18 14.00 13.44 13.71 13.95 HH ppm 21.18 2.33 2.04 18.5 1.78 1.90 2.00 2.17 1.92 2.18 1.99 2.04 2.04 2.04 2.16 2.20 2.13 19.9																				
Nb ppm 20.13 20.35 18.44 14.21 14.32 14.31 14.51 14.53 13.20 14.18 14.00 13.44 13.71 13.95 Cr ppm 21.46 13.200 12.96 136.43 13.24 137.0 125.00 13.70 125.25 12.64 12.00 12.70 127.00 <th< th=""><th>-</th><th></th><th></th><th></th><th></th><th></th><th></th><th></th><th></th><th></th><th></th><th></th><th></th><th></th><th></th><th></th><th></th><th></th><th></th><th></th></th<>	-																			
Hr ppm 2.18 2.33 2.04 1.85 1.78 1.90 2.00 2.17 1.92 2.18 1.99 2.04 2.04 2.16 2.20 2.14 Cr ppm 124.60 132.00 129.06 136.43 132.40 137.10 135.61 125.00 132.05 137.70 125.5 126.48 120.90 137.22 127.00 127.00 V ppm 121.50 122.00 115.96 137.25 130.50 133.30 124.30 118.20 133.30 124.10 116.02 119.78 121.50 112.22 132.40 118.40 Co ppm 25.64 39.19 23.23 25.61 26.57 26.02 24.29 23.45 23.95 26.54 26.22 24.12 24.42 28.77 24.38 34.66 Cu ppm 83.74 55.63 44.22 54.41 54.62 53.41 45.92 65.63 45.00 55.24 58.82 78.34 60.40 47.69 118.20 140.90 Ni ppm 80.30 104.40 75.74 79.07 79.65 81.82 75.79 73.67 71.72 82.90 75.35 69.75 71.71 87.57 86.41 91.90 Ppm 150.70 160.00 150.15																				
Gr ppm 124,60 132,00 129,06 136,43 132,40 137,10 125,50 132,05 137,70 125,25 126,68 120,90 137,22 127,00 127,00 Co ppm 121,50 122,00 115,96 137,25 135,00 133,30 118,20 130,30 124,10 116,02 119,78 121,50 112,22 132,40 118,40 Co ppm 83,74 55,63 44,22 56,41 56,62 53,41 45,92 65,63 45,00 55,24 58,82 78,34 60,40 47,69 118,20 40,40 40,90 40,34 45,19 45,62 53,41 45,92 65,63 45,00 55,24 58,82 78,34 60,40 47,69 118,20 44,09 Ni ppm 80,30 104,00 150,15 170,59 149,70 155,80 216,52 138,50 143,96 146,60 143,63 134,73 137,60 145,06 160,00 151,30<																				
V ppm 121.50 122.00 115.96 137.25 130.50 133.30 124.30 118.20 130.30 124.10 116.02 119.78 121.50 112.22 132.40 118.40 Co ppm 25.64 39.19 23.23 25.61 26.57 26.02 23.45 23.95 26.54 26.22 24.12 24.42 22.877 24.38 34.66 NI ppm 80.30 104.40 75.74 79.07 79.65 81.82 75.79 73.67 71.72 82.90 75.35 69.75 71.71 87.57 86.41 91.79 Zh ppm 150.70 160.00 150.15 170.59 149.70 155.80 216.52 138.50 143.96 146.60 143.63 134.73 137.60 145.06 160.00 151.30 La ppm 46.34 45.19 44.39 44.27 48.04 412.93 39.35 40.17 35.49 35.72 37.44 28.6<																				
Co ppm 25.64 39.19 23.23 25.61 26.57 26.02 24.29 23.45 23.95 26.54 26.22 24.12 24.42 28.77 24.38 34.66 Cu ppm 83.74 55.63 44.22 54.41 54.62 53.41 45.92 65.63 45.00 55.24 58.82 78.34 60.40 47.69 118.20 44.09 Ni ppm 80.30 104.40 75.74 79.07 79.65 81.82 75.79 73.67 71.72 82.90 75.35 69.75 71.71 87.57 86.41 91.79 Zn ppm 150.70 160.00 150.15 170.59 149.70 155.80 216.52 138.50 143.96 146.60 143.63 134.73 137.60 145.06 160.80 151.30 La ppm 46.34 45.19 45.59 44.39 42.67 46.04 41.29 45.23 39.35 40.17 35.49 35.62 39.25 37.38 45.88 40.82 Ce ppm 89.02 87.47 87.68 86.10 79.35 87.67 82.46 83.48 70.52 74.42 64.85 65.41 73.99 73.44 83.36 74.53 Pr ppm 10.22 10.15 10.14 10.10 9.65 10.30 9.60 10.25 8.77 9.38 7.69 7.89 9.22 87.2 10.35 9.46 Nd ppm 36.98 37.16 36.31 37.18 35.04 37.48 35.67 37.48 35.67 37.17 32.11 34.48 28.27 28.97 33.78 32.40 38.05 34.83 Sm ppm 6.86 6.86 6.52 6.93 6.46 6.94 6.64 6.75 6.00 6.51 5.33 5.41 6.48 6.01 7.18 6.57 Eu ppm 1.33 1.36 1.21 1.30 1.19 1.30 1.27 1.25 1.09 1.22 1.03 1.09 1.22 1.18 1.39 1.27 Gd ppm 5.29 5.44 4.91 5.52 4.88 5.35 5.13 5.33 4.62 5.06 4.23 4.40 4.96 4.82 5.66 5.18 Tb ppm 0.77 0.80 0.70 0.78 0.71 0.78 0.73 0.77 0.65 0.73 0.61 0.63 0.73 0.70 0.84 0.75 Dy ppm 4.29 4.55 4.00 4.52 3.92 4.42 4.22 4.38 3.69 4.15 3.47 3.76 4.22 4.00 4.88 4.41 Ho ppm 0.83 0.90 0.76 0.90 0.75 0.84 0.82 0.85 0.71 0.80 0.68 0.75 0.81 0.79 0.96 0.85 Er ppm 1.90 2.00 1.58 1.96 1.66 1.90 1.74 1.87 1.50 1.65 1.48 1.65 1.84 1.65 1.84 1.68 2.18 1.94 Tm ppm 0.29 0.31 0.25 0.30 0.22 0.27 0.24 0.27 0.29 0.24 0.26 0.24 0.26 0.24 0.26 0.28 0.26 0.33 0.30 Yb ppm 1.90 2.00 1.58 1.96 1.66 1.90 1.74 1.87 1.50 1.15 1.16 1.09 1.13 1.13 1.14 1.12 1.11 Eu/Eu* 0.68 0.68 0.68 0.65 0.66 0.66 0.66 0.66 0.66 0.66 0.66																				
Cu ppm 83.74 55.63 44.22 54.41 54.62 53.41 45.92 65.63 45.00 55.24 58.82 78.34 60.00 47.69 118.20 44.09 Ni ppm 80.30 104.40 75.74 79.07 79.65 81.82 75.79 71.72 82.90 75.35 69.75 71.71 87.57 86.41 91.79 Zn ppm 150.70 160.00 150.15 170.59 149.70 155.80 216.52 188.50 143.96 146.60 143.63 131.473 131.760 145.06 160.00 151.30 Lu ppm 46.34 45.19 45.59 44.39 42.67 46.04 41.29 45.23 39.35 40.17 35.49 35.62 39.25 37.38 45.88 40.82 Pr ppm 10.22 10.15 10.14 10.10 9.60 10.25 8.77 9.38 7.69 7.89 9.22 8.72 37.	•																			
Ni ppm 80.30 104.40 75.74 79.07 79.65 81.82 75.79 73.67 71.72 82.90 75.35 69.75 71.71 87.57 86.41 91.79 Zn ppm 150.70 160.00 150.15 170.59 149.70 155.80 216.52 138.50 143.96 146.60 143.63 134.73 137.60 145.06 160.80 151.30 La ppm 46.34 45.19 45.59 44.39 42.67 46.04 41.29 45.23 39.35 40.17 35.49 35.62 39.25 37.38 45.88 40.82 Ce ppm 89.02 87.47 87.68 86.10 79.35 87.67 82.46 83.48 70.52 74.42 64.85 65.41 73.99 73.44 83.36 74.53 Pr ppm 10.22 10.15 10.14 10.10 9.65 10.30 9.60 10.25 8.77 9.38 7.69 7.89 9.22 8.72 10.35 9.46 Nd ppm 36.98 37.16 36.31 37.18 35.04 6.64 6.94 6.64 6.75 6.00 6.51 5.33 5.41 6.48 6.01 7.18 6.57 Eu ppm 1.33 1.36 1.21 1.30 1.19 1.30 1.27 1.25 1.09 1.22 1.03 1.09 1.22 1.18 1.39 1.27 Ed ppm 5.29 5.44 4.91 5.52 4.88 5.35 5.13 5.33 4.62 5.06 4.23 4.40 4.96 4.82 5.66 5.18 Th ppm 0.77 0.80 0.70 0.78 0.71 0.78 0.73 0.77 0.65 0.73 0.61 0.63 0.73 0.70 0.84 0.75 Dy ppm 4.29 4.55 4.00 4.52 3.92 4.42 4.22 4.38 3.69 4.15 3.47 3.76 4.22 4.00 4.88 4.41 Ho ppm 0.83 0.90 0.76 0.90 0.75 0.84 0.82 0.85 0.71 0.80 0.65 Er ppm 2.11 2.30 1.89 2.25 1.92 2.15 2.06 2.14 1.81 1.98 1.75 1.91 2.06 2.00 2.48 2.20 Tm ppm 0.27 0.30 0.22 0.27 0.24 0.27 0.26 0.27 0.21 0.24 0.26 0.24 0.26 0.24 0.26 0.28 0.26 0.33 0.30 Yb ppm 1.90 0.27 0.30 0.22 0.27 0.24 0.27 0.26 0.27 0.21 0.24 0.22 0.24 0.25 0.26 0.28 0.35 0.87 Eu/Eu* 0.68 0.68 0.68 0.66 0.66 0.66 0.66 0.66																				
Zn ppm 150.70 160.00 150.15 170.59 149.70 155.80 216.52 138.50 143.96 146.60 143.63 134.73 137.60 145.06 160.80 151.30 La ppm 46.34 45.19 45.59 44.39 45.23 39.35 40.17 35.49 35.62 39.25 37.38 45.88 40.82 Ce ppm 89.02 87.47 87.68 86.10 79.38 87.67 82.46 83.48 70.52 74.42 64.85 65.41 73.99 73.44 83.36 74.53 Pr ppm 10.22 10.15 10.14 10.10 9.65 10.30 9.60 10.25 8.77 9.38 7.69 7.89 9.22 8.72 10.35 9.46 Nd ppm 6.86 6.86 6.52 6.93 6.46 6.94 6.64 6.75 6.00 6.51 5.33 5.41 6.48 6.01 7.18 6																				
La ppm 46.34 45.19 45.59 44.39 42.67 46.04 41.29 45.23 39.35 40.17 35.49 35.62 39.25 37.38 45.88 40.82 Ce ppm 89.02 87.47 87.68 86.10 79.35 87.67 82.46 83.48 70.52 74.42 64.85 65.41 73.99 73.44 83.36 74.53 Nd ppm 10.22 10.15 10.14 10.10 9.65 10.30 9.60 10.25 8.77 9.38 7.69 7.89 9.22 8.72 9.38 7.69 7.89 9.22 8.72 9.46 8.86 8.66 6.63 6.52 6.93 6.46 6.94 6.64 6.75 6.00 6.51 5.33 5.41 6.48 6.01 7.18 6.57 Eu ppm 5.29 5.44 4.91 5.52 4.88 5.35 5.13 5.33 4.62 5.06 4.23 4.0																				
Ce ppm 89.02 87.47 87.68 86.10 79.35 87.67 82.46 83.48 70.52 74.42 64.85 65.41 73.99 73.44 83.36 74.53 Pr ppm 10.22 10.15 10.14 10.10 9.65 10.30 9.60 10.25 8.77 9.38 7.69 7.89 9.22 8.72 10.35 9.46 Nd ppm 36.98 37.16 36.31 37.18 35.04 37.48 35.67 37.17 32.11 34.48 28.27 28.97 33.78 32.40 38.05 34.83 Sm ppm 6.86 6.86 6.52 6.93 6.46 6.94 6.64 6.75 6.00 6.51 5.33 5.41 6.48 6.01 7.18 6.57 Eu ppm 1.33 1.36 1.21 1.30 1.19 1.30 1.27 1.25 1.09 1.22 1.03 1.09 1.22 1.18																				
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Tb ppm 0.77 0.80 0.70 0.78 0.71 0.78 0.73 0.77 0.65 0.73 0.61 0.63 0.73 0.70 0.84 0.75 Dy ppm 4.29 4.55 4.00 4.52 3.92 4.42 4.22 4.38 3.69 4.15 3.47 3.76 4.22 4.00 4.88 4.41 Ho ppm 0.83 0.90 0.76 0.90 0.75 0.84 0.82 0.85 0.71 0.80 0.68 0.75 0.81 0.79 0.96 0.85 Er ppm 2.11 2.30 1.89 2.25 1.92 2.15 2.06 2.14 1.81 1.98 1.75 1.91 2.06 2.00 2.18 1.96 0.6 0.30 0.27 0.29 0.24 0.26 0.24 0.26 0.24 0.26 0.28 0.26 0.33 0.30 Yb ppm 1.90 2.0	Eu	ppm	1.33	1.3	36 1	.21 1	.30 1.19	1.30	1.27	7 1.	25 1.09	1.22	1.03	1.09	1.22	1.18	8 1.39	9 1.27	7 1.41	1.2
Dy ppm 4.29 4.55 4.00 4.52 3.92 4.42 4.22 4.38 3.69 4.15 3.47 3.76 4.22 4.00 4.88 4.41 Ho ppm 0.83 0.90 0.76 0.90 0.75 0.84 0.82 0.85 0.71 0.80 0.68 0.75 0.81 0.79 0.96 0.85 Er ppm 2.11 2.30 1.89 2.25 1.92 2.15 2.06 2.14 1.81 1.98 1.75 1.91 2.06 2.00 2.48 2.20 Tm ppm 0.29 0.31 0.25 0.30 0.26 0.30 0.27 0.29 0.24 0.26 0.24 0.26 0.24 0.26 0.28 0.26 0.33 0.30 Yb ppm 1.90 2.00 1.58 1.96 1.66 1.90 1.74 1.87 1.50 1.65 1.48 1.65 1.84 1.	Gd	ppm	5.29	5.4	14 4	.91 5	.52 4.88	5.3	5 5.13	5.	33 4.62	5.06	4.23	3 4.40	4.96	4.82	2 5.66	6 5.18	5.76	5.2
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Er ppm 2.11 2.30 1.89 2.25 1.92 2.15 2.06 2.14 1.81 1.98 1.75 1.91 2.06 2.00 2.48 2.20 Tm ppm 0.29 0.31 0.25 0.30 0.26 0.30 0.27 0.29 0.24 0.26 0.24 0.26 0.24 0.26 0.24 0.26 0.28 0.26 0.33 0.30 Yb ppm 1.90 2.00 1.58 1.96 1.66 1.90 1.74 1.87 1.50 1.65 1.48 1.65 1.84 1.68 2.18 1.94 Lu ppm 0.27 0.30 0.22 0.27 0.24 0.27 0.21 0.24 0.22 0.24 0.27 0.25 0.31 0.28 Ta ppm 1.41 1.52 1.40 1.15 1.17 1.16 1.20 1.15 1.16 1.09 1.13 1.13 1.14 1.12	-	ppm																		
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	ıa	ppm	1.41	1.5	oz 1	.40 1	.15 1.15	1.1	/ 1.16) 1.	20 1.15	1.16	1.09	1.13	1.13	1.14	4 1.12	z 1.1:	1 1.10) 1.1
	Eu/Eu*		0.68	3 0.6	58 N	.65 0	.64 0.65	0.69	5 0.66	5 n	63 0.63	0.65	0.66	0.68	0.66	0.6	7 0.66	6 06	7 0.67	7 0.6
	ΩCe		0.99				.98 0.94													
Cr/Th 8.98 10.08 9.00 10.06 9.40 9.18 10.34 7.73 10.58 8.95 11.37 11.34 8.77 11.05 8.82 9.87																				
Th/Sc 0.62 0.57 0.70 0.63 0.59 0.64 0.63 0.72 0.59 0.74 0.56 0.55 0.65 0.61 0.60 0.59																				

Table 2 (end)

Sample	age	¹⁴⁷ Sm/ ¹⁴⁴ Nd	TDm	¹⁴³ Nd/ ¹⁴⁴ Nd	± 2σ	$\epsilon_{\text{Nd}(0)}$	± 2σ	°′Sr/°°Sr	± 2σ	% SeA	Laboratory
1R2W	Quaternary			0.512031	0.000015	-11.8	0.3	0.715226	0.000009	0.8	GET
1R4W	Quaternary			0.511891	0.000010	-14.6	0.2	0.715925	0.000014	71.9	GET
2R1W	Late Pliocene	0.1085	1.45	0.512034	0.000003	-11.8	0.1	0.71869	0.000010	0.0	Geochronos
2R3W	Late Pliocene	0.1035	1.54	0.511923	0.000008	-13.9	0.2	0.72052	0.000090	52.8	Geochronos
2RCC	Late Pliocene	0.1079		0.512025	0.000007	-12.0	0.1	0.71826	0.000010	4.6	Geochronos
3R3W	Early Pliocene	0.1077	1.5	0.511994	0.000010	-12.6	0.2	0.71737	0.000080	21.4	Geochronos
3R5W	Early Pliocene	0.1060	1.51	0.511966	0.000003	-13.1	0.1	0.7165	0.000020	35.6	Geochronos
4R5W	Late Miocene	0.1012	1.43	0.511977	0.000009	-12.9	0.2	0.714955	0.000015	28.7	Geochronos
5R2W	Late Miocene	0.1069	1.45	0.512017	0.000006	-12.1	0.1	0.71671	0.000010	9.5	Geochronos
5R4W	Late Miocene			0.511954		-13.3	0.0	0.715143	0.000010	44.3	GET
6R2W	Late Miocene	0.1068	1.49	0.511987	0.000005	-12.7	0.1	0.71632	0.000010	22.3	Geochronos
6R3W	Middle Miocene	0.1063	1.46	0.512008	0.000005	-12.3	0.1	0.71647	0.000040	12.6	Geochronos
7R2W	Middle Miocene	0.1068	1.5	0.511983	0.000009	-12.8	0.2	0.7179	0.000010	29.5	Geochronos
7R3W	Middle Miocene	0.1064	1.51	0.511974	0.000007	-13.0	0.1	0.71903	0.000004	35.9	Geochronos
7R5W	Middle Miocene	0.1080	1.48	0.512008	0.000004	-12.3	0.1	0.7187	0.000090	15.7	Geochronos
8R1W	Early Miocene										
8R3W	Early Miocene	0.1005	1.47	0.511945	0.000004	-13.5	0.1	0.71848	0.000020	51.2	Geochronos
8R4W	Early Miocene										
8R5W	Late Oligocene										
9R2W	Late Oligocene										
9R5W	Late Oligocene	0.1059	1.51	0.511968	0.000005	-13.1	0.1	0.71777	0.000080	37.4	Geochronos
10R3W	Late Oligocene			0.511905	0.000007	-14.3	0.1	0.716928	0.000018	75.4	GET
10R4W	Late Oligocene										
10R5W	Late Oligocene										
11RCCW	Early Oligocene			0.511919	0.000012	-14.0	0.2	0.715875	0.000012	64.4	GET
12R2W	Early Oligocene	0.1059	1.51	0.511965	0.000008	-13.1	0.2	0.7207	0.000210	40.1	Geochronos
13R1W	Early Oligocene			0.511894	0.000008	-14.5	0.2	0.71682	0.000018	70.7	GET
13R2W	Early Oligocene	0.1074	1.57	0.511939	0.000008	-13.6	0.2	0.7307	0.000100	51.4	Geochronos
13R4W	Early Oligocene	0.1038	1.49	0.511959	0.000004	-13.2	0.1	0.71866	0.000050	33.2	Geochronos
14R4W	Early Oligocene	0.1076	1.58	0.511933	0.000014	-13.8	0.3	0.717634	0.000012	45.9	Geochronos
14R3W	Early Oligocene	0.1084	1.54	0.511969	0.000007	-13.1	0.1			34.3	Geochronos
15RO2W	Late Eocene			0.511908	0.000023	-14.2	0.4	0.717244	0.000014	64.2	GET
16R1W	Late Eocene	0.1028	1.51	0.511937	0.000007	-13.7	0.1	0.72016	0.000020	57.9	Geochronos
17R1W	Late Eocene							0.72048	0.000020		GET
18R1W	Late Eocene	0.1091	1.59	0.511939	0.000004	-13.6	0.1	0.72104	0.000020	55.9	Geochronos
19R3W	Late Eocene	0.1001	1.00	0.511881	0.000004	-14.8	0.3	0.719207	0.000010	84.7	GET

Table 3: Nd-Sr isotopic systematics of the analyzed sediments; SeA: Southeast Africa; GET: Géosciences Environnement Toulouse and Geochronos;

Age	CIA	N	S.D.	C.I.	Eu/Eu*	N	S.D.	C.I.	ΩCe	N	S.D.	C.I.	Cr/Th	N	S.D.	C.I.	Th/Sc	N	S.D.	C.I.	8 _{Nd(∅)}	N	S.D.	C.I.	87Sr/86Sr	N	S.D.	C.I.	% SeA	S.D.
Quaternary	80	2	7	10	0.69	2	0.04	0.06	0.98	2	0.01	0.02	10.90	2	0.01	0.02	0.54	2	0.06	0.09	-13.2	2	1.9	2.7	0.715576	2	0.000494	0.000685	36.4	50.3
Late Pliocene	76	3	5	6	0.69	3	0.00	0.01	0.88	3	0.02	0.02	10.34	3	1.36	1.54	0.57	3	0.03	0.04	-12.6	3	1.2	1.4	0.719157	3	0.001200	0.001358	19.1	29.2
Early Pliocene	61	2	25	34	0.70	2	0.01	0.02	0.93	2	0.01	0.01	10.77	2	1.70	2.35	0.60	2	0.05	0.08	-12.8	2	0.4	0.5	0.716935	2	0.000615	0.000853	28.5	10.0
Late Miocene	80	4	1	1	0.69	4	0.01	0.01	0.91	4	0.02	0.02	11.95	4	1.66	1.63	0.55	4	0.07	0.06	-12.8	4	0.5	0.5	0.715782	4	0.000865	0.000847	26.2	14.5
Middle Miocene	71	4	15	14	0.68	4	0.01	0.01	0.95	4	0.03	0.03	9.17	4	0.73	0.72	0.62	4	0.04	0.04	-12.6	4	0.3	0.3	0.718025	4	0.001140	0.001117	23.4	11.1
Early Miocene	80	3	1	1	0.67	3	0.01	0.01	1.01	3	0.02	0.02	8.78	3	1.12	1.27	0.67	3	0.04	0.04	-13.5	1	N.A.		0.718480	1	N.A.		51.2	N.A.
Late Oligocene	78	6	2	1	0.66	6	0.02	0.01	0.97	6	0.02	0.01	9.45	6	0.50	0.40	0.62	6	0.05	0.04	-13.7	2	0.9	1.2	0.717349	2	0.000595	0.000825	56.4	26.9
Early Oligocene	74	7	7	5	0.65	7	0.02	0.01	0.94	7	0.03	0.02	9.87	7	1.40	1.04	0.63	7	0.07	0.05	-13.6	7	0.5	0.4	0.720065	6	0.005468	0.004375	48.6	14.6
Late Eocene	72	5	5	4	0.66	5	0.01	0.01	0.93	5	0.03	0.03	9.52	5	0.95	0.84	0.61	5	0.02	0.02	-14.1	4	0.5	0.5	0.719626	5	0.001489	0.001305	65.7	13.2

Table 4: mean and standard deviation (S.D) of selected proxies and isotopic ratios. N: number of sample used to calculate the mean and standard deviation for each age period. C.L.: Confidence Interval calculated using a normal law at 95%. N.A.: not applicable. Sea: Southeast Africa