

Deep EMI source signature inferred from geochemistry of ultrapotassic volcanic rocks in eastern CAOB: evidences of Re-Os and Li isotopes

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It has been a long-standing debate as to whether the origin of enriched mantle end-member EMI is attributed to metasomatized ancient subcontinental lithosphere, recycled upper and lower crust, or deep mantle (transition zone and/or lower mantle) mixed with recycled ancient sediments. Previous study has shown that a suite of Pleistocene volcanic rocks from northwest part of NE China constructs a spectrum of potassic-rich to ultrapotassic in lithology, in which Ma'anshan leucitite does show the geochemical characteristics as an end-member. Major chemistry shows its $K_2O > 9\%$, $K_2O/Na_2O > 3$. The geochemical data indicate that these rocks are highly enriched in LILE and REE, with extremely fractionated LREE/HREE ratios, and relative depletion of U, Th, Nb, Ta, Sr. The $^{87}Sr/^{86}Sr$ (~ 0.70558), ϵNd (~ -12) and very low $^{206}Pb/^{204}Pb$ (~ 16.34), $^{207}Pb/^{204}Pb$ (~ 15.27) demonstrate a typical EM I affinity, or to define as a LOMU signature, which is similar to other leucite-bearing ultrapotassic volcanic rocks elsewhere in the world (Leucite Hill, Crazy Mts. and Smoky Butte). Recent Re-Os isotopic study on leucitites reveals that most $^{187}Os/^{188}Os$ ratios are higher than 0.1270, the expected value for SCLM. Meanwhile, preliminary study on Li and Mg isotopes, together with $^3He/^4He$ values (Lai *et al.*, 2005) indicate a clear signature of dehydration in the source region, and multiple metasomatism. Based on above geochemical observations and thermodynamic calculation (Kuritani *et al.*, 2013), as proposed by Murphy *et al.*, (2002) and Rapp *et al.*, (2008), we prefer a deep mantle source for these ultrapotassic rocks, which has been isolated for 2-3 Ga and experienced multiple and distinctive mantle processes, such as ancient deep subduction, dehydration, melts/fluids metasomatism, and recycling geodynamics involved.

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