

# Reply on the comment by X. Xue on "Towards the reconciliation of viscosity change and CO2-induced polymerization in silicate melt" by Yann Morizet, Michael Paris, David Sifré, Ida Di Carlo, Sandra Ory, and Fabrice Gaillard [chemical Geology 458, 38-47]

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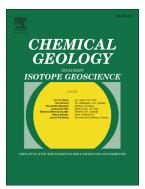
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#### 1. Introduction

There is a growing interest into the study of carbon dioxide (CO<sub>2</sub>) in magmatic systems and capturing the dissolution mechanisms of CO<sub>2</sub> and how it affects the structure of silicate melts constitutes a prime milestone. The comment by Xue and this reply illustrate well the difficulties in addressing this fundamental quest. In the comment, Xue (2019) provides corrected NBO/T values using the <sup>17</sup>O NMR data produced in Morizet et al. (2017c). The comment also shows that the quantitative relationship in between theoretical model for CO<sub>2</sub> dissolution in silicate melt structure and existing NMR data can be difficult to apprehend. Finally, the comment points out the lack of agreement in between the XCO<sub>3</sub><sup>2-</sup> from <sup>17</sup>O NMR and XCO<sub>2</sub> from Raman spectroscopy as reported in Morizet et al. (2017b). In the present reply, we propose a corrigendum to our work (Morizet et al., 2017c) and an addendum pointing towards the future scientific questions to be answered.

Since the seventies, many scientific questions arose concerning the  $CO_2$  behaviour in magmatic systems: 1) the  $CO_2$  solubility in silicate melts as a function of intensive conditions (P,T and  $fO_2$ ) and compositions (e.g. silica content), 2) the way  $CO_2$  molecules dissolve into the silicate melt structure (i.e. speciation) and 3) the influence  $CO_2$  dissolution induces on the silicate melt physical properties (e.g. viscosity, density or electrical conductivity).

For the first two points, many scientific studies have produced large database for  $CO_2$ solubility and subsequent thermodynamic modelling have been built to constrain  $CO_2$ solubility (and degassing) in silicate melts and taking into account the  $CO_2$  dissolution mechanisms (Newman and Lowenstern, 2002; Papale et al., 2006; Iacono-Marziano et al., 2012; Massuyeau et al., 2015). For the third point, the effect  $CO_2$  induces on melt physical properties is still highly debated. Recent works addressed the effect of  $CO_2$  onto silicate melt

density (Seifert et al., 2013) and silicate melt electrical conductivity (Sifré et al., 2014). For viscosity, the effect of  $CO_2$  is unclear and requires further investigations to decipher the exact effect of  $CO_2$  onto silicate melt viscosity. One way to address the effect  $CO_2$  induces onto silicate melt viscosity is to investigate in detail the effect  $CO_2$  dissolution induces onto the silicate melt structure which is considered as a proxy for melt physical properties and in particular viscosity. Our recent research has focused on this aspect and we believe to have reached a substantial answer to the effect of  $CO_2$  onto silicate melt viscosity (Morizet et al., 2017c). The comment by Xue (2019) and our accompanying reply constitute a cornerstone on this hot scientific topic and emphasise that there is a need for future work in the subject.

2. Corrigendum to the evolution of the silicate melt degree of polymerization as a function of  $CO_2$  content from <sup>17</sup>O and <sup>29</sup>Si NMR

We have to admit that the accompanying comment by Xue (2019) is extremely wellconducted from the technical point of view and we recognize that we have not been cautious enough in the calculation of the NBO/T from the obtained <sup>17</sup>O NMR spectra; as well as the apprehension of the theoretical equation for the polymerization of the silicate network upon CO<sub>2</sub> dissolution. In her comment Xue (2019) adequately recalculates the NBO/T values from the original <sup>17</sup>O NMR data published in Morizet et al. (2017c).

In a recent paper (Morizet et al., 2014b), we conducted a similar work on a more polymerized melt composition (nephelinite with NBO/T ~ 1.3) than the one studied in Morizet et al. (2017c; NBO/T ~ 2). Using <sup>29</sup>Si NMR, we showed that CO<sub>2</sub> was polymerizing the aluminosilicate network; however, there was a glitch in the interpretation because mass balance calculation was not satisfied with respect to the initial reaction for CO<sub>2</sub> dissolution mechanism:

$$CO_2^{fluid} + 2Q^n \xrightarrow{melt} \longleftrightarrow CO_3^{2-melt} + 2Q^{n+1-melt}$$
 Eq. 1

Hence, it was then suspected that a more complex mechanism occurred and that <sup>29</sup>Si NMR was not able to identify it. We concluded that <sup>29</sup>Si NMR was probably not the best appropriate method to determine the effect CO<sub>2</sub> induces on the aluminosilicate network. For instance, the distinction in between the effects of the topological changes (Q<sup>n</sup> distribution) and geometrical changes (bond length and angles) (as defined in Massiot et al., 2013) appears difficult to determine with the sole use of <sup>29</sup>Si NMR. We suggested that CO<sub>2</sub> could induce unquantified topological changes and that the measured polymerization using <sup>29</sup>Si NMR did not reflect the true change in the degree of polymerization upon CO<sub>2</sub> solubility. Consequently, we changed our strategy and considered that it would be more appropriate to conduct <sup>17</sup>O NMR investigation (i.e. probing BO and NBO concentrations) in conjunction to <sup>29</sup>Si NMR (i.e. probing Q<sup>n</sup> distribution). As such, the combined use of <sup>17</sup>O and <sup>29</sup>Si NMR spectroscopy would answer the initial problem as to whether or not the Q<sup>n</sup> distribution measured by <sup>29</sup>Si NMR

From our work (Morizet et al., 2017c), we showed that both <sup>17</sup>O and <sup>29</sup>Si NMR results are totally consistent. However, the approach we employed to determine the NBO/T from the <sup>17</sup>O NMR results is not correct as shown by Xue (2019) who proposes the right method to calculate the NBO/T from the subsequent <sup>17</sup>O NMR deconvolution results. Anyhow, our results and interpretations supported the previous recent investigations on the effect of CO<sub>2</sub> onto silicate melt structure (Morizet et al., 2014b; Moussallam et al., 2016).

The second mistake pointed out by Xue (2019) is with regards to the theoretical equation (Eq. 1) for  $CO_2$  dissolution. We lacked clear sighting in handling this theoretical equation. It has two consequences (see Figure 1): 1) the NMR results are in agreement with the theoretical

equation proposed in early work for  $CO_2$  dissolution and corresponding to Eq. 1; and 2)  $CO_2$  does not have a more polymerizing effect than the effect quantified with Eq. 1.

Consequently, we propose a corrigendum to the work in Morizet et al. (2017b), section 5.2 Fig. 3. We followed the description made by Xue (2019) and results are shown in Figure 1. We choose to represent the data in a NBO/T vs  $XCO_2/T$  as it is more appropriate for apprehending the change in the degree of polymerization as a function of  $CO_2$  content: different compositions (with different NBO/T) can be readily compared and the slope for  $CO_2$ dissolution theoretical equation is -2.

For the RB8 sample, we report the NBO/T calculated from the <sup>29</sup>Si NMR data and corrected values of NBO/T from <sup>17</sup>O NMR data; we also added the data for nephelinite (Morizet et al., 2014b) and haplokimberlite (Morizet et al., 2017a). From Figure 1, it can be observed that the whole dataset agrees and that CO<sub>2</sub> incorporation in the melt induces a decreasing in the NBO/T whatever the considered melt compositions. Using the correct calculation, the NBO/T derived from <sup>17</sup>O NMR is in agreement with the <sup>29</sup>Si NMR results. Hence, the distribution of Q<sup>n</sup> species as determined by <sup>29</sup>Si NMR can be used to constrain the change in the silicate glass degree of polymerization as a function of CO<sub>2</sub> content dissolved as Free Ionic Carbonate  $CO_3^{2-}$  molecular groups. Based on the Q<sup>n</sup> distribution, we proposed the following equation for CO<sub>2</sub> dissolution mechanism in silica undersaturated melts:

$$2.25CO_2^{fluid} + 2.5Q^{1} \xrightarrow{melt} \longleftrightarrow 2.25CO_3^{2-melt} + 0.5Q^{2-melt} + 2.0Q^{3-melt}$$
Eq. 2

We emphasise that <sup>17</sup>O NMR results were only used in a qualitative manner in Morizet et al. (2017c) and the reaction for CO<sub>2</sub> dissolution was constrained by <sup>29</sup>Si NMR results. The Eq. 2 is a combination of several individual reactions such as the one reported in Eq. 1. We think that the use of this global reaction is more representative of the thermodynamic at stake: each individual reaction may not have the same equilibrium constant; the configuration entropy

(i.e. related to viscosity) is proportional to the sum of all  $Q^n$  species molar fraction. The Eq. 2 constitutes a real and unambiguous advance in our knowledge of the CO<sub>2</sub> dissolution in the silicate network structure for depolymerized silicate glass compositions.

The proposed Eq. 2 is equivalent to the theoretical Eq. 1. In Figure 1, we report the linear trends derived from Eq. 1 and for three different compositions (i.e. NBO/T = 1, 2 and 3). The change in the NBO/T as a function of CO<sub>2</sub> content from the theory and NMR data are in agreement; at least for RB8 (Morizet et al., 2017c) and HK (Morizet et al., 2017a) data. It suggests that the measured change in the glass polymerization degree strictly follows the theoretical change in the degree of polymerization imposed by Eq. 1. It seems that data obtained for nephelinite composition (<sup>29</sup>Si NMR, Morizet et al., 2014b) departs from the theory imposed by Eq. 1; which is only valid if CO<sub>2</sub> dissolves as Free Ionic Carbonates that is the case for strongly depolymerized RB8 and HK compositions. The off-trend pattern for nephelinite data could be explained by a different CO<sub>2</sub> dissolution mechanism owing to the higher degree of polymerization for this glass composition (NBO/T  $\sim$  1). For moderately depolymerized compositions, several CO<sub>2</sub> dissolution mechanisms can be evoked. We suspect that Free Ionic Carbonates are formed but accompanied by other types of  $CO_3^{2-}$  such as network carbonates and NBO carbonates (Brooker et al., 2001); however, these different types of  $CO_3^{2-}$  molecular groups do not necessarily involve an increase in the degree of polymerization and therefore cannot explain the steeper slope observed in Figure 1 for nephelinite <sup>29</sup>Si NMR data. One alternative explanation lies in the difficulty to interpret quantitatively the <sup>29</sup>Si NMR data: for example, we know that the  $\delta_{iso}$  of each individual  $Q^n$ species is modified by the presence of surrounding Al atoms (Engelhardt et al., 1985; Hiet et al., 2009). As a result, if the question of the  $CO_2$  effect onto the degree of polymerization is clearly sorted out for depolymerized glass composition and that reconciliation to the change in

physical properties has been proposed; we believe that the question still remains opened for glass compositions demonstrating several CO<sub>2</sub> dissolution mechanisms.

# 3. Quantification of $CO_3^{2-}$ species with <sup>17</sup>O NMR spectroscopy

In our recent work (Morizet et al., 2017b), using <sup>17</sup>O NMR spectroscopy, we have shown that the peak intensity at  $\sim$ +150 ppm was linearly correlated to the total CO<sub>2</sub> content determined by Raman spectroscopy (Morizet et al., 2013). The observed correlation helped us to assign the <sup>17</sup>O NMR peak located at ~+150 ppm to carbonate environments in an unambiguous manner. We should point out that this was the first piece of evidence for  $CO_3^{2-}$  dissolved in silicate glasses by <sup>17</sup>O NMR. The proposed assignment was later confirmed in our work (Morizet et al., 2017a) using  $\{^{13}C\}$ - $^{17}O$  J-HMQC NMR technique. In this correlation (based on 7 data points), we determined a slope of 0.8. As reported in Morizet et al. (2017b) the XCO<sub>3</sub><sup>2-17</sup>O NMR cannot be straightforwardly related to XCO<sub>2</sub> Raman mainly because the quantitative calculations are dependent on too many parameters as pointed out by Xue (2019). We doubt that the CO<sub>2</sub> concentration determined by Raman can be put into question as most of the samples were also analysed using elemental analyser which is an absolute method (Moussallam et al., 2015). Instead, we believe that the quantification of  $CO_3^{2-}$  species using <sup>17</sup>O NMR spectra acquired on a 500 MHz NMR spectrometer is inherently limited. Due to the broadening of the peak at ~+150 ppm and its relatively low intensity, the quantification of  $CO_3^{2-}$  species is certainly accompanied by a substantial error. The corollary problem is that it will induce an error in the derived oxygen species concentrations (NBO and BO); however, considering the large scatter observed in Figure 1 and the range of investigated  $CO_2$ , the quantitative error will have a limited impact. The reported trend in Morizet et al. (2017a)

using <sup>17</sup>O NMR can only be used in a qualitative manner and quantitative application will necessitate more thorough investigation.

#### 4. Future work and perspectives

As pointed out in the introduction, obtaining a clear vision of the  $CO_2$  dissolution mechanisms in silicate melt structure is a prerequisite for the understanding of the effect  $CO_2$  dissolution induces on the silicate melt physical properties such as viscosity or electrical conductivity. It appears clear from the subsequent discussion, and the recent work we have published as well as the decades of knowledge on the  $CO_2$  behaviour in silicate melts that there are still discoveries to be made. In particular, the work we accomplished (Morizet et al., 2017a, b, c) is only a small part of the iceberg as it focuses on particular melt compositions which are depolymerized (i.e. low silica content),  $CO_2$ -rich and analogous to natural magmas which are very peculiar on Earth. As mentioned earlier, the question of the effect of  $CO_2$  dissolution on the silicate melt structure for compositions having multiple  $CO_2$  dissolution mechanisms (Free Ionic Carbonates, Network Carbonates and NBO Carbonates) is still to be determined.

Previous NMR studies (Kohn et al., 1991; Brooker et al., 1999; Morizet et al., 2014a, 2015) have commonly used <sup>13</sup>C NMR approach to investigate the CO<sub>2</sub> behaviour in silicate melt. Pioneer work by Kohn et al. (1991) used the <sup>13</sup>C NMR to investigate the CO<sub>2</sub> speciation in glasses when CO<sub>2</sub> is dissolved as both  $CO_2^{mol}$  and  $CO_3^{2-}$  groups. Considerable progresses were conducted with the work of Brooker et al. (1999, 2001) which stated that several types of carbonates groups are present in silicate glasses; with the type of  $CO_3^{2-}$  depending on the silicate glass composition. They clearly demonstrated that network carbonates (T-Carb.-T) are identified below +165 ppm in the <sup>13</sup>C NMR spectrum and the chemical shift being dependent on the nature of the cation (e.g. Si or Al) in the tetrahedral unit. This assignment was also later

confirmed for CO<sub>2</sub>-bearing phonolitic glasses (Morizet et al., 2002). We proposed an assignment for <sup>13</sup>C NMR signature above +165 ppm to NBO-Carb.  $M^{n+}$  or free carbonate groups (Morizet et al., 2014a). These carbonate groups were also suggested to be present in depolymerized composition from Brooker et al. (2001). However, the discrimination in between NBO-Carb  $M^{n+}$  and  $CO_3^{2-} M^{n+}$  was impossible using <sup>13</sup>C NMR and we had to access this information by alternative means: {<sup>13</sup>C}-<sup>17</sup>O J-HMQC NMR technique (Morizet et al., 2017a). The recent work by Xue et al. (2017) combining <sup>13</sup>C NMR and Ab-Initio calculations went further in identifying the different  $CO_3^{2-}$  species and investigated the change in the  $CO_3^{2-}$  molecular configuration for a wide range of glass compositions (i.e. from fully polymerized to strongly depolymerized).

We have conducted a simple approach which is to report the change in the peak centroid of the carbonate <sup>13</sup>C NMR peak (Figure 2) as a function of composition expressed as wt.% Al<sub>2</sub>O<sub>3</sub> + SiO<sub>2</sub> for simplicity. We considered <sup>13</sup>C NMR spectra for compositions ranging from molten carbonates to depolymerized haplokimberlites (Moussallam et al., 2016) and to fully polymerized jadeite (Morizet et al., 2001), nepheline and albite (Brooker et al., 1999) having varying Al/Si ratio. We observe that the <sup>13</sup>C peak centroid is slightly changing (~2 ppm) with increasing Al<sub>2</sub>O<sub>3</sub> + SiO<sub>2</sub> content up to 60 wt.% Al<sub>2</sub>O<sub>3</sub> + SiO<sub>2</sub>. The slight decrease in the <sup>13</sup>C ppm shift from 0 to 60 wt.% Al<sub>2</sub>O<sub>3</sub> + SiO<sub>2</sub> is then followed by a brutal decrease in the <sup>13</sup>C peak centre. This dichotomy might indicate the change in the CO<sub>3</sub><sup>2-</sup> dissolution mechanisms from Free Ionic Carbonates (and possibly NBO-Carb. M<sup>n+</sup>) to Network Carbonates (T-Carb.-T). The brutal change in the <sup>13</sup>C peak centre also seems to mark the limit between slightly depolymerized (NBO/T ~ 0.2 for phonolite, Morizet et al., 2002) to fully polymerized (NBO/T ~ 0 for jadeite, Morizet et al., 2001; 2007). In the Network carbonate region, Brooker et al. (1999) made the distinction in between the different types of carbonates for nepheline, jadeite and albite which have different Al/Si ratio. However, within the Free Ionic Carbonates

region in Figure 2 which represents the majority of natural magmatic systems in fact, a variation of ~2 ppm in the <sup>13</sup>C NMR spectrum (the assumed analytical error is on the order of  $\pm 0.5$  ppm) is subtle. Using in conjunction Ab-Initio calculations for NMR parameters and various NMR approaches (i.e. <sup>17</sup>O and <sup>13</sup>C) represents the ideal solution to investigate the  $CO_3^{2-}$  molecular environments and the recent works accomplished have demonstrated it (Morizet et al., 2017a,b,c; Xue et al., 2017).

#### References:

Brooker, R.A., Kohn, S.C., Holloway, J.R., McMillan, P.F., Carroll, M.R., 1999. Solubility, speciation and dissolution mechanisms for CO<sub>2</sub> in melts on the NaAlO<sub>2</sub>-SiO<sub>2</sub> join. Geochim. Cosmochim. Acta 63, 3549-3565.

Brooker, R.A., Kohn, S.C., Holloway, J.R., McMillan, P.F., 2001. Structural controls on the solubility of  $CO_2$  in silicate melts. Part II: IR characteristics of carbonate groups in silicate glasses. Chem. Geol. 174, 241–254.

Engelhardt, G., Nofz, M., Forkel, K., Wihsmann, F.G., Magi, M., Samoson, A., Lippmaa, E., 1985. Structural studies of calcium aluminosilicate glasses by high-resolution solid-state <sup>29</sup>Si and <sup>27</sup>Al magic angle spinning nuclear magnetic resonance. Phys. Chem. Glasses 26, 157–165.

Hiet, J., Deschamps, M., Pellerin, N., Fayon, F., Massiot, D., 2009. Probing chemical disorder in glasses using silicon-29 NMR spectral editing. Phys. Chem. Chem. Phys. 11, 6935–6940.

Iacono-Marziano, G., Morizet, Y., Le Trong, E., Gaillard, F., 2012. New experimental data and semi-empirical parameterization of H<sub>2</sub>O-CO<sub>2</sub> solubility in mafic melts. Geochim. Cosmochim. Acta 97, 1-23.

Kohn, S.C., Brooker, R.A., Dupree, R., 1991. <sup>13</sup>C MAS NMR: A method for studying CO<sub>2</sub> speciation in glasses. Geochim. Cosmochim. Acta 55, 3879-3884.

Massiot, D., Messinger, R.J., Cadars, S., Deschamps, M., Montouillout, V., Pellerin, N., Veron, E., Allix, M., Florian, P., Fayon, F., 2013. Topological, geometric, and chemical order in materials: Insights from Solid-State NMR. Acc. Chem. Res. 46, 1975–1984.

Massuyeau, M., Gardès, E., Morizet, Y., Gaillard, F., 2015. A model for the activity of silica along the carbonatite-kimberlite-melilitite-basanite melt compositional joint. Chem. Geol. 418, 206–216.

Morizet, Y., Kohn, S.C., Brooker, R.A., 2001. Annealing experiments on CO<sub>2</sub>-bearing jadeite glass: an insight into the true temperature dependence of CO<sub>2</sub> speciation in silicate melts. Min. Mag. 65, 701-707.

Morizet, Y., Brooker, R.A., Kohn, S.C., 2002. CO<sub>2</sub> in haplo-phonolite Melt: Solubility, speciation and carbonate complexation. Geochim. Cosmochim. Acta 66, 1809-1820.

Morizet, Y., Nichols, A.R.L., Kohn, S.C., Brooker, R.A., Dingwell, D.B., 2007. The influence of H<sub>2</sub>O and CO<sub>2</sub> on the glass transition temperature: insights into the effects of volatiles on magma viscosity. Eur. J. Mineral. 19, 657–669.

Morizet, Y., Brooker, R.A., Iacono-Marziano, G., Kjarsgaard, B., 2013. Quantification of CO<sub>2</sub> dissolved in silicate glasses of various compositions with Micro-Raman spectroscopy. Am. Mineral. 98, 1788–1802.

Morizet, Y., Paris, M., Gaillard, F., Scaillet, B., 2014a. Carbon dioxide in silicaundersaturated melt. Part I: The effect of mixed alkalis (K and Na) on CO<sub>2</sub> solubility and speciation. Geochim. et Cosmochim. Acta 141, 45–61.

Morizet, Y., Paris, M., Gaillard, F., Scaillet, B., 2014b. Carbon dioxide in silicaundersaturated melt. Part II: Effect of  $CO_2$  on quenched glass structure. Geochim. et Cosmochim. Acta 144, 202–216.

Morizet, Y., Vuilleumier, R., Paris, M., 2015. A NMR and molecular dynamics study of CO<sub>2</sub>bearing basaltic melts and glasses. Chem. Geol. 418, 89–103.

Morizet, Y., Florian, P., Paris, M., Gailllard, F., 2017a. <sup>17</sup>O NMR evidence of free ionic clusters M<sup>n+</sup> CO<sub>3</sub><sup>2-</sup> in silicate glasses: Precursors for carbonate-silicate liquids immiscibility. Am. Mineral. 102, 1561-1564.

Morizet, Y., Paris, M., Sifré, D., Di Carlo, I., Gaillard, F., 2017b. The effect of Mg concentration in silicate glasses on CO<sub>2</sub> solubility and solution mechanism: Implication for natural magmatic systems. Geochim. et Cosmochim. Acta 198, 115-130.

Morizet, Y., Paris, M., Sifré, D., Di Carlo, I., Gaillard, F., 2017c. Towards the reconciliation of viscosity change and CO<sub>2</sub>-induced polymerization in silicate melts. Chem. Geol. 458, 38-47.

Moussallam, Y., Morizet, Y., Massuyeau, M., Laumonier, M., Gaillard, F., 2015. CO<sub>2</sub> solubility in kimberlite melts. Chem. Geol. 418, 198–205.

Moussallam, Y., Florian, P., Corradini, D., Morizet, Y., Sator, N., Vuilleumier, R., Guillot, B., Iacono-Marziano, G., Schmidt, B.C., Gaillard, F., 2016. The molecular structure of melts along the carbonatite–kimberlite–basalt compositional joint: CO<sub>2</sub> and polymerization. Earth Planet. Science Lett. 434, 129-140.

Newman, S., Lowenstern, J.B., 2002. VOLATILECALC: a silicate melt-H<sub>2</sub>O–CO<sub>2</sub> solution model written in Visual Basic for excel, Source. Comput. Geosci. 28, 597–604.

Papale, P., Moretti, R., Barbato, D., 2006. The compositional dependence of the saturation surface of  $H_2O+CO_2$  fluids in silicate melts. Chem. Geol. 229, 78-95.

Seifert, R., Malfait, W.J., Lerch, P., Sanchez-Valle, C., 2013. Partial molar volume and compressibility of dissolved CO<sub>2</sub> in glasses with magmatic compositions. Chem. Geol. 358, 119-130.

Sifré, D., Gardès, E., Massuyeau, M., Hashim, L., Hier-Majumder, S., Gaillard, F., 2014. Electrical conductivity during incipient melting in the oceanic low-velocity zone. Nature 509, 81–87.

Xue, X., 2019. Comment on "Towards the reconciliation of viscosity change and CO<sub>2</sub>induced polymerization in silicate melts" by Yann Morizet, Michael Paris, David Sifré, Ida Di Carlo, Sandra Ory, and Fabrice Gaillard [Chemical Geology 458 (2017) 38-47] Figure caption:

Figure 1: Change in the NBO/T as a function of  $CO_2$  content (expressed as the molar fraction of  $CO_2$  dissolved in melt divided by the molar fraction of tetrahedral cations). The NBO/T is calculated either from <sup>17</sup>O or <sup>29</sup>Si NMR. Data are from Morizet et al. (2014b) and Morizet et al. (2017a,b,c).

Figure 2: Change in the peak centroid position for carbonate molecular groups determined by  $^{13}$ C MAS NMR spectroscopy as a function of wt.% Al<sub>2</sub>O<sub>3</sub> + SiO<sub>2</sub>. Peak position in ppm has been determined for various melt compositions ranging from molten carbonates to silica-rich jadeite and albite.

Solution

Figure 1

 Figure 2

#### **Declaration of interests**

 $\boxtimes$  The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

The authors declare the following financial interests/personal relationships which may be considered as potential competing interests: