

1 Volatiles and Redox along the East African Rift

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12 13 **Abstract**

14
15 The upper mantle under the Afar Depression in the East African Rift displays some of the
16 slowest seismic wave speeds observed globally. Despite the extreme nature of the geophysical
17 anomaly, lavas erupted along the East African Rift record modest thermal anomalies. We
18 present measurements of major elements, H₂O, S, and CO₂, and Fe³⁺/ΣFe and S₆₊/ΣS in
19 submarine glasses from the Gulf of Aden seafloor spreading center and olivine-, plagioclase-,
20 and pyroxene-hosted melt inclusions from Ertä Ale volcano in the Afar Depression. We combine
21 these measurements with literature data to place constraints on the temperature, H₂O,
22 and fO₂ of the mantle sources of these lavas, as well as initial and final pressures of melting.
23 The Afar mantle plume is C/FOZO/PHEM in isotopic composition, and we suggest that this
24 mantle component is damp, with 852 ± 167 ppm H₂O, not elevated in fO₂ compared to the
25 depleted MORB mantle, and has temperatures of ~1401-1458°C. This is similar in fO₂ and H₂O
26 to estimates of C/FOZO/PHEM in other locations. Using the moderate H₂O contents of the
27 mantle together with the moderate thermal anomaly, we find that melting begins around 93 km
28 depth and ceases around 63 km depth under the Afar Depression and around 37 km depth
29 under the Gulf of Aden, and that ~1-29% partial melts of the mantle can be generated in these

conditions. We speculate that the presence of melt, and not elevated temperatures or high H₂O contents, are the cause for the prominent geophysical anomaly observed in this region.

Plain Language Summary

The mantle under the Afar Depression and Gulf of Aden, in Northeastern Africa is geophysically distinct from the mantle elsewhere on Earth. Typically, these geophysical distinctions are thought to arise from elevated temperatures, but the composition of lavas erupted in this region demonstrate that the mantle is only moderately warm and cannot fully explain the geophysical nature of the mantle in this region. We produce new measurements of submarine pillow glasses erupted from Erta Ale volcano and find that in addition to being somewhat warm, the mantle in this region is also somewhat hydrated compared to the mantle that feeds mid-ocean ridge volcanoes, but is not substantially different in bulk oxidation state from the mantle. These conditions together produce a region of partial melt that exists between 93 km and 63-36 km depth under the surface. We speculate that this lens of melt can explain the geophysical observations of the mantle in this region.

1. Introduction

Continental rifting is a primary component of the plate tectonic cycle. It records the onset on continental fragmentation and the progression to the production of new oceanic crust and ocean basins. The East African Rift is one modern example that includes incipient continental extension in the southern termini of the Eastern and Western branches, well-developed continental rifting in the Main Ethiopian rift and Afar Depression, and full oceanic spreading and the production of new oceanic crust in the Gulf of Aden (Figure 1). Despite the importance of continental rifting to plate tectonic cycles, the physical mechanisms that drive the initiation and development of continental rifts remain uncertain. The expected magnitudes of the major tectonic forces such as slab pull, asthenospheric drag, and ridge push (Forsyth & Uyeda, 1975) may be insufficient to overcome the expected strength of continental lithosphere, suggesting that continental lithosphere is weakened prior to rifting. One way this could be accomplished is through the injection of magma or other fluids into the continental lithosphere,

and indeed, some continental rifts are associated with significant magmatism at the time of initiation of the rift (e.g., East African Rift and the Ethiopian Flood Basalt Province; Hofmann et al., 1997). However, the production of this magmatism through mantle melting presents new challenges - continental lithosphere ranges from 40 km to 280 km in thickness (Pasyanos, 2010) and there is significantly lower heat flow beneath continents (a mean value of 64.7 mW m^{-2}) than beneath oceans (a mean value of $\sim 95.9 \text{ mW m}^{-2}$; Davies, 2013; Jaupart and Mareschal, 2007). Thus, one would expect limited extents of melting in a mantle at high pressures and cool ambient temperatures as predicted to exist beneath pre-rifted continental lithosphere. This suggests that if continental rifting is magma assisted from the onset, it requires elevated mantle temperatures and/or hydrous and/or carbonated mantle lithologies that melt at lower temperatures than nominally dry, carbon-free peridotite.

The challenge of understanding the role of magmatism in continental rifting is displayed in the East African Rift. Tomographic models of P- and S-wave speeds along the rift present one of the most prominent geophysical anomalies in Earth's upper mantle, with seismic wave speeds of $\delta V_p \sim -6\%$, $\delta V_s \sim -4\%$ relative to standard Earth models (Bastow et al., 2008; Emry et al., 2018). Elevated mantle potential temperatures are expected to slow seismic wave velocities by reducing the shear and bulk moduli of peridotite (Karato & Jung, 2003). If the observed slowness is due to increased mantle temperatures alone, it requires lavas that record mantle potential temperatures near 1700°C (Gallacher et al., 2016). However, the major element compositions of relatively unevolved lavas erupted throughout the Ethiopian/Afar triangle (where seismic wave speeds are slowest) in the last 10 my suggest moderate thermal anomalies of 1490°C (Ferguson et al., 2013; Rooney et al., 2012). This is not only low compared to mantle

potential temperature estimates for the mantle sources of other flood basalts and ocean island basalts which range from nominal ambient mantle temperatures near 1350°C for some Azores lavas to in excess of 1600°C for some Hawaiian lavas (Rooney et al., 2012), but also at odds with a thermal-only explanation for present day geophysical observations of the upper mantle in this region. As suggested by Rooney et al. (2012) the combination of very slow seismic wave speeds and moderate thermal anomaly for the mantle along the East African Rift may require the influence of other factors hypothesized to change the bulk and shear moduli of peridotite, such as melt (Hammond & Humphreys, 2000), H₂O (Karato & Jung, 1998), or high fO_2 (Cline II et al., 2018) in the mantle under the East African Rift.

The H₂O and CO₂ contents and fO_2 of the mantle beneath the ridge axis are largely unconstrained, all properties which influence the extent of melting of peridotite (Dasgupta et al., 2013; Stagno et al., 2013; Till et al., 2012). This uncertainty exists in part because the sources of lavas along the rift are complicated by the potential presence of the depleted upper mantle, material from the Afar plume/African superplume that may extend from the base of the continental lithosphere in this region to the core-mantle boundary (Mulibo & Nyblade, 2013), and contamination by some degree of the assimilation of a wide variety of materials contained within the continental lithosphere that record long histories of plate tectonic cycles (Hutchison et al., 2018). Each of these materials may vary in their H₂O, CO₂ contents and fO_2 , making reasonable predictions of their importance to the observed geophysical characteristics of the upper mantle and the role of each in rifting difficult. Additionally, the volatile elements CO₂ and H₂O are typically quantitatively degassed from subaerially erupted lavas (such as those erupted in the Afar Depression and along the Main Ethiopian Rift, where the geophysical anomaly is

most pronounced), and to constrain the CO₂ and H₂O contents of the undegassed magmas requires either (1) submarine erupted glasses where the confining pressure of the water column limits/prohibits degassing or (2) in the case of subaerially erupted lavas, analysis of naturally glassy phenocryst-hosted melt inclusions. Submarine erupted glasses are rare in continental settings by definition, and melt inclusions are complex, integrated records of (1) the magma from which the phenocryst crystallized and thus the mantle sources of those magmas (e.g., Kelley et al., 2010), (2) crystallization and diffusion processes within the melt inclusion after entrapment in the phenocryst host (Newcombe et al., 2014; Saper & Stolper, 2020), and (3) the evolving host magma composition, which can be communicated through the phenocryst to the melt inclusions by rapid diffusion (Brounce et al., 2021; Bucholz et al., 2013; Humphreys et al., 2022). Further, preservation of naturally glassy melt inclusions is not guaranteed, as many phenocrysts erupt and cool relatively slowly in large volcanoclastic blocks/bombs and/or lava flows, causing the melt pocket contained in the phenocryst to crystallize (Lloyd et al., 2013), at which point spectroscopic measurements of H₂O and CO₂ and Fe³⁺/ΣFe and S⁶⁺/ΣS are not feasible. The result is that there are relatively few datasets available to assess parental and primary melt H₂O, CO₂ and *f*O₂.

The East African Rift is comprised of the Main Ethiopian Rift and Eastern and Western branches. The Main Ethiopian Rift forms a triple junction along with spreading centers in the Red Sea and Gulf of Aden, the latter of which continues eastward and forms the Central Indian Ridge of the Indian Ocean mid-ocean ridge spreading center (Figure 1). The radiogenic isotopic (Sr-Nd-Hf-Pb) compositions of Quaternary-aged Gulf of Aden submarine pillow glasses (dredged by the R/V Vema cruise 33-07; Schilling et al., 1992) and subaerial lavas of the Main Ethiopian

Rift have been used to elucidate the complex contributions from three distinct sources to the magmas that erupt along the Gulf of Aden and into the Main Ethiopian Rift: the depleted upper mantle, the Afar mantle plume, and the Pan-African lithosphere (Rooney et al., 2012; Schilling et al., 1992). Gulf of Aden submarine glasses can be described as predominantly (i.e., >88% contribution) melts of the depleted upper mantle (sample V3307-64D-3g; Schilling et al., 1992) or predominantly (i.e., >98% contribution) melts of the Afar mantle plume (sample V3307-50D-1g; Schilling et al., 1992), all with some small contributions (<5%) of melts of the Pan-African lithosphere. This framework was extended to include the lavas of the Main Ethiopian Rift, where contributions from the Pan-African lithosphere increase, and the influence of the Afar mantle plume appears to have a toroidal surface expression (Rooney, et al., 2012). These glassy pillow basalt samples are critical to constraining the composition, including H_2O , CO_2 , and $f\text{O}_2$, each of the main mantle sources for lavas along the East African Rift, and thus in improving our understanding of the geophysical anomaly present in the upper mantle under the rift.

Here we present new measurements of H_2O , CO_2 , $\text{Fe}^{3+}/\Sigma\text{Fe}$, and $\text{S}^{6+}/\Sigma\text{S}$ of the same submarine Gulf of Aden glasses studied by Schilling et al. (1992) shown in Supplementary Data Table 1, and together with published major and trace element and radiogenic isotopic compositions, place constraints on the H_2O content and $f\text{O}_2$ of the Afar mantle plume (Table 1, Supplementary Data Table 2), depleted upper mantle, and Pan-African lithosphere. We also calculate the temperatures and pressures of melting along the Gulf of Aden, and melt fractions represented by the erupted submarine lavas (Supplementary Data Table 2). We combine these new data on the pillow basalts with new measurements of the major and trace element compositions of naturally glassy, olivine- and plagioclase-hosted melt inclusions and their hosts,

along with dissolved S and H₂O, Fe³⁺/ΣFe, and S⁶⁺/ΣS ratios in the glassy melt inclusions from Erta Ale volcano (Supplementary Data Table 3). We integrate previously collected melt inclusion datasets from the same volcano (de Moor et al., 2013; Field, et al., 2012), and nearby Dabbahu (Field et al., 2012) and Nabro volcanoes (Donovan et al., 2018) to assess the relative importance of various differentiation processes active prior to and during eruption of the host tephra, and to constrain pre-erupted water concentrations of magmas erupted subaerially in the Afar Depression. As for Gulf of Aden submarine glasses, we place constraints on the H₂O and *f*O₂ of the mantle sources of these magmas, temperatures and pressures of melting, and melt fractions represented by the erupted lavas (Table 1; Supplementary Data Table 4). From this combined data set, we assess the importance for the range of the observed slowness and attenuated nature of seismic waves in the region.

2. Geologic Background and prior work

The northern terminus of the East African Rift is where the Main Ethiopian Rift meets the Afar Depression, a broad low-lying land region that includes northern Ethiopia, Djibouti, Eritrea, and northwestern Somalia (Figure 1). The Red Sea spreading center continues away from Afar to the northwest and the Gulf of Aden spreading center continues to the east, where new oceanic lithosphere is actively produced. Though in detail there are complex micro-tectonic processes and structures in this area, the region encompassing the Afar Depression and Red Sea and Gulf of Aden spreading centers is thought to broadly be the final transition away from continental rifting to the development of true oceanic spreading in the eastern Gulf of Aden and the Red Sea. The lavas erupted here have clear trace element and radiogenic isotopic influences

from melts of the depleted upper mantle, the Afar plume, and the Pan-African lithosphere (Hutchison et al., 2018; Rooney et al., 2012); the proportions of each component present have been calculated using three component mixing models and Sr-Nd-Pb isotopic compositions (see Supplementary Data Table 5 for calculation reproduction), and these proportions have been shown to vary spatially in recent magmatism and through time as the rift matured in this region (Rooney, 2020).

Gulf of Aden submarine pillow lavas dredged to the east of 47.1°E have negatively sloped rare earth element patterns (i.e., $\text{La/Sm} < 1$) and Sr-Nd-Pb isotopic compositions that indicate that these lavas are predominantly melts of the depleted mid-ocean ridge mantle (DMM; Schilling et al., 1992). Submarine pillow lavas dredged to the west, between 43.9-46.7°E, have more steeply positively sloped rare earth patterns (i.e., $\text{La/Sm} = 2.5 - 4$), and trace element patterns and Sr-Nd-Pb isotopic compositions that indicate that these lavas are mixtures of melts of the DMM and the mantle endmember C/FOZO/PREMA, thought to be transported into the melting region by the Afar plume (Rooney et al., 2012; Schilling et al., 1992). The lithological identity of the C/FOZO/PREMA mantle endmember is debated, possibly representing portions of the transition zone or lowermost mantle (Hanan & Graham, 1996; Hart et al., 1992; Hauri et al., 1994). It may sample the long-lived residue of the extraction of continental crust from a chondritic mantle (Giuliani et al., 2021), or may contain recycled oceanic lithosphere of a composition that is not currently present at Earth's surface (Castillo, 2015). Whatever the origin of this isotopic endmember, the radiogenic isotopic composition of Gulf of Aden submarine lavas between 43.9-46.7°E, and along the West Sheba Ridge in particular, can be explained as being 25-99% comprised of melts of C/FOZO/PREMA (Schilling et al., 1992). Submarine lavas

further west from 43.9°E require greater contributions from melts of the African lithosphere to explain their flat to gently positively sloped rare earth element patterns and radiogenic Sr and Nd but unradiogenic Pb isotopic compositions (Schilling et al., 1992).

There are three locations in the Afar Depression where melt inclusion studies place constraints on the volatile contents of magmas prior to eruption: Erta Ale, Dabahu, and Nabro volcanoes (Figure 1). Erta Ale is associated with the Red Sea spreading center that extends to the north, though it is offset to the west relative to the Red Sea spreading center by the Danakil Block, which is being rifted from Afar. It is one in a series of aligned stratovolcanoes that mark the edge of the Danakil Block and have trace element and radiogenic isotopic compositions that indicate that the lavas erupted here are predominantly melts of material from the Afar plume, with minimal crustal assimilation and contributions from melts of DMM (Barrat et al., 1998; Rooney, 2020). Two melt inclusion studies from Erta Ale reveal that pre-eruptive magmas have low H₂O and CO₂ contents (<0.13 wt% H₂O, <200 ppm CO₂; de Moor et al., 2013; Field et al., 2012), and suggest shallow crystallization of the host phenocrysts from relatively dry (<0.15 wt% H₂O) magmas near 1150°C and fO_2 of $\Delta QFM \approx 0$.

The lineament of volcanoes in which Erta Ale resides is connected to the Main Ethiopian Rift by a series of rift sectors, one of which includes Dabbahu volcano. Dabbahu lavas range widely in composition from basalt to rhyolite, and the only melt inclusion study available is on the 2011 eruption of evolved magmas with melt inclusion glass compositions containing 68-75 wt% SiO₂ (Field et al., 2012). These glass inclusions contain between 3-5 wt% H₂O and 0-400 ppm CO₂ (Field et al., 2012) and the parental basaltic magma to these evolved compositions is thought to contain <1 wt% H₂O (Field et al., 2012). Magnetite-ilmenite pairs in the basaltic

trachy-andesite lavas from Dabbahu indicate crystallization at $\Delta QFM = 0$ to $+0.7$ (Field et al., 2012).

To the northeast of Dabbahu is Nabro volcano, which sits atop the Precambrian-aged Danakil metamorphic rocks and whose magmas undergo substantial magma mixing in a large crystal mush zone in the crust (Donovan et al., 2018). Basaltic trachyandesitic tephra containing olivine- and plagioclase-hosted melt inclusions are basaltic to trachy-basaltic, with as high as ~ 7 wt% MgO, and record pre-eruptive magmas with between 0.25-2.0 wt% H₂O and 0-3,000 ppm CO₂ (Donovan et al., 2018). The parental magma to these melt inclusion compositions is thought to have major element compositions like those of lavas from the Edd Volcanic Field, that contain approximately 1.3 wt% H₂O, 2000 ppm CO₂, and fO_2 between $\Delta QFM = 0$ to $+0.7$ (Donovan et al., 2018).

3. Samples and Methods

3.1 Sample Descriptions

3.1.1 Erta Ale

The tephra sampled in this study was collected from a cinder/spatter cone during the November 2010 overflow and is the same tephra studied by de Moor et al. (2013). The tephra consists of vesicular, glassy scoria clasts that are < 2 cm in the longest dimension. Olivine, plagioclase, and pyroxene are found throughout the tephra; however, the pyroxene is somewhat less abundant than olivine and plagioclase with $\sim 10\%$ of the crystal load consisting of pyroxene, $\sim 40\%$ olivine, and $\sim 50\%$ plagioclase. Olivine (~ 1 -2mm) in this tephra are subhedral to anhedral and are encrusted with matrix glass. These olivine contain several spherical to oblate,

pale-brown naturally glassy silicate melt inclusions that are ~150-250 μm in diameter. The pyroxene (~1-2mm) are deep green in color, are subhedral to anhedral and are encrusted in matrix glass. Typically, one to two naturally glassy spherical to oblate silicate melt inclusions are found towards the center of each pyroxene grain; these melt inclusions are ~90-250 μm in diameter. The plagioclase (~1-2mm) are milky white in color, and anhedral. These plagioclase grains contain several spherical to oblate pale brown naturally glassy silicate melt inclusions that are ~60-300 μm in diameter.

3.1.2 Gulf of Aden

The samples in this study were dredged in 1976 during R/V Vema cruise 33-07, where sampling of the ridge axis took place along with physiographic, structural, and magnetic anomaly mapping (Schilling et al., 1992). Most of the basalts collected in the Gulf of Aden were from fresh pillows and sheet flows, with fresh glass present on many of the basalts (Schilling et al., 1992). The samples provided consist of naturally glassy, pale brown, submarine glass chips (~2-4mm). Some samples contain small olivine and plagioclase phenocrysts (~60-100 μm), that are anhedral to subhedral.

3.2 Electron probe micro-analysis

Olivine-, plagioclase-, and pyroxene-hosted melt inclusions from Erta Ale and matrix glasses adhered to the outside of these mineral grains were exposed on a single side and polished for electron probe micro-analysis (EPMA) using a JEOL-JXA 8200 Superprobe at the University of California Los Angeles for major element analyses of glass inclusions and their phenocryst hosts. During major element analyses of both the glass and the phenocrysts, the

beam was focused and operated at a current of 15 nA, an accelerating voltage of 15 keV. For measurements of the phenocryst hosts, sodium and potassium were measured first with 10 second peak and 5 second background counting times to minimize alkali loss. Calcium, silicon, and total iron were also measured in the first round with 20 second peak and 5 second background counting times. Titanium, aluminum, manganese, magnesium, and phosphorus were measured in a second round with 20 second peak and 5 second background counting times. For measurements of the glass inclusions and matrix glasses, sodium and potassium were measured first with 20 second peak and 10 second background counting times to minimize alkali loss (i.e., no corrections were made). Calcium, silicon, and total iron were also measured in the first round, with 20 second peak and 5 second background counting times for silicon and 30 second peak and 15 second background counting times for calcium and total iron. Titanium, aluminum, manganese, magnesium, and phosphorus were measured in a second round with 40 second peak and 20 second background counting times.

All data were subject to ZAF correction procedures. Primary calibration standards include forsterite, magnetite, anorthite, Ti-albite, K -feldspar, sphene, manganese, and Durango apatite. The VG-A99 glass was monitored as secondary standard during each run. Sulfur and chlorine were measured separately on Erta Ale glass inclusions and matrix glasses, as well as on chips of Gulf of Aden submarine glasses using a 10 μm beam operated at 80 nA and an accelerating voltage of 15 kV. Both sulfur and chlorine were measured with 100 second peak and 25 second background counting times. The peak position for sulfur was searched for on unknown samples because the position of the k-alpha peak for sulfur is known to vary as the oxidation state of sulfur changes from S^{2-} to S^{6+} (Carroll and Rutherford, 1988). Pyrite, barite, Ba-

Cl apatite, and synthetic BAAP were used as the primary calibration standards. The VG-A99 glass was monitored as a secondary standard during each run. The major element compositions of the olivine, plagioclase, and pyroxene hosts were measured adjacent to the glass inclusions.

3.3 FTIR analysis

After EPMA of melt inclusions and their phenocryst hosts, all sample surfaces were polished to remove possible beam damage within the activation volume of each EPMA spot. Melt inclusions were then polished from the opposite side until doubly exposed, and Gulf of Aden glasses were wafered to thicknesses of 17-125 μm to create wafers with analyzable pools of optically clear glass. All wafered samples were washed gently with acetone to remove epoxy residues. Dissolved H_2O and CO_2 concentrations in glass inclusions and Gulf of Aden submarine glasses were analyzed by Fourier-transform infrared (FTIR) spectroscopy at the University of California, Riverside using a Thermo Scientific Nicolet iS50 Fourier-transform infrared spectrometer with a Nicolet Continuum microscope attachment. Spectra for all samples were collected between 1000 and 6000 cm^{-1} using a tungsten-halogen source, KBr beamsplitter and a liquid nitrogen cooled MCT-A detector. The bench, microscope, and samples were continuously purged by air free of water and carbon dioxide using a Whatman purge-gas generator. Aperture dimensions were selected for each sample depending on the geometry of free glass pathway, ranging in size from 11x14 μm to as large as 100x145 μm . The thicknesses of each sample were measured using a piezometric digimatic indicator with a precision of $\pm 1 \mu\text{m}$.

3.4 XANES analysis

303 The $S^{6+}/\Sigma S$ ratios of melt inclusions and Gulf of Aden submarine glasses, and $Fe^{3+}/\Sigma Fe$
304 ratios of Gulf of Aden submarine glasses, were determined by micro-X-ray absorption near-
305 edge structure (μ -XANES) spectroscopy at beamline 13-IDE, Advanced Photon Source, Argonne
306 National Laboratory. For S measurements, spectra were collected in fluorescence mode from
307 2447 eV to 2547 eV, with a dwell time of two seconds on each point, using a Si [111]
308 monochromator and a defocused beam, with effective diameter of 15 μm . Counts were
309 recorded on a multi-element silicon drift detector X-ray spectrometer, equipped with two Si
310 drift diode detectors. All analyses were done in a helium atmosphere to avoid interaction
311 between the incident photon beam and atmosphere. Incident beam intensity was on the order
312 of 10^7 photons per second per μm^2 , reflecting a balance between the intensity required to
313 produce interpretable S-XANES spectra from materials with low S-abundances (i.e., <2000 ppm)
314 and the mounting evidence that very high photon density fluxes electronically damage Fe and S
315 in silicate materials (e.g., S^{6+} , when present, is reduced to S^{4+} : Brounce et al., 2017; Fe^{3+} is
316 reduced to Fe^{2+} : Cottrell et al., 2018). Each analysis was performed using a stationary beam.
317 Spectral merging, background subtraction, and normalization for these spectra was done using
318 the X-ray absorption spectroscopy data software package ATHENA (Ravel and Newville, 2005),
319 applied uniformly to all spectra so that the region from 2447-2462 eV varies around a value of 0
320 and region from 2485-2457 varies about a value of 1. These normalized spectra were then
321 subject to spectral fitting routines using the Peak ANalysis (PAN) software package. Each
322 normalized spectrum was fit between 2462-2487 eV with four Gaussian curves – one for the
323 background (peak center fixed at 2485 eV) and one each for sulfate (peak center fixed at 2481
324 eV), the broad sulfide feature (peak center fixed at 2477 eV), and the narrow sulfide feature

(peak center fixed at 2470 eV). The integrated $S^{6+}/\Sigma S$ ratios were calculated using the area under the curve of the 2485 eV peak divided by the sum of the areas under the curves of the 2477 and 2485 eV peaks (after Brounce et al., 2022). Alternative methods for calculating $S^{6+}/\Sigma S$ ratios from these spectra (i.e., Nash et al., 2019; Brounce et al., 2017) are provided in the supplement.

For Fe measurements, spectra were collected in fluorescence mode from 7012 eV to 7485 eV using a Si [111] monochromator and a defocused beam diameter of $\sim 10\ \mu\text{m}$. Counts were recorded on a multi-element silicon drift detector x-ray spectrometer, equipped with two Si drift diode detectors. 100 μm of aluminum foil was placed in the path of the incident photon beam to decrease the intensity of the photon beam prior to interaction with the sample surface, which could lead to auto-oxidation of Fe species dissolved in the glass. The incident photon beam intensity resulted in on the order of 2×10^7 photons/second/ μm^2 . The Fe-XANES spectra were normalized, and the pre-edge features were fit following the techniques of Brounce et al. (2017), using two background functions and two Gaussian curves to fit the Fe^{2+} and Fe^{3+} peaks. The calibration glasses of Cottrell et al. (2009) recalibrated according to Zhang et al. (2018) were used to calculate $\text{Fe}^{3+}/\Sigma\text{Fe}$ ratios from the ratio of the areas of the two Gaussian features fit to the pre-edge peaks.

4. Results

4.1 Gulf of Aden

The new Gulf of Aden data presented in this manuscript are available in EarthChem Library (Brounce et al., 2025b) and are available as supplementary data tables in this publication. The major element compositions of Gulf of Aden submarine dredged glasses are

348 reported by Kelley et al. (2013) and summarized here. They are basaltic in composition and
349 range in composition from 6.7-11.7 wt% MgO, 8.6-13.4 wt% FeO*, 10.2-12.8 wt% CaO, 0.04-0.7
350 wt% K₂O, and 0.9-3.4 wt% TiO₂ (Figure 2a-d). The FeO*, K₂O, and TiO₂ contents are loosely
351 negatively correlated with MgO, as expected for magmas with variable extents of crystallization
352 of variable proportions of olivine, plagioclase, and/or pyroxene (Figure 2a, c, d). There is no
353 correlation between CaO with MgO. The K₂O contents of Gulf of Aden glasses vary from values <
354 0.1 wt % to as high as 0.7 wt% K₂O, forming three distinct groups. “Low K₂O” glasses contain less
355 than 0.2 wt % K₂O (light gray box, Figure 2d), “Medium K₂O” glasses contain between 0.2 and
356 0.5 wt % K₂O (medium gray box, Figure 2d), and a single “High K₂O” glass contains 0.7 wt% (dark
357 gray box, Figure 2d). The “Low K₂O”, “Medium K₂O”, and “High K₂O” glasses are each found in
358 specific geographic regions - “Low K₂O” glasses are mostly found east of 49°E, “Medium K₂O”
359 glasses are found between 43°E and 48°E, and the single “High K₂O” glass is found at 46°E. One
360 sample, V60 (indicated by italic font on Figure 2), a glassy fragment recovered by a core aboard
361 R/V Valdivia (originally called sample VA3-302P by Bäcker et al. 1973) and renamed V60 by
362 Schilling et al. (1992) has anomalously high FeO* and TiO₂ compared to the rest of the sample
363 suite and has K₂O contents that put it in the “Medium K₂O” group. The high FeO* and TiO₂
364 suggest higher pressure and lower extent of melting of a mantle source with much lower K₂O
365 contents for this glassy fragment compared to the rest of the suite, and the overall lack of
366 correspondence between K₂O and TiO₂ in the sample suites taken together suggests that the
367 variation in K₂O is not driven by variably extents of melting of a mantle source of constant
368 composition.

369 Our FTIR measurements of these same Gulf of Aden submarine glasses have 0.2-0.8 wt%
370 H₂O, CO₂ from below detection limits via FTIR (i.e., < 30 ppm CO₂ for thinned wafers <75 μm
371 thick, such as used in this study) to 158 ppm CO₂, 800-1300 ppm S, and 20-415 ppm Cl (Figure
372 3a-d; Supplementary Data Table 1). The sulfur and FeO* contents of these glasses cluster
373 around the sulfide saturation curve, consistent with saturation with a free sulfide phase (Figure
374 3d). The sulfur contents of these glasses are uncorrelated with H₂O contents (Figure 3c). The
375 H₂O contents of Gulf of Aden glasses are also uncorrelated with MgO but are positively
376 correlated with K₂O within the “Low K₂O” and “Medium K₂O” groups (Figure 3b).

377 Measured Fe³⁺/ΣFe values range between 0.136-0.189, and S⁶⁺/ΣS values range from
378 0.06-0.27. There are negative correlations between both Fe³⁺/ΣFe and S⁶⁺/ΣS and MgO contents
379 (Figure 4a), consistent with the observed slight increase in *f*O₂ in silicate magmas during low
380 pressure crystallization of olivine +/- plagioclase (Cottrell and Kelley, 2011; Brounce et al., 2014;
381 Brounce et al., 2021; Birner et al., 2018; Shorttle et al., 2015; Le Voyer et al., 2014; O’Neill et al.,
382 2018). There are two glasses with anomalously high Fe³⁺/ΣFe compared to the rest of the sample
383 suite – one is sample V60 and the other is sample V3307-51D-1g (labelled on Figure 4a for
384 clarity). Sample V60 also has the highest FeO* and TiO₂ contents of all the Gulf of Aden glasses
385 and has “Medium K₂O”. Sample V3307-51D-1g is indistinguishable in FeO* and TiO₂ from the
386 other Gulf of Aden glasses, and like V60, has “Medium K₂O”. Measured Fe³⁺/ΣFe ratios are
387 positively correlated with S⁶⁺/ΣS (Figure 4b). Both Fe³⁺/ΣFe and S⁶⁺/ΣS are uncorrelated with their
388 radial distance from Lake Abhe – in particular, most samples have Fe³⁺/ΣFe ~0.147 and S⁶⁺/ΣS
389 ~0.11 (Figure 5). However, samples V60 and V3307-51D-1g have anomalously high S⁶⁺/ΣS and

are higher by ~5 times the standard deviation of the rest of the measurements (standard deviation = +/- 0.03) (Figure 4a).

4.2 Erta Ale

The new Erta Ale data presented in this manuscript are available in EarthChem Library (Brounce et al., 2025a) and are available as supplementary data tables in this publication. Erta Ale melt inclusions are trapped in 2 olivine grains with compositions of Fo79 and Fo80, 10 plagioclase grains that range in composition from An71 to An82, and 5 pyroxene grains that range in composition from Di₈₉ to Di₉₂. The major element compositions of these inclusions were assessed for the effects of post-entrapment crystallization of the host mineral on the edges of the melt inclusions as follows. For olivine grains, we predicted the composition of olivine in equilibrium with our measured melt inclusions assuming $\text{Fe}^{3+}/\Sigma\text{Fe} = 0.16$ (de Moor et al., 2013) and $\text{Fe}^{2+}/\text{Mg } K_D^{\text{ol/liq}}$ as calculated according to Toplis et al. (2005). This yielded and $\text{Fe}^{2+}/\text{Mg } K_D^{\text{ol/liq}}$ of 0.298 and predicted equilibrium forsterite number of 81.0 for Erta Ale-10, compared to measured forsterite number of 80.1 from measurements of the olivine host, and $\text{Fe}^{2+}/\text{Mg } K_D^{\text{ol/liq}}$ of 0.300 and predicted equilibrium forsterite number of 79.2 and 79.5 for Erta Ale-14A and B respectively, compared to measured forsterite number of 79.8 from measurements of the olivine host. We consider these within the range of uncertainties of the value of $\text{Fe}^{3+}/\Sigma\text{Fe}$ for these specific melt inclusions and we opted to apply no correction for post-entrapment crystallization for these inclusions. For pyroxene grains, the Diopside-Hedenbergite component of a modeled pyroxene that is in equilibrium with the measured melt inclusion composition was calculated according to the model of Putirka (1999). This predicted equilibrium composition was then compared to the measured composition of the pyroxene host. Any melt inclusion-host pair

that was >4 units apart was disregarded from further consideration. For plagioclase grains, the anorthite component of a modeled plagioclase that is in equilibrium with the measured melt inclusion composition was calculated using the Post-Entrapment CrystallizationELTS calculator of Kress & Ghiorso (2004). This predicted equilibrium composition was then compared to the measured composition of the plagioclase host. Any melt inclusion-host pair that was >4 units apart was disregarded from further consideration. In this way, we limit the effects of post-entrapment crystallization in our data consideration and narrow our dataset from 51 discrete melt inclusion measurements in 17 grains to 16 discrete melt inclusion measurements in 7 grains.

The major element compositions of the accepted inclusions are basaltic with 4.9-6.8 wt% MgO, 10.5-12.9 wt% FeO*, 9.0-11.3 wt% CaO, 0.6-0.9 wt% K₂O, and 2.1-2.7 wt% TiO₂ (Figure 3a-d; Supplementary Data Table 3). The K₂O, FeO*, and TiO₂ contents of Erta Ale inclusions are negatively correlated with MgO (Figure 3a, b, d), while CaO/Al₂O₃ is positively correlated with MgO (Figure 3c). Matrix glass adhered to the outside of olivine, plagioclase, and pyroxene grains that contain the melt inclusions measured here was also analyzed, and these matrix glass compositions range from 6.2-6.5 wt% MgO, 11.0-12.9 wt% FeO*, 10.6-11.1 wt% CaO, 0.59-0.66 wt% K₂O, and 2.4-2.5 wt% TiO₂ (light green circles Figure 3a-d; Supplementary Data Table 3). There is no distinction in major element compositions of melt inclusions according to the identity of the mineral host (plagioclase, pyroxene, and olivine).

Erta Ale melt inclusions range from 0.05 to 0.4 wt% H₂O and 30 to 1220 ppm S, and CO₂ below detection via FTIR. The S contents of Erta Ale inclusions are uncorrelated, or perhaps loosely negatively correlated, with FeO* as FeO* concentrations range between 10.6 and 12.9

wt% FeO* while S concentrations change by ~25x (Figure 4d). Olivine hosted melt inclusions extend to higher sulfur concentrations (~1218 ppm) than plagioclase or pyroxene hosted inclusions, and pyroxene hosted inclusions have the lowest sulfur concentrations (~136 ppm), overlapping with those of the matrix glass.

The measured $S^{6+}/\Sigma S$ ratios for these melt inclusions range between 0.06 and 0.14, and one measurement of the matrix glass adhered to the outside of an olivine phenocryst containing one of the melt inclusions discussed above has $S^{6+}/\Sigma S$ of 0.17 (Figure 4a; Supplementary Data Table 3). The $Fe^{3+}/\Sigma Fe$ ratios of these inclusions were not measured.

5. Discussion

5.1 Parental magmas for Gulf of Aden and Erta Ale from new measurements

To estimate the effects of fractional crystallization on major element chemistry of Gulf of Aden glasses, we used Rhyolite-MELTS v.1.2.1 (Gualda et al., 2012) at a pressure equal to 300 bar (i.e., the pressure indicated by volatile saturation and their eruption pressure on the seafloor, see next paragraph), a starting fO_2 of $\Delta QFM = 0$ and $H_2O = 0.5$ wt%. At this pressure and starting fO_2 , a modelled melt that begins with a composition equal to that of sample V3307-66D-1g crystallizes olivine, then olivine and plagioclase, then olivine, plagioclase and clinopyroxene, as well as small amounts of spinel and apatite as it cools from a calculated liquidus temperature of 1278 °C to 900 °C. This model (solid curve, Figure 2) is broadly consistent with the measured major element compositions of Gulf of Aden glasses (except for K_2O , see below) and indicates that samples with $MgO > 8.5$ wt% are separated from their parental and primary melt compositions only by crystallization of olivine.

We calculated the pressure of volatile saturation of Gulf of Aden glasses using VolatileCalc2 (Newman & Lowenstern, 2002). For the five glasses where CO₂ contents could be resolved using FTIR (V3307-64D, -66D, -69D, -42D, and -46D), the saturation pressure of the volatile contents (92-357 bars) correspond closely to the pressure of collection on the seafloor (130-355 bars; supplementary data table 1). In the remaining glasses, there were no resolvable CO₂ peaks in the FTIR spectra, and these H₂O-only volatile saturation pressures are much lower than the pressure of collection on the seafloor. This, and positive correlations between H₂O and K₂O in these samples, lead us to assume that, while CO₂ was lost during degassing, significant loss of H₂O from these magmas during volcanic degassing did not occur, following in the style of Dixon and Stolper (1995) on other mid-ocean ridge basaltic magmas. We therefore use the measured values of H₂O and fO_2 as parental magma values from which we calculate primary melt compositions.

We also estimated the effects of fractional crystallization on major element chemistry of Erta Ale glass inclusions using Rhyolite-MELTS v.1.2.1 (Gualda et al., 2012), this time at a pressure equal to 770 bar (see paragraph below), a starting fO_2 of $\Delta QFM = -0.5$ and H₂O = 0.2 wt%. At this pressure and starting fO_2 , a modelled melt that begins with a composition equal to sample BS-h2-MI1 (de Moor et al., 2013) crystallizes as it cools from its liquidus temperature of 1180°C and decompresses to 10 bar, beginning with clinopyroxene and plagioclase, then also olivine at 6 wt% MgO. This model is broadly consistent with the measured compositions of Erta Ale melt inclusions in this study and previous works and indicates that the melts trapped by the inclusions studied here can be produced by 5-41% crystallization from a parental magma similar in composition to sample BS-h2-MI1. We also model the effects of fractional crystallization on

major element chemistry, beginning with a composition equal to sample G-111 (Castillo et al., 2020) which has higher MgO contents than any melt inclusion measured. This model was run under the identical parameters described above except with a starting fO_2 of $\Delta QFM = -0.15$, $H_2O = 0.2$ wt%, and a liquidus temperature of $1198^\circ C$. This melt cools to $900^\circ C$ and decompresses to 10 bar, beginning with the crystallization of olivine. Plagioclase begins to crystallize along with olivine when the melt reaches 7.56 wt% MgO, and clinopyroxene joins the crystallizing assemblage when the melt reaches 7.26 wt% MgO. This model (dashed curve, Figure 2) is consistent with the major element composition of whole rock and melt inclusions from Erta Ale and demonstrates that olivine is the only phase crystallizing from magmas with $MgO > 8.0$ wt% (Figure 2).

To assess the possible variation in magma composition (including fO_2) that would result from degassing, we calculated a degassing trajectory for the same parent magma (BS-h2-MI1; the highest MgO sample measured for Erta Ale; de Moor et al., 2013) one starting with 0.20 wt% H_2O (informed from H_2O measurements of Erta Ale melt inclusions in this study, see supplementary data tables) and one starting with 0.1 wt% H_2O (the highest H_2O measurements from Field et al., 2012), 200 ppm CO_2 , and 1200 ppm S. We chose this volatile composition as most representative of the highest volatile contents measured in melt inclusions at Erta Ale from a combination of studies (this study; de Moor et al., 2013; Field et al., 2012), though it remains unclear whether magmas at depth may have been more volatile rich. We ran the model at a starting fO_2 of $\Delta QFM = -0.5$ (the same fO_2 as used in the crystallization model, corresponding to $Fe^{3+}/\Sigma Fe = 0.135$ and $S^{6+}/\Sigma S = 0.098$) and $1180^\circ C$, neglecting the effect of crystallization on H_2O . Because we have measured $Fe^{3+}/\Sigma Fe$ and $S^{6+}/\Sigma S$ directly in our Gulf of

Aden glasses, following Muth & Wallace (2021) we choose a value for B in $\log K = A/T + B$ for which Sulfur_X (Ding et al., 2023) returned the measured $\text{Fe}^{3+}/\Sigma\text{Fe}$ and $\text{S}^{6+}/\Sigma\text{S}$ of our Gulf of Aden glasses. This results in an expression for the reaction $8\text{Fe}^{3+} + \text{S}^{2-} = \text{S}^{6+} + 8\text{Fe}^{2+}$ of $\log K = -2863/T + 7.5$. This modelled magma composition is vapor saturated at 770 bars, and proceeds to degas CO_2 immediately, then also S beginning substantially near 250 bars total pressure, and H_2O does not much change to 2 bars total pressure at these temperatures and compositions. The $f\text{O}_2$ of this modelled melt decreases slightly from its starting value of $\Delta\text{QFM} = -0.5$ to $\Delta\text{QFM} = -0.65$ (corresponding to $\text{Fe}^{3+}/\Sigma\text{Fe} = 0.127$ and $\text{S}^{6+}/\Sigma\text{S} = 0.05$) by 2 bars total pressure, and degasses S from the residual melt down to 830 ppm S. Our melt inclusion analyses and those of previous studies are consistent with this degassing trajectory with respect to H_2O and CO_2 measurements, but we find that S remains more soluble in the model than measurements suggest. Nonetheless, degassing in these conditions (namely starting at relatively low $f\text{O}_2$, low volatile contents, and low pressures), and like previous studies of Erta Ale magmas (de Moor et al., 2013; Field et al., 2012), we will use the most volatile rich compositions and highest measured $f\text{O}_2$ s as parental melt compositions from which to calculate primary melt compositions and the mantle source.

We note that it is highly likely that all samples measured in this study and prior studies reflect some amount of CO_2 lost from a parental magma to a gas phase. Estimates for the CO_2 content of an undegassed magma in the Afar region are 1000-1200 ppm CO_2 (Gerlach, 1989). The degassing of CO_2 is slightly oxidizing to residual magmas – loss of ~1000 ppm CO_2 has been shown to result in an increase in the residual magma $f\text{O}_2$ by ~0.1 log unit (Brounce et al., 2017). This is small, and we do not correct for it here.

522

523 5.2 Fe-S redox

524 The $\text{Fe}^{3+}/\Sigma\text{Fe}$ and $\text{S}^{6+}/\Sigma\text{S}$ were both measured via XANES in the Gulf of Aden submarine
525 glasses (Figure 4b). The two are positively correlated, however the $\text{S}^{6+}/\Sigma\text{S}$ ratios reported for
526 these Gulf of Aden submarine glasses are higher than recent models predict for a given major
527 element composition, S content, temperature, and $\text{Fe}^{3+}/\Sigma\text{Fe}$ (Boulliung & Wood, 2022; O'Neill &
528 Mavrogenes, 2022; Supplementary Data Table 1). The difference between measured and
529 modeled $\text{S}^{6+}/\Sigma\text{S}$ ratios is large – on average the measured values are 11% (absolute) higher than
530 models predict. However, if one assumes that major element composition, S content, and
531 $\text{Fe}^{3+}/\Sigma\text{Fe}$ are known and temperature is varied, we find that relatively modest changes in
532 assumed temperature away from the MgO magmatic temperature (calculated using Helz and
533 Thornber, 1987) is required to reproduce the measured $\text{S}^{6+}/\Sigma\text{S}$ ratios. All but one sample
534 required a decrease of between 5-49°C relative to the MgO magmatic temperature and the one
535 sample required a 16°C increase. The average change in temperature required across all
536 samples with both $\text{Fe}^{3+}/\Sigma\text{Fe}$ and $\text{S}^{6+}/\Sigma\text{S}$ measurements is a 32°C decrease in the temperature
537 recorded by $\text{Fe}^{3+}/\Sigma\text{Fe}$ and $\text{S}^{6+}/\Sigma\text{S}$ ratios compared to the MgO thermometer magmatic
538 temperature (Supplementary Data Table 1; Fig. 4b). This uncertainty in temperature is small.
539

540 5.3 Primary magmas and mantle sources under the northern terminus of the East African Rift

541 We estimated primary melt compositions, defined as compositions immediately before
542 their segregation from their mantle residues, prior to crystallization-differentiation and
543 degassing, for the Gulf of Aden lavas, Erta Ale, Nabro, and Dabbahu. We used two approaches:

(1) by adding equilibrium olivine back to measured compositions until we obtained melt compositions in equilibrium with olivine of various compositions typically assumed to be representative of mantle peridotite olivine – Fo₈₉, Fo₉₀, and Fo₉₁, and (2) using the PRIMELT-3P software, which combines the inverse model of olivine addition approach with forward models of batch and fractional peridotite partial melting to inform at what extent of olivine addition should the inverse model stop (Herzberg et al., 2023). We describe the results of these calculations and compare them below.

5.3.1 Primary melts from olivine addition

For Gulf of Aden glasses, we incrementally added equilibrium olivine back to the compositions of Gulf of Aden glasses with MgO > 8 wt%. For Fo₉₀ compositions, this required between 2-15% olivine addition and produces model melt compositions with 11.4-13.5 wt% MgO, 8.8-10.4 wt% FeO*, 0.13-0.73 wt% H₂O, and Fe³⁺/ΣFe ratios of 0.136-0.185 (see supplement for full report of these calculations results, and Table 1 for a summary).

Because models of fractional crystallization for Erta Ale magma conditions recorded by melt inclusion and whole rock studies indicate that these magmas are multiply saturated with olivine, plagioclase and/or clinopyroxene below 8.0 wt% MgO (see above, section 5.1), we use the major element composition of whole rock lavas from the GeoROC database for Erta Ale volcano, combined with parental magma H₂O=0.2 wt% and Fe³⁺/ΣFe = 0.145 (constraints from melt inclusions in this study and de Moor et al. 2013; Field et al. 2012) to calculate primary melt compositions, only for GeoROC compositions with >8.0 wt% MgO (Barberi et al., 1971; Barrat et al., 1998). As above, we incrementally added equilibrium olivine back to these compositions

until we obtained melt compositions in equilibrium with Fo₈₉, Fo₉₀, and Fo₉₁ olivine (see supplement for full report of these calculation results). For Fo₉₀ compositions, this required between 5-23% olivine addition and produces model melt compositions with 12.5-15.5 wt% MgO, 9.6-11.6 wt% FeO*, 0.16-0.19 wt% H₂O, and Fe³⁺/ΣFe ratios of 0.117-0.137 (Table 1).

Similarly, we use literature whole rock data for Nabro (De Fino et al., 1978) and Dabbahu (Barberi et al., 1975) for lavas that have compositions that are plausible parental melts for those volcanoes. These samples have greater than 8.0 wt% MgO, and we use the recommendations of melt inclusion studies at each location for parental melt H₂O equal to 1.3 wt% (Nabro; Donovan et al., 2018) and 1 wt% (Dabbahu; Field et al., 2012). Both studies estimate that magmas at each volcanic center crystallized at fO_2 between ΔQFM = 0 and = 0.7, so we calculate primary melts assuming Fe³⁺/ΣFe = 0.145, and Fe³⁺/ΣFe = 0.190. Again, we incrementally added equilibrium olivine back to these compositions until we obtained melt compositions in equilibrium with Fo₈₉, Fo₉₀, and Fo₉₁ olivine (see supplement for full report of these calculation results). For Fo₉₀ compositions at Nabro volcano, in the oxidized scenario (parental melt Fe³⁺/ΣFe = 0.190) this required between 0-16% olivine addition and produces model melt compositions with 10.4-13.9 wt% MgO, 8.2-10.7 wt% FeO*, 1.1-1.3 wt% H₂O, and Fe³⁺/ΣFe ratios of 0.167-0.190 (Table 1). In the reduced scenario (parental melt Fe³⁺/ΣFe = 0.145) this required between 2-18% olivine addition and produces model melt compositions with 11.0-14.6 wt% MgO, 8.2-10.6 wt% FeO*, 1.1-1.3 wt% H₂O, and Fe³⁺/ΣFe ratios of 0.121-0.143 (Table 1). For Fo₉₀ compositions at Dabbahu volcano, in the oxidized scenario (parental melt Fe³⁺/ΣFe = 0.190) this requires 12% olivine addition and produces a model melt composition with 13.7 wt% MgO, 10.9 wt% FeO*, 0.9 wt% H₂O, and Fe³⁺/ΣFe ratios of 0.170 (Table 1). In the reduced scenario (parental melt

Fe³⁺/ΣFe = 0.145) this required 14% olivine addition and produces a model melt composition with 14.5 wt% MgO, 11.0 wt% FeO*, 0.9 wt% H₂O, and Fe³⁺/ΣFe ratios of 0.126 (Table 1).

We use these primary melt compositions to calculate the fraction of melt required to produce those compositions and the H₂O content of the mantle source following methods described by Kelley et al. (2006). The following describes primary melts in equilibrium with Fo₉₀ olivine (see Table 1 for summary); the full details for modeled primary melts in equilibrium with Fo₈₉, Fo₉₀, and Fo₉₁ olivine can be found in the supplemental materials (Supplementary Data Tables 2 and 4). We calculate mantle source TiO₂ for our samples by comparing the TiO₂/Y ratios of our samples to that of MORB, bulk partition coefficient during mantle melting for TiO₂ of 0.04, and an assumed TiO₂ content for DMM of 0.133, following equation 11 from Kelley et al. (2006). Using this approach, the primary melt compositions described in the previous paragraphs correspond to melt fractions of 8-16% for the Gulf of Aden glasses. In the absence of trace element compositions for samples used to constrain primary melt compositions at Erta Ale, Nabro, and Dabbahu volcanoes, we calculated melt fractions H₂O contents of the mantle sources three ways – one assuming the mantle source has a value equal to the lowest calculated mantle source TiO₂ from the Gulf of Aden (0.128 wt%, from sample 64D; supplementary data table 2), one assuming the mantle source has a value equal to DMM (0.133 wt%; Kelley et al., 2006), and one assuming the mantle source has a value equal to the average calculated mantle source TiO₂ of the most Afar mantle plume influenced Gulf of Aden samples (0.191 wt%, from samples 51D, 48D, and 50D; supplementary data table 2). For Fo₉₀ magmas, this resulted in calculated melt fractions of 2-11% for Erta Ale magmas, 1-4% for Nabro magmas, and 1-4% for Dabbahu magmas (Table 1). Using these melt fractions and assuming a bulk partition coefficient

610 during mantle melting for H₂O of 0.012 (Kelley et al., 2006), these calculations suggest that the
611 mantle sources of Gulf of Aden glasses have H₂O contents from 304 ± 105 ppm H₂O to the east
612 of 49°E (i.e., in normal mid-ocean ridge spreading scenario and approaching the Central Indian
613 Ridge), 852 ppm ± 167 ppm H₂O between 45°E and 49°E (i.e., along the West Sheba Ridge), and
614 ~330 ppm H₂O in the Gulf of Tadjoura (i.e., approaching the subaerial Afar Depression; Table 1).
615 For the subaerial volcanic centers, these parameters suggest that the mantle sources of: (1) Erta
616 Ale have H₂O contents of 113 ppm H₂O ± 46 ppm H₂O, (2) Nabro have H₂O contents of 397 ppm
617 ± 152 ppm H₂O, and (3) Dabbahu have H₂O contents of 288 ± 114 ppm H₂O (Table 1).

618 We also calculate the temperatures of these modeled primary melts from MgO contents
619 according to the olivine liquidus relations (Herzberg et al., 2023), as well as the fO_2 indicated by
620 the calculated Fe³⁺/ΣFe ratios at these temperatures and 1 atm pressure (Borisov et al., 2018;
621 Jayasuriya et al., 2004; O'Neill et al., 2018) as well as at 1.5 GPa pressure (Kress & Carmichael,
622 1991; other oxybarometer results can be found in the supplemental materials). East of 49°E
623 along the Gulf of Aden, modeled primary melts have temperatures of 1378°C ± 24° and record
624 fO_2 s of ΔQFM = -0.02 ± 0.12 at 1.5 GPa (Kress & Carmichael, 1991), or ΔQFM = -0.17 ± 0.11 at 1
625 atm (Borisov et al., 2018; Table 1). Between 45°E and 49°E, temperatures and fO_2 s of modeled
626 primary melts increase somewhat, to 1401°C ± 33° and ΔQFM = +0.20 ± 0.43 at 1.5 GPa (Kress &
627 Carmichael, 1991), or ΔQFM = -0.03 ± 0.51 at 1 atm (Borisov et al., 2018), driven strongly by
628 sample V3307-51D-1g (Table 1). In the Gulf of Tadjoura, the temperature and fO_2 of the one
629 modeled primary melt in this study in this location drops to 1387°C and ΔQFM = +0.08 ± 0.43 at
630 1.5 GPa (Kress & Carmichael, 1991), or ΔQFM = -0.11 at 1 atm (Borisov et al., 2018). For the
631 subaerial volcanic centers, modeled primary melts record temperatures and fO_2 s of 1412°C ±

25° and $\Delta\text{QFM} = -0.08 \pm 0.08$ at 1.5 GPa (Kress & Carmichael, 1991), or $\Delta\text{QFM} = -0.39 \pm 0.16$ at 1 atm (Borisov et al., 2018) at Erta Ale, and $1375^\circ\text{C} \pm 43^\circ$ and $\Delta\text{QFM} = +0.30 \pm 0.41$ at 1.5 GPa (Kress & Carmichael, 1991), or $\Delta\text{QFM} = +0.04 \pm 0.47$ at 1 atm (Borisov et al., 2018) at Nabro (Table 1). For Dabbahu, we calculated primary melts assuming a parental melt at $\Delta\text{QFM} = 0$ and $\Delta\text{QFM} = +0.7$. Accordingly, the temperatures calculated for primary melt from the one sample available for this calculation with $\text{MgO} > 8.0$ wt% are 1402°C ($\Delta\text{QFM} = 0$) or 1422°C ($\Delta\text{QFM} = +0.7$), and the primary melt $f\text{O}_2$ is either $\Delta\text{QFM} = 0.09$ at 1.5 GPa (for parental melt $\Delta\text{QFM} = 0$) (Kress & Carmichael, 1991), or $\Delta\text{QFM} = 0.85$ at 1.5 GPa (for parental melt $\Delta\text{QFM} = +0.7$; Table 1). Using the Borisov calibration, the primary melt $f\text{O}_2$ at 1 atm would be $\Delta\text{QFM} = -0.37$ (for parental melt $\Delta\text{QFM} = 0$) or $\Delta\text{QFM} = 0.39$ (for parental melt $\Delta\text{QFM} = +0.7$; Table 1).

5.3.2 Primary melts from PriMELT3

We use the same samples as described in the previous section in the excel calculator to constrain the composition of primary melts using PriMELT-3P (Herzberg et al., 2023). Importantly, though we have tried to identify and avoid compositions that are multiply saturated with olivine +/- pyroxene +/- plagioclase, the PriMELT3 calculator identifies several samples from which there is evidence of pyroxene fractionation at high pressures as indicated by inappropriately low CaO at a given MgO concentration. This warning eliminates the single sample constraint for Dabbahu, but constraints for the Gulf of Aden submarine pillows, Erta Ale, and Nabro remain.

The PriMELT-3P calculations proceeded in general with greater extents of olivine addition to olivine compositions of higher fosterite content, yielding primary melt compositions

654 with higher MgO. For example, for Gulf of Aden glasses, in the calculation combined with batch
655 melting forward model, PriMELT-3P proceeded with 16-24% olivine addition until equilibrium
656 with $\text{Fo}_{91}\text{-Fo}_{92}$ olivine was reached. This produced primary melts with 14-18 wt% MgO, 8.9-10
657 wt% FeO^* , and $\text{Fe}^{3+}/\Sigma\text{Fe}$ ratios of 0.107-0.150 (compare to 2-15% olivine addition to obtain a
658 melt with 11.6-13.8 wt% MgO, 8.8-10.4 wt% FeO^* , and $\text{Fe}^{3+}/\Sigma\text{Fe}$ ratios of 0.136-0.185 for the
659 same parental/initial melt compositions when stopping at Fo_{90} as described in the previous
660 section; Table 1). The batch melting approach by PriMELT-3P predicts substantially higher
661 degrees of melting – 20-29% melting of harzburgitic mantle source – than calculated using
662 primary TiO_2 as described in the previous section (8-15% melting; Table 1). When combined
663 with a fractional melting forward model, PriMELT-3P proceeded with somewhat less olivine
664 addition – 13-19% - and returns primary melt compositions with 13-16 wt% MgO, 8.9-10 wt%
665 FeO^* , and $\text{Fe}^{3+}/\Sigma\text{Fe}$ ratios of 0.111-0.156. Whether combined with batch or fractional melting
666 forward models, because PriMELT-3P predicts primary melts with higher MgO contents than
667 olivine addition to assumed olivine compositions, it also calculates higher temperatures for
668 primary melts along the Gulf of Aden of $1469 \pm 43^\circ\text{C}$ (batch melting) or $1431 \pm 32^\circ\text{C}$ (fractional
669 melting). At these conditions, PriMELT-3P predicts that melting under the spreading ridge of the
670 Gulf of Aden begins at 2.8 ± 0.3 GPa ($\sim 85 \pm 9$ km depth) and stops at 1.3 ± 0.3 GPa ($\sim 40 \pm 9$ km
671 depth; Figure 6, Table 1). These pressures for the start of melting are very close to the spinel-
672 garnet transition (Figure 6), though most Gulf of Aden samples do not display obvious signs of
673 garnet as a residual phase during melting (Figure 7), e.g., they have flat sloped heavy rare earth
674 element patterns. The exception to this is sample V60, which has a Dy/Yb_N value of 1.5. This
675 sample however does not pass our filtering methods for calculating primary melt compositions,

and thus is not involved in calculating the initial pressures of melting described in this paragraph.

The PriMELT-3P calculation similarly proceeded with greater extents of olivine addition to higher forsterite number olivines for Erta Ale. In the case of combining olivine addition with the forward batch melting model, 8-30% olivine addition was done, yielding primary melts with 13.4-18.9 wt% MgO, 9.8-10 wt% FeO*, and $\text{Fe}^{3+}/\Sigma\text{Fe}$ ratios of 0.108-0.133 (compare to 5-23% olivine addition and model melt compositions with 12.5-15.5 wt% MgO, 9.6-11.6 wt% FeO*, and $\text{Fe}^{3+}/\Sigma\text{Fe}$ ratios of 0.117-0.137 for the same parental/initial melt compositions when stopping at Fo₉₀ as described in the previous section; Table 1). These compositions are obtained through 21-28% melting of harzburgitic mantle source. When combined with a fractional melting forward model, PriMELT3 added 8-24% olivine to the parental/initial compositions, yielding primary melt compositions with 13-17 wt% MgO, 9.8-10 wt% FeO*, and $\text{Fe}^{3+}/\Sigma\text{Fe}$ ratios of 0.112-0.132, obtained through 19-26% melting of a harzburgitic mantle source (Table 1). These primary melts yield higher temperatures than the method described in the previous section of $1458 \pm 35^\circ\text{C}$ (Table 1). At these conditions, PriMELT-3P predicts that melting under Erta Ale begins at 3.1 ± 0.3 GPa (93 ± 10 km depth) and stops at 2.1 ± 0.2 GPa (63 ± 7 km depth; Figure 6; Table 1).

For Nabro, the two styles of calculation are more similar in part because there are fewer samples to constrain the parental melt composition, and as is the case with the olivine addition method described in the previous section, the result of the PriMELT-3P calculations depends on the $\text{Fe}^{3+}/\Sigma\text{Fe}$ of the parent/initial magma composition, which changes depending on the $f\text{O}_2$ of the parent magma within the reported range of $\Delta\text{QFM} = 0$ to $+0.7$. In the case that the parent

magma has fO_2 of $\Delta QFM = 0$, PriMELT3 proceeds with 0-17% olivine addition to reach olivine compositions of Fo90.2-90.7. This results in primary melts with 10.5-14.7 wt% MgO, 8.6-10.8 wt% FeO*, and $Fe^{3+}/\Sigma Fe$ ratios of 0.158-0.190 (compare to 2-18% olivine addition and melt compositions with 11.0-14.6 wt% MgO, 8.2-10.6 wt% FeO*, and $Fe^{3+}/\Sigma Fe$ ratios of 0.121-0.143 for the same parental/initial melt compositions when stopping at Fo90 as described in the previous section, Table 1). In the case that the parent magma has fO_2 of $\Delta QFM = +0.7$, PriMELT-3P adds somewhat more olivine (0-21%) to reach somewhat more forsteritic mantle olivine compositions (Fo90.4-90.9), resulting in primary melts with somewhat higher MgO (11.3-15.7 wt% MgO) and lower $Fe^{3+}/\Sigma Fe$ ratios (0.116-0.145; Table 1). These primary melts can be obtained by 7-13% melting of a harzburgitic mantle source at $1384 \pm 66^\circ C$ (Table 1). PriMELT-3P predicts a narrow range for melting, beginning at 2.3 ± 0.6 GPa (71 ± 18 km depth) and stopping within the uncertainty of 18 km (Figure 6), for both the lower and higher estimates parent magma starting fO_2 s (assuming the parent/initial magma has fO_2 of $\Delta QFM = +0.7$ yields estimates of the pressure of the start of melting of 2.5 ± 0.6 GPa).

5.4 The H_2O and fO_2 of the Afar plume

To summarize, the Gulf of Aden samples collected east of $49^\circ E$ have Sr-Nd-Pb-Hf isotopic compositions that are characteristic of the depleted upper mantle (Schilling et al., 1992). Our new measurements of H_2O , $Fe^{3+}/\Sigma Fe$, and $S^{6+}/\Sigma S$ indicate that these samples, specifically their modeled primary melts, have low water contents (calculated via method 1 described above 0.27 ± 0.09 wt% H_2O), and $Fe^{3+}/\Sigma Fe$ characteristic of MORB primary melts (0.126 ± 0.01 using method 1, or 0.112 ± 0.01 using method 2). The Gulf of Aden samples collected between $45^\circ E$ and $49^\circ E$

have Sr-Nd-Pb-Hf isotopic compositions along with enriched trace element patterns that have been interpreted as arising due to major contributions in their mantle sources from the Afar mantle plume and minor contributions from the Pan African lithosphere and the depleted upper mantle (Schilling et al., 1992). The modeled primary melts for these samples have elevated concentrations of H₂O (0.6 ± 0.11 wt% H₂O calculated via method 1 described above), but Fe³⁺/ΣFe still characteristic of MORB primary melts (0.140 ± 0.02 using method 1, or 0.132 ± 0.02 using method 2), corresponding to *f*O₂s near the QFM oxygen buffer. These redox measurements indicate that the Afar mantle plume is not substantially different in *f*O₂ from that of DMM (Zhang et al., 2018; O'Neill et al., 2018).

Using the radiogenic isotopic studies of Schilling et al. (1992) and Rooney et al. (2012) as templates and using constraints on primary melt compositions from our new measurements of H₂O and Fe³⁺/ΣFe of Gulf of Aden glasses, we can calculate H₂O contents and Fe³⁺/ΣFe ratios of the depleted upper mantle, the Afar mantle plume, and the Pan-African lithosphere. We use the same endmember compositions as Rooney et al. (2012), which differ from those of Schilling et al. (1992) in the composition of the isotopic endmember of the Afar plume, which is now thought of as a C/FOZO/PHEM plume (Figure 8). We calculate that the isotopic compositions of Gulf of Aden glasses east of 49°E require 79-88% contribution from melts of the depleted upper mantle, 10-17% contribution from melts of the Afar plume, and 2-4% contribution from melts of the Pan African lithosphere (Supplementary Data Table 5). The contributions from the Afar plume increase and contributions from the depleted upper mantle decrease in some samples collected between 45°E and 49°E, requiring 0-70% contribution from melts of the depleted upper mantle, 26-100% contribution from melts of the Afar plume, and 0-5% contribution from

melts of the Pan African lithosphere. Importantly, the isotopic composition of sample V3307-50D-1g can be entirely described as a melt of the Afar plume with a composition indicated by Rooney et al. (2012) and from this we assign the H_2O and $\text{Fe}^{3+}/\Sigma\text{Fe}$ of the modeled primary melt from this sample as representative of primary melts of the Afar plume (Table 1). These values are 0.7 wt% H_2O and $\text{Fe}^{3+}/\Sigma\text{Fe} = 0.123$ (Table 1), both calculated using method 1, corresponding to an $f\text{O}_2$ of $\Delta\text{QFM} = -0.27$ using Borisov et al. (2018). Combined with the estimates of ~13% melting of the mantle to produce this primary melt composition, estimated from primary melt TiO_2 contents as described above (see above, *5.3.1 Primary magmas and mantle sources*) and the simple batch melting equation, this suggests that the Afar plume contains ~1082 ppm H_2O .

Because PriMELT-3P requires greater extents of olivine addition to reach equilibrium with higher forsterite number olivine to satisfy both the inverse olivine addition and forward mantle melting models simultaneously, it is likely that the primary melt H_2O contents calculated using method 1 are higher compared to the PriMELT-3P approach. If we take sample V3307-51D-1g, which passes PriMELT3 calculation requirements, and continue adding olivine as in method 1 to the PriMELT-3P suggested olivine composition of Fo_{92} , we obtain a primary melt composition with 0.91 wt% TiO_2 and 0.45 wt% H_2O (compare to the Fo_{90} composition stopping point from method 1 of 1.00 wt% TiO_2 and 0.50 wt% H_2O). Following the same approach to calculate degree of melting and mantle source water contents described above, this yields a mantle source with 924 ppm H_2O (Table 1). This is lower but not substantially different (i.e., does not lead to large differences in interpretation of the tectonic setting) than the estimate of 1082 ppm H_2O , obtained using sample V3307-50D-1g, for which PriMELT-3P indicates the CaO content of the primary melt is too low to be both derived from peridotite and experienced only

764 olivine fractionation prior to eruption, and method 1 for primary melt calculations. This
765 illustrates well the level of uncertainty in various approaches to the “primary melt problem” and
766 using erupted basaltic liquids to place constraints on mantle rock compositions.

767 We can also compare H₂O/Ce ratios of samples in this study (133-537; Figure 5d) to
768 those from other locations. The H₂O/Ce of MORB range between 150-500 (Dixon et al., 2002,
769 2017; Michael, 1995; Wang et al., 2021), with the highest of these values occurring in the
770 Southwest Indian Ridge (Wang et al., 2021). The highest H₂O/Ce ratios in Gulf of Aden samples
771 in this study occur to the east of 49°E, in samples that are far from the influence of the Afar
772 mantle plume. These high H₂O/Ce ratios are driven by low Ce concentrations that are not
773 accompanied by depletions in H₂O. The reason for these high H₂O/Ce ratios in MORB has been
774 proposed to be related to ancient subduction zone mantle wedge material in the melting region
775 of mid-ocean ridge spreading centers (Wang et al., 2021), and warrant further study in the
776 context of mid-ocean ridge source mantle. Here, we focus on the H₂O/Ce ratios of the samples
777 most influenced by the Afar mantle plume (i.e., from radiogenic isotopes require 47-99% of the
778 Afar plume; 48D-2, 50D-1, and 51D-1; Figure 5), which have H₂O/Ce ratios of 234-244. These
779 H₂O/Ce ratios are similar to values for the isolated component FOZO at Hawaii (~200; Shimizu et
780 al., 2019), the Azores mantle plume (210-279; Dixon et al., 2002, 2017; Asimow et al., 2004),
781 and the Easter Salas y Gomez mantle plume (223; Simons et al., 2002). Using the batch melting
782 equation, bulk D_{Ce} = 0.01 (following from Wang et al., 2021), and melt fractions calculated in
783 this study (from Fo₉₀ olivine addition calculations), these H₂O/Ce values predict mantle source
784 H₂O contents of 904 ppm (51D-1), 851 ppm (48D-2) and 1182 ppm (50D-1), and an average of

979 ppm H₂O. In summary, our full range of constraints on the H₂O content of the Afar mantle plume is 697-1182 ppm H₂O, or 951 ± 169 .

5.5 Geophysical and Geochemical models of the mantle

As described in the introduction, the mantle under the East African Rift presents one of the most prominent geophysical anomalies present in the upper mantle and it has been challenging to understand the (1) the primary reasons for these anomalous seismic wave behaviors and (2) the importance of the characteristics of the anomaly to rifting strong continental lithosphere more broadly.

We present constraints on the mantle potential temperature, water content, fO_2 , degrees of melting and initial and final pressures of melting for the Afar Depression, where the anomaly is the most prominent, in the preceding sections. The average of all constraints (Gulf of Aden, Erta Ale, Nabro, and Dabbahu) yield potential temperatures of $1458 \pm 68^\circ\text{C}$, in good agreement with previous studies in this area (Figure 6). This is warmer than ambient mantle that feeds the typical, global mid-ocean ridge spreading center system ($1280\text{-}1400^\circ\text{C}$ using the model presented here, Herzberg et al., 2023, Figure 6), but not remarkably hot in the context of global mantle plume potential temperatures (e.g., the Hawaiian mantle plume is estimated to be 1510 to nearly 1600°C ; Rooney et al., 2012; Figure 6). Using the Gulf of Aden submarine glasses and three component mixing models to match radiogenic isotopic compositions, we suggest that the Afar plume has $\sim 852 \pm 167$ ppm H₂O – higher than estimates for the combined mantle sources of Hawaiian lavas, which vary from 350-450 ppm H₂O (Wallace, 1998, Dixon et al., 2008) and substantially lower than e.g., the mantle wedge in subduction zones, which have

2000-8000 ppm H₂O (Kelley et al., 2010). The estimate of $\sim 852 \pm 167$ ppm H₂O for the Afar mantle plume is broadly consistent with estimates for the isolated FOZO/C/PHEM mantle component in other FOZO-dominant plume or plume component of 620-920 ppm, including Jan Mayen, Iceland, Azores, Easter Salas y Gomez, and the FOZO component in the Hawaiian plume (Asimow et al., 2004; Dixon et al., 2017; Nichols et al., 2002; Shimizu et al., 2019; Simons et al., 2002). We show that existing melt inclusion constraints on the pre-eruptive water contents of subaerially erupted lavas in the Afar Depression suggest that the combined mantle sources of recently erupted Erta Ale, Nabro, and Dabbahu lavas (mixtures of the depleted upper mantle, the Afar plume, and the Pan-African lithosphere) have 100-300 ppm H₂O, intermediate between values for DMM (50-100 ppm H₂O; Shimizu et al., 2019; Dixon et al., 2008) and the Afar plume (this study). Additionally, we show that the mantle sources of all samples studied here have fO_2 at or ~ 0.25 orders of magnitude within $\Delta QFM = 0$. PRiMELT-3P predicts that melting begins as deep as ~ 93 km depth and proceeds as shallow as ~ 63 km depth under Afar and ~ 37 km depth under the Gulf of Aden (Figure 6). We emphasize that the mantle temperatures for melt generation, mantle source H₂O contents, and mantle source fO_2 constraints presented here are not extreme examples of these values in the upper mantle in any tectonic setting, and thus can be eliminated as the sole explanation for the extremely slow seismic wave speeds. Of variables otherwise suggested to impact the bulk and shear moduli of the Earth's mantle and thus impact seismic wave behaviors, we are left to evaluate the role of the presence of partial melt.

These results are broadly consistent with other geochemical models of melting in the region, based on major and trace element compositions of lavas erupted to the surface in the East African Rift (Rooney et al., 2012; Ferguson et al., 2013; Beccaluva et al., 2009; Furman et

829 al., 2016.). We highlight key points from Figure 6: 1) The initial pressures of melting at Erta Ale
830 and the most heavily plume-influenced Gulf of Aden spreading center ridge segment occur
831 slightly deeper and cooler than the dry peridotite solidus. This requires that melt generation is
832 fundamentally driven by the presence of fusible mantle components, in this case, the Afar
833 plume transporting some material that contains higher-than-typical H_2O contents; 2) Under the
834 Gulf of Aden spreading center, melting under the most heavily plume-influenced ridge segment
835 proceeds to shallower depths (~27 km depth, pink circle, Figure 6) than under the ridge
836 segments not influenced by the Afar plume (~42 km depth, black circle, Figure 6). The Afar
837 plume has a clear role in enabling melt generation to lower pressures within the oceanic
838 spreading regime; 3) At Erta Ale, where the Afar plume is present like it is under the heavily
839 plume-influenced segment of the Gulf of Aden spreading center, but melt generation occurs
840 under the continental lithosphere, melting stops at deeper depths (~63 km, green circle, Figure
841 6) than any location under the Gulf of Aden spreading ridge. The lithosphere, or some other
842 thermomechanical boundary, stops melting at rather high pressures. These observations outline
843 a clear role of the Afar plume in enabling melt generation. These results also support
844 hypotheses that there is a thick and diffuse thermomechanical boundary layer between the
845 asthenosphere and lithosphere and that this may explain for instance observed differences in
846 the position of the lithosphere-asthenosphere boundary as placed by seismic tomography (60
847 km depth; Emry et al., 2018) and receiver functions (30 km depth; Rychert et al., 2012). Our
848 results suggest that a ~30 km thick melt-rich (~10-20% melt fraction) layer exists under the Afar
849 Depression beginning ~93 km depth (the predicted depth of the start of melting under Erta Ale,
850 Table 1) and extending up the final depth of melting of ~63 km at Erta Ale, and about ~27 km in

the Gulf of Aden where the Afar mantle plume has the strongest influence, and ~42 km in the Gulf of Aden far from influence from the Afar mantle plume today (Table 1). This final depth of melting in each region may correspond to the lithosphere-asthenosphere boundary in the region or may reside below that boundary.

Because the presence of melt has a first order impact on the speed of seismic wave speeds, we hypothesize that this melt layer plays a primary role in defining the nature of the geophysical anomaly under the Afar Depression and may reconcile geophysical and geochemical models of the mantle in this region, as proposed by previous seismological studies in the region (e.g., Bastow et al., 2005; Kendell et al., 2005). The thermal anomaly is modest, water contents are elevated but not remarkable, fO_2 values are like those of the upper mantle that feeds the mid-ocean ridge spreading system – these variables independently cannot drive the remarkable nature of the geophysical anomaly under the Afar rift. The modest thermal anomaly and somewhat elevated H_2O contents are characteristics of the Afar mantle plume and in combination, do however provide a mechanism for generating and sustaining the presence of partial melts in the mantle in this region. While there are several competing models for both the shear-wave velocity of melt-free peridotite and the effect of melt on shear-wave velocity, no current model can explain shear-wave velocities at the temperatures inferred in this study below ~4.1 km/s without recourse to some effect of melt (Byrnes et al., 2023). Our PriMELT-3P calculations report that the mantle residue for samples in this study is peridotitic to harzburgitic in composition (see supplement), and at 2 GPa and 1400°C this rock is expected to have $V_s \sim 4.3$ km/s (Hacker and Abers, 2004) for typical grain sizes of ~1 cm. Observed V_s at this depth in Afar are ~4.0 km/s or lower, representing a 7% or greater decrease in V_s (Emry et al., 2018). This can

be achieved by the persistent presence of ~1% or slightly less of partial melt (1% melt produces a decrease in V_s of 7.9%; Hammond and Humphreys, 2000). In a broad sense, this is supported by our PriMELT-3P calculations that suggest for instance, that Erta Ale lavas are 21-28% partial melts of a harzburgitic mantle source produced over a range of ~30 km in the mantle from 93 km to 63 km depth. This is a simplification, but if this melt were distributed evenly within that range of melting, it would correspond to ~0.7-0.9% melt per km of mantle rock below the edifice. Our work supports recent similar calculations from seismological perspectives (e.g., Chambers et al., 2019); chemical and physical models for the crust and the mantle in the East African Rift converge. Importantly, the persistent presence of broadly distributed melt is likely supported by the presence of the Afar plume and argues for the importance of the plume and magmatism more broadly in the initiation and continued development of the rift.

6.0 Conclusions

Gulf of Aden submarine glasses range in H_2O contents from 0.14 to 0.84 wt% and have 0.06 to 0.30 $S^{6+}/\Sigma S$ and 0.135 to 0.189 $Fe^{3+}/\Sigma Fe$ ratios. The glasses recovered east of 49°E have radiogenic isotopic compositions most like melts of the depleted MORB mantle and have the lowest $S^{6+}/\Sigma S$, $Fe^{3+}/\Sigma Fe$ and H_2O contents. Glasses erupted between 45°E and 49°E have radiogenic isotopic compositions most like melts of the Afar mantle plume and have higher H_2O contents (average = 0.61 wt%) but low $Fe^{3+}/\Sigma Fe$ (average = 0.158) and moderate $S^{6+}/\Sigma S$ (average = 0.18). Erta Ale melt inclusions in plagioclase, pyroxene, and olivine all have $H_2O < 0.67$ wt% and $S^{6+}/\Sigma S \sim 0.12$, consistent with two previous melt inclusion studies of the same eruption. Combined with previous studies, we model the primary melt and mantle source characteristics

of the mantle along the Gulf of Aden and into the Afar Depression and find that the Afar mantle plume has moderate H₂O contents of $\sim 852 \pm 167$ ppm H₂O and fO_2 of $\Delta QFM \sim -0.2$, similar to that of the depleted MORB mantle. This is consistent with its radiogenic isotopic character as a C/FOZO/PHEM plume which has been shown to not produce lavas substantially elevated in fO_2 at Reunion Island (Brounce et al., 2022; Nicklas et al., 2022). The mantle sources of Afar Depression volcanoes Erta Ale, Dabbahu, and Nabro have 113-397 ppm H₂O and fO_2 of $\Delta QFM \sim 0$ to +0.8. Melting is estimated to begin under the Afar Depression and the Gulf of Aden around 93 km and end at 37 km under the Gulf of Aden and at 63 km under the Afar Depression. This occurs in a mantle with average region wide potential temperature of 1458°C, producing melt fractions of 11-16% (simple olivine addition) or 27-29% (PriMELT3P) along the Gulf of Aden most influenced by the Afar plume, to melt fractions of ~ 1 -11% (simple olivine addition) or 7-28% (PriMELT3P) under the Afar Depression. We find our results are consistent with recent geophysical models of seismic wave speeds that suggest a melt rich lens in the asthenosphere under the Afar Depression to explain the extreme nature of the present-day geophysical anomaly. Taken together, these works reconcile seemingly disparate views of the mantle under the East African Rift – moderate geochemical anomalies (i.e., slightly elevated mantle potential temperatures and a damp mantle plume) generate melt, which has a pronounced impact on seismic wave behaviors. It emphasizes the continued importance of the role of the Afar mantle plume in the East African Rift through to the present day.

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OPEN RESEARCH: Data availability Statement

The major and volatile element, and Fe and S redox data used in the study are available as EarthChem libraries (Brounce et al., 2025a; 2025b).

Figure Captions

Figure 1. Map of the East African Rift with major segment names indicated. The location of samples for which new data are presented here are marked as stars, the location of samples for which we rely on literature data are marked as circles. Important geographical features are labeled, including the position of Lake Abhe, the suggested center of the Afar mantle plume. The basemap was created using GeoMapApp (<http://geomapapp.org>; Ryan et al., 2009)

941 Figure 2. Major element variations for phenocryst-hosted melt inclusions and matrix glasses
942 from Erta Ale (large dark and light green circles with black outlines) and submarine pillow
943 glasses from the Gulf of Aden (large black circles), as well as phenocryst-hosted melt inclusions
944 from Erta Ale and Nabro volcano from the literature (Field et al., 2012; de Moor et al., 2013;
945 Donovan et al., 2018) and submarine pillow glasses from the Central Indian Ridge mid-ocean
946 ridge spreading system (Gale et al., 2013). Black curves show the trajectory of a cooling basaltic
947 liquid during crystallization produced using MELTs (see main text for details).

948
949 Figure 3. Volatile element variations for phenocryst-hosted melt inclusions from Erta Ale and
950 submarine pillow glasses from the Gulf of Aden, as well as phenocryst-hosted melt inclusions
951 from Erta Ale and Nabro volcano. Symbols are as in figure 2. The black curve in panel (d) shows
952 the calculated sulfur content at sulfide saturation for a selected Gulf of Aden glass, calculated
953 using the model of O'Neill & Mavrogenes (2022), using measured $\text{Fe}^{3+}/\Sigma\text{Fe}$, major element
954 composition, Ni and Cu abundances, and pressure of seafloor at the point of sampling for that
955 sample.

956
957 Figure 4. (a) S oxidation states for phenocryst-hosted melt inclusions and matrix glass from Erta
958 Ale (large dark and light green circles with black outlines), Fe oxidation states for phenocryst-
959 hosted melt inclusions from Erta Ale from de Moor et al. (2013; small tan circles) and paired Fe
960 oxidation states (large black circles) and S oxidation states (large gray circles with black outlines)
961 on Gulf of Aden submarine glasses. Note that values for S oxidation states are shown on the
962 right hand y-axis, and values for Fe oxidation states are shown on the left hand y-axis. (b) Fe and

S oxidation states measured in the same glass chips for submarine pillow glasses from the Gulf of Aden. The black curve shows the line of best fit through model calculations of the S oxidation states of these glasses, given their major element composition, temperature, and measured Fe oxidation states, using the model of O'Neill & Mavrogenes (2022; note that the model of Boullind & Wood, 2023 produces a very similar curve). Gray arrow indicates the expected shift in S oxidation state of a basaltic silicate glass as a function of temperature at a fixed Fe oxidation state.

Figure 5. Geochemical compositions of Gulf of Aden submarine glasses as a function of their distance from Lake Abhe, used by Rooney et al. (2012) as a marker of the presumed center of the Afar Mantle plume under the Afar Depression. The shaded gray region in panel (a) indicates the composition of the “C” mantle component by Hanan & Graham (1996), in which the Afar Mantle Plume is thought to be abundant. Panel (d) plots H₂O contents (left y-axis, black circles) and H₂O/Ce ratios (right y-axis, gray circles). The dark gray, light gray, and pink rectangles demarcate the H₂O/Ce ratios measured in MORB (light gray; Dixon et al. 2002, Michael, 1995), proposed value for FOZO based on measurements in Hawaii (dark gray; Shimizu), and measured in SWIR (pink; Wang et al., 2021). Panel (e) plots Fe³⁺/ΣFe (left y axis, black circles) and S⁶⁺/ΣS (right y axis, gray circles). The fO₂ shown in panel (f) is calculated using the calibration of Kress & Carmichael (1991) at 1 atmosphere and the magmatic temperature calculated using MgO glass compositions according to Helz and Thornber (1987; see supplement).

984 Figure 6. Summary of proposed pressures and temperatures of melting for lavas erupted in the
985 Afar Depression. New constraints from this study are the fractional melting models from
986 PriMELT3, with initial pressures (P_i) and final pressures (P_f) of melting indicated by two circles
987 (large white, green, black, and pink circles). Results from other studies are shown in gray circles
988 (Ferguson et al., 2013; Furman et al., 2016), gray rectangles with black outlines (Rooney et al.,
989 2012); black crosses (Beccaluva et al., 2009), and black vertical lines, also with initial and final
990 pressure of melting indicated by the length of the line (Beccaluva et al., 2009). Also shown are
991 equilibration pressures and temperatures of lithospheric xenoliths (light gray shapes with no
992 outline; Beccaluva et al., 2009; Conticelli et al., 1999). We include temperature estimates for
993 melting along the mid-ocean ridge spreading system, at the Azores, and at Hawaii, calculated
994 using an earlier version of PriMELT (black horizontal lines, Rooney et al., 2012). The spinel to
995 garnet transition is marked by the thick dashed gray curve (Roobinson and Wood, 1998). Two
996 dry peridotite solidi are shown (Hirschmann, 2000, thin black curve; Sarafian et al., 2017, thin
997 dashed black curve), as well as the damp peridotite solidus (Sarafian et al., 2017) and the dry
998 pyroxenite solidus (Pertermann and Hirschmann, 2003).

999

1000 Figure 7. Rare earth element diagrams for Gulf of Aden glasses (normalized to chondrite from,
1001 Sun and McDonough, 1995). Bold curves in the top panel mark two samples with exceptionally
1002 high H_2O/Ce ratios. Bold curves in the middle panel mark three samples with exceptionally high
1003 contributions from the Afar plume (as calculated from radiogenic isotopic mixing calculations,
1004 see main text and supplement). Bold curves in the bottom panel mark a fourth sample with

exceptionally high contributions from the Afar plume (red curve) and the only sample with high Dy/Yb_N (green curve).

Figure 8. Plot of ⁸⁷Sr/⁸⁶Sr and ²⁰⁶Pb/²⁰⁴Pb isotopic compositions of oceanic basalts. Small gray circles are global MORB and OIB from Stracke (2012). Gulf of Aden samples from this study are large black circles, isotopic compositions reported by Schilling et al. (1992). Colored circles are isotopic compositions at OIB locations where some samples have Fe-XANES constraints for *f*O₂ at the time of writing (Brounce et al., 2021 and references therein).

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