Diffusive processes in aqueous glass dissolution.

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High level nuclear waste is often immobilised in a borosilicate glass for disposal. However, this glass corrodes in contact with aqueous solutions. To predict radionuclide releases from wasteforms, their dissolution mechanisms must be understood. Understanding glass dissolution mechanisms presents a challenge across numerous other disciplines and many glass dissolution models still remain conflicted. Here we show that diffusion was a significant process during the later stages of dissolution of a simplified waste glass but was not evidenced during the initial stages of dissolution. The absence of measurable isotopic fractionation in solution initially supports models of congruent dissolution. However, the solution becoming isotopically lighter at later times evidences diffusive isotopic fractionation and opposes models that exclude diffusive transport as a significant mechanism. The periodically sampled isotopic methodologies outlined here provide an additional dimension with which to understand glass dissolution mechanisms beyond the usual measurement of solution concentrations and, post-process, nano-scale analysis of the altered glass.

Introduction

The safe disposal of vitrified nuclear waste arising from the reprocessing of spent nuclear fuel requires that dissolution rates within geological disposal facilities are accurately modelled and aqueous dissolution mechanisms are well understood1. In recent years, a model of silicate mineral weathering, herein referred to as the ‘interfacial dissolution-precipitation model’, has been applied to silicate glasses2–4. This model challenged not only the structure of the alteration layers formed as predicted by established models of glass dissolution5,6, but also the long-held hypothesis that interdiffusion (hydration of the glass and ion-exchange reactions between protonated water species and glass species) is a rate-limiting mechanism during the initial moments of dissolution and again at high reaction progress7. Considering a timespan of at least one million years8, the rates of residual glass alteration would be different if dissolution was congruent through network hydrolysis from a supersaturated interfacial film of water, hypothesised by the interfacial dissolution-precipitation model8,9, or if dissolution was incongruent as predicted by diffusion-based models; some of which hypothesise interdiffusion reactions and the precipitation of secondary phases control the residual rate of glass dissolution1,7,9. In this case, interdiffusion reactions across a highly passivating, hydrated, altered glass interphase would be rate limiting.

In the weathering of natural systems, Li potentially fractionates through two isotope effects: kinetic isotopic effects during dissolution of the primary phase or equilibrium isotope effects where Li is incorporated into or adsorbed onto secondary phases10–12. Kinetic isotope effects are associated with rapid, unidirectional processes wherein the lighter isotope is preferentially transported across a phase boundary (such as in diffusion) or through a phase (such as in evaporation) due to its higher velocity13. Whilst the large mass difference between 6Li and 7Li (approximately 15 %) should be very sensitive to kinetic isotope effects, no such effects have been observed during the dissolution of the primary phase during basaltic glass dissolution11,14–16. Equilibrium isotope effects occur due to differences in isotopic masses effecting differences in bond energy preferences. Molecules containing heavier isotopes will have a higher reduced mass, a lower frequency as defined by Hooke’s law, a lower zero-point energy and so are more stable than molecules containing lighter isotopes13. Consequently, the heavier isotope is preferentially partitioned into higher bond energy sites17. For Li, longer bond lengths, less stiff bonds and lower bond energies are associated with higher coordination number sites10,13. As such, when there is precipitation, 6Li is preferentially partitioned into the higher CN octahedral and pseudo-hexagonal sites (CN of six) of clay secondary phases, 7Li preferentially remains in solution (CN of four)10,18,19 and the solution becomes isotopically heavier with time20–24. Further, Li can be adsorbed into the interlayer sites of smectites or onto the adsorption sites of gibbsite and ferrihydrite, although the former is not expected to be associated with any isotopic fractionation10,14,18.

Here, isotope fingerprinting techniques25 have been applied to glass leachates to probe whether the dissolution of a simplified simulant nuclear waste glass was rate-limited by diffusion. A seven-oxide borosilicate simplified glass analogue, 6Li-Mg-EM, was synthesised based on an existing Li-free analogue of a complex simulant Magnox (Mg-Al alloy fuel cladding) waste glass of 25 wt.% waste loading26. 6Li-Mg-EM was synthesised with a measured δ7Li of ‑916.5 ± 0.3 ‰ (7Li/6Li ratio of 1.029 ± 0.004).

Glass powders were leached at 90.0 ± 0.1 °C in deionised water, with leachate aliquots being taken in triplicate at intervals of 6 hours, 12 hours, 7 days, 28 days and 98 days. After the 28-day aliquots were taken, the leachant in one of the leaching vessels was replaced with fresh deionised water before the vessel was placed back into the heaters for a further 70 days. The pH of the experiments was allowed to vary freely. Leachate aliquots underwent matrix purification and then Li isotope analysis to determine how the isotopic composition of the solutions evolved with time relative to the pristine glass.

Results

Initially congruent dissolution

## After 6 hours of dissolution, significant amounts of Li were leached into solution (Fig. 1a), the alteration layers were estimated to be 0.3 µm thick (from boron release) and an estimated 1.6 % of the glass was altered (Fig. 2). However, the release of Li did not correlate with any measurable isotopic fractionation (Fig. 1b), with the leachate’s isotopic composition being within error of the pristine glass isotopic composition. If the nature of dissolution were incongruent dominantly through interdiffusion reactions during this initial regime, the leachates would be isotopically lighter than the pristine glass (δ7Li would decrease) due to kinetically enhanced 6Li diffusion through the hydrated glass layer. This demonstrated that the initial stage of dissolution was congruent.

## Secondary phase precipitation

Between 6 hours and 7 days, the average estimated alteration layer thickness grew from 0.3 to 3.1 µm and the estimated amount of glass altered rose from 1.6 to 15.4 % (Fig. 2). The leachates becoming isotopically heavier with time (increase of the δ7Li) during the initial rate regime is consistent with congruent dissolution alongside the dominant removal of Li from solution. During this time, the Li to B and Li to Na ratios in solution decreased (Fig. 3a and Fig. 3c respectively). Neither B nor Na are completely retained by the leachant. Although B is known to adsorb onto the surface of clays with a strong pH dependency27, significant masses of B are not expected to co-precipitate from solution during glass dissolution28. However, some Na is also expected to be incorporated into a sodium silicate hydrate phase29,30. The combined decreasing Li to Na and Li to B ratios with time are consistent with a Li-bearing phase being precipitated from solution onto the surface of the glass. This phenomenon was confirmed through static 6Li-1H cross polarisation nuclear magnetic resonance (CP-NMR) experiments on a glass sample of the same elemental composition leached under similar conditions30,31. This technique transfers proton nuclear magnetic polarisation via a through space coupling to adjacent 6Li atoms for NMR detection at the 6Li Larmor frequency and so it selectively probes Li in the alteration layers or secondary phases. A small but measurable signal was observed from the protonated alteration layers and demonstrated that Li was being incorporated into a secondary phase or the alteration layers after just 7 days (Fig. 4). This was consistent with the observation of a Li-bearing clay secondary phase layer forming on the surface of an immobilised low-activity waste glass during long-term dissolution in a pressurised, undersaturated environment at a high glass surface area to leachant volume ratio32.

The preferential incorporation of 6Li into these secondary phases resulted in the leachates becoming significantly enriched in 7Li with time between 6 hours and 7 days through an equilibrium isotope effect (Fig. 1b)14,15. The increasing δ7Li in solution, preceded by no measurable isotopic fractionation, suggested dissolution continued to be congruent during this initial dissolution regime and no measurable diffusive isotopic fractionation was present to offset this increase. Alternatively, diffusive isotopic fraction could have been masked by secondary phase precipitation, although this was considered unlikely due to the initial absence of diffusive isotopic fractionation. During this time, the Li to Mg ratio in solution increased (Fig. 3b), suggesting that Li is incorporated into secondary phases at a much slower rate than Mg. Based upon observations of a macroporous Mg-silicate layer wholly covering the surfaces of glass particles of the same elemental composition (not enriched in 6Li) after dissolution in a similar study under similar conditions to those used herein30, it could be expected that Li is incorporated into such a secondary phase; either through Li substituting for Mg into this secondary phase due to their similar ionic radii or due to this secondary phase inherently containing significant fractions of Li by being the amorphous precursor to a Li-containing clay such as a hectorite33.

Diffusive processes

A different process began operating in the later stage of dissolution. After 7 days, the δ7Li in solution had reached a maximum and the solution then started to become enriched in 6Li with time (Fig. 1b). If dissolution were congruent and secondary phase precipitation continued, then only equilibrium isotope effects would be occurring and the solution would continue to be enriched in 7Li with time. In such dissolution models of congruent glass dissolution which exclude diffusion as a rate-limiting mechanism, only the dissolution of the already formed 6Li-rich secondary phases or dissolution of the pristine glass unaccompanied by further equilibrium isotope effects through secondary phase precipitation could make the solution isotopically lighter with time. As these secondary phases were formed from the leachant and no notable changes in the leaching vessel environment took place between 7 and 28 days, it seems implausible that these secondary phases would spontaneously dissolve without an external driving force. Additionally, although the Li to Na and Li to B ratios (Fig. 3) decreased at a slower rate during this period compared to the interval between 6 hours and 7 days, this observed decreasing trend suggested that the removal of Li from solution continued during the later dissolution regimes. The continued removal of Li from solution was confirmed through 6Li-1H CP-NMR of a sample leached for 112 days at 90 °C in a similar study31, which showed Li continued to be incorporated into secondary phases between 7 and 112 days (Fig. 4). Therefore, the enrichment of 6Li in solution was attributed to kinetic isotopic fractionation due to the enhanced diffusive transport of 6Li out of the glass over 7Li.

## The magnitude of kinetic isotopic fractionation due to the diffusion of Li was large enough to entirely offset the heavy equilibrium isotopic fractionation due to the continued precipitation of secondary phases and enrich the solution in 6Li with time. This is despite a slowed rate of glass dissolution as shown by the Li concentrations in solution (Fig. 1a). The evolution of the leachates back towards the bulk isotopic composition, coupled with only a small change in Li concentration between 7 and 28 days (also 98 days in the non-refreshed experiment) is well illustrated using an isotope-concentration cross plot (Fig. 5). During this time the average estimated alteration layer thickness and estimated amount of glass altered grew from 3.1 µm and 15.4 % to 4.9 µm and 23.8 % respectively (Fig. 2).

During the extended leaching for an additional 70 days, the leachates from the ‘refreshed leachant experiment’ had lower Li concentration than the levels observed after 7 days of initial dissolution of the pristine glass (Fig. 1a and Fig. 5). This demonstrates that a passivating altered layer component was formed after 28 days which was able to retain its passivating nature after being significantly disturbed through leachant renewal. As such, this observation is consistent with other dissolution experiments involving solution renewal34. According to this study, this passivating component is expected to form between 7 and 28 days, as is consistent with the Si concentration in solution reaching an apparent steady-state value during this time (Supplementary Figure 1). In this time, the estimated alteration layer thickness in the refreshed leachant experiment grew significantly by 2.8 µm from 4.8 to 7.6 µm, compared to an increase of 0.3 µm from 5.0 to 5.3 µm in the non-refreshed leachant experiment (Fig. 2a).

Discussion

Fig. 6 displays theoretical Li isotopic signatures of the leachates alongside theoretical cross-sections of leached glass after dissolution for the two conflicting models of dissolution. The diffusion-based model (Fig. 6a) assumes the hydrated glass is a relict glass structure denuded of mobile glass species and at long-duration is expected to contain the highly passivating, nanoporous alteration layer component. If dissolution strictly followed this model, diffusive isotopic fractionation would be visible after 6 hours of dissolution. At long-duration, the isotopic signature of the leachates could be expected to evolve following the observed trend due to the passivating alteration layer component inhibiting dissolution1,7. Contrastingly, the interfacial dissolution-precipitation model predicts consistently congruent dissolution with alteration layers which have precipitated from a supersaturated interfacial film of water (Fig. 6b). In this model, the isotopic signatures of the leachates would initially be within error of that of the pristine glass, but the leachates would continue to become isotopically heavier with time after 7 days as Li-bearing secondary phase continued to precipitate2,3.

The passivating altered layer component inhibited glass dissolution. The significant diffusive isotopic fractionation measured in solution coinciding with this layers’ formation demonstrated that glass and water species are forced to diffuse across this layer to and from the pristine glass interface; presumably through its tight nanoporous structure7,28. This process results in the lighter isotopes of glass species diffusing out of this layer faster due to their lower mass. That is, the leachates becoming enriched in 6Li with time after 7 days are consistent with models of glass dissolution which predict the residual rate of dissolution is controlled by diffusion across a highly passivating altered layer component and the consumption of this hydrated glass layer through hydrolysis fuelled by secondary phase precipitation7. In this view, whilst the glass alters at its residual rate, the inward diffusive transport of water species to the pristine glass interface presents a rate limiting mechanism and the isotopes of all glass species which outwardly diffuse through this layer leach from the glass incongruently. As such, this dissolution behaviour is consistent with the isotopic signatures which could be expected to be produced by a diffusion-based dissolution model at long-duration (Fig. 6a) and is in agreement with isotope tracing experiments performed on the simplified analogue glass composition ‘International Simple Glass’35,36.

Appearing to contradict the diffusive isotopic fractionation observed in the later dissolution regimes, the absence of isotopic fractionation at short duration was consistent with models of congruent glass dissolution (Fig. 6b). However, the high dissolution temperature and rapid leaching of species from the glass during the initial moments of dissolution might have resulted in interdiffusion only controlling the rate of alteration for mere seconds or minutes. Consequently, whilst it was shown that dissolution during the initial 6 hours was congruent or virtually congruent, it is not known whether this was preceded by a negligibly short period of incongruent dissolution through interdiffusion reactions insufficient to produce measurable diffusive isotopic fractionation.

The decreasing isotope ratios in solution after 7 days of dissolution provided evidence for diffusion partly controlling the residual rate of alteration. However, this experiment alone does not allow us to rank models of glass dissolution. Whilst the interfacial dissolution-precipitation model excluded interdiffusion reactions and the formation of a passivating relict hydrated glass structure as rate limiting mechanisms3,4, a later version of this model hypothesised a passivating altered layer component may precipitate behind a congruent dissolution front which could limit transport due to diffusion2. Therefore, despite challenging models which exclude diffusion and the formation of a passivating alteration layer as significant dissolution mechanisms at high reaction progress, this evidence for diffusion being a rate-limiting mechanism is insufficient to elucidate whether this diffusive isotopic fractionation is the result of interdiffusion reactions across a passivating densified hydrated glass layer, or if it is caused by rate-limiting diffusion across a passivating precipitated layer behind an interfacial thin-film of water and a congruent dissolution front. However, it should be noted that the techniques described here afforded clear demonstration of diffusive processes, which otherwise would be difficult to detect with leachate concentration measurements and nano-scale measurements of elemental profiles within the alteration layers3,9,28. As such, these techniques may be applied to investigate the isotopic signatures of other elements, such as B, in order to distinguish between these two conflicting dissolution models.

# Methods

Sample preparation

The 6Li-Mg-EM glass (of nominal mol% composition 3.24% Al2O3, 18.33% B2O3, 1.14% La2O3, 2.56% 6Li2O, 2.56% 7Li2O, 8.69% MgO, 5.11% Na2O and 58.37% SiO2) was prepared as a single batch from precursor powders which were dried at 90 °C: Al2O3 (99.99%), B2O3 (99.98%), La2O3 (99.999%), Li2CO3 (99.998%, assumed natural abundance of δ7Li), Li2CO3 (Euriso-top, 95.77% 6Li atoms), MgO (99.99%), anhydrous Na2B4O7 (99.95%) and SiO2 (99.9 %). These precursors were melted in a 90% Pt 10% Rh 200 ml crucible inside a laboratory chamber furnace (Lenton UAF 16/10) at a rate of 10 °C/min from room temperature to 750 °C, held for 3 hours, then heated to 1500 °C. The glass was poured onto room temperature graphite-coated steel plates then, upon cooling to room temperature, the glass was crushed to a fine powder with an agate mortar and pestle. This powder then underwent the same heating procedure again but without a 3 hour hold and was poured onto 250 °C graphite-coated steel plates then cooled for 2 hours. After which, the plates were no longer heated and the glass cooled for another 2 hours before being annealed at 550 °C for 2 hours.

The glass samples were crushed in an agate mortar and pestle, and sieved according the hand sieving procedure outlined in ASTM PCT-B C1285-141437 using 75 µm and 150 µm stainless steel test sieves. Visible contaminants and magnetic particles were removed, before the glass was washed in absolute ethanol: the glass powder was placed into a 60 ml PTFE beaker and forcefully squirted with ethanol in a circular motion to approximately 1.5 times the volume of the glass powder, swirled for 10 seconds, left to settle for 15 seconds and then the ethanol was decanted. This was repeated twice more but on the third time the ethanol was not decanted, but instead the beaker was placed into an ultrasonic bath for 2 minutes. The ethanol was subsequently decanted and the entire process was repeated until the decanted ethanol appeared clear or a minimum of four times in total. The washed glass was then dried in a box oven at 90 °C for over 16 hours before being checked again with a magnet.

Density measurements

The density of 6Li-Mg-EM was characterised using Archimedes’ Principle in triplicate using approximately 1 g pieces of glass with no visible defects. An analytical balance (Ohaus Pioneer) with a weigh-below hook and a length of wire with a beaker of room temperature DI were used in the measurements. The density of 6Li-Mg-EM was calculated to be 2.47 ± 0.03 gcm-3.

X-ray powder diffraction

Approximately 100 mg of washed and sized 6Li-Mg-EM was analysed for crystalline phases using an X-ray diffractometer (Bruker D8 Advance). A Cu K-α wavelength was used with a near sample aperture slit size of 0.600 mm. The powder was mounted onto a circular glass slide using acetone, which was mounted onto a PMMA sample holder. A range of 10.0000 ° to 60.0123 ° (2θ) was analysed in 978 steps of 0.0511 °. The sample was found to be amorphous (Supplementary Figure 2).

Laser diffraction analysis

A laser diffraction analyser (Malvern Instruments Mastersizer E) was used for particle size analysis. An enclosed cell with PTFE magnetic stirrer filled with absolute ethanol was placed in front of a 300 mm focal length lens. A beam length of 14.3 mm was used to analyse a distribution of 1.2 µm to 600 µm. Approximately 75 mg of sample was added to the cell (45000 particles) and the particle size distribution was measured. An obscuration of 0.125 % and residual less than 2.000% were targeted. A uniform particle size distribution was obtained for 6Li-Mg-EM with a median particle size of 132.75 µm (Supplementary Figure 3).

Pristine glass composition

Electron probe microanalysis (EPMA) was used to obtain the concentrations of all elements within the pristine glass except B and Li. Three pieces of 6Li-Mg-EM were mounted in a two-part epoxy resin and polished using successive diamond pastes down to 3 µm before being carbon coated. An electron microprobe (Cameca SX 100) was used with a beam diameter of 10 µm, acceleration voltage of 15 keV, and beam current of 20 nA. 12 spots were analysed per piece of glass, selected in lines of 6 points each. Standards of jade for Na, diopside for Si, periclase for Mg, corundrum for Al, and lanthanum hexaboride for La were used.

Laser ablation inductively coupled plasma mass spectrometry (LA-ICP-MS) was used to determine the B and Li concentrations within the pristine glass. NIST610 was analysed to calibrate the elemental sensitivity of the run, with NIST614, NIST612 and BCR2-G analysed to veryify this calibration. A laser ablation system (ESI NWR193UC) with 193 nm Excimer laser using NeF gas connected to an ICP-MS (Perkin-Elmer NexION 350D) was used to analyse the resin mounted glass samples. Five 50 by 50 µm squares were ablated on each of the three glass pieces. A fluence of 10 Jcm-2, integration time of 5000 ms, and dwell of 10 ms were used, taking 41 s to analyse each spot. Li and B concentrations were analysed assuming the glass had a 7Li/6Li ratio of 1.029 (equal to that measured), NIST610 had a 7Li/6Li ratio at natural abundance and the 11B/10B ratios of the glass and NIST610 were the same (Supplementary Table 1). To correct for analytical bias, concentrations were normalised to the SiO2 concentration measured using EPMA as an internal standard. All elements were well above their respective detection limits.

Dissolution experiments

A glass surface area to leachant volume ratio (SA/V) of 2000 m-1 was targeted with 4.00 ± 0.02 g of washed and sized 75 µm to 150 µm 6Li‑Mg‑EM powder and 42.5 ± 0.4 ml of type 1 deionised water (18.2 MΩ.cm at 25 °C) leachant; a V/m (leachant volume to glass mass ratio) of 10.0 would yield an SA/V of 2159 m-1 assuming a spherical particle diameter of 112.5 µm, and so more leachant was used. Static-batch dissolution experiments took place in duplicate at 90 °C with duplicate blanks. Four acid cleaned 60 ml PFA (Savillex) standard jars with fluid transfer port (3.175 mm inner diameter) lids (herein collectively referred to as leaching vessels) were sealed using a piece of acid cleaned PFA tubing (3.175 mm outer diameter) held in place with ferrule nuts. The experiment was set up following ASTM PCT Test Method B25 protocol. Al blocks were pre-heated to 90.0 ± 0.1 °C inside two dry block heaters (QBH2, Grant Instruments). The fluid transfer lid design afforded rapid access to the leachant by acid cleaned PFA tubes attached to the end of 100 µl pipette tips. At intervals of 0.25, 0.5, 7, 28 and 98 days, the seals were temporarily removed, PFA tubes were inserted and 100 µl aliquots in triplicate were taken from each leaching vessel. A fresh piece of acid-washed PFA tubing and pipette tip were used for each aliquot.

After the 28 day aliquots were taken, all four leaching vessels were weighed then one set of sample and blank vessels were quickly returned to the heaters. The other set had all their leachant replaced with approximately the same volume of fresh deionised water, then all four vessels were leached for a further 70 days. The glass was not dried before the leachant was refreshed. The SA/V stayed within 10% of the target 2000 m-1 throughout dissolution. The pH of the leachant before the experiment, the leachates after 28 days in the refreshed leachant experiment, and after a total leaching time of 98 days in all experiments were measured (Supplementary Figure 4). Measurements took place on a FE20 FiveEasy benchtop pH meter with Mettler Toledo LE438 pH electrode calibrated using 7.00 and 10.01 ± 0.01 pH NIST traceable buffer solutions.

Leachate concentration analysis

The aliquots in 1.5 ml vials were weighed on an analytical balance (Sartorius MC1 Analytic AC 210s). All dilution acids were double distilled in a Teflon sub-boiling still. 1.0 ml of 0.1 M HNO3 and 0.3 M HF was added to each vial, which were then mixed using a benchtop mixer (Fisher Scientific FB15013 Topmix) and inspected for precipitates before being weighed again. These masses were used to calculate a gravimetric dilution factor of approximately 10, depending upon the exact aliquot volume and degree of sample evaporation. Samples were diluted further using two 25 fold dilutions with 2 % HNO3 before undergoing a final 10 fold dilution with 0.1 M HNO3 and 0.3 M HF for a final dilution factor of approximately 62500. All samples and blanks were analysed for the constituent elements of 6Li-Mg-EM using ICP-MS (Thermo Fisher Scientific Element XR); with isotopes 6Li, 7Li, 10B, 11B, 25Mg and 139La being analysed at low resolution, and 7Li, 11B, Na, 25Mg, Al and 28Si analysed at medium resolution. A set of four multielement standards containing 0.0, 0.5, 5.0 and 50.0 ng/ml of B and Li, and 0.0, 0.05, 0.5 and 5.0 ng/ml of Al, La, Mg, Na and Si were analysed before and after the samples. All standards contained isotopes at natural abundance.

Samples were analysed in sets of seven, with each set of samples and set of four multi-element standards being bracketed by two multi-element standard measurements of 5.0 ng/ml of B and Li, and 0.5 ng/ml of all other elements. The first set of four multi-element standards was used to create calibration curves. All blanks contained less than 0.05 ng/ml of Li and B, 0.5 ng/ml Al, 1.5 ng/mlSi excluding 2 outliers, 0.15 ng/ml Na and 0.75 ng/ml Mg. Sample concentrations were corrected for background concentrations, drift corrected, normalised to their first bracketing standard concentration and then corrected for the dilution factor used. The high background associated with Si meant that the 0.05 and 0.5 ng/ml standards were below detection limits. Therefore, although many of the Si readings were above the detection limit, these values were considered unreliable as they were created from a single-point calibration. The concentrations of Li, B Na and Mg in the leachates were well above their respective detection limits, whilst the concentrations of La and Al were not.

Leachate concentration data treatment

The corrected concentrations of element *i* in solution, *CiSample*¸ were then used with the initial leachant volume, *Vi*, the mass of glass powder leached, *m*, and the elemental mass fraction of *i* within the glass, *fi*, to calculate the estimated fraction of *i* leached from the glass, *LFi*, using Equation 1.

From the leached fraction of B, an average alteration layer thickness, *t,* could be calculated using Equation 2. This equation was based on the shrinking core model assuming pristine spherical glass particles of median radius (*r0* = 56.25 µm) were uniformly altered38.

Alkali fusion and HF digestion

Four sets of 50 mg of pristine washed and sized 6Li-Mg-EM powders underwent alkali fusion using 250 mg of K2CO3 following a fusion methodology originally designed for natural samples39. These fusion supernatants were then analysed via ICP-OES (Agilent 5100) and Li recoveries for the alkali fusion process alone were found to be between 84% and 95%. These supernatants were then diluted 100 times using 2% HNO3 before being prepared for column chromatography.

The accuracy of the pristine glass isotopic composition obtained through the alkali fusions was verified through HF digestion: 15.0 mg of washed and sized 6Li-Mg-EM powder was digested in 0.7 ml single distilled HF and 0.5 ml double distilled HNO3 at 150 °C for 24 hours. In this time, the glass was noted to have completely dissolved. The digest was then dried down, before being taken up in 1 ml double distilled aqua regia at 150 °C for 24 hours then dried down again. This procedure was then repeated twice more using double distilled 6 N HCl, but on the final time was not dried down.

Column chromatography

The samples were subjected to cation exchange column chromatography to prepare purified mono-elemental Li solutions through matrix separation for MC-ICP-MS analysis. All HCl used for dilutions and column chromatography had been double distilled using a Teflon sub-boiling still and calibrated at 0.7 N using titrations with 1.0 M of NaOH (Fisher Scientific, certified grade). Twelve 3 ml Teflon high aspect ratio ion exchange columns, with a length of 250 mm and an internal diameter of 4 mm, containing AG MP-50 macroporous cation exchange resin were used. For the leachates and alkali fusions, sample volumes containing 300 ng Mg (leachates) or 4 ng Li (alkali fusions) were dried down in PFA vials at 80 °C, refluxed in double distilled concentrated HNO3, dried down again and taken up in 200 µl of 0.7 N HCl for loading on the column. The HF digest in 6 N HCl was instead diluted using 0.7 N HCl to a nominal Li concentration of 20 ng/ml, before 200 µl (4 ng Li) was loaded onto the columns in duplicate. The methodology used for column chromatography did not differ from that described in the original methodology for the leachates25, but a 15 ml Li cut (2 ml more than the original methodology, with the initial matrix elution being 2 ml less) was used for the alkali fusions and HF digests. After the column elutions with 0.7 N HCl, the methodology of which is described in great detail in the original publication25, the Li fractions were again dried down, refluxed in concentrated HNO3 and after drying down once more were taken up in 0.5 ml (leachates) or 1.0 ml (alkali fusions and digests) 2% HNO3 for isotopic analysis.

δ7Li analyses

For the leachates, the volumes of the Li fractions and 2% HNO3 required to yield 9 ng/ml of Li were dispensed into acid cleaned 2 ml PFA ICP vials. δ7Li analysis took place on a multicollector ICP-MS (MC-ICP-MS, Thermo Scientific Neptune Plus) at low resolution. Samples were introduced using a quartz Apex-IR (ESI) system with 140 °C spray chamber with 2 °C Peltier coil. A Savillex PFA 100 µl/min nebuliser with Ni-Jet sample and Ni-X type skimmer cones were used. Cups L4 and H4 were used with 1011Ω pre-amplifiers to measure 6Li and 7Li respectively. Measurements were composed one block of 30 cycles with an integration time of 8.4 s, and were taken with respect to and bracketed by NIST SRM 8545 L-SVEC40 at a concentration of 9 ng/ml. Analyte measurements were corrected for instrumental mass bias and drift during analysis using a standard-sample bracketing technique with the L-SVEC standard10,25,41. Li7-N42, also at a concentration of 9 ng/ml, was analysed at the beginning and end of each set of nine samples and L-SVEC standard measurements as a secondary standard. A blank (2 % HNO3) was measured after each standard or sample measurement, with typical blank concentrations of approximately 0.015 ng/ml. Each sample was measured in duplicate, with samples being rerun again throughout the analysis to confirm reproducibility.

For the alkali fusions, the same instrument, Apex-IR (ESI) nebuliser and cones were used as for the leachates. However, 7Li and 6Li were instead measured using 1013 Ω amplifiers. Samples were diluted to a target concentration of 0.7 ng/ml using 2 % HNO3 and analysed alongside NIST SRM 8545 L-SVEC at the same concentration to bracket each sample measurement, with each sample analysed in triplicate. A 5 ng/ml Ca wash was used between each measurement to minimise the Li background, with typical blank backgrounds of approximately 0.007 ng/ml; two orders of magnitude lower than those of the standards and samples. Li6-N and Li7-N standards were again analysed at the beginning of the run to assess external reproducibility at a concentration of 0.7 ng/ml42. The HF digests were analysed following the same procedure as the alkali fusions, except each Li fraction was analysed in duplicate at 0.56 ng/ml rather than in triplicate at 0.7 ng/ml.

The blank corrected sample 7Li/6Li were converted to δ7Li (‰) with respect to the average of the two blank corrected bracketing L-SVEC standard 7Li/6Li measurements using Equation 3:

A δ7Li for each sample was then calculated as the average δ7Li of the duplicate or triplicate sample measurements and interpolated δ7Li values calculated between each pair of blank corrected sample measurements and the blank corrected L-SVEC standard 7Li/6Li measured between them. Errors (2σ) were calculated as twice the standard deviation of the blank-corrected duplicate sample measurements and the interpolated-blanks δ7Li value measured between them. An average δ7Li of -916.5 ± 0.3 ‰ was measured for the four full-procedure replicate K2CO3 fusions of the pristine 6Li-Mg-EM glass. The δ7Li of the digest was found to be in excellent agreement with that of K2CO3 fusions at -916.4 ± 0.1 ‰ and confirmed the accuracy of the K2CO3 methodology39.

6Li-1H cross polarisation nuclear magnetic resonance.

A sample of the same target composition as 6Li-Mg-EM, without isotopic enrichment which had been leached in a separate study, underwent 6Li-1H cross polarisation nuclear magnetic resonance (CP-NMR) analysis31. This sample had been leached for 7 days in deionised water at 90 °C at a SA/V of 2000 m-1. However, experiments were ended entirely rather than just sampled at each sampling interval. A pristine sample was also analysed to demonstrate there was no measurable signal (polarisation) transferred from ambient proton absorption on to the glass. Approximately 130 mg of sample was loaded into a PTFE liner inside a 7.5 mm zirconia rotor, which was placed into a 7.5 mm double resonance MAS probe but not spun. A spectrometer frequency of 499.72 MHz for 1H and 73.54 MHz for 6Li were used. A π/2 pulse (8.0 µs) was found for 1H using RTV silicone and a pulse delay of 3 s was used. Powdered LiOH was used to optimise matching conditions and contact time. Spectra were collected with between 75000 and 250000 repetitions under the conditions above and were referenced to 6Li in a 1.0 M 6LiCl (95.77% 6Li atoms) solution at 0.0 ppm.

Data availability

The data that support the findings of this study are available in Apollo, the University of Cambridge institutional repository with the identifier doi:10.17863/CAM.45721 (ref. 43).

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Competing interests

The authors declare no competing financial or non-financial interests.

# Author Contributions

T.G. fabricated the glass, began the dissolution experiments, sampled the aliquots, alkali fused the glass, HF digested the glass, prepared samples for concentration and isotope analyses, assisted in analysing the leachates, interpreted the results and wrote the manuscript. S.M. supervised the isotope work, analysed the leachates and assisted in interpreting the results. E.T. arranged and supervised the alkali fusion, analysed the digested samples and supervised their digestion and assisted in interpreting the results. M.B. analysed the alkali fused glass, digested the glass in the first instance, supervised and assisted in column chromatography, and assisted in interpreting the results. R.G. assisted in acquiring the NMR spectrum and interpreting the results. I.F. conceived the experiments, supervised the study and assisted in interpreting the results.

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Figure legends

Figure 1 | Temporal evolution of the Li concentration and the Li isotopic composition of the leachates. The mass spectrometric (ICP-MS) results display the total concentration of Li in the leachates (background, drift and dilution factor corrected) with one s.d. errors (a) and the MC-ICP-MS results show the Li isotopic composition (δ7Li) of the same leachates and the pristine glass with two s.d. errors (b). In (a), the solid and dotted lines and markers represent the mean values. In (b), individual analyses are plotted for each of the triplicate leachates with the mean values indicated by the solid and dotted lines. The blue solid line with diamond markers and the orange dotted line with square markers in both figures represent the two datasets of the duplicated experiments. The open diamond markers represent the same experiment as the blue filled diamond markers in which the leachant was refreshed in one vessel after the sampling of the 28 day aliquots followed by a further 70 days of leaching in fresh DI water (98 days total dissolution). The central black dashed line in (b) denotes the measured pristine Li isotopic composition of 6Li-Mg-EM, with the lines above and below this denoting the two s.d. error on this measurement.

Figure 2 | Temporal evolution of the estimated thickness of the alteration layers and amount of glass altered. The average estimated thickness of the alteration layers (a) and estimated amount of glass altered (b) from the total concentrations of B leached into solution. Points are plotted as the average of triplicate leachate measurements with errors calculated as one s.d. of each set of triplicate aliquot measurements taken from each vessel at each time. The blue solid line with diamond markers and the orange dotted line with square markers in both figures represent the two datasets of the duplicated experiments. The open diamond markers represent the same experiment as the filled blue diamond markers in which the leachant was refreshed in one vessel after the sampling of the 28 day aliquots followed by a further 70 days of leaching in fresh DI water (98 days total dissolution).

Figure 3 | Temporal evolution of select elemental ratios of the leachates. The Li/Na (a), Li/Mg (b), Li/B (c) ratios in the leachates, as calculated from their respective concentrations in solution in µg/ml measured using ICP-MS. Each data point represents the average of the triplicate leachate measurements with errors given to one s.d. The blue solid line with diamond markers and the orange dotted line with square markers in all figures represent the two datasets of the duplicated experiments. The open diamond markers indicate the same experiment as the filled diamond markers in which in which the leachant was refreshed in one vessel after the 28 day aliquots were taken, before the glass was leached for further 70 days (98 days total dissolution).

Figure 4 | Static 6Li-1H CP-NMR spectra of a sample leached for 7 days at 90 °C (Red) and a sample leached for 112 days at 90 °C (Purple) in static batch dissolution experiments31 compared with the spectrum of the pristine glass (Cyan). The glass composition was the same as the glass used in the isotopic study but was not enriched in 6Li31. Spectra were normalised to the mass of sample analysed and the number of repetitions.

Figure 5 | Evolution of the leachate Li isotopic composition as a function of Li concentration change. Cross plot of the average Li isotopic composition of the leachates (δ7Li) measured using MC-ICP-MS for each triplicate (also given in Figure 1b) against the total Li concentration in solution (also given in Figure 1a). Errors are given to two s.d. on the isotopic data and one s.d. on the concentration data, as calculated from the triplicate aliquot measurements. The blue line with diamond markers and the orange line with square markers represent the two datasets of the duplicated experiments. The open diamond marker indicates the same experiment as the filled diamond markers in which the leachant was refreshed in one vessel after the 28-day aliquots were taken, before the glass was leached for further 70 days (98 days total dissolution). The central dashed black line indicates the pristine glass isotopic composition with the lines above and below this representing the two s.d. error on this measurement, as calculated from the triplicate MC-ICP-MS measurements of the four full-procedure alkali fusion replicates of the pristine glass.

Figure 6 | Theoretical plots of the Li isotopic compositions of the leachates against time accompanied by theoretical elemental profiles within cross sections of leached glass at high reaction progress (approximately 28 days here). The theoretical structures of alteration layers and the cartoons of the evolution of the Li isotopic composition of the leachates according to a diffusion-based model (a) and the interfacial dissolution-precipitation model (b) are given1–3,7. The cartoons of the isotopic evolution of the leachates in (a) assumes the isotopic composition of the leachate at 6 hours is purely the result of kinetic isotopic fractionation from diffusion (pristine glass isotopic composition multiplied by the square root of the ratio of the masses of 6Li and 7Li), after which time the leachates evolve following the observed profile. The cartoon of the evolution of the isotopic signature of the leachate in (b) follows the observed profile until 7 d, whereafter it assumes secondary phase precipitation continues in the absence of diffusive isotopic fractionation at a rate depending upon the rate that Li is released into solution; in doing so, it is assumed the fraction of leached Li incorporated into the secondary phases and the composition of the secondary phases is constant throughout dissolution.