## Heads and tails: 30 million years of the Afar plume

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Abstract: Primitive recent mafic lavas from the Main Ethiopian Rift provide insight into the structure, composition and long-term history of the Afar plume. Modern rift basalts are mildly alkalic in composition, and were derived by moderate degrees of melting of fertile peridotite at depths corresponding to the base of the modern lithosphere (c.100 km). They are typically more silica-undersaturated than Oligocene lavas from the Ethiopia–Yemen continental flood basalt province, indicating derivation by generally smaller degrees of melting than were prevalent during the onset of plume head activity in this region. Major and trace element differences between the Oligocene and modern suites can be interpreted in terms of melting processes, including melt-induced binary mixing of melts from the Afar plume and those from three mantle end-member composition itself has remained essentially constant over the past 30 million years, indicating that the plume is a long-lived feature of the mantle. The geochemical and isotopic compositions of mafic lavas derived from the Afar plume support a modified single plume model in which multiple plume stems rise from a common large plume originating at great depth in the mantle (i.e. the South African superplume).

Large igneous provinces, including both continental flood basalts (CFB) and oceanic plateaus, represent the largest outpourings of mafic igneous material on our planet. These provinces are attributed to melting associated with the initiation of mantle plume activity (Morgan 1972; White & McKenzie 1995; Richards et al. 1989; Hill et al. 1992). Several major flood basalt provinces (e.g. Deccan, Paraná-Etendeka, Greenland) have been studied intensively in recent years, leading to the recognition that flood basalt lavas often include contributions from both lithospheric mantle and deep mantle sources, in some cases overprinted by crustal contamination (Hawkesworth et al. 1998; Peate et al. 1992; Arndt et al. 1993; Baker et al. 1996a, 2000; Fram & Lesher 1997; Janney & Castillo 2001; Ewart et al. 2004). It is generally difficult to determine how these source components vary either temporally or spatially, as most flood basalt provinces are either (a) too old/altered to preserve sufficient fresh lavas for detailed geochemical analysis; or (b) deeply dissected by erosion and tectonism associated with the final stages of continental rifting. These difficulties make evaluation of temporal geochemical trends, particularly those associated with the change from plume head magmatism to plume tail activity (e.g. Farnetani et al. 2002), unfeasible. The lone exception to these limitations is the Cenozoic Ethiopia–Yemen continental flood basalt province (Fig. 1), the majority of which was emplaced at c. 30 Ma and has not yet been compromised by extensive faulting and alteration.

Most studies attribute Ethiopia-Yemen flood basalt activity to impact of the Afar plume head beneath the Ethiopian plateau (e.g. Baker et al. 1996a; Pik et al. 1999; Kieffer et al. 2004). The relationship between this copious volcanism at 30 Ma and sparser modern activity in the region is not entirely clear. Most problematic in this regard is the lack of an identifiable track linking the flood basalt province to the modern rift setting that corresponds to plate motion reconstructions. Specifically, it is unresolved whether Africa has been essentially stationary over the past 30 million years (Silver et al. 1998; Gripp & Gordon 2002) or whether it has moved north-northeastward over this time period (Gordon & Jurdy 1986; Burke 1996). If modern Afar volcanism is derived from the same plume that fed the Oligocene flood basalts, then geodynamic models must incorporate physical coupling between the plume and the overlying lithosphere (see Calais et al. 2006). This issue arises, in part, because Cenozoic volcanism in the East African rift is not limited to the CFB province

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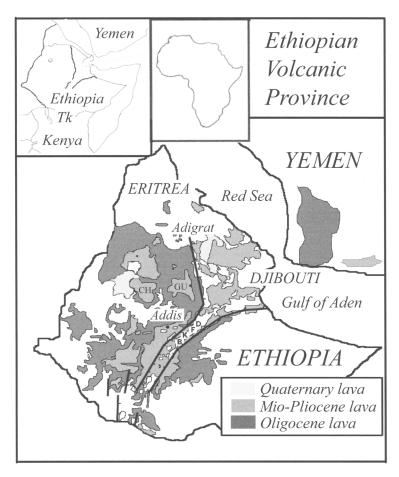


Fig. 1. Map of the Ethiopian Volcanic Province showing ages of volcanic rocks and sample localities mentioned in the text. Recent volcanic centres in the Main Ethiopian Rift are labelled: D = Dofan, F = Fantale, K = Kone, B = Boseti. CH and GU are the 23 Ma volcanic centres of Choke and Guguftu. On the inset map, Tk indicates the location of the Turkana province.

in Ethiopia and Yemen. In southernmost Ethiopia, voluminous basaltic activity began at c. 45 Ma, about 15 million years prior to the inferred arrival of the Oligocene plume head at the base of the Ethiopian lithosphere (Davidson & Rex 1980; Haileab et al. 2004; Vetel & Le Gall 2006). This dual-focus distribution of thick basalt sequences characterized by a variety of compositional differences has led to a range of geodynamic models involving both single (Ebinger & Sleep 1998), modified single (Furman et al. 2004, 2006) and double (George et al. 1998; Rogers et al. 2000; Pik et al. 2006; Rogers 2006) plumes, the validity of which remain unresolved. Most recently, volcanism in Afar and throughout East and South Africa has been attributed to the South African superplume (Janney et al. 2002; Kieffer et al. 2004; Furman

*et al.* 2004, 2006), the largest seismic anomaly in the deep mantle (e.g. Ritsema *et al.* 1999; Gurnis *et al.* 2000; Nyblade *et al.* 2000; Zhao 2001; Ni *et al.* 2002). If our analysis indicates that the Afar plume has been a consistent feature of the deep mantle for 30 million years, it will provide a new window into the thermochemical structure of the South African superplume and its long-term evolution.

In this paper, we present new geochemical data on young mafic lavas from the Main Ethiopian Rift. These data permit characterization of the melting conditions and source components as they vary temporally. We then integrate our findings with the geochemical signatures from Oligocene and recent mafic lavas over the spatial extent of the CFB province. We combine data from several

well-constrained regional studies from the Yemeni and Ethiopian portions of the Oligocene flood basalt sequences, and compare them to recent volcanics from Djibouti, the Afar region, the Main Ethiopian Rift and the Turkana province. The new data, taken in context with the CFB data, provide constraints on several outstanding questions relevant to flood basalt petrogenesis, namely: (a) What are the conditions of mantle melting at the onset of plume activity? (b) What source domains are involved in the genesis of the Oligocene magmas? (c) Do these same characteristics persist until the present time? (d) Is the Afar plume a long-lived feature of the mantle?

### Geodynamic setting

The Cenozoic Ethiopia-Yemen flood basalt province is located at the junction of three rifts: the Red Sea, the Gulf of Aden, and the East African rift system. Within central Ethiopia and Eritrea, the flood basalts cover an area of about 600 000 km<sup>2</sup>. with an estimated total volume of about 350 000 km<sup>3</sup> (Mohr 1983; Mohr & Zanettin 1988). The lava pile thickness varies, reaching up to 2000 m along the margins of the southern Red Sea, and thinning to about 500 m towards both north and south. Most of the basalts and associated felsic rocks were erupted in a period of c.1 million years at around 30 Ma (Zumbo et al. 1985; Baker et al. 1996b; Hofmann et al. 1997; Pik et al. 1998), concomitant with the onset of continental rifting by 29 Ma (Wolfenden et al. 2004). This continental flood basalt province contains the youngest and best-exposed sequence of mafic volcanic rocks associated with incipient continental break-up. It has been an area of episodic magmatic activity since the Oligocene, and is today a site of mafic volcanism attributed to the Afar plume (Vidal et al. 1991; Schilling et al. 1994; Marty et al. 1993, 1996; Deniel et al. 1994; Barrat et al. 1998). The Ethiopian continental flood basalt province thus provides an excellent opportunity to evaluate questions related to plume magmatism throughout the process of incipient continental rifting.

Detailed geochemical studies of modern hotspot volcanism in Hawaii (Lassiter & Hauri 1998; Blichert-Toft et al. 1999; DePaolo et al. 2001; Abouchami et al. 2005; Bryce et al. 2005), the Galápagos (Hoernle et al. 2000; Harpp & White 2001; Geldmacher et al. 2003; Thompson et al. 2004) and Iceland (Thirlwall 1995; Fitton et al. 1997; Hanan & Schilling 1997; Hanan et al. 1999; Kempton et al. 2000) have demonstrated convincingly that long-lived heterogeneities are preserved within mantle plumes located in ocean basins. These heterogeneities are expressed, to first order, spatially and/or temporally on these oceanic islands and thus are thought to represent melting of long-lived chemical and mineralogical domains located within the source region of the plume tail. No comparable study of spatial and temporal variations in plume volcanism has been attempted in a continental setting. The Afar plume provides an ideal laboratory to test these ideas using fairly young lavas (<30 Ma) from a tectonic environment that is well-constrained in both structural and geophysical terms (e.g. Mahatsente *et al.* 1999; Benoit *et al.* 2003; Keranen *et al.* 2004; Dugda *et al.* 2005; Kendall *et al.* 2005; Mackenzie *et al.* 2006; Bastow *et al.* 2005).

The mafic lavas in this study were collected from four volcanic centres within the Wonji Fault Belt (Mohr 1962; WoldeGabriel et al. 1990); from north to south they are Dofan, Fantale, Kone and Boseti (Fig. 1). Each volcano has a well-developed central silicic volcanic edifice, with basaltic lavas erupted primarily from cinder cones and along fissures towards the volcano flanks and between the volcanoes. At Kone, basaltic fissures cut through the nested Quaternary caldera structures and continue to the SW. These silicic volcanoes have been active for about three million years (WoldeGabriel et al. 1990; Chernet & Hart 1999) and have had several catastrophic eruptions which produced the pumice and ash that cover the rift valley floor.

## Sampling

Samples in this study were collected during field excursions in 2002 and 2003 and were chosen to obtain representative coverage of mafic lavas from the individual Quaternary volcanoes, augmented by post-Miocene syn-rift basalts emplaced along the border faults. This investigation is thus more comprehensive than previous studies that have broader regional coverage and/or include samples that range from Oligocene to Recent (Hart et al. 1989; Boccaletti et al. 1999; Chernet & Hart 1999). We focus here on fresh, young mafic lavas with >6 wt% MgO in order to elucidate the history of mantle melting and the nature of the source domains that contribute to this modern volcanism. The origin of more-evolved lavas (samples with <6 wt% MgO), including a discussion of shallow-level fractionation processes and its relationship to magmatic segmentation of the rift axis, will be presented elsewhere. The basalts are sparsely phyric, typically with <5vol.% phenocrysts of olivine, accompanied in more-evolved basalts by clinopyroxene and plagioclase in variable proportions.

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In order to assess the chemical heterogeneity within the modern volcanic province, we incorporate studies of mafic lavas from Djibouti (Vidal et al. 1991; Deniel et al. 1994), Erta 'Ale (Barrat et al. 1998; Ayalew et al. unpublished data) and the West Sheba Ridge in the Gulf of Aden (Schilling) et al. 1992). To evaluate possible temporal evolution of the Afar plume, we compare these recent basalt suites to well-documented Oligocene continental flood basalts from Ethiopia (Pik et al. 1998, 1999; Kieffer et al. 2004) and Yemen (Chazot & Bertrand 1993: Baker et al. 1996a, 2000). There are limited data available for lavas emplaced between 30 and 3 Ma; we include analyses of the c.7 Ma Addis Ababa series (this study) and 23 Ma basalts from Choke and Guguftu volcanoes (Kieffer et al. 2004; Fig. 1). To broaden our regional scope, we compare the Afar province lavas to basalts from Turkana, nothern Kenya (Furman et al. 2004, 2006); geochemical characteristics of the Kenya rift are described elsewhere (Rogers 2006).

## **Analytical methods**

Centimetre-thick slabs were polished to remove saw marks, and then crushed into millimetre-sized chips. These chips were then powdered in a WC disc mill. Bulk rock analyses for major and trace elements were completed by DCP and ICP-MS analysis at Duke University and by ICP-MS at the Centre Récherches Pétrographiques et Géochimiques, Nancy, France (Table 1). Bulk rock Sr– Nd–Pb isotopic analyses (Table 2) were performed at UCLA and San Diego State University. Details of the analytical procedures are in the Appendix.

## Results

## Major elements

The modern Ethiopian rift lavas comprise primarily transitional tholeiitic basalts and trachybasalts (Fig. 2). Their bulk compositions are hyperstheneto mildly nepheline-normative, and evolved lavas tend towards silica-undersaturated compositions. The Ethiopian lavas are thus compositionally similar to contemporaneous mafic lavas from Turkana, northern Kenya (Furman *et al.* 2004) and 23 Ma basalts from Choke and Guguftu, but more alkali-rich than the Oligocene flood basalts which are dominated by quartz-normative tholeiites (Pik *et al.* 1999; Kieffer *et al.* 2004). The degree of differentiation of recent lavas varies between the individual volcanoes, with the most primitive basalts erupted from fissures immediately south of Kone. Each of the central volcanic complexes is characterized by low-MgO basalts through rhyolitic lavas.

The majority of lavas described in this study contain between 6.5 and 9.5 wt% MgO; no picritic lavas have been sampled in the modern rift setting although rare Fe-rich picrites (ferropicrites of Gibson et al. 2000) occur in both Ethiopian and Yemeni flood basalt sequences. Recent Ethiopian basalts have TiO<sub>2</sub> contents that range from 1.8-2.3 wt%, somewhat higher than values observed among Oligocene low-Ti lavas (1.1-2.1 wt%) but substantially lower than the primitive high-Ti suite (2.7-5.1 wt%). Pik et al. (1999) identified these coeval high- and low-titanium basalt series within the continental flood basalt sequence, and proposed a geographic control on eruptive composition. The presence of sparse high-Ti lavas among the Yemeni flood basalt sequence (Baker et al. 1996b), however, suggests that this lava type is more widespread than recognized previously.

## Trace elements

Abundances of compatible trace elements approach those expected for primary mantle melts (c. 200 ppm Ni, c. 750 ppm Cr, c. 33 ppm Sc; Table 1), suggesting that the most magnesian modern Ethiopian rift lavas have undergone only minor olivine fractionation during melt ascent and storage. Variation of these elements with MgO content among primitive Ethiopian basalts of all ages suggests a dominant role for olivine crystallization; in contrast, co-precipitation of olivine and clinopyroxene is inferred for Quaternary basalts from Turkana, nothern Kenya (Furman *et al.* 2004).

Chondrite-normalized rare earth element (REE) patterns of the Ethiopian rift basalts have moderate negative slopes and show little variation within or between eruptive centres (Fig. 3). Oligocene flood basalts from throughout Ethiopia (Pik et al. 1999) have lower overall abundances and consistently flatter rare earth element patterns, whereas primitive high-Ti samples have steeper patterns and substantially higher abundances, despite their higher MgO contents (Fig. 3). Two Oligocene high-Ti lavas from Adigrat (NW Ethiopia) have moderately steep, concave-down patterns that set them apart from the other flood basalts. In contrast to all of the Oligocene lavas, the 23 Ma lavas from Choke and Guguftu have rare earth element abundances that fall within the range of the modern rift lavas (Fig. 3), consistent with major element evidence (Fig. 2) suggesting these lavas are more closely related to Quaternary Turkana basalts than to the CFB and Main Ethiopian Rift products.

Incompatible trace element abundances in the young Ethiopian rift basalts are moderately

ecision and g to Hart	1030 Kone	$\begin{array}{c} 8.8\\ 39.79\\ 16.33\\ 16.33\\ 16.33\\ 15.00\\ 13.00\\ 13.00\\ 1.43\\ 1.5\\ 1.43\\ 1.5\\ 1.43\\ 1.6\\ 1.43\\ 1.6\\ 1.43\\ 1.6\\ 1.43\\ 1.6\\ 1.43\\ 1.6\\ 1.43\\ 1.6\\ 1.43\\ 1.6\\ 1.43\\ 1.6\\ 1.43\\ 1.6\\ 1.43\\ 1.6\\ 1.43\\ 1.6\\ 1.43\\ 1.6\\ 1.6\\ 1.2\\ 1.2\\ 1.2\\ 1.2\\ 1.2\\ 1.2\\ 1.2\\ 1.2$	5.5
echniques, pr uted accordin	E99-3 Kone	$\begin{array}{c} 46.68\\ 16.58\\ 16.58\\ 16.58\\ 13.23\\ 15.53\\ 10.41\\ 10$	5.2
or analytical t tes are compi	N-17 Kone	8.88 39.79 46.64 16.37 17.47 17.57 1	6.5
Appendix fc e ∆8/4 valı	1035A Fantale	$\begin{array}{c} 8.80\\ 8.86\\ 15.64\\ 15.64\\ 15.64\\ 12.34\\ 12.34\\ 12.80\\ 0.68\\ 0.68\\ 0.68\\ 0.68\\ 0.68\\ 0.68\\ 0.68\\ 0.68\\ 0.42\\$	5.1
<i>i</i> isotopic data for Ethiopian rift basalts. See supplementary data section in the Appendix for analytical techniques, precision and for the 7 Ma Addis Ababa samples (denoted with <sup>*</sup> ) represent initial values. The $\Delta 8/4$ values are computed according to Hart or the 7 Ma values using $\kappa = 2.5$ .	1035 Fantale	8.80 39.68 47.28 15.48 7.68 7.68 7.68 7.68 7.68 0.19 0.71 0.71 0.71 0.71 0.71 0.71 0.71 0.71	4.5
lementary data *) represent ini	N-18 Fantale	$\begin{array}{c} 8.91\\ 8.91\\ 15.63\\ 15.63\\ 12.24\\ 12.06\\ 12.35\\ 10.19\\ 12.06\\ 12.35\\ 10.11\\ 10.19\\ 0.50\\ 0.50\\ 0.51\\ 1.9\\ 1.9\\ 1.9\\ 1.9\\ 1.9\\ 1.9\\ 2.01\\ 1.9\\ 2.01\\ 1.9\\ 2.01\\ 1.9\\ 2.01\\ 2.01\\ 2.01\\ 2.01\\ 2.02$	6.6
c data for Ethiopian rift basalts. See suppl. 7 Ma Addis Ababa samples (denoted with* 10 7 Ma values using $\kappa = 2.5$ .	1032 Fantale	$\begin{array}{c} 8.87\\ 8.87\\ 16.25\\ 16.25\\ 16.25\\ 0.20\\ 0.57\\ 0.57\\ 0.57\\ 0.57\\ 0.57\\ 0.57\\ 0.57\\ 0.57\\ 0.57\\ 0.57\\ 0.57\\ 0.57\\ 0.57\\ 0.57\\ 0.5\\ 0.5\\ 0.5\\ 0.5\\ 0.5\\ 0.5\\ 0.5\\ 0.5$	6.7
thiopian rift bc Ababa samples ues using κ =	E99-2 Fantale	$\begin{array}{c} 46.52\\ 16.72\\ 16.72\\ 16.72\\ 15.35\\ 0.23\\ 0.23\\ 0.52\\ 0.52\\ 0.5\\ 0.5\\ 0.5\\ 0.5\\ 0.5\\ 0.5\\ 0.5\\ 0.5$	6.9
i isotopic data for Ethiopian r for the 7 Ma Addis Ababa san rrected to 7 Ma values using	1018 Fantale	$\begin{array}{c} 8.92\\ 8.92\\ 16.26\\ 16.28\\ 16.28\\ 16.28\\ 1.01\\ 1$	6.2
	1023 Dofan	$\begin{array}{c} 9.14\\ 9.14\\ 15.75\\ 15.75\\ 15.75\\ 15.75\\ 15.75\\ 12.45\\ 0.18\\ 0.18\\ 0.18\\ 0.18\\ 0.18\\ 0.18\\ 0.18\\ 0.18\\ 0.24\\ 0.2\\ 0.9\\ 0.9\\ 0.9\\ 0.9\\ 0.9\\ 0.9\\ 0.9\\ 0.9$	5.3
Table 1. Major and trace element and iccuracy. The isotopic data presented j           1984); the NHRL model values are co	1027 Dofan	$\begin{array}{c} 9.48\\ 40.20\\ 15.73\\ 15.73\\ 15.73\\ 12.77\\ 12.77\\ 12.77\\ 12.77\\ 12.73\\ 12.73\\ 12.73\\ 12.73\\ 12.73\\ 12.75\\ 12.66\\ 12.85\\ 12.7\\ 1.7\\ 1.7\\ 1.7\\ 1.7\\ 1.7\\ 1.7\\ 1.7\\ 1$	3.0
<b>Table 1.</b> Major and trace element an accuracy. The isotopic data presented (1984); the NHRL model values are co	Sample Location	Latitude $(N)$ Longitude $(N)$ SiO <sub>2</sub> SiO <sub>2</sub> SiO <sub>2</sub> SiO <sub>2</sub> SiO <sub>2</sub> Si MgOO Si MgOO SiO <sub>2</sub> SiO	Pr

99

(continued)

Sample Location	1027 Dofan	1023 Dofan	1018 Fantale	E99-2 Fantale	1032 Fantale	N-18 Fantale	1035 Fantale	1035A Fantale	N-17 Kone	E99-3 Kone	1030 Kone
pN	25.4	21.9	25.9	30	28.5	28.4	19.4	21.3	28.5	23.3	23.5
Sm	5.67	4.82	5.50	6.34	6.20	5.97	4.31	4.69	6.24	5.17	5.26
Eu	2.02	1.74	2.09	2.36	2.32	2.13	1.54	1.73	2.27	1.89	1.98
Gd	5.96	5.10	5.88	6.17	6.35	6.46	4.56	5.09	6.68	4.79	5.34
Tb	0.96	0.73	0.84	0.89	0.92	1.01	0.68	0.75	1.07	0.72	0.84
Dy	5.34	4.02	4.47	4.84	5.05	5.05	3.73	4.16	5.35	4.17	4.58
Ho	1.07	0.78	0.87	0.93	0.97	0.99	0.71	0.82	1.02	0.78	0.93
Er	2.82	2.03	2.28	2.37	2.45	2.61	1.86	2.11	2.75	2.00	2.32
Yb	2.51	1.77	1.96	2.14	2.18	2.25	1.67	1.88	2.37	1.90	2.06
Lu	0.37		0.29	0.32	0.33	0.34	0.25	0.27	0.36	0.28	0.32
8/Sr/80Sr	0.703722	0.704102	ļ	0.704117		0.703961	0.704198	}	ł	ļ	l
<sup>143</sup> Nd/ <sup>144</sup> Nd	0.512862	<b>C</b> 1	ł	0.512798	1	0.512799	0.512767				-
ENd	4.4	3.5	]	3.1		3.1	2.5		ļ		
$^{206}Pb/^{204}Pb$	18.827	18.752		ļ	I	18.732	18.627			l	١
$^{207}\text{Pb}/^{204}\text{Pb}$	15.581	15.584		ł	-	15.597	15.585			l	I
$^{208}Pb/^{204}Pb$	38.896	38.782		ł		38.834	38.748			1	ł
$\Delta 8/4$	50.72	48.38		1	-	56.00	60.10	ł			
·											
Sample	N-16	N-22	K0-12	N-20	1037	N-15	1036	E99-11	KO-4B	N-21	
Location	Kone	Kone	Kone	Kone	Kone	Kone	Kone	Kone	Kone	Kone	
Latitude (N)	8.84	8.72		8.83	8.86	8.78	8.84			8.78	
Longitude (E)	39.70	39.68		39.69	39.75	39.63	39.71			39.68	
SiO <sub>2</sub>	46.14	45.42	46.53	45.86	47.14	46.35	46.50	46.28	46.43	46.86	
$Al_2O_3$	15.91	15.29	15.58	14.86	15.06	14.89	14.78	15.03	14.75	14.10	
TiO <sub>2</sub>	2.46	2.31	2.27	2.21	1.95	2.22	2.20	2.14	2.17	1.90	
$Fe_2O_3$	12.82	12.43	12.65	12.46	11.92	12.18	12.56	12.42	12.55	11.73	
MgO	6.83	7.97	8.14	8.33	8.35	8.64	8.51	8.78	8.85	10.44	
MnO	0.19	0.19	0.18	0.19	0.19	0.19	0.20	0.17	0.19	0.18	
CaO	10.95	11.19	10.78	10.58	10.17	10.91	10.69	11.37	10.74	11.02	
$Na_2O$	3.06	3.12	2.89	3.25	2.87	2.73	3.21	2.70	3.22	2.82	
$K_2O$	0.82	1.13	0.64	1.12	0.88	1.06	1.07	0.65	1.00	0.67	
$P_2O_5$	0.45	0.55	0.60	0.54	0.46	0.45	0.55	0.33	0.60	0.37	
Sum	99.63	99.61	100.26	99.41	98.99	99.61	100.27	99.87	100.50	100.09	
Cs	0.13	0.34	0.15	0.26	0.09	0.15	0.19	0.12	0.33	0.13	

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12.7 246	1.41	0.4	6.27 1 3	3.2	392	141	3.6	23.5	54.1	556	165	287	24.7	17.0	37.1	4.8	21.1	4.50	1.61	4.89	0.81	4.21	0.81	2.21	2.04	0.29	0.704228	0.512793	3.0	18.624	15.586	38.794	65.06
26.7 357	2.97	0.7	0.00 2.0	3.0	526	148	3.5	23.2	50.1	433	141	283	33.5	27.9	57.3	7.0	28	5.73	1.82	5.53	0.76	4.52	0.84	2.30	1.90	0.31	0.704015	0.512801	3.2	18.801	15.596	38.872	51.46
11.2 241	1.84	0.4	15	1.7	427	125	3.0	21.2	54.0	465	160	287	29.4	17.7	40.7	4.7	20.6	4.64	1.71	4.45	0.67	3.73	0.76	2.01	1.78	0.28	I	Ι	I				
22.2 411																																	
18.4 324	2.04	0.5 20.2	2.0C	2.0	468	138	3.2	24.1	49.6	345	145	293	17.7	20.2	42.8	5.3	23.2	4.84	1.67	5.22	0.85	4.31	0.85	2.27	1.96	0.30	0.704203	0.512800	3.2	18.680	15.588	38.879	66.79
12.7 401	1.63	0.4 20.6	18.0	2.0	479	148	3.7	27.0	61.3	292	118	256	32.4	23.9	51.7	6.8	28.8	6.28	2.12	5.81	0.93	5.10	1.00	2.63	2.34	0.35	0.704251	0.512768	2.5	18.557	15.584	38.632	56.96
23.6 391	2.69	0.8	40.0 2.3	2.6	586	164	3.7	25.5	50.2	339	135	259	30.7	28.2	57.9	7.0	29.3	5.70	1.99	6.20	0.95	4.65	0.90	2.44	2.05	0.31	0.703996	l	1	18.811	15.592	38.873	50.35
12.2 272	1.57	0.3 10.7	10./	3.0	488	122	3.1	21.4	51.3	289	109	300	29.4	18.6	40.2	5.2	22.6	5.28	1.85	4.75	0.70	3.90	0.80	1.91	1.73	0.31	0.704303	0.512765	2.5	18.658	15.593	38.826	64.15
28.2 405	3.25	0.0	6.74 C.C.C	2.3	619	175	3.7	27.1	52.0	204	96	287	31.3	31.5	64.4	T.T	31.6	6.16	2.05	6.51	1.01	4.98	0.96	2.53	2.09	0.32	0.703768	0.512838	3.9	18.945	15.595	38.896	36.45
13.1 299	1.57	0.4	15	2.0	521	141	3.4	25.7	48.8	143	73	302	31.0	19.2	42.2	5.5	24.2	5.19	1.87	5.73	0.91	4.74	0.92	2.38	2.17	0.31	-	I					
Rb Ba	Th		Ta Ta	Pb	Sr	Zr	Hf	Y	C	Ċ	iz	^	Sc	La	Ce	Pr	PN	Sm	Eu	Gd	Tb	Dy	Ho	Er	$\mathbf{Y}\mathbf{b}$	Lu	$^{87}Sr/^{86}Sr$	<sup>143</sup> Nd/ <sup>144</sup> Nd	ENd	$^{206}Pb/^{204}Pb$	$^{207}$ Pb/ $^{204}$ Pb	$^{208}Pb/^{204}Pb$	$\Delta 8/4$

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(continued)

.02		· · · · · · · · · · · · · · · · · · ·
	E01-15A W. Rift Flank	8.92 39.55 47.55 15.60 15.60 5.74 5.74 0.22 0.55 99.29 99.29 0.55 0.55 1.3 1.3 1.3 1.3 1.3 1.3 1.3 1.3 1.3 1.3
	E01. W.	
	W-04 E. Rift Flank	$\begin{array}{c} 45.05\\ 16.94\\ 15.05\\ 12.63\\ 12.63\\ 12.63\\ 12.63\\ 0.18\\ 0.18\\ 0.18\\ 0.16\\ 0.16\\ 0.16\\ 0.05\\ 0.05\\ 0.05\\ 1.62$
	N-09 E. Rift Wall	$\begin{array}{c} 8.26\\ 8.26\\ 15.49\\ 1.54\\ 1.54\\ 1.54\\ 1.54\\ 1.54\\ 1.52\\ 2.58\\ 3.36\\ 1.02$
	N-12 E. Rift Wall	$\begin{array}{c} 8.28\\ 3.2659\\ 50.59\\ 50.59\\ 50.595\\ 50.595\\ 5.95\\ 5.95\\ 5.97\\ 5.95\\ 5.97\\ 1.83\\ 5.97\\ 5.95\\ 5.96\\ 1.98\\ 1.6\\ 1.98\\ 0.16\\ 0.16\\ 0.16\\ 0.16\\ 0.16\\ 0.16\\ 0.16\\ 0.16\\ 0.16\\ 0.2\\ 2.13\\$
	E99-20 Boseti	$\begin{array}{c} 46.87\\ 46.87\\ 15.09\\ 12.16\\ 0.18\\ 0.13\\ 0.18\\ 0.13\\ 0.16\\ 0.39\\ 0.36\\ 0.36\\ 0.36\\ 0.3\\ 0.06\\ 0.3\\ 0.06\\ 0.3\\ 0.06\\ 0.3\\ 0.06\\ 0.3\\ 0.06\\ 0.3\\ 0.06\\ 0.3\\ 0.3\\ 0.3\\ 0.3\\ 0.3\\ 0.3\\ 0.3\\ 0.3$
	E99-13 Boseti	$\begin{array}{c} 47.37\\ 15.15\\ 15.15\\ 15.16\\ 15.16\\ 15.16\\ 12.19\\ 8.46\\ 0.77\\ 0.77\\ 0.18\\ 0.11\\ 0.77\\ 0.18\\ 0.13\\ 0.41\\ 0.77\\ 0.62\\ 0.9\\ 0.9\\ 0.9\\ 0.9\\ 0.9\\ 0.9\\ 0.9\\ 0.9$
	N-01 Boseti	$\begin{array}{c} 8.39\\ 39.33\\ 46.78\\ 14.65\\ 11.96\\ 11.96\\ 11.96\\ 0.18\\ 0.18\\ 0.18\\ 0.16\\ 0.50\\ 0.50\\ 0.50\\ 0.50\\ 0.50\\ 0.50\\ 0.50\\ 0.53\\ 373\\ 2.5\\ 1.6\\ 1.6\\ 1.6\\ 0.5\\ 0.5\\ 373\\ 3.2\\ 2.5\\ 1.6\\ 1.6\\ 1.6\\ 1.6\\ 0.5\\ 2.5\\ 3.3\\ 2.5\\ 3.3\\ 2.5\\ 1.6\\ 1.6\\ 1.6\\ 1.6\\ 1.6\\ 1.6\\ 1.6\\ 1.6$
	N-02 Boseti	$\begin{array}{c} 8.44\\ 8.44\\ 39.40\\ 11.98\\ 7.60\\ 7.60\\ 7.60\\ 7.60\\ 0.11\\ 1.04\\ 1.04\\ 1.04\\ 1.04\\ 1.04\\ 1.04\\ 1.04\\ 1.04\\ 1.04\\ 1.04\\ 1.04\\ 1.02\\ 0.58\\ 0.5\\ 2.0\\ 2.0\\ 2.0\\ 2.0\\ 2.0\\ 2.0\\ 2.0\\ 2.0$
inued.	E99-6 Kone	$\begin{array}{c} 47.11\\ 14.51\\ 14.51\\ 1.82\\ 10.66\\ 0.18\\ 0.63\\ 0.63\\ 0.55\\ 0.53\\ 0.53\\ 0.53\\ 0.53\\ 0.53\\ 0.54\\ 1.9\\ 0.4\\ 1.9\\ 1.9\\ 0.4\\ 1.9\\ 1.9\\ 0.4\\ 1.9\\ 1.9\\ 0.4\\ 1.9\\ 0.4\\ 1.9\\ 0.4\\ 1.5\\ 20.5\\$
Table 1. Continued.	Sample Location	Latitude (N) Longitude (N) SiO <sup>2</sup> SiO <sup>2</sup> SiO <sup>2</sup> SiO <sup>2</sup> Sum MgO MgO MgO MgO MgO Na <sub>2</sub> O Sum Na <sub>2</sub> O CaO CaO CaO CaO CaO CaO CaO CaO CaO Ca

Af	FAR PLU	ME HEADS AND TAILS	103
			(continued)
$\begin{array}{c} 7.43 \\ 1.19 \\ 6.80 \\ 3.81 \\ 3.81 \\ 3.40 \\ 0.52 \\ 0.52 \\ \end{array}$			
5.22 0.822 0.85 0.85 1.87 1.87 1.87 1.87 0.28	AA99-19 Addis Ababa	$\begin{array}{c} 48.49\\ 17.17\\ 1.94\\ 1.94\\ 1.59\\ 0.16\\ 0.36\\ 0.36\\ 0.38\\ 0.38\\ 0.38\\ 0.38\\ 0.38\\ 0.38\\ 0.38\\ 0.38\\ 0.23\\ 0.38\\ 0.38\\ 0.23\\ 0.38\\ 0.23\\ 0.23\\ 0.38\\ 0.23\\ 0.2$	
	AA99-03* Addis Ababa	$\begin{array}{c} 47.67\\ 15.11\\ 15.11\\ 2.26\\ 12.53\\ 8.40\\ 9.80\\ 9.80\\ 9.80\\ 9.80\\ 0.18\\ 9.80\\ 0.18\\ 0.18\\ 1.08\\ 1.08\\ 1.08\\ 1.28\\ 1.28\\ 1.28\\ 3$	
6.94 1.11 5.76 5.76 1.12 1.12 3.03 2.79 0.42 0.704982 0.704982 0.7104982 0.512585 -1.0 17.853 15.578 38.181 96.97	AA99-6* Addis Ababa	46.14 14.11 12.79 12.14 0.18 9.11 0.74 0.74 0.74 0.32 0.32 1.73 1.73 1.73 1.73 1.73 1.73 1.73 1.73	
4.61 0.63 3.76 0.71 1.89 1.76 0.24 0.704117 	AA99-10* Addis Ababa	45.12 13.05 11.25 11.25 14.38 14.38 0.16 0.16 0.28 0.26 0.28 0.28 0.28 0.28 0.28 0.28 0.28 0.28	
$\begin{array}{c} 7.17 \\ 0.95 \\ 0.92 \\ 0.92 \\ 0.30 \\ 0.30 \\ 0.30 \\ \end{array}$			
5.64 0.90 4.45 0.83 0.83 0.83 0.83 0.83 0.83 0.83 0.704485 0.704485 0.704485 0.704485 0.7124485 0.712488 1.85 1.8 1.85 1.8 1.85 1.8 1.85 1.8 1.85 1.85	E-43 W. Rift Flank	9.13 39.73 47.74 15.37 15.37 7.20 0.18 0.18 0.18 0.18 0.18 0.43 0.43 0.43 0.43 0.43 0.43 0.43 0.43	
6.31 6.31 6.31 6.31 6.31 6.31 6.31 6.31 6.31 6.31 6.31 6.31 6.31 6.31 6.31 6.31 6.31 6.31 6.31 6.0 6.0 6.0 6.0 6.0 6.85 6.0 6.95 6.0 6.95 6.0 6.95 6.0 6.95 6.0 6.0 6.85 6.0 6.95 6.0 6.97 6.0 6.0 6.0 6.0 6.0 6.0 6.0 6.0	E01-32 Awash Gorge	8.88 40.10 47.09 16.20 2.22 13.06 6.49 6.49 0.23 11.09 2.25 0.23 11.09 2.25 0.37 0.30 0.37 0.010.07 0.3 1.14 1.14 1.14 1.14 1.14 1.14 1.14 1.	
4.16 0.68 3.74 0.77 1.78 1.78 1.78 1.78 1.78	EUI-20 W. Rift Flank	$\begin{array}{c} 9.02\\ 39.56\\ 48.93\\ 15.01\\ 15.01\\ 2.50\\ 13.60\\ 2.57\\ 0.21\\ 10.35\\ 3.25\\ 0.21\\ 0.21\\ 0.21\\ 0.21\\ 0.21\\ 0.21\\ 0.21\\ 0.25\\ 0.72\\ 0.11\\ 0.46\\ 0.10\\ 0.11\\ 0.46\\ 0.11\\ 0.12\\ 1.5\\ 1.5\\ 1.5\\ 1.5\\ 1.5\\ 1.5\\ 1.5\\ 1.5$	
$\begin{array}{c} Gd \\ Tb \\ Dy \\ Ho \\ Er \\ Yb \\ Lu \\ ^{87}Sr / ^{86}Sr \\ ^{143}Nd / ^{144}Nd \\ ^{209}Pb / ^{204}Pb \\ ^{207}Pb / ^{204}Pb / ^{204}Pb / ^{204}Pb \\ ^{207}Pb / ^{204}Pb / ^{20$	Sample Location	Latitude (N) Longitude (E) SiO2 Al2O3 TiO2 Fe2O3 MnO CaO Na2O CaO Na2O Sum CaO Na2O Ca Sum Ca Sum Ca Sum Ca Sum Ca Sum Ca Sum Ca Sum Ca Sum Ca Sum Ca Sum Ca Sum Ca Sum Ca Su Su Su SiO2 SiO2 SiO2 SiO2 SiO2 SiO2 SiO2 SiO2	

nomination it areas	3							
Sample Location	E01-20 W. Rift Flank	E01-32 Awash Gorge	E-43 W. Rift Flank	AA99-10* Addis Ababa	AA99-6* Addis Ababa	AA99-03* Addis Ababa	AA99-19 Addis Ababa	
	3.6	127	269	66	144	174	133	
	28.3	2.9	5.9	2.5	2.9	4.0	2.9	
	45.1	24.7	30.8	17.9	26.2	28.4	22.4	
	52	47.5	54.0	64.7	64.3	58.2	44.9	
	50	45	109	1136	663	391	64	
Ni	313	73	72	410	389	124	35	
	32.5	281	281	234	218	258	272	
	19.8	29.9	23.9	30.8		30.8		
	45.0	15.9	33.9	17.4	19.6	27.4	20.7	
	5.9	35.3	47.8	36.7	36.1	61.5	44.2	
	25.8	4.8	9.7	4.3	4.4	7.5	5.3	
	8.53	20.9	39.7	17.8	18.3	30.9	23	
	2.01	4.80	8.27	3.67	3.93	6.33	4.86	
	6.06	1.76	2.71	1.27	1.33	2.11	1.63	
	0.96	5.12	8.15	4.01	3.94	6.54	4.53	
	5.21	0.82	1.17	0.57	0.61	0.99	0.65	
	1.02	4.40	5.93	3.06	3.47	5.21	3.92	
	2.69	0.00	1.13	0.70	0.67	1.02	0.74	
	2.36	2.37	2.86	1.60	1.83	2.57	1.93	
	0.34	2.17	2.32	1.64	1.81	2.33	1.96	
Ľu	0.704237	0.32	0.33	0.21	0.28	0.35	0.28	
a/Sr/ <sup>ao</sup> Sr	0.512753	0.704321	0.703841	0.703643	0.703880	0.704064		
DN <sup>441</sup> /DN <sup>641</sup>	2.2	0.512767	0.512842	0.512886	0.512780	0.512748		
8Nd		2.5	4.0	4.8	2.8	2.1		
$^{206}Pb/^{204}Pb$		18.429	18.837	18.779	-	18.212	l	
$dd_{PD}/dd_{DD}$	1	15.580	15.597	15.594		15.570		
$^{208}Pb/^{204}Pb$	1	38.540	38.851	38.769		38.398		
$\Delta 8/4$		63.23	45.01	43.74		75.19	1	

Table 1. Continued

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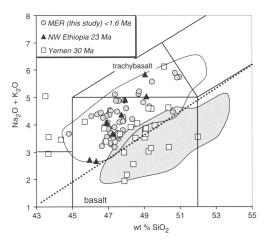


Fig. 2. The total alkalis-silica classification diagram for mafic volcanics of the Ethiopia-Yemen CFB province shows consistent temporal variations. Data are recalculated to 100% on an anhydrous basis with  $Fe^{3+} = 15\%$  of total iron; the dotted line is the alkalictholeiitic division of Macdonald & Katsura (1964) for Hawaiian lavas. Mildly alkalic basalts from the modern Main Ethiopian Rift (MER samples from this study shown as data points; field includes data from Hart et al. 1989; Boccaletti et al. 1999; Chernet & Hart 1999) define a range distinct from that of tholeiitic 30 Ma Ethiopian lavas (shaded field; Hart et al. 1989; Pik et al. 1999; Kieffer et al. 2004). Yemeni CFBs (Chazot & Bertrand 1993; Baker et al. 1996b) generally overlap the Ethiopian suite but include samples with higher alkali contents. Note that 23 Ma basalts from NW Ethiopia (Kieffer et al. 2004) plot within the modern MER field.

enriched over primitive mantle (Sun & Mcdonough 1989), and generally overlap values measured in recent Djibouti lavas attributed to the Afar plume (Vidal et al. 1991; Deniel et al. 1994). Normalized incompatible trace element variation diagrams (Fig. 3) show generally parallel patterns within the Ethiopian rift suite: all samples have prominent positive Ba and Nb-Ta anomalies and weakly negative K anomalies. In detail, however, some consistent differences are found between lavas from individual volcanic centres that require distinct histories of melting and/or a limited degree of source heterogeneity along the rift axis. For example, MgO-rich Kone basalts have the highest abundances of most highly incompatible trace elements, suggesting these lavas are derived by a lower degree of melting than occurs beneath neighbouring volcanoes; this observation is consistent with the higher normative nepheline contents of those same lavas.

Regional and temporal differences observed among incompatible trace element abundances

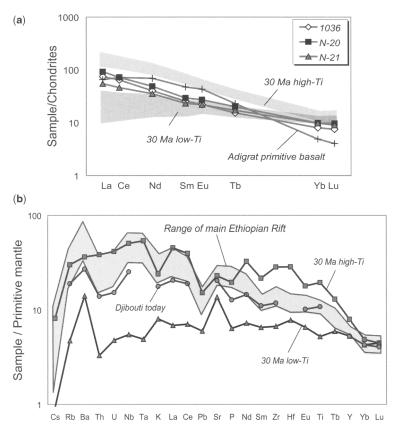
provide insight into the source regions of the mafic lavas through space and time (Fig. 3). Recent Djibouti basalts have slightly lower abundances of the most highly incompatible elements and P, but otherwise define patterns that parallel those of modern Main Ethiopian Rift lavas. Primitive Oligocene high-Ti lavas from Ethiopia and Yemen, including Adigrat, have abundances of the most highly incompatible trace elements that generally overlap the Ethiopian rift suite, but the older samples lack Ba anomalies and have consistently higher abundances of the middle and heavy REE, Zr, Hf, Ti and Y (Pik et al. 1999). In contrast, Oligocene lavas with <7 wt% MgO have lower contents of the more highly incompatible elements than the Ethiopian rift lavas, and overlapping abundances of the moderately incompatible elements (Fig. 4). Most of these suites are characterized by lavas with positive Ba anomalies (leading to elevated Ba/Rb) and negative K anomalies that suggest melting in the presence of minor amphibole.

Mafic Ethiopian rift lavas define a restricted range in La/Nb-Ba/Nb that is close to primitive mantle values, indicating a limited role for enriched mantle or mantle lithosphere sources (Fig. 4). Many of the Ethiopian rift basalts plot within the field defined by recent lavas derived from the Afar plume as sampled in Djibouti and Erta' Ale. Primitive high-Ti Oligocene lavas overlap these fields at low Ba/Nb values, but moderate- and low-Ti flood basalts from both Ethiopia and Yemen extend to much higher values of La/Nb and Ba/Nb, indicating a substantial contribution from enriched mantle sources. Similarly, Ce/Pb values of the Ethiopian rift and primitive high-Ti lavas plot primarily within the range of mantle-derived basalts, whereas the other suites trend towards both crustal and lithospheric end-members (Fig. 4). This ratio is influenced by crustal contributions, as Pb is elevated in crustal rocks relative to mantle lithologies, and the Ce/Pb value of mantle-derived liquids is well constrained ( $25 \pm 5$ ; Hofmann *et al.* 1986). It is interesting to note that Yemen flood basalts that plot within the field of modern rift lavas in Fig. 2 have consistently low values of Ce/Pb (<10) that suggest crustal contamination has played a role in elevating their alkali abundances.

## Radiogenic isotopic data

Radiogenic isotope signatures of recent Ethiopian rift basalts (Table 1) indicate along-axis variations in source composition and/or melt generation processes, and probably also reflect variations in both crustal stretching (Maguire *et al.* 2006) and lithospheric thinning (Bastow *et al.* 2005). Lavas from

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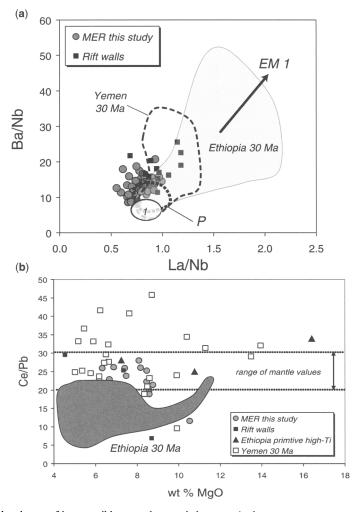
**Fig. 3.** Incompatible element abundances of selected Ethiopian lavas. (a) Rare earth elements. Modern rift basalts have abundances intermediate to those of low- and high-TiO<sub>2</sub> continental flood basalts (data from Pik *et al.* 1999; Kieffer *et al.* 2004); three representative samples are shown (see Table 1). A small number of primitive basalts from Adigrat have more highly concave REE patterns (Pik *et al.* 1999). Recent mafic lavas from Djibouti (Deniel *et al.* 1994) and 23 Ma Ethiopian basalts (Kieffer *et al.* 2004) have REE patterns (not shown) that fall within the range of the modern rift basalts (normalizing values of Boynton, 1984). (b) Primitive mantle-normalized incompatible trace element diagrams (normalizing values of Sun & McDonough 1989) for selected mafic lavas. The range of modern rift basalts is shaded; representative samples from Djibouti and the Oligocene low- and high-TiO<sub>2</sub> suites are shown for comparison. See text for discussion.

Dofan, the northernmost volcanic centre studied, have Sr-Nd-Pb isotope values that fall within the range of young plume-derived Djibouti and Erta  $^{87}{
m Sr}/^{86}{
m Sr} \sim 0.7037,$ 'Ale basalts (Fig. 5):  $^{206}$ Pb/ $^{204}$ Pb ~ 18.83,  $^{143}$ Nd/ $^{144}$ Nd ~ 0.51286,  $^{208}$ Pb/ $^{204}$ Pb ~ 38.90. Isotopic signatures trend slightly towards more enriched values from Dofan southward to Fantale and Kone. The most radiogenic values are seen at Boseti and Gedemsa, i.e.  ${}^{87}Sr/{}^{86}Sr$  up to 0.7053. Post-rift basalts from the Addis Ababa series have isotopic compositions that overlap the Dofan lavas and extend to compositions similar to those observed at Fantale and Kone; syn-rift basalts sampled along the rift walls and away from the main rift axis have more

enriched isotopic signatures similar to those at Boseti and Gedemsa.

At a regional scale, Sr-Nd isotope analyses do not allow individual source components to be distinguished reliably. Ethiopian rift basalts and the modern Afar plume both lie within the broad range defined by 30 million-year-old lavas from Ethiopia and Yemen; 23 Ma basalts from Choke and Guguftu plot within this field as well. The exceptions to this overall uniform picture are the two Oligocene high-Ti lavas from Adigrat noted previously to have unusual rare earth element abundances: these two samples have Sr-Nd isotopic compositions that are more depleted than any other Ethiopian lavas, and fall

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**Fig. 4.** Relative abundances of incompatible trace elements help constrain the source components that contribute to mafic volcanism. (a) La/Nb-Ba/Nb variations among modern rift basalts define a restricted range with La/Nb values similar to those of the primitive mantle, and Ba/Nb values somewhat elevated. Previously published analyses from the same region (Boccaletti *et al.* 1999, not shown) and our new data on syn-extensional basalts from the rift walls extend to higher values, requiring greater contributions from non-primitive source components. Most of the rift basalts plot near the field of young mafic lavas from Djibouti (Deniel *et al.* 1994) and Erta'Ale (Barrat *et al.* 1998), encircled by the dotted line and believed to represent the modern Afar plume. The Ethiopian (Pik *et al.* 1998); Kieffer *et al.* 2004) and Yemeni (Chazot & Bertrand 1993; Baker *et al.* 1996*a*) flood basalts define distinct ranges, but both groups are substantially enriched relative to the modern samples. The field labelled '1' encompasses primitive high-Ti Ethiopian lavas. (b) Ce/Pb-MgO variations show that most of the Main Ethiopian Rift lavas plot within the range of mantle-derived basalts, whereas the rift wall lavas and both Yemeni and Ethiopian flood basalts record contributions from lithospheric or crustal sources. Primitive high-Ti Ethiopian lavas plot within the range of mantle-derived basalts (sources of data: Baker *et al.* 1996*a*; Pik *et al.* 1999; Kieffer *et al.*).

near the MORB suite from the nearby West Sheba Ridge (Fig. 5).

Pb isotope signatures of the recent Ethiopian rift lavas form an array between the modern Afar plume and less radiogenic compositions broadly similar to those observed in 30-million-year-old continental flood basalts from both northern Ethiopia (including Adigrat) and Yemen (Fig. 5). These isotopic values are clearly distinct from compositions observed in Turkana, Kenya from 45 Ma through the present (Furman *et al.* 2004, 2006). Interestingly, the 23 Ma Ethiopian basalts from Choke and Guguftu with unusual major and trace element abundances have Pb isotopic

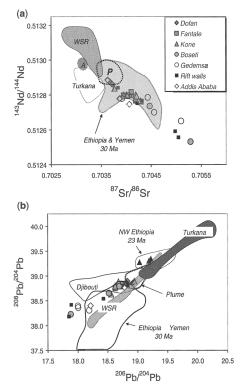


Fig. 5. Radiogenic isotope variations among Ethiopian lavas. (a) <sup>87</sup>Sr/<sup>86</sup>Sr-<sup>143</sup>Nd/<sup>144</sup>Nd variations are strongly correlated among the recent rift lavas, and define a trend that extends from the Afar plume (labelled **P**), defined by mafic lavas from Djibouti (Deniel et al. 1994) and Erta' Ale (Barrat et al. 1998), towards radiogenic compositions more typical of crustal and lithospheric sources (Gedemsa data from Trua et al. 1999; Peccerillo et al. 2003). Lavas from Dofan, the northernmost volcano examined here, and the Addis Ababa series have isotopic compositions that overlap the Afar plume. Oligocene flood basalts (Chazot & Bertrand 1993; Baker et al. 1996a; Pik et al. 1999; Kieffer et al. 2004) define a broad range that encompasses many of the modern lavas but is distinct from Turkana basalts erupted 45 Ma through to the present (Furman et al. 2004, 2005). Two primitive high-Ti basalts from Adigrat (field labelled A) are compositionally distinct from all other basalts reported here. (b) The  ${}^{206}$ Pb/ ${}^{204}$ Pb- ${}^{208}$ Pb/ ${}^{204}$ Pb systematics of these sample suites further define the source components involved. Modern rift basalts vary from compositions overlapping the Afar plume to lower Pb isotope values, and generally overlap the range of 30 Ma flood basalts from both Ethiopia (including Adigrat) and Yemen. Lavas from Turkana have consistently higher Pb isotope compositions, ranging from a high- $\mu$  component sampled between 40–17 Ma to the modern Afar plume, and there is no overlap between the flood and rift basalts and the Turkana suite. Two 23 Ma lavas from Choke, Ethiopia (Kieffer et al. 2004) have elevated Pb isotope compositions relative to all other Ethiopian rocks (see text for discussion).

compositions that fall within the Turkana array rather than the long-term Ethiopia/Afar array (Fig. 5).

# Discussion: melting histories and basaltic source regions

## Depth and extent of melting

Modern Ethiopian rift basalts can be distinguished from the Oligocene continental flood basalts on the basis of major and trace element geochemistry, whilst the radiogenic isotopic compositions of the two groups show greater variability and, perhaps as a result, greater overlap. We first explore the extent to which the bulk geochemical differences reflect spatial and temporal variations in processes associated with melt extraction and emplacement, or whether they instead require distinct suites of lithospheric and sublithospheric source domains.

Primitive recent Ethiopian rift basalts have bulk compositions that can be derived by melting fertile spinel peridotite as suggested, for example, on the basis of their REE profiles (Fig. 3). The SiO<sub>2</sub> and FeO\* contents of these lavas (backcorrected for olivine fractionation; Fig. 6) plot near the range of experimentally derived melts of fertile peridotite at a pressure of c. 30 kb (Baker & Stolper 1994; Kushiro 1996). The associated depth of c. 100 km is greater than the lithospheric thickness inferred on the basis of mantle tomography (e.g. Bastow et al. 2005; Tiberi et al. 2005). The melting pressures inferred for nepheline-normative lavas in this manner are slightly but consistently greater than those obtained for hypersthene-normative basalts (Fig. 6), although there is some overlap between the groups. A similar pattern is observed among young Djibouti lavas: with lower degrees of silica samples saturation indicate melting at greater depths than those inferred for hypersthene- and quartznormative MgO-rich lavas (Fig. 6).

Among the Oligocene flood basalts, inferred pressures of melting range from 15 to  $\geq$  30 kb for hypersthene- and quartz-normative samples, and from 30 to >40 kb for nepheline-normative lavas. The more highly silica-saturated lavas also yield calculated primary melt compositions with higher SiO<sub>2</sub> and FeO\* relative to the nepheline-normative lavas, consistent with their derivation by greater degrees of melting (cf. Kushiro 1996; Wasylenki *et al.* 2003). A more realistic picture of the melting process provides for polybaric melting, i.e. vertical aggregates of melts derived over a range of depths within a melting column that extends over tens of kilometres. The pressure



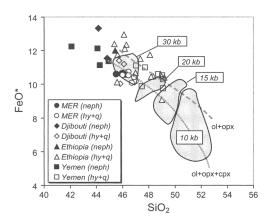


Fig. 6. FeO\* versus SiO<sub>2</sub> for selected mafic lavas and experimental melts in equilibrium with mantle peridotite (Baker & Stolper 1994; Kushiro 1996). The dotted line connects experimental runs for melts coexisting with cpx + oliv + opx, and the dashed line connects experimental melts coexisting with oliv + opx after higher degrees of melt removal. Major element compositions were renormalized with total iron as FeO, then corrected for olivine fractionation to Mg#  $\sim$  72; only samples with Mg# > 62 were corrected using this procedure. Experimental data are plotted as reported, and are shown in the enclosed fields. Filled symbols indicate calculated primary melts for lavas with nepheline-normative compositions; open symbols indicate starting compositions that are quartz- or hypersthene-normative. Calculated primary melts for nepheline-normative lavas indicate consistently higher pressures and lower extents of melting than the more silica-saturated compositions. Sources of data: Deniel et al. 1994; Baker et al. 1996a; Pik et al. 1999; Kieffer et al. 2004.

estimates presented here provide simply a framework for comparison. Interestingly, there is no geographical control on the calculated melting depths, as lavas from Ethiopia and Yemen indicate comparable melting conditions. Most of the low-Ti Oligocene lavas are too evolved to be evaluated using this calculation scheme, as they have MgO contents below 7 wt.% and have undergone extensive fractionation of olivine  $\pm$  clinopyroxene  $\pm$  plagioclase feldspar (Pik *et al.* 1998, 1999).

## Identifying lithospheric and sub-lithospheric source regions

Prior studies of the Ethiopia–Yemen continental flood basalts and Plio-Quaternary lavas from the Afar region have identified several distinct source domains that contribute to magma genesis. To first order, these components include the Afar plume, one or more additional mantle reservoir(s), and a subordinate contribution from the continental crust or mantle lithosphere (Hart et al. 1984; Chazot & Bertrand 1993; Baker et al. 1996a, 2000; Pik et al. 1999; Kieffer et al. 2004). The Afar plume is broadly thought to include a component with  $^{87}$ Sr/ $^{86}$ Sr ~0.704,  $^{143}$ Nd/ $^{144}$ Nd  $\sim 0.51295$ ,  $^{206}$ Pb/ $^{204}$ Pb  $\sim 18.8$  and  $^{3}$ He/ $^{4}$ He up to 19 R/R<sub>A</sub> (Hart et al. 1989; Schilling et al. 1992; Vidal et al. 1991; Marty et al. 1993, 1996; Deniel et al. 1994; Scarsi & Craig 1996; Pik et al. 1999), but the compositions of the remaining source regions are poorly constrained. For example, Hart et al. (1989) infer an enriched mantle component, while Pik et al. (1999) infer the presence of both MORB-like mantle and a second depleted component.

One reason for the lack of consensus is that Pb isotope compositions vary widely among Ethiopian plateau and rift lavas, and tend not to correlate well with Sr–Nd systematics (Fig. 5). Although Pb isotopic compositions of mafic melts are often heterogeneous at all scales, presumably reflecting melt segregation and aggregation processes (Saal *et al.* 1998; Kent *et al.* 2002; Bryce & DePaolo 2004), there are consistent spatial and temporal patterns among the lavas that correlate with both geographical and geochemical parameters, bringing us closer to an integrated understanding of the source regions.

Following Hart (1984), we use the dimensionless parameter  $\Delta 8/4$  to systematize variations in Pb isotopic composition ( $\Delta 8/4$  indicates the difference in <sup>208</sup>Pb/<sup>204</sup>Pb of a sample relative to the Northern Hemisphere Reference Line; see Table 1); this parameter itself has no physical meaning but provides a framework within which to evaluate chemical distinctions among and between sample suites. Variations in  $\Delta 8/4$ -  $^{87}$ Sr/ $^{86}$ Sr among the flood basalts and recent mafic lavas show clear regional trends that can be interpreted in terms of source regions sampled by the melting process (Fig. 7; Baker et al. 1996a). The modern Afar plume is well-constrained by the isotopic compositions of MgO-rich lavas from Erta 'Ale and Djibouti, and the remaining samples form linear arrays that radiate from this region of isotopic space.

Recent lavas from the Ethiopian rift and Djibouti define a trend towards high, EM1-like <sup>87</sup>Sr/<sup>86</sup>Sr and  $\Delta 8/4$ . Samples in this group have La/Nb-Ba/Nb values that are not dramatically elevated relative to primitive mantle, so they indicate a radiogenic source region with only moderately enriched incompatible trace element abundances, interpreted as the lower crust (Baker *et al.* 1996*a*; Trua *et al.* 1999). A second well-defined trend among flood basalts from Ethiopia and Yemen extends towards high, EMII-like <sup>87</sup>Sr/<sup>86</sup>Sr and low  $\Delta 8/4$  values. This trend was interpreted by Baker *et al.* (1996a) as indicating an upper crustal component. The



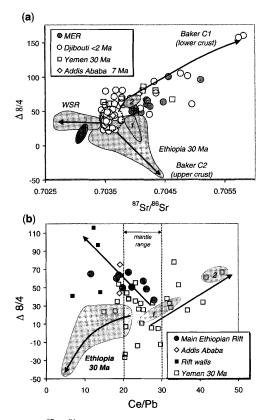


Fig. 7.  $^{87}Sr/^{86}Sr$  and Ce/Pb vs.  $\Delta8/4$  require regional and temporal differences in source components contributing to mafic volcanism in the Afar region. Prior to plotting, samples from Yemen where crustal contamination could be demonstrated through anomalously high values of  ${}^{207}\text{Pb}/{}^{204}\text{Pb}$  and/or  $\delta^{18}\text{O}$ (Baker et al. 1996a, 2000) were removed. (a) Regional sample suites define tight arrays that extend from the Afar plume (dotted circle; data from Djibouti (Deniel et al. 1994) and Erta' Ale (Barrat et al. 1998) towards distinct additional source regions. Labelled components from Baker et al. (1996a) are discussed in the text. Fields labelled 1 and 2 indicate primitive high-Ti Oligocene lavas from Ethiopia (Pik et al. 1999) and Yemen (Baker et al. 1996a), respectively; field 3 encircles primitive Oligocene Adigrat basalts (Pik et al. 1999). Data on mid-ocean basalts from the nearby West Sheba Ridge (WSR) are from Schilling et al. (1992); Main Ethiopian Rift (MER) samples include lavas in Table 1 and data from Trua et al. (1999) and Peccarillo *et al.* (2003). (b) High  $\Delta 8/4$ components recorded in young Main Ethiopian Rift (MER) lavas and Oligocene Yemen continental flood basalts (Baker et al. 1996a) can be resolved by their distinct Ce/Pb values. These data indicate distinct source components in each region that lie outside the range of mantle-derived liquids (Hofmann et al. 1996). All samples plotted here have >6.5 wt% MgO. Fields labelled 1 and 2 indicate primitive high-Ti Oligocene lavas from Ethiopia (Pik et al. 1999) and Yemen (Baker et al. 1996a), respectively. See text for discussion.

high Ba/Nb-La/Nb values of these samples suggest instead that the high <sup>87</sup>Sr/<sup>86</sup>Sr component may reside in the enriched mantle lithosphere; low degrees of melting can also produce variations in these parameters, but the quartz- and hypersthene-normative compositions of these lavas are not consistent with low degrees of mantle melting. There is evidence that the source region was metasomatized to a small degree by hydrous fluids: the high Ba and low K contents of the lavas relative to Kone basalts suggest the presence of amphibole. Interestingly, the primitive high-Ti lavas interpreted by Pik et al. (1999) as representing the essential Oligocene plume (HT2 series) also plot within this array, suggesting they include a contribution from hydrated mantle lithosphere. The high degrees of melting inferred for both of these sample groups makes assimilation of surrounding lithologies during ascent physically plausible.

The third distinct trend observed among the mafic lavas examined here is defined by recent sea-floor basalts from the West Sheba Ridge near Djibouti (Schilling et al. 1992). These samples define an array trending from the modern plume composition towards lower <sup>87</sup>Sr/<sup>86</sup>Sr at near-constant  $\Delta 8/4$ . We interpret this trend to indicate involvement of the convecting upper-mantle source region for mid-ocean ridge basalts. Note that the high-Ti lavas from Adigrat plot along this array and are clearly isotopically different from all other lavas observed in the Oligocene through Recent Ethiopian volcanic provinces. We emphasize that the volume of these lavas is small, and their occurrence rare, but they provide insight into a source component apparently sampled only during flood basalt volcanism.

To resolve the additional components involved in long-term Afar volcanism we incorporate trace element evidence, as relative abundances of incompatible trace elements are sensitive to the mineralogy of specific source regions. Variations in Ce/Pb against  $\Delta 8/4$  (Fig. 7) show clearly the trend of crustal involvement for Ethiopian Oligocene basalts identified above. Interestingly, trends for modern Ethiopian rift lavas and Oligocene Yemen flood basalts can be distinguished from one another on the basis of their distinct Ce/Pb values. From this diagram, we conclude that the crustal source sampled by the modern Ethiopian rift lavas has been available throughout development of the rift, whereas the Yemen rocks record involvement of a separate crustal component that is absent in Ethiopia.

We can also combine our understanding of the melting conditions with the isotopic characteristics of the basalts. In each area, basalts derived at higher pressures and/or through lower degrees of melting

(i.e. nepheline-normative compositions) have more depleted Sr and Nd isotopic signatures than corresponding lavas that are derived at more shallow depths and/or through greater extents of melting (Fig. 8). This observation reinforces the existence of an extended melting column within the mantle, and helps constrain the occurrence of different source domains in space and time.

The integrated geochemical evidence allows us to identify the source components involved in regional volcanism over the past 30 million years. First, Oligocene through Recent basalt volcanism in Ethiopia, Yemen and Djibouti includes a contribution from a common composition which we interpret as the Afar plume (Fig. 8). This component is identifiable on the basis of Sr-Nd-Pb isotopes and incompatible trace element abundances in

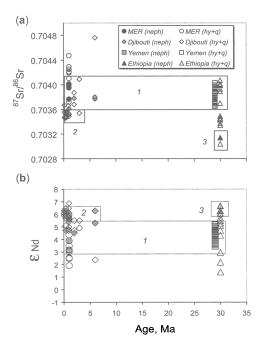


Fig. 8. Temporal and spatial variations in Sr–Nd isotopic compositions of primitive (>6.5 wt% MgO) lavas from Ethiopia, Yemen and the modern Afar plume indicate three source components present at low degrees of mantle melting. Each of these source components is isotopically unradiogenic relative to those recorded in lavas derived by higher degrees of melting. Box 1 encompasses a compositional range similar to bulk silicate earth that is found in lavas of all ages, interpreted here as representing the core of the Afar plume. Box 2 is a MORB-like composition found in ocean floor basalts and lavas from Djibouti (Deniel *et al.* 1994) and Erta' Ale (Barrat *et al.* 1998). Box 3 encloses lavas from Adigrat, Ethiopia (Pik *et al.* 1999). See text for discussion.

basalts of all ages from throughout the region (Figs 4 and 5). Its persistence through time suggests that it forms the core of the upwelling mantle plume, and its occurrence in melts formed at high pressures indicates that it is a fertile composition. This plume composition appears to be nearly constant in both space and time, and that all of the regional trace element and isotopic data form binary arrays that radiate from this composition towards geochemically distinct end-members (cf. Schilling *et al.* 1992).

Three compositionally distinct source domains can be identified on the basis of the mixing trajectories. One is the depleted mantle (DMM), observed both in recent lavas from Djibouti and Erta 'Ale and in primitive Oligocene lavas from Adigrat, Ethiopia (Fig. 8). We also identify an enriched mantle (EMI-like) or crustal component in both low- and high-Ti mafic lavas from the entire Ethiopian plateau; these lavas are characterized by higher La/Nb-Ba/Nb values and require higher timeintegrated parent-daughter ratios to account for their more radiogenic isotope compositions relative to the plume or DMM. A second enriched component (EMII-like) is recorded in within-rift and rift wall basalts from central Ethiopia and Djibouti, the off-rift volcanics from the Addis Ababa series, and the Oligocene Yemen basalts. We concur with Baker et al. (1996a) and Trua et al. (1999) that this source is located in the lower crust; alternatively the enriched source may represent mafic magmas emplaced during the earliest stage of flood basalt activity (e.g. Mackenzie et al. 2005; Maguire et al. 2006).

Two observations about the enriched source domains and their occurrence are significant. First, the enriched mantle contributions are found only in lavas derived from moderate degrees of mantle melting. Secondly, each enriched mantle composition forms a unique mixing array with the Afar plume. Together, these observations indicate that the plume records melting of chemically distinct source domains in the lithosphere and asthenosphere. This conclusion is consistent with numerical models that suggest an ascending plume will melt or displace, rather than entrain, surrounding and overlying mantle material (Farnetani *et al.* 2002).

#### Geodynamic evolution of the region

#### Documenting the head-to-tail transition

Numerical modelling of thermochemical plumes derived from a deep mantle thermal boundary layer predicts that geochemical domains present within a plume head will also be preserved in magmatism associated with the plume tail (Farnetani *et al.* 

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2002). This prediction results in part from the observation that ascending plume material displaces, rather than entrains, surrounding ambient upper mantle, so heterogeneities are thinned and stretched, preserving discrete flow paths within the plume core. We use this numerical prediction, derived for vigorous plumes emplaced beneath oceanic lithosphere, as a framework to assess geochemical heterogeneity in mafic lavas derived from the Afar plume throughout its 30 million-year history. Long-lived spatial variability in plume tails has been documented in Hawaii (e.g. Lassiter & Hauri 1998; DePaolo et al. 2001; Blichert-Toft et al. 2003; Mukhopadhyay et al. 2003; Kurz et al. 2004; Abouachami et al. 2005; Bryce et al. 2005) and the Galápagos (Geist 1992; White et al. 1993; Hoernle et al. 2000; Harpp & White 2001; Geldmacher et al. 2003; Thompson et al. 2004). There are good reasons to expect that the results from Afar may differ from those observed in the oceanic setting: the Afar plume has a substantially lower buoyancy flux than those calculated for Hawaii, Galápagos and Iceland (Ebinger & Sleep 1998) and was emplaced beneath thick continental lithosphere. In addition, the sampling density in the Ethiopia-Yemen flood basalt province is not yet sufficient to resolve consistent spatial variations in lava chemistry. Nonetheless, our results will help test the robustness of the numerical models developed for oceanic hot spots.

The new data presented here indicate temporal trends in source availability and mixing that in turn provide insight into the long-term evolution of the Afar plume system. The Oligocene flood basalts in Ethiopia form the thickest sequence, indicating the highest flux of both material and heat. Significantly, these lavas show the greatest and most uniform contribution from the lithospheric mantle, indicating high degrees of melting over a broad region. This lithospheric source region apparently does not play a major role in generating the recent MER basalts, suggesting it has been thinned and/or partially removed under the rift axis itself. In contrast, lithospheric geochemical signatures dominate the chemistry of lavas from highly extended portions of southwestern Ethiopia (Stewart & Rogers 1996; George et al. 1998; Rogers et al. 2000) and much of the Kenya rift (e.g. Macdonald 1994; Späth et al. 2001), indicating that lithospheric removal is not widespread.

The geochemical evidence suggests that the total magmatic and thermal flux of the Afar plume has waned through time. Modern mafic lavas from Djibouti, Erta' Ale and the Main Ethiopian Rift often show evidence of crustal interaction, presumably occurring in magma chambers within the crust (e.g. Ayalew *et al.* 2006). In the modern rift, low magma supply rates over the past 3 million years led to development of nested silicic calderas at,

for example, Kone and Gedemsa. We suspect that the low magma flux calculated for the modern Afar plume relative to Hawaii and the Galápagos (Ebinger & Sleep 1998) means that it may not be feasible to correlate quantitatively any spatial variations in the chemical domains over the life of the Afar plume.

# Geochemical evidence for one- and two-plume models

The recognition of long-lived geochemically consistent contributions to magmatism in the Ethiopia-Yemen region supports models that call upon the Afar plume to be the source for both Oligocene (plume head) and recent (plume tail) magmatism. The broad distribution of long-lived volcanism in Ethiopia and portions of the East African rift has led to various models of plume structure. The suggestion that there are two discrete plumes (George et al. 1998; Rogers et al. 2000) was based on Sr-Nd isotopic compositions of lavas from southern Ethiopia and the Kenya rift. Although appealing in some ways, this model has two difficulties that have not been overcome: (1) it does not adequately account for the distribution of lavas within the EARS, as no viable plume track exists for the proposed Kenya plume; and (2) the south Ethiopian lavas were primarily derived from the lithosphere and thus provide little insight into sub-lithospheric processes (Furman et al. 2004).

The single-plume model proposed by Ebinger & Sleep (1998) places a plume head beneath southern Ethiopia at *c*. 45 Ma and attributes widespread Cenozoic magmatism (e.g. Hoggar and Tibesti massifs, the modern Afar area) to the lateral transport of plume material along zones of thinned lithosphere. This model benefits from physical simplicity, but is not immediately consistent with recent geophysical studies indicating possible plume stems beneath both Afar and the Tanzania craton (Debayle *et al.* 2001; Nyblade *et al.* 2000; Weeraratne *et al.* 2003; Montelli *et al.* 2004).

Similarly, the consistent Pb isotope differences between contemporaneous sub-lithospheric magmas erupted in Ethiopia and Turkana (Furman *et al.* 2004, 2005) are compatible with the geophysical evidence for separate plume stems beneath these two areas. Turkana lavas have a distinctive high  $^{206}$ Pb/ $^{204}$ Pb signature that is not observed in mafic lavas associated with the Afar plume. This characteristic Pb isotopic signature is, however, found in other areas associated with the South African superplume (e.g. St Helena – Chaffey *et al.* 1989; South Africa – Janney *et al.* 2002). Over the long history of volcanism in Ethiopia

and Turkana, only basalts from the 23 Ma Choke shield volcano have Pb isotope compositions that plot within the Turkana array (Fig. 5; Kieffer *et al.* 2004). These data are intriguing because they may indicate northward magma transport associated with volcano-tectonic activity at 23 Ma in the Turkana area (Furman *et al.* 2006).

The isotopic data thus cannot discriminate between models invoking two unrelated plumes (e.g. George et al. 1998) or one large heterogeneous plume with multiple stems as predicted by numerical and experimental results (Ishida et al. 1999; Davaille et al. 2003). The clear geophysical evidence for the South African superplume (Ritsema et al. 1999; Gurnis et al. 2000; Zhao 2001; Ni et al. 2002) appears to favour the latter scenario, although the shallow mantle structure cannot yet be resolved in sufficient spatial detail. The two small isotopically distinct plume stems may rise from the superplume and interact with regionally unique crustal and sublithospheric sources during ascent. Tomographic investigation of plume pairs (Azores and Canaries, Ascension and St Helena) indicates that two areas of upwelling can merge at mantle depths over 1000 km, thus tapping a common source at depth (Montelli et al. 2004). The large physical dimension of the South African superplume-it represents the largest-known heat and mass transfer from the lowermost portion of the mantle (e.g. Romanowicz & Gung 2002) makes it likely that a variety of source materials coexist within the ascending region, and hence two or more chemically distinct plume heads could come from the same major deep mantle thermal and chemical anomaly.

## Conclusions

Mafic lavas from Ethiopia and Yemen record 30 million years of volcanism associated with the Afar plume. Modern Ethiopian rift basalts have major and trace element geochemical features that are distinct from those of the Oligocene continental flood basalts, whereas the isotopic characteristics of these groups show greater overlap. The geochemical differences can be interpreted in terms of melting processes (depth and degree of melting), as well as mixing of melts of the Afar plume and from mantle sources with three end-member compositions. Taken together, the geochemical evidence indicates that the Afar plume is a long-lived feature of the mantle and that its fundamental isotopic composition has effectively not changed during the transition from Oligocene plume head volcanism to modern plume tail activity. Temporal changes in Ethiopian volcanic geochemistry thus appear to reflect the evolving tectonic environment

rather than a significant change in sub-lithospheric source composition.

Specifically, we find that:

1. Oligocene lavas are generally derived by greater degrees of melting than the recent basalts. Most of the flood basalts are hypersthene- and quartz-normative, while the younger lavas include a higher proportion of nepheline-normative compositions.

2. The isotopic compositions of Ethiopian and Yemeni basalts of all ages are linked with the depth and degree of melting: the Sr and Nd isotopic signatures of lavas derived from smaller degrees of melting and/or melting at higher pressure are more depleted than those signatures found in lavas derived at shallower depths and/or through greater extents of melting.

3. The Sr- and Pb-isotopic signatures of Oligocene through Recent mafic lavas define three distinct binary mixing arrays that radiate from a common composition interpreted as the Afar plume. The end-members involved in mixing include both depleted (DM, interpreted as the convecting upper mantle source region for mid-ocean ridge basalts) and enriched (EMI-like, inferred to be within the lower crust and EMII-like, inferred to reside within the upper crust) mantle components, each of which is observed in a geographically restricted area.

4. Lack of ternary mixing among these components suggests that the ascending plume melts chemically distinct local source domains in the overlying asthenosphere and lithosphere, rather than entraining and incorporating ambient mantle on a regional scale.

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## Appendix

## Supplemental material: analytical procedures

E99- and KO-series lavas and samples AA99-6, -19 were analysed at the Centre Récherches

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Pétrographiques et Géochimiques, Nancy, France by G. Yirgu (major elements by ICP-AES and trace elements by ICP-MS). All other lavas were analysed for major and minor elements (including Ba and Sr) by DCP on an ARL-Fisons Spectraspan 7 and for  $P_2O_5$  and all trace elements by ICP-MS using a VG PlasmaQuad-3 at Duke University (M. Rudnicki and G. Dwyer, analysts). Precision based on replicate analyses of samples and natural basalt standards are generally <1% for SiO<sub>2</sub>, Sr, Y, Zr, Nb, La, Ce; <3% for other major elements, Ba, Sr, Rb, Cs, Cr, Sc, V, Co, Pr, Nd, Sm, Eu, Gd, Tb, Dy, Ho, Er, Hf and Ta; <5% Ni, Yb, Lu, Pb, Th and <8% for U. All oxides are presented as wt %; trace elements are given as ppm.

All Pb analyses, a majority of the Nd analyses, and several Sr analyses were carried out in the clean laboratories and analytical facilities at San Diego State University. These Sr, Nd, and Pb isotope ratios were determined from a single HF-HNO<sub>3</sub> dissolution of approximately 0.4 g of rock powder. Pb was separated according to a twocolumn anion exchange chromatographic procedure described by Hanan & Schilling (1989). Nd and Sr elemental concentrations were subsequently carried out using procedures modified from Schilling et al. (1994), with Eichrom Ln-Spec resin for the rare earth separation, and Eichrom Sr-Spec as a final step for cleaning up the Sr. The total analytical blanks for these procedures were <90 pg Pb, <200 pg Nd, and <300 pg Sr. Thus no blank corrections were made to the data reported in Table 1. At SDSU, the Nd and Pb isotope ratios were measured on the Nu Plasma HR by multicollector inductively coupled plasma mass spectrometry (MC-ICP-MS). Sr isotope ratios were measured using VG Instruments Sector 54 sevencollector thermal ionization mass spectrometer. The <sup>143</sup>Nd/<sup>144</sup>Nd and <sup>87</sup>Sr/<sup>86</sup>Sr ratios were normalized to correct for mass fractionation using values of  ${}^{146}$ Nd/ ${}^{144}$ Nd = 0.7219 and  ${}^{88}$ Sr/ ${}^{86}$ Sr = 0.1194, respectively. The <sup>143</sup>Nd/<sup>144</sup>Nd of AMES Nd standard =  $0.512118 \pm 0.000003$ , and the NIST SRM 987 Sr standard =  $0.710219 \pm 0.000005$ (all  $2\sigma/n$ ). Sr isotopic data in Table 1 are renormalized to NIST SRM 987 = 0.710250. The Nd reported in Table 1 is given relative to La Jolla Nd = 0.511 816. Pb isotope compositions analysed by MC-ICP-MS on the Nu Plasma HR at SDSU using the Tl-doping technique with NIST SRM 997 Tl and  $^{205}$ Tl $^{203}$ Tl = 2.3889 (White *et al.* 2000; Thirlwall 2002; Albarède et al. 2004). NIST SRM981 was used to correct the Pb isotope data for fractionation and machine bias (White et al. 2000). The NIST 981 standard was run every second or third analysis. The final Pb data were adjusted to the Todt et al. (1996) values for NIST 981 relative to the bracketing standard runs. The

NIST SRM 981 Pb standard averaged (with  $2\sigma/n$ ) <sup>206</sup>Pb/<sup>204</sup>Pb = 16.9372 (0.0006), <sup>207</sup>Pb/<sup>204</sup>Pb = 15.4915 (0.0006), and <sup>208</sup>Pb/<sup>204</sup>Pb = 36.6957 (0.0016) during the analysis period. The uncertainties for the Pb isotope ratios, representing error propagation of the in-run  $2\sigma/n$  analytical errors plus the  $2\sigma/n$  reproducibility of the standards, were less than 0.002 for <sup>206</sup>Pb/<sup>204</sup>Pb and <sup>207</sup>Pb/<sup>204</sup>Pb and <0.006 for <sup>208</sup>Pb/<sup>204</sup>Pb.

A portion of the Sr and Nd isotopic measurements presented in Table 1 were carried out in the W. M. Keck Foundation Center for Isotope Geochemistry at UCLA. Approximately 200-250 mg aliquots of whole rock powders were digested in a HF-HNO3 mixture. Samples for Sr isotopic study were prepared using standard ion chromatographic separation techniques employing cation exchange resins. Rare earth separations were carried out using a procedure involving a second cation ion exchange resin. Purified Sr salts were loaded in Ta<sub>2</sub>O<sub>5</sub> on rhenium filaments for analysis. The strontium isotopic measurements were made via thermal ionization mass spectrometry (TIMS) on a VG sector-54 multicollector mass spectrometer operating in dynamic mode. Isotopic fractionation was corrected with an internal normalization of  ${}^{86}$ Sr/ ${}^{88}$ Sr = 0.1194. The average for NIST SRM 987 measured during the course of these analyses was  $0.710238 \pm 0.000027$  (2 $\sigma$ ). The  ${}^{87}\text{Sr}/{}^{86}\text{Sr}$ values reported in Table 1 are normalized to NBS-987 values of 0.710 250. Neodymium isotopic measurements were also carried out TIMS on a Sector 54-30 multicollector operating in dynamic mode; Nd isotopic compositions were normalized to  $^{146}$ Nd/ $^{144}$ Nd = 0.7219 to correct for within-run fractionation. Standards run over the course of the analyses provided mean values (with  $2\sigma$ ) of the La Jolla standard (n = 5) of  $^{143}$ Nd/ $^{144}$ Nd = 0.511 830 ± 0.000 017 (2 $\sigma$ ). Two replicate samples were measured for Nd isotopic compositions in both the UCLA and SDSU facilities. These samples (N22 and E43) agreed in  $^{143}$ Nd/ $^{144}$ Nd values within 0.000 022, or well within the overlapping uncertainties. The values reported in the table for these two samples are those evaluated at SDSU. For the sake of consistency, the UCLA Nd isotopic data are normalized to the SDSU values for the La Jolla Nd standard (i.e.  ${}^{143}$ Nd/ ${}^{144}$ Nd = 0.511 816).

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