Multitracer determination of apparent groundwater ages in peridotite aquifers within the Samail ophiolite, Sultanate of Oman

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A B S T R A C T

CO2 sequestration in the form of carbonate minerals via alteration of oceanic crust and upper mantle is an important part of the global carbon cycle, but the annual rate of CO2 mineralization is not well quantified. This study aimed to constrain groundwater ages within the Samail ophiolite, Sultanate of Oman. Such ages could provide upper bounds on the time required for ongoing low temperature CO2 mineralization. While we were able to estimate apparent groundwater ages for modern waters, results from hyperalkaline boreholes and springs were disappointing. Waters from boreholes and hyperalkaline springs within the ophiolite were characterized using multiple environmental tracers including tritium (3H), noble gases (3He, 4He, Ne, Ar, Kr, Xe), stable isotopes (δ18O, δD), and chemical parameters (pH, Ca, Mg, DIC, etc.). Shallow peridotite groundwater and samples from boreholes near the mantle transition zone have a pH < 9.3, are 4-40 yrs old, have little to no non-atmospheric He accumulation, NGTs (noble gas temperatures) equivalent to the modern mean annual ground temperature, and stable isotopes within the range of current local precipitation. In contrast, hyperalkaline springs and deeper samples from peridotite boreholes have pH > 10, are pre-H-bomb (older than 1952), have significant non-atmospheric helium accumulation (30-70% of dissolved helium), often are isotopically heavier (enriched in δ18O), and can have NGTs 6-7°C lower than the modern ground temperature. These differences suggest that groundwater in deep (>50 m) peridotite aquifers is considerably older than shallow groundwater in peridotite and water in deeper aquifers near the mantle transition zone. Unfortunately, how much older remains an open question. The low NGT of groundwater from one deep (300 m) peridotite borehole indicates it is probably glacial in origin. If so, it must date back to at least the late Pleistocene, the most recent glacial period; He accumulation suggests it could be from 20-220 ka. The inefficacy of this suite of environmental tracers to quantitatively estimate apparent groundwater age for hyperalkaline fluids necessitates the use of different techniques. Future work to constrain groundwater ages should utilize a packer system to isolate discrete depth intervals within boreholes and less common environmental tracers such as 35Ar and 81Kr.

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1. Introduction

1.1. Natural carbon storage via seafloor alteration

Quantifying the natural global carbon budget is a key piece of the climate change puzzle, as we must understand the natural system in order to quantify the impact of human influence. One aspect of the global carbon budget is sequestration in the form of carbonate minerals due to alteration of oceanic crust and upper mantle by reacting with circulating ocean water. During this process, alteration of peridotite to form serpentine and carbonate minerals results in an average concentration on the order of 0.1 to 0.2 wt% C (e.g., Keleman and Manning, 2015; Keleman et al., 2011 and references therein), though the process can generate calcites with up to 9.6 wt% C (e.g., Agrinier et al., 1988; Schwarzenbach et al., 2013). Because of the large volumes of peridotite exposed to seawater alteration, the process is estimated to sequester 1.5-150 × 10^6 metric tons CO2/yr (tCO2/yr) (e.g.,
Alt and Teagle, 1999; Kelemen et al., 2011; Alt et al., 2013; Schwarzenbach et al., 2013; Kelemen and Manning, 2015.

The rate of CO₂ mineralization – conversion of CO₂ to carbonate minerals for permanent storage – is controlled by the rate of water-peridotite reaction and depends on a variety of factors including availability of dissolved CO₂, reactive surface area of peridotite, and kinetics of geochemical reactions. Fluid transport affects all three of these factors, and thus it is critical to characterize the flow regime beneath the surface. Studies have provided valuable insight into flow and transport properties of the upper portion of oceanic crust (e.g., Wheat et al., 2000; Fisher et al., 2003; Hutnak et al., 2006; Winslow et al., 2016; Neira et al., 2016). However, it is unlikely that fluid flow through volcanic basalt is representative of that through fractured peridotite. It is not yet technologically feasible to conduct field studies of reaction rates within peridotite of the upper mantle, but ophiolites – portions of oceanic crust and upper mantle thrust onto a continent during tectonic collisions – offer more accessible field laboratories in which to measure CO₂ mineralization.

Aqueous in the peridotite section of the Samail ophiolite in the Sultanate of Oman are sites of ongoing low-temperature alteration: circulation of meteoric water containing atmospheric CO₂ causes both serpentinization and CO₂ mineralization. Natural CO₂ mineralization within the Samail ophiolite is thought to sequester up to 10² tCO₂/yr of atmospheric carbon dioxide in the form of carbonate minerals (Kelemen and Matter, 2008). This estimate for CO₂ mineralization rate is based on the abundance and age of carbonate terraces and veins; it doesn’t include the flow rate of groundwater and the reactive surface area of the peridotite.

Groundwater age and chemistry could provide additional constraints on the rate of natural CO₂ mineralization. When water enters the groundwater system, it carries dissolved carbon from interaction with the atmosphere and soil CO₂. Along its flow path, as the water reacts with peridotite to form serpentine, the reaction releases Mg²⁺ and Ca²⁺ into the water (Eq. (1)), which then combine with the dissolved carbon to form magnesite, dolomite and calcite (e.g., Eq. (2)). This process continues until dissolved carbon is depleted, at which point Ca²⁺ begins to accumulate and greater extents of water-rock reaction produce high concentrations of dissolved Ca²⁺ and high pH, often reaching hyperalkaline values (pH 11-12).

\[ 4\text{Mg}_2\text{Si}_2\text{O}_5 + \text{CaMgSi}_2\text{O}_6 + 7\text{H}_2\text{O} \Rightarrow 3\text{Mg}_2\text{Si}_2\text{O}_5(\text{OH})_2 + \text{Ca}^{2+} + 2\text{OH}^- \]  \hspace{1cm} (1)

\[ \text{Ca}^{2+} + \text{HCO}_3^- \rightarrow \text{CaCO}_3 + \text{H}^+ \]  \hspace{1cm} (2)

The chemistry of hyperalkaline spring waters in the Samail ophiolite (pH > 11, high Ca²⁺, depleted in dissolved inorganic carbon) suggests that all dissolved CO₂ in the water has been mineralized (e.g., Neal and Stanger, 1985; Paukert et al., 2012). Groundwater flow controls CO₂ transport, so groundwater age provides the maximum amount of time during which mineralization of dissolved CO₂ could have taken place. This leaves reactive surface area and reaction kinetics as unknown variables controlling reaction rates. If we assume that reaction kinetics in the aquifers are comparable to those measured in laboratory settings, geochemical modeling can be used to calculate the minimum amount of reactive surface area necessary to produce the reaction rate per kg of peridotite that achieves complete mineralization of dissolved CO₂ in the timeframe of the apparent groundwater age.

Most CO₂ mineralization appears to occur in the upper 100 m of the peridotite, where the groundwater is closer to the atmosphere and may absorb atmospheric and soil CO₂ (Kelemen and Matter, 2008; Kelemen et al., 2018). However, additional mineralization occurs in veins at depth and in carbonate terraces when hyperalkaline water emerges at the surface and absorbs CO₂ from the atmosphere. Reaction path modeling of water-rock interaction in Oman peridotite conducted by Paukert et al. (2012) suggests that natural CO₂ mineralization at depth in aquifers closed to atmospheric exchange is limited by the amount of dissolved CO₂ in infiltrating groundwater. Thus, flow rate controls CO₂ mineralization at depth, as well as at the surface through the amount of hyperalkaline fluid emitted by springs and therefore available to react with atmospheric CO₂. The modeling also showed CO₂ mineralization may occur relatively quickly – with dissolved CO₂ being removed from the groundwater after 600 yrs of water-rock reaction – after which the groundwater continues to interact with and serpentinize the peridotite for another ~6,000 yrs before it attains a pH of 12. The ubiquity of hyperalkaline springs emerging from the Oman peridotite (Neal and Stanger, 1983, 1985) is thus inferred to result from generally long residence times for waters within the peridotite aquifer. This presents a complication: if CO₂ mineralization is concentrated in the early years of groundwater circulation, the minimum reaction rate for CO₂ mineralization (and associated reactive surface area) calculated from the apparent groundwater age would be a significant underestimate.

Because serpentinization continues to occur after dissolved CO₂ has all been mineralized, serpentinization rate estimates may provide better constraints on reactive surface area. Serpentinization in the absence of dissolved CO₂ causes a rise in pH and accumulation of Ca²⁺ in the fluid (Eq. (1)). Again, using the apparent groundwater age as a maximum duration of water-rock reaction, together with laboratory-determined kinetic rates, geochemical modeling can be used to calculate the minimum serpentinization rate and reactive surface area necessary to generate the observed pH and concentration of dissolved Ca²⁺ in the groundwater. The reactive surface area calculated from serpentinization can then be used as an input for geochemical models to better estimate the CO₂ mineralization rate.

Either method for estimating alteration rate and peridotite reactive surface area requires accurate determination of groundwater age to constrain the maximum reaction time. Thus, a better understanding of groundwater ages in the Samail ophiolite–aquifer could offer valuable information on natural CO₂ mineralization rates and reactive surface area in peridotite that could be applicable both to other ophiolites and at the seafloor.

1.2. Natural and engineered carbon storage via CO₂ mineralization in ophiolites

In addition to providing insight into the natural CO₂ mineralization process, the Samail ophiolite has been proposed as a site for an engineered geological CO₂ storage project (e.g., Kelemen and Matter, 2008; Kelemen et al., 2011). With approximately 50 × 10¹² metric tons of peridotite (Kelemen and Matter, 2008), the Samail ophiolite has the capacity to mineralize about 30 × 10¹² tCO₂ if fully carbonated. At the 2015 global rate of emissions of 36.3 × 10⁹ tCO₂/yr (Le Quéré et al., 2016), that would theoretically be enough to sequester over 800 years’ worth of anthropogenic CO₂ emissions. Though it is unlikely that the Samail ophiolite will be fully carbonated, there is still the potential for significant CO₂ storage if the rate of CO₂ mineralization can be increased.

Engineered in situ CO₂ mineralization in mafic geologic formations shows promise. Pilot projects in Wallula, WA and near Reykjavik, Iceland have demonstrated rapid (<2 yrs) CO₂ mineralization in basaltic rocks (e.g., Matter et al., 2016; McGrail et al., 2017; Snæbjörnsdóttir et al., 2017). With equivalent reactive surface area, peridotite could produce even faster reactions and CO₂ mineralization than basaltic rocks (Kelemen et al., 2011). However, as with natural CO₂ mineralization, the reaction rate will be constrained by the amount of reactive surface area among other
factors (e.g. temperature, CO₂ supply rate). Reactive transport modeling of dissolved CO₂ injection into the peridotite section of the Samail ophiolite suggests that maximizing CO₂ mineralization rates would require careful tuning of the CO₂ injection rate to the reactive surface area (Paukert, 2014). Thus, in order to accurately predict the CO₂ storage potential and design a pilot project, it is necessary to first quantify the amount of reactive surface area.

1.3. Geologic setting

The Samail ophiolite is the largest in the world, with approximately 15,000 km² of peridotite within a few kilometers of the surface (Nicolas et al., 2000; Kelemen and Matter, 2008). The Samail ophiolite aquifer is hosted by fractured crystalline rock with most of the porosity and permeability in fractures (Dewandel et al., 2005) and much of the matrix potentially shielded from interaction with aquifer fluid. Peridotite from the Samail ophiolite contains veins and fractures on a wide variety of scales, from larger fractures and veins with meter spacing down to serpentine-filled veins with 10 micron spacing (Kelemen et al., 2011). Water flow within peridotite aquifers of the Samail ophiolite is dominated by larger fractures and fissures (Dewandel et al., 2005). However, the reactive surface area is likely dominated by the smaller, 10 to 100 micron scale fractures and veins (hereafter referred to as microcracks). Microcracks also constitute pathways for fluid circulation, allowing less flow than larger fractures, but more flow than the peridotite matrix.

1.4. Apparent groundwater ages

Previous attempts to constrain the age of water in aquifers in the peridotite section of the Samail ophiolite have been unsuccessful. ³H/³He dating of hyperalkaline springs provides a minimum age, as spring waters have <0.6 TU and therefore must have recharged prior to 1952. Radiocarbon dating of spring water has been unsuccessful, as the hyperalkaline water is so depleted in dissolved carbon (e.g., Neal and Stanger, 1983; Neal and Stanger, 1985; Paukert et al., 2012) that it is difficult to get enough material to date. Also, when exposed to air these carbon-poor hyperalkaline waters rapidly absorb atmospheric carbon dioxide with modern ¹⁴C values so samples are easily contaminated by the modern signal (see ¹⁴C data in the Supplementary Material). Additionally, if the spring waters experience mixing with shallow groundwater during their migration to the surface, their ³H and carbon content may be dominated by the modern component.

Due to the limited success of some groundwater dating techniques, this study used a variety of environmental tracers: ³H, noble gases (He, Ne, Ar, Kr, Xe), and stable isotopes (¹⁸O, ²⁰H) as well as water chemistry to constrain apparent groundwater age (detailed discussion of hyperalkaline spring and shallow groundwater chemistry can be found in Paukert et al., 2012). Additionally, water samples were collected from boreholes in the peridotite to avoid the effects of mixing between deep groundwater and waters in the shallow subsurface, as may occur along the flow path of spring water. These data are used to estimate the ages of waters in different settings in the ophiolite, which may then be used to constrain natural CO₂ mineralization rates.

2. Methods

2.1. Environmental tracers

2.1.1. ³H/³He dating

³H is commonly used for identifying modern groundwater: water recharged after the advent of atmospheric thermonuclear testing in 1952. ³H concentration in precipitation peaked at more than 1,000 TU in 1963 (Fig. 1), but subsequent radioactive decay to ³He and attenuation from mixing with oceans and groundwater caused ³H in precipitation in the northern hemisphere to approach steady state by the early-mid 1990s (Rozanski et al., 1991). ³H ages can be calculated using Eq. (3):

\[
³H\text{Age} = - (t_{1/2}/\ln(2)) \times \ln(³H/³H_{\text{init}})
\]

where \(t_{1/2}\) is the half-life of ³H (12.3 yrs), ³H is the concentration of ³H measured in the sample, ³H_{init} is the concentration of ³H at the time of recharge. However, due to decay, current ³H concentrations in groundwater recharged at any point in the last few decades would show little variation (Fig. 1). Fortunately, ³H/³He dating allows more accurate determination of apparent groundwater age (Eq. (4); Tolstikhin and Kamensky, 1969):

\[
³H/³He\text{Age} = (t_{1/2}/\ln(2)) \times \ln(1 + ³H_{\text{init}}/³H)
\]

where ³H_{init} is the concentration of ³He in the sample produced by ³H decay.

2.1.2. Noble gas thermometer

Dissolved noble gases Ne, Ar, Kr, Xe can be used as a thermometer for the mean annual ground temperature at the time of recharge (e.g., Mazor, 1972; Stute and Schlosser, 1993; Aeschbach-Hertig et al., 1999). Typically, the apparent groundwater age is established through other dating methods (e.g., radiocarbon) and the noble gas temperature (NGT) is used to reconstruct paleotemperatures. However, in the case of the peridotite aquifers in Oman where other dating techniques have been unsuccessful, we may be able to compare NGT to the estimated variation of atmospheric temperature with time (e.g., Clark et al., 1997; Weyhenmeyer et al., 2000; Fleitmann and Matter, 2009; Morrissey et al., 2010) in order to constrain the apparent groundwater age.

![Fig. 1. ³H in local precipitation and borehole water samples. Solid black line is initial ³H concentration in precipitation measured at Bahrain IAEA/WMO GNIP station 4115001, with inferred concentration of 3 TU in recent precipitation. Dashed line is ³H in precipitation decay corrected for January 1, 2014. Note that decay corrected ³H concentrations would be similar for precipitation that fell between 1980 and 2000. Symbols show calculated ³H_{initial} vs recharge year determined by ³H/³He dating of modern groundwater samples.](image-url)
Table 1
Field data for borehole samples.

<table>
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<tr>
<th>Borehole</th>
<th>UTM coordinates WGS 84 datum</th>
<th>Well depth (mbtoc)</th>
<th>Screened interval depth (mbtoc)</th>
<th>Sample ID</th>
<th>Sampling depth (mbtoc)</th>
<th>pH</th>
<th>Temp. (°C)</th>
<th>EC (μS/cm)</th>
<th>ORP (mV)</th>
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<td>203</td>
<td>23-27, 39-47, 101-148, open &gt; 156</td>
<td>12.08Y</td>
<td>65</td>
<td>9.05</td>
<td>33.1</td>
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</tr>
</tbody>
</table>

Shallow boreholes

|                  |                               |                   |                               |           |                       |    |            |            |          |
|                  | 2.2B                           | 2582899           | 53                             | 17-49    | 10.11AX                | 18 | 7.23       | 33.8       | 598      | 201      |
|                  | 2.2A                           | 2582864           | 20                             | 4-16     | 10.11AY                | 14 | 7.57       | 34.0       | 491      | 164      |
|                  | 2.2B                           | 2582068           | 38                             | 20-36    | 10.11AZ                | 18 | 9.36       | 34.1       | 586      | 94       |

a All depth measurements given in meters below top of casing.
b Samples collected with point-source bailer.
c Samples collected with typhon pump.
d Previously reported in Paukert et al. (2012).

2.1.3. Non-atmospheric helium accumulation

He concentration in groundwater typically increases with time and can be used to estimate apparent groundwater age when combined with a He flux (e.g., Torgersen and Ivey, 1985; Stute et al., 1992; Torgersen and Stute, 2013). This technique has been used qualitatively in other areas of Oman (Müller et al., 2016). He can accumulate in groundwater from a variety of sources: in situ production, crustal flux, mantle flux, tritiogenic production, and release from aquifer formation rocks by enhanced diffusion or mineral dissolution (e.g., Stute et al., 1992; Solomon et al., 1996; Torgersen and Stute, 2013). The 3He/4He ratio is different in different sources, and can be used to differentiate between the different He components in some cases (e.g., Torgersen and Stute, 2013 and references therein).

2.1.4. Stable isotopes

Stable isotopes (δ18O, δ2H) in water are commonly used to determine water vapor source. Stable isotopes in meteoric water generally become depleted in heavy isotopes with increasing latitude, distance from the vapor source, and amount of rainout (e.g., Rozanski et al., 1993 and references therein). In Oman, there are two primary vapor sources: a northern, Mediterranean source that dominates during the winter, and a heavier, δ18O enriched southern, Indian Ocean source that dominates during the summer (Weyhenmeyer et al., 2002). Though the northerly source is currently the predominant one, a weakening of that system during the last glacial maximum is thought to have shifted the primary vapor source to that from the south (Weyhenmeyer et al., 2000). Stable isotopes in Oman groundwater may be used to differentiate between modern water primarily sourced from the north vs. glacial-aged water primarily sourced from the south.

2.2. Field sampling methods

Sampling was conducted over multiple field seasons, from 2009 to 2012. Water samples and dissolved noble gas samples were collected from five deep (200-350 m) boreholes set in the ophiolite (four in peridotite and one in gabbro), three shallow boreholes (<60 m) in peridotite or peridotite alluvium, and five hyperalkaline spring outlets in peridotite, comprising a total of 12 different field sites (two shallow boreholes were co-located) (Table 1, Fig. 2).

For deep boreholes, water samples for 3H, water chemistry, and stable isotope analysis were collected without purging the borehole, using a point-source stainless steel bailer with a check-valve both above and below the sample to prevent mixing with shallower water as the sample is raised to the surface. Shallow ground water samples were collected using a submersible pump and spring samples were drawn directly from spring outlets with syringes. Stable isotope and water chemistry samples were filtered to 0.2 μm in the field and samples for major cation and trace element analysis were acidified. Alkalinity was measured in the field using a two-step titration process to distinguish between carbonate alkalinity and hydroxide alkalinity.

Noble gas samples from deep boreholes were collected in airtight copper tubes (Weiss, 1968) using a borehole in situ sampling device modified from Solomon (1992) with a simple check valve operated by an ultrahigh purity nitrogen cylinder. After lowering the copper tube down the borehole to the sampling depth, pressure in the nitrogen line was reduced to open the check valve and allow water to enter the sampling line to the height of the water column. The system was then repressurized and brought to the surface to be sealed. Noble gas samples from shallow boreholes and hyperalkaline springs were collected using a submersible typhon pump and peristaltic pump, respectively.

2.3. Analytical methods

3H and noble gases were measured at Lamont-Doherty Earth Observatory (Ludin et al., 1997; Stute et al., 1995). Analytical precision was better than 2% (standard deviation 1σ) for samples with >1 TU, 15-30% for samples with <0.1 TU, 3% for helium, and 1% for neon, argon, krypton, and xenon.

Major ions and dissolved inorganic carbon were measured at Columbia University, Barnard College, and Queens College. Analytical precision was 4% or better (standard deviation 1σ). Trace elements were measured at Arizona State University, and stable isotopes (δ18O and δ2H) were measured in the Environmental Isotopes Laboratory at the University of Waterloo.

2.4. Statistical methods

Noble gas concentrations were modeled using NOBLE90, a non-linear, error-weighted least squares inversion program that fits measured noble gas concentrations with respect to pressure, salinity, excess air, and fractionation (Aeschbach-Hertig et al., 1999). Heavier noble gases were used to calculate helium due to excess
air and non-atmospheric sources. See Supplementary Material for more detailed information about model inputs.

3. Results and discussion

3.1. $^3$H/$^3$He dating

Waters with pH < 9.3 have $^3$H concentrations >0.6 TU, and waters with pH > 9.3 have <0.6 TU (Table S1, Fig. 3). Given the $^3$H concentrations in local precipitation over time, any water recharged post-1952 would have a $^3$H concentration of 0.6 TU or higher (Fig. 1). All waters with pH > 9.3 can therefore be classified as pre-bomb recharge mixed with varying fractions of modern water.

The $^3$H concentration in Oman precipitation has not been directly measured, but can be inferred from the nearest IAEA/WMO Global Network of Isotopes in Precipitation monitoring station, in Bahrain. Bahrain is less than 1000 km to the northwest of Oman and at similar elevation. Precipitation in Bahrain and northern Oman appears to have originated from the same water source and have experienced a similar transport history (Weyhenmeyer et al., 2002; Macumber, 2003; Matter et al., 2005). Bahrain modern precipitation varies between 1.6 and 4 TU interannually, with an average value of 3 TU (IAEA, 2018). It is not possible to identify the exact source function for $^3$H in precipitation in Oman, but for calculations of apparent groundwater age the modern value is taken to be 3 TU.

$^3$H/$^3$He dating (Eq. (4)) was applied to samples with >0.6 TU. To isolate $^3$He from solubility equilibrium and excess air were calculated using NOBLE90 and subtracted from total $^3$He (Aeschbach-Hertig et al., 1999). Very young samples had negative surplus $^3$He estimates, but these values were within error. For these samples, apparent groundwater ages were calculated using Eq. (3) with a $^3$He init value of 3.

Waters from gabbro boreholes (WDA 05, WDA 17, SJA 4B) had $^3$He init (calculated from $^3$H + $^3$He init) values of 3 to 11 TU and apparent ages ranging from 13 to 37 yrs, while waters from the gabbro borehole (WDA 16) were younger, with an apparent age of 4 yrs. When $^3$He init versus $^3$H/$^3$He age is plotted on the Bahrain $^3$H curve, the samples fall along the local precipitation line indicating that they have not undergone significant mixing and that the $^3$H/$^3$He apparent ages are reliable (Fig. 1).

For waters with pre-bomb $^3$H concentrations, mixing calculations were performed assuming a modern $^3$H value of 3 TU and an old $^3$H value of 0 TU. Hyperalkaline waters contain $^3$H concentrations at 0.2-6% of the modern value, suggesting small amounts of mixing with modern water. One pre-bomb borehole sample, SJA 3B, has a $^3$H concentration that is 18% of the modern value. The higher degree of mixing with modern water at this site may be due to the shallow depth of the borehole (30 m). Surface water in
Oman frequently alternates between flowing in wadis (ephemeral streams) and infiltrating into the shallow subsurface where it could mix with older groundwater.

Within boreholes, there does not appear to be a relationship between \(^3\)H/\(^4\)He apparent age and sample depth, either within the same borehole or in aggregate. Within the same borehole, there may not be much variation in apparent age with depth if water in the borehole is dominated by input from a single fracture. When one fracture is significantly more conductive than surrounding ones, most fluid in the borehole will originate from that fracture and reflect the apparent age of that source fluid. A shallow, highly conductive fracture could be responsible for the young apparent age of water at all three depths in borehole WDA 16, which is the deepest borehole and yet has the youngest \(^3\)H/\(^4\)He apparent ages (less than 5 years old). Alternatively, the young apparent age of WDA 16 water may be due to the lithology of the borehole: WDA 16 is in gabro while all other sampled boreholes are in peridotite. In the Samail ophiolite, gabro is estimated to have hydraulic conductivity 1 to 2 orders of magnitude higher than peridotite (Dewandel et al., 2005), which would potentially allow correspondingly faster flow.

3.2. Water chemistry

Waters generally group into one of two compositions commonly seen in ultramafic settings: Type I Mg\(^{2+}\)-HCO\(_3\^-\) water and Type II Ca\(^{2+}\)-OH\(^-\) water (e.g., Barnes and O'Neil, 1969; Neal and Stanger, 1985; Bruni et al., 2002). Our Type I water has pH < 10, relatively low Na\(^+\) and Cl\(^-\) (0.7-2 mmol/l) and Ca\(^{2+}\) (<0.5 mmol/l), and high Mg\(^{2+}\) and DIC (1.5-3 mmol/l) (Tables 1 and 2). Our Type II water has pH > 10.5, relatively high Na\(^+\) (10-14 mmol/l), Cl\(^-\) (14-40 mmol/l) and Ca\(^{2+}\) (4-14 mmol/l), but low Mg\(^{2+}\) and DIC (<0.5 mmol/l). The shallowest sample from borehole NSHQ 14 is intermediate between these two types.

Almost all Type I waters have modern \(^3\)H/\(^4\)He apparent ages, while all Type II water samples are pre-bomb (Fig. 3). The relationship between groundwater apparent age and water composition fits water-rock interaction models in which increasing extent of reaction with olivine and pyroxene causes an increase in pH, as in Eq. (1).

All waters with pH less than 8 are young, recharged within the last 20 years. Waters with pH between 9 and 9.3 are modern, recharged between 20 and 40 yrs ago. This suggests that less than 20 yrs of water-rock interaction are required in order to generate a pH of 9, consistent with the evolution of pH with time predicted by reaction path modeling of peridotite-water interaction (Paukert et al., 2012). All waters with pH above 9.3 are pre-bomb, suggesting that it requires at least 60 yrs for water-rock interaction to produce pH 11 water. This number is a minimum estimate: reaction path modeling predicted that it might take 500-6,500 yrs to develop pH 11-12 water (Paukert et al., 2012), though the kinetic data used to constrain that modeling are highly uncertain.

3.3. Dissolved noble gases

Noble gas concentrations for each sample (Table 3) were fitted with models in NOBLE90 (Aeschbach-Hertig et al., 1999). Calculated NGTs and excess air components, as well as selected NOBLE90 models, probabilities, and model uncertainties are given in Table 4. Modeled NGT uncertainties were all below 1°C (Table 4). Uncertainties in salinity, recharge elevation, and model choice in NOBLE90 added additional uncertainties in recharge temperature of up to 2°C (for all samples except Misbit), but as these factors each push the entire dataset in the same direction, they do not affect relative differences between samples (see Supplementary Material for detailed examination of uncertainties). Misbit spring had greater uncertainty in recharge elevation and thus a larger uncertainty in recharge temperature. Three of the five hyperalkaline spring samples and one borehole sample did not have an acceptable fit, so recharge temperatures from these samples are not presented.

3.3.1. Excess air

All borehole water samples contained excess air, as evidenced by neon being present in quantities 1.21-1.93 times solubility equi-
The two hyperalkaline spring samples that had acceptable model probabilities contained excess air, though less than the borehole samples, with Ne at 1.02 to 1.20 times solubility. The three hyperalkaline spring samples that could not be successfully modeled had no excess air and neon concentrations 0.27-0.87 times solubility. This provides evidence that these three samples must have undergone extensive degassing. Though a closed-system equilibration model has been shown to be effective for some degassed water samples (e.g., Aeschbach-Hertig et al., 2008), unfortunately our degassed spring samples could not be fitted by the model.

librium ($R_{Ne}$, Table 3). This range is typical of groundwater, which commonly contains up to 85% excess air (Heaton and Vogel, 1981).

Excess air in the Samail ophiolite aquifer may be particularly high due to the recharge pattern. Northern Oman is an arid environment that sporadically experiences heavy rainfall. Such intermittent recharge events considerably raise the water table and lead to entrapment and (partial) dissolution of air bubbles (Heaton and Vogel, 1981). Large amounts of excess air could also be produced by rapid infiltration of water without a dramatic increase in the water table: in artificial recharge areas the accelerated transport of water to depth is thought to carry entrapped bubbles and cause greater dissolution of excess air (Cey et al., 2008).
Hyperalkaline spring waters probably originally contained excess air amounts similar to the borehole samples, but lost dissolved gases as they returned to the surface, prior to reaching the spring outlet. Hyperalkaline springs in areas of peridotite serpentinization are known to have active bubbling of hydrogen and/or methane (e.g., Neal and Stanger, 1983; Chavagnac et al., 2013; Etope et al., 2013; Miller et al., 2016), and these bubbles could have stripped noble gases from the water in the shallow subsurface. Degassing could also have been caused by pressure reduction during sampling due to the use of a peristaltic pump to retrieve the sample from the spring outlet.

3.3.2. Noble gas recharge temperatures

Modeled estimates of temperature indicate that most samples recharged at 30-33°C, close to the mean annual ground temperature (Table 4). The mean annual air temperature near sample locations is 28.5-30°C (TuTiempo, 2014); ground temperatures are expected to be approximately 3°C higher as that difference is often recorded in arid locations with little vegetative cover (Smith et al., 1964; Beyerle et al., 2003). Wintertime groundwater temperatures at the water table within boreholes have been measured by geophysical well logging at 31.6°C (borehole WDA 16) and 34.8°C (borehole NSHQ 14). These estimates are similar to the previously established 33°C mean annual ground temperature for a more coastal location in Oman (Weyhenmeyer et al., 2000).

Fig. 4. Recharge temperatures modeled from noble gases. The larger and asymmetric error associated with the Misbit spring value is due to a larger uncertainty in recharge elevation. Mean annual ground temperature is estimated at 33°C.

Fig. 5. 3He/4He ratio in samples corrected for excess air as a function of the amount of excess helium, where 3He, and 4He are the concentrations measured in the sample, 3He and 4He are the concentrations due to excess air, and 3He/4He is the concentration at solubility equilibrium with air (after Stute et al., 1992). Line 1 is a mixing line between water saturated with atmospheric helium and crustal helium or helium produced in situ. Addition of tritogenic 3He from decay of 3 TU (as in pre-bomb recharge) shifts line 1 to line 2. Addition of 3He from decay of 12 TU (as in 1977 recharge, the age of our oldest sample datable by 3H-3He) shifts line 1 to line 3. Addition of 30% mantle helium shifts line 2 to line 4.

3.4. Non-atmospheric helium

Fig. 5 displays 3He/4He corrected for addition of excess air as a function of the amount of excess helium. All modern samples plot close to the value for water in equilibrium with the atmosphere between lines 1 and 3. These samples have addition of tritogenic helium but little to no accumulation of crustal or mantle helium.

One sample – the Al Hilayw hyperalkaline spring – plots below line 1, which can only be caused by removal of helium through degassing. The removal of helium by degassing in this sample is further supported by the fact that the NGT could only be fit using a model that incorporated fractionation by degassing. As discussed earlier, spring samples may have degassed due to bubbling of hydrogen or methane through the water as it approached the surface.

The two samples plotting above line 3 and to the left of solubility equilibrium – borehole NSHQ 04 and the Misbit spring – show significant helium in excess of solubility equilibrium. As both of
these samples are pre-bomb, their excess $^3$He is not from decay of bomb $^3$H and must therefore be mantle helium. NSHQ 04 has approximately 4% mantle helium, while the Misbit spring has almost 30% mantle helium. The source of mantle helium is likely alteration of the peridotite; some olivine in peridotite samples from the Samail ophiolite contains $7 \times 10^{-10}$ to $2 \times 10^{-7}$ ccSTP/g of He (Mervine, 2012). Mervine (2012) analyzed three olivine samples that had been shielded from cosmogenic radiation and hence production of cosmogenic $^3$He: all three were enriched in $^3$He relative to the atmospheric $^3$He/$^4$He ratio and one had a helium ratio similar to mantle helium ($1.16 \times 10^{-5}$). We do not have a source function for mantle helium in the region, so cannot accurately assess the time it took to accumulate.

Samples from borehole NSHQ 14 also have significant helium in excess of solubility equilibrium. The samples plot on line 1, the mixing line between atmospheric helium and in situ production or crustal flux of helium. In situ production of $^4$He occurs through the generation of alpha particles during radioactive decay of uranium and thorium in a rock. The average concentration of uranium and thorium in Oman peridotite is 2.3 ppb and 1.2 ppb, respectively (Hanghøj et al., 2010), about one thousand times lower than average crustal concentrations (Taylor, 1964). Given these uranium and thorium concentrations, the in situ helium production rate is extremely low. Assuming a porosity of 1% and that all helium produced within the rock is released to aqueous water, accumulation of helium in the peridotite occurs at $1 \times 10^{-13}$ ccSTP/jr/g water (e.g., Ballentine et al., 1991). At that rate, it would take 200–220 kys to accumulate the levels of excess helium measured in NSHQ 14 samples. If instead the effective porosity is 0.1%, the upper bound measured for upper oceanic crust and a value that implies that flow is isolated within a few channels (Neira et al., 2016), then the amount of He in the water of NSHQ 14 could accumulate in 20–22 kys. These age estimates are considered to be upper bounds. Younger water could obtain this level of non-atmospheric He through $^4$He production and accumulation within peridotite minerals over a long period of time followed by rapid release due to peridotite alteration (cracking of mantle-derived fluid inclusions, dissolution, diffusion). Relatively rapid diffusion of He from aquifer formation rock can occur due to increased surface area (Solomon et al., 1996). In the case of peridotite, surface area may be increased by alteration that produces microfractures at the 10 micron scale (Kelemen et al., 2011). Though $^3$He/$^4$He ratios measured in olivine from the Samail ophiolite were enriched in $^3$He (Mervine, 2012), local variation did occur and it is possible that in other locations the peridotite contains He enriched in $^4$He. If peridotite dissolution is primarily responsible for the addition of non-atmospheric He into groundwater, the varying $^3$He/$^4$He ratios in the rock could be responsible for varying $^3$He/$^4$He ratios in the older groundwater samples. Alternatively, crustal fluxes from underlying rock formations with higher uranium and thorium concentrations could also increase $^4$He in the groundwater more rapidly than in situ production. Again, without a source function for local crustal and mantle helium fluxes it is not possible to quantitatively calculate apparent groundwater ages.

However, these helium data are still useful for qualitatively determining relative ages. We can conclude that the five samples with significant helium in excess of solubility equilibrium are older than the samples that have not accumulated non-atmospheric helium. This is particularly useful for water from NSHQ 04, which has no age constraints other than the fact that it is pre-bomb. The water chemistry supports this relative age relationship—the only samples with significant amounts of helium in excess of solubility equilibrium are Type II waters. These waters must be relatively old in order to have attained the level of water-rock interaction necessary to develop this composition. In contrast, all samples with

3.5. Stable isotopes

$\delta^{18}$O values in groundwater vary from −2.7 to 1.9‰ VSMOW and $\delta^2$H values vary from −11.2 to 6.3‰ VSMOW (Fig. 6). Almost all borehole samples and some hyperalkaline spring samples plot between the Local Meteoric Water Line-North (LMWL-N) and Local Meteoric Water Line-South (LMWL-S) from Weyhenmeyer et al. (2002). Additional hyperalkaline spring samples and borehole samples with $\text{pH} > 11$ plot to the right of the LMWL-S, suggesting enrichment in $\delta^{18}$O. It is possible that $\delta^{18}$O values in groundwater have been altered by peridotite dissolution and CO$_2$ mineralization, and both $\delta^{18}$O and $\delta^2$H values could have been altered by serpentinitization. The effect of low-temperature water-rock interaction on stable isotopes in these fluids remains an open question, and a thorough investigation of this sort is beyond the scope of this paper. Though the following discussion is limited to the effects of water vapor source and evaporation, we do not exclude the possibility of effects from water-rock interaction, particularly in hyperalkaline waters where an extensive amount of water-rock interaction is necessary to generate water of this composition (high Ca$^{2+}$, pH > 11).

The linear regression line through all samples intersects the LMWL-N at −6.1‰ $\delta^{18}$O and −20.0‰ $\delta^2$H. This intersection is taken as the starting isotope composition for calculations to estimate the possible effect of evaporation. Rayleigh distillation curves from the starting point for evaporation at 30°C and humidities of 0% and 50% are shown in Fig. 6. While some borehole and spring samples plot along the curve for evaporation at 0% humidity, it is unlikely that the humidity was that low even in an arid environment such as Oman. Weyhenmeyer et al. (2002) estimate the humidity in Northern Oman at 50%. Generating the level of $\delta^{18}$O enrichment seen in our samples by evaporation at 50% humidity
from the starting point along the LMWL-N results in $\delta^2$H enrichment that 5 to 15% higher than observed. Thus, it is not possible to reach the stable isotope composition of most of our borehole samples through evaporation from a starting point along the LMWL-N. Instead, these waters were likely recharged by a combination of northern and southern vapor sources, as is generally the case for modern Northern Oman groundwater (e.g., Weyhenmeyer et al., 2002; Matter et al., 2005).

Many hyperalkaline spring samples plot around or to the right of the LMWL-S, indicating they have either undergone extensive evaporation from a mixed northern and southern water vapor source, or they have a predominantly southern water vapor source. All samples with NGT < 28°C (hereafter referred to as the low NGTs) plot to the right of the LMWL-S, close to other hyperalkaline waters. This suggests these low NGT samples also may have had a southern water vapor source, and in addition have some enrichment in $\delta^{18}$O and $\delta^2$H due to evaporation. During the late Pleistocene, the dominant vapor source in northern Oman shifted from the northern, Mediterranean vapor source to the more enriched southernly Indian Ocean vapor source (Weyhenmeyer et al., 2000). This shift may apply to glacial periods in general. If so, groundwater recharged during glacial periods when the southern vapor source prevailed would show a more enriched isotopic signature. A glacial origin is one possible explanation for the $\delta^{18}$O enrichment of low NGT samples.

4. Conclusions

The application of multiple environmental tracers has allowed for some constraints to be placed on the apparent age of groundwater in the Samail ophiolite. $^3$H-$^3$He dating shows Type I Mg$^{2+}$-HCO$_3^-$ waters are modern and are inferred to be 0 to 40 yrs old. Waters with pH > 9.3 (generally Type II Ca$^{2+}$-OH$^-$ water) are pre-bomb with minimal modern mixing. The Type II waters in boreholes and one hyperalkaline spring showed accumulation of helium from both crustal and mantle origins. These apparent ages and water compositions match the expected chemical evolution during serpentinitization and CO$_2$ mineralization predicted by reaction path modeling: samples reflecting greater water-rock interaction have older inferred ages.

NGTs for all samples with pH < 10 are between 30°C and 33°C with an average of 32°C, which is consistent with modern recharge. The NGT for the Al Hilayw hyperalkaline spring is 5°C lower, or 27°C. NGTs for samples from the only hyperalkaline borehole for which there were noble gas data to fit – NSHQ 14 – modeled without degassing, range from 24.7 to 25.1°C, 7°C lower than modern recharge. The NGTs for NSHQ 14 models without degassing correspond to the estimated temperature in Oman during a glacial period in the late Pleistocene (Weyhenmeyer et al., 2000) and suggest these waters are glacial in origin. Stable isotopes for low NGT samples and many hyperalkaline springs are enriched in $\delta^{18}$O, suggesting that they were primarily supplied by the southern vapor source. The southern vapor source was dominant during the last glacial maximum, consistent with the idea of glacial recharge of NSHQ 14 samples with low NGTs, and suggesting that water discharged by the hyperalkaline spring at Al Hilayw is also glacial in origin. However, our other hyperalkaline spring sample from Misbit has a modern NGT, indicating that thousands of years are not required to produce Type II waters. Stable isotope data show that most groundwater samples in the ophiolite are comprised of a mixture between water vapor sources along the LMWL-N and LMWL-S that has lost 0-30% of its water mass due to evaporation.

Unfortunately, it was not possible to assign apparent groundwater ages to any of the samples with pH > 9.3, so the minimum time necessary to generate Type II water by water-rock interaction, and hence the reactive surface area of the peridotite, remains unknown.

5. Future work

Quantitatively determining apparent groundwater age will require more exotic environmental tracers and more sophisticated sampling, e.g., the use of a packer system. Better constraints on apparent groundwater age in the peridotite aquifer may be possible with the collection of a suite of helium data with high spatial and vertical resolution. Establishing a source function for crustal and mantle helium might allow quantitative apparent age determination from helium concentrations in boreholes, rather than just relative age relationships (e.g., Stute et al., 1992). This would be particularly valuable for understanding samples from boreholes NSHQ 14 and NSHQ 04, which have significant excess helium.

The application of $^{39}$Ar and $^{81}$Kr dating could also help constrain the apparent age of peridotite aquifers in the Samail ophiolite. $^{39}$Ar has a half-life of 269 yrs, making it ideal for dating waters from a few decades to 1,000 yrs of age while $^{81}$Kr has a half-life of 229,000 yrs and can be used to date groundwater that is up to 1 Ma (e.g., Loosli et al., 2000; Ritterbusch et al., 2014; Jiang et al., 2017).

Additionally, more noble gas data from borehole and hyperalkaline spring samples could provide greater insight on apparent groundwater age from NGTs. In our dataset, three out of four samples with pH > 10 have low NGTs, but two of these three are from the same borehole. A larger dataset would allow us to determine if this is a coincidence, or if most pH > 10 waters have low NGTs and are glacial in origin.

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Appendix A. Supplementary material

Supplementary material related to this article can be found online at https://doi.org/10.1016/j.epsl.2019.03.007. The Supplementary Material file contains a table of stable isotope ($\delta^{18}$O, $\delta^2$H, $^3$H, and $^{14}$C) data for boreholes and hyperalkaline springs, as well as a detailed discussion of the sensitivities and uncertainties associated with our dissolved noble gas temperatures.


