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1 Osmium isotope evidence for rapid melt migration towards the Moho in the

2 Oman ophiolite

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10 **6499 words**

Abstract

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The oceanic crust, covering two-thirds of the Earth's surface, is formed along mid-oceanic ridges by crystallization at shallow levels of melts formed at depth by partial melting of mantle peridotite. Yet, the process of melt transport to the ridge axis remains poorly understood. Ophiolites, which provide a window into the uppermost mantle, contain dunite bodies often interpreted as relics of melt flow conduits, formed by pyroxene dissolution during melt-peridotite interaction. Here, we present structural and geochemical data on peridotites from the southeastern Oman ophiolite where three types of dunite, corresponding to the Moho transition zone (MTZ), the main and basal mantle sections, are identified. We focus on osmium isotopes, which are particularly well-adapted to tracing melt flow through peridotites. Osmium isotope signatures from host harzburgites accord with abyssal peridotite values and do not vary systematically with setting. In contrast, dunite Os compositions depend on structural context. Basal dunites display compositions similar to harzburgite values, while MTZ dunites have highly radiogenic compositions similar to those of the overlying crust, requiring extensive interaction with melts more radiogenic than MORB. Modeling shows that melts percolating through and equilibrating with dunite channels would acquire unradiogenic compositions, inconsistent with the observed Os signatures of MTZ dunites and lower crust. Thus, our findings require melt transport without equilibration with dunite or harzburgite, arguing for rapid or at least chemically isolated melt migration from the mantle source to the Moho.

233 words

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31 Oman ophiolite, Mantle, Osmium isotopes, Melt migration, Dunite

Highlights

- 1. Distinction of three types of dunite in the mantle section of SE Oman ophiolite
- 2. Strong correlation between dunite Os isotope composition and structural context
- 35 3. MTZ dunites interacted extensively with melts more radiogenic than MORB
- 4. High ¹⁸⁷Os/¹⁸⁸Os at odds with melt transport by percolation in dunite channels
 - 5. Rapid/chemically isolated melt transport needed to maintain radiogenic signatures

1. Introduction

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Melt migration processes beneath mid-oceanic ridges remain controversial, engendering a diversity of models based on structural (Nicolas, 1986), geochemical (Kelemen et al., 1995), petrological (Morgan and Liang, 2003) and numerical studies (Ahern and Turcotte, 1979; Stevenson, 1989). One strong constraint is the existence of disequilibrium between shallow mantle peridotites and mid-ocean ridge basalts (MORBs) (O'Hara, 1965; Stolper, 1980), manifested notably by undersaturation of orthopyroxene (opx) in MORBs. Melt-peridotite interactions can result in pyroxene dissolution in the host peridotite (Kelemen, 1990; Lambart et al., 2009; Morgan and Liang, 2003), forming dunite if continued to completion. The origin of ophiolitic dunite bodies is commonly attributed to this process (Boudier and Nicolas, 1977, 1995; Büchl et al., 2002, 2004; O'Driscoll et al., 2012), which may generate dunite channels providing transport pathways for ascending melts (Kelemen et al., 1995). In such channels, melts are isolated from equilibration with the surrounding peridotite, but may equilibrate with the dunite. The Oman ophiolite is well-suited for the study of melt migration beneath mid-oceanic ridges. This large (>20 000 km²), well-exposed segment of oceanic lithosphere, formed ~95 Ma ago along a rapid spreading center (Boudier et al., 1997; Nicolas et al., 2000a; Rioux et al., 2012), provides a window into the uppermost mantle, which outcrops over >50% of its surface. The Re-Os radiometric system, in which ¹⁸⁷Re decays to ¹⁸⁷Os ($t_{1/2}$ = 41.6 Ga, λ = 1.666 × 10⁻¹¹) (Smoliar et al., 1996), is a particularly useful tool for studying mantle melt transport. The contrasting behavior of these two elements during partial melting (rhenium is moderately incompatible while osmium is compatible) creates Earth reservoirs with strikingly different Re/Os ratios, producing over time contrasting Os isotope ratios. Recycled oceanic crust and sediments develop highly radiogenic Os compositions while mantle peridotites remain unradiogenic. When melt migrates through dunite, the isotopic and elemental compositions of incompatible elements (e.g. light rare earth elements) are rapidly overprinted in the dunite by equilibration with the magma, which has much higher concentrations of these elements. For Os, the mass balance is reversed, allowing gradual changes in Os isotope composition to be recorded in the dunite as melt progressively passes through.

Here, we present structural observations and geochemical data on melt migration and dunite formation from peridotites from the southeastern Oman ophiolite. We focus on Os isotopes, tracers exceptionally well-adapted to studying melt-peridotite interactions. Implications of our results for melt transport mechanisms are modeled in light of the structural context of each sample.

2. The Oman ophiolite

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2.1. Internal structure

As documented by early studies (e.g. Boudier and Coleman (1981); Lippard et al. (1986); Nicolas et al. (2000a)), the Oman ophiolite consists of a >15 kilometer thick dominantly harzburgitic mantle section, overlain by a complete mafic crust. Our study focuses on the mantle section. This domain was first detailed by Boudier and Coleman (1981), based on a cross-section in the Wadi Tayin massif. The uppermost few hundred meters beneath the crust correspond to the Moho Transition Zone (MTZ), a primarily dunitic layer of variable thickness. The MTZ has been extensively studied (e. g. Boudier and Nicolas (1995); Godard et al. (2000); Koga et al. (2001); Rospabé et al. (2018)) and is interpreted as a melt storage and reaction zone, where ascending melts stagnate beneath the crust-mantle interface (Moho). These melts dissolve orthopyroxene of the host harzburgite, producing both pure dunite and "melt-impregnated" dunite with interstitial plagioclase, pyroxene and occasional amphibole, as well as rare cumulate gabbro lenses (<5%). In the lowermost MTZ, interfingering relationships between harzburgite and dunite with orthopyroxene relics suggest a melt-peridotite reactional origin for the dunite (Nicolas and Prinzhofer, 1983). Furthermore, the olivine Mg# [= Mg/(Mg+Fe_{total})_{at}] is similar in the MTZ and underlying harzburgite, whereas spinel TiO₂ content in the MTZ is higher, indicating dunite arose from equilibration with TiO2-rich primary melts rather than crystal fractionation (Koga et al., 2001). Beneath the MTZ, the primarily harzburgitic main mantle section (MMS) comprises most of the ophiolite volume. The exact distribution of dunite within the MMS is unclear and its estimated proportion varies between studies (e.g. <5% for Boudier and Coleman (1981) and 5-15% for Lippard et al. (1986)). Dunite occurs either as planar and irregular bodies with dimensions up to ~20 meters (Bartholomew, 1983), crosscutting the mantle fabric, or more rarely, as thin concordant bands. Crosscutting gabbro and pyroxenite dikes are occasionally observed.

At the bottom of the mantle section near the basal thrust, a highly deformed, basal banded unit (BBU) is often observed. In this section, Boudier and Coleman (1981) report subequal proportions of harzburgite (or occasionally lherzolite) and concordant dunite bands, varying in thickness from 0.1 to several meters. Lippard et al. (1986) also note significantly increased dunite abundance near the base. These basal dunites have never been thoroughly nor systematically studied.

Braun and Kelemen (2002) determined a power law relationship between dunite width and cumulative abundance, which they used to support a model of melt transport to the ridge in coalescing dunite channels. The mid to large-sized dunites in their study were measured in the Muscat massif (Fig. 1). The proximity with the basal thrust and the exceptionally high dunite abundance (18-46%) of this small massif suggest that it is derived from the BBU.

2.2. Geodynamic context

The geodynamic context of the Oman ophiolite has long been controversial. The ophiolite presents several characteristics akin to those of fast spreading ridges (e. g. magma chamber shape, mantle structure, crustal continuity, structure and thickness variation...) (Boudier et al., 1997; Nicolas et al., 2000a). Moreover, basalts above the sheeted dike complex (V1, Geotimes unit) display MORB-like compositions (Ernewein et al., 1988; Godard et al., 2006). However, upper lavas (V2, Lasail unit, found in northern Oman) overlying hydrothermal deposits display island arc affinities (Alabaster et al., 1982; Ernewein et al., 1988), suggesting ophiolite formation in a supra-subduction setting. As these upper units clearly postdate most of the crustal accretion, and probably the recorded mantle flow, this origin has been disputed. However, reappraisal of the Geotimes unit (MacLeod et al., 2013) and dating of the metamorphic sole (Guilmette et al., 2018) strongly argue for accretion of the entire ophiolite from a hydrated mantle above a subducting slab. Nevertheless, the critical parameters controlling magmatic accretion at the ridge are spreading rate and presence or absence of a magma chamber, regardless of

the broader tectonic context. Thus, the Oman ophiolite remains a well-suited laboratory for studying melt migration in the mantle beneath accretion centers, as long as we focus on features contemporaneous with the asthenospheric mantle flow. Systematic petrostructural mapping (Nicolas et al. (2000a) and references therein) revealed the deformation temperatures of the peridotite and their spatial variation. Coarse-grained granular and porphyroclastic textures with well-recovered sub-structures are ascribed to asthenospheric deformation at the spreading center. Finer-grained porphyroclastic textures with varying degrees of recrystallization into neoblasts, mostly restricted to the borders of the massifs, reflect overprinting low-temperature deformation, associated with ophiolite emplacement. Later faulting is limited. Thus, the ophiolite can be seen as a fossilized piece of oceanic lithosphere, dissected by erosion and slightly tilted following the dip of the Moho. In the southeastern part of the ophiolite, the focus of our study, a NW-SE paleo-spreading center was identified from dike orientations, which trend NW-SE in the center of the Sumail massif and perpendicularly (NE-SW) to the north and south of this central zone (Fig. 1) (Boudier et al., 1997). The position of this spreading center is inferred from the alignment of several harzburgitic mantle diapirs, defined by steeply plunging lineations with radial horizontal lineations at their outskirts (Jousselin et al., 1998). Solid-state flow lineations in the harzburgite and MTZ, and magmatic lineations in the overlying crust are remarkably parallel, indicating that the mantle lineations formed before the gabbro was frozen. These observations show that harzburgite diapirs, MTZ formation and ridge accretion were contemporaneous, with diapirs being likely feeding centers for the overlying ridge segments.

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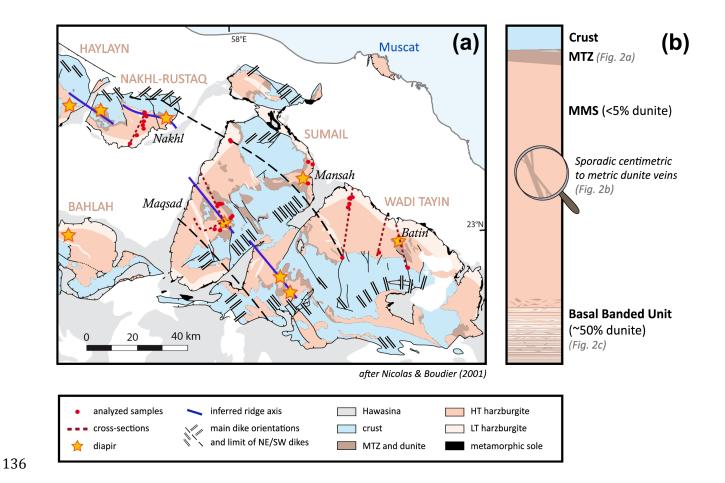


Fig. 1. Sample locations and structure of the southeastern part of the Oman ophiolite. (a) Simplified geological map of the southeastern massifs (Nakhl-Rustaq, Sumail, Wadi Tayin and Haylayn) of the Oman ophiolite (after Nicolas and Boudier (2001)) showing the sample locations (red dots) and the spatial distribution of the crust, Moho Transition Zone (MTZ), mantle and metamorphic sole sections. Most samples were collected along cross-sections (bold burgundy dotted lines) in on-axis and off-axis settings. Hawasina sediments, inferred ridge axis (bold blue lines), mantle diapirs (stars), main dike orientations (double black lines) and limits of NE-SW dikes (black dashed curves) are also shown. Batin dunite (in dark brown) is visible in the Batin area in the eastern part of the Wadi Tayin massif, below the diapir. (b) Schematic mantle cross-section based on field observations, showing the distribution of the three types of dunite described in the text: Mantle transition zone (MTZ), sporadic centimetric to metric veins in the main mantle section (MMS) and abundant bands in the basal banded unit (BBU). (For interpretation of the colors in the figure(s), the reader is referred to the web version of this article.)

2.3. Definition of three types of dunite in the mantle section

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To clarify dunite distribution, we made observations and collected samples along six cross-sections traversing the mantle section, in diapiric and non-diapiric areas in the Nakhl-Rustaq, Sumail (Maqsad diapir) and Wadi Tayin massifs (Fig. 1 and Fig. A.1). Complementary observations were made at sites near Muscat, at the base of the Haylayn massif and in the Batin area in the eastern part of Wadi Tayin. Our observations distinguish three types of dunite, each associated with a specific part of the mantle section (Fig. 1-2; dunite thicknesses in Table A.1): (1) Moho Transition Zone (MTZ) (Fig. 2a): As noted by previous studies (Boudier and Nicolas, 1995; Koga et al., 2001; Rospabé et al., 2018), the MTZ is composed of pure and plagioclase/clinopyroxeneimpregnated dunites, and reaches thicknesses of 50-500 meters above harzburgite diapirs, and usually <30 meters far from diapirs. Dunitic lenses several meters wide, interlayered with harzburgite, are sometimes found in the hundred meters underlying the MTZ. The MTZ volume is so large that it can be represented on maps including several massifs (scales of 1/1 000 000 to 1/500 000) (Fig. 1). (2) Harzburgitic main mantle section (MMS): Discordant gabbroic and pyroxenitic dikes rimmed by dunitic reaction zones crosscut sporadically the MMS harzburgite (Fig. 2b left), as described by Boudier and Coleman (1981). In some cases, all of the melt was extracted from the dike, leaving only the dunitic reaction zones, several centimeters to 5 meters wide, crosscutting the mantle fabric (henceforth such features referred to as "dunite veins") (Nicolas et al. (2000b) and references therein). Angular contacts, harzburgite blocks trapped between dunite veins, and stockwork-like geometry suggest a hydrofracturing origin for the magmatic dikes from which the dunitic reaction zones formed (Fig. 2b right). The crosscutting nature of these veins indicates they post-date asthenospheric flow, but several examples of partially tectonically transposed dunite veins suggest they also could form in the flowing mantle. Centimetric dunites parallel to the foliation are also observed in the MMS (henceforth "dunite bands") (insert in Fig. 2b), possibly representing extreme examples of transposed veins. All the MMS dunites, even those in harzburgite diapirs, represent a negligible, unmappable fraction of the mantle volume.

(3) Basal banded unit (BBU): This zone often hosts abundant, mainly concordant dunitic bands (Fig. 2c). These are usually centimeters to meters thick, rarely up to 20 meters, with the largest being mappable locally at the 1/10 000 scale. They are sometimes composed of opx-poor harzburgite instead of dunite. Coarse-grained samples sometimes occur between fine-grained ones (Table A.2) and at the top of the unit, suggesting that the transition from the low-temperature to the hightemperature domain is irregular, and that the BBU is thicker than the low-temperature domain. In the Wadi Tayin and Nakhl-Rustaq massifs, the top of this unit, where coarse-grained asthenospheric deformation is preserved, includes 3-20 meter "blobs" of dunite, often with several dunite veins branching into them, exactly as drawn by Bartholomew (1983). Such blobs are rare in the MMS (none in our cross-sections). The concentration of dunitic bands in the BBU is poorly understood. Lippard et al. (1986) suggested that ophiolite thrust-related extreme deformation of scattered dunite bodies in the mantle section could explain the basal dunite banding; however, this would not explain the higher dunite content in this section relative to the MMS, and the fact that parts of the banding, as well as the dunitic blobs, display asthenospheric deformation textures. An exceptional fourth type of dunite is exposed in the Batin area, consisting of a dunite body >9 kilometers long, 2 kilometers wide and several hundred meters thick (Fig. 1), evident on the USGS Wadi Tayin geological map and clearly visible in the field and on aerial photographs. This body was sampled by the ICDP Oman Drilling Project and several dedicated studies, including our own, are ongoing. Apart from the Batin area, no dunite larger than 30 meters, or similar to those at the base of the ophiolite, was found in the MMS. This is true even in the diapiric areas, thought to represent feeding zones for the overlying crust.

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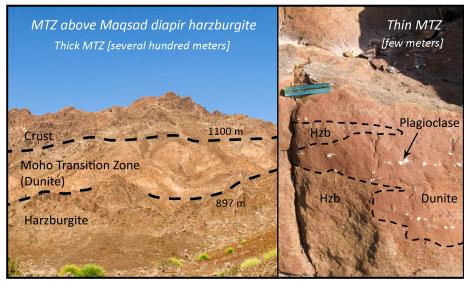
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(a) Moho Transition zone (MTZ) at the summit of the mantle section



(b) Sporadic dykes in the main mantle section



(c) Abundant concordant dunite bands at the base of the ophiolite

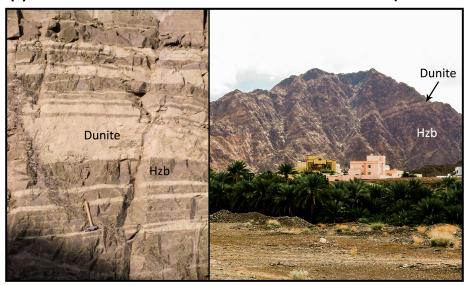


Fig. 2. Field photographs illustrating the three types of dunite observed in the mantle section of the SE Oman ophiolite. (a) Moho Transition Zone (MTZ): Left, landscape-view of the thick dunitic MTZ just beneath the crust in the Maqsad diapir region; Right, plagioclase-impregnated dunite of the thin MTZ from the Nakhl-Rustaq massif in contact with residual harzburgite. (b) Sporadic veins in the main mantle section (MMS) of the Nakhl-Rustaq massif: Left, thin pyroxenite dike with dunite reaction zone; insert, concordant dunite band. Right, cross-cutting plurimetric dunite veins (160D1) to compare with insert showing sketch of a breccia hydrofracture network from Jébrak (1997). (c) Abundant concordant dunitic bands at the base of the ophiolite: Left, succession of dunitic and harzburgitic bands in the basal section from the northern Sumail massif; Right, landscape-view of the basal banded unit in the Wadi Tayin massif. Hzb, harzburgite.

3. Samples and analytical techniques

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Samples were selected from the three types of dunite (19 from 81 collected dunites, chosen for freshness and variety) and host harzburgite (20 from 140 collected harzburgites) in the mantle sections of the Nakhl-Rustaq, Sumail, Wadi Tayin and Haylayn massifs (Fig. 1; UTM coordinates in Tables A.2-3). In six cases, dunite and harzburgite were collected in close proximity (<1 meter) (Table A.1). Sampling localities include both off-axis (e.g. Nakhl-Rustaq and Wadi Tayin massifs, Mansah diapir in Sumail massif) and on-axis (e.g. Magsad diapir in Sumail massif) settings. Dunite structural data and petrographic descriptions of all analyzed samples are given in Tables A.1-2 (complementary descriptions of non-analyzed samples in Table A.9). Harzburgites are moderately to highly serpentinized and have coarse-grained porphyroclastic textures, except for four fine-grained samples from the BBU. Dunites are moderately to highly serpentinized, with higher degrees of serpentinization in the MTZ and the BBU. Sulfides are very rare and small in both lithologies. Medium to high-temperature porphyroclastic and granular textures are observed among all three types of dunite. Grain size, spinel abundance, crystal shape and serpentinization degree do not vary systematically, making it difficult to distinguish different dunite types simply from examination of hand specimens or thin sections. Nevertheless, basal dunites (1) never contain plagioclase, and (2) often contain orthopyroxene relics. Our dunitic samples include seven pure dunites (98.0-99.5% and <0.5-2.0%_{spl}), eight containing pyroxene relics (<1.5%_{Px}), three impregnated with plagioclase and/or pyroxene, and one olivine troctolite. Whole-rock major and trace element compositions (Tables A.3-5) were determined respectively by the SARM at CRPG using an ICP-OES iCap6500 and at Géosciences Montpellier (AETE-ISO University of Montpellier, France) using an Agilent 7700X quadrupole ICP-MS. After chemical separation, Os isotopic analyses (Tables A.6-8) were done at CRPG by NTIMS using a Finnigan MAT262 mass spectrometer.

Details of analytical methods are provided in the Supplementary Material.

4. Variations in major element and Os isotope compositions with structural

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Harzburgites from both the main and basal sections have Mg# mostly from 0.900 to 0.916 and Os concentrations from 2.27 to 6.29 ppb (Fig. 3b, 4a; Table 1 and Tables A.3, 6-7). Basal harzburgites generally have lower Al₂O₃ (0.39-0.63 wt.%) than MMS harzburgites (0.48-0.96 wt.%), except for two samples (16OL27B and 16OL29), whose compositions (0.92 and 1.21 wt.%) approach those of clinopyroxene-bearing harzburgites from Godard et al. (2000) (Fig. 3a). Plagioclase-impregnated, orthopyroxene-poor harzburgite 16OD5C from the uppermost MMS displays the highest FeO (8.76 wt.%) and lowest Os content of all of the harzburgites (Fig. 3b, 4a). Values of ¹⁸⁷Os/¹⁸⁸Os_{95Ma} vary from 0.1131 to 0.1345, with most between 0.12 and 0.13, and no systematic differences are observed between MMS and basal harzburgites (Fig. 4a and Table 1). These values span nearly the entire global abyssal peridotite range (Becker and Dale, 2016), with the least radiogenic samples having Redepletion model ages (T_{RD}) ~2.2 Ga (using Primitive Upper Mantle (PUM) evolution curve of Meisel et al. (2001)). T_{RD} ages are obtained by comparing measured ¹⁸⁷Os/¹⁸⁸Os values directly with the mantle evolution curve, implicitly assuming Re concentration equals zero in residual peridotite after melt extraction. However, this assumption is often false, as moderate melt extraction leaves peridotites with non-zero Re contents. Therefore, T_{RD} ages are always minimum values that could dramatically underestimate true melt extraction ages. Nevertheless, the T_{RD} ages >0.5 Ga of most of the harzburgites imply that they experienced some melt extraction hundreds of million years before possible remelting beneath the Oman spreading center.

Dunites have highly variable compositions correlating with their structural contexts (Fig. 3, 4b-c):

(1) MTZ dunites have generally low Mg# (0.897-0.920; mean: 0.904) with higher Al_2O_3 (0.09-1.71 wt.%) and lower Os concentration (0.39 to 1.17 ppb) than most other dunites. Their $^{187}Os/^{188}Os_{95Ma}$ ratios (0.1415 to 0.1490) are higher than the estimated PUM value (~0.1289 at 95 Ma; Meisel et al. (2001)) and coherent with those of Oman MTZ chromitites (Ahmed et al., 2006). Plagioclase-impregnated dunite 100M25 and especially troctolite 160L34B are enriched in magmatophile elements (Fe, Al, Ca)

- (Fig. 3). The Os concentration and ¹⁸⁷Os/¹⁸⁸Os_{95Ma} of dunite 10OM25 do not differ from those of other 257 258 MTZ dunites, but troctolite 16OL34B is less radiogenic (0.1319) and has a higher Os concentration 259 (2.30 ppb) (Fig. 4b). 260 (2) Relative to MTZ dunites, most MMS dunites display a restricted range of Mg# (0.893-0.908; mean: 261 0.902), lower Al₂O₃ (mostly 0.11-0.51 wt.%), higher Os concentration (1.12 to 5.95 ppb) and lower 262 187 Os/ 188 Os $_{95Ma}$ (0.1249 to 0.1330). One unusual sample (16OD9A) located ~1 km below the MTZ has atypical Os characterístics (11.80 ppb Os; ¹⁸⁷Os/¹⁸⁸Os_{95Ma}= 0.1483) but an unexceptional major element 263 264 composition (Mg#= 0.905; Al₂O₃= 0.33 wt.%). In MMS dunite/harzburgite pairs, the dunite is always 265 much more radiogenic than the harzburgite (Fig. 4c). 266 (3) Basal dunites display higher overall Mg# (0.908-0.923; mean: 0.914) and lower Al₂O₃ (≤0.8 wt.%) 267 than the other dunites. Their Os concentrations vary substantially (0.77 to 7.17 ppb), while their ¹⁸⁷Os/¹⁸⁸Os_{95Ma} ratios (0.1165 to 0.1280) are similar to abyssal peridotite values. Unlike in the MMS, 268
- When placed in the structural framework defined here, previous Mg#, Al₂O₃, Os concentration and Os isotope results from harzburgites and dunites of the Oman ophiolite (Godard et al., 2000; Hanghøj et al., 2010; Senda et al., 2020; Xiong et al., 2020a,b) are consistent with these observations.

paired dunites and harzburgites collected in the BBU have very similar isotope ratios.

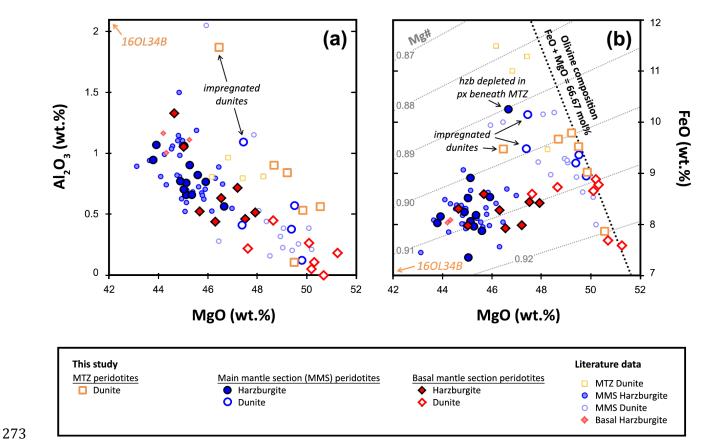


Fig. 3. Whole-rock major element compositions of harzburgites and the three types of dunite recalculated on a volatile-free basis in (a) Al_2O_3 vs. MgO and (b) FeO vs. MgO diagrams. In panel (b), dotted grey lines represent constant Mg# values and the bold dotted black line shows the evolution of olivine composition constrained by FeO + MgO = 66.67 mol% following Godard et al. (2000). In all panels, olivine troctolite (16OL34B) plots outside of the graph area, with higher Al_2O_3 and lower FeO and MgO concentrations. Published Oman peridotite data are shown for comparison: MTZ dunites (Godard et al., 2000); MMS dunites (Godard et al., 2000; Hanghøj et al., 2010); MMS harzburgites (Godard et al., 2000; Hanghøj et al., 2010) and basal harzburgites (Godard et al., 2000). Attribution of samples reported in the literature to the three domains defined in our study was based on their geographic positions in the mantle section.

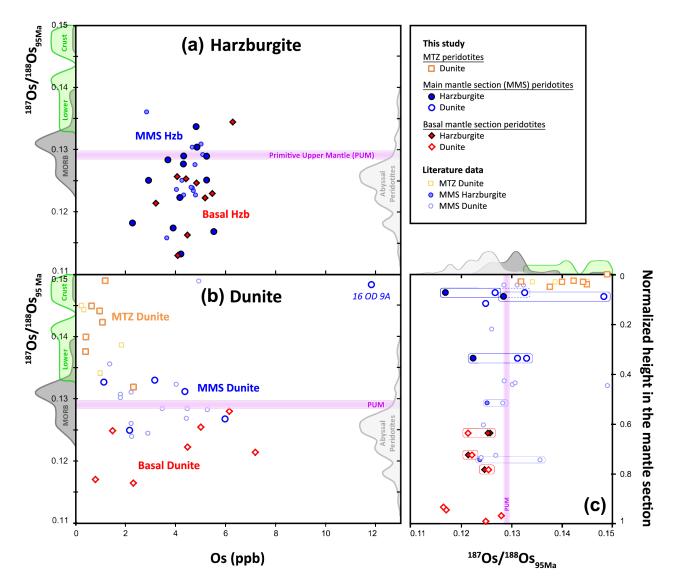


Fig. 4. Os isotope compositions of harzburgites and the three types of dunite. (a) ¹⁸⁷Os/¹⁸⁸Os_{95Ma} vs. [Os] in harzburgites; (b) ¹⁸⁷Os/¹⁸⁸Os_{95Ma} vs. [Os] in dunites; (c) Variation in dunite ¹⁸⁷Os/¹⁸⁸Os_{95Ma} with approximate height in the stratigraphic section, normalized to the total height (Moho to base) of each section to allow comparison of samples from different cross-sections. Harzburgites taken adjacent to six of the dunites are also shown, with blue and red contours delimiting harzburgite-dunite pairs in the MMS and the BBU, respectively. Dunite 16OD4C (dashed blue contour) is associated with the 16OD4D_{Dun/Hzb} pair but was collected at a greater distance from the harzburgite (30 cm). In all panels, light purple fields represent the estimated ¹⁸⁷Os/¹⁸⁸Os ratio for the Primitive Upper Mantle (PUM) (~0.1289 at 95 Ma) (Meisel et al., 2001). Light grey, dark grey and green curves on both sides of the diagrams show the probability density distributions of ¹⁸⁷Os/¹⁸⁸Os ratios for abyssal peridotites (compilation in Becker and Dale (2016)), MORB (MORB sulfide data compiled in Gannoun et al. (2016))

and Oman lower crust (Peucker-Ehrenbrink et al., 2012) respectively. Published Oman peridotite data are shown for comparison: MTZ dunites (Xiong et al., 2020a,b); MMS dunites (Hanghøj et al., 2010; Xiong et al., 2020a,b) and MMS harzburgites (Hanghøj et al., 2010) (symbols as in Fig. 3), with attribution to each mantle domain based on geographic positions in the mantle section.

Table 1. Mg#, Re and Os concentrations and Os isotope compositions of the studied Oman peridotites

Sample	Massif	Lithology	Hzb/Dun pairs	Normalized height in the section	Mg#	[Os] (ppb)	[Re] (ppb)	¹⁸⁷ Re/ ¹⁸⁸ Os	(¹⁸⁷ Os/ ¹⁸⁸ Os) _m	(¹⁸⁷ Os/ ¹⁸⁸ Os) _{95 Ma}	2σ error
10 OM 25	Sumail (Maqsad)	impregnated dunite		0.030	0.897	0.399	0.114	1.376	0.14214	0.13996	0.0005
10 OM 45C	Sumail (Maqsad)	dunite		0.030	0.900	0.947	0.050	0.255	0.14456	0.14415	0.0006
18 OK 26	Sumail (Maqsad)	dunite		0.040	0.920	0.628	0.409	3.149	0.14993	0.14494	0.0008
10 OM 5C*	Sumail (Mansah)	dunite		0.025	0.908	1.063			0.14235	0.14115	0.0005
10 OM 5C-d							0.168	0.760			
07 OD 1	Sumail (Mansah)	dunite		0.050	0.900	0.390	0.199	2.462	0.14151	0.13761	0.0005
16 OL 11B	Wadi Tayin	pure dunite		0.000	0.903	1.173	0.033	0.137	0.14922	0.14900	0.0008
16 OL 34B	Wadi Tayin	olivine troctolite		0.028	0.899	2.305	0.053	0.111	0.13209	0.13192	0.0020
Main mantle se	ction (MMS) peridotite	s									
16 OD 5C	Nakhl-Rustaq	depleted harzburgite		0.007	0.890	2.274	0.448	0.948	0.11976	0.11826	0.0019
16 OD 5D	Nakhl-Rustaq	harzburgite		0.007	0.916	4.211	0.152	0.174	0.11362	0.11334	0.0011
16 OD 7	Nakhl-Rustaq	harzburgite		0.053	0.909	2.921	0.025	0.042	0.12517	0.12510	0.0016
16 OD 4D _{Hzb}	Nakhl-Rustaq	harzburgite	16 OD 4D _{Dun}	0.073	0.900	5.519	0.358	0.312	0.11741	0.11691	0.000
16 OD 9C	Nakhl-Rustaq	harzburgite	16 OD 9A	0.089	0.908	3.691	0.130	0.170	0.12871	0.12844	0.001
16 OD 2	Nakhl-Rustaq	harzburgite		0.241	0.907	4.314	0.500	0.559	0.12990	0.12902	0.001
L6 OD 1A	Nakhl-Rustaq	harzburgite	16 OD 1B	0.337	0.906	4.164	0.599	0.693	0.12350	0.12241	0.000
L6 OD 10	Nakhl-Rustaq	harzburgite		0.419	0.906	4.855	0.381	0.378	0.13112	0.13052	0.001
l1 OD 15A	Sumail	harzburgite		0.280	0.912	3.894	0.152	0.187	0.11782	0.11753	0.000
07 OD 24	Sumail	harzburgite		0.254	0.907	4.826	0.460	0.459	0.13451	0.13378	0.000
07 OD 27B	Sumail	harzburgite		0.344	0.904	4.314	0.221	0.247	0.12817	0.12778	0.000
07 OD 8C	Sumail	harzburgite		0.150	0.910	5.246	0.379	0.348	0.12961	0.12906	0.000
07 OD 8C-d					_	5.228	0.321	0.295	0.12570	0.12523	0.000
16 OD 4C	Nakhl-Rustaq	impregnated dunite (vein)	(16 OD 4D _{Hzb})	0.073	0.893	1.120	0.305	1.312	0.13474	0.13267	0.001
16 OD 4D _{Dun}	Nakhl-Rustaq	impregnated dunite (vein)	16 OD 4D _{Hzb}	0.073	0.899	5.954	0.414	0.335	0.12729	0.12676	0.000
16 OD 9A	Nakhl-Rustaq	pure dunite (band)	16 OD 9C	0.089	0.905	11.802	0.431	0.177	0.14862	0.14834	0.000
16 OD 1B	Nakhl-Rustaq	dunite (vein)	16 OD 1A	0.337	0.904	3.145	0.087	0.134	0.13324	0.13303	0.0006
16 OD 1B-d						4.353	0.094	0.104	0.13135	0.13119	0.000
16 OD 31	Wadi Tayin	dunite (band)		0.116	0.908	2.147	0.910	2.043	0.12814	0.12491	0.002
Basal peridotite	25										
16 OD 13C	Nakhl-Rustaq	harzburgite	16 OD 13B	0.637	0.909	4.415	0.415	0.453	0.12607	0.12536	0.000
16 OD 13C-d						4.076	0.379	0.448	0.12647	0.12577	0.000
16 OL 25	Wadi Tayin	harzburgite		0.724	0.905	6.289	0.081	0.062	0.13464	0.13454	0.000
16 OL 26B	Wadi Tayin	harzburgite	16 OL 26A	0.724	0.910	3.214	0.022	0.032	0.12151	0.12146	0.001
16 OL 27B	Wadi Tayin	harzburgite	16 OL 27A	0.784	0.910	4.843	0.314	0.312	0.12522	0.12472	0.001
16 OL 27C	Wadi Tayin	harzburgite		0.784	0.913	4.097	0.046	0.053	0.11314	0.11306	0.001
16 OL 28B	Wadi Tayin	harzburgite		0.860	0.909	4.481	0.168	0.180	0.11659	0.11630	0.001
16 OL 29	Wadi Tayin	harzburgite		0.872	0.906	5.187	0.169	0.157	0.12256	0.12231	0.001
16 OL 24	Wadi Tayin	harzburgite		0.878	0.913	5.475	0.221	0.194	0.12335	0.12305	0.000
16 OD 13B	Nakhl-Rustaq	dunite (band)	16 OD 13C	0.637	0.923	7.169	0.158	0.106	0.12155	0.12139	0.000
16 OD 15E	Nakhl-Rustaq	dunite (band)		0.934	0.911	2.291	0.106	0.223	0.11688	0.11653	0.000
16 OD 18B	Sumail	dunite (band)		0.969	0.908	6.134	0.018	0.014	0.12800	0.12798	0.000
l6 OD 19A	Sumail	pure dunite (band)		0.946	0.910	0.774	0.066	0.413	0.11773	0.11707	0.001
16 OL 26A	Wadi Tayin	dunite (band)	16 OL 26B	0.724	0.909	4.460	0.014	0.015	0.12227	0.12225	0.001
16 OL 27A	Wadi Tayin	dunite (band)	16 OL 27B	0.784	0.912	4.998	0.079	0.077	0.12560	0.12548	0.001
L4 11E	Haylayn	pure dunite (band)		0.990	0.922	1.474	0.157	0.512	0.12573	0.12492	0.000

Os isotopic compositions in whole rock were analyzed at the CRPG laboratory in Nancy (France) using N-TIMS (Finnigan MAT 262) and MC-ICP-MS (Neptune +). All uncertainties are 2 \u03c3 and include all recognized sources of uncertainty, including in-run analytical precision, reproducibility of standard analyses, uncertainties related to the variability of the blank and its isotope composition, uncertainties related to the age correction (95 Ma). Samples belonging to harzburgite/dunite pairs collected in close proximity (<1 meter) are indicated. For each sample, an approximate estimation of its height in the mantle section was obtained by normalizing to the total section height to compare samples from different cross-sections. Duplicate analyses were performed for several samples (noted "-d") and UB-N standard. More details are presented in the Supplementary Material. * For 10 OM 5C sample, (¹⁸⁷Os,¹⁸⁸Os)_{95 Ma} was calculated using Re compositions of its duplicate.

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Mg# = Mg/(Mg+Fe)at; Hzb = harzburgite; Dun = dunite.

5. Discussion

- Our coupled geochemical and structural observations underscore three findings:
- 304 (1) Throughout the mantle section, harzburgite Mg#, Al₂O₃ contents, Os concentrations and isotopic
- compositions span a range comparable to that of abyssal peridotites (Becker and Dale, 2016) (Fig. 3,
- 306 4a). Dunites display more variable compositions linked to their structural contexts, with ¹⁸⁷Os/¹⁸⁸Os_{95Ma}
- 307 and Al_2O_3 content increasing and Mg# generally decreasing from the BBU through the MMS to the
- 308 MTZ (Fig. 3, 4b-c).

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- 309 (2) In the BBU, dunite Os isotope compositions closely resemble those of adjacent harzburgites (Fig.
- 310 4c). In contrast, dunites in the overlying MMS have more radiogenic compositions than associated
- 311 harzburgites.

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- 312 (3) In the MTZ, dunite 187 Os/ 188 Os_{95Ma} ratios are much more radiogenic than those of abyssal
- peridotites or even MORB (Gannoun et al., 2016), and dunite Os concentrations are low (Fig. 4b-c).

5.1. Evidence against seawater contamination

- 315 We first consider whether the radiogenic compositions of the MTZ dunites result from hydrothermal
- 316 circulation of seawater, which has high ¹⁸⁷Os/¹⁸⁸Os (~0.51–0.57 in the mid-Cretaceous; Ravizza et al.
- 317 (2001)) and could modify ¹⁸⁷Os/¹⁸⁸Os in oceanic peridotites (discussion in Becker and Dale (2016)).
- 318 Given the very low Os concentration of seawater (~10 fg/g; Levasseur et al. (1998)), water/rock ratios
- 319 required to significantly change Os compositions of peridotites are extremely high (>1000). In
- ophiolitic peridotites, water/rock ratios are almost always <100 (McCulloch et al., 1981), so significant
- 321 modification of 187 Os/ 188 Os should be rare.
- Our samples have water contents (LOI < 15.42%, most <12%) typical of serpentinized peridotites. No
- 323 correlation is observed between LOI values, highly mobile element (Rb, Cs, Ba and U) abundances or
- 324 Th/U ratios (Fig. A.2 and Tables A.3-4) and Os isotope composition. As shown in Fig. A.2b, interaction
- of typical peridotite (187 Os/ 188 Os= 0.124; [Os] from 0.4 to 6 ppb) with seawater cannot produce the
- 326 radiogenic Os compositions found in MTZ dunites without decreasing Th/U ratios far below observed
- 327 levels, implying that the Os isotope signatures of our samples reflect essentially magmatic processes.

5.2. Os isotope evolution during melt-rock interaction

5.2.1. First order considerations

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values of MMS (\sim 0.130) or MTZ (\sim 0.140) dunites.

Dunite bodies are generally viewed as reaction products formed as melts migrate through host harzburgite, dissolving pyroxene and precipitating olivine. Focused melting experiments show that melt fractions of 0.25-0.30 are required to dissolve all pyroxenes at 0.5 GPa (Lambart et al., 2009). However, once all pyroxenes have been removed, the amount of additional melt migrating through a dunite body is difficult to quantify. Signatures of incompatible element tracers, which are dramatically less abundant in dunite than in melt, will be rapidly overprinted by equilibration with the liquid during melt percolation. In contrast, peridotites, including dunites, have higher Os concentrations than mantle melts. Thus, as melts equilibrate with peridotite during melt percolation, dunite Os concentration and $^{187}\mathrm{Os}/^{188}\mathrm{Os}$ should change slowly and progressively, allowing this element to serve as a tracer of the extent of melt-peridotite interaction. To roughly estimate the proportion of melt needed to obtain the initial isotopic compositions of the three types of dunite, we first consider a two-component mixture between peridotite (187Os/188Os= 0.124, typical of Oman harzburgites) and melt (0.2 ppb Os; 187 Os/ 188 Os= 0.150). The melt 187 Os/ 188 Os was chosen to be slightly more radiogenic than the MTZ dunites. This value is higher than those of all MORB and most arc lavas (except for extrusives with extremely low Os contents, whose isotope compositions are easily modified by assimilation of radiogenic crustal material; see data compilation in Bezard et al. (2021)). Results are shown for two peridotite Os concentrations (Fig. 5; details in Supplementary Material), representing typical harzburgite (4 ppb) and MTZ dunite (0.5 ppb). The peridotite ¹⁸⁷Os/¹⁸⁸Os ratio rises slowly with increasing melt fraction and is nearly unchanged at the melt fraction (0.25-0.30) needed to fully transform harzburgite to dunite. Much higher melt fractions, 0.75-0.96 and ≥0.94 respectively using the assumed parameters, are needed to attain ¹⁸⁷Os/ ¹⁸⁸Os

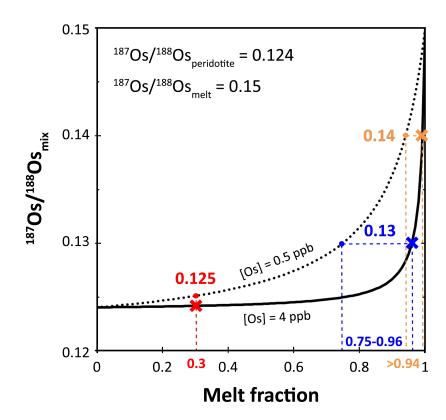


Fig. 5. Two-component mixture between peridotite ($^{187}Os/^{188}Os = 0.124$) and melt ($^{187}Os/^{188}Os = 0.150$); [Os]= 0.2 ppb) (cf. Supplementary Material). The two starting Os concentrations for the peridotite are used to simulate a mixture of a typical harzburgite (4 ppb; continuous curve) and of a MTZ dunite (0.5 ppb; dotted curve) with the melt. Basal dunites (red) have Os isotope compositions similar to those of harzburgites, suggesting low melt fractions. Higher melt fractions, 0.75-0.96 and ≥0.94 respectively, increase $^{187}Os/^{188}Os$ to 0.130 (MMS dunite; blue) or 0.140 (MTZ dunite; orange).

5.2.2. Melt-rock interaction during melt percolation

5.2.2.1. Single-cell percolation model

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While two-component mixing models allow qualitative assessment of the melt fractions needed to produce the radiogenic compositions of MMS and MTZ dunites, melt percolation models taking the Os partition coefficients (k_d) of the pertinent phases into account are more realistic. One critical point is the compatible behavior of Os in common mantle rocks partly due to its extremely chalcophile nature (k_{d in sulfide} ~10⁶) (Mungall and Brenan, 2014). During percolation of sulfide-undersaturated melts in the uppermost mantle, the sulfide fraction in the peridotite decreases progressively. As long as the peridotite contains sulfide, Os is trapped in the residue. When the peridotite loses its final trace of sulfide, the Os partition coefficient of the peridotite is likely to be lowered significantly causing Os concentration to decrease with continuing melt percolation. Concomitantly, if the percolating melts have a more radiogenic composition, the ¹⁸⁷Os/¹⁸⁸Os of the host peridotite increases gradually. In a simple model, a representative elementary volume (cell) of harzburgite is flushed repeatedly by increments of 2% melt of a constant radiogenic composition (details in Supplementary Material). As in most more complex models (e.g. Vernières et al. (1997)), chemical and isotopic equilibrium with the peridotite is assumed to be attained after each melt increment is added. After equilibration, the melt moves out of the cell and is replaced by fresh melt of the original composition. We suppose that the initial Os compositions of melt and harzburgite are the same as those used in the two-component mixture, and that transformation of harzburgite to dunite occurs after interaction with 25-30% of melt, corresponding to 10-15 melt increments. For simplicity, we also consider that complete sulfide removal is coincident with full transformation of harzburgite into dunite, i.e. when opx depletion is complete. If sulfide exhaustion occurs slightly before or after transformation to dunite, the results will not change substantively. In our model, the harzburgite contains 100 ppm sulfide, equivalent to ~40 ppm S, assumed to decrease to zero over 10 increments of 2% melt. After each increment of new melt equilibrates with the peridotite, the Os isotope composition is controlled by mass balance, while the new Os concentrations of the melt and the residual peridotite are controlled by the Os partition coefficients of the phases present. Osmium is strongly retained by the residue as long as sulfides remain. As sulfur is removed into the silicate melt, the IPGE (Iridiumgroup PGE: Os, Ir, Ru, Rh) are increasingly concentrated in the refractory sulfide residues, leading eventually to exsolution of platinum group minerals (PGM) such as Ir-Os alloys (Fonseca et al., 2012). Ir-Os alloys, laurites (RuS₂) and other PGM are probably the main hosts of Os in dunites and other sulfide-poor mantle peridotites (reviewed in Lorand and Luguet (2016) and O'Driscoll and González-Jiménez (2016)). In addition to PGM, experimental data (summarized in Brenan et al. (2016)) indicate that Ir and Ru are compatible in Cr-rich spinel, which constitutes ~1-3% of the dunite (Table A.2), and to a lesser extent in olivine, suggesting that Os could also be hosted by these phases. Thus, Os may behave as a moderately compatible element in dunite even after all sulfides have been removed. Nevertheless, dunites that have been fluxed by large quantities of magma have lower Os concentrations than most mantle harzburgites (Büchl et al., 2002), suggesting unsurprisingly that the compatibility of Os in peridotite is lowered in the absence of sulfide. Taking these observations into consideration, we consider a bulk partition coefficient (Dos) of 2 for dunites, which are assumed to have lost all of their original sulfide content. The effect of varying this value is explored below. Further constraint of Dos will require a better understanding of the behavior and stability of Os-bearing PGM during melt fluxing.

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Results of the model are compared with the data in Fig. 6. The model is obviously simplistic. The harzburgitic protolith is treated as a single composition rather than the wide range of harzburgite compositions observed in the ophiolite, likely explaining much of the mismatch with the data. Similarly, incoming melt composition and Dos, treated as constants, most probably vary with time. Nevertheless, this model provides a conceptual framework for examining the extent of melt-rock reaction experienced by each type of dunite. While the geometries and contexts of the three types of dunite are quite different, the melt-rock reaction process can be modeled in each case by melt fluxing through a volume of rock, using the parameters given in the caption to Fig. 6.

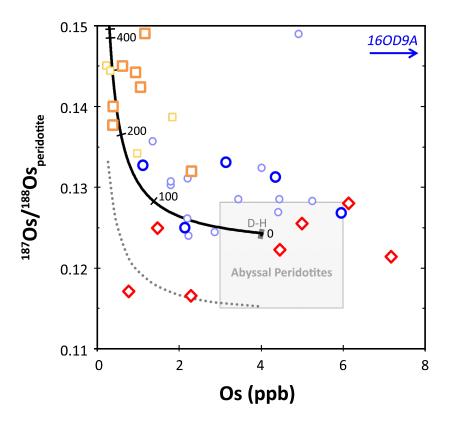


Fig. 6. Effect of percolation of radiogenic melt (¹⁸⁷Os/¹⁸⁸Os= 0.150; [Os]= 0.2 ppb) through a representative elementary volume (i.e., a single cell) of typical harzburgite (¹⁸⁷Os/¹⁸⁸Os= 0.124; [Os]= 4 ppb). Black curve shows evolution of peridotite ¹⁸⁷Os/¹⁸⁸Os vs. [Os] during this process. As repeated increments of melt (taken here as 2%) traverse the harzburgite, the pyroxenes will dissolve progressively, resulting in dunite formation after ~10-15 increments have passed (zone indicated as D-H on diagram). Concomitantly, any sulfides present in the harzburgite will also be dissolved in the

percolating melt, with complete sulfide dissolution assumed to approximately coincide with the transition from harzburgite to dunite. As Os will be retained as long as sulfide is present, Os concentrations will not start to decrease before this transition is nearly complete. In the example shown, the starting harzburgite is assumed to contain 100 ppm sulfide (~40 ppm S), $k_{d,0s}$ = 10^6 for sulfide and D_{Os} = 2 for dunite (see discussion of parameter choices in text and Supplementary Material). After transition to dunite, continued fluxing with repeated increments of melt (tick marks placed at 100, 200, 300 and 400 increments on the curve) will cause the Os isotopic ratio and concentration to move from the starting position in the abyssal peridotite field along the black curve towards the higher $^{187}\text{Os}/^{188}\text{Os}_{95\text{ Ma}}$ ratios and lower Os concentrations found in the MTZ dunites. Dotted grey curve represents percolation of a different melt ([Os]= 0.2 ppb; MORB-like $^{187}\text{Os}/^{188}\text{Os}$ = 0.133 - Gannoun et al. (2016)), interacting with harzburgite with $^{187}\text{Os}/^{188}\text{Os}$ towards the bottom of the abyssal peridotite range (Becker and Dale, 2016), possibly explaining low [Os] and unradiogenic compositions of two of the basal dunites. Os concentrations and initial Os isotope ratios for the three types of dunite are shown for comparison with the results of the melt percolation model (symbols as in Fig. 3).

In the MTZ, about 300 increments of melt fluxing, equivalent to a melt/rock ratio $^{\circ}$ 6, would be needed to lower the Os contents and raise the 187 Os/ 188 Os ratios of the initial harzburgite to the values ($^{\circ}$ 0.2-1.0 ppb and 187 Os/ 188 Os_{95Ma} $^{\circ}$ 0.138-0.149) observed in MTZ dunites. Such high melt/rock ratios are consistent with the intense melt-peridotite interactions thought to occur in this zone (Boudier and Nicolas, 1995; Rospabé et al., 2018).

pair studied, the dunites are more radiogenic than the associated harzburgite. For the same melt composition, the degree of melt-rock interaction required to obtain MMS-like Os isotopic signatures is smaller than for MTZ dunites (less than 150 increments of melt, corresponding to melt/rock ratios <3; Fig. 6). This is consistent with dunite formation by wall-rock reaction in melt dikes traversing the MMS (Fig. 2b, 7a), a process reported in other peridotitic bodies (Becker et al., 2001; Büchl et al., 2002). Wall-rock reaction would not have significantly modified the composition of melt flowing rapidly

through the center of the dike, because of the limited contact with the peridotite relative to the large volume of melt. In one case, two dunites were collected in the reaction zone of the same dike. Dunite 16OD4C, sampled ~30 cm from the dike edge and impregnated with plagioclase and clinopyroxene, has a moderately radiogenic $^{187}Os/^{188}Os_{95~Ma}$ ratio (0.1327). Clinopyroxene-impregnated dunite 16OD4D_{Dun}, from the border with the host harzburgite (16OD4D_{Hzb}), is less radiogenic (0.1268), suggesting lesser degrees of melt-rock interaction than near the dike center (Fig. 7a). This dunite is nevertheless more radiogenic than the harzburgite (0.1169). The unusual MMS dunite band 160D9A, with very high Os concentration (11.80 ppb) and very radiogenic ¹⁸⁷Os/¹⁸⁸Os_{95Ma} (0.1483) cannot be explained by this model and instead requires a specific mechanism of PGM or radiogenic sulfide precipitation. The Os isotopic composition of this sample is similar to the melt end-member used in the model, suggesting that such phases could have precipitated from magmas similar to those that interacted with harzburgite to form the MTZ and MMS dunites. Individual basal dunites show Os isotope compositions in the abyssal peridotite range, indicating that each experienced much lower extents of melt-rock interaction than MTZ or MMS dunites (<<100 melt increments; Fig. 6), though, given the high proportion of dunite, the total melt flow through the BBU may have been quite high. Unlike the MMS dunites, BBU dunites have ¹⁸⁷Os/¹⁸⁸Os_{95Ma} ratios very similar to those of adjacent harzburgites (Fig. 4c). They also have higher Mg#s and lower Al₂O₃ contents than MMS and MTZ dunites (Fig. 3), arguing for lesser degrees of interaction with fertile melt. Furthermore, some sections include opx-bearing dunite and/or opx-poor harzburgite bands, likely formed by small degrees of melt-rock interaction insufficient to fully dissolve orthopyroxene. Two dunites have low Os concentrations coupled with very unradiogenic compositions, implying formation from harzburgites with very low ¹⁸⁷Os/¹⁸⁸Os (though within the abyssal peridotite range) and perhaps interaction with less radiogenic melts than those required for MTZ dunites.

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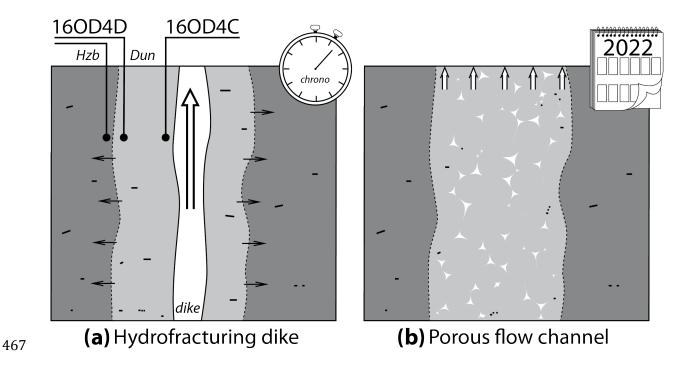
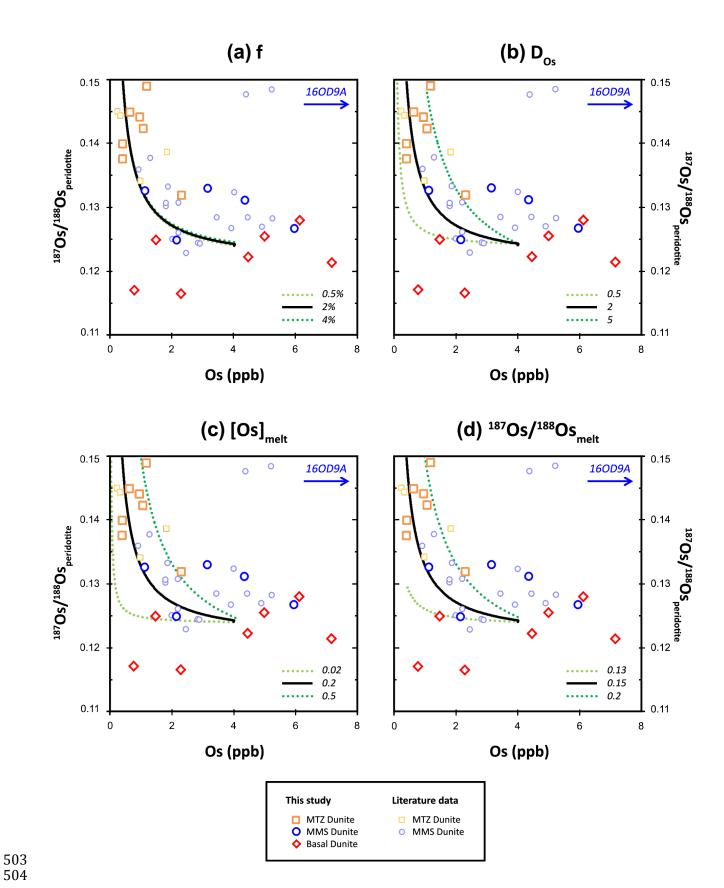


Fig. 7. Schematic diagrams of two end-member melt transport mechanisms: hydrofracturing (a) vs. porous flow (b). White: melt; light grey: dunite; dark grey: harzburgite; small black lines: spinel grains marking the foliation. In panel (a), a hydrofracturing dike contains 100% melt which travels fast (large arrow) and has a short life-span (weeks or months). Dikes produce a dunite reaction zone at their margins, that grows with time, and where the trace of foliation is preserved. In this case, samples at the center of the reaction zone (ie, closer to the dike) may have reacted with melt for a longer time and be more radiogenic than samples near the outside border of the reaction zone. This fits observations for MMS dunite samples 160D4D_{Dun} and 160D4C, both associated with host harzburgite 16D4D_{Hzb}.In panel (b), a porous flow channel is represented with a homogeneous distribution of 1-3% melt, which travels slowly through the solid matrix along grain boundaries (short arrows) and has a long life-span (possibly steady-state). In this case, the dunite may fully equilibrate with the melt.

5.2.2.3. Insensitivity to parameter choices

The pertinence of the model parameter choices was evaluated by changing each parameter individually while holding the others constant (Fig. 8). The instantaneous melt fraction modifies the number of melt increments needed to obtain a given ¹⁸⁷Os/¹⁸⁸Os ratio, but has little effect on the corresponding melt/rock ratio (Fig. A.3) or on the relationship between ¹⁸⁷Os/¹⁸⁸Os and [Os] (Fig. 8a). The Os bulk-rock partition coefficient after sulfide exhaustion affects the rapidity with which Os in the dunite is lost to the passing melts, and accordingly the rate at which ¹⁸⁷Os/¹⁸⁸Os is modified by repeated introduction of new melt (Fig. 8b and Fig. A.4). As discussed above, this parameter depends on the behavior and stability of rare PGM, and though modeled as a constant, probably changes during melt fluxing. To match both the Os isotope compositions and concentrations of the MTZ dunites, Dos values between about 0.5 and 5 are required (Fig. 8b). For input melt of a fixed Os isotope composition, the Os concentration will control the number of iterations required to obtain a given ¹⁸⁷Os/¹⁸⁸Os ratio. The data field can be matched by melt Os concentrations ranging from about 0.02 to 0.5 ppb (Fig. 8c and Fig. A.5). To produce MTZ dunites, the starting melt must have a ¹⁸⁷Os/¹⁸⁸Os ratio of at least 0.15 (Fig. 8d and Fig. A.6), but melts with much higher ratios do not fit the data. The reasonableness of all of the successful parameters supports the plausibility of the model.

Fig. 8. Effect of varying (a) instantaneous melt fraction (f) (0.5%, 2% and 4%), (b) Os bulk partition coefficient for dunite (D_{OS}) (0.5, 2, 5), (c) melt Os concentration (0.02, 0.2 and 0.5) and (d) melt $^{187}Os/^{188}Os$ (0.13, 0.15 and 0.2) on [Os] and $^{187}Os/^{188}Os$ of the peridotite in the single-cell melt percolation model shown in $^{187}Os/^{188}Os$ vs. Os diagrams similar to Fig. 6. Data from our study for the three types of dunite, as well as published Oman dunite data are shown for comparison with initial Os isotope ratios (MTZ dunites - Xiong et al. (2020a,b); MMS dunites - Hanghøj et al. (2010) and Xiong et al. (2020a,b); attribution of literature data to dunite types based on geographic positions) (symbols as in Fig. 3). The chosen parameters in the model (black continuous curves) best fit the data.



5.3. Need for rapid transport of radiogenic melt through the mantle section

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A melt more radiogenic than MORB is needed to explain the radiogenic character of many MMS and nearly all MTZ dunites. The high ¹⁸⁷Os/¹⁸⁸Os_{95Ma} ratios of the MTZ dunites are in good agreement with the ¹⁸⁷Os/¹⁸⁸Os_{95Ma} (~0.142) of the Oman lower crust (Peucker-Ehrenbrink et al., 2012). Two sources of radiogenic melt are possible: ancient recycled oceanic crust (cf. pyroxenites; Sergeev et al. (2014)) or a downgoing slab (Becker et al., 2004). We cannot exclude either possibility, but the first case would require an unusual concentration of ancient pyroxenites in the mantle source, probably unrepresentative of heterogeneity in the upper mantle, as radiogenic values similar to those of MTZ dunites are rare in MORBs (Fig. 4). A slab-related source (effective during the main accretion phase of the Oman oceanic crust) seems more consistent with a suprasubduction setting for the Oman ophiolite, as proposed by Guilmette et al. (2018) and MacLeod et al. (2013). Regardless of its origin, the radiogenic nature of the melt is fortunate, as this allows us to trace the extent of melt-rock reaction in the different types of dunite. A mechanism is needed to transport the radiogenic melt to the upper MMS and the MTZ. In one commonly accepted model (Kelemen et al., 1995), the melt is transported by porous flow in dunite channels. Once the olivine-rich channels are established, the melt is protected from equilibration with the surrounding peridotite, but equilibrates with the dunite through which it travels (Fig. 7b). To consider whether this mechanism is consistent with our Os results, we model a dunite channel as a stack of reaction cells, each similar to that described above (Fig. 9a and details in Supplementary material). Each increment of fresh radiogenic melt enters the bottom cell and equilibrates with the peridotite, and then migrates upwards into the overlying cell and is replaced by a new melt increment with the original composition. In this idealized model, the melt leaving this bottom cell will have a much less radiogenic Os composition than the entering melt, because of the strong buffering effect of the Os-rich peridotite (Fig. 9b). Similarly, the ¹⁸⁷Os/¹⁸⁸Os ratios of the overlying cells will be successively less affected by melt interaction, with the ratio of the uppermost cell being hardly modified at all (Fig. 9c). Therefore, melt leaving the dunite channel will have a ¹⁸⁷Os/¹⁸⁸Os ratio close to that of the original harzburgite from which the dunite formed (Fig. 9c). This implies that melts that have traveled through such channels, equilibrating with the dunite along the way, cannot retain the highly radiogenic compositions needed to explain the high ¹⁸⁷Os/¹⁸⁸Os_{95Ma} ratios of the MTZ dunites and the overlying lower crust. Instead, a mechanism without (or with little) equilibration with dunite or harzburgite is needed to bring melts with a radiogenic Os signature from depth towards the Moho. This lack of equilibration suggests rapid transfer of large melt volumes through the upper mantle section, also proposed by Kogiso et al. (2004) to explain the existence of linear correlations between Os and Sr, Nd and Pb isotopes in some ocean island basalts. The quantitative constraint that preservation of radiogenic Os compositions places on melt migration rates will depend on the geometry of melt pathways and on Os diffusion coefficients. While there are few experimental constraints on the latter, on theoretical grounds, Kogiso et al. (2004) estimated Os diffusion coefficients in olivine to be similar to those of Sr and Pb. In addition to the melt brought to the MTZ through a rapid process, some melt may have percolated through the harzburgite in mantle diapirs (Godard et al., 2000; Lorand et al., 2009). Coexistence of two transport processes could be consistent with modeling of melt migration beneath mid-ocean ridges based on U-series isotope systematics (Elkins et al., 2019). However, any melt percolating slowly through harzburgite will acquire the non-radiogenic isotope composition of the peridotite, of which no trace is found in the MTZ dunites. Such a scenario would require the melts traveling rapidly to have had an even more radiogenic composition before mixing in the MTZ with melts that may have traveled slowly.

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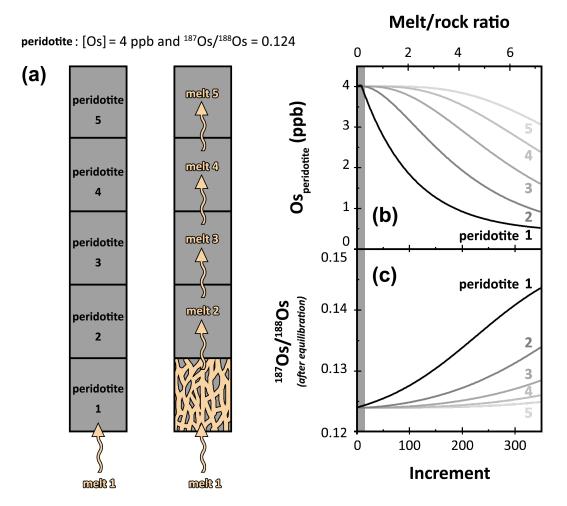
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 $melc1: [Os] = 0.20 ppb and <math>^{187}Os/^{188}Os = 0.15$

Fig. 9. Multi-cell model (see text) showing Os evolution during melt percolation through dunite channel developed in harzburgite. Starting Os compositions and sulfide fractions for harzburgite and melt are the same as those in Fig. 6. Panel (a) depicts melt percolation through a five-cell peridotite column. Melt enters the bottom cell and equilibrates with peridotite. After equilibration the melt and peridotite will have the same isotopic composition. The melt then moves into the overlying cell as a fresh melt increment enters the bottom cell; this process continues up the column. Compositions of peridotite cells (black to light grey curves) vs. melt increment are shown in (b) [Os] and (c) 187 Os/188 Os; grey band on left represents harzburgite not yet transformed into dunite. The results of the model for the lowermost cell correspond to those of the single-cell melt percolation model shown in Fig. 6. Due to the strong buffering effect of the Os-rich peridotite, the Os isotope compositions of the peridotite cells will be successively less affected by melt interaction during the progressive ascent of the melt through the channel. The compositions of the uppermost cell are hardly modified at all.

5.4. Concordance of model with geological observations

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The radiogenic Os composition of the MTZ dunites and overlying lower crust implies that they cannot have been fed by melts percolating through and equilibrating with a network of coalescing dunite channels. This inference is consistent with the scarcity of several meter thick dunites throughout the MMS. These observations contrast with the common view that large tabular dunites are frequent throughout the mantle section, as first expressed in the schematic diagram of Lippard et al. (1986). In the southern massifs of the Oman ophiolite, dunites are abundant in only three areas: (1) the MTZ, which structural, petrographic and geochemical evidence (Boudier and Nicolas, 1995; Godard et al., 2000; Koga et al., 2001; Rospabé et al., 2018) suggest formed in situ, while the overlying crust was still partially molten; (2) the BBU above the metamorphic sole; (3) the vicinity of the spectacular, yet atypical, giant Batin dunite body. In contrast, dunites are very rare even in the cores of mantle diapirs, thought to represent magmatic feeder zones for the overlying crust. The model of porous melt transport in vertical dunite channels (Kelemen et al., 1995) would predict the existence of large vertical dunite roots beneath the thick MTZ in these areas. Yet, such dunite conduits have not been found within any of the mantle diapirs. Alternatively, it has been proposed that upwardly migrating melts pool along a permeability barrier (e.g. the top of the melting region, the base of the lithosphere or the locus of plagioclase crystallization) and then migrate laterally, "uphill", along this boundary towards the ridge axis through porous flow channels or as "porosity waves" (Hebert and Montési, 2010; Sparks and Parmentier, 1991). At first glance, the basal dunite bands could correspond to the geometry suggested by this model. However, their Os isotope compositions are equivalent to those of the surrounding harzburgite, indicating they did not serve as pathways for the much more radiogenic melts that reached the MTZ and the lower crust. If, as the geochemical and structural data indicate, melt did not migrate to the Moho through dunite channels, either vertically or along a putative permeability barrier, how was it transported? It appears to have traveled rapidly across the mantle section, possibly leaving very little chemical and isotopic imprint, except perhaps as small dunite reaction zones similar to those developed at the margins of the later cross-cutting veins (Fig. 2b, 7a). If such dunites formed while the mantle was still flowing under asthenospheric conditions, they could have been transposed and stretched. These dunites may be difficult to identify now, or may correspond to the rare thin concordant bands observed in the MMS (insert in Fig. 2b and Table A.1). The modeling of Stevenson (1989) and the experimental results of Daines and Kohlstedt (1994) predict the formation of veins in the melting mantle. Rapid melt transport through hydrofractures in the asthenosphere is a physically complex process (Richardson, 1998; Rubin, 1995), whose viability remains to be better established, but this process would fit our structural and geochemical data.

6. Conclusions

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Dunite is distributed in the mantle section of the southeastern Oman ophiolite in three distinct domains: (1) the Moho Transition Zone, a horizontal dunitic melt storage and reaction zone at the summit of the mantle section, representing by far the largest volume of dunite; (2) the harzburgitic main mantle section, just below the MTZ, crosscut by sporadic dikes rimmed by dunitic reaction zones and rarer concordant bands; and (3) the basal banded unit, with many concordant dunitic bands above the metamorphic sole. Though these three types of dunite cannot be distinguished petrographically, their Os isotope compositions are strongly correlated with their structural context. Basal dunites, like harzburgites throughout the massifs, have ¹⁸⁷Os/¹⁸⁸Os_{95Ma} ratios in the range of abyssal peridotites, while the dunitic veins and bands of the MMS have variable but somewhat more radiogenic compositions. Critically, MTZ dunites have highly radiogenic Os isotope compositions, similar to those of the overlying lower crust. Such compositions are inconsistent with models that call on melt transport to the ridge by porous flow through dunite channels, as the high ¹⁸⁷Os/¹⁸⁸Os ratios of the Os-poor melts would be erased by equilibration with the unradiogenic compositions of the Os-rich dunites through which they travel. Furthermore, the absence of thick vertical dunite bodies in mantle diapirs, thought to represent magmatic feeder zones for the overlying crust, also argues that dunite channels are not the dominant pathway of melt transport to the ridge. Thus, our results reveal the need for a process, possibly hydrofracturing, that can rapidly bring radiogenic melts from the source region to the Moho

without significant equilibration with dunite or harzburgite of the surrounding mantle.

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Author Contributions

D. Klaessens, D. Jousselin and L. Reisberg participated in fieldwork. D. Jousselin provided geological context. C. Aupart completed initial analyses. D. Klaessens completed most of the analyses and wrote the initial manuscript. L. Reisberg supervised Re-Os analyses and modeling. M. Godard oversaw trace element analyses. D. Klaessens, L. Reisberg and D. Jousselin discussed the results and their implications. All authors assisted in the final revisions to the manuscript.

CRediT author statement

Delphine Klaessens: Conceptualization, Methodology, Formal analysis, Validation, Investigation, Visualization, Writing - Original Draft, Writing - Review & Editing

Laurie Reisberg: Conceptualization, Methodology, Investigation, Writing - Review & Editing, Supervision, Funding acquisition

- David Jousselin: Conceptualization, Investigation, Writing Review & Editing, Supervision, Funding
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- 644 Marguerite Godard: Validation, Writing Review & Editing, Supervision
- 645 Claire Aupart: Formal analysis, Investigation, Writing Review & Editing

Competing Interests statement

None of the authors have competing interests.

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