Phase relations and chemistry of richteritic amphiboles in the system $K_2O-Na_2O-CaO-MgO-Al_2O_3 SiO_2-H_2O$ (KNCMASH)

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Introduction

Knowledge of alkali-amphibole and phlogopite stability plays a key-role in the understanding of water storage and recycling in the upper mantle. The few experimental data so far available at P >5.0 GPa yield an upper pressure stability of < 9GPa at 1200°C for pure phlogopite and 13-14 GPa at 1200°C for pure KNaCa-richterite KNaCaMg₅Si₈O₂₂(OH)₂ (Trønnes 1990). In phlogopite-bearing natural peridotites Trønnes et al. (1988) observed growth of KNaCa-richterite at the expense of phlogopite at about 10 GPa and 1000-1200°C. Sudo & Tatsumi (1990) stabilized pure K₂CaMg₅Si₈O₂₂(OH)₂ at 13 GPa and 1000°C. The present study has been undertaken to better understand phase relations of richterite in the presence of Na in P, T and composition space as this may reduce the upper stability limit of

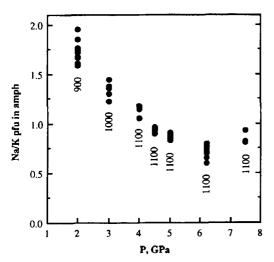


FIG. 1a. Na/K ratio of amphiboles from 2.0 to 7.5 GPa data points are individual amphibole analyses normalized to 24 O and 2 OH. Numbers below datapoints refer to experimental temperatures.

richterite (Sweeney *et al.* 1993). We also investigate any PT-dependent changes in chemistry of amphibole and coexisting phases.

Methods

Experiments were carried out with a Boyd & England-type piston cylinder at $P \leq 3.0$ GPa and a Walker-type multi anvil device from 3.5 to 8.0 GPa. Mineral/phase transformations used for calibrations include α -quartz-coesite, coesite-stishovite, fayalite- γ -spinel, garnet-perovskite in CaGeO₃ and garnet-ilmenite in CdGeO₃. For cold calibration Bi phase transitions were used. Pressure accuracy is believed to be \pm 0.2 GPa between 3.0 and 8.0 GPa and T-precision to be \pm 20°C at 1250°C.

Two different starting compositions were used: An oxide-mix with $Na_2O + K_2O/Al_2O_3 = 1.71$ representative of MARID (Dawson & Smith 1977) and metasomatic phlogopite-K-richterite perido-

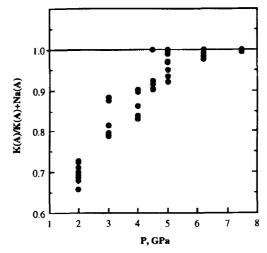


FIG. 1b. Alkali-ratio on A-site of amphiboles from 2.0 to 7.5 GPa. Experimental temperatures as in fig. 1a.

tites (mix1) and a mix of prc-synthesized phlogopite + omphacite + diopside + orthopyroxene with $Na_2O + K_2O/Al_2O_3 = 1.0$ which is thought to represent slightly metasomatically altered peridotites (mix2). These bulk systems are labelled peralkaline and subalkaline respectively.

Results

Preliminary results show KNaCa-richterite stability to be critically dependent on the alkalialumina ratio. In the peralkaline starting composition (mix1) amphibole is stable from 2.0 to 7.5 GPa at 1100°C and coexists with phlogopite + clinopyroxene \pm olivine. At 6.2 to 7.5 GPa Carich garnet appears as an additional phase stable with KNaCa-richterite. At 4 GPa amphibole disappears at 1200°C slightly below the solidus; melting starts between 1200°C and 1225°C, at 1250°C 30-50 % strongly peralkaline melt is present as extremely fine-grained quench. Broadbeam analyses yield approx. 7 wt.% Al₂O₃, 15 wt.% K₂O, 5 wt.% Na₂O and 42-46 wt.% SiO₂.

Amphibole and coexisting phases show systematic changes in chemistry throughout the pressure-interval investigated: most obvious is a decrease in Na/K ratios of amphiboles from > 1.5 at 2.0 GPa to < 0.85 at 6.2 GPa (Fig.1a); with the incoming of garnet at 6.2 GPa Na/K increases slightly. With increasing pressure Na is continuously replaced by K on the A-site (Fig.1b) with K also entering the M(4)-site at 6.2 GPa. With increasing pressure coexisting phlogopite shows a systematic decrease in Na and a slight increase in total Al, clinopyroxene exhibits an increase in jadeite-component.

In mix2 KNaCa-richterite first appears at 6.0 GPa/1100°C. At P < 6.0 GPa the stable assemblage is phlogopite + clinopyroxene + orthopyroxene + garnet. Amphibole is formed by phlogopite-breakdown according to

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phlogopite + enstatite + clinopyroxene =
KNaCa-richterite + garnet
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Preliminary data yield a negative slope for this reaction which is in accordance with results from Sudo & Tatsumi (1990).

References

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