## The T-X phase diagram $Na_2CO_3$ -CaCO<sub>3</sub> at 3 GPa

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Phase relations in the system Na<sub>2</sub>CO<sub>3</sub>-CaCO<sub>3</sub> have been studied at 3 GPa and 800–1525 °C. The system has one intermediate compound, Na<sub>2</sub>Ca<sub>3</sub>(CO<sub>3</sub>)<sub>4</sub>, at 800 °C, and two intermediate compounds, Na<sub>2</sub>Ca(CO<sub>3</sub>)<sub>2</sub> and Na<sub>2</sub>Ca<sub>3</sub>(CO<sub>3</sub>)<sub>4</sub>, at 850 °C (Fig. 1a). CaCO<sub>3</sub> crystals recovered from experiments at 950 °C and 1000 °C are aragonite and calcite, respectively. Maximum solid solution of CaCO<sub>3</sub> in Na<sub>2</sub>CO<sub>3</sub> is 20 mol% at 850 °C. The Na-carbonate–Na<sub>2</sub>Ca(CO<sub>3</sub>)<sub>2</sub> eutectic locates near 860 °C and 56 mol% Na<sub>2</sub>CO<sub>3</sub>. Na<sub>2</sub>Ca(CO<sub>3</sub>)<sub>2</sub> melts incongruently to Na<sub>2</sub>Ca<sub>3</sub>(CO<sub>3</sub>)<sub>4</sub> and a liquid containing about 51 mol% Na<sub>2</sub>CO<sub>3</sub> at ~ 880 °C. Na<sub>2</sub>Ca<sub>3</sub> (CO<sub>3</sub>)<sub>4</sub> disappears above 1000 °C via incongruent melting to calcite and a liquid containing about 43 mol% Na<sub>2</sub>CO<sub>3</sub>. At 1050 °C, the liquid, coexisting with Na-carbonate, contains 87 mol% Na<sub>2</sub>CO<sub>3</sub>. Na<sub>2</sub>CO<sub>3</sub>. Na-carbonate remains solid up to 1150 °C and melts at 1200 °C (Fig. 1c). The Na<sub>2</sub>CO<sub>3</sub> content in the liquid coexisting with calcite decreases to 15 mol% as temperature increases to 1300 °C. CaCO<sub>3</sub> remains solid up to 1500 °C and melts at 1525 °C (Fig. 1d).

Considering the present and previous data, a range of the intermediate compounds on the CaCO<sub>3</sub>-Na<sub>2</sub>CO<sub>3</sub> join changes as pressure increases in the following sequence: Na<sub>2</sub>Ca(CO<sub>3</sub>)<sub>2</sub>, Na<sub>2</sub>Ca<sub>2</sub>(CO<sub>3</sub>)<sub>3</sub> (0.1 GPa)  $\rightarrow$  Na<sub>2</sub>Ca(CO<sub>3</sub>)<sub>2</sub>, Na<sub>2</sub>Ca<sub>3</sub>(CO<sub>3</sub>)<sub>4</sub> (3 GPa)  $\rightarrow$  Na<sub>4</sub>Ca(CO<sub>3</sub>)<sub>3</sub>, Na<sub>2</sub>Ca<sub>3</sub>(CO<sub>3</sub>)<sub>4</sub>, Na<sub>2</sub>Ca<sub>4</sub>(CO<sub>3</sub>)<sub>5</sub> (6 GPa) (Fig. 1b). Thus, the nyerereite stability field extends to the shallow mantle pressures, whereas the shortite stability field terminates somewhere between 0.1 and 3 GPa. Consequently, findings of nyerereite and shortite among daughter phases in the melt inclusions in olivine from the sheared garnet peridotites are consistent with their mantle origin.

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Fig. 1. (a) The system  $Na_2CO_3$ -CaCO<sub>3</sub> at 3 GPa. (b) Comparison with previous data at 0.1 GPa (Cooper et al. 1975) and 6 GPa (Shatskiy et al. 2013). (c)  $Na_2CO_3$  melting. (d) CaCO<sub>3</sub> melting. Arg = aragonite; Cal = calcite;  $Na_2$  = solid solution of CaCO<sub>3</sub> in  $Na_2CO_3$ ;  $Na_4Ca = Na_4Ca(CO_3)_3$ ;  $Na_2Ca = Na_2Ca(CO_3)_2$ ;  $Na_2Ca_3 = Na_2Ca_3(CO_3)_4$ ;  $Na_2Ca_4 = Na_2Ca_4(CO_3)_5$ ; L = liquid; F = CO<sub>2</sub> fluid.

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