



INSTITUT DE FRANCE  
Académie des sciences

# *Comptes Rendus*

---

## *Géoscience*

### *Sciences de la Planète*

Daniel R. Neuville

**Glass, an ubiquitous material**

Volume 354, Special Issue S1 (2022), p. 1-14


Published online: 29 May 2023

Issue date: 1 December 2022

<https://doi.org/10.5802/crgeos.171>

**Part of Special Issue:** Glass, an ubiquitous material

**Guest editor:** Daniel Neuville (Université de Paris, Institut de physique du globe de Paris, CNRS)

 This article is licensed under the  
CREATIVE COMMONS ATTRIBUTION 4.0 INTERNATIONAL LICENSE.  
<http://creativecommons.org/licenses/by/4.0/>



*Les Comptes Rendus. Géoscience — Sciences de la Planète sont membres du  
Centre Mersenne pour l'édition scientifique ouverte*

[www.centre-mersenne.org](http://www.centre-mersenne.org)

e-ISSN : 1778-7025



---

Glass, an ubiquitous material / *Le verre, un matériau omniprésent*

# Glass, an ubiquitous material

## *Le verre, un matériau omniprésent*

Daniel R. Neuville<sup>® a</sup>

<sup>a</sup> IPGP, CNRS, Université de Paris, 1 rue Jussieu, 75005 Paris, France  
E-mail: [neuville@ipgp.fr](mailto:neuville@ipgp.fr)

**Abstract.** Glasses play a fundamental role in our daily lives at the economic, cultural, societal, energy and geological levels. Geological glasses testify to igneous activity of the Earth and represent an important material for tools and ornamental objects from the Paleolithic to the present day. Glasses are also used to manufacture technical materials, such as containers (dishes, glasses, jars, bottles...), screens (television, computer, smartphone...), fibers with multiple applications (reinforcement, information, transport, energy, health...), to ensure the storage of domestic or nuclear waste and, more recently, biomaterials (dental or bone implants...). Thus, silica-based glasses are at the heart of the history of the Earth and humanity. The variation in composition of natural and industrial glasses is vast but its structure is generally based on a tetrahedral frame of SiO<sub>4</sub> units, the backbone of more than 90% of the glasses that surround us in our daily lives. Around this silica frame, other chemical elements are well constrained to form a single unit structure for use as network modifying elements, charge compensator, dyes, volatiles, and other substances. This article is a short introduction on glass.

**Résumé.** Les verres jouent un rôle fondamental dans notre vie quotidienne aux niveaux économique, culturel, sociétal, énergétique et géologique. Les verres géologiques témoignent de l'activité ignée de la Terre et représentent une source importante d'outils et d'objets décoratifs du paléolithique à nos jours. Les verres sont également utilisés pour fabriquer des matériaux techniques, tels que des récipients (plats, verres, bocaux, bouteille...), des écrans (télévision, ordinateur, smartphone...), des fibres aux applications multiples (renforcement, transport d'information, énergie, santé...), pour assurer le stockage des déchets domestiques ou nucléaires et, plus récemment, des biomatériaux (implants dentaires ou osseux...). Les verres à base de silice sont donc au cur de l'histoire de la Terre et de l'humanité. La variation de la composition des verres naturels et industriels est vaste mais sa structure est généralement basée sur une trame tétraédrique d'unités SiO<sub>4</sub>, épine dorsale de plus de 90% des verres qui nous entourent au quotidien. Autour de cette trame de silice, les autres éléments chimiques se répartissent en éléments modificateurs de réseau, compensateur de charge, colorants, volatiles, et l'ensemble constitue un seul matériau ou substance. Cet article est une brève introduction au verre.

**Keywords.** Glass, Structure, Properties, Disordered matter, Liquid.

**Mots-clés.** Verre, Structure, Propriétés, Matière désordonnée, Liquide.

Published online: 29 May 2023, Issue date: 1 December 2022

### Prologue

The United Nations has proclaimed 2022 as the International Year of Glass, highlighting the role of glass in

the scientific, economic, artistic and cultural fields. Glass is essential to many vital technologies, as it facilitates the transition to a more sustainable world and beautifies our lives.

This United Nations resolution is a unique and historic event for glass. It is the result of a long journey initiated in 2018 which has been made possible thanks to the support of 1500 universities and research centers, learned societies and associations, museums, artists, manufacturers and by companies in 79 countries and across 5 continents. The formal resolution was adopted at the United Nations General Assembly on 18 May 2021.

Glass is obviously the material of our daily lives that we find in tableware, food packaging, bottles for beverage, cosmetic perfumery and pharmacy. We are surrounded by glass in our homes, we use it for transport (float glass, glass wool, rockwool, or textile glassfiber) and technological applications in optics and energy (photonics, battery). Glass is also an asset for prostheses (bioglasses for bone implants, for example) and we must not forget that glass is also an important repository for memories of humanity and the history of our planet and the universe. This special issue of *C. R. Geosciences* brings together several papers on these various aspects after a short introduction on this material, glass. Cormier et al. [2022] describe the glass structure and focus on cationic mixing based on X-ray and neutron diffraction at high temperature. Their paper shows also the link between glass and liquid. Next, the article edited by Bauchy team details challenges and opportunities of the computational techniques needed to understand the structure and properties of glasses and liquids [Liu et al., 2022]. Glass is not only our daily glass as shown by the paper by Pradel and Piarristeguy [2022] who show that glass is not only composed of oxide, but that the world of glasses is more complex because it is possible to obtain glasses with all the elements [Pradel and Piarristeguy, 2022] and with all the chemical bonds; a complementary article on metal glasses will be published in the *CR Physique* [Champion, 2022]. Glass has always accompanied humanity and it allows us to transcend our beliefs and fears, as shown by symbols, stories and myths around decorated and stained glasses. The paper by Hunault et al. [2022] presents an emblematic stained glass, and how its restoration has allowed to learn more about its origin and history, the rose of the Sainte-Chapelle. McCloy and Schuller [2022] discuss waste vitrification. Glass has multiple uses in our daily lives and it will become more and more present tomorrow, as shown in the article by Blanc et al. on

optical fibers and sensor development [Blanc et al., 2022].

Brauer and Hupa [2022] demonstrate that glass can also use to rebuild parts of the human body. They explain how since the first work made by Hench et al. [1971], it is possible to use glass for regenerating bone tissues. Actually, several glass compositions are used in medicine and dentistry [Brauer and Hupa, 2022] and it is easy to imagine that in the coming years, the proportion of bioglasses will further increase.

Glass is also a witness to our past, because the investigation of glasses found either in meteorites or in crystals in the form of glass inclusions enable us to trace the history of our planet. These glasses may have different water content which is the main volatile component in magmas and plays an important role in eruptive mechanisms. The paper by Le Losq et al. focuses on how water affects the polymerization of glasses and liquids [Le Losq et al., 2022]. Another key to understanding eruptive processes is the evolution of magma crystallization [Vetere et al., 2022]. Finally, Moretti [2022] focuses on the volatile-melt interaction, showing how these affect the redox state, hence the speciation state of gases released by active volcanic systems.

## 1. Introduction

Glass is a homogeneous material that can form from almost all elements of the periodic table (Figure 1) and all types of chemical bonds. This material differs from other states of matter (crystalline solid, liquid, gas and plasma) firstly by its solid state which is characterized by an absence of order at a great distance and, secondly, by the presence of a second-order transition, called glass temperature transition. The reader will find detailed and recent publications on all families of glasses and their properties in the publication “Handbook of Glass [Musgraves et al., 2019]”. In this introduction, we will present some of the general concepts accepted by the entire scientific community and taught in universities and engineering schools.

A glass is an amorphous solid. The term solid implies high viscosity, usually greater than  $10^{11}$  Pa·s. This viscosity limits the flow of the material. The term amorphous implies the absence of long-range order which reveals an analogy with the liquid state. A glass is therefore a solid whose certain properties are close

Legend:

- Network Former, NF (Orange)
- Network Modifier, NM or Charge Compensator, CC (Light Blue)
- Intermediate (Dark Blue)
- Ligand (Yellow)
- Ligand or NF (Orange-Yellow)
- Gas (Green)

1 1.008* <b>H</b> hydrogen																	2 4.003 <b>He</b> helium															
3 6.94* <b>Li</b> lithium	4 9.012 <b>Be</b> beryllium																	5 10.81* <b>B</b> boron	6 12.01* <b>C</b> carbon	7 14.01* <b>N</b> nitrogen	8 16.00* <b>O</b> oxygen	9 19.00 <b>F</b> fluorine	10 20.18 <b>Ne</b> neon									
11 22.99 <b>Na</b> sodium	12 24.31* <b>Mg</b> magnesium																	13 26.98 <b>Al</b> aluminium	14 28.09* <b>Si</b> silicon	15 30.97 <b>P</b> phosphorus	16 32.06* <b>S</b> sulfur	17 35.45* <b>Cl</b> chlorine	18 39.95 <b>Ar</b> argon									
19 39.10 <b>K</b> potassium	20 40.08 <b>Ca</b> calcium	21 44.96 <b>Sc</b> scandium	22 47.87 <b>Ti</b> titanium	23 50.94 <b>V</b> vanadium	24 52.00 <b>Cr</b> chromium	25 54.94 <b>Mn</b> manganese	26 55.85 <b>Fe</b> iron	27 58.93 <b>Co</b> cobalt	28 58.69 <b>Ni</b> nickel	29 63.55 <b>Cu</b> copper	30 65.38 <b>Zn</b> zinc	31 69.72 <b>Ga</b> gallium	32 72.63 <b>Ge</b> germanium	33 74.92 <b>As</b> arsenic	34 78.97 <b>Se</b> selenium	35 79.90* <b>Br</b> bromine	36 83.80 <b>Kr</b> krypton															
37 85.47 <b>Rb</b> rubidium	38 87.62 <b>Sr</b> strontium	39 88.91 <b>Y</b> yttrium	40 91.22 <b>Zr</b> zirconium	41 92.91 <b>Nb</b> niobium	42 95.95* <b>Mo</b> molybdenum	43 [98] <b>Tc</b> technetium	44 101.1 <b>Ru</b> ruthenium	45 102.9 <b>Rh</b> rhodium	46 106.4 <b>Pd</b> palladium	47 107.9 <b>Ag</b> silver	48 112.4 <b>Cd</b> cadmium	49 114.8 <b>In</b> indium	50 118.7 <b>Sn</b> tin	51 121.8 <b>Sb</b> antimony	52 127.6 <b>Te</b> tellurium	53 126.9 <b>I</b> iodine	54 131.3 <b>Xe</b> xenon															
55 132.9 <b>Cs</b> caesium	56 137.3 <b>Ba</b> barium	57-71 lanthanides	72 178.5 <b>Hf</b> hafnium	73 180.9 <b>Ta</b> tantalum	74 183.8 <b>W</b> tungsten	75 186.2 <b>Re</b> rhenium	76 190.2 <b>Os</b> osmium	77 192.2 <b>Ir</b> iridium	78 195.1 <b>Pt</b> platinum	79 197.0 <b>Au</b> gold	80 200.6 <b>Hg</b> mercury	81 204.4* <b>Tl</b> thallium	82 207.2 <b>Pb</b> lead	83 209.0 <b>Bi</b> bismuth	84 [209] <b>Po</b> polonium	85 [210] <b>At</b> astatine	86 [222] <b>Rn</b> radon															
87 [223] <b>Fr</b> francium	88 [226] <b>Ra</b> radium																	89 [227] <b>La</b> lanthanum	90 232.0 <b>Ce</b> cerium	91 231.0 <b>Pr</b> praseodymium	92 238.0 <b>Nd</b> neodymium	93 [237] <b>Pm</b> promethium	94 [244] <b>Pu</b> plutonium	95 [243] <b>Am</b> americium	96 [247] <b>Cm</b> curium	97 [247] <b>Bk</b> berkelium	98 [251] <b>Cf</b> californium	99 [252] <b>Es</b> einsteinium	100 [257] <b>Fm</b> fermium	101 [258] <b>Md</b> mendelevium	102 [259] <b>No</b> nobelium	103 [266] <b>Lr</b> lawrencium

**Figure 1.** Periodic table with possible role of elements in glass or melt. Elements in white can be in glass without any specific role. Elements in white are almost never investigated in glass structure, but they can be present in a glass probably as a gas or network modifier.

to those of liquids. Parks and Huffman even speak of “a fourth state of matter” [Parks and Huffman, 1926].

The glass state can occur for many chemical compositions as long as the crystallization phenomenon is avoided. There are mineral glasses and other common glasses based on silicate compounds, oxides, salts, metal, organic glasses such as gels, caramels, candies. In fact, it is possible to obtain glasses regardless of the type of chemical bond: covalent, ionic, metallic, Van der Waals or hydrogen. The state of glass therefore is characteristic of condensed matter and there can be an infinite palette of chemical composition of glasses, each with its own structure. As we will see, each of them, individually, can be defined as a material or substance in its own right with its own physical and chemical characteristics such as glass transition temperature ( $T_g$ ), X-ray pattern or Raman spectrum, elemental chemical analysis, density, refractive index, etc.

Many methods of preparation make it possible to obtain amorphous materials. They are generally developed by rapid cooling of a liquid phase, but it is also possible to obtain them from a gas phase, by amorphization of a crystalline phase or by sol-gel methods [Descamps, 2017]. The most common method of glass forming is by freezing of a liquid during rapid cooling. During the cooling of a liquid, there is a continuous increase in viscosity to such a value that the material can be considered a solid. We can therefore imagine the structure of the glass as being similar to that of a liquid, whose entire atomic movements are blocked. Glasses have, like liquids, a disordered structure with only a short-range order. For example, silica, a vitreous phase, consists of  $\text{SiO}_4$  tetrahedra, associated with each other by an oxygen atom but without long-range periodicity of the lattice, seen in crystals. The oxygen atom connecting two silicon atoms is said to bridge. Figure 2 shows

a 2D schematic representation of  $\text{SiO}_4$  tetrahedra for crystal and glass.

## 2. A disordered structure?

In Section 1, we mentioned a glass composed only of silicon and oxygen whose respective roles are that of network former and ligand, but how to differentiate the structure of silica from that of a quartz crystal? Figure 3 clearly shows for  $\text{SiO}_2$ , amorphous or crystalline (quartz or low-cristobalite), the spectral differences obtained by two different structural probes (X-ray diffraction, Raman spectrometry).

Regardless of the type of glass the experimental technique used, the crystalline phase data show fine peaks while those of glass show broad peaks. More precisely, X-ray diffraction shows very intense Bragg peaks up to a distance of  $8 \text{ \AA}^{-1}$  for quartz crystal and very low intensity wide strips for glass. The short-range order for distances less than  $3 \text{ \AA}$  characterizes the  $\text{SiO}_4$  coordination polyhedron, interatomic distances and bond angles, as well as the possibility of having heteropolar bonds (Si–O). To summarize, at a short distance, there is a local order that concerns the first layer of coordination. This is governed by the chemical bond that imposes the number of neighbors (coordination), the interatomic distances, and the symmetry of the site, for example a tetrahedron  $\text{SiO}_4$  in silicates.

The region at the longest distance between  $3\text{--}20 \text{ \AA}$  corresponds to the sequences and arrangements between polyhedra. It is on this scale that the distinction is made between crystalline and amorphous phase. A detailed analysis of this region provides important information on the knowledge of the angles between polyhedra, the type of connectivity (corner-, edges- or faces-sharing), the 2D or 3D dimensionality of the network and, potentially, the existence of structural/compositional heterogeneities. For X-ray diffraction, long-range or high-energy signal loss materializes the difference between an ordered and disordered structural state [Cormier and Neuville, 2022]. Raman spectrometry also highlights the differences between the spectrum of quartz and that of silica glass. In particular, the Raman spectrum of silica reveals glass-specific structures, which do not exist in the crystalline phases: these are rings with three or four  $\text{SiO}_4$  tetrahedra characterized by peaks at 490

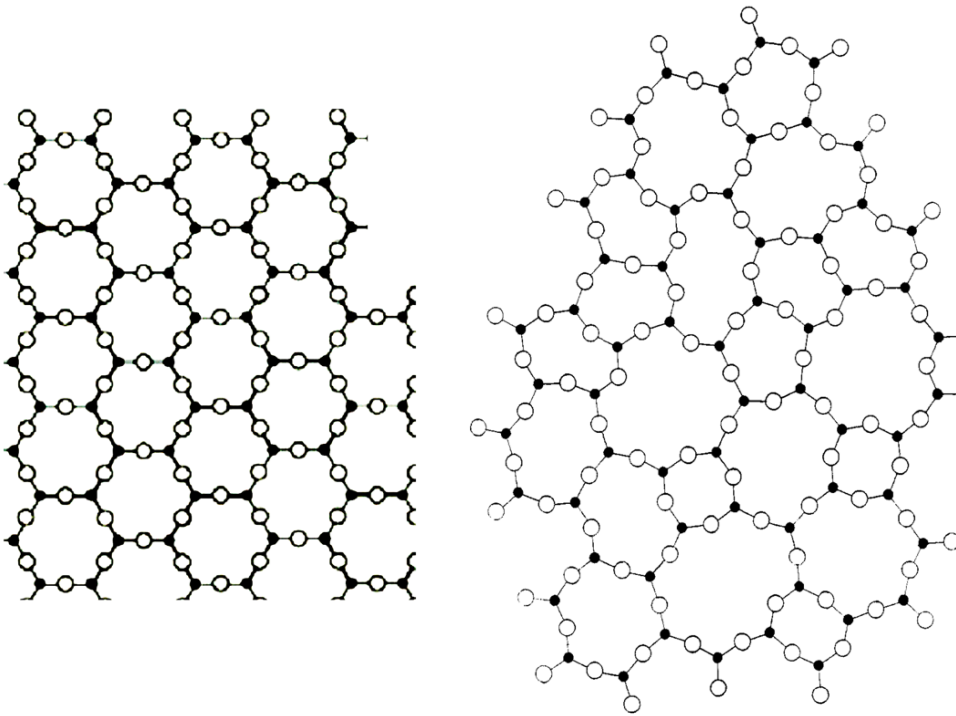
and  $605 \text{ cm}^{-1}$ , called D1 and D2, respectively. In addition, it should be noted that the Raman spectral efficiency of a quartz crystal and amorphous silica is strongly different. Indeed, the Raman spectrum of quartz is acquired in 10 s against 100 s for that of silica glass.

To summarize, at medium distance, the order is defined by the relations between the different sites, for example, the angle between two tetrahedra  $\text{SiO}_4$  in silicates. This order in glasses is difficult to characterize experimentally and is the subject of numerous experimental studies and numerical simulations [Liu et al., 2022]. Its characterization is essential because this scale of organization is directly related to the origin of most of the properties of glasses, such as density, ion conduction or chemical durability. Finally, at a great distance, there is no periodicity of structures, unlike crystals.

This short paragraph shows that, whatever the structural technique used, it is particularly difficult to grasp the nature and extent of the medium-distance order characteristic of the structural disorder of glass. In addition, this deficit of information is one of the major current challenges in the structural knowledge of glasses.

## 3. The glass transition temperature

Unlike the case of a crystallized solid, glass does not have a melting temperature. Conversely, the solidification of a glass corresponds to a continuous and gradual increase in viscosity during the cooling of a liquid, without the appearance of a crystalline structure. In Figure 4, we reported the variations in viscosity as a function of reciprocal temperature, for several molten silicates. There is no break in the viscosity curves between the values obtained at high temperature in the liquid state and those at low temperature in the glass state. This shows the continuous transition from glass to liquid for a property such as viscosity. The glass transition is a kinetic phenomenon that characterizes the loss of internal thermodynamic equilibrium. Indeed, the properties of a glass no longer depend only on pressure and temperature, but also on the speed,  $|q|$ , at which the glass transition temperature occurs (for a detailed discussion refer to Neuville and Le Losq [2022]).



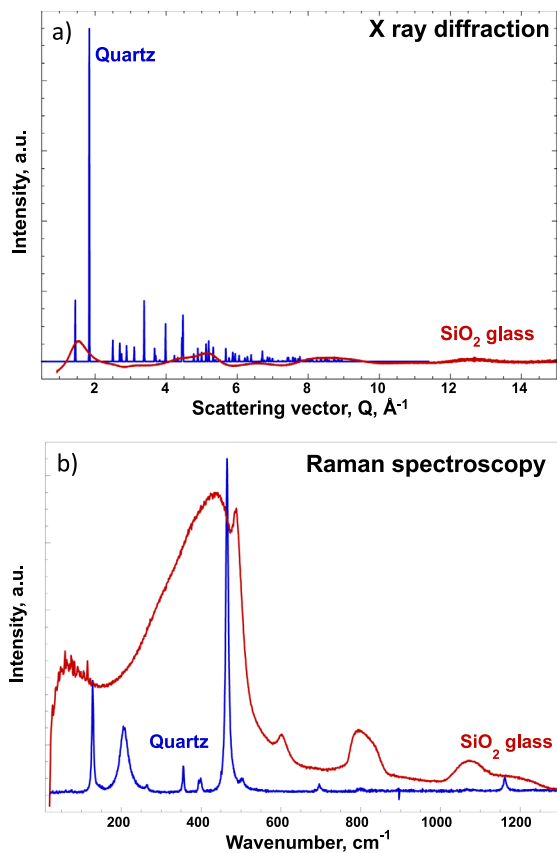
**Figure 2.** 2D representation of: (a) a crystal structure consisting of Si atom (black circle) and oxygen (white circle), there is a geometric pattern (hexagon) and a periodicity while in (b) atomic arrangements are present (rings with 3, 4, 5, 6 tetrahedra) but these arrangements are not regular or periodic [according to Zachariasen, 1932]. Note that both structures have the same number of Si atoms, but do not occupy the same surface area due to a lower density for glass than for crystal [modified from Neville and Cormier, 2022].

The simplest and oldest characterization of this transition has been described by Parks and Huffman [1927]. They note that by heating a glass, “it continuously passes from the amorphous state to the liquid state, its properties changing regularly with temperature, without any discontinuity”. In fact, as they write, “while there is no precise temperature comparable to the melting point of a crystal, there is nevertheless a temperature interval, net and reproducible, in which many properties change with a speed that approaches that is observed in the fusion of a crystal. In short, there is a softening interval instead of a melting point. Glass as it exists below this softening interval differs so markedly from the liquid that exists above that it could well be considered a different state from the substance.”

The glass transition corresponds to a zone of temperature and pressure in which properties such

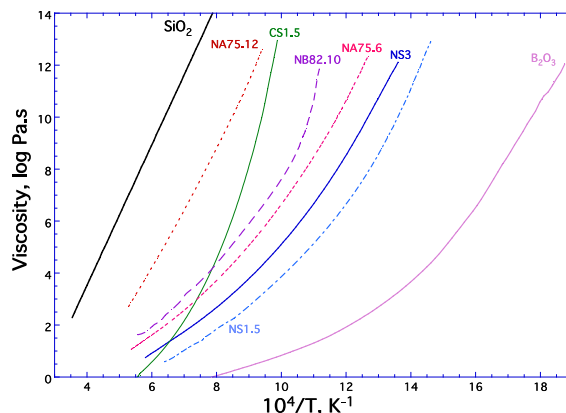
as specific heat undergo a second-order transition [Moynihan et al., 1976b,a]. Therefore, the physico-chemical properties of a glass depend on the speed at which the temperature varies and therefore on its thermal history.

It is theoretically possible to obtain identical temporal variations for all properties, such as thermal expansion or viscosity. It is therefore clear that there is a response time to the temperature variation. We call this time  $\tau$ , relaxation time. We define it as follows: a liquid has a large number of different configurational states [Goldstein, 1976, 1969, 2011]. Each state corresponds to a minimum of potential energy. When the temperature decreases, the time of structural relaxation of a liquid,  $\tau$ , increases. When  $\tau$  is large enough, the liquid can no longer carry out cooperative structural rearrangements [Goldstein, 1976, 1969]. Thus, the number of configurations that



**Figure 3.** (a) X-ray diffraction, (b) Raman spectrometry in blue (quartz or low-cristobalite) and SiO<sub>2</sub> amorphous in red. For practical reasons, spectra are reported in arbitrary units. The conditions of acquisition of the spectra of the crystalline and amorphous phases are different for the same technique and from one technique to another [redrafted from Cormier and Neuville, 2022].

can be reached by a system decreases with temperature and, at sufficiently low temperature ( $T < T_g$ ), the liquid freezes: the system can no longer change its configurational state (a more detailed discussion can be found in Neuville and Le Losq [2022]). In addition, the configuration state of the glass does not change on cooling from  $T_g$  to 0 K. It can be noted that Raman and infrared vibrational spectroscopy suggest that the structure of a glass is an image of the instantaneous configuration of the liquid [Kashio et al., 1980, Kusabiraki and Shiraishi, 1981a,b, Neuville and Mysen, 1996].



**Figure 4.** Viscosity as a function of reciprocal temperature for a silica glass, SiO<sub>2</sub>, sodium silicate glasses, NS3 and NS2, containing respectively 75 and 66 mol% of SiO<sub>2</sub> and the difference to 100 being Na<sub>2</sub>O, the compositions NS1.5 and CS1.5 correspond respectively to 60 SiO<sub>2</sub>–40 Na<sub>2</sub>O and 60 SiO<sub>2</sub>–40 CaO (mol%) from Neuville [2006] for CS1.5, NS1.5, NS2. The compositions NA75.6 and NA75.12 contain 75 mol% SiO<sub>2</sub> each and 6 or 12 mol% Al<sub>2</sub>O<sub>3</sub>, the difference to 100 being Na<sub>2</sub>O [data of Le Losq et al., 2014] for NS3 and NAX. Y compositions, Napolitano et al. [1965] for B<sub>2</sub>O<sub>3</sub>; from Sipp et al. [1997] for NB82.10 = Pyrex.

Viscosity is an important and widely used property by the glass industry. It has therefore been, for the sake of uniformity, standardized. The temperature of the glass transition is industrially defined as the temperature at which the viscosity is 10<sup>12</sup> Pa·s, for a cooling rate of the order of a few °C·min<sup>-1</sup>, which corresponds to the cooling rates of the glass in industrial processes. This viscosity value corresponds to relaxation times of about 15 min.

From a thermodynamic point of view, we can state that temperature and pressure are not enough to characterize the glass state. Tool and Eichlin [1931] introduced the concept of fictive temperature,  $T_{fic}$ , to characterize glass at constant pressure. We can define it as the temperature at which the glass would be in a steady state if it could be heated instantly. The fictive temperature can be defined during cooling in dilatometry or calorimetry, as being the glass transition temperature.

It is possible to give a formal thermodynamic

definition of the fictive temperature, considering it as an order parameter. It is recalled that the glass transition resembles a second-order transition for which  $V$ ,  $H$ ,  $\eta$  are continuous functions but not  $C_p$ ,  $\alpha$ ,  $K$  [see the reviews Neuville and Le Losq, 2022].

three-dimensional network of silica (about 14 orders of magnitude near 1000 °C, Figure 3). In addition to reducing viscosity, boron oxide is added to glasses because it greatly improves the thermal and mechanical properties of glass as in the case of Pyrex<sup>®</sup> glass (composition NB82.10, Figure 4).

## 4. Chemical role of elements in glass

### 4.1. Cations

The structure of glasses involves a large family of elements divided into several groups, network formers, network modifiers or charge compensators, intermediate elements, coloring elements, refining.

(i) “*Network formers*” are the oxides that can form a glass alone. These elements are, mainly, in their cationic forms:  $\text{Si}^{4+}$ ,  $\text{B}^{3+}$ ,  $\text{Ge}^{4+}$  . . . . These network former cations have very covalent, and therefore, directional cation–oxygen bonds that ensure the three-dimensional framework of the glass. These elements form small size polyhedral with a ligand-like oxygen (for example, tetrahedral for Si and Ge or triangular for B). In the case of silicate glasses, two cations are connected to each other by so-called bridging oxygens but in thio-silicate glasses, they are connected by a bridging sulfur [Pradel and Piaristeguy, 2022]. These connected polyhedra form a three-dimensional disordered lattice. So, it is possible to build a glass, with several ligands, O, S, Se, Te. For this short introduction, we will only talk about ligand O in oxide glasses.

Note that silicon oxide,  $\text{SiO}_2$ , or silica, is the majority oxide of commercial glasses, and is the main former of the vitreous network. In the case of silica, the tetrahedral entities are denoted  $Q^n$  where  $n$  represents the number of bridging oxygens carried by the central atom Q (typically Si). The silica glass thus consists of Si surrounded by 4 bridging oxygens, all connected to other tetrahedra to form the three-dimensional network of species  $Q^4$  with Si–O–Si bridges (Figure 2b). This structure is characterized by very high viscosity on the one hand and an Arrhenian dependence of viscosity with temperature on the other hand (Figure 3). Boron oxide  $\text{B}_2\text{O}_3$  also acts as a lattice former, but it forms  $\text{BO}_3$  plane triangles. Its structure therefore has a strongly two-dimensional character, which explains why the viscosity of  $\text{B}_2\text{O}_3$  is significantly lower than that of the



(ii) “*Network modifiers*” are oxides that do not form glass by themselves. These are mainly alkaline and alkaline-earth elements. The introduction of  $R^{n+}O_{n/2}$  ( $n = 1$  or  $2$ ) changes the continuity of the three-dimensional network constituted by the network former. The network is broken with formation of non-bridging oxygens, which are no longer bound only to network formers. Alkaline and alkaline-earth elements play a similar role with regard to the structure and main properties of glasses and melts, but have some different characteristics. Thus, sodium oxide,  $Na_2O$ , transforms silica into silicate by chemical attack and is mainly used to reduce the temperature necessary for the fusion of  $SiO_2$  compound. On the contrary, calcium oxide, because it forms slightly stronger Ca–O bonds than Na–O bonds, increases the melting temperature of the mixture and also improves several properties such as the durability of the glass. These 3 oxides,  $SiO_2$ ,  $Na_2O$  and  $CaO$  represent more than 95% of the weight composition of industrial glasses and are, therefore, generally called soda-lime silicate glasses. From a practical point of view, a large majority of commercial glasses also contain potassium oxide,  $K_2O$ , and magnesium oxide,  $MgO$ . Magnesium oxide is usually added because it improves the mechanical properties and the hydrolytic strength of glasses like aluminum oxide. However, these two oxides also tend to increase the temperature of the silicate bath and the working temperatures of the glass. By adding  $R^{n+}O_{n/2}$  to silica, non-bridging oxygens are created, which establish  $R \cdots O$  ionic bonds with the  $R^{n+}$  cation. The  $R \cdots O$  bond is weaker than that involved in Si–O–Si bridges, resulting in a strong decrease in viscosity. This is clearly visible by comparing the viscosity curve of  $SiO_2$  and NS3 glass compositions (Figure 4). The formation of non-bridging oxygens can be easily investigated by several techniques like Raman spectroscopy or nuclear magnetic resonance (NMR). Their proportions increase with sodium content which increases the weakly connected entities,  $Q^3$ ,  $Q^2 \dots$ , contributing to the decrease in viscosity. In Figure 4, NS1.5 has a lower viscosity than NS3, which results from an increase of the sodium content from 25 to 40 mol%. As we have seen with NS3, non-bridging oxygens are formed when lattice modifying elements ( $Li^+$ ,  $Na^+$ ,  $K^+$ ,  $Mg^{2+}$ ,  $Ca^{2+}$ ,  $Fe^{2+} \dots$ ) are added to depolymerize the 3D network made by the network former. In Figure 4, comparing the viscosity curve of NS1.5 which

is lower than that of CS1.5, it is clear that sodium participates more in the breaking of the network than calcium.

(iii) “*Charge compensators*”, alkaline and/or alkaline earth elements can change their roles and become charge compensators in presence of  $(AlO_4)^-$  and  $(BO_4)^-$  which are tetrahedra with a deficit charge with respect to  $SiO_4$  tetrahedra. The aluminum fits into the network in a tetrahedral site, similar to  $Si^{4+}$ . But, being an  $Al^{3+}$  cation, the  $AlO_4$  tetrahedron has a charge deficit that requires compensation to stabilize it energetically, which is accomplished by the proximity of an alkaline or alkaline earth element. The case of boron,  $B^{3+}$ , is different because it is the result of a change in boron coordination.  $B_2O_3$  glass is a network of  $BO_3$  triangles, fully stabilized with a low viscosity curve, which can be transformed into a network  $(BO_4)^-$  tetrahedra by the addition of an alkali or alkaline-earth oxides, which requires a charge compensation around  $(BO_4)^-$  similar to  $(AlO_4)^-$ . The formation of these  $(BO_4)^-$  tetrahedra increases the connectivity of the network, which is evident from the fact that the viscosity curve of the NB82.10 glass is more viscous than that is observed for  $B_2O_3$  melt (Figure 4).

(iv) “*Intermediate elements*”, display a behavior as network formers but in general cannot form glass alone. It should be noted that Al, can form glasses, called inverted glasses [Neuville et al., 2008b,a, 2010, Licheron et al., 2011, Drewitt et al., 2012, 2017]. The main intermediate elements are  $Al^{3+}$ ,  $Pb^{4+}$ ,  $Ti^{4+}$  or  $Fe^{3+}$ . Aluminum is mainly in coordination 4 [Stebbins and Xu, 1997, Neuville et al., 2004c,a,b, Neuville, 2006, Neuville et al., 2008b,a, Stebbins, 2008, Le Losq et al., 2014, Drewitt et al., 2022, Neuville and Le Losq, 2022]. By adding  $Al_2O_3$  to the NS3 glass composition, viscosity increases strongly and becomes quasi-Arrhenian again, as illustrated in Figure 3 for the viscosity of NA75.6 and NA75.12 (albite glass composition) which contain 6 and 12 mol% of  $Al_2O_3$  respectively. A similar behavior is observed when considering the viscosity of molten wollastonite,  $CaSiO_3$ , and anorthite,  $CaAl_2Si_2O_8$ . Both albite and anorthite glass compositions are fully polymerized systems and form a three-dimensional network of  $AlO_4^-$  and  $SiO_4$  tetrahedra with  $Na^+$  or  $Ca^{2+}$  in the holes of the three-dimensional network, playing the role charge compensator of aluminum.  $Al^{3+}$  in four-fold coordination then plays a role of network former.  $Al^{3+}$

can also be in five-fold coordination [Neuville et al., 2004c,a,b, Neuville, 2006, Neuville et al., 2008b,a, Le Losq et al., 2014, Drewitt et al., 2022, Neuville and Le Losq, 2022]; in this condition, it is even possible to define a role of reticator (i.e. an element that ensures a strong connection between network former elements, according to Patrick Royall et al. [2008]).

Intermediate elements can change their role as a function of the chemical composition of the glass and their proportions in the glass [i.e. Neuville, 2006, Neuville et al., 2008b], temperature [i.e. Stebbins et al., 1992, Stebbins and Farnan, 1992, Stebbins, 2008, Neuville et al., 2008a] and pressure [i.e. Allwardt et al., 2005, 2007] but also according to oxygen fugacity [i.e. Moretti, 2005].

(v) *Transition elements*, can play the role of coloring or decoloring agents [Hunault et al., 2022]. A more detailed discussion about colored glasses can be found in Cormier et al. [1999], Galois et al. [2005], Hunault et al. [2014, 2016], and in the review of de Ligny and Möncke [2019].

#### 4.2. Anions

Cations make up at most 50% of a glass, the rest being anions. In the anion subnetwork, the most important element is usually oxygen which occupies different positions. We distinguish in fact (Figure 5):

- bridging oxygens, BO, which link tetrahedra together and which have no effective charge because they are polarized by two covalent bonds;
- non-bridging oxygens, NBO, associated with a single tetrahedron. They therefore have a residual charge because they are partially polarized by a single covalent bond;
- free oxygens, FO, associated with no tetrahedron and which are therefore less polarized;
- tricluster oxygens, connected to three polyhedra, usually one or two tetrahedra and one pentahedron. These tricluster oxygens represent only a small percentage and play a minor role in the structure and properties of glasses and liquids [Iuga et al., 2005].

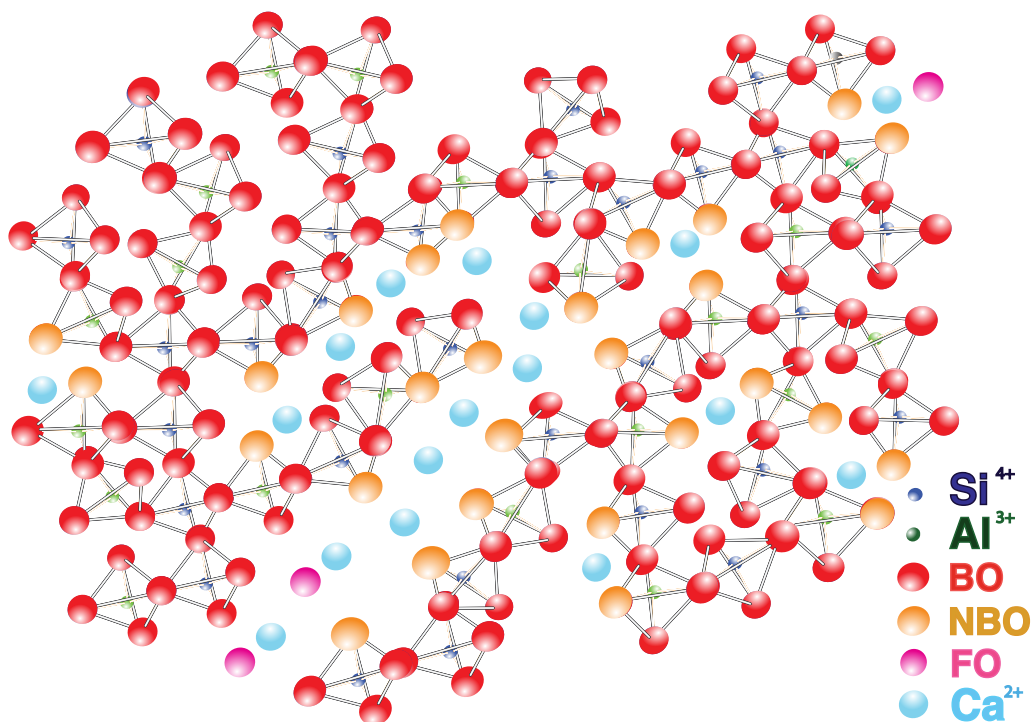
This vision of a network made by oxygens with different roles is complementary to that described by the cations. Some experimental techniques such as  $^{17}\text{O}$  NMR, make it possible to visualize this anionic

network and highlight the connectivity of the network [Eden, 2012, Stebbins et al., 2001].

This anionic network is also observed in chalcogenide glasses in which sulfur or selenium play a role of ligand identical to that of oxygen. In these glasses, it is possible to show, as already mentioned, the existence of bridging or non-bridging sulfur [Pradel and Piarristeguy, 2022].

### 5. A universal idea of the glass structure?

The ideas developed in the previous paragraphs show that a glass, seen by the cationic or anionic network, has a disordered character. However, Greaves et al. [1981] showed using X-ray absorption near edge structure (XANES) at the Na K-edge that the short-range order around  $\text{Na}^+$  in silicate glasses was relatively well established. Moreover, these cations have a non-random distribution within the silicate network [Greaves, 1985]. Following these pioneering studies, spectroscopic (Extended X-ray absorption fine structure, NMR, neutron diffraction...) or rheological (viscosity and calorimetry) studies have shown that cation–oxygen distances and cation coordination are well defined [Taylor and Brown, 1979a,b, Calas and Petiau, 1983, Cormier et al., 2001a,b, 1998, 2022, Eckersley et al., 1988, Le Losq et al., 2017, Neuville et al., 2007, Neuville, 2006, Neuville et al., 2004c,a]. Other studies have confirmed the existence of medium range orders around modifying cations in the structuring of the silicate or aluminosilicate network [Gaskell et al., 1991, Greaves, 1985, Cormier et al., 2001a,b, Meyer et al., 2004, Greaves and Sen, 2007, Kargl and Meyer, 2008, Caurant et al., 2010, Le Losq et al., 2017]. According to these studies, cations are not only randomly distributed within the structure of glass, which implies a modification of the classical model proposed by Warren et al. [Warren, 1941, Warren and Bischof, 1938]. Greaves [1985] proposed the Modified Random Network (MRN) distributed into domains enriched in modifying cations, separated from domains enriched in network former. It is possible to distinguish channels or clusters, more or less connected where network modifiers, non-bridging and free oxygens, and also volatile elements are concentrated. Molecular dynamic simulations on sodium and potassium silicates also highlight the existence of such structures in the liquid state [Kargl and Meyer, 2008, Meyer



**Figure 5.** Structural sketch of a calcium aluminosilicate glass.

et al., 2004]. The existence of percolation channels can explain the rapid ion diffusion decoupled from the network in glasses and liquids near  $T_g$  [Greaves and Sen, 2007] which can also explain nucleation and growth as proposed by Neuville and Le Losq [2022]. In addition, it is in agreement with the observation of unmixing zones along the MO–SiO<sub>2</sub> binary joints (with M = Mg, Ca, Na), which can be explained by the immiscibility between M–Q<sup>3</sup> pairs and Q<sup>4</sup> species [Neuville, 2006].

Greaves and Ngai proposed a “modified” version of the MRN model for vitreous tectosilicates [Greaves and Ngai, 1995]: the Compensated Continuous Random Network (CCRN) model predicts that charge compensating cations are also present in channels but at the limits and close to elements or entities such as (AlO<sub>4</sub>)<sup>−</sup> or (BO<sub>4</sub>)<sup>−</sup> that require a charge compensation. The existence of subnetworks, one enriched in network former and one in the network modifier is in agreement with the simulations of Shintani and Tanaka [2006], which suggest that structures showing a quasi-crystalline order exist in the supercooled liquid close to  $T_g$  and strongly influence the dynamics

of the system. There is therefore probably a link between structural and compositional heterogeneities [Moesgaard et al., 2010]. These percolation channels enriched in network modifier elements can also be the starting points of unmixing and/or nucleation zones [Dargaud et al., 2011, 2012]. This picture of a glass and melt organized in percolation channels is now recognized as a universal approach to the structure of oxide glasses and also chalcogenide glasses.

## 6. Conclusion

Whether by its structure or properties, a glass with a given chemical composition is a unique material. The oldest glasses in our solar system are several billion years old and could be observed on the Moon surface (4.53 billion years). On Earth, the oldest natural glasses are younger, because glass is a highly alterable material in the presence of water, which causes its alteration in a few million years: this makes it difficult on Earth to find glasses older than a few hundred million years.

Regardless of the place and time of manufacture, whether natural or man-made, if the chemical composition and formation conditions are identical, the glass manufactured will be identical in terms of structure and properties. The chemical composition and the conditions of formation are therefore the criteria that will determine the kind of glass to be obtained. It is important to emphasize that glass is not the addition of several oxides but a new material resulting from the reaction of the mixture of raw materials.

In this short introduction, we have shown that glass is a material with a disordered structure, characterized by the absence of order at a long distance, and presenting a phenomenon of glass transition. The structural role of chemical elements in a glass is relatively well constrained between network former, network modifier, charge compensator, intermediate and transition elements. The role of the elements can vary depending on their concentration, and the conditions of formation (pressure, temperature, oxygen fugacity etc.). All these features make glass a unique and ubiquitous material.

## Conflicts of interest

The author has no conflict of interest to declare.

## Acknowledgments

A big thank you to François Chabaud who gave me the opportunity to make this issue on glass within the framework of the International Year of Glass and thanks also to all the authors who made very beautiful papers which allow to show the universality of glass. Ongoing friendly discussions with Laurent Cormier, Annie Pradel, Delia Brauer and Roberto Moretti are very much appreciated.

## References

- Allwardt, J. R., Poe, B. T., and Stebbins, J. F. (2005). The effect of fictive temperature on Al-coordination in high-pressure (10 GPa) Na aluminosilicate glasses. *Am. Mineral.*, 90, 1453–1457.
- Allwardt, J. R., Stebbins, J. F., Terasaki, H., Du, L. D., Frost, D. J., Withers, A. C., Hirschmann, M. M., Suzuki, A., and Ohtani, E. (2007). Effect of structural transitions on properties of high-pressure silicate melts:  $^{27}\text{Al}$  NMR, glass densities, and melt viscosities. *Am. Mineral.*, 92, 1093–1104.
- Blanc, W., Schenato, L., Molardi, C., Palmieri, L., Galtarossa, A., and Tosi, D. (2022). Distributed fiber optics strain sensors: from long to short distance. *C. R. Géosci.*, 354(S1), 121–143.
- Brauer, D. S. and Hupa, L. (2022). Glass as a biomaterial: strategies for optimising bioactive glasses for clinical applications. *C. R. Géosci.*, 354(S1), 185–197.
- Calas, G. and Petiau, J. (1983). Structure of oxides glasses. Spectroscopic studies of local order and crystallochemistry. Geochemical implications. *Bull. Mineral.*, 106, 33–55.
- Caurant, D., Majèrus, O., Fadel, E., Quintas, A., Gervais, C., Charpentier, T., and Neuville, D. R. (2010). Structural investigation of borosilicate glasses containing  $\text{MoO}_3$  by MAS NMR and Raman spectroscopy. *J. Nucl. Mater.*, 396, 94–101.
- Champion, Y. (2022). Entropy of metallic glasses and the size effect on glass transition. *C. R. Phys.*, 24. Forthcoming.
- Cormier, L., Calas, G., and Gaskell, P. H. (2001a). Cationic environment in silicate glasses studied by neutron diffraction with isotopic substitution. *Chem. Geol.*, 174, 349–363.
- Cormier, L., Galois, L., and Calas, G. (1999). Evidence of Ni-containing ordered domains in low alkali borate glasses. *Europhys. Lett.*, 45, 572–578.
- Cormier, L., Galois, L., Delaye, J.-M., Ghaleb, D., and Calas, G. (2001b). Short- and medium-range structural order around cations in glasses: a multidisciplinary approach. *C. R. Acad. Sci. Sér. IV*, 2, 249–262.
- Cormier, L., Gaskell, P. H., Calas, G., and Soper, A. K. (1998). Medium-range order around titanium in a silicate glass studied by neutron diffraction with isotopic substitution. *Phys. Rev. B*, 58, 11322–11330.
- Cormier, L., Hennet, L., Lelong, G., Cuello, G., and Bytchkov, A. (2022). Structure from glass to melt: a case study along the  $\text{MgSiO}_3$ – $\text{CaSiO}_3$  join using neutron and X-ray diffraction. *C. R. Géosci.*, 354(S1), 15–34.
- Cormier, L. and Neuville, D. R. (2022). Les verres, quel désordre? *Reflets Phys.* (in press).
- Dargaud, O., Cormier, L., Menguy, N., and Patriarche,

- G. (2012). Multi-scale structuration of glasses: Observations of phase separation and nanoscale heterogeneities in glasses by Z-contrast scanning electron transmission microscopy. *J. Non-Cryst. Solids*, 358(10), 1257–1262.
- Dargaud, O., Cormier, L., Menguy, N., Patriarche, G., and Calas, G. (2011). Mesoscopic scale description of nucleation processes in glasses. *Appl. Phys. Lett.*, 99(2), article no. 021904.
- de Ligny, D. and Möncke, D. (2019). Colors in glasses. In Musgraves, J. D., Hu, J., and Calvez, L., editors, *Springer Handbook of Glass, Springer Handbooks*, pages 297–342. Springer International Publishing, Cham.
- Descamps, M. (2017). États amorphe et vitreux des composés moléculaires et pharmaceutiques - Propriétés générales. *Tech. Ing.*, article no. PHA2030.
- Drewitt, J. W. E., Barnes, A. C., Jahn, S., Kohn, S. C., Walter, M. J., Novikov, A., Neuville, D. R., Fischer, H. E., and Hennet, L. (2017). Structure of liquid tri-calcium aluminate. *Phys. Rev. B*, 95, 064203–064214.
- Drewitt, J. W. E., Hennet, L., and Neuville, D. R. (2022). From short to medium range order in glasses and melts by diffraction and Raman spectroscopy. *Rev. Mineral. Geochem.*, 87, 55–103.
- Drewitt, J. W. E., Hennet, L., Zeidler, A., Jahn, S., Salmon, P. S., Neuville, D. R., and Fischer, H. E. (2012). Structural transformations on vitrification in the fragile glass forming system  $\text{CaAl}_2\text{O}_4$ . *Phys. Rev. Lett.*, 109, 235501–235506.
- Eckersley, M. C., Gaskell, P. H., Barnes, A. C., and Chieux, P. (1988). Structural ordering in a calcium silicate glass. *Nature*, 335, 525–527.
- Eden, M. (2012). NMR studies of oxide-based glasses. *Ann. Rep. Prog. Chem. Sect. C*, 108, 177–221.
- Galoisy, L., Calas, G., Cormier, L., Marcq, B., and Thibault, M. H. (2005). Overview of the environment of Ni in oxide glasses in relation to the glass colouration. *Phys. Chem. Glass.*, 46, 394–399.
- Gaskell, P. H., Eckersley, M. C., Barnes, A. C., and Chieux, P. (1991). Medium-range order in the cation distribution of a calcium silicate glass. *Nature*, 350, 675–677.
- Goldstein, M. (1969). Viscous liquids and the glass transition: a potential energy barrier picture. *J. Chem. Phys.*, 51, 3728–3739.
- Goldstein, M. (1976). Viscous liquids and the glass transition. V. Sources of the excess specific heat of the liquid. *J. Chem. Phys.*, 64, 4767–4774.
- Goldstein, M. (2011). On the reality of the residual entropy of glasses and disordered crystals: The entropy of mixing. *J. Non-Cryst. Solids*, 357, 463–465.
- Greaves, G. N. (1985). EXAFS and the structure of glass. *J. Non-Cryst. Solids*, 71, 203–217.
- Greaves, G. N., Fontaine, A., Lagarde, P., Raoux, D., and Gurman, S. J. (1981). Local structure of silicate glasses. *Nature*, 293, 611–616.
- Greaves, G. N. and Ngai, K. L. (1995). Reconciling ionic-transport properties with atomic structure in oxide glasses. *Phys. Rev. B*, 52, 6358–6380.
- Greaves, G. N. and Sen, S. (2007). Inorganic glasses, glass-forming liquids and amorphizing solids. *Adv. Phys.*, 56, 1–166.
- Hench, L. L., Splinter, R. J., Allen, W. C., and Greenlee, T. K. (1971). Bonding mechanisms at the interface of ceramic prosthetic materials. *J. Biomed. Mater. Res.*, 5, 117–141.
- Hunault, M., Calas, G., Galoisy, L., Lelong, G., and Neuville, M. (2014). Local ordering around tetrahedral  $\text{Co}^{2+}$  in silicate glasses. *J. Am. Ceram. Soc.*, 97, 60–62.
- Hunault, M. O. J. Y., Bauchau, F., Boulanger, K., Hérold, M., Calas, G., Lemasson, Q., Pacheco, C., and Loisel, C. (2022). The rose of the Sainte-Chapelle in Paris: sophisticated stained glasses for late medieval painters. *C. R. Géosci.*, 354(S1), 101–120.
- Hunault, M. O. J. Y., Galoisy, L., Lelong, G., Neuville, M., and Calas, G. (2016). Effect of cation field strength on  $\text{Co}^{2+}$  speciation in alkali-borate glasses. *J. Non-Cryst. Solids*, 451, 101–110.
- Iuga, D., Morais, C., Gan, Z., Neuville, D. R., Cormier, L., and Massiot, D. (2005). NMR heteronuclear correlation between quadrupolar nuclei in solids. *J. Am. Chem. Soc.*, 127, 11540–11541.
- Kargl, F. and Meyer, A. (2008). Na-relaxation and intermediate range structure in sodium-potassium silicate melts. *Chem. Geol.*, 256, 278–285.
- Kashio, S., Iguchi, I., Goto, T., Nishina, Y., and Fuwa, T. (1980). Raman spectroscopic study on the structure of silicate slag. *Trans. Iron Steel Inst. Japan.*, 20, 251–253.
- Kusabiraki, K. and Shiraishi, Y. (1981a). Infrared emission spectra of molten alkaline metal silicates. *J. Japan. Inst. Metals*, 45, 250–258.
- Kusabiraki, K. and Shiraishi, Y. (1981b). On the infrared emission of the molten  $\text{Na}_2\text{O-Al}_2\text{O}_3\text{-SiO}_2$ .

- J. Japan. Inst. Metals*, 45, 888–895.
- Le Losq, C., Mysen, B., and Cody, G. (2022). Water solution mechanism in calcium aluminosilicate glasses and melts: insights from *in* and *ex situ* Raman and  $^{29}\text{Si}$  NMR spectroscopy. *C. R. Géosci.*, 354(S1), 199–225.
- Le Losq, C., Neuville, D. R., Florian, P., Henderson, G. S., and Massiot, D. (2014). Role of  $\text{Al}^{3+}$  on rheology and nano-structural changes of sodium silicate and aluminosilicate glasses and melts. *Geochim. Cosmochim. Acta*, 126, 495–517.
- Le Losq, C., Neuville, D. R., Florian, P., Massiot, D., Zhou, Z., Chen, W., and Greaves, N. (2017). Percolation channels: a universal idea to describe the atomic structure of glasses and melts. *Sci. Rep.*, 7, article no. 16490.
- Licheron, M., Montouillout, V., Millot, F., and Neuville, D. R. (2011). Raman,  $^{27}\text{Al}$  NMR structure investigations of aluminate glasses:  $(1 - x)\text{Al}_2\text{O}_3 - x\text{MO}$ , with  $\text{M} = \text{Ca}, \text{Sr}, \text{Ba}$  and  $0.5 < x < 0.75$ . *J. Non-Cryst. Solids*, 257, 2796–2801.
- Liu, H., Zhao, Z., Zhou, Q., Chen, R., Yang, K., Wang, Z., Tang, L., and Bauchy, M. (2022). Challenges and opportunities in atomistic simulations of glasses: a review. *C. R. Géosci.*, 354(S1), 35–77.
- McCloy, J. S. and Schuller, S. (2022). Vitrification of wastes: from unwanted to controlled crystallization, a review. *C. R. Géosci.*, 354(S1), 145–184.
- Meyer, A., Horbach, J., Kob, W., Kargl, F., and Schober, H. (2004). Channel formation and intermediate range order in sodium silicate melts and glasses. *Phys. Rev. Lett.*, 93, article no. 027801.
- Moesgaard, M., Keding, R., Skibsted, J., and Yue, Y. (2010). Evidence of intermediate-range order heterogeneity in calcium aluminosilicate glasses. *Chem. Mater.*, 22, 4471–4483.
- Moretti, R. (2005). Polymerisation, basicity, oxidation state and their role in ionic modelling of silicate melts. *Ann. Geophys.*, 48, 583–608.
- Moretti, R. (2022). Redox behavior of degassing magmas: critical review and comparison of glass-based oxybarometers with application to Etna volcano. *C. R. Géosci.*, 354(S1), 249–279.
- Moynihan, C. T., Easteal, A. J., Tran, D. C., Wilder, J. A., and Donovan, E. P. (1976a). Heat capacity and structural relaxation of mixed-alkali glasses. *J. Am. Ceram. Soc.*, 59, 137–140.
- Moynihan, C. T., Sasabe, H., and Tucker, J. (1976b). Kinetics of the glass transition in a calcium-potassium nitrate melt. *Proc. Electrochem. Soc.*, 6, 182–194.
- Musgraves, J. D., Hu, J., and Calvez, L., editors (2019). *Springer Handbook of Glass*, Springer Handbooks. Springer, Cham.
- Napolitano, A., Macedo, P. B., and Hawkins, E. G. (1965). Viscosity and density of boron trioxide. *J. Am. Ceram. Soc.*, 48, 613–616.
- Neuville, D. R. (2006). Viscosity, structure and mixing in (Ca, Na) silicate melts. *Chem. Geol.*, 229, 28–42.
- Neuville, D. R. and Cormier, L. (2022). Le verre : un matériau d’hier, d’aujourd’hui et de demain. *Matér. Tech.*, 110(4), article no. 404.
- Neuville, D. R., Cormier, L., Flank, A. M., de Ligny, D., Roux, J., and Lagarde, P. (2008a). Environment around Al, Si and Ca in aluminate and aluminosilicate melts by X-ray absorption spectroscopy at high temperature. *Am. Mineral.*, 93, 228–234.
- Neuville, D. R., Cormier, L., Flank, A. M., Lagarde, P., and Massiot, D. (2004a). Aluminum X-ray absorption near edges structure in minerals and glasses. *Chem. Geol.*, 213, 153–163.
- Neuville, D. R., Cormier, L., and Massiot, D. (2004b). Role of aluminium in peraluminous region in the CAS system. *Geochim. Cosmochim. Acta*, 68, 5071–5079.
- Neuville, D. R., Cormier, L., Montouillout, V., Florian, P., Millot, F., Rifflet, J. C., and Massiot, D. (2008b). Structure of Mg- and Mg/Ca aluminosilicate glasses:  $^{27}\text{Al}$  NMR and Raman spectroscopy investigations. *Am. Mineral.*, 93, 1721–1731.
- Neuville, D. R., Cormier, L., Montouillout, V., and Massiot, D. (2007). Local environment of Al in aluminosilicate glasses: a NMR point of view. *J. Non-Cryst. Solids*, 353, 180–185.
- Neuville, D. R., Cormier, L. R., Flank, A. M., Prado, R. J., and Lagarde, P. (2004c). Na K-edge XANES spectra of minerals and glasses. *Eur. J. Mineral.*, 16, 809–816.
- Neuville, D. R., Henderson, G. S., Cormier, L., and Massiot, D. (2010). Structure of  $\text{CaO}-\text{Al}_2\text{O}_3$  crystal, glasses and liquids, using X-ray absorption at Al L and K edges and NMR spectroscopy. *Am. Mineral.*, 95, 1580–1589.
- Neuville, D. R. and Le Losq, C. (2022). Link between Long range order to macroscopic properties of silicate glasses. In Neuville, D. R., Henderson, G. S., and Dingwell, D. B., editors, *Magmas, Melts, Liquids and Glasses: Experimental Insights. Reviews*

- in Mineralogy & Geochemistry*. GeoScienceWorld, McLean, VA.
- Neuville, D. R. and Mysen, B. O. (1996). Role of aluminium in the silicate network: *in situ*, high-temperature study of glasses and melts on the join  $\text{SiO}_2\text{-NaAlO}_2$ . *Geochim. Cosmochim. Acta*, 60, 1727–1737.
- Parks, G. S. and Huffman, H. M. (1926). Glass as a fourth state of matter. *Science*, 64, 363–364.
- Parks, G. S. and Huffman, H. M. (1927). Studies on glass. I. The transition between the glassy and liquid states in the case of some simple organic compound. *J. Phys. Chem.*, 31, 1842–1855.
- Patrick Royall, C., Williams, S. R., Ohtsuka, T., and Tanaka, H. (2008). Direct observation of a local structural mechanism for dynamic arrest. *Nat. Mater.*, 7(7), 556–561.
- Pradel, A. and Piarristeguy, A. (2022). Thio and selenosilicates, sulfide and selenide counterparts of silicates: similarities and differences. *C. R. Géosci.*, 354(S1), 79–99.
- Shintani, H. and Tanaka, H. (2006). Frustration on the way to crystallization in glass. *Nat. Phys.*, 2, 200–206.
- Sipp, A., Neuville, D. R., and Richet, P. (1997). Viscosity, configurational entropy and relaxation kinetics of borosilicate melts. *J. Non-Cryst. Solids*, 211, 281–293.
- Stebbins, J. F. (2008). Temperature effects on the network structure of oxide melts and their consequences for configurational heat capacity. *Chem. Geol.*, 256, 80–91.
- Stebbins, J. F. and Farnan, I. (1992). Effects of high temperature on silicate liquid structure: a multinuclear NMR study. *Science*, 255, 586–589.
- Stebbins, J. F., Farnan, I., and Xue, X. (1992). The structure, dynamics of alkali silicate liquids: a view from NMR spectroscopy. *Chem. Geol.*, 96, 371–385.
- Stebbins, J. F. and Xu, Z. (1997). NMR evidence for excess non-bridging oxygen in an aluminosilicate glass. *Nature*, 390, 60–62.
- Stebbins, J. F., Zhao, P., Lee, S. K., and Oglesby, J. V. (2001). Direct observation of multiple oxygen sites in oxide glasses: recent advances from triple-quantum magic-angle spinning nuclear magnetic resonance. *J. Non-Cryst. Solids*, 293–295, 67–73.
- Taylor, M. and Brown, G. E. (1979a). Structure of mineral glasses: I. The feldspar glasses  $\text{NaAlSi}_3\text{O}_8$ ,  $\text{KAlSi}_3\text{O}_8$ ,  $\text{CaAl}_2\text{Si}_2\text{O}_8$ . *Geochim. Cosmochim. Acta*, 43, 61–75.
- Taylor, M. and Brown, G. E. (1979b). Structure of mineral glasses. II. The  $\text{SiO}_2\text{-NaAlSiO}_4$  join. *Geochim. Cosmochim. Acta*, 43, 1467–1475.
- Tool, A. Q. and Eichlin, C. G. (1931). Variations caused in the heating curves of glass by heat treatment. *J. Am. Ceram. Soc.*, 14(4), 276–308.
- Vetere, F., Iezzi, G., Perugini, D., and Holtz, F. (2022). Rheological changes in melts and magmas induced by crystallization and strain rate. *C. R. Géosci.*, 354(S1), 227–248.
- Warren, B. E. (1941). Summary of work on atomic arrangement in glass. *J. Am. Ceram. Soc.*, 24, 256–261.
- Warren, B. E. and Bischof, J. (1938). Fourier analysis of X-ray patterns of soda-silica glass. *J. Am. Ceram. Soc.*, 21, 259–265.
- Zachariasen, W. H. (1932). The atomic arrangement in glass. *J. Am. Ceram. Soc.*, 54, 3841–3851.