Calcium isotope compositions of mantle pyroxenites and peridotites in Hannuoba, China: Insights into mantle Ca isotope heterogeneity

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Melt-peridotite interaction, caused by asthenosphere upwelling or recycling of crustal materials, could redistribute major elements such as Ca and form pyroxenites that produces a highly inhomogeneous mantle in chemical compositions. Whether and to which extent melt-peridotite reaction leads to heterogeneity in Ca isotopic compositions $(\delta^{44/40}Ca)$ of the mantle rocks are unclear but important to understand the origins of considerable variation of $\delta^{44/40}$ Ca in mantle-derived rocks. Here we report $\delta^{44/40}$ Ca of diverse types of pyroxenite xenoliths (spinel pyroxenites, phlogopitebearing spinel clinopyroxenites and garnet pyroxenites) and surrounding peridotites from Hannuoba, North China Craton. The spinel pyroxenites are cumulates of asthenospherederived silicate melts, and the garnet pyroxenites are the reaction product of peridotite with silicate melt derived from recycled crust material while Phl-bearing spinel clinopyroxenites were formed by metasomatism of peridotite with fluid-rich silicate melt. Each pair (N=8) of garnet pyroxenite and reactive lherzolite show no measurable difference in $\delta^{44/40}$ Ca (0.86‰ to 0.98‰ versus 0.87‰ to 0.93‰), implying that addition of recycled silicate component could not fractionate Ca isotopes within the analytical uncertainty (0.14‰, 2sd). The spinel pyroxenites and Phl-bearing spinel clinopyroxenites have similar $\delta^{44/40} Ca$ (0.90±0.06‰ and 0.96±0.05‰, respectively). The indistinguishable $\delta^{44/40}Ca$ values among these different pyroxenites suggest no measurable Ca isotope variations during silicate melt-peridotite interaction and fractional crystallization, even if recycled silicate materials are involved. The results support the notion that mantle source irrespective of peridotites and pyroxenites typically show uniform $\delta^{44/40}$ Ca. The observed Ca isotope variations in the mantle-derived rocks may result from other factors such as kinetic disequilibrium processes or carbonate components in the source.