Diffusion of molybdenum and tungsten in anhydrous and hydrous granitic melts

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ABSTRACT

To better understand the transport of Mo and W in granitic melts and the formation mechanism of porphyry ore deposits, we have investigated the diffusivities of Mo and W in granitic melts with 0.04-5.1 wt% H₂O at 1000-1600 °C and 1 GPa using a diffusion couple approach and a Mo saturation approach with Mo sheet serving as the source. The Mo and W diffusivities obtained from diffusion profiles measured by LA-ICP-MS can be described as:

\[ D_{\text{Mo, anhy}} = 10^{-1.47\pm0.73} \exp\left[-\frac{(387 \pm 25)}{RT}\right], \]
\[ D_{\text{W, anhy}} = 10^{-1.28\pm1.05} \exp\left[-\frac{(396 \pm 35)}{RT}\right], \]
\[ D_{\text{Mo, 2.7 wt\% H₂O}} = 10^{-5.37\pm0.52} \exp\left[-\frac{(211 \pm 18)}{RT}\right], \]
\[ D_{\text{Mo, 5.1 wt\% H₂O}} = 10^{-6.87\pm0.69} \exp\left[-\frac{(133 \pm 20)}{RT}\right], \]

where \( D \) is diffusivity in m²/s (with the subscripts denoting water contents and “anhy” representing nominally anhydrous melt), R is the gas constant, T is temperature in K, and the activation energies in the exponential are in kJ/mol. When the influence of H₂O is incorporated, Mo diffusivity in granitic melts with <5.1 wt% H₂O can be modeled as:

\[ \log D_{\text{Mo}} = - (1.94 \pm 1.58) - (0.87 \pm 0.36)w - \frac{((19341 \pm 2784) - (2312 \pm 620)w)}{T} \]

where \( w \) is H₂O content in the melt in wt%. The diffusion behavior (low diffusivities, high activation energies, and strong H₂O effects) of Mo and W indicates that they exist and diffuse in the melt in the form of hexavalent cations. Their low diffusivities imply that the bulk concentrations of Mo and W in exsolved hydrothermal fluid and those in the
melt are probably not in equilibrium. However, because of the large fluid-melt partition
coefficients of Mo and W, they can still be enriched, although to a lesser extent than
equilibrium partitioning would allow, in the hydrothermal fluid. Slow Mo and W
diffusion can be a significant rate-limiting step for the formation of porphyry Mo/W
deposits.

**Keywords:** Porphyry deposits, molybdenum, tungsten, diffusivity, granitic melt
INTRODUCTION

Porphyry-type ore deposits, of which the majority were formed in the Phanerozoic (Kesler and Wilkinson 2008), are commonly found at oceanic or continental arcs above subduction zones (Wilkinson 2013). They are the primary source of copper and molybdenum (approximately accounting for 60% of all mined Cu and 99% of all mined Mo) and an important source of gold, silver and tungsten (Singer 1995; Simon and Ripley 2011). Porphyry deposits are typically large-tonnage and low-grade (e.g., Cu, Mo and W generally less than 1 wt%, 0.2 wt% and 0.2 wt%, respectively; Kirkham and Sinclair 1995). They are associated with intermediate to felsic hypabyssal intrusions (granite porphyry and granodiorite porphyry) located in mid- to shallow crust (Sedorff et al. 2005; Sillitoe 2010).

Porphyry deposits are a cumulative product of a series of geological processes starting from the dehydration of subducted slab (Richards 2005). Within crustal depths, the formation of a porphyry deposit generally involves the following steps (e.g., Annen et al. 2005; Candela and Holland 1986; Candela 1997; Landtwing et al. 2010; Stoffell et al. 2004; Wilkinson 2013): (1) production of metal-containing hydrous magma in the deep crust; (2) emplacement of the magma in the mid- to shallow crust, and enrichment of metals into exsolved magmatic hydrothermal fluid, partly driven by crystallization; (3) deposition of ore minerals locally from the hydrothermal fluid. The second step requires not only that the metals are partitioned into the fluid relative to the silicate melt, but also
that the transport of metals in the melt is rather efficient. Huber et al. (2012) demonstrated with quantitative modeling that rapidly diffusing metals tended to establish equilibrium fluid-melt partition and hence be efficiently scavenged from the melt, whereas slowly diffusing metals would be left behind in the melt. A great deal of effort has been devoted to investigating the partitioning behavior of metals between fluid and melt (e.g., Candela and Holland 1984; Manning and Henderson 1984; Keppler and Wyllie 1991; Bai and Van Groos 1999; Zajacz et al. 2008; Frank et al. 2011; Tattitch and Blundy 2017). By contrast, studies of diffusion kinetics of ore-forming metals in silicate melts are scarce. Among the three major metals of porphyry deposits (Cu, Mo, W; Robb 2004), several recent experimental studies already reported Cu diffusivities (Ni and Zhang 2016; Ni et al. 2017; Ni et al. 2018), but the diffusivity data for W and Mo are much fewer. There was only a single study on W diffusion in anhydrous haplogranitic melt and in that melt doped with extra Na$_2$O (Mungall et al. 1999). For Mo, there are no measurements to our knowledge.

Due to the lack of Mo and W diffusivity data that are needed for quantitative modeling of processes in the formation of porphyry deposits, the present study experimentally investigates the diffusion of Mo and W in silicate melts of granitic composition. The results are used to discuss the transfer and enrichment of magmatic Mo and W into ore-forming hydrothermal fluids.
EXPERIMENTAL AND ANALYTICAL METHODS

Starting material

The composition of granitic melt was based upon that of the east porphyry of the Chuquicamata porphyry complex, Chile (Ballard 2001), which hosts the largest open pit Cu deposit worldwide. Cu diffusivity in this melt has been reported elsewhere (Ni et al. 2018). Chuquicamata also produces a significant amount of molybdenum (Ossandon et al., 2001).

The anhydrous granitic glass Anhy-Gran was synthesized by fusing a mixture of oxides (SiO$_2$, Al$_2$O$_3$, MgO, Fe$_2$O$_3$, TiO$_2$) and carbonates (K$_2$CO$_3$, Na$_2$CO$_3$, CaCO$_3$) at 1600 °C in a muffle furnace for 4 h, after decarbonation at 1000 °C for 10 h. Then glass powder of Anhy-Gran was divided into three parts. Two parts were doped with different amounts of MoO$_3$ and WO$_3$ and remelted at 1600 °C for 4 h to obtain Mo-rich, W-poor (Anhy-Gran-Mo) and W-rich, Mo-poor (Anhy-Gran-W) granitic glasses. The doped glasses were crushed and remelted at 1600 °C for 4 h to ensure homogeneous distribution of Mo and W. Electron microprobe analyses show that the major compositions of the synthesized glasses closely resembled the east Chuquicamata porphyry (Table 1). LA-ICP-MS analyses indicate that Mo and W were distributed homogeneously in the Mo and W doped glasses. Anhy-Gran-Mo contains ~1900 ppm and ~300 ppm of Mo and W, respectively. Anhy-Gran-W contains ~1900 ppm and ~300 ppm W and Mo, respectively (Table 1). Water contents of these glasses were found to be ~110 ppm based upon FTIR
measurements. To synthesize hydrous granitic glasses, the mixture of anhydrous glass powder and distilled water was sealed in a Pt capsule and fused at 1200 °C and 0.5 GPa in a piston cylinder apparatus for 12 h. The hydrous glasses synthesized from the third part of Anhy-Gran (undoped with Mo or W) contained ~2.8 wt% or ~5.1 wt% H$_2$O, whereas those synthesized from Anhy-Gran-Mo and Anhy-Gran-W contained ~3.9 wt% H$_2$O. All of the synthesized glasses had a light brown color and were free of crystals. The anhydrous glasses contained a small amount of air bubbles.

**Diffusion experiments**

The diffusion experiments were carried out in a 3/4" end-loaded piston cylinder apparatus at the University of Science and Technology of China (USTC). Two different approaches were adopted. The diffusion couple approach, aimed at obtaining Mo and W diffusivities together, involved two halves, with one being Mo-rich and W-poor and the other being W-rich and Mo-poor. Two granitic glass cylinders with a diameter of 2.7 mm and a height of 1.5 mm were doubly polished and welded shut in a Pt capsule of 3 mm outer diameter and 0.15 mm wall thickness, with the W-rich half being placed at bottom (Fig. 1a). In Mo-saturated diffusion experiments aimed at obtaining Mo diffusivity, an undoped granitic glass cylinder of 1.5-2.5 mm height and 2.7 mm (for the samples with ~2.9 wt% H$_2$O) or 4.7 mm (for the samples with ~5.1 wt% H$_2$O) diameter was placed on top of a Mo sheet of 25 μm thickness, with the latter serving as the source of Mo (Fig.
1b). They were welded shut in a Pt capsule of 3 mm or 5 mm outer diameter and 0.15 mm wall thickness.

From the outside to the inside, the sample assembly comprised of talc, Pyrex glass, graphite heater and crushable MgO with the Pt capsule fit inside. A type-S thermocouple at the top of the Pt capsule served to monitor the temperature during the experiment. The thermocouple tip was only \( \sim 2 \) mm away from the portion of the sample containing the effective diffusion profile. The thermal gradient over that distance was found to be small, and therefore no temperature correction was applied. A piston-out procedure was used and a pressure correction of 15\% was applied (Guo et al. 2016). The sample assembly was first relaxed at 473 K and 1 GPa for at least 2 hours and was then rapidly heated (at 100 °C/s) to the target temperature ranging from 1000-1600 °C by a programmed procedure. The 1 GPa pressure guaranteed that the hydrous melts remained to be water undersaturated. Rapid quench by turning off the power was applied after a dwell time of 1-93 h to achieve appropriate diffusion distances.

After the diffusion experiments, the recovered samples were cut into two halves longitudinally using a diamond saw. One half was mounted into epoxy resin and polished for LA-ICP-MS analysis of Mo and W concentrations, and the other half was doubly polished for FTIR analysis to measure H\(_2\)O contents.

**Analytical methods**
The major element compositions of the granitic glasses were analyzed with a Shimadzu EPMA-1600 electron microprobe at USTC using a beam of 15 kV, 10 nA and 10 μm diameter. Twelve points were measured on each glass to obtain an average composition.

Water contents of the starting glass cylinders and the experimental products were measured by a Spotlight 200 microscope system attached to a PerkinElmer Frontier FTIR spectrometer at USTC, using an MIR or NIR source, a CaF$_2$ or KBr beamsplitter, and an MCT detector. FTIR spectra were collected using 64 scans. Multiple measurements confirmed that water was homogeneously distributed in the samples. For the nominally anhydrous glasses, the calibration of Leschik et al. (2004) for the O-H stretching band at 3550 cm$^{-1}$ (a linear molar absorptivity of 8.0 L mol$^{-1}$ mm$^{-1}$) was used. For the hydrous glasses, the calibration of Withers and Behrens (1999) for the combination bands at 5200 cm$^{-1}$ (0.166 L mol$^{-1}$ mm$^{-1}$) and 4500 cm$^{-1}$ (0.141 L mol$^{-1}$ mm$^{-1}$) was used to obtain molecular H$_2$O and OH concentrations, where the baselines were fit by two tangential lines (TT calibration). The total water contents were obtained by summing the concentrations of the two hydrous species.

Concentrations of Mo and W of granitic glasses before and after diffusion experiments were measured with LA-ICP-MS at USTC (Coherent GeoLas HD Excite excimer 193nm short pulse width attached to Agilent 7900 quadrupole mass spectrometer). The sample chamber was flushed with helium at a rate of 0.5 L/min, with
1 L/min Ar being added on the way to the ICP-MS. Spot analyses were performed with a beam diameter of 32 μm and repetition rate of 6 Hz. For each analysis, the sample was measured for 45 s (7 ms dwell time for each mass) after measurement of the gas background for 20 s. The analytical results were processed with the ICPMSDataCal routine (Liu et al. 2008; Chen et al. 2011). External standardization was performed on the NIST SRM 610 and SRM 612 silicate glass (used for Mo and W), BCR-2G, BHVO-2G and BIR-1G. Internal standardization was achieved by normalizing the total of all of the measured elements to 100%. Detection limits were of the order of 1 ppm, and typical analytical precision was 2% to 4%. For each experimental product, 3 diffusion profiles consisting of 30-80 points were measured along the centerline and 250 μm apart from it on both sides (Fig. 2).

RESULTS

Four diffusion couple experiments and nine Mo saturation experiments have been conducted, with the conditions and results presented in Table 2. The recovered samples have largely preserved their original geometry (Figure 2). Although the samples usually contained several cracks quasi-perpendicular to the cylindrical axis, diffusion profiles across the cracks appeared to be smooth. As such, we decided not to correct for the widths of cracks, which are much shorter than the diffusion profiles.

LA-ICP-MS analyses indicate that the major element compositions of experimental
products remain essentially unchanged except for FeO. For the samples from diffusion
couple experiments, FeO contents adjacent to capsule walls dropped to 0.5-1.2 wt%, due
to the widely known problem of Fe loss to Pt (Holloway and Wood 1988), but this did
not affect melt interior where Mo and W diffusion profiles reside. The samples from Mo
saturation experiments suffered more severe Fe loss. In most of them, FeO contents
decreased from 0.5-1.0 wt% at the Mo end to 0.04-0.2 wt% at the opposite end,
indicating a weaker affinity of Fe with Mo than with Pt. One exception is sample
Hydr-Gran-9, with FeO content being lower (~0.5 wt%) at the Mo end and higher (~1.0
wt%) at the opposite end.

FTIR analyses show that the anhydrous samples have gained a small amount of water
after experiments, with H$_2$O content increased to 0.06-0.19 wt% (Table 2). Most hydrous
samples have preserved their original water (Table 2). However, the H$_2$O content of
Hydr-Gran-4 decreased from 3.0 to 2.3 wt%. More severe water loss was found for
Hydr-Gran-9, the H$_2$O content of which dropped from 5.1 wt% to 3.6 wt%. To keep
consistency, the mean value of initial and final water contents was used to characterize
the sample regardless of the degree of water loss or gain (Table 2).

**Fitting of diffusion profiles**

The Mo and W concentration profiles measured along the centerlines of samples
were used for fitting and extraction of diffusivity. Those from diffusion couple
experiments (Fig. 3) were fit by the analytical solution for one-dimensional diffusion in
an infinite medium (Crank 1975):

\[
C = \frac{C_+\infty + C_-\infty}{2} + \frac{C_+\infty - C_-\infty}{2} \text{erf} \left( \frac{x - x_0}{\sqrt{4Dt}} \right),
\]

(1)

where \(C_+\infty\) and \(C_-\infty\) are the initial concentrations of Mo or W of the two halves of the
diffusion couple respectively, \(x_0\) is the position of the interface, \(t\) is time, and \(D\) is
diffusivity of Mo or W. Given that the locus of the original interface cannot be
determined precisely, \(x_0\) is not constrained in fitting.

The concentration profiles of Mo saturation experiments (Fig. 4) were fit by the
analytical solution for one-dimensional diffusion in semi-infinite medium (Crank 1975):

\[
C = C_0 + (C_\infty - C_0) \text{erf} \left( \frac{x}{\sqrt{4Dt}} \right),
\]

(2)

where \(C_\infty\) is the initial Mo concentration of the undoped granitic glass and is taken to be
zero, and \(C_0\) is the Mo concentration at the melt-Mo interface (i.e., Mo solubility) and is
not constrained.

The values of \(C_0\) from fitting are summarized in Table 2, ranging from 0.25 wt\% to
1.8 wt\%. Interface Mo concentration increases with increasing temperature and H\(_2\)O
content (Fig. 5). However, Hydr-Gran-9, the sample that experienced the most significant
water loss, shows abnormally high \(C_0\) at 1000 °C which is inconsistent with the general
trend.
Diffusivities of Mo and W in granitic melts

The diffusivities of Mo and W obtained from least-squares regression of measured profiles with error function curves (equation 1 and equation 2) are summarized in Table 2 and Figure 6. Results from the four diffusion couple experiments indicate similar diffusivity for Mo and W at both anhydrous and hydrous conditions. In addition, the two experimental approaches, diffusion couple and Mo saturation, yield consistent diffusivity values.

The data sets collected at a series of temperatures but roughly the same H$_2$O content show good agreement with the Arrhenius relationship $D = D_0 \exp(-E_a/R T)$ with $D_0$ being the pre-exponential factor, $E_a$ being the activation energy, $R$ being the gas constant, and $T$ being temperature. Diffusivities of Mo and W in nominally anhydrous granitic melt (with H$_2$O <0.1 wt%) at 1 GPa and 1400-1600 °C can be expressed by the following Arrhenius relations:

$$D_{Mo,anhy} = 10^{-1.47\pm0.73} \exp[-(387\pm25)kJ/mol/R T] \quad (3)$$

$$D_{W,anhy} = 10^{-1.28\pm1.05} \exp[-(396\pm35)kJ/mol/R T] \quad (4)$$

where $D$ is in m$^2$/s and $T$ is in K.

Diffusivities of Mo in hydrous granitic melts with ~2.7 wt% and ~5.1 wt% H$_2$O at 1 GPa and 1400-1600 °C and 1100-1400 °C can be expressed by the following expressions, respectively:

$$D_{Mo,2.7wt\%H_2O} = 10^{-5.37\pm0.52} \exp[-(211\pm18)kJ/mol/R T] \quad (5)$$
$D_{\text{Mo, 5.1 wt\% H}_2\text{O}} = 10^{-6.87\pm0.69} \exp[-(133\pm20)\text{kJ/mol}/RT] \quad (6)$

where $D$ is in m$^2$/s and $T$ is in K.

Water has a significant impact on Mo and W diffusion in granitic melt. Using nominally anhydrous melt as the benchmark, at 1400 °C, Mo diffusivity increases by a factor of 40 with the addition of 2.7 wt\% H$_2$O, and by a factor of 300 with 5.1 wt\% H$_2$O. If extrapolated to 800 °C, a more realistic temperature for felsic magmatic systems, the influence of water would be much more dramatic: Mo diffusivity would increase by 7 orders of magnitude with the addition of 5.1 wt\% H$_2$O. The strong influence of water is also evident with regard to activation energy: $E_a$ for Mo diffusion decreases from 387 kJ/mol in anhydrous granitic melt to 211 kJ/mol at 2.7 wt\% H$_2$O, and to 133 kJ/mol at 5.1 wt\% H$_2$O. When compared to the diffusivities of other cations (e.g., Mungall et al. 1999), the influence of water on Mo and W is stronger than that on rapidly diffusing species such as Ca and Sr, but is weaker than that on slowly diffusing species such as Zr and Hf.

The logarithm of Mo diffusivity at 1400 °C increases roughly linearly with H$_2$O content (Fig. 7a). Closer inspection reveals that both the logarithm of the pre-exponential factor $D_0$ (Fig. 7b) and the activation energy $E_a$ (Fig. 7c) decrease roughly linearly with H$_2$O content. Based on the data sets of Mo diffusivity at three different H$_2$O contents (<0.1 wt\%, ~2.7 wt\%, and ~5.1 wt\%), the following general expression of Mo diffusivity at 1 GPa in granitic melts containing <5.1 wt\% H$_2$O was obtained:
\[
\log D_{\text{Mo}} = - (1.94 \pm 1.58) - (0.87 \pm 0.36)w - [(19341 \pm 2784) - (2312 \pm 620)w]/T \quad (7)
\]

where \( D \) is in m\(^2\)/s, \( w \) is water content in granitic melt in wt\%, and \( T \) is in K. Equation (7) reproduces the majority of \( \log D \) values to within 0.16 \( \log D \) units. When the mean values of initial and final H\(_2\)O contents are used, the Mo diffusivity data obtained from Hydr-Gran-1 and Hydr-Gran-9 are roughly consistent with equation (7), despite that these data were not used for developing the general expression and that Hydr-Gran-9 experienced severe H\(_2\)O loss and has an abnormally high interface Mo concentration. In view of the similarity between W diffusivity and Mo diffusivity, equation (7) is also considered to be applicable to describing W diffusion.

DISCUSSION AND IMPLICATIONS

Complications

One may question whether convection has shaped the measured Mo and W concentration profiles in granitic melts. In their study on Cu diffusion, Ni et al. (2017) dismissed a significant convection effect based upon three lines of reasoning: (1) granitic melts have high viscosity; (2) multiple traverses give consistent diffusion profiles; (3) experiments with different dwell durations yield consistent diffusivity. These arguments are also applicable to our experiments. For example, three traverses, one along the centerline and the other two 250 \( \mu \)m apart from it on each side (Fig. 2), show good consistency (Fig. 3 and Fig. 4). The regular variation of diffusivity with temperature and
H$_2$O content also imply that convection was unlikely a serious problem in our experiments.

At the effective diffusion profiles, the limited changes in FeO contents (negligible in diffusion couple experiments but from 1.7 wt% to 0.5-1.0 wt% in Mo saturation experiments) are not expected to have any major effect on Mo and W diffusion. Sample Hydr-Gran-9 shows severe H$_2$O loss (5.1 wt% dropped to 3.6 wt%), abnormally high interface Mo concentration (1.8 wt%), and atypical Fe-loss pattern (more severe at the Mo end than at the opposite Pt end). H$_2$O loss was probably due to failure of the Pt capsule during the long dwell duration (43 h). We infer that H$_2$O escaped from the melt in the form of H$_2$, leaving behind an increased Fe$^{3+}$/Fe$^{2+}$ ratio and hence increased intrinsic oxygen fugacity. A more oxidized environment would mitigate Fe loss to Pt (Ford 1978). On the other hand, it probably enhanced the dissolution of Mo into the melt, giving rise to an abnormally high $C_0$. To be on the safe side, we excluded this experiment when developing the general expression of Mo diffusivity, but the Mo diffusivity obtained from Hydr-Gran-9 appears to be roughly consistent with equation (7) when the mean value of the initial and final water contents is used for calculation.

Oxygen fugacity was not controlled in our experiments. The intrinsic redox state of our sample assembly was probably close to the fayalite-magnetite-quartz (FMQ) buffer (Jakobsson 1997), but the starting anhydrous granitic melts were more oxidized because they were synthesized in air. The speciation of Mo and W in silicate melts is not sensitive
to redox state: these two elements exist predominantly as Mo\(^{6+}\) and W\(^{6+}\) in silicate glasses and melts over a broad range of oxygen fugacity, from air to the iron-wüsite buffer (Farges et al. 2006a, 2006b; O’Neill et al. 2008; Wade et al. 2013). Therefore, the diffusion of Mo and W is unlikely to be affected by oxygen fugacity in a significant way.

One may also wonder whether the similar Mo and W diffusivities obtained from diffusion couple experiments were an artifact of interdiffusion. In other words, Mo diffusion and W diffusion were coupled and what was measured is actually Mo-W interdiffusivity. However, Mo and W are generally trace, or at most minor, elements in the melts, since even the interface Mo concentrations are typically only several thousand ppm (Table 2). In Mo saturation experiments, despite the decrease of Mo activity from ~1 at the interface to ~0 at far interior of the melt, the diffusion profiles can be well fit by error function curves, indicating trace element behavior of Mo. Furthermore, in both anhydrous and hydrous melts, the Mo diffusivities obtained from diffusion couple experiments are in good agreement with those from Mo saturation experiments. We therefore conclude that Mo and W behave as trace elements in silicate melts and should not interfere much with each other with respect to diffusion. The similarity between Mo diffusivity and W diffusivity is real.

Comparison with previous work

Mungall et al. (1999) measured diffusivities of W alongside other 17 trace elements
in anhydrous haplogranitic melt containing 80 wt% SiO$_2$ and 12% Al$_2$O$_3$, at 1600, 1400, 1137 °C and 1 atm. Compared to their composition, our granitic melt is less siliceous and more aluminous. Considering that the pressure effect for element diffusion in silicate melt is generally negative and is less significant compared to the influence of temperature and H$_2$O content (Zhang et al. 2010), the small difference in W diffusivity between Mungall et al. (1999) and this study (Fig. 8) is attributed mainly to the difference in melt composition rather than to the 1 GPa pressure gap. The W diffusivity data in haplogranitic melt doped with 20 wt% Na$_2$O are comparable to Mo diffusivity in granitic melt with 5 wt% H$_2$O, indicating a stronger influence of H$_2$O than Na$_2$O on a wt% basis (but comparable on a molar basis).

In anhydrous granitic melts, diffusivities of W and Mo are much lower than those of univalent and divalent cations such as Cu$^+$ and Ca$^+$, but are similar to those of pentavalent cations such as Nb$^{5+}$ and Ta$^{5+}$ (Fig. 8). This is consistent with that Mo and W exist and diffuse in the form of hexavalent cations in silicate melts, since Mo$^{6+}$, W$^{6+}$, Nb$^{5+}$ and Ta$^{5+}$ not only are similar with respect to valence but also have almost identical ionic radii (0.073-0.078 nm, Shannon 1976) in octahedral sites.

**Implications**

Compared to porphyry Cu deposits, porphyry Mo deposits and porphyry W deposits generally formed at greater depths, at a minimum of 4-8 km (Simon and Ripley 2011;
Robb 2004). In the formation of porphyry Mo and W deposits, before reaching the point of water saturation, a significant degree of crystallization probably took place. As Mo and W are incompatible elements, crystallization tended to enrich Mo and W in silicate melt (Robb 2004; Adam and Green 2006, 2011; Wilkinson 2013), unless a sulfide phase formed which strongly sequestered Mo (Mengason et al. 2011; Li and Audétat 2012).

For the growth of hydrothermal fluid bubbles from silicate melt, H$_2$O is the principal equilibrium-determining component as defined in Zhang (2008). In a hydrous granitic melt with ~5 wt% H$_2$O, Mo diffusivity at 800 °C is estimated to be 2 orders of magnitude lower than H$_2$O diffusivity (Ni and Zhang 2008; Fig. 8). Therefore, the concentrations of Mo and W in the fluid are unlikely in equilibrium with their bulk concentrations in the melt. The degree of disequilibrium is also controlled by their partitioning behavior. The fluid-melt partition coefficients ($K$) of Mo and W increase with the salinity of the fluid and mostly fall in the range of 5 to 100 (Manning and Henderson, 1984; Zajacz et al., 2008; Tattitch and Blundy 2017 and references therein). According to equation (4-145) in Zhang (2008), the enrichment factor (the ratio of element concentration in the fluid over that in the melt) of a trace element can be estimated by

$$EF = \frac{K}{1 + \sqrt{\pi} \gamma e^{\gamma^2} \left[1 - \text{erf}(\gamma)\right](K-1)}, \quad (8)$$

where $\gamma$ is a dimensionless factor proportional to the square root of the diffusivity ratio $D/D_{H2O}$. When $D \ll D_{H2O}$, $\gamma \gg 1$ and $\sqrt{\pi} \gamma e^{\gamma^2} \left[1 - \text{erf}(\gamma)\right] \approx 1$, leading to an enrichment factor of 1. In this limiting case, the hydrothermal fluid will have the same concentration.
as the bulk melt (i.e., no enrichment). For Mo and W, the difference between their diffusivity and $D_{H2O}$ may result in $\gamma$ in the range of 0.1 to 2, leading to enrichment factors smaller than $K$ but larger than 1 (Fig. 9).

In summary, owing to the large partition coefficients of Mo and W between fluid and melt, they can still be enriched, although to a lesser extent than equilibrium partitioning would allow, in the hydrothermal fluid. However, the slow diffusive transport of Mo and W in silicate melts can be a significant rate-limiting step for the formation of porphyry deposits.

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FIGURE CAPTIONS

FIGURE 1. Sketches of samples in diffusion experiments: (a) diffusion couple composed of a Mo-rich, W-poor glass cylinder (with ~1900 ppm Mo and ~300 ppm W) and a W-rich, Mo-poor glass cylinder (with ~1900 ppm W and ~300 ppm Mo), sealed in a Pt capsule; (b) Mo saturation method involving an undoped glass cylinder and a Mo sheet serving as the source of Mo.

FIGURE 2. Photomicrograph of experimental products: (a) sample Anhy-Gran-2; the two halves coalesced after the experiment, but the interface distinguishes itself by tiny intrusion of Pt capsule walls; (b) sample Hydr-Gran-7 after Mo saturation experiment. For each sample, laser ablation pits along three profiles are indicated.

FIGURE 3. Diffusion profiles of Mo and W from four diffusion couple experiments. (a), (b) and (c) anhydrous granitic melts; (d) hydrous granitic melt with 3.9 wt% H_2O, with three points of original concentrations (squares) added at the W-rich end due to damage of the sample during polishing. Solid circles denote data of the central profile, and open circles and crosses denote data from two side profiles. The curves are error function fits of the central profiles.
FIGURE 4. Diffusion profiles of Mo from nine Mo saturation experiments. (a) anhydrous granitic melt; (b), (c) and (d) hydrous granitic melt with ~2.7 wt% H₂O; (e), (f), (g) and (h) hydrous granitic melt with ~5.1 wt% H₂O; (i) hydrous granitic melt with ~4.4 wt% H₂O (5.1 wt% dropped to 3.6 wt%). Solid circles denote data of the central profile, and open circles and crosses denote data from side profiles. A few odd points near the interface (marked in grey) are excluded from fitting. The curves are error function fits of the central profiles.

FIGURE 5. Variation of Mo concentration at the melt-Mo interface in Mo saturation experiments with temperature and H₂O content. Anomalous interface concentration is observed for Hydr-Gran-9 at 1000 °C in which H₂O content dropped from 5.1 wt% to 3.6 wt%. See text for detailed interpretation.

FIGURE 6. Arrhenius plot of the diffusivity data of Mo (solid symbols) and W (open symbols) in granitic melts at 1 GPa obtained in the present study. For Mo diffusivity in anhydrous melt, the data from diffusion couple experiments are shown in circles and a single datum from a Mo saturation experiment is shown in cross. The squares and triangles represent Mo diffusivities from Mo saturation experiments, and the diamonds represent Mo and W diffusivities from a diffusion couple experiment, with H₂O contents of melts marked in the labels. Error bars are smaller than the symbol sizes.
FIGURE 7. Variation of (a) Mo diffusivity at 1400 °C; (b) the pre-exponential factor $D_0$; and (c) the activation energy for Mo diffusion in granitic melts with H$_2$O content. Error bars are at 1σ level.

FIGURE 8. Comparison of diffusivities of Mo and W obtained in this study (solid lines) with literature diffusivity data of W and other species (dashed lines and symbols) in granitic melts. Water contents are given in the labels for hydrous melts. The triangles denote W diffusivity data in Mungall et al. (1999) for both a granitic melt and that melt with 20 wt% added Na$_2$O. Ca, Nb, Ta, Mungall et al. (1999); Cu, Ni et al. (2017); H$_2$O, Ni and Zhang (2008).

FIGURE 9. Variation of enrichment factor of Mo and W (concentration in the fluid over that in the melt) for a range of fluid-melt partition coefficients ($K$) and parameter $\gamma$ depends on the ratio of Mo or W diffusivity over water diffusivity in the melt. Calculated with the trace element diffusion-partitioning model in Zhang (2008).
TABLE 1. Chemical compositions of the synthesized granitic glasses in wt%

<table>
<thead>
<tr>
<th></th>
<th>Chuquicamata&lt;sup&gt;a&lt;/sup&gt;</th>
<th>Starting material&lt;sup&gt;b&lt;/sup&gt;</th>
<th>After experiment&lt;sup&gt;c&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Anhy-Gran</td>
<td>Anhy-Gran-Mo</td>
<td>Anhy-Gran-W</td>
</tr>
<tr>
<td>SiO&lt;sub&gt;2&lt;/sub&gt;</td>
<td>69.83</td>
<td>70.96±0.48</td>
<td>71.08±0.44</td>
</tr>
<tr>
<td>TiO&lt;sub&gt;2&lt;/sub&gt;</td>
<td>0.27</td>
<td>0.23±0.03</td>
<td>0.25±0.04</td>
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<tr>
<td>Al&lt;sub&gt;2&lt;/sub&gt;O&lt;sub&gt;3&lt;/sub&gt;</td>
<td>16.54</td>
<td>15.59±0.15</td>
<td>15.44±0.31</td>
</tr>
<tr>
<td>FeO&lt;sub&gt;t&lt;/sub&gt;</td>
<td>1.67</td>
<td>1.64±0.11</td>
<td>1.74±0.09</td>
</tr>
<tr>
<td>MgO</td>
<td>0.58</td>
<td>0.56±0.03</td>
<td>0.60±0.04</td>
</tr>
<tr>
<td>CaO</td>
<td>1.74</td>
<td>1.36±0.06</td>
<td>1.38±0.06</td>
</tr>
<tr>
<td>Na&lt;sub&gt;2&lt;/sub&gt;O</td>
<td>4.04</td>
<td>3.80±0.09</td>
<td>3.82±0.12</td>
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<tr>
<td>K&lt;sub&gt;2&lt;/sub&gt;O</td>
<td>5.34</td>
<td>5.14±0.07</td>
<td>5.01±0.13</td>
</tr>
<tr>
<td>Total</td>
<td>99.28</td>
<td>99.24</td>
<td>99.15</td>
</tr>
<tr>
<td>Mo</td>
<td>0.189±0.003</td>
<td>0.030±0.001</td>
<td>0.199±0.004</td>
</tr>
<tr>
<td>W</td>
<td>0.030±0.001</td>
<td>0.199±0.004</td>
<td></td>
</tr>
<tr>
<td>H&lt;sub&gt;2&lt;/sub&gt;O</td>
<td>0.01</td>
<td>0.01</td>
<td>0.01</td>
</tr>
</tbody>
</table>

<sup>a</sup> The target composition: East porphyry of the Chuquicamata complex, Chile (Ballard, 2001), normalized to 100%.

<sup>b</sup> Composition of initial anhydrous granitic glasses, with the major element contents from electron microprobe, Mo and W concentrations from LA-ICP-MS, and H<sub>2</sub>O contents from FTIR. Standard deviations are at 1σ level.

<sup>c</sup> Composition of experimental products, with the major element contents from LA-ICP-MS (normalized to 100%) and H<sub>2</sub>O contents from FTIR. Standard deviations are at 1σ level.
TABLE 2. Summary of conditions and results of Mo and W diffusion experiments

<table>
<thead>
<tr>
<th>Run No.</th>
<th>Type</th>
<th>T (°C)</th>
<th>t (h)</th>
<th>$D^c$ ($10^{-12}$ m²/s)</th>
<th>$C_0^d$ (ppm)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Mo</td>
<td>W</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>initial</td>
<td>final</td>
</tr>
<tr>
<td>Anhy-Gran-1</td>
<td>Diff-cp</td>
<td>1600 ± 20</td>
<td>4</td>
<td>0.572 ± 0.010</td>
<td>0.536 ± 0.015</td>
</tr>
<tr>
<td>Anhy-Gran-2</td>
<td>Diff-cp</td>
<td>1500 ± 20</td>
<td>22</td>
<td>0.119 ± 0.004</td>
<td>0.101 ± 0.004</td>
</tr>
<tr>
<td>Anhy-Gran-3</td>
<td>Diff-cp</td>
<td>1400 ± 15</td>
<td>93</td>
<td>0.029 ± 0.001</td>
<td>0.026 ± 0.001</td>
</tr>
<tr>
<td>Anhy-Gran-4</td>
<td>Mo-sat</td>
<td>1600 ± 20</td>
<td>4</td>
<td>0.65 ± 0.01</td>
<td></td>
</tr>
<tr>
<td>Hydr-Gran-1</td>
<td>Diff-cp</td>
<td>1600 ± 20</td>
<td>1</td>
<td>8.79 ± 0.36</td>
<td>7.00 ± 0.34</td>
</tr>
<tr>
<td>Hydr-Gran-2</td>
<td>Mo-sat</td>
<td>1600 ± 20</td>
<td>1</td>
<td>5.97 ± 0.13</td>
<td></td>
</tr>
<tr>
<td>Hydr-Gran-3</td>
<td>Mo-sat</td>
<td>1500 ± 20</td>
<td>3</td>
<td>2.47 ± 0.14</td>
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<tr>
<td>Hydr-Gran-4</td>
<td>Mo-sat</td>
<td>1400 ± 15</td>
<td>7</td>
<td>1.18 ± 0.05</td>
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<tr>
<td>Hydr-Gran-5</td>
<td>Mo-sat</td>
<td>1400 ± 15</td>
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<td>8.99 ± 0.42</td>
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<tr>
<td>Hydr-Gran-6</td>
<td>Mo-sat</td>
<td>1300 ± 15</td>
<td>2</td>
<td>4.69 ± 0.19</td>
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<tr>
<td>Hydr-Gran-7</td>
<td>Mo-sat</td>
<td>1200 ± 15</td>
<td>3</td>
<td>3.40 ± 0.12</td>
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</tr>
<tr>
<td>Hydr-Gran-8</td>
<td>Mo-sat</td>
<td>1100 ± 15</td>
<td>11</td>
<td>1.00 ± 0.05</td>
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<tr>
<td>Hydr-Gran-9</td>
<td>Mo-sat</td>
<td>1000 ± 15</td>
<td>43</td>
<td>0.052 ± 0.001</td>
<td></td>
</tr>
</tbody>
</table>

$^a$ “Diff-cp” and “Mo-sat” represent experiment approach of diffusion couple and Mo saturation, respectively.

$^b$ These two runs show notable water loss.

$^c$ Diffusivities (with errors at 1σ level) extracted from fitting diffusion couple or Mo saturation profiles.

$^d$ Mo concentrations (with errors at 1σ level) at the interface in Mo saturation experiments, obtained from fitting.
Figure 1
Figure 2
Figure 3

(a) Anhy-Gran-1
- $T = 1600^\circ C$
- $P = 1$ GPa
- $t = 4$ h
- $C_{H_2O} = 0.04$ wt%

- $D_{Mo} = 5.72 \times 10^{-13}$ m$^2$/s
- $D_{W} = 5.36 \times 10^{-13}$ m$^2$/s
- $R^2 = 0.9997$

(b) Anhy-Gran-2
- $T = 1500^\circ C$
- $P = 1$ GPa
- $t = 22$ h
- $C_{H_2O} = 0.10$ wt%

- $D_{Mo} = 1.19 \times 10^{-13}$ m$^2$/s
- $D_{W} = 1.01 \times 10^{-13}$ m$^2$/s
- $R^2 = 0.9996$

(c) Anhy-Gran-3
- $T = 1400^\circ C$
- $P = 1$ GPa
- $t = 93$ h
- $C_{H_2O} = 0.06$ wt%

- $D_{Mo} = 2.90 \times 10^{-14}$ m$^2$/s
- $D_{W} = 2.55 \times 10^{-14}$ m$^2$/s
- $R^2 = 0.9995$

(d) Hydr-Gran-1
- $T = 1600^\circ C$
- $P = 1$ GPa
- $t = 1$ h
- $C_{H_2O} = 3.92$ wt%

- $D_{Mo} = 8.79 \times 10^{-12}$ m$^2$/s
- $D_{W} = 7.00 \times 10^{-12}$ m$^2$/s
- $R^2 = 0.9970$
Figure 4
Figure 5
Figure 6
Figure 7

(a) Log D (m² s⁻¹) vs. water content (wt%) at T = 1400 °C.

(b) Log D₀ (m² s⁻¹) vs. water content (wt%).

(c) Ea (kJ/mol) vs. water content (wt%).
Figure 9