Mesozoic high-Mg andesites from the Daohugou area, Inner Mongolia: Upper-crustal fractional crystallization of parental melt derived from metasomatized lithospheric mantle wedge

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Abstract

Mineral chemistry, major- and trace-element data, zircon U–Pb ages, and Sr–Nd isotopic data are presented for a suite of volcanic rocks from the Daohugou area, Ningcheng City, Inner Mongolia, on the northern margin of the North China Craton. Samples from the suite are of basaltic andesite to rhyolite in composition, with the rocks containing ~60 wt% SiO2. High MgO, Cr, and Ni contents, classify as high-Mg andesites (HMAs). Zircons from a rhyolite yielded weighted mean 206Pb/238U age of 164 ± 1 Ma, indicating that the Daohugou volcanic suite is coeval with the Tiaojishan Formation of northern Hebei and western Liaoning Province. The HMAs have similar enriched-mantle I (EMI)-type isotopic compositions to each other, with low εNd (t) values, moderate (87Sr/86Sr)i ratios, enrichment in LREEs relative to LILEs, and depletion in HFSEs (e.g., Nb, Ta, Ti), indicating formation through upper-crustal fractional crystallization of a common parental magma. The unusually low CaO contents and CaO/FeO ratios of olivine phenocrysts in the HMAs suggest that the parental melt was subduction-related. The results of Rhyolite-MELTS modelling indicates that HMAs may form through upper-crustal fractional crystallization of a parental melt derived from metasomatized lithospheric mantle at crustal depths. The addition of “water” to the cratonic keel may have played a key role in the destruction of the North China Craton.

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1. Introduction

The ancient, cratonic mantle lithosphere, similar to that beneath the Kaapvaal, Siberian, and other Archean cratons, was removed from the base of the Eastern Block of the North China Craton (NCC) during the Mesozoic, and partly replenished with younger, less refractory lithospheric mantle (Fan et al., 2000; Gao et al., 2002, 2004; Griffin et al., 1998; Menzies et al., 1993; Menzies and Xu, 1998; O'Reilly et al., 2001; Xu, 2001; Zhang et al., 2003; Zheng, 1999; Zheng et al., 2001, 2007). As a consequence, the composition, thermal structure, and thickness of the sub-continental lithospheric mantle (SCLM) of the North China Craton changed dramatically, although the mechanisms and timing of the changes are debated (Niu, 2005). The widespread Mesozoic volcanic rocks in eastern China, especially basalt and high-Mg andesites (HMAs), provide a window into mantle sources and processes, including the evolution of the lithospheric mantle beneath the NCC. The term “high-Mg andesite” was first used in reference to a suite of andesitic volcanic rocks from Adak Island in the Aleutians (Kay, 1978). HMAs have elevated Mg numbers (Mg# = 100 × Mg / (Mg + Fe2+)), high abundances of large-ion lithophile elements (LILEs), high Sr/Y ratios, and were previously thought to be derived from partial melting of subducted oceanic crust where the melt interacted with mantle peridotite during ascent (e.g., Defant and Drummond, 1990; Kay, 1978; Liu et al., 2012; Sato et al., 2014). HMAs can, however, be generated by other processes in a range of tectonic settings (e.g., Castillo, 2012), including melting of a metasomatized mantle peridotite (Calmus et al., 2003; Hirose, 1997; Kelemen, 1995; Shirley and Hanson, 1984), and partial melting of dehydrated lower continental crust, with the melts rising and interacting with mantle peridotite (Chung et al., 2003; Gao et al., 2004; Huang et al., 2008; Kay and Kay, 1993; Rapp et al., 2010; Xu et al., 2002). In all three of the above models, the initial partial melting of the mafic crust and/or assimilation of ultramafic rocks occur at mantle depths. However, some HMAs are considered to be derived from crustal-level magma mixing or fractional crystallization
2. Geological setting and sample description

The North China Craton is one of the world’s oldest Archean cratons, preserving crustal remnants as old as 3800 Ma (Liu et al., 1992, 2008; Song et al., 1996). It is surrounded by the Mesozoic Dabie and Sulu orogenic belts to the south and east (Li et al., 1993; Meng and Zhang, 2000), and by the Central Asian Orogenic Belt to the north (Sengör et al., 1999). The craton is divided into the Eastern and Western Blocks by the Trans-North China Orogen/Central Orogenic Belt (Santosh, 2010; Zhai and Santosh, 2011; Zhao et al., 2005; Zhao and Zhai, 2013) (Fig. 1).

Ningcheng City, in Inner Mongolia, is located at the northern margin of the NCC, within the central part of the Yanshanian orogenic belt. The Ningcheng basin basement comprises predominantly Archean and Mesoproterozoic rocks that are unconformably overlain by late Mesozoic Jurassic–Cretaceous volcanic sedimentary rocks (Fig. 1). These well studied late Mesozoic volcano-sedimentary sequences comprise, from the stratigraphic base upwards, the Jiujiangshan Formation, Tiaojishan Formation, and Lower Cretaceous Tuchengzi and Yixian Formations (Liu et al., 2006). The Tuchengzi Formation in western Liaoning is considered to be Late Jurassic (J3tjc) in age (Hong et al., 2017).

Samples were collected from the Daohugou area of Ningcheng, ~80 km south of Chifeng City, Inner Mongolia (Wang et al., 2006), and ~30 km north of Lingyuan City, Liaoning Province, China (Ji and Yuan, 2002). Previous studies focused mainly on the geochemistry of Daohugou strata because of their abundant fauna and fossil assemblages. Therefore, it is suitable for U–Pb dating (Streck et al., 2007).

3. Analytical methods

3.1. Mineral analyses

The major element compositions of minerals were measured at the State Key Laboratory of Geological Processes and Mineral Resources, China University of Geosciences in Wuhan, China, with a JEOL JXA-8100 Electron Microprobe Analyzer equipped with four wavelength-dispersive spectrometers (WDS). The polished thin-sections were firstly coated with a thin conductive carbon film prior to analysis. The precautions suggested by Zhang and Yang (2016) were used to minimize the difference of carbon film thickness between samples and obtain a ca. 20 nm approximately uniform coating. An accelerating voltage of 20 kV, a beam current of 20 nA, and a 2 μm spot diameter were used to analyze minerals. Data were corrected on-line using a modified ZAF (atomic number, absorption, fluorescence) correction procedure. The peak counting time was 10 s for Na, Mg, Al, Si, K, Ca, Fe and 20 s for Ti and Mn. The background counting time was one-half of the peak counting time on the high- and low-energy background positions.

3.2. Zircon U–Pb dating

Zircons were separated from samples by traditional heavy–liquid and magnetic methods and then purified by hand picking under a binocular microscope. Zircon grains were selected and mounted on a double-sided tape, cast in epoxy resin, and polished to expose surfaces suitable for U–Pb dating.

The Cathodoluminescence (CL) images were used to show the internal textures of zircons and to select optimum spots for U–Pb dating. The CL images of zircons were obtained using a CarlZeiss Merlin Compact high resolution scanning electron microscope connected with an Oxford IE250 energy dispersive system and a Gatan Mono CL4 CL system in the Beijing SHRIMP Center, Chinese Academy of Geological Sciences (CAGS), China.

Zircons were dated using the SHRIMP II ion microprobe at the Beijing SHRIMP Center, CAGS, China. Analytical procedures and conditions were similar to those described by Williams (1998). Mass resolution during the analytical sessions was 5000 (1% peak height on the UO-peak of the standard). The intensity of the primary O− ion beam was 3−4 nA. Spot size was ~30 μm in diameter, and each site was rastered for 180 s prior to analysis to remove surface contamination. Reference zircons M257 (U = 840 ppm, Nasdala et al., 2008) and TEMORA 1 (206Pb/238U age = 417 Ma, Black et al., 2003) were used to calibrate the U abundance and U−Pb dating, respectively. TEMORA 1 was analyzed once every 3−4 unknown analyses. Mass analysis at each site consisted of five scans through the mass stations. Common lead correction was applied using the measured 204Pb abundances. Data processing was carried out using the SQUID and ISOPLOT programs (Ludwig, 2001; Ludwig, 2003). Uncertainties of individual analyses are reported at 1σ; mean weighted 207Pb,206Pb ages were calculated at 95% confidence.

3.3. Major and trace element analysis

Rock samples were trimmed to remove altered surfaces and crushed to a grain size of ~200 mesh for major and trace element, and isotopic analyses. Major elements were analyzed using an X−ray fluorescence (XRF; Rikagu RIX 2100) on fused glass disks at Northwest University, China, following Rudnick et al. (2004).

For trace element analyses, samples were digested using a mixture of HF + HNO3 in Teflon bombs and analyzed with an Agilent 7500a ICP-MS at Northwest University, China following Rudnick et al. (2004). Results of the USGS rock standards (BHVO−1, BCR−2 and TEB−1) were used as standards for calibration.
Fig. 1. (a) Sketch map of the major tectonic divisions of China. WB, TNCO and EB denote the Western Block, Trans-North China Orogen and Eastern Block, respectively, of the North China Craton (Zhao et al., 2005). SC, South China Block; YZ, Yangtze Block. (b) Simplified map of the Ningcheng-Lingyuan Basin. (c,) Simplified geological map showing sample locations of the Daohugou volcanic rocks.
AGV–1) indicate accuracies and precisions better than 5% for all elements except for Ni, Ta and Th, which are 5%–10%. The analytical results of these standards are given in Table S2.

3.4. Sr–Nd isotope analysis

Sr, Nd isotopic compositions were determined using MC–ICP–MS (Nu Plasma HR, Nu Instruments, Wrexham, UK) in static mode at Northwest University, China.

For Sr–Nd isotopes, ~100 mg rock power was digested in a sealed Teflon bomb with a mixture of concentrated HNO₃, HF and HClO₄. The sealed bombs were kept in an oven at 190 °C for 48 h. The decomposed samples were then dried at 140 °C followed by adding concentrated HNO₃ and HCl. Sr and Nd (and other REEs) were separated/concentrated using standard chromatographic columns with AG50W–x8 and HDEHP resins following Gao et al. (2004). The measured 143Nd/144Nd and 87Sr/86Sr ratios were normalized to 146Nd/144Nd = 0.7219 and 86Sr/88Sr = 0.1194, respectively. External reproducibility of the isotopic measurement was monitored by repeated analysis of international standards. Repeated analyses of La Jolla along with unknown samples gave an average 143Nd/144Nd of 0.511835 ± 0.000001 (2σ, n = 13), and NBS 987 gave 87Sr/86Sr = 0.710191 ± 0.000002 (2σ, n = 12), which is identical to reference values within analytical error. Analysis of BCR–2 gave 87Sr/86Sr = 0.705003 ± 0.00001 (2σ), 143Nd/144Nd = 0.512622 ± 0.000001 (2σ). Procedural blanks are about ~20 pg and ~40 pg for Nd and Sr, respectively.

4. Analytical data

4.1. Mineral chemistry

4.1.1. Olivine

HMA olivine compositions are listed in Table A1. Olivine crystals have Fo (Fo = 100 × Mg / (Mg + Fe)) values of 78–83, and relatively narrow ranges of CaO contents and CaO/FeO ratios, similar to those within the Xinkailing HMAs of western Liaoning (Fig. 2). They generally exhibit low CaO contents and CaO/FeO ratios. In olivine compositional diagrams (Fig. 2), they plot within the field of subduction-related lavas, and outside the field of non-subduction-related lavas (including mid-ocean-ridge and ocean-island basalts; MORB and OIB) (Fig. 2).

4.1.2. Clinopyroxene

Clinopyroxene phenocrysts in the HMAs are mostly augite (En₄₁–₄₅Fs₁₂–₁₅Wo₄₃–₄₄), with some exhibiting the reverse compositional zoning with high-Fe and low-Mg in the core and high-Mg in the rim (Table S1; Fig. A2). Reverse compositional zoning has also been reported in orthopyroxene phenocrysts in Xinglonggou HMAs from western Liaoning (Gao et al., 2004).

4.2. Geochronology

Seventeen zircon grains from one rhyolite sample located at the upper unit of the Daohugou sequence (sample DHG–57) were dated using SHRIMP II ion microprobe. U–Pb zircon data are summarized in Table S3. The zircons are typically euhedral to subhedral, have Th/U ratios of >0.5, and are characterized by oscillatory zoning (Fig. 3), suggesting a magmatic origin. The zircons have a weighted mean 206Pb/238U age of 164 ± 1 Ma (MSWD = 0.27, n = 17) (Fig. 4).

4.3. Major and trace elements

Representative major- and trace-element data are given in Table 1, with full data provided in Supplementary Table S4. In the total alkalis vs. silica (TAS) diagram (Le Maitre et al., 1989), the samples plot in...
the alkaline field, and consist mainly of trachyandesite and trachyte with minor basaltic trachyandesite and rhyolite (Fig. 5). They have SiO$_2$ and MgO contents of 54.44–74.57 wt% and 0.03–6.05 wt%, respectively. Most samples with SiO$_2$ contents of <60 wt% have high MgO content (>-3 wt%) and Mg$#$ values, similar to the HMAs (Gao et al., 2008). On major-element vs. SiO$_2$ diagrams (Fig. 6), they exhibit broad trends except for an abrupt decrease in Na$_2$O and Al$_2$O$_3$ contents with increasing SiO$_2$ content.

Chondrite-normalized REE patterns are characterized by enrichment in light rare-earth elements (LREEs) relative to heavy REEs (HREEs). Eu anomalies are insignificant (Eu/Eu$^*$ = 0.92–1.17) for all the samples except the rhyolites (Fig. 7a). In the primitive-mantle normalized trace-element diagram (Fig. 7b), the samples display a pronounced depletion in high field strength elements (HFSEs; e.g., Nb, Ta, Ti), and enrichment in large ion lithophile elements (LILEs) relative to LREEs and HFSEs. The samples exhibit positive Ba, K, and Pb anomalies. HMAs display positive Sr anomalies, and samples with SiO$_2$ contents above 60 wt%, show negative Sr anomalies. This can be explained by fractional crystallization of plagioclase, as supported by the positive correlation between Sr and SiO$_2$ at SiO$_2$ contents below ~60 wt% and negative correlation above ~60 wt% (Fig. 8). The Sr content is 21–1489 ppm over the entire sample suite, with the HMAs having the higher concentrations (981–1489 ppm), and Sr/Y ratios are 53–112 (Table 2), similar to adakites (Defant and Drummond, 1990; Kay and
Fig. 6. Major-element vs. SiO$_2$ variation diagrams for the Daohugou volcanic rocks. Fe$_2$O$_3$ is expressed as total Fe. The data show linear trends except for a sharp decrease in Al$_2$O$_3$ and Na$_2$O contents at SiO$_2$ contents above ~60 wt%, indicating the appearance of plagioclase as a liquidus phase.
Fig. 7. (a) Chondrite normalized REE patterns and (b) primitive mantle normalized trace-element patterns for the Daohugou volcanic rocks. Chondrite values are from Taylor and McLennan (1985), and primitive-mantle values are from Sun and McDonough (1989).
However, in a plot of Sr/Y vs. Y (Fig. 11), the HMAs plot in the field of typical arc rocks with higher Y contents (16.6–19.7 ppm). They also have high Cr (173–230 ppm) and Ni (73–133 ppm) contents, similar to those documented from the type locality of HMA by Kay (1978).

4.4. Sr–Nd isotopes

Sr and Nd isotope data are given in Table 2. Initial Sr and Nd isotope compositions were calculated to the time of formation at 160 Ma using the measured Sr–Nd isotopic composition and parent/daughter ratios. The εNd (t) are calculated using \((10,000 \times \frac{[\text{143} \text{Nd}/\text{144} \text{Nd}]_{\text{sample}}}{[\text{143} \text{Nd}/\text{144} \text{Nd}]_{\text{chondrite}} - 1})\), in which, subscript sample and chondrite denote values of sample and chondrite at the time of sample formation, and the \([\text{143} \text{Nd}/\text{144} \text{Nd}]_{\text{chondrite}} = 0.512638\) (Dickin, 1997). The samples display moderate Sr and low Nd isotopic ratios, similar to EMI-type isotope features, with \((87 \text{Sr}/86 \text{Sr}) = 0.705230–0.706402, [\text{147} \text{Nd}/\text{144} \text{Nd}] = 0.511735–0.511944,\) and \(\varepsilon_{\text{Nd}} (t) = -11.8\) to -15.4 (Fig. 12).

5. Discussion

5.1. Fractional crystallization

The Daohugou samples consist of basaltic trachyandesite, trachyandesite, and rhyolite. The strong correlations displayed in the variation diagrams (Fig. 6) could be due either to varying degrees of fractional crystallization from a common parental magma, or to mixing of basaltic melts with crustal melts. The similarities between isotopic compositions and trace-element systematics of the samples do not favor a mixing model. However, their similar isotopic compositions indicate a genetic relationship, with the felsic melts derived from protracted fractional crystallization. The Rhyolite-MELTS (Gualda and Ghiorso, 2015) was used to model Daohugou HMA evolution at varying pressures (1 and 10 kbar), water contents (4% and 0.5%) and oxygen fugacities (NNO, NNO + 1, and NNO + 2, where NNO is the nickel–nickel oxide buffer) (Figs. 9 and 10). The most primitive basaltic rocks (DHG 67) in this study was used as the starting composition (Appendix 1). The results are broadly consistent with fractional crystallization along the liquid lines of descent of olivine, spinel and clinopyroxene, and then feldspar, orthopyroxene, accessory Fe–Ti oxides andapatite at lower pressure (1 kbar), -4 wt% H2O content and higher oxygen fugacities (NNO + 1, and NNO + 2, (Fig. 9). The proportions of crystallized minerals at 1 kbar, with NNO + 2 fugacity and 4 wt% H2O are listed in Appendix 1.

Trace-element variation diagrams (Fig. 8), are also consistent with the fractionation modelling. The trace-element contents of the most primitive basalt (DHG67) were used as the starting composition, with mineral/melt partition coefficients as summarized in Appendix 1. The SiO2 content was on basis of the Rhyolite-MELTS modelling at a pressure of 1 kbar, 4 wt% H2O, and high fO2 (NNO + 2). The fractional crystallization trend is shown in Fig. 8 and the tick marks at 20% intervals represent percentage of crystallization. The modelling results are in good agreement with the HMA compositions. The highly compatible elements, Cr and Ni, are rapidly depleted as MgO contents approach zero as a result of the removal of 8% spinel, 6% olivine, 14% clinopyroxene, 7% orthopyroxene, and 45% plagioclase (Fig. 8). Sr contents and Sr/Y ratios increase with increasing SiO2, reaching a maxima at SiO2 content of ~60 wt%, as a result of the fractionation of olivine, clinopyroxene, and spinel prior to the crystallization of plagioclase, because Sr is
Fig. 9. Chemical variations of rock suites including HMAs from the Daohugou area and results of modeling at low pressure (1 kbar). Rhyolite-MELTS fractionation models of primitive high-Mg basaltic composition (sample DHG67) are given at various oxygen fugacities (NNO, NNO + 1, NNO + 2), and initial H$_2$O contents of 0.5 wt% (undersaturated) and 4 wt% (saturated at 1 kbar). Tick marks at 10% intervals represent the percentage of crystallization.
Fig. 10. Chemical variations in rock suites at high pressure (10 kbar). Rhyolite-MELTS fractionation models of primitive high-Mg basaltic composition (sample DHG67) are given at various oxygen fugacities (NNO, NNO + 1, NNO + 2), and initial H₂O contents of 0.5 wt% (undersaturated) and 4 wt% (undersaturated at 10 kbar). Tick marks at 10% intervals represent the percentage of crystallization. The results of modeling do not reproduce the HMA compositions.
incompatible in spinel, olivine and clinopyroxene, but highly compatible in plagioclase. The observation that Nb content decreases with increasing SiO₂ content may be due to the high compatibility of Nb in Fe–Ti oxides such as ilmenite (Green and Pearson, 1987).

5.2. Petrogenesis of HMAs

Our geochemical data demonstrate that the samples with SiO₂ <60 wt% mostly have relatively high MgO (>3 wt%), Cr and Ni contents and are termed as HMAs. It has been suggested that HMAs reflect the interaction of the partial melt of subducted eclogite with the mantle wedge (Kay, 1978; Defant and Drummond, 1990; Kelemen et al., 2003). However, the relatively high (⁸⁷Sr/⁸⁶Sr)₀ ratios, negative εNd(t) values, and non-ragiogenic Pb isotopic compositions of the Daohugou HMAs do not support their derivation from subducted oceanic crust. Furthermore, the “adakitic” signatures of the Daohugou HMAs are relatively weak in terms of HREE and Y depletion (Fig. 11), indicating that little or no garnet was involved in their petrogenesis. Alternatively, the HMAs could represent primary magmas in equilibrium with mantle peridotite (Calmus et al., 2003; Hirose, 1997; Kelemen, 1995; Shirey and Hanson, 1984; Tatsumi, 1981) or the interaction of partial melts of delaminated lower continental crust and mantle peridotite (Chung et al., 2003; Gao et al., 2004; Huang et al., 2008; Kay and Kay, 1993; Rapp et al., 2010; Xu et al., 2002). It is also possible that the HMAs do not represent primary magma. Fractional crystallization of basaltic magmas has previously been used to explain the petrogenesis of HMAs (e.g., Baja, California Peninsula; Castillo, 2008). HMAs could form by the fractionation of basalts at crustal depths (Chen et al., 2013; Humphreys et al., 2006). Shellnutt and Zellmer (2010) demonstrate that HMAs, which have been regarded as particularly hydrous primary melts generated in equilibrium with mantle peridotite, can form by crystal fractionation from low-H₂O primitive arc basalts in the upper crust as inferred from MELTS modelling. Both of these models may yield the observed EMI-type isotopic compositions of the Daohugou HMAs, with highly fractionated REE patterns, pronounced depletions in HFSEs (e.g., Nb, Ta, and Ti), and enrichments in LILEs relative to LREEs and HFSEs (Fig. 7). The petrogenetic pressure may have influenced the petrogenesis of the Daohugou HMAs. Rhyolite-MELTS modelling indicates that HMAs cannot be produced by fractional crystallization at high pressure (10 kbar) regardless of the relative oxidation state (fO₂) and water content (Fig. 10). Lower pressure (1 kbar) conditions with ~4%H₂O content, and higher fO₂ (NNO + 2) are better able to produce the Daohugou HMA melts (Fig. 9). Shellnutt and Zellmer (2010) have shown that HMAs may be generated by fractional crystallization at NNO + 2 ± 1 log10 unit. Shallow fractionation is also consistent with the observation that the Daohugou HMAs do not display residual garnet signatures. A relatively high H₂O content is also supported by their mineral composition, with olivine phenocrysts of the Daohugou HMAs having unusually low CaO contents and CaO/FeO ratios (Fig. 2), similar to water-rich subduction-related lavas (Fig. 2; Kamenetsky et al., 2006). Comparable characteristics are observed in the Xinkailing HMAs, for which the water contents are estimated to be ~6 wt% and 2.6–6.0 wt%, respectively, based on the effects of water pressure on mineral/melt partition coefficients for Ca and on the olivine liquidus (Hong et al., 2017). The modelling results predict the fractionation of about 15% spinel, 3% olivine, and 11% clinopyroxene (En₄₃₋₄₅Fa₁₂₋₁₅Wo₄₃₋₄₄) to produce a magma with the composition of the Daohugou HMAs. Early fractionation of spinel-structured oxides is key to the formation of HMAs, as it causes an increase in SiO₂ content and associated depletion of iron oxides, with minimal effect on the remaining major elements, producing HMA melts with 58.5 wt% SiO₂, 3.5 wt% MgO. The Daohugou HMAs were therefore most likely derived from shallow-level fractional crystallization from island arc basalts, consequently, crustal differentiation processes may have been important in petrogenesis of HMAs.

The similar isotopic compositions and trace-element pattern of the samples of the present study, combined with the results of quantitative Rhyolite-MELTS modelling, indicate that the samples may be genetically related to each other, and that the magmas which formed the mafic rocks underwent fractionation to produce silicic rocks. The samples show highly fractionated REE patterns with pronounced depletion in HFSE (e.g., Nb, Ta, and Ti) and enrichment in LILE relative to LREE and HFSEs (Fig. 7). Compared with mantle-derived magmatic rocks globally, depletions in Nb, Ta, and Ti are a characteristic of volcanic arc rocks or the “arc signature” exhibited by many continental crustal materials. The negative εNd(t) values and variable initial ⁸⁷Sr/⁸⁶Sr ratios of the samples are consistent with derivation from an EMI-type isotopically enriched mantle source. EMI-type isotopic compositions are evident in >110 Ma basalts (e.g., Gao et al., 2008; Meng et al., 2015; Xu, 2001; Yang and Li, 2008; Zhang et al., 2002) and HMAs (Gao et al., 2004) in the NCC, while the <110 Ma basalts were derived from the asthenosphere (e.g., the Jianguo basalts; Zhang et al., 2003).
the results of Rhyolite-MELTS modelling, the parental melt was most likely hydrous and formed in a subduction-zone setting. Hydrous SiO2-rich fluids or melts released from the subducted slab would have reacted with peridotite in the mantle wedge and induced partial melting, with the melts being enriched in LILEs but depleted in HFSEs (e.g., Nb, Ta and Ti).

The sub-continental mantle lithosphere (SCLM) may become enriched in incompatible elements through protracted mantle metasomatism (O’Reilly and Griffin, 1988). Such metasomatism would not significantly affect major elements, but may result in enrichments in volatiles and the more incompatible elements, leading to enriched isotopic signatures (Niu, 2005). A series of successive subduction events have occurred near the eastern NCC since the Paleozoic, resulting in the addition of large amounts of water into the overlying SCLM (Windley et al., 2010). This is indicated by the presence of Early Cretaceous mafic lavas at the northern, eastern, and southeastern margins of the NCC that originated from a hydrous SCLM (e.g., Hong et al., 2017; Ma et al., 2016; Xia et al., 2013).

In summary, the Daohugou HMAs may have formed by upper-crustal fractional crystallization of arc basalt derived from metasomatized lithospheric mantle.

5.3. Geodynamic significance

The ancient, cratonic mantle lithosphere, similar to that present beneath the Kaapvaal, Siberian, and other Archaean cratons, is inferred to have been removed from the base of the Eastern Block of the NCC in the Mesozoic, and was partly replenished with younger, less refractory lithospheric mantle (Fan et al., 2000; Gao et al., 2002, 2004; Griffin et al., 1998; Menzies et al., 1993; Menzies and Xu, 1998; O’Reilly et al., 2001; Xu, 2001; Zhang et al., 2003; Zheng, 1999; Zheng et al., 2001, 2007). As a consequence, the composition, thermal structure, and thickness of the sub-continental lithospheric mantle (SCLM) of the North China Craton have changed dramatically. However, the mechanisms of this process remain controversial.

The “arc-like”, EMI Sr–Nd isotopic compositions, the composition of olivine, and the results of Rhyolite-MELTS modelling lead to the conclusion that the Daohugou HMAs were produced by upper-crustal fractional crystallization of arc basalt derived from a metasomatized mantle wedge. Previous studies have suggested a hydrous lithosphere, based on the high water content of early Cretaceous mafic lavas and HMAs at the southeastern, eastern, and northern margins of the NCC (Hong et al., 2017; Ma et al., 2016; Xia et al., 2013).

The final closure of the Paleo-Asian Ocean between the North China Craton and Siberian Craton took place in the late Permian (Xiao et al., 2003; Xu et al., 2013), and eastern China has been affected by subduction of the Paleo-Pacific Plate since the late Mesozoic (Niu, 2005; Zhao et al., 1994). Successive subduction events around the eastern NCC may have introduced large amounts of water into the overlying SCLM (Windley et al., 2010; Zhang et al., 2003), resulting in the production of hydrous melts that ascended, weakening the base of the lithosphere and, converting basal lithospheric mantle into asthenospheric mantle (Niu, 2005). These hydrous melts re-fertilized the overlying SCLM (Tang et al., 2013; Hong et al., 2017), possibly accelerating the removal of the SCLM, and leading to destabilization of the craton (Wang et al., 2006; Zhu et al., 2012).

6. Conclusions

Mineral chemistry, geochronology, major- and trace-element compositions and Sr–Nd isotopic compositions of the Mesozoic lavas in Inner Mongolia lead to the following conclusions.

1) Zircon U–Pb dating of a Daohugou rhyolite yields an age of 164 ± 1 Ma, suggesting that the Daohugou volcanic sequence is coeval with the Tiaojishan Formation in northern Hebei and western Liaoning.

2) The Daohugou volcanic rock suite constitutes a complete compositional spectrum from basaltic trachyandesite to rhyolite. All the rocks are enriched in LILEs and LREEs, and depleted in HFSEs, with similar EMI-type Sr–Nd isotopic compositions.
3) All samples have similar isotopic compositions, and trace-element systems, consistent with a genetic link by shallow-level (at 1 kbar pressure) fractional crystallization.

4) Samples containing <60 wt% SiO₂ have the characteristics of HMAs, with high MgO, Cr, and Ni contents. The results of MELTS modeling predicts the fractionation of about 15% spinel, 3% olivine, and 11% clinopyroxene (Erta1–45Fs12–15Wo43–44), to yield the magmatic composition of the Daohugou HMAs under conditions of low pressure (1 kbar), a high H₂O content (4 wt%), and high H₂O (NNO + 2). This indicates upper-crustal fractionation of arc basalt derived from metasomatized lithospheric mantle.

5) The addition of “water” to the crustal keel may have played a major role in the destruction of the NCC.

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