Subduction-modified oceanic crust mixed with a depleted mantle reservoir in the sources of the Karoo continental flood basalt province

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Abstract

The great majority of continental flood basalts (CFBs) have a marked lithospheric geochemical signature, suggesting derivation from the continental lithosphere, or contamination by it. Here we present new Pb and Os isotopic data and review previously published major element, trace element, mineral chemical, and Sr and Nd isotopic data for geochemically unusual mafic and ultramafic dikes located in the Antarctic segment (Ahlmannryggen, western Dronning Maud Land) of the Karoo CFB province. Some of the dikes show evidence of minor contamination with continental crust, but the least contaminated dikes exhibit depleted mantle –like initial εNd (+9) and 187Os/188Os (0.1244–0.1251) at 180 Ma. In contrast, their initial Sr and Pb isotopic compositions (87Sr/86Sr = 0.7035–0.7062, 206Pb/204Pb = 18.2–18.4, 207Pb/204Pb = 15.49–15.52, 208Pb/204Pb = 37.7–37.9 at 180 Ma) are more enriched than expected for depleted mantle, and the major element and mineral chemical evidence indicate contribution from (recycled) pyroxenite sources. Our Sr, Nd, Pb, and Os isotopic and trace element modeling indicate mixed peridotite-pyroxenite sources that contain ~10–30 % of seawater-altered and subduction-modified MORB with a recycling age of less than 1.0 Ga entrained in a depleted Os-rich peridotite matrix. Such a source would explain the unusual combination of elevated initial 87Sr/86Sr and Pb isotopic ratios and relative depletion in LILE, U, Th, Pb and LREE, high initial εNd, and low initial 187Os/188Os. Although the sources of the dikes probably did not play a major part in the generation of the Karoo CFBs in general, different kind of recycled source components (e.g., sediment-influenced) would be more difficult to distinguish from lithospheric CFB geochemical signatures. In addition to underlying continental lithosphere, the involvement of recycled sources in causing the apparent lithospheric geochemical affinity of CFBs should thus be carefully assessed in every case.

Keywords: Large igneous province; Continental flood basalt; Karoo; Picrite; Mantle source; Crustal recycling

1. Introduction

Continental flood basalts (CFBs) represent the most voluminous magmatic activity on the continents. They are commonly associated with the early stages of continental breakup, but whether they arise due to processes related to the continental lithosphere (e.g., thinning, delamination, and insulation) or instead derive from melting of a deep mantle plume, remains an issue of discussion (e.g., Anderson, 2005; Beccaluva et al., 2009; Campbell, 2005; Coltice et al.,
2. Geological and geochemical context

The Karoo CFBs erupted on the landmasses of Africa and Antarctica, both then part of the Gondwana supercontinent, at 184–178 Ma (Fig. 1; Jourdan et al., 2005). The magmas intruded through thick continental lithosphere that consists of a variety of Archean to Paleozoic rocks.
2.1 Pre-Jurassic geology of western Dronning Maud Land

In western Dronning Maud Land, the NW portion of the area is dominated by the Archean Grunehogna craton (Fig. 1; Krynauw et al., 1991; Wolmarans and Kent, 1982). The Archean basement is only exposed at Annandagstoppane (Fig. 1; Marschall et al., 2010) and elsewhere is covered by metamorphosed Mesoproterozoic supracrustal rock types belonging to the Ritscherflya Supergroup and/or by Borgmassivet mafic intrusions (Krynauw et al., 1988, 1991; Riley and Millar, in press; Wolmarans and Kent, 1982). The southern and eastern parts of the Precambrian basement of western Dronning Maud Land belong to the Proterozoic Maud Belt (Fig. 1; Groenewald et al., 1995). Late Paleozoic sedimentary rocks that overlay the basement are exposed at Vestfjella, Heimefrontfjella, and southwest Kirwanveggen (Fig. 1; e.g., Juckes, 1972; Wolmarans and Kent, 1982).

2.2. The Karoo CFBs and related intrusive rocks of Antarctica

The Karoo CFBs, exposed at Vestfjella, Kirwanveggen, and Heimefrontfjella, represent the youngest preserved rock unit in western Dronning Maud Land (Fig. 1). Associated intrusive rocks are more widespread and can also be found crosscutting the basement at Ahlmanryggen, Mannefallknausane, and H. U. Sverdrupfjella (Fig. 1). The mafic to ultramafic CFBs and intrusive rocks show notable geochemical heterogeneity and can be grouped into various low-Ti and high-Ti magma types in terms of their trace element and isotopic composition (Luttinen and Furnes, 2000; Luttinen et al., 2010; Riley et al., 2005). Unlike from the African parts of the Karoo CFB province, several dikes of unusual sublithospheric geochemical character have been described from Antarctica: the Sr, Nd, Pb, and Os isotopic compositions of the Vestfjella

Fig. 1. Outcrop map of western Dronning Maud Land from Vestfjella to H. U. Sverdrupfjella. Distribution of Karoo flood basalts and Ahlmannryggen Group 3 dikes is shown. Lithospheric boundary between Grunehogna craton and Maud belt is after Corner (1994). Distribution of Karoo flood basalts and related intrusive rocks (outside the flood basin areas) in reconstructed Gondwana supercontinent (cf. Heinonen et al., 2010) is shown in the inset.
depleted ferropicrite suite (Fig. 2; Heinonen and Luttinen, 2008, 2010) are indistinguishable from those of SWIR MORBs and imply derivation from an upper mantle source (Heinonen et al., 2010), although non-chondritic primitive mantle sources may also be plausible (Jackson and Carlson, 2011). Incompatible element depleted low-Nb basaltic and picritic dikes also identified from Vestfjella have been interpreted to represent low-pressure, high-degree melting of this same source (Heinonen et al., 2010). The OIB-like Vestfjella enriched ferropicrite suite (Fig. 2; Heinonen and Luttinen, 2008) has been ascribed to either an anomalous lithospheric source or a recycled sediment-influenced pyroxenite mantle source (Heinonen et al., 2010). The depleted Group 3 dikes from Ahlmannryggen are considered in more detail in the following section.

**Fig. 2.** Sr and Nd isotopic characteristics of the Ahlmannryggen Group 3 dikes (Riley et al., 2005) shown at 180 Ma. Data for Vestfjella depleted (D-FP) and enriched (E-FP) ferropicrite suites (Heinonen and Luttinen, 2008; Heinonen et al., 2010), Karoo CFBs (Ellam and Cox, 1989, 1991; Harris et al., 1990; Hawkesworth et al., 1984; Jourdan et al., 2007; Luttinen and Furnes, 2000; Luttinen et al., 1998, 2010; Riley et al., 2005; Sweeney et al., 1994), SWIR MORB (le Roex et al., 1983, 1992; Mahoney et al., 1992), and depleted MORB mantle (DMM; Workman and Hart, 2005) are also presented. The isotopic compositions of SWIR MORB sources and DMM were back-calculated using DMM isotopic ratios after Workman and Hart (2005). Lithospheric contamination models are after Heinonen et al. (2010).

### 2.2.1 Group 3 dikes of Ahlmannryggen

The Karoo dikes of Ahlmannryggen crosscut the Precambrian Ritscherflya metasupracrustal rocks and can be grouped into four geochemical types (Groups 1–4; Riley et al., 2005). Most of the dikes are fairly evolved basalts, but Group 3 and Group 4 include several samples with high MgO (> 8 wt. %). Whereas Group 4 exhibits relatively unradiogenic εNd (from -5 to +2 at 180 Ma), Group 3 shows highly radiogenic εNd (from +5 to +9 at 180 Ma) indicative of derivation from sublithospheric sources (Fig. 2). The Group 3 dikes show only negligible secondary alteration (e.g., average LOI = 1 wt. %) and are generally porphyritic with olivine and/or pyroxene (orthopyroxene and/or augite) phenocrysts surrounded by groundmass consisting of pyroxene, plagioclase, and Fe-Ti oxides. The olivine phenocrysts show Mg-rich (Fo16–90) compositions indicating that the dikes crystallized from primitive magmas (Heinonen et al., 2013). The Group 3 dikes are characterized by high FeOtot (12–14 wt. %) and TiO2 (3.3–4.9 wt. %), low CaO (9–11 wt. %), La/Sm (0.5–0.8 chondrite-normalized), and Nb/Y (0.1–0.3), high εNd (from +5 to +9), and slightly elevated 87Sr/86Sr (0.7035–0.7062) at 180 Ma (Table 1; Fig. 2; Riley et al., 2005). They also show general depletion of strongly incompatible elements, and depletion of large-ion lithophile elements (LILE), U, and Th relative to Nb and Ta. The dikes are not

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significantly altered and LILEs (and $^{87}\text{Sr}/^{86}\text{Sr}$) show coherent behaviour with more immobile elements in general, indicating that post-crystallization modification has been negligible (cf. Riley et al., 2005).

The Group 3 dikes were further divided into two subgroups on the basis of Sr and Nd isotopic composition (Fig. 2) by Riley et al. (2005): the high-$\varepsilon_{\text{Nd}}$ subgroup ($\varepsilon_{\text{Nd}} = +7.0$ to $+9.0$) and $^{87}\text{Sr}/^{86}\text{Sr} = 0.7035-0.7041$ at 180 Ma) was thought to have crystallized from uncontaminated mantle-derived magmas whereas the composition of the low-$\varepsilon_{\text{Nd}}$ subgroup ($\varepsilon_{\text{Nd}} = +5.0$ to $+5.5$ and $^{87}\text{Sr}/^{86}\text{Sr} = 0.7054-0.7062$ at 180 Ma) was thought to have been affected by minor contamination with continental crust (possibly Precambrian Borgmassivet intrusions; Riley et al., 2005). The subgroups are spatially separated by a distance of ~20 km (Fig. 1).

Riley et al. (2005) hypothesized that the Group 3 magmas represent partial melts of a strongly depleted mantle component possibly entrained in a mantle plume. Recently, Heinonen et al. (2013) interpreted the overall geochemical (high Ti, low Ca) and mineral chemical (high-Ni olivine) characteristics of the dikes to indicate a significant role for pyroxene-rich sources in their petrogenesis. $^{40}\text{Ar}/^{39}\text{Ar}$ whole-rock data indicate a disturbed and thus somewhat unreliable plateau age of 187.3 ± 3.6 Ma (sample Z1816.1; Riley et al., 2005). More detailed information on the Group 3 and other Jurassic Ahlmannryggen dikes and their comparisons with other Karoo CFBs are provided in Riley et al. (2005) and Heinonen et al. (2013).

3. Sample selection and analytical methods

Nine Group 3 samples from distinct dike outcrops were selected for Pb and Os isotopic analyses (Table 1). The isotopic measurements were performed at the Department of Terrestrial Magnetism (DTM) of the Carnegie Institution of Washington. The rock samples were extracted with a hammer from the bedrock and subsequently chopped to smaller pieces with a hydraulic press and by hammering the samples under a cloth. Sample pieces that had metal marks on them or were in direct contact with the press were not included. The samples were further crushed in a ceramic jaw crusher and the resulting chips were hand-picked and powdered in an agate or ceramic mill to further avoid contamination with metals. The crushing devices were purified with aliquots of clean quartz between runs. For the chemical treatment of Pb and Os at the DTM, the reader is referred to Heinonen et al. (2010) with exceptions that are listed in Table S1.

Isotopic compositions of Pb and Re for isotope-dilution concentration determinations were measured on the DTM multiple-collector Nu Plasma high resolution inductively coupled plasma mass spectrometer (HR ICP-MS). Pb was measured statically using Faraday cups. Mass fractionation was corrected by comparing bracketing runs of the NBS-981 standard to values reported by Todt et al. (1996). Four standard runs gave the following average isotopic ratios: $^{206}\text{Pb}^{204}\text{Pb} = 16.937 ± 0.005$ (2σ), $^{207}\text{Pb}^{206}\text{Pb} = 15.491 ± 0.005$, and $^{208}\text{Pb}^{204}\text{Pb} = 36.70 ± 0.01$. The uncertainty for Pb for the sample analyses is assigned the external errors measured for the multiple standard analyses, because they are larger than the in-run precisions. The low Re concentrations were measured by simultaneous collection in electron multipliers. Instrument fractionation for Re was estimated by normalizing to bracketing standard runs. Analytical precision for Re is estimated to be 3 %.

The isotopic composition of Os was measured by thermal ionization mass spectrometry (TIMS) using the Thermo-Fisher Triton of DTM. Osmium was loaded on Pt filaments, covered with Ba(OH)$_2$ and run as OsO$_3$$^-$ ions. The measurements were obtained on the single electron multiplier, monitoring interferences from Re, and correcting for fractionation to $^{192}\text{Os}^{188}\text{Os} = 3.083$. Four intervening in-house standard (DTM Johnson Matthey Os) runs gave an average $^{187}\text{Os}^{188}\text{Os}$ ratio of 0.17381 ± 0.00004 (2σ). The uncertainty for Os is assigned to the in-run precisions, because they are larger than the external error.

The extraction techniques and mass spectrometry resulted in total blanks of 100 pg for Pb and < 2 pg for Re and Os that posed negligible corrections for concentrations and isotopic ratios in all cases.
Table 1
Whole-rock major element (normalized to 100% volatile free) and Sr, Nd, Pb, and Os isotopic composition of Group 3 samples from Ahlmannryggen (western Dronning Maud Land, Antarctica). For previously published spatial and chemical data, the reader is referred to Riley et al. (2005) and Heinonen et al. (2013).

<table>
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<th>Sample</th>
<th>Z1812.3</th>
<th>Z1816.1</th>
<th>Z1816.2</th>
<th>Z1813.1</th>
<th>Z1816.3</th>
<th>Z1817.2</th>
<th>Z1803.1</th>
<th>Z1803.5</th>
<th>Z1834.3</th>
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<td>high-εNd</td>
<td>high-εNd</td>
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<td>high-εNd</td>
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<tr>
<td>TiO₂ (wt. %)</td>
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<td>3.34</td>
<td>3.91</td>
<td>3.65</td>
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<td>4.06</td>
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<td>12.87</td>
<td>13.67</td>
<td>13.81</td>
<td>11.08</td>
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<td>12.96</td>
<td>13.04</td>
<td>12.83</td>
</tr>
<tr>
<td>MgO (wt. %)</td>
<td>11.91</td>
<td>14.73</td>
<td>14.83</td>
<td>12.70</td>
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<td>1.77</td>
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<td>0.63</td>
<td>0.97</td>
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<td>0.85</td>
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<td>0.13329</td>
<td>0.13042</td>
<td>0.00006</td>
<td>0.13331</td>
<td>0.13545</td>
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<tr>
<td>Pb (i)</td>
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<td>Pb (l)</td>
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<td>0.13807</td>
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<td>0.13780</td>
<td>0.13777</td>
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<td>Pb (g)</td>
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<td>0.15492</td>
<td>0.15511</td>
<td>0.15347</td>
<td>0.15313</td>
<td>0.15394</td>
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<tr>
<td>Pb (t)</td>
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<td>0.13688</td>
<td>0.13737</td>
<td>0.13749</td>
<td>0.13745</td>
<td>0.13735</td>
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4. Pb and Os isotopic composition of the Group 3 dikes

Pb and Os isotopic data for the Ahlmannryggen Group 3 dikes are shown in Table 1 and illustrated in Figs. 3 and 4. Hereafter, the 206Pb/204Pb, 207Pb/204Pb, 208Pb/204Pb, and 187Os/188Os (and 87Sr/86Sr and εNd) of the Ahlmannryggen dikes and other Karoo CFB-related rocks refer to initial ratios calculated at 180 Ma unless otherwise stated. The radiogenic ingrowth of Pb was calculated using U and Th concentration data from Riley et al. (2005).

The high-εNd subgroup shows a wide range of Pb isotopic compositions (206Pb/204Pb = 17.4–18.4, 207Pb/204Pb = 15.3–15.5, and 208Pb/204Pb = 37.0–37.9). The three samples with the most radiogenic εNd (+9) are the most radiogenic also in terms of Pb isotopic ratios: 206Pb/204Pb of 18.4 is the highest recorded for Karoo CFBs and related intrusive rocks (Fig. 3). The 208Pb/204Pb and 207Pb/204Pb of these three samples are similar to those found in some SWIR MORB (Fig. 3c), but 206Pb/204Pb is lower (and εNd higher) at a given 208Pb/204Pb (Fig. 3d). Their 206Pb/204Pb and 207Pb/204Pb (and 87Sr/86Sr and εNd) are also rather similar to the prevalent mantle component (PREMA) of Zindler and Hart (1986). The low-εNd subgroup shows a more restricted range of Pb isotopic compositions (206Pb/204Pb = 17.7–17.8, 207Pb/204Pb = 15.4, and 208Pb/204Pb = 37.4–37.5) that overlap those of the Karoo CFBs.

The Os isotopic composition correlates negatively with εNd; the high-εNd subgroup shows 187Os/188Os (0.124–0.125) akin to DMM, whereas the low-εNd subgroup shows marginally higher 187Os/188Os of 0.127–0.128 (Fig. 4).

Fig. 3. $^{87}\text{Sr}/^{86}\text{Sr}$ vs. $^{206}\text{Pb}/^{204}\text{Pb}$ (a), $\varepsilon_{\text{Nd}}$ vs. $^{206}\text{Pb}/^{204}\text{Pb}$ (b), $^{207}\text{Pb}/^{205}\text{Pb}$ vs. $^{206}\text{Pb}/^{204}\text{Pb}$ (c), and $^{208}\text{Pb}/^{204}\text{Pb}$ vs. $^{206}\text{Pb}/^{204}\text{Pb}$ (d) compositions of the Ahlmannryggen Group 3 dikes in comparison with Karoo CFBs (Ellam and Cox, 1989, 1991; Jourdan et al., 2007), Vestfjella depleted and enriched ferropicrite suites (Heinonen and Luttinen, 2008; Heinonen et al., 2010), SWIR MORB (le Roex et al., 1983, 1992; Mahoney et al., 1992), depleted MORB mantle (DMM; Workman and Hart, 2005), prevalent mantle component (PREMA, $^{208}\text{Pb}/^{204}\text{Pb}$ not defined; Zindler and Hart, 1986) and non-chondritic primitive mantle (NCPM; cf. Jackson and Carlson, 2011; Jackson and Jellinek, 2013; Pb isotope composition constrained by 4.50 Ga and 4.43 Ga isochrons) at 180 Ma. The compositions of SWIR MORB sources and DMM were back-calculated using DMM isotopic ratios after Workman and Hart (2005) and the Sr and Nd composition of NCPM was back-calculated following Jackson and Jellinek (2013). PREMA at 180 Ma was approximated using E-DMM isotopic ratios after Workman and Hart (2005).

5. Discussion

5.1. Crustal contamination of the Group 3 magmas

The high-$\varepsilon_{\text{Nd}}$ subgroup exhibits the highest $\varepsilon_{\text{Nd}}$ values (up to +9.0) recorded for Karoo CFBs, suggesting that their primary melts were generated in the sublithospheric mantle. Samples with the highest $\varepsilon_{\text{Nd}}$ show elevated $^{87}\text{Sr}/^{86}\text{Sr}$ and $^{208}\text{Pb}/^{204}\text{Pb}$ relative to DMM at 180 Ma (Fig. 3). These are considered primary features (and not caused by in-situ reactions with wall rock or hydrothermal alteration), because $^{87}\text{Sr}/^{86}\text{Sr}$ and $^{208}\text{Pb}/^{204}\text{Pb}$ correlate negatively with, e.g., Th/Ta and La/Nb, and because the samples having $\varepsilon_{\text{Nd}}$ of +9 exhibit homogeneous compositions in terms of mobile trace elements and Sr and Pb isotopic ratios (cf. Riley et al., 2005). In the case of secondary alteration, distinct samples from different dikes would not be expected to form such coherent compositional groups (cf. Fig. 2 and 3).

The new Pb isotopic data reveal two possible contamination trends for the Group 3 dikes: (1) The high-$\varepsilon_{\text{Nd}}$ subgroup shows a rather wide range of Pb isotopic compositions that correlate
negatively with $^{87}$Sr/$^{86}$Sr and positively with $^{143}$Nd (Fig. 5); (2) The low-$^{143}$Nd subgroup exhibits relatively lower $^{143}$Nd and higher $^{87}$Sr/$^{86}$Sr at a given $^{206}$Pb/$^{204}$Pb (Fig. 5). Importantly, the relatively high $^{187}$Os/$^{188}$Os of the low-$^{143}$Nd subgroup (Fig. 4), and the Nd-Pb isotope systematics of the high-$^{143}$Nd subgroup (Fig. 5b; trend not directed towards the lithosphere-signatured CFBs) make SCLM an unlikely contaminant in both cases.

![Fig. 4. $^{143}$Nd vs. $^{187}$Os/$^{188}$Os compositions of the Ahlmannryggen Group 3 dikes in comparison with the Vestfjella depleted and enriched ferropicrites suites (Heinonen and Luttinen, 2008; Heinonen et al., 2010), Mwenezi picrites (Ellam and Cox, 1989; Ellam et al., 1992), OIBs that sample enriched mantle (EM) domains (Eisele et al., 2002; Woodhead and Devey, 1993; Workman et al., 2004), Gondwana SCLM (estimated after mantle xenoliths; Simon et al., 2007; Walker et al., 1989), and depleted MORB mantle (DMM; Shirey and Walker, 1998; Workman and Hart, 2005) at 180 Ma. In the case of mantle reservoirs, the isotopic compositions were back-calculated using $^{187}$Re/$^{188}$Os of 0.06 (DMM) and 0.4 (EM) (cf. Shirey and Walker, 1998) and $^{147}$Sm/$^{144}$Nd of 0.2485 (DMM; after Workman and Hart, 2005), 0.2138 (EM1; after Eisele et al., 2002), and 0.1840 (EM2; after Workman et al., 2004). EC-AFC models for a Group 3 high-$^{143}$Nd parental magma with high-Os and low-Os upper (Archean) crustal contaminant (UC) also illustrated (see Table S2 for parameters).]

In order to constrain crustal contamination of Group 3 dikes, we performed energy-constrained assimilation-fractional crystallization (EC-AFC) modeling (Bohrson and Spera, 2001; Spera and Bohrson, 2001) using a primitive high-$^{143}$Nd sample Z1816.2 as a parental melt composition and a diverse suite of Archean Kaapvaal TTGs, shales, and amphibolites (Kreissig et al., 2000), and Proterozoic Ritscherflya sedimentary rocks (Moyes et al., 1995; Pb after Wareham et al., 1998), Maud Belt gneisses (H.U. Sverdrupfjella; Wareham et al., 1998), and Borgmassivet mafic intrusive rocks (Sr and Nd data after the model of Riley et al., 2005; Pb data by T.R. Riley, unpublished) as contaminants. Details of the contamination modeling are presented in Table S2.

Our EC-AFC modeling indicates that minor (~1 %) contamination of a high-$^{143}$Nd parental magma with an Archean crustal contaminant could plausibly explain the Sr, Nd, and Pb isotopic composition of the three samples that belong to the high-$^{143}$Nd subgroup and show $^{143}$Nd of +7 and relatively unradiogenic Pb isotopic compositions (Fig. 5; cf. Table 1). The low-$^{143}$Nd subgroup, on the other hand, has overly high $^{87}$Sr/$^{86}$Sr and low $^{143}$Nd at a given $^{206}$Pb/$^{204}$Pb to be explained by contamination of a high-$^{143}$Nd parental magma with the majority of the Archean crustal contaminants (Fig. 5); an anomalous TTG contaminant (sample 96/228) is able to produce similar $^{87}$Sr/$^{86}$Sr, $^{143}$Nd, and $^{206}$Pb/$^{204}$Pb at ~1 % of contamination, but cannot explain the higher $^{207}$Pb/$^{204}$Pb and $^{206}$Pb/$^{204}$Pb of the low-$^{143}$Nd subgroup (Fig. 5). Models with felsic Proterozoic contaminants (H. U. Sverdrupfjella gneiss and Ritscherflya sedimentary rock) show a slightly better fit in terms of $^{87}$Sr/$^{86}$Sr, $^{143}$Nd, $^{206}$Pb/$^{204}$Pb, and $^{208}$Pb/$^{204}$Pb, but are not compatible with the lower $^{207}$Pb/$^{204}$Pb of the low-$^{143}$Nd subgroup. Based on our model, the Borgmassivet intrusive suite with relatively high $^{208}$Pb/$^{204}$Pb also appears to be an unsuitable contaminant.
Although the Sr, Nd, and Pb isotopic composition of the low-\(\epsilon_{\text{Nd}}\) subgroup cannot be satisfactorily explained by contaminating a high-\(\epsilon_{\text{Nd}}\) parental magma with aforementioned crustal materials, such a scenario cannot be completely ruled out: our modeling is hampered by limited Pb data on the Proterozoic contaminants (e.g., H.U. Sverdrufpjella and Borgmassivet models based only on a single analysis). The spatially distinct high-\(\epsilon_{\text{Nd}}\) and low-\(\epsilon_{\text{Nd}}\) subgroups may thus have fractionated in separate magma chambers at different crustal levels (cf. Fig. 1). On the other hand, the higher olivine Ni contents and \(^{187}\text{Os}/^{188}\text{Os}\) (Fig. 4) of the low-\(\epsilon_{\text{Nd}}\) subgroup imply that their parental magmas may in fact have been derived from more pyroxene-rich sources than the high-\(\epsilon_{\text{Nd}}\) parental magmas (Heinonen et al., 2013; cf. section 5.2.3).

Given the likelihood of Archean crustal contamination in the case of the high-\(\epsilon_{\text{Nd}}\) samples with \(\epsilon_{\text{Nd}}\) of +7 and the possibility of combined crustal contamination and source heterogeneity in the case of the low-\(\epsilon_{\text{Nd}}\) subgroup, we concentrate on the three uncontaminated high-\(\epsilon_{\text{Nd}}\) samples (Z1812.3, Z1816.1, and Z1816.2 with \(\epsilon_{\text{Nd}}\) of +9) in the following discussion on Group 3 mantle sources.
5.2. Sublithospheric mantle sources of the Group 3 dikes

5.2.1. Evidence for a pyroxene-rich source and its origin

The major element and mineral chemical characteristics of the Group 3 dikes, discussed in detail by Riley et al. (2005) and Heinenon et al. (2013), provide evidence for contribution from pyroxenite sources. Summarizing, picrites with such high TiO₂ (3–5 wt. %) and FeO₁₀ (12–14 wt. %) and low CaO (9–11 wt. %) cannot derive from melting of solely peridotitic mantle (Heinenon et al., 2013; cf. Herzberg and Asimow, 2008; Prytulak and Elliott, 2007; Tuff et al., 2005). The high Ni contents in olivine (0.5–0.6 wt. % at Fo₉₀) and high whole-rock Zn/Fe (1.2–1.5 × 10⁻³) are also indicative of pyroxene-rich sources (Heinenon et al., 2013; cf. Le Roux et al., 2010; Sobolev et al., 2007), although the former may also partly reflect high pressures beneath the Gondwanan lithosphere (cf. Li and Ripley, 2010). Importantly, the negative correlation of whole-rock CaO (at a given MgO) and olivine Ni to primary compositional variation that is compatible with some degree of mixing of pyroxenitic and peridotitic source components (Heinenon et al., 2013).

The pyroxene content of a mantle section can be influenced by local-to-regional scale melt infiltration and metasomatism in the lithospheric mantle (e.g., Bodinier et al., 2008; Liu et al., 2005) or by reactions between mantle peridotite and partial melts of subducted oceanic crust (e.g., Mallik and Dasgupta, 2012; Yaxley and Green, 1998). The low La/Sm and highly radiogenic Nd isotopic signature of the high-εNd subgroup (Figs. 6 and 7) are not readily explained by melting of metasomatized lithospheric mantle (e.g., Obata and Nagahara, 1987) that is expected to be relatively enriched in the most incompatible elements and develop relatively unradiogenic εNd over time. On the other hand, lithospheric mantle pyroxenites with strongly depleted incompatible trace element compositions have been reported from, e.g., the Ronda orogenic peridotite (e.g., Bodinier et al., 2008). These pyroxenites show strong relative depletion of Nb and Ta (e.g., Bodinier et al., 2008; Garrido and Bodinier, 1999), however, and thus cannot be a primary source for the high-εNd subgroup that shows enrichment of Nb and Ta relative to similarly incompatible elements (cf. Fig. 6).

Oceanic crust generally exhibits low Sm/Nd and would thus also develop relatively low εNd over time, unless it had been modified by, e.g., dehydration and/or partial melting related to subduction (e.g., Kogiso et al., 1997; Sakuyama et al., 2013), and does not contain significant amounts of pelagic sediments (Fig. 7; Stracke et al., 2003; cf. Plank and Langmuir, 1998). Given that convergent-margin processing of subducted crust is a widely recognized process, that Nd may be up to three times more mobile in subduction fluids than Sm (Kogiso et al., 1997), and that fluid-immobile Nb and Ta are effectively held in subducted igneous oceanic crust (Kogiso et al., 1997; cf. Rudnick et al., 2000), we conclude that Group 3 pyroxenite sources most likely contained recycled igneous crustal materials. The nature and significance of this recycled component is evaluated in detail in the following sections.

5.2.2. Trace element constraints on the recycled mantle component

In order to model the trace element composition of the recycled component, we used the mean MORB of Gale et al. (2013) as the igneous crust composition. The effects of subduction modification were calculated on the basis of fluid-rock distribution coefficients obtained in dehydration experiments on an MORB-like amphibolite (Kogiso et al., 1997; Table S3). Such a completely modified recycled MORB is referred to as sm-MORB 1, whereas a more mildly (50% less effectively) modified recycled MORB is referred to as sm-MORB 2 (Fig. 6). The high-εNd signature shows a better fit with the sm-MORB 2 component (especially in the case of Pb; Fig. 6) and thus we concentrate on it in the following discussion.

The trace element pattern of sm-MORB 2 (10%) + DMM peridotite (90%) mixture illustrates that a subduction-modified signature is effectively transferred to the ambient mantle even at low portions of entrained recycled material due to low incompatible element contents of DMM (Fig. 6). Furthermore, a partial melt model of the mixture indicates that such a source is capable of producing a melt with trace element characteristics akin to the high-εNd subgroup (Fig. 6). The positive Ba anomaly and more incompatible-element-depleted trace element pattern of the high-εNd subgroup relative to the model could be related to an additional Ba-rich component
(or lower mobility of Ba during subduction) and more depleted compositions of the crustal and/or mantle components than those used in the model, respectively. Nevertheless, given the overall similarities (Fig. 6), we suggest that the incompatible trace element signature of the high-$\varepsilon_{\text{Nd}}$ subgroup can be viably explained by sources that contain subduction-modified MORB.

![Fig. 6](image)

**Fig. 6.** Primitive mantle–normalized (Sun and McDonough, 1989) incompatible trace element diagrams of the uncontaminated ($\varepsilon_{\text{Nd}} = +9$) Group 3 dikes. Average MORB (Gale et al., 2013), variably subduction-modified MORB (sm-MORB 1 with 100% modification, sm-MORB 2 with 50% modification; Kogiso et al., 1997; cf. section 5.2.2.), mixture (9:1) of DMM (Workman and Hart, 2005) and sm-MORB 2, simple modal partial melt model of the mixture (details given in Table S3), and average Ronda Group C websterite (Bodinier et al., 2008) also shown.

### 5.2.3. Isotopic constraints on the recycled mantle component

In order to model the Sr, Nd, and Pb isotopic composition of the recycled MORB component (cf. section 5.2.2.), we used the spreadsheet and standard input parameters provided by Stracke et al. (2003). Due to uncertainties related to initial concentrations, isotopic compositions and behaviour of Re and Os (Carlson, 2005; Stracke et al., 2003), Os isotopes were modeled by simple binary mixing of DMM and MORB, compositions of which were constrained on the basis of data compilations presented in Shirey and Walker (1998). We emphasize that all the model parameters represent average or recommended values; full details of the modeling are presented in Figs. 7 and 8, and in Table S4.

Modeling of isotopes provides further constraints on the petrogenesis of Group 3 dikes (Fig. 7). Using recommended values for the isotopic evolution of the mantle and crustal components (Table S4; cf. Stracke et al., 2003), the best-fit in terms of Nd and Pb isotopic compositions is attained with a mixture of 0.7 Ga recycled sm-MORB 2 (~10–30 %) and DMM (~70–90 %) (Fig. 7b). In the case of Nd isotopes, an even better fit would be attained with a more depleted DMM (cf. Workman and Hart, 2005) or MORB composition (cf. Gale et al., 2013) or if Nd is assumed to be more mobile during subduction (cf. grey curve with sm-MORB 1 component in Fig. 7). The high $^87\text{Sr}/^{86}\text{Sr}$ of the high-$\varepsilon_{\text{Nd}}$ subgroup suggests that the Sr isotopic signature of the recycled component was more radiogenic than in our model (Fig. 7a).
Radiogenic Sr is readily introduced into the upper oceanic crust via replacement by seawater Sr (30% on average; Kawahata et al., 1987) that has also been incorporated into the equations of Stracke et al. (2003). Seawater-influenced (30%) Sr isotope model is compatible with trace element and Nd and Pb isotope models (cf. Figs. 6 and 7).

Fig. 7. $^{87}/^{86}$Sr vs. $^{206}/^{204}$Pb (a), $\varepsilon_{Nd}$ vs. $^{206}/^{204}$Pb (b), $^{207}/^{204}$Pb vs. $^{206}/^{204}$Pb (c), and $^{208}/^{204}$Pb vs. $^{206}/^{204}$Pb (d) compositions of the Ahlmannryggen Group 3 dikes with the results of isotopic modeling of recycled subduction-modified MORB (cf. Fig. 6) at 180 Ma. Black lines illustrate binary mixing curves between DMM and 0.7 Ga and 1.0 Ga sm-MORB 2 (cf. Fig. 6) with an effect of 30% of seawater alteration (sw) shown in Fig. 7a (seawater alteration has negligible effect on other isotope ratios); square dots indicate 10% and 30% of recycled material in the mixture (only shown for 1.0 Ga curve for clarity). Gray unbroken line illustrates binary mixing curve between DMM and 0.7 Ga seawater-altered (30% of Sr replaced) sm-MORB 1 (cf. Fig. 6); square dots indicate 2% and 5% of recycled material in the mixture. Gray stippled lines illustrate the composition of 1 Ga recycled unmodified MORB with an additional sedimentary component (sed; Plank and Langmuir, 1998; cf. Stracke et al., 2003) and its mixing with DMM. See Table S4 for detailed model parameters and references. The compositions of the Vestfjella depleted and enriched ferropicrite suites, Karoo CFBs, non-chondritic primitive mantle (NCPM) and PREMA as in Fig. 3.

For comparison, models with unmodified recycled MORB, additional sedimentary component, as well as pervasively fluid-modified MORB (sm-MORB 1) fail to produce the Sr, Nd, and Pb isotopic composition of the uncontaminated high-$\varepsilon_{Nd}$ subgroup (Fig. 7). Although the model with sm-MORB 1 produces a better-fit for $\varepsilon_{Nd}$ than the model with sm-MORB 2, Sr shows a poor fit even with seawater Sr included. Furthermore, a model with sm-MORB 1 shows radiogenic Pb isotopic composition that would constrain the amount of recycled crust to be less than 2% in the mixture. Such a small portion of recycled component would not be enough to produce the pyroxenite fingerprint observed in the major element geochemistry of the Group 3 dikes (Heinonen et al., 2013).

The low $^{187}/^{188}$Os of the high-$\varepsilon_{Nd}$ subgroup (Fig. 4) is not directly compatible with recycled crustal sources. Recycled MORB sources should develop high $^{187}/^{188}$Os over time (>
because MORBs exhibit high Re (0.5–2 ppb) and low Os (0.001–0.05 ppb) relative to depleted mantle peridotite (0.05–0.14 ppb and 0.8–9 ppb, respectively; Shirey and Walker, 1998). In the case of a mixed source, however, the Os-rich peridotite component will control the Os isotopic composition (e.g., Shirey and Walker, 1998), unless the MORB component is old (≥ 2 Ga) or constitutes a major fraction of the mixture. Importantly, the high Os contents of the high-εNd subgroup (1–2 ppb; Table 1) require a predominantly peridotitic source and our Sr, Nd, and Pb isotopic modeling suggests that the recycled component must have been quite young (< 1 Ga; cf. section 5.2.4). In addition, studies of subducted portions of oceanic crust (i.e. eclogites and blueschists; Becker, 2000; Dale et al., 2007) have indicated Os to be relatively immobile and Re relatively mobile (similar to Rb, Ba, and K; Dale et al., 2007) during subduction-related dehydration of the basaltic oceanic crust. Therefore, subduction-related loss of Re would tend to decrease the radiogenic production of $^{187}$Os in a mixed mantle source. Our mixing models demonstrate that the Os isotopic composition is indeed highly dependent on the Os content of the peridotite component and that the effect of possible Re loss is negligible (Fig. 8). The Os isotopic composition of the high-εNd subgroup can be best explained by melting of a mixture of Os-rich DMM (~70–90%) and Re-poor MORB (~10–30%; Fig. 8, models A and B) that is compatible with the trace element and Sr, Nd, and Pb isotopic modeling (cf. Figs. 6 and 7).

![Fig. 8. $^{206}$Pb/$^{204}$Pb vs. $^{187}$Os/$^{188}$Os compositions of the uncontaminated (εNd = +9) Group 3 dikes and hypothetical mixtures between depleted MORB mantle peridotite (DMM) and 0.7 Ga sm-MORB 2 (cf. Figs. 6 and 7). A and B models with Os-rich peridotite and Re-poor MORB (A with 30% loss of Re) and C and D models with Os-poor peridotite and Re-rich MORB (C with 30% loss of Re). See Table S4 for more detailed parameters. Pb isotopic modeling performed as in Fig. 7 (Table S4). It is important to note that due to the low Os content of the recycled MORB, the models involving Os-rich peridotite (A and C) are not notably different whether Re has been subduction-modified or not.

5.2.4. Constraints on the age of the recycled mantle component

We emphasize that the recycling age (i.e. the age at which the recycled crust started to evolve as a closed system until melting at 180 Ma; cf. Stracke et al., 2003) of 0.7 Ga suggested by our isotopic models (Figs. 7 and 8) should not be considered definitive given all the possible uncertainties in model parameters. Nevertheless, the recycled component is not likely to be older than 1.0 Ga, because such a component would result in low εNd and would require higher input of seawater Sr to produce the high-εNd signature at a given Pb isotopic composition (Fig. 7). In addition, due to the greater divergence in isotopic composition with time, the amount of ≥1 Ga recycled component in the mixture would have to be significantly below 10% for it to result in high-εNd signature (Fig. 7). This would not likely be enough to produce the pyroxenite fingerprint observed in the major element geochemistry of the Group 3 dikes (Heinonen et al., 2013).
The aforementioned effects of recycling age are not significantly affected by reasonable modifications in the other model parameters and, therefore, we conclude that the Group 3 dikes sampled a recycled component that was less than 1 Ga old.

5.2.5. Nature of the peridotitic mantle component

We used DMM as the peridotitic source component in the trace element and isotopic modeling (Figs. 6–8), because its composition is relatively well constrained (Workman and Hart, 2005). This depleted peridotite component could also represent the mantle portion of the subducted oceanic lithosphere, which would be difficult to distinguish geochemically from DMM if mixed with recycled crustal sources. As an alternative, Jackson and Carlson (2011) recently suggested that CFBs could derive from non-chondritic primitive mantle sources that had remained untapped deep in the Earth’s mantle for over 4 Ga. Their primary arguments were related to the isotopic characteristics of primitive CFBs: many of them exhibit high ³He/⁴He, non-chondritic primitive mantle-like εNd (+5 to +9 at present), and Pb isotopic compositions that plot near the geochron in ²⁰⁷Pb/²⁰⁴Pb vs. ²⁰⁶Pb/²⁰⁶Pb space (Fig. 7). Jackson and Carlson (2011) further suggested that the PREMA component that seems to dominate in OIB sources (Zindler and Hart, 1986) and closely corresponds to the isotopic composition of the uncontaminated high-εNd subgroup (Fig. 7) could represent mixing of non-chondritic primitive mantle with recycled materials.

It is difficult to distinguish between DMM and non-chondritic primitive mantle peridotite source without He isotopic data (cf. Fig. 7). Using the most recently reported trace element compositions for the peridotite components, we modeled the partial melting of mixtures consisting of DMM (Workman and Hart, 2005) or non-chondritic primitive mantle (Jackson and Jellinek, 2013) and subduction-modified MORB (sm-MORB 2; cf. Fig. 6). The most notable differences between the modeled melts are the positive Pb anomaly and relative enrichment in the most incompatible elements in the case of the non-chondritic primitive mantle mixture (Fig. 9). Given that formation of mantle pyroxenite is a complex process involving several mineral-melt reactions (e.g., Bodinier et al., 2008; Mallik and Dasgupta, 2012), that Pb is the most mobile of the concerned elements during subduction (cf. Fig. 6), and that the overall shape of the trace element signature is dependent on melting conditions, it is difficult to uniquely identify the peridotite component entrained in Group 3 dikes on the basis of trace element compositions. If the peridotitic component is non-chondritic primitive mantle, it would have to be from the high-εNd compositional end of the spectrum proposed for this component (cf. Fig. 7b).
5.3. **Group 3 sources and CFB magmatism**

The compositions of the Ahlmannryggen Group 3 dikes provide evidence for the involvement of a recycled MORB component in Karoo magmatism. In addition, sediment-bearing recycled sources have been suggested for the Vestfjella enriched ferropicrite suite (Heinonen et al., 2010). Interestingly, isotopic models imply similar recycling ages in both cases (best-fit at 0.7 and 0.8 Ga, respectively; Fig. 7; Heinonen et al., 2010). At ~0.7–1.0 Ga, oceanic crust was being subducted along the margins of the supercontinent Rodinia (e.g., Murphy et al., 2004). Whether the subducted crust remained in the upper mantle or was recycled via lower mantle and entrained by a deep-seated mantle plume before being incorporated into the sources of the Karoo CFBs is unclear (cf. Heinonen et al., 2010). Although the suggested recycling ages are compatible with both options (cf. Sobolev et al., 2011a), we consider that whole-mantle circulation of the oceanic crust would probably have led to a series of metamorphic and melting events causing strong compositional modifications in the subducted material, whereas our models infer that the compositional effects of the recycling process were limited to subduction modification (Figs. 6–8). We therefore suggest that the purported recycled components rather resided in the upper mantle and were entrained into a rising mantle plume (cf. Richards et al., 1989; Riley et al., 2005; scenario 2 of Jourdan et al., 2007) or were heated within the ambient depleted upper mantle beneath the insulating Gondwana supercontinent (cf. Coltice et al., 2009; Hastie et al., 2014; Heinonen et al., 2010; scenario 1 of Jourdan et al., 2007).

The role of lithospheric sources in the generation of the Karoo CFBs is another outstanding question (cf. Heinonen et al., 2010; Jourdan et al., 2007; Luttinen et al., 2010). Importantly, the Sr-Nd isotopic compositions of the Karoo CFBs can be reproduced by lithospheric contamination of melts derived from DMM sources (cf. Fig. 2; Heinonen et al., 2010). In contrast, the Group 3 high-εNd dikes with high 87Sr/86Sr appear to be unsuitable to represent parental magmas for the majority of the Karoo CFBs (cf. Fig. 2), implying that their sources did not have a major role in Karoo magmatism. The anomalous geochemical signature of the high-εNd subgroup does not exclude the possibility that Karoo CFBs carry geochemical traces of some other recycled components, however. For example, less intensive subduction-modification or seawater alteration, or additional sedimentary component in the recycled crust would lead to less radiogenic εNd in the recycled component (cf. Fig. 7). Such a mild fingerprint, combined with the effect of additional contamination with the continental lithosphere, would be difficult to distinguish from the range of common CFB signatures (cf. low-εNd subgroup; Figs. 5 and 7). Accordingly, our results lend support to the possibility that the lithospheric geochemical signature typical of most CFBs could have been inherited at least partly from recycled lithospheric materials that melted in the deep mantle. Similar conclusions have recently been drawn from geochemical and isotopic data (Ewart et al., 2004; Luttinen et al., 2010; Rocha-Júnior et al., 2012), olivine and melt inclusion chemistry (Kent et al., 2002; Sobolev et al., 2007), and geophysical modeling (Cordery et al., 1997; Leitch and Davies, 2001) of CFBs. Our study substantiates the view that the potential of recycled sources in creating chemical heterogeneity of CFBs should be carefully assessed in every case.

6. **Conclusions**

Lead and Os isotopic data and previously published geochemical data for the Ahlmannryggen mafic and ultramafic dikes (Group 3) from the Antarctic extension of the Jurassic Karoo CFB province provide a geochemical window into the deep mantle beneath the Gondwana supercontinent. The radiogenic initial εNd of the Group 3 dikes (from +5 to +9 at 180 Ma) indicates that their source was in the sublithospheric mantle. Correlations of Sr, Nd, and Pb isotopes indicate that some of the Group 3 magmas experienced minor contamination with continental crust. The Group 3 dikes that show the most radiogenic εNd (+9) and derive from least contaminated magmas exhibit relatively radiogenic initial 206Pb/204Pb (18.2–18.4) and 87Sr/86Sr (0.7035–0.7037), indicating that they did not originate solely from ambient depleted upper mantle source (206Pb/204Pb = 18.0 and 87Sr/86Sr = 0.7026 at 180 Ma). Isotopic and trace element
modeling indicates that the source contained ~10–30% of seawater-altered and subduction-modified MORB that was affected by the loss of LILEs, U, Pb, LREE, Nd, and possibly Re, and had a recycling age of ≤ 1.0 Ga. This pyroxenitic component was entrained in an Os-rich peridotite matrix that either represents depleted mantle–like material (ambient upper mantle or recycled oceanic mantle lithosphere) or non-chondritic primitive mantle. Although the recycled MORB sources suggested for the Group 3 dikes were not likely the predominant source of Karoo magmatism, broadly similar but less subduction-modified or more sediment-influenced recycled components would not be readily recognized in evolved CFBs. Therefore, the role of recycled source components in influencing magma chemistry and petrogenesis in Karoo and other CFB provinces should be carefully assessed.

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Appendix A. Supplementary material
Supplementary material related to this article can be found on-line at http://dx.doi.org/10.1016/j.epsl.2014.03.012.

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