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and atomistic simulations, we correlate the dissolution rates-ranging over 4 orders of magnitude-of a selection of silicate glasses and crystals to the number of chemical topological constraints acting between the atoms of the dissolving solid. The number of such constraints serves as an indicator of the effective activation energy, which arises from steric effects, and prevents the network from reorganizing locally to accommodate intermediate units forming over the course of the dissolution.

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Topological Control on Silicates Dissolution Kinetics

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Like many others, silicate solids dissolve when placed in contact with water. In a given aqueous environment, the dissolution rate depends highly on the composition and the structure of the solid, and can span several orders of magnitude. Although the kinetics of dissolution depends on the complexities of both the dissolving solid and the solvent, a clear understanding of which critical structural descriptors of the solid control its dissolution rate is lacking. Through pioneering dissolution experiments and atomistic simulations, we correlate the dissolution rates – ranging over four orders of magnitude – of a selection of silicate glasses and crystals, to the number of chemical topological constraints acting between the atoms of the dissolving solid. The number of such constraints serves as an indicator of the effective activation energy, which arises from steric effects, and prevents the network from reorganizing locally to accommodate intermediate units forming over the course of the dissolution.

I. INTRODUCTION

Due to their abundance in Earth’s crust, silicate minerals and glasses are the base components of numerous industrial materials including glasses, ceramics, and concrete. As such, understanding and predicting the kinetics of silicate dissolution in the presence of water has fundamental and industrial implications, with examples including bioactive glasses, nuclear waste confinement matrices, and the reaction of cement with water, the binding phase in concrete, all of which require accurate predictions of dissolution rates, in the context of microstructure evolution, phase stability and chemical durability.

Predicting the kinetics of dissolution of materials under given thermodynamic conditions is, a priori, a complex problem as it depends, amongst others, on the solvent chemistry and the geometry of the surface. Another difficulty arises from the fact that the dissolution of minerals can feature leaching, or incongruency, that is, occurring in a nonstoichiometric fashion. However, this latter feature typically is true only during the initial stages of contact with water, with steady-state dissolution usually showing congruency. Nevertheless, for a given solvent and thermodynamic condition, both the composition and structure of a material dictate its dissolution kinetic.

Mere knowledge of the composition of a material is insufficient to predict its dissolution rate as, e.g., glassy silica dissolves between one and three orders of magnitude faster than crystalline α-quartz depending on the solvent pH. While it may be argued that glasses should indeed be less stable that their crystalline equivalents, such a trend is not systematic as, e.g., albite glass and crystal show comparable dissolution rates. In fact, the dissolution of silicate glasses and minerals has recently been shown to occur by similar mechanisms and to be mostly controlled by short-range atomic order, rather than the long-range disorder. Indeed, the dissolution rates of silicate glasses are generally thought to decrease with the connectedness, or the degree of polymerization of the atomic network described by the ratio of terminating non-bridging oxygen (NBO) per tetrahedral unit (T) formed by silicon or aluminum atoms (NBO/T).

However, using such a metric is far too restrictive as, e.g., silica, albite, jadeite, and nepheline glasses all feature a fully polymerized network (NBO/T = 0), but show vastly differing dissolution kinetics. As such, given the disordered nature of glass, it is a challenge to identify a metric that is simple enough to enable practical predictions of dissolution rates, but that takes into account enough structural detail to provide realistic results.

By capturing the topology of atomic networks while filtering out chemical details that ultimately do not affect the macroscopic properties, topological constraint theory (TCT) is a promising tool that can elucidate such a metric. Within the TCT framework, atomic networks are described as mechanical trusses, in which the atoms experience topological constraints, as imposed by radial and angular chemical bonds. Hence, following Maxwell’s stability criterion, an atomic network is described as flexible, stressed-rigid, or isostatic, if the number of topological constraints per atom (n_c) is lower, higher, or equal to three, i.e., the number of degrees of freedom per atom in three dimensions (see Figure 1). In this paper, we establish that the dissolution rates of glassy and crystalline silicates in caustic environments are dictated by n_c, which serves as an indicator of the effective activation energy of the dissolution process, and, in turn, of the activation energies of solid state ion-diffusion and conduction. This suggests that, simply, the
mechanical stability of the bulk atomic network controls the dissolution behavior, and the transport properties of silicate solids.

II. RESULTS

A. Dissolution rates

The dissolution rates of glassy silica and α-quartz were measured under isothermal conditions, and at different temperatures using vertical scanning interferometry (VSI, see the Supplemental Material). This technique has been extensively applied to measure the dissolution rates of minerals of geological relevance. By directly tracking the evolution of the surface topography in time, with sub-nanometer vertical resolution, VSI accesses the true dissolution rate of a solid dissolving in a given solvent. Unlike dissolution assessments that are based on analysis of solution compositions, which may be affected due to aspects including metastable barrier formation, incongruency in dissolution or, ion adsorption, VSI analytics are not influenced by such complexities. Dissolution rates of the solids were quantified using a rain-drop dissolution method, wherein both the solution pH and composition (i.e., the under-saturation level with respect to the dissolving solids) are kept constant over the course of a given experiment (see Figure 2). In addition to our measurements, to assess the compositional dependence of dissolution kinetics, the dissolution rates of sodium trisilicate (Na2O–3SiO2), albite (NaAlSi3O8), jadeite (NaAlSi2O6), and nepheline (NaAlSi2O6) glasses were sourced from the tabulations of Hamilton et al. Note that, for consistent comparisons between solids of different compositions, all the dissolution rates presented herein are normalized in terms of moles of O2 dissolved per unit of surface and time.

B. Relationship between composition and dissolution rates

Figure 3a shows dissolution rates for a range of solids in basic aqueous solvents. As expected, all the dissolution rates increase with pH, due to the increasing abundance of HO− species available for nucleophilic attacks of Si or Al tetrahedra. Although the slope of the dissolution rates versus pH curve varies for each material, we note that the ranking of dissolution kinetics remains consistent across a wide pH range and scales as α-quartz < albite < jadeite < nepheline < glassy silica < sodium silicate.

The structural origin of the dissolution rates can be elucidated by comparing dissolution rates for a representative pH, in this case pH 12. First, as shown in Figure 3c, relying on the degree of depolymerization, NBO/T, is insufficient. For example, if sodium silicate (NBO/T = 0.67) dissolves faster than glassy silica (NBO/T = 0), this metric does not distinguish the structures of nepheline, jadeite, albite and α-quartz (NBO/T = 0) either, although their dissolution rates span three orders of magnitude. Second, one cannot rely on composition either. The ranking of the energies of Si–O (444 kJ/mol), Al–O (423 kJ/mol), and Na–O (83 kJ/mol) bonds suggests that the stability of a given silicate should increase with the fraction of silicon present, thereby decreasing the dissolution rate. Although the slope of the dissolution rates span three orders of magnitude. Second, one cannot rely on composition either. The ranking of the energies of Si–O (444 kJ/mol), Al–O (423 kJ/mol), and Na–O (83 kJ/mol) bonds suggests that the stability of a given silicate should increase with the fraction of silicon present, thereby decreasing the dissolution rate. Although, as shown in Figure 3a, this trend is true for nepheline, jadeite, and albite, it does not allow one to explain differences in dissolution rates between glassy and crystalline silica. In summary, none of the typical, and expected metrics are able to capture sufficient structural detail to permit quantitative estimation of the dissolution kinetics.

We now calculate the rigidity of the atomic networks of the different silicate solids to evaluate the relevance of this metric for dissolution. In a fully connected network, like, e.g., chalcogenide glasses, an atom of coordination number r experiences r/2 radial bond-stretching (BS) constraints, and 2r−3 angular bond-bending (BB)
Figure 3. Influence of composition and structure on dissolution rate. (a) Room-temperature dissolution rates ($K$) of sodium silicate,\cite{22} glassy silica, nepheline,\cite{13} jadeite,\cite{14} albite,\cite{15} and α-quartz (normalized with respect to the amount of O$_2$ dissolved) versus solvent pH. (b) Dissolution rates at pH = 12 with respect to the ratio of non-bridging oxygen and Si or Al tetrahedra (NBO/T). (c) Same data as b), plotted with respect to the ratio of Si and cations atoms ($Si/(Si+Al+Na)$). (d) Same data as b), plotted with respect to the number of constraints per atom $n_c$. The data are fitted by an Arrhenius-like law $K = K_0 \exp(-n_c E_0 RT)$, with $K_0 = 1.4 \times 10^{11}$ µmol/m$^2$/s and $E_0 = 25.5$ kJ/mol (dashed line). The inset shows the same data as d), in linear scale. The grey area indicates the flexible domain ($n_c < 3$).

This allows one to enumerate $n_c = 3.67$ constraints per atom for quartz (stressed-rigid). In glassy silica, the Si–O–Si angle typically shows a much broader angular excursion,\cite{22} so that this angle appears unconstrained. This results in $n_c = 3.00$ for glassy silica (isostatic), in agreement with the observation that silica is an excellent glass forming system.\cite{22} The insertion of sodium cations typically substantially decreases the glass transition temperature of the glass and, thereby, restores the Si–O–Si BB constraint. In addition, each Na cation depolymerizes the base silica network by creating one NBO. The number of constraints per atom of (Na$_2$O)$_x$(SiO$_2$)$_{1-x}$ can then be calculated\cite{22} as being $n_c = (11 - 10x)/3$. This is in agreement with experimental evidence\cite{22} of a rigidity transition ($n_c = 3$) at $x = 0.20$. Then, from a simplistic viewpoint, starting from sodium silicate, each added aluminum cation consumes one NBO. Na cations then act as charge compensators around Al$^{IV}$ cations\cite{22} and are, therefore, not considered any further in the constraints enumeration. Note that the tetrahedral angular environment of Al cations is not as well defined as that of Si, so that the O–Al–O angles are considered unconstrained.\cite{22} Consequently, the number of constraints per atom for (Na$_2$O)$_x$(Al$_2$O$_3$)$_x$(SiO$_2$)$_{1-2x}$ is given by $n_c = (11 - 10x)/3$. However, this enumeration under-estimates $n_c$ as a small fraction of over-coordinated three-fold oxygen tricluster and five-fold aluminum species are found.\cite{22,20} Hence, the rigidity of the aluminosilicate glasses was evaluated by a careful analysis of their structure, as predicted by the molecular dynamics (MD) simulations. This allows clear differentiation of intact from thermally broken BS (BB) constraints by computing the radial (angular) excursion of each neighbor, using a previously established methodology (see the Supplemental Material). As such, we find $n_c = 3.33$, 3.22, and 3.00 for albite, jadeite, and nepheline, respectively. Note that the latter value implies that nepheline features an isostatic network ($n_c = 3$), which is consistent with the fact that its dissolution rate is similar to that of glassy silica.

C. Topology controls the kinetics of dissolution

Figure 3d shows the dissolution rates $K$ as a function of the number of constraints per atom $n_c$. As shown in the inset, $K$ significantly increases in the flexible domain ($n_c < 3$), in agreement with the idea that, due to the presence
of internal low energy modes of deformation (i.e., floppy modes), a flexible network is less stable than its rigid counterparts, and should, therefore, dissolve faster. The observed evolution of $K$ with respect to the number of constraints $n_c$ suggests an Arrhenius-like relationship, of the form:

$$K = K_0 \exp \left( \frac{-n_c E_0}{RT} \right)$$

(1)

where $K_0$ is a rate constant that depends on the solution phase chemistry, and yields the barrier-less dissolution rate of a completely depolymerized material (i.e., for which $n_c = 0$) and $E_0 = 25.5$ kJ/mol is an energy barrier that needs to be overcome to break a unit atomic constraint. This suggests that:

$$E_{A}^{\text{eff}} = n_c E_0$$

(2)

serves as an effective activation energy that controls the kinetics of the dissolution. To validate this hypothesis, we evaluated the low-temperature activation energy ($E_A$) of dissolution of α-quartz and glassy silica by measuring their dissolution rates at 3.5, 25, and 45 °C at pH 13. We chose this higher pH to achieve appreciable dissolution of quartz at the lowest temperature. Note that these materials feature a different $n_c$ (3.67 and 3.00 for quartz and glassy silica, respectively) but similar composition, which allows us to isolate the singular effect of atomic topology on the activation energy, with no compositional effects. As illustrated in Figure 4, both materials show an Arrhenius-like evolution of their dissolution rate with temperature. As such, we find $E_A = 101±33$ kJ/mol and $52±20$ kJ/mol for α-quartz and glassy silica, respectively, in agreement with prior studies – an expected outcome since α-quartz shows a higher $n_c$ than glassy silica (i.e., on account of its stressed-rigid nature). Interestingly, as shown in Figure 4, these values are in fair agreement with the effective activation energy predicted simply from the knowledge of the number of constraints per atom ($E_{A}^{\text{eff}} = n_c E_0$), which yields to $E_{A}^{\text{eff}} = 93.6$ kJ/mol and 76.5 kJ/mol for α-quartz and glassy silica, respectively. This suggests the following atomistic picture: starting from $n_c = 0$, which would correspond to a fully depolymerized material, each new constraint per atom effectively reduce the dissolution kinetics by increasing the associated activation energy needed for bond rupture.

The activation energies for dissolution are compared to the activation energies of other thermally-activated processes, namely, for sodium silicate (self-diffusion and conduction of Na), potassium silicate (self-diffusion and conduction of K), fused silica (diffusion of water), and quartz (diffusion of water). First, we note that, interestingly, for a given material, the activation energies associated with dissolution, diffusion and conduction are consistent with each other, which suggests that these energy barriers all arise from a common atomistic origin – which we identify as the rigidity of the network. Second, the activation energies all increase with the number of constraints per atom and are similar to the values predicted by equation 2, that is, the effective activation energy induced by the number of constraints per atom.

III. DISCUSSION

The question remains regarding the origin of the relationship that is found herein between the activation energy $E_A$ and the number of topological constraints per atom $n_c$. As such, we propose that $n_c$ serves as an indicator of steric effects in the atomic network, which prevent reorganization and internal motion of the constituent species. The reaction mechanism of a silicate with water can take different forms: (1) hydration, where intact water molecules enter the glass network (2) hydrolysis, where water reacts with Si–O bonds to form hydroxyl groups, and (3) ion-exchange, where network modifying cations, e.g. Na⁺, are replaced by H⁺ cations. The three mechanisms can occur simultaneously, and influence process kinetics. For (1), the rate of water diffusion is primarily controlled by steric constraints, i.e., atomic packing, imposed by the network. However, in typical silicates, the holes in the structure, as defined by the ring statistics, are too small to allow direct diffusion of intact water molecules into the network. Therefore, water would preferentially penetrate through hydrolysis and condensation reactions. In mechanism (2), in basic solvents, Si tetrahedra are subjected to nucleophilic attack by HO⁻ species, thereby forming five-fold coordinated Si intermediate units. On the contrary, in acidic environments, inter-tetrahedral bridging oxygen atoms are subjected to electrophilic attack by H⁺ species, thereby forming three-fold coordinated O intermediate units. In these two cases, process kinetics depend on the ability of the network to accommodate such over-coordinated units by local rearrangement. For (3), the rate-limiting step is suggested to be the penetration of the glass surface by water. However, the three mechanisms are strongly coupled as the depolymerization of the silicate network due to (2) can open up the atomic network, e.g., due to ring opening thus enhancing water mobility for (1) and (3) to occur at an expedited rate.

In these three mechanisms, the kinetics of dissolution are limited by the ability of the atomic network to locally rearrange in order to: (a) accommodate intermediate over-coordinated atoms (SiV or O¹H) or (b) enable the creation of pathways, or channels, by enlarging preexisting holes, thereby allowing a penetrating species (water molecule or H⁺) to transfer between two sites. As such, the activation energy associated with such reorganizations takes the form of a strain energy that characterizes the ability of the network to locally resist, that is, to prevent the formation of energetic instabilities (departure from an equilibrium configuration) that are induced due to a dissolution process. Then, it becomes
clear that the development of a strain energy, and its follow on dissipation, is linked to the number of constraints per atom $n_c$, since each constraint acts as a little spring between the atoms. Therefore, $n_c$ characterizes the steric effect imposed by the architecture of the atomic network, which prevents the accommodation of defects (i.e., over-coordinated atom or enlarged hole to create a channel). This picture is in line with result from density functional theory, which have shown that the activation energy of the hydrolysis of the inter-tetrahedra bridging oxygen increases with the extent of the steric constraints imposed by the network, that is, its connectivity and therefore, rigidity[21].

IV. CONCLUSIONS

Our results demonstrate that the dissolution rate of silicate-based glassy and crystalline solids in caustic solutions is correlated to the number of topological constraints per atom present in the network. Such atomic constraints are indicative of the effective activation energy that needs to be overcome for dissolution to occur. Moreover, $n_c$ characterizes the ease with which the atomic network can reorganize to accommodate the intermediate (metastable) defects states that are formed over the course of the net dissolution process. The rate at which such reorganization can occur, appears to be the primary rate-controlling step during dissolution, that is, which controls its kinetics. This leads to the idea that dissolution kinetics, a response through to be controlled by surface properties, is paradoxically primarily controlled by the topology of the bulk architecture of the atomic network of the dissolving material. In addition, it is remarkable that a number of processes that share a common thermally activated origin, e.g., dissolution, diffusion and transport show similar trends in activation energy with respect to the number of topological constraints per atom. If, indeed, it is well known that diffusion and conductivity are related to each other via the Nernst-Einstein equation, it is likely that dissolution shares a similar basis, although perhaps better described by a Butler-Volmer type of formulation[22]. Nevertheless, our results suggest that dissolution, diffusion, and conduction could potentially be treated within the same atomistic, and thermodynamic framework – enabling a priori estimation of such phenomena. This has important implications for better understanding and modeling processes ranging from cement hydration, to the chemical durability of glasses used to encapsulate radioactive wastes, to rate controls on silicate mineral dissolution that accompanies ocean acidification[23].

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1 Julian R. Jones, “Review of bioactive glass: From Hench to hybrids,” *Acta Biomaterialia* 9, 4457-4486 (2013)
2 K.b. Harvey and C.d. Litke, “Model for Leaching Behavior of Aluminosilicate Glasses Developed as Matrices for Immobilizing High-Level Wastes,” *Journal of the American Ceramic Society* 67, 553–556 (1984)
3 Bernd Grambow, “Nuclear Waste Glasses - How Durable?” *Elements* 2, 357–364 (2006)
4 H. F. W. Taylor, *Cement Chemistry* (Thomas Telford, 1997).
5 Céline Cailleteau, Frédéric Angeli, François Devreux, Stéphane Gin, Jacques Jestin, Patrick Jollivet, and Olivier Spalla, “Insight into silicate-glass corrosion mechanisms,” *Nature Materials* 7, 978–983 (2008)
6 Chris Anbeek, “Surface roughness of minerals and implications for dissolution studies,” *Geochemica et Cosmochimica Acta* 56, 1461–1469 (1992)
7 Susan L. Brantley, “Kinetics of Mineral Dissolution,” in *Kinetics of Water-Rock Interaction*, edited by Susan L. Brantley, James D. Kubicki, and Art F. White (Springer New York, 2008) pp. 151–210.
8 W. A. Lanford, K. Davis, P. Lamarche, T. Laursen, R. Groege, and R. H. Doremus, “Hydration of soda–line glass,” *Journal of Non-Crystalline Solids* 33, 249–266 (1979)
9 William H. Casey and Henry R. Westrich, “Control of dissolution rates of orthosilicate minerals by divalent metal–oxygen bonds,” *Nature* 355, 157–159 (1992)
10 C. André Ohlin, Eric M. Villa, James R. Rustad, and William H. Casey, “Dissolution of insulating oxide materials at the molecular scale,” *Nature Materials* 9, 11–19 (2010)
11 James P. Hamilton, Carlo G. Pantano, and Susan L. Brantley, “Dissolution of albite glass and crystal,” *Geochimica et Cosmochimica Acta* 64, 2605–2615 (2000)
12 Roland Hellmann, Stéphane Cotte, Emmanuel Cadel, Saarain Malladi, Lisa S. Karlsson, Sergio Lozano-Perez, Martiane Cabié, and Antoine Seyeux, “Nanometre-scale evidence for interfacial dissolution–reprecipitation control of silicate glass corrosion,” *Nature Materials* 14, 307–311 (2015)
13 James P. Hamilton and Carlo G. Pantano, “Effects of glass structure on the corrosion behavior of sodium-aluminosilicate glasses,” *Journal of Non-Crystalline Solids: Practical Implications of Glass Structure*, 222, 167–174 (1997)
14 James P. Hamilton, Susan L. Brantley, Carlo G. Pantano, Louise J. Criscenti, and James D. Kubicki, “Dissolution of nepheline, jadeite and albite glasses: toward better models for aluminosilicate dissolution,” *Geochemica et Cosmochimica Acta* 65, 3683–3702 (2001)
15 J. C. Phillips, “Topology of covalent non-crystalline solids I. Short-range order in chalcogenide alloys,” *Journal of Non-Crystalline Solids* 34, 153–181 (1979)
16 M. F. Thorpe, “Continuous deformations in random networks,” *Journal of Non-Crystalline Solids* 57, 355–370 (1983)
17 J. C. Mauro, “Topological constraint theory of glass,” *American Ceramic Society Bulletin* 90, 31–37 (2011).
18 J. Clerk Maxwell, “L. On the calculation of the equilibrium and stiffness of frames,” *Philosophical Magazine Series* 4 27, 294–299 (1864).
19 Aditya Kumar, Jason Reed, and Gaurav Sant, “Vertical Scanning Interferometry: A New Method to Measure the Dissolution Dynamics of Cementitious Minerals,” *Journal of the American Ceramic Society* 96, 2766–2778 (2013)
20 Antonio C. Lasaga and Andreas Lutgge, “Variation of Crystal Dissolution Rate Based on a Dissolution Stepwave Model,” *Science* 291, 2400–2404 (2001)
21 Patricia M. Dove, Nizhou Han, and James J. De Yoreo, “Mechanisms of classical crystall growth theory explain quartz and silicate dissolution behavior,” *Proceedings of the National Academy of Sciences* 102, 15357–15362 (2005)
22 James Patrick Hamilton, *Corrosion behavior of sodium aluminosilicate glasses and crystals* (Academic Press Inc, 1993).
23 A. K. Varshneya, *Fundamentals of Inorganic Glasses* (Academic Press Inc, 1993).
24 M. Micoulaut, A. Kachmar, M. Bauchy, S. Le Roux, C. Massobrio, and M. Boero, “Structure, topology, rings, and vibrational and electronic properties of Ge$_x$S$_{1-x}$ glasses across the rigidity transition: A numerical study,” *Physical Review B* 88, 054203 (2013)
25 Mathieu Bauchy, Mohammad Javad Abdolhosseini Qomi, Christophe Bichara, Franz-Josef Ulm, and Roland J.-M. Pellenq, “Rigidity Transition in Materials: Hardness is Driven by Weak Atomic Constraints,” *Physical Review Letters* 114, 125502 (2015).
26 X.-L. Yuan and A. Cormack, “Si-O-Si bond angle and torsion angle distribution in vitreous silica and sodium silicate glasses,” *Journal of Non-Crystalline Solids* 319, 31–43 (2005).
27 M. Bauchy and M. Micoulaut, “Atomic scale foundation of temperature-dependent bonding constraints in network glasses and liquids,” *Journal of Non-Crystalline Solids* 357, 2530–2537 (2011).
28 Y. Vaill, T. Qu, M. Micoulaut, F. Chaimault, and P. Boolchand, “Direct evidence of rigidity loss and self-organization in silicate glasses,” *Journal of Physics: Condensed Matter* 17, 4889–4896 (2005)
29 Ye Xiang, Jincheng Du, Morten M. Smedskjaer, and John C. Mauro, “Structure and properties of sodium aluminosilicate glasses from molecular dynamics simulations,” *The Journal of Chemical Physics* 139, 044507 (2013)
M. Bauchy, “Structural, vibrational, and elastic properties of a calcium aluminosilicate glass from molecular dynamics simulations: The role of the potential,” The Journal of Chemical Physics 141, 024507 (2014).

M. Bauchy, “Topological constraints and rigidity of network glasses from molecular dynamics simulations,” American Ceramic Society Bulletin 91, 34–38A (2012).

M. Bauchy, M. Micoulaut, M. Celino, S. Le Roux, M. Boero, and C. Massobrio, “Angular rigidity in tetrahedral network glasses with changing composition,” Physical Review B 84, 054201 (2011).

B. C. Bunker, “Molecular mechanisms for corrosion of silica and silicate glasses,” Journal of Non-Crystalline Solids Proceedings of the First PAC RIM Meeting on Glass and Optical Materials, 179, 300–308 (1994).

Siting Zhang and Yun Liu, “Molecular-level mechanisms of quartz dissolution under neutral and alkaline conditions in the presence of electrolytes,” Geochemical Journal 48, 189–205 (2014).

Günther Heinz Frischat, Ionic diffusion in oxide glasses (Trans Tech Publications, 1975).

K. L. Ngai and S. W. Martin, “Correlation between the activation enthalpy and Kohlrausch exponent for ionic conductivity in oxide glasses,” Physical Review B 40, 10550–10556 (1989).

A. J. Moulson and J. P. Roberts, “Water in silica glass,” Trans. Faraday Soc. 57, 1208–1216 (1961).

E. W. Shaffer, J. Shi-Lan Sang, A. R. Cooper, and A. H. Heuer, “Diffusion of tritiated water in a-quartz,” Sam by BJ Gjdleto H, A Yoder, RA 1, 131 (1974).

Bruno MJ Smets and T. P. Lommen, “Role of molecular water in the leaching of glass,” Physics and Chemistry of Glasses 24, 35–6 (1983).

M. Bauchy and M. Micoulaut, “From pockets to channels: Density-controlled diffusion in sodium silicates,” Physical Review B 83, 184118 (2011).

Alexander Pelmenschikov, Helene Strandh, Lars G. M. Pettersson, and Jerzy Leszczynski, “Lattice Resistance to Hydrolysis of Si-O-Si Bonds of Silicate Minerals: Ab Initio Calculations of a Single Water Attack onto the (001) and (111) β-Cristobalite Surfaces,” The Journal of Physical Chemistry B 104, 5779–5783 (2000).

Kai Kristiansen, Markus Valtiner, George W. Greene, James R. Boles, and Jacob N. Israelachvili, “Pressure solution – The importance of the electrochemical surface potentials,” Geochimica et Cosmochimica Acta 75, 6882–6892 (2011).

James C. Orr, Victoria J. Fabry, Olivier Aumont, Laurent Bopp, Scott C. Doney, Richard A. Feely, Anand Gnanadesikan, Nicolas Gruber, Akio Ishida, Fortunat Joos, Robert M. Key, Keith Lindsay, Ernst Maier-Reimer, Richard Matear, Patrick Monfray, Anne Mouchet, Raymond G. Najjar, Gian-Kasper Plattner, Keith B. Rodgers, Christopher L. Sabine, Jorge L. Sarmiento, Reiner Schlitzer, Richard D. Slater, Ian J. Totterdell, Marie-France Weirig, Yasuhiro Yamanaka, and Andrew Yool, “Anthropogenic ocean acidification over the twenty-first century and its impact on calcifying organisms,” Nature 437, 681–686 (2005).