



## **The influence of CO<sub>2</sub> and H<sub>2</sub>O on the glass transition in synthetic phonolite and jadeite.**

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CO<sub>2</sub> plays an important role in degassing and eruptive processes. Recent spectroscopic studies established that CO<sub>2</sub> speciation is strongly temperature dependent and that CO<sub>2</sub> speciation preserved in quenched glasses is different from the true CO<sub>2</sub> speciation observed in the melts. It is yet not clear what is the effect of CO<sub>2</sub> on the glass transition, T<sub>g</sub>: either CO<sub>2</sub> has an influence on T<sub>g</sub>, or the CO<sub>2</sub> speciation is decoupled from the bulk silicate melt structural relaxation.

We conducted calorimetric measurements to investigate T<sub>g</sub> on synthetic phonolite and jadeite glasses synthesised in piston-cylinder at 1300 < T < 1550 °C, 1.0 < P < 2.5 GPa. CO<sub>2</sub> and H<sub>2</sub>O concentrations were measured using LECO bulk analyser and FTIR. Volatile concentrations studied is CO<sub>2</sub> up to 2.29 wt.%, H<sub>2</sub>O up to 5.49 wt.%.

For both compositions, H<sub>2</sub>O has a large effect in reducing T<sub>g</sub>, but CO<sub>2</sub> appears to have little or no effect on T<sub>g</sub>. For all range of H<sub>2</sub>O content, T<sub>g</sub> is decreased exponentially from 870 to 523K and 1036 to 636K for phonolite and jadeite respectively, regardless of the CO<sub>2</sub> content. For all range of CO<sub>2</sub> content and almost constant H<sub>2</sub>O content, only small variations in T<sub>g</sub> are observed and an average value of 880K and 903 was derived for phonolite and jadeite respectively.

These results suggest CO<sub>2</sub> contribution to the change in physical properties is negligible compared to H<sub>2</sub>O. It is also a strong evidence of the decoupling of CO<sub>2</sub> speciation from the bulk silicate melt structural relaxation.