



# Magma mixing origin for high Ba–Sr granitic pluton in the Bayankhongor area, central Mongolia: Response to slab roll-back



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## ABSTRACT

Petrogenesis of high Ba–Sr granitoids is a matter of debate. This paper presents whole-rock geochemical, Sr–Nd isotopic and zircon U–Pb and Hf isotopic data for a suite of high Ba–Sr biotite monzogranite and associated mafic microgranular enclaves (MMEs) from the Ulaan Uul batholith in the Bayankhongor area, central Mongolia, aiming at elucidating the complex petrogenesis of the high Ba–Sr granitoids. The data manifest that magma mixing between a felsic magma derived from sedimentary rocks and a mafic magma derived from Sub-Continental lithospheric mantle (SCLM) may account for the formation of high Ba–Sr granitoids. The host granite and MMEs yielded U–Pb ages of  $546 \pm 3$  Ma and  $547 \pm 3$  Ma, respectively, suggesting that the mafic and felsic magmas were coeval. Rocks of the host granitoids possess high SiO<sub>2</sub> and low MgO contents and display peraluminous high-K calc-alkaline and shoshonitic characteristics. Their relatively high Ba, Sr but low HREE (Yb < 2 ppm) resemble those of typical high Ba–Sr granitoids. Rocks from the granitoids have crust-like Sr–Nd isotopic compositions [ $(^{87}\text{Sr}/^{86}\text{Sr})_i = 0.7059\text{--}0.7065$ , and  $\epsilon\text{Nd}(t) = -1.8$  to  $-2.1$ ], and most zircons from the host rocks show negative  $\epsilon\text{Hf}(t)$  values ( $-0.63$  to  $-11.9$ ) with two-stage Hf model ages of 1.54–2.25 Ga. The MMEs have low SiO<sub>2</sub> and high MgO concentrations, enriched Sr–Nd isotopic compositions [ $(^{87}\text{Sr}/^{86}\text{Sr})_i = 0.7060\text{--}0.7066$ , and  $\epsilon\text{Nd}(t) = -1.6$  to  $-2.1$ ], and most zircons from the MMEs have positive  $\epsilon\text{Hf}(t)$  values ( $+0.7$  to  $+2.6$ ), with late Mesoproterozoic single-stage Hf model ages (1.05–1.12 Ga), suggesting an origin of late Mesoproterozoic SCLM. We interpret that the Ulaan Uul batholith has a mixed origin, i.e. partial melting of sedimentary rocks triggered by upwelling of lithospheric mantle-derived magma (represented by MMEs) under an extensional scheme, and subsequent mixing of the mantle- and crust-derived melts resulted in the high Ba–Sr granitoids. We speculate that the late Neoproterozoic to early Cambrian extensional environment in the Bayankhongor area was caused by the roll-back of subducting Bayankhongor oceanic lithosphere.

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## 1. Introduction

High Ba–Sr granitoids were originally recognized by Tarney and Jones (1994) in the Scottish Caledonides and received wide attention (Eklund et al., 1998; Fowler et al., 2001, 2008; Qian et al., 2003; Choi et al., 2009; Yuan et al., 2010; Bruand et al., 2014). Rocks of this kind are characterized by high Ba (>500 ppm) and Sr (>300 ppm), which are distinct from typical I-, S- and A-type granitoids (Tarney and Jones, 1994). Petrogenesis of high Ba–Sr granitoids has been extensively studied, but no consensus has been

reached, and a variety of petrogenetic models have been proposed, including: (1) partial melting of subducted ocean plateaus (Tarney and Jones, 1994); (2) partial melting of mafic lower crust (Ye et al., 2008; Choi et al., 2009); (3) partial melting of enriched lithospheric mantle metasomatized by asthenosphere-derived carbonatitic melts or subduction-related fluids and/or melts (Eklund et al., 1998; Fowler et al., 2001, 2008; Peng et al., 2013). High Ba–Sr granitoids commonly contain mafic microgranular enclaves (MMEs) which have complex origins such as restites (Chappell and White, 1992; White et al., 1999), cumulates formed by early-stage crystallization (Didier and Barbarin, 1991; Dahlquist, 2002; Donaire et al., 2005) and inclusions of mantle-derived magma (Vernon, 1984; Griffin et al., 2002; Barbarin, 2005; Yang et al.,

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2007). The MMEs can provide important information on the source nature and geodynamic setting as well as crust–mantle interaction processes of the granitoids.

The Central Asian Orogenic Belt (CAOB) (Sengör et al., 1993; Sengör and Natal'in, 1996; Jahn et al., 2000; Khain et al., 2003; Xiao et al., 2004) extends from the Urals in the west, through Kazakhstan, Kyrgyzstan, NW China, and southern Mongolia to the Pacific coast in the east. It experienced multiple subduction–accretion and collisional processes and consists mainly of accreted fragments including island arcs, accretionary complexes, seamounts, ophiolites and microcontinents (Kröner et al., 2007; Windley et al., 2007; Xiao et al., 2008). Distinct from the past cognition that the CAOB was formed by southward growth of juvenile terranes through subduction–accretion (e.g. Sengör and Natal'in, 1996; Windley et al., 2007; Xiao et al., 2009), a mechanism of northeastward growth of accretionary complexes as a result of southward subduction has been proposed to account for the formation of the Bayankhongor region in the central Mongolia (Figs. 1 and 2) (Buchan et al., 2001). During the evolution of the Bayankhongor region, extensive felsic and mafic magmas were produced, including high Ba–Sr granitoids and mafic enclaves (Buchan et al., 2002; Jahn et al., 2004; Demoux et al., 2009), and can provide important constraints on the evolutionary history of the Bayankhongor region. Recent research in the Bayankhongor region mostly focused on the accretionary complexes, however, little is known regarding the geodynamic and genetic relationships between high Ba–Sr granitoids and associated MMEs.

In this contribution, we report new U–Pb zircon ages, whole rock major and trace element and Sr–Nd–Hf isotopic compositions for a Precambrian high Ba–Sr granitoid pluton (the Ulaan Uul batholith) and associated MMEs, aiming to (1) investigate the petrogenesis and genetic relationship between the host granite and associated MMEs; (2) constrain the accretionary process of the Bayankhongor region in more detail.

## 2. Geological background and sample descriptions

### 2.1. Geological background

As a part of the CAOB, Mongolia experienced multiple subduction and collisional events as a consequence of the consumption of the Paleo–Asian Ocean (Fig. 1) (Badarch et al., 2002). Tectonically, Mongolia can be divided into the southern and northern domains separated by the Main Mongolian Lineament which marked a late Orodovician plate boundary (Lehmann et al., 2010). The southern domain is dominated by middle to late Paleozoic accretionary complexes, whereas the northern domain mainly comprises Precambrian to early Paleozoic metamorphic rocks, Neoproterozoic ophiolitic belts and early Paleozoic arc-related volcanic rocks (Buchan et al., 2001, 2002; Jiang et al., 2012). The Bayankhongor region in the northern domain is sandwiched between the Baydrag microcontinent in the south and the Hangay microcontinent in the north (Fig. 2) (Jahn et al., 2004; Jian et al., 2010). Previous workers divided the Bayankhongor region into several tectonic subunits, i.e., from south to north the Baydrag microcontinent, the Burd Gol mélange, the South Volcanic Belt, the Bayankhongor suture zone and the Dzag zone of the Hangay microcontinent (Fig. 2) (Buchan et al., 2001, 2002).

The basement of the Baydrag microcontinent is mainly composed of Neoproterozoic to Paleoproterozoic tonalitic to granitic gneisses which have underwent amphibolite- and granulite-facies metamorphism (Mitrofanov et al., 1985; Kotov et al., 1995; Teraoka et al., 1996; Kozakov et al., 2001; Demoux et al., 2009). To the northeast of the Baydrag block, the Burd Gol mélange represents a subduction–accretion complex and comprises phyllites, slates, chlorite–biotite schists which probably deposited in the earliest Neoproterozoic (Demoux et al., 2009). Moreover, this mélange underwent Barrovian-type metamorphism with metamorphic grade increasing northwards, and the time of metamorphism can

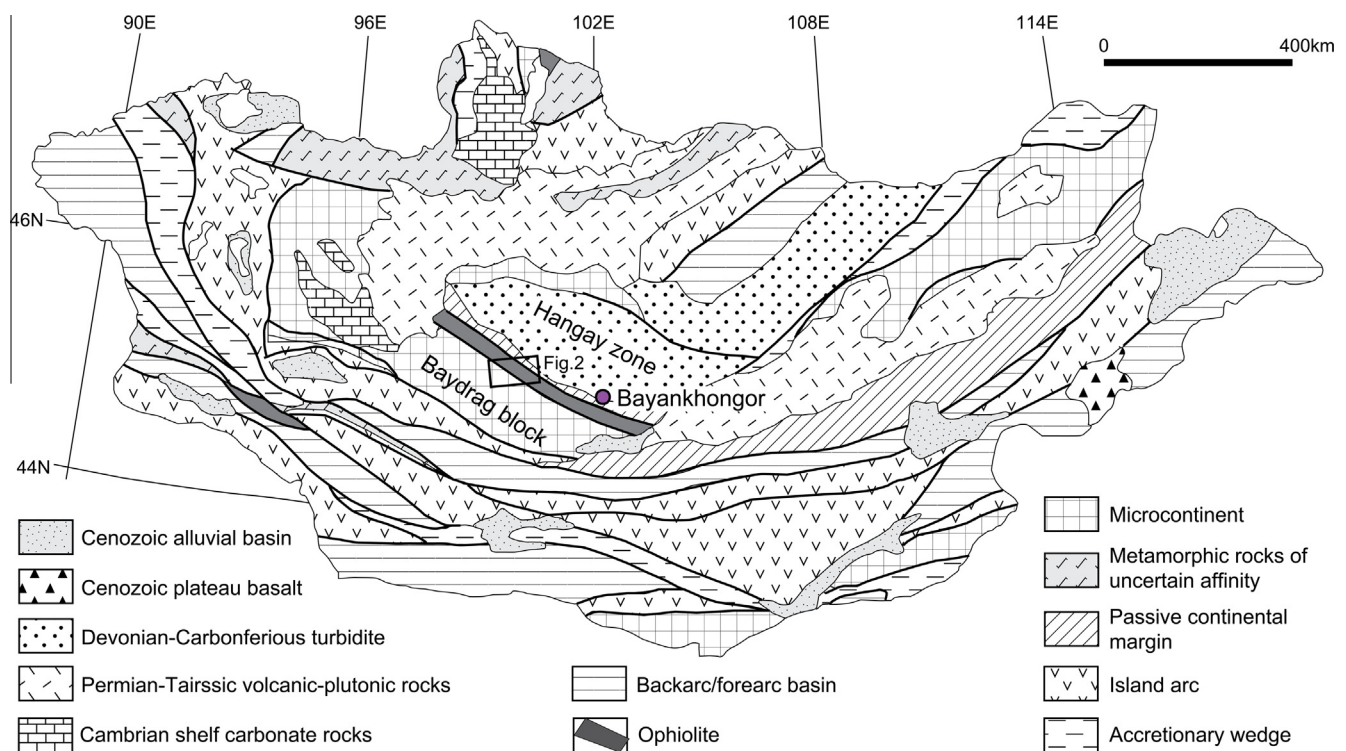


Fig. 1. Simplified terrane map of Mongolia (modified after Badarch et al., 2002).

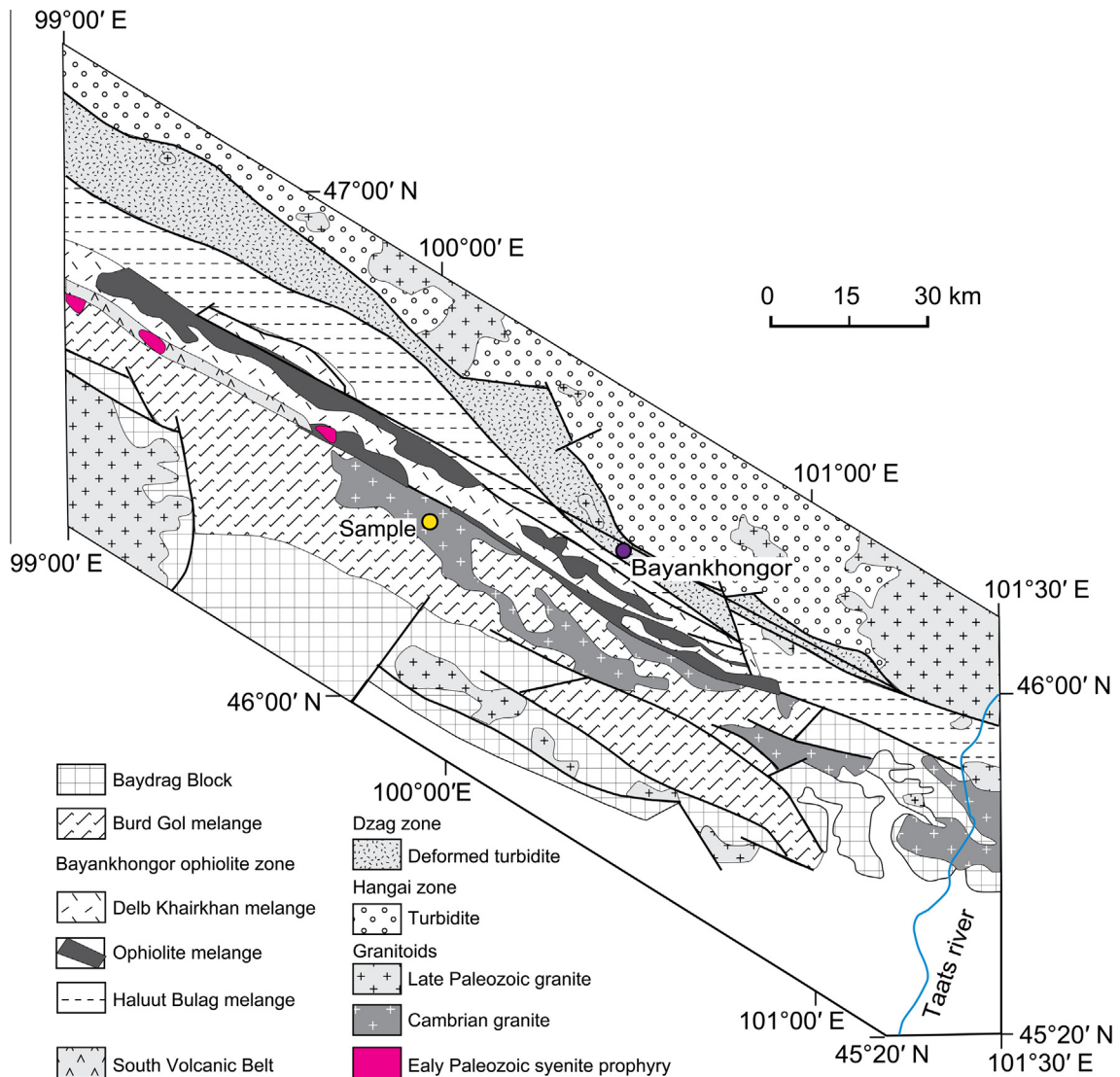


Fig. 2. Geological map of the Bayankhongor region (modified after Jian et al., 2010). The position of analyzed samples is also marked.

be roughly bracketed between ca. 562 and ca. 533 Ma (Höck et al., 2000; Kozakov et al., 2006). The South Volcanic Belt consists mainly of late Neoproterozoic to early Paleozoic arc-related volcanic rocks (Kröner et al., 2011). The Bayankhongor suture zone can be divided into three subunits, which are, from south to north, the Delb Khairkhan mélangé, the Bayankhongor ophiolitic mélangé and the Haluut Bulag mélangé (Buchan et al., 2002) (Fig. 2). The Delb Khairkhan mélangé is dominated by pelitic schists and has been considered to be middle Neoproterozoic to early Paleozoic in age based on the presence of Neoproterozoic stromatolites and Cambrian fossils (Dergunov et al., 1997). The Bayankhongor ophiolitic mélangé comprises ultramafic cumulates, gabbros, pillow lavas and sheeted dykes which have been tectonically dismembered into slices enclosed in a matrix of serpentinite (Buchan et al., 2002). Zircons from ophiolitic anorthosite, gabbro and plagiogranite yielded late Neoproterozoic  $^{207}\text{Pb}/^{206}\text{Pb}$  ages ranging from  $655 \pm 4$  to  $636 \pm 6$  Ma (Jian et al., 2010). The Haluut Bulag mélangé contains lenticular mudstone, limestone, sandstone and locally vesicular basalt enclosed in a matrix of pelitic schist (Buchan et al., 2001). Although no precise age is reported, the presence of radiolarian fossils in Cambrian sediments indicates that the depositional age of the Haluut Bulag mélangé is no older than Cambrian (542 Ma) (Osozawa et al., 2008). The Dzag zone of the

Hangay microcontinent comprises greenschist-facies psammitic and pelitic schists which probably deposited on the passive margin of the Hangay microcontinent to the north (Kurimoto et al., 1998; Buchan et al., 2001).

## 2.2. Granitoid intrusions

Voluminous late Neoproterozoic to early Cambrian granitoids intruded the Bayankhongor region (Fig. 2). For example, Buchan et al. (2002) reported zircon evaporation  $^{207}\text{Pb}/^{206}\text{Pb}$  ages of  $539 \pm 5$  Ma for a granitic pluton that intruded the Burd Gol mélangé. A syenite porphyry intruding the Burd Gol mélangé yielded a SHRIMP zircon age of  $523 \pm 2$  Ma (Jian et al., 2010). In addition, several zircon U–Pb ages (520–565 Ma) have been reported for granitic intrusions along the Taats River (Fig. 2), which has been considered as the eastern extension of the Baydrag block (Kozakov et al., 2006, 2008; Demoux et al., 2009).

In this study, we focus on the Ulaan Uul batholith, whose majority intrudes the northern margin of the Burd Gol mélangé (Fig. 2). Outcrops of this batholith can also be found in the Delb Khairkhan mélangé and the Bayankhongor ophiolitic mélangé. This batholith has been considered to be early Cambrian in age based on two zircon evaporation  $^{207}\text{Pb}/^{206}\text{Pb}$  ages of  $539 \pm 1$  and  $545 \pm 2$  Ma (Buchan

et al., 2002). The body is composed mainly of biotite monzogranite with rounded mafic microgranular enclaves, and has medium-grained granular textures (Fig. 3a). The main minerals are plagioclase (35–45 vol.%), alkali-feldspar (25–35 vol.%), quartz (20–25 vol.%), and biotite (~5 vol.%), with minor titanite, apatite and zircon. Enclaves within the batholith are fine-grained and ellipsoidal with size ranging from 10 to 30 cm, and generally show porphyritic or equigranular textures (Fig. 3a and b). The contact between the enclaves and host rocks are mostly sharp, though gradational contact can be observed locally. The enclaves are mainly dioritic in lithology and consist of plagioclase (35–40 vol.%), hornblende (30–35 vol.%), K-feldspar (10–15 vol.%), biotite (5–10 vol.%), with minor apatite (Fig. 3a and b). Except for a few that show gradational contact with the host, most enclaves do not contain K-feldspar megacrysts.

### 3. Analytical methods

All geochemical analysis of zircon U–Pb dating, whole rock geochemistry, Sr–Nd–Hf isotopic compositions, were carried out at the State Key Laboratory of Isotope Geochemistry, Guangzhou Institute of Geochemistry, Chinese Academy of Sciences (SKLIG GIG CAS).

#### 3.1. Zircon U–Pb dating

Zircons were separated from two samples (one from the Ulaan Uul batholith and one from mafic enclave) using conventional heavy liquid and magnetic separation techniques and then were hand-picked under a binocular microscope. Representative zircon grains were mounted in epoxy resin discs, then polished to about half of their thickness and coated with carbon. Cathodoluminescence (CL) images of zircons were obtained at SKLIG GIG CAS using a JEOL JXA-8100 Electron Probe Microanalyzer with a Mono CL3 CL System for inspecting internal morphology of zircons. Zircon U–Pb dating was carried out by using Laser ablation ICP–MS. NIST SRM 610 and Temora zircon standards were used as external standards. Detailed operating conditions for the laser ablation system and the ICP–MS instrument and data reduction can be found in Li et al. (2011).

#### 3.2. Whole-rock geochemistry and Sr–Nd isotope analyses

Major element compositions were measured on fused glass disks using X-ray fluorescence spectrometry (Rigaku ZSX100e)

following analytical procedures described by Li et al. (2006) and analytical precision for major elements was between 1% and 5%. Trace elements analyses were performed using a Perkin–Elmer Sciex ELAN 6000 inductively coupled plasma mass spectrometry (ICP–MS). Sample powders were digested using an HF + HNO<sub>3</sub> mixture in high pressure Teflon bombs for two days in order to assure complete dissolution of the refractory minerals. USGS rock standards and Chinese national rock standards (GSR-1, GSR-2, GSR-3, GSD-9) were used to calibrate elemental concentrations of the measured samples. The analytical procedures are referred to Li et al. (2002), and analytical precision was generally better than 5%. Sr and Nd isotopic analyses were performed on a Micromass Isoprobe multi-collector mass spectrometer (MC–ICP–MS) at SKLIG GIG CAS, following the procedures described by Li et al. (2004). Measured <sup>143</sup>Nd/<sup>144</sup>Nd and <sup>87</sup>Sr/<sup>86</sup>Sr ratios were normalized to <sup>146</sup>Nd/<sup>144</sup>Nd = 0.7219 and <sup>86</sup>Sr/<sup>88</sup>Sr = 0.1194, respectively. The <sup>87</sup>Sr/<sup>86</sup>Sr ratio of the NBS987 standard and <sup>143</sup>Nd/<sup>144</sup>Nd ratio of the Shin Etsu JNdi-1 standard measured during this study were 0.710256 ± 22 (2σ) and 0.512110 ± 8 (2σ), respectively.

#### 3.3. Zircon Lu–Hf isotope analyses

In situ zircon Hf isotopic analysis was carried out using the Neptune Plus MC–ICP–MS, attached to the Resonetics RESOLUTION M-50-LR Excimer Laser Ablation System at SKLIG GIG CAS. During the analyses, a laser repetition rate of 8 Hz at 80 mJ was used for ablating zircons, and spot size was 45 μm. The ablated material was transported in a helium carrier gas with addition of a small flow of nitrogen. Data acquisition for each analysis consists of 30 s gas background collection and 30 s signal collection for laser ablation. A signal collection model for one block includes 200 cycles, in which one cycle has 0.131 s integration time. The measured isotopic ratios of <sup>176</sup>Hf/<sup>177</sup>Hf were normalized to <sup>176</sup>Hf/<sup>177</sup>Hf = 0.7325, using an exponential correction for mass bias. Correction for isobaric interference of <sup>176</sup>Lu on <sup>176</sup>Hf and <sup>176</sup>Yb on <sup>176</sup>Hf was performed using the recommended <sup>176</sup>Lu/<sup>175</sup>Lu ratio of 0.02655 (Machado and Simonetti, 2001) and <sup>176</sup>Yb/<sup>171</sup>Yb ratio of 0.90184. Penglai zircons were used as reference standards and were analyzed twice before and after every 5 analyses. Repeated measurements of this zircon standard during our study yielded <sup>176</sup>Hf/<sup>177</sup>Hf = 0.282891 ± 0.000012 (2σ, n = 22), which is identical to the previously reported literature values (Li et al., 2010). The <sup>176</sup>Lu decay constant of 1.867 × 10<sup>-11</sup> year<sup>-1</sup> (Soderlund et al.,

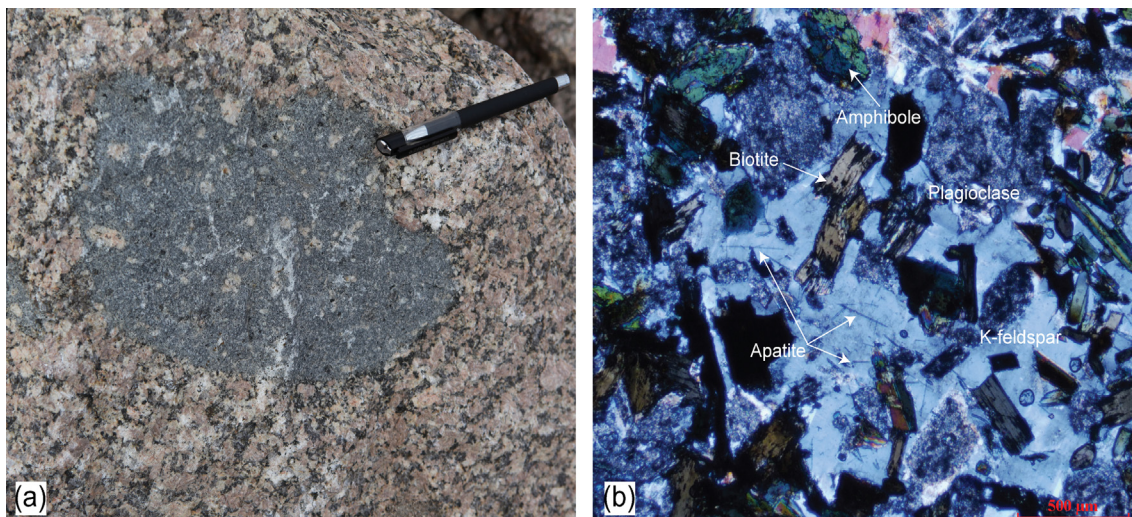


Fig. 3. (a) Field photograph showing the relationship between the host granite and MMEs; (b) photomicrograph showing the main textures and mineral assemblages of the MME.

2004) was used to calculate initial  $^{176}\text{Hf}/^{177}\text{Hf}$  ratios. The chondritic values of  $^{176}\text{Hf}/^{177}\text{Hf}$  (0.282772) and  $^{176}\text{Lu}/^{177}\text{Hf}$  (0.0332) reported by (Blichert-Toft and Albarede, 1997) were adopted for the calculation of  $\varepsilon\text{Hf}$  values. Single-stage Hf model ages ( $T_{\text{DM1}}$ ) were calculated relative to the depleted mantle present-day value of  $^{176}\text{Hf}/^{177}\text{Hf} = 0.283250$  and  $^{176}\text{Lu}/^{177}\text{Hf} = 0.0384$  (Griffin et al., 2000). Two-stages “crustal” model ages ( $T_{\text{DM2}}$ ) were calculated using the mean  $^{176}\text{Lu}/^{177}\text{Hf}$  ratio of 0.015 for the average continental crust (Griffin et al., 2002).

## 4. Analytical results

### 4.1. Geochronology

LA-ICP-MS zircon U–Pb isotope analytical results of the host granite and associated MME can be found in Appendix 1. Zircons from the host granite (MG11-89-11) are prismatic and transparent with well-developed oscillatory zoning, indicating a magmatic origin. Size of the zircons ranges from 100 to 200  $\mu\text{m}$  in length, with length/width ratios of 2:1 to 4:1 (Fig. 4a). The analyzed zircons have intermediate Th (27–523 ppm) and U (41–421 ppm) contents with Th/U ratios of 0.29 to 1.33, consistent with an igneous origin. Twenty-seven analyses were obtained and form a coherent group with a weighted mean  $^{206}\text{Pb}/^{238}\text{U}$  age of  $546 \pm 3$  Ma (Fig. 5b). The age is consistent with previous zircon evaporation  $^{207}\text{Pb}/^{206}\text{Pb}$  ages ( $539 \pm 1$  Ma and  $545 \pm 2$  Ma) (Buchan et al., 2002), and is considered to be the crystallization age of the Ulaan Uul batholith.

Zircons from the MME (MG11-89-1) are stubby to prismatic and grayish dark in color, and have weakly oscillatory zoning. The zircons range in size from 50 to 200  $\mu\text{m}$  in length, with length/width ratios of 1:1 to 3:1 (Fig. 4b). The analyzed zircons have variable Th (29–814 ppm) and U (69–537 ppm) contents with Th/U ratios of

0.28–1.51, indicating a magmatic origin. No inherited cores were found. Seventeen analyses of zircons form a coherent group on the Concordia diagram with a weighted mean  $^{206}\text{Pb}/^{238}\text{U}$  age of  $547 \pm 3$  Ma (Fig. 5a), which represents the crystallization age of the mafic enclaves. This age is consistent with the U–Pb ages for the host biotite monzogranite, suggesting that the MMEs and their host were coeval mafic and felsic magmatism.

### 4.2. Major and trace elements

Major and trace element data for the host granite and MMEs are listed in Table 1. Samples from the host biotite monzogranite display high  $\text{SiO}_2$  (66.0–75.4 wt.%), low  $\text{MgO}$  (0.55–1.47 wt.%) and  $\text{TiO}_2$  (0.23–0.64 wt.%) contents (Fig. 7). The host samples are characterized by high total alkali ( $\text{K}_2\text{O} + \text{Na}_2\text{O} = 7.1$ –10.0 wt.%) (Fig. 6a) and  $\text{K}_2\text{O}/\text{Na}_2\text{O}$  (0.7–1.7), showing high-K calc-alkaline to shoshonitic characteristics (Fig. 6b). Rocks from the host granite are strongly peraluminous with ASI values ranging from 1.10 to 1.17 (Fig. 6c). Rocks of the biotite monzogranite contain high Ba (1189–2544 ppm), Sr (440–742 ppm) and low Yb (<2.1 ppm) and Ta (<2.2 ppm) contents. In addition, they exhibit high K/Rb (354–418), Sr/Y (30–75) and La/Yb (21–91), and low Rb/Ba (<0.07) and Rb/Sr (<0.27) ratios. The rocks are characterized by enrichment of LREE and display slight HREE fractionation ( $(\text{La}/\text{Yb})_N = 14.8$ –64.9;  $(\text{Gd}/\text{Yb})_N = 2.0$ –2.6) (Fig. 8b), with slightly to moderate negative Eu anomalies ( $\text{Eu}/\text{Eu}^* = 0.65$ –0.98). On the primitive mantle-normalized trace element diagram, the biotite monzogranite samples show relative enrichment in LILEs (such as Rb, Ba) and depletion in HFSEs (such as Nb, Ta, Ti) (Fig. 8d).

The MMEs are mafic to intermediate in composition with  $\text{SiO}_2$  ranging from 52.1 to 67.6 wt.% (Fig. 6a). The MME samples have variable  $\text{MgO}$  (1.3–3.7 wt.%),  $\text{Al}_2\text{O}_3$  (14.7–19.1 wt.%) and  $\text{TiO}_2$

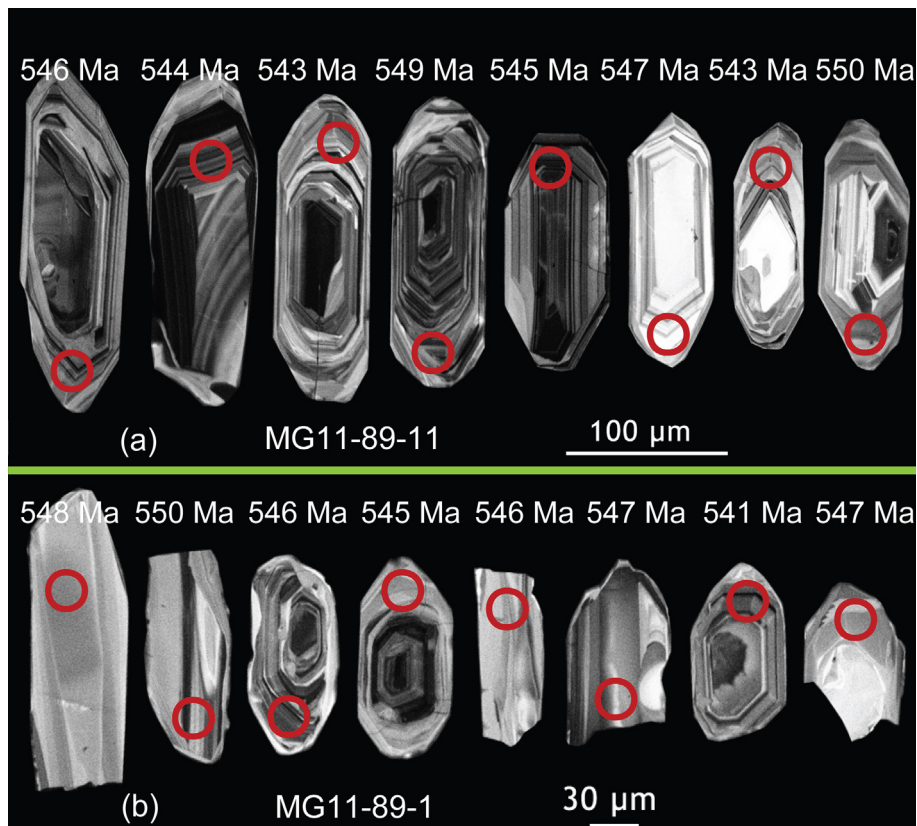


Fig. 4. CL images of representative zircons from the host granite (a) and associated MME (b).

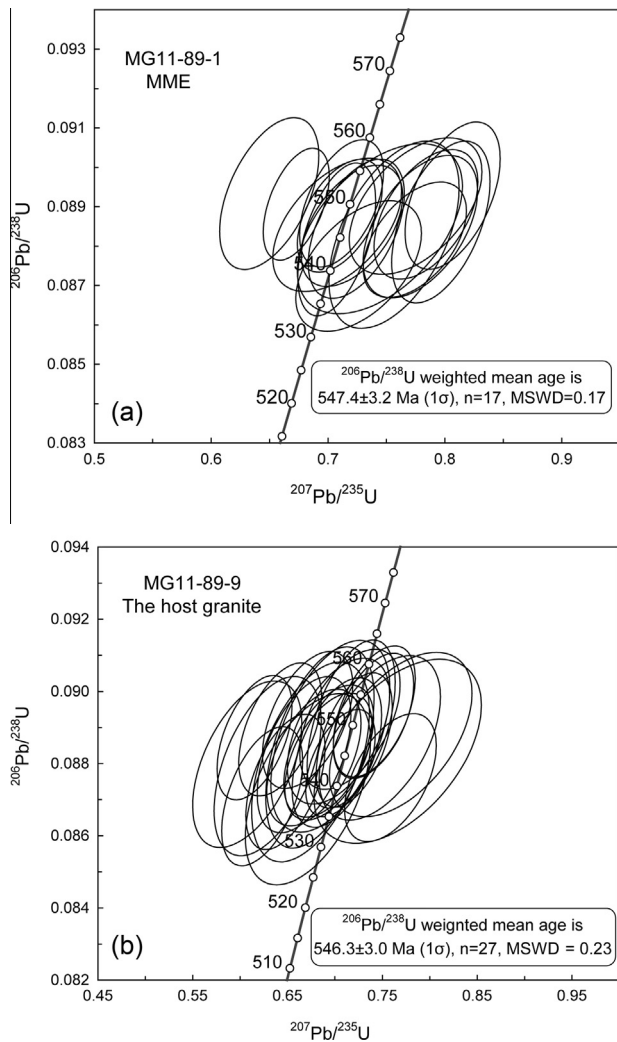


Fig. 5. Zircon U–Pb Concordia diagrams and weighted mean  $^{206}\text{Pb}/^{238}\text{U}$  ages for the MME (a) and host granite (b).

(0.7–1.6 wt.%) contents (Fig. 7), characterized by high total alkali contents ( $\text{K}_2\text{O} + \text{Na}_2\text{O} = 5.8\text{--}8.3$  wt.%) (Fig. 6a). The MME rocks are metaluminous to slightly peraluminous ( $\text{ASI} = 0.88\text{--}1.05$ , mostly  $<1$ ) (Fig. 6c), and relatively Na-enriched as indicated by their high  $\text{Na}_2\text{O}/\text{K}_2\text{O}$  ratios (1.1–2.5, mostly  $>1.8$ ). Rocks of the MMEs have high concentrations of Sr (772–907 ppm), and highly variable Y (9.1–64.5 ppm), Nb (10.5–52.5) contents and Sr/Y (12–87) ratios. The MME samples are characterized by variable LREE enrichment ( $(\text{La}/\text{Yb})_N = 4.8\text{--}39.4$ ) and HREE fractionation ( $(\text{Gd}/\text{Yb})_N = 1.4\text{--}3.1$ ), with moderate negative Eu anomalies ( $\text{Eu}/\text{Eu}^* = 0.51\text{--}0.75$ ) (Fig. 8a). On the primitive mantle-normalized trace element diagram, the MME samples show relative enrichment in large ion lithophile element (LILE, e.g. Rb, Ba), and variable depletion in high field-strength element (HFSE, e.g. Nb, Ta, Zr) as suggested by their variable Nb/La ratios (0.15–1.27) (Fig. 8c).

#### 4.3. Sr–Nd isotopic compositions

The Sr–Nd isotope compositions of the host granite and MMEs are presented in Table 2 and Fig. 9a. Samples from the host granite have a narrow range of initial  $^{87}\text{Sr}/^{86}\text{Sr}$  ratios (0.7059–0.7065) and  $\epsilon\text{Nd}(t)$  (–1.8 to –2.1) with Nd model ages ( $T_{\text{DM}}$ ) of 1.43–1.45 Ga. The MMEs also have uniform initial  $^{87}\text{Sr}/^{86}\text{Sr}$  ratios (0.7060–

0.7066) and  $\epsilon\text{Nd}(t)$  (–1.6 to –2.1), and Nd model ages ( $T_{\text{DM}}$ ) of 1.41–1.45 Ga, overlapping the range of the host granite samples.

#### 4.4. Zircon Hf isotopic compositions

In situ zircon Hf isotopic composition for the host granite and MME samples are listed in Appendix 2 and presented in Fig. 10. Except for two zircons from the host granite display positive  $\epsilon\text{Hf}(t)$  values (+0.12 and +1.4) with single-stage Hf model ages of 1.08 and 1.13 Ga, the rest twenty-four zircons from the host rock all show variably negative  $\epsilon\text{Hf}(t)$  values (–0.63 to –11.9) with two-stage Hf model ages of 1.54–2.25 Ga. Zircons from the MME have higher  $^{176}\text{Hf}/^{177}\text{Hf}$  ratios and  $\epsilon\text{Hf}(t)$  values than those of the host granite samples. Ten of the seventeen analyzed zircons from MME have positive  $\epsilon\text{Hf}(t)$  values (+0.7 to +2.6) with single-stage Hf model ages of 1.05–1.12 Ga, and seven MME zircons have negative  $\epsilon\text{Hf}(t)$  values (–0.75 to –3.5) with two-stage Hf model ages of 1.51–1.72 Ga.

### 5. Discussion

#### 5.1. Magma mixing

The igneous texture of the MMEs indicates a magmatic origin (Fig. 3b). In addition, zircon U–Pb dating suggests that the MMEs were coeval with the host granite, thus an origin of xenolith captured from the wall–rock can be precluded. Recently, a cognate origin of MMEs (e.g., early-stage cumulates or differentiation) has been proposed (Dahlquist, 2002; Donaire et al., 2005). Although the similar zircon U–Pb ages of the MMEs and host granite may imply a cogenetic suite, their different zircon Hf isotopic compositions suggest different origins for the MMEs and host granite. In fact, the MMEs have higher total REE contents (mostly  $>230$  ppm) than the host granite samples (mostly  $<230$  ppm), thus precluding an origin of early-stage differentiation product from a cognitive magma. Moreover, some workers proposed that MMEs represent residual materials (restite) successively immixing with the melt (Chappell and White, 1992; White et al., 1999). In this case, MMEs should share similar isotopic compositions to the host granitoids. Although having Sr–Nd isotopic compositions similar to their host (Fig. 9a), the MMEs in the Ulaan Uul batholith display different  $^{176}\text{Hf}/^{177}\text{Hf}$  ratios and  $\epsilon\text{Hf}(t)$  values from those of the host granite (Fig. 10), which, together with their igneous texture (Fig. 3b), argues against a restite origin. Similar whole-rock Sr–Nd isotopes and distinct zircon Hf isotopes between the host granite and associated MMEs, pointing to somewhat Nd–Hf isotope decoupling, have been served as a powerful evidence for crust–mantle magma mixing (Kemp et al., 2005, 2007; Yang et al., 2007; Qin et al., 2009). Generally, chemical equilibrium and isotopic equilibrium can be achieved by elemental diffusion during the magma mixing process (Karsli et al., 2007). The MMEs share similar Sr–Nd isotopic compositions to the host (Fig. 9a), which can be explained by sufficient diffusion exchange with the granitic melt (Qin et al., 2009). However, as a resistant and refractory mineral, zircon commonly crystallizes in the early stage of magmatism, probably prior to the homogenization of mixed magma, thus may retain its primary Hf isotopic signature acquired from the magma and survive post-crystallization thermal disturbances (Kinny and Maas, 2003; Zheng et al., 2007).

Magma mixing process has long been considered as an important mechanism in producing granitoids. When small volumes of mafic magma injects into a felsic magma chamber, the mafic magma would cool rapidly and become more viscous to form discrete magma globules (Vernon, 1984; Griffin et al., 2002; Barbarin, 2005; Yang et al., 2007). In the case of the Ulaan Uul batholith and

**Table 1**

Major oxides (wt.%) and trace element (ppm) concentrations of the Ulaan Uul batholith and associated Mafic enclaves in the Bayankhongor area, central Mongolia.

Sample	Ulaan Uul batholith										MME	
	MG11-89-9	MG11-89-10	MG11-89-11	MG11-89-12	MG11-89-13	MG11-89-14	MG11-89-15	MG11-89-16	MG11-89-17	MG11-89-18	MG11-89-1	MG11-89-2
<i>Major oxides (wt.%)</i>												
SiO <sub>2</sub>	73.3	67.3	69.5	68.0	70.2	68.8	70.1	75.4	69.3	66.0	54.1	55.9
TiO <sub>2</sub>	0.23	0.64	0.31	0.51	0.42	0.44	0.45	0.27	0.40	0.64	1.56	1.39
Al <sub>2</sub> O <sub>3</sub>	14.1	15.4	15.9	15.9	15.5	15.2	15.1	12.3	15.9	16.3	18.0	18.1
Fe <sub>2</sub> O <sub>3</sub>	1.60	3.74	1.97	2.89	2.57	3.54	2.73	1.98	2.17	3.59	8.19	7.51
MnO	0.02	0.06	0.03	0.04	0.04	0.04	0.04	0.03	0.04	0.06	0.17	0.15
MgO	0.55	1.47	0.60	1.02	0.99	0.90	1.10	0.59	0.95	1.45	3.56	3.25
CaO	0.75	2.04	0.71	1.37	0.87	0.98	0.86	0.82	0.80	1.63	5.40	4.73
Na <sub>2</sub> O	3.29	4.35	3.74	4.01	4.35	4.02	4.46	3.06	3.96	3.89	4.81	4.87
K <sub>2</sub> O	5.13	2.94	6.23	4.72	4.18	4.59	3.78	4.07	5.31	4.79	2.13	2.33
P <sub>2</sub> O <sub>5</sub>	0.05	0.22	0.08	0.17	0.15	0.11	0.15	0.07	0.12	0.21	0.48	0.42
L.O.I	0.87	1.63	0.81	1.18	0.65	1.22	1.11	1.27	0.85	1.27	1.39	1.18
Total	99.9	99.8	99.9	99.9	99.8	99.8	99.6	99.9	99.8	99.8	99.8	99.8
<i>Trace elements (ppm)</i>												
Sc	0.58	4.71	0.66	1.08	1.50	1.16	2.94	0.73	1.25	3.93	17.3	15.0
Ti	1223	3127	1638	2634	2145	2363	2352	1443	2151	3324	8915	7711
V	27.4	41.3	27.0	38.7	31.3	52.8	33.1	29.8	30.7	48.3	116	104
Cr	7.93	8.33	7.27	10.0	9.32	8.70	7.16	8.34	9.92	13.6	35.8	30.5
Mn	177	425	220	316	304	314	330	197	257	438	1294	1145
Co	2.99	6.82	2.84	4.90	4.85	4.76	4.63	3.22	4.17	7.25	17.6	16.1
Ni	1.58	3.88	1.74	2.10	2.87	2.25	2.70	1.64	2.04	5.69	22.2	18.1
Cu	2.25	4.37	1.98	2.78	2.47	2.68	2.09	1.79	2.14	6.38	9.36	8.23
Zn	23.7	50.6	26.6	40.6	37.8	42.9	40.4	30.2	32.6	53.5	122	117
Ga	14.2	18.8	15.6	17.4	16.7	18.0	17.0	13.1	16.4	19.1	26.1	26.2
Ge	1.10	1.32	1.20	1.30	1.16	1.54	1.11	1.25	1.12	1.47	1.92	1.99
Rb	105	68.8	130	97.9	93.5	92.0	91.3	80.7	110	107	94.2	104
Sr	614	742	475	689	532	554	440	519	584	736	784	791
Y	8.14	22.7	11.7	19.3	12.9	17.2	14.5	7.98	9.07	22.4	42.6	34.9
Zr	170	250	207	213	177	280	167	234	140	215	134	150
Nb	5.17	18.8	10.3	16.6	9.67	12.4	12.0	6.23	9.66	18.6	36.6	33.5
Cs	0.50	1.31	0.35	0.72	0.52	0.38	0.40	0.46	0.48	1.53	1.38	1.55
Ba	2375	1189	2544	1984	1655	1955	1240	1876	2354	2056	475	425
La	62.4	60.7	46.1	46.7	40.1	131	34.1	71.7	39.9	39.9	41.7	38.9
Ce	107	120	87.5	101	76.7	238	71.5	123	70.0	94.8	117	109
Pr	10.2	14.3	9.74	12.6	8.73	23.9	8.69	12.1	7.87	11.9	17.6	15.9
Nd	30.6	51.8	32.8	44.9	29.8	73.2	31.2	36.2	26.8	44.4	73.3	63.8
Sm	3.47	8.48	4.78	7.49	4.780	8.70	5.37	4.08	4.04	7.81	14.3	11.9
Eu	0.43	1.56	0.78	1.24	0.83	1.22	1.00	0.54	0.79	1.43	3.02	2.52
Gd	1.89	6.05	3.06	5.26	3.31	4.30	3.72	2.01	2.57	5.48	11.0	9.00
Tb	0.29	0.93	0.46	0.77	0.51	0.73	0.57	0.32	0.39	0.84	1.66	1.34
Dy	1.58	4.86	2.33	4.05	2.68	3.71	3.13	1.67	2.04	4.64	8.89	7.13
Ho	0.30	0.90	0.45	0.75	0.51	0.68	0.59	0.31	0.37	0.88	1.68	1.36
Er	0.80	2.32	1.15	1.93	1.26	1.70	1.49	0.83	0.92	2.17	4.27	3.48
Tm	0.12	0.33	0.16	0.27	0.19	0.24	0.21	0.12	0.13	0.31	0.60	0.49
Yb	0.78	2.11	1.09	1.72	1.16	1.54	1.32	0.79	0.82	1.94	3.83	3.22
Lu	0.12	0.30	0.16	0.24	0.17	0.22	0.19	0.13	0.12	0.28	0.53	0.46
Hf	3.75	5.82	4.88	4.98	4.32	6.29	4.16	5.16	3.48	4.95	3.25	3.65
Ta	0.53	2.21	1.10	1.95	1.10	1.40	1.29	0.64	1.03	1.86	3.38	3.43
Pb	18.0	11.4	15.2	17.4	15.1	15.9	10.9	15.5	16.3	15.7	9.56	9.27
Th	8.28	7.91	6.13	7.36	5.81	15.2	4.73	8.58	4.71	5.81	6.17	5.85
U	0.35	0.91	0.58	0.85	0.86	0.60	1.01	0.41	0.65	0.86	1.33	1.36
Mg <sup>#</sup>	44.4	47.8	41.4	45.2	47.4	37.1	48.5	41.1	50.4	48.5	50.3	50.2

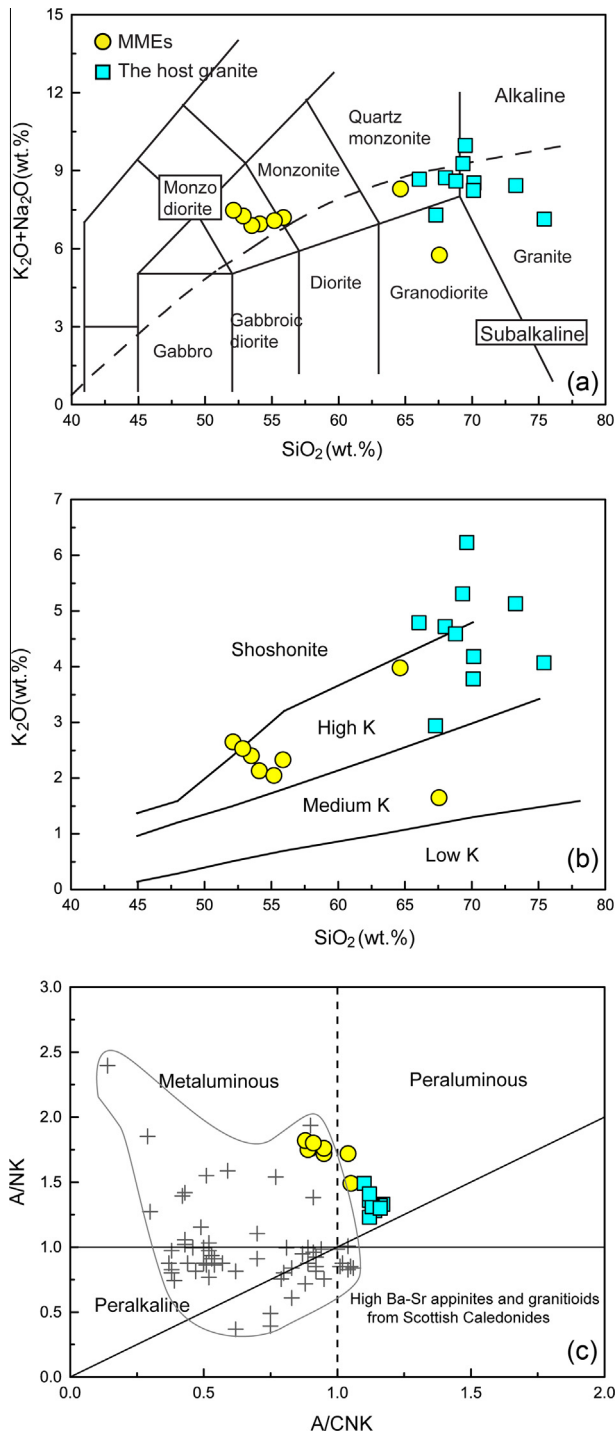
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Table 1 (continued)

Sample	MME					
	MG11-89-3	MG11-89-4	MG11-89-5	MG11-89-6	MG11-89-7	MG11-89-8
<i>Major oxides (wt.%)</i>						
SiO <sub>2</sub>	67.5	55.2	52.1	53.5	52.8	64.6
TiO <sub>2</sub>	0.66	1.38	1.56	1.47	1.53	0.71
Al <sub>2</sub> O <sub>3</sub>	14.7	18.3	19.1	18.2	18.9	17.0
Fe <sub>2</sub> O <sub>3</sub> <sup>T</sup>	4.87	7.46	8.50	8.00	8.35	4.03
MnO	0.06	0.16	0.17	0.12	0.15	0.07
MgO	1.31	3.06	3.74	4.03	3.71	1.42
CaO	3.08	5.57	5.14	5.83	5.71	2.64
Na <sub>2</sub> O	4.10	5.02	4.83	4.50	4.73	4.33
K <sub>2</sub> O	1.65	2.05	2.65	2.40	2.53	3.98
P <sub>2</sub> O <sub>5</sub>	0.20	0.37	0.41	0.39	0.40	0.25
L.O.I	1.71	1.21	1.60	1.31	0.86	0.78
Total	99.9	99.8	99.8	99.8	99.8	99.8
<i>Trace elements (ppm)</i>						
Sc	1.26	18.4	16.5	11.6	11.7	3.16
Ti	3586	7614	8288	7831	8059	4237
V	86.9	105	119	136	129	56.3
Cr	15.1	22.0	23.1	35.5	19.5	17.2
Mn	445	1200	1257	892	1120	538
Co	8.37	17.9	21.6	23.4	21.1	8.45
Ni	5.09	12.8	16.1	23.8	16.1	6.81
Cu	5.02	61.9	45.3	26.5	15.7	3.07
Zn	61.3	107	127	93.6	113	66
Ga	19.4	26.2	26.8	22.8	25.0	22.7
Ge	1.27	2.13	1.81	1.48	1.49	1.69
Rb	65.2	49.9	118	78.1	111	123
Sr	791	772	854	907	794	854
Y	9.12	64.5	40.6	13.6	18.6	23.7
Zr	180	224	184	159	189	269
Nb	10.3	52.5	34.9	10.5	15.6	21.7
Cs	0.89	1.00	1.60	1.09	1.53	1.67
Ba	435	423	592	777	518	1547
La	65.5	41.1	39.0	32.5	33.0	63.2
Ce	113	123	109	61.7	69.7	123
Pr	11.6	20.6	16.5	7.22	9.20	14.6
Nd	35.1	89.2	67.4	28.6	36.4	51.1
Sm	3.87	18.8	13.0	5.30	6.45	8.22
Eu	0.80	3.59	2.68	1.52	1.63	1.62
Gd	1.95	15.2	10.0	4.26	5.04	5.55
Tb	0.28	2.36	1.55	0.62	0.72	0.87
Dy	1.53	13.0	8.49	3.17	3.84	4.62
Ho	0.32	2.54	1.63	0.60	0.74	0.89
Er	0.91	6.64	4.09	1.37	1.89	2.29
Tm	0.15	0.96	0.60	0.19	0.26	0.33
Yb	1.19	6.17	3.86	1.16	1.63	2.11
Lu	0.21	0.86	0.54	0.17	0.25	0.30
Hf	7.34	5.72	4.23	3.74	4.32	6.16
Ta	0.69	6.54	3.61	0.43	0.94	2.14
Pb	9.19	10.8	9.67	9.19	13.7	16.5
Th	8.33	7.60	4.77	3.67	2.63	9.19
U	0.86	1.79	1.22	1.00	0.71	1.51
Mg <sup>#</sup>	38.6	48.9	50.6	54.0	50.8	45.1

Mg<sup>#</sup> = 100 \* Mg/(Mg + Fe<sup>2+</sup>), assuming Fe<sup>2+</sup>/Fe<sup>total</sup> = 0.85; LOI = loss of ignition; Fe<sub>2</sub>O<sub>3</sub><sup>T</sup> = total Fe oxides as Fe<sub>2</sub>O<sub>3</sub>.





**Fig. 6.** Petrochemical diagrams of the host granite and MMEs. (a) Total alkali–silica (TAS) diagram (Le Bas et al., 1986); (b)  $\text{SiO}_2$ – $\text{K}_2\text{O}$  diagram (Peccerillo and Taylor, 1976); (c) ANK–ACNK diagram (Maniar and Piccoli, 1989). Data for the high Ba–Sr apinites and granitoids in the Scottish Caledonides from Fowler and Henney (1996) and Fowler et al. (2001, 2008).

associated MMEs, there are some characteristics in support of magma mixing. For example, some MMEs within the Ulaan Uul batholith contain large K-feldspar phenocrysts, suggesting a non-consolidation state which allows physical transfer of crystals from the host felsic magma into the mafic magma (Fig. 3a) (Waight et al., 2000; Perugini et al., 2003). In addition, the finer grained texture of the MMEs was caused by the undercooling of a hot mafic magma in a relatively cool, felsic magma (Vernon, 1984). As a

mineral crystallized in the early stage of magma, zircon has high Lu–Hf closure temperature ( $>800^\circ\text{C}$ ) and hence can record primitive isotopic signatures of the magmatic components and subsequent isotopic variations during the mixing process (Cherniak and Waston, 2001; Griffin et al., 2002; Kemp and Hawkesworth, 2006; Yang et al., 2007; Li et al., 2009). Most zircons from the MMEs have positive  $\varepsilon\text{Hf}(t)$  values (0.7–2.6) with single-stage Hf model ages of 1.05–1.12 Ga, while others have negative  $\varepsilon\text{Hf}(t)$  values ( $-0.75$  to  $-3.5$ ) with two-stage Hf model ages of 1.51–1.72 Ga. In contrast, most of the zircons from the host granite have negative  $\varepsilon\text{Hf}(t)$  values ( $-0.63$  to  $-11.9$ ) with two-stage Hf model ages of 1.54–2.25 Ga, and only two spots possess positive  $\varepsilon\text{Hf}(t)$  values of 0.12 and 1.4 with single-stage Hf model ages of 1.13 and 1.08 Ga, respectively (Fig. 10). The distinct Hf isotopic compositions of zircons from the MMEs and host granite does not reflect a simple, cogenetic evolution through closed-system fractionation. Rather, the different Hf isotopic compositions of the zircons suggest that the MMEs and their host granite were probably derived from different sources, and that magma mixing/mingling process is the most likely regime to form the Ulaan Uul batholith. Such a mixing mechanism can be further testified by the hyperbolic curves in diagrams involving two ratios of incompatible elements (Langmuir et al., 1978). In the plot of Zr/Sm vs. Th/La, rocks from the Ulaan Uul batholith and associated MMEs define a characteristic hyperbolic mixing line (Fig. 12a), consistent with a two-component mixing process. In addition, the binary diagrams of  $^{143}\text{Nd}/^{144}\text{Nd}$  vs.  $^{147}\text{Sm}/^{144}\text{Nd}$  and Rb/Sr vs.  $^{87}\text{Sr}/^{86}\text{Sr}$  (Fig. 12b and c) confirm that the magma mixing process is a valid mechanism in the genesis of the host rocks.

## 5.2. Source nature of the MMEs and host granite

The MMEs possess low  $\text{SiO}_2$  (mostly  $<56$  wt.%) contents, high  $\text{Mg}^\#$  (up to 55) as well as relatively high V (up to 136 ppm) and Cr (up to 36 ppm) concentrations, suggesting that a mantle component played a prominent role in their genesis (Wilson, 1989; Rudnick and Gao, 2003). Several MME samples with high  $\text{Mg}^\#$  (50–55) may represent more primitive magmatic components, and display enriched Sr–Nd isotopic compositions (initial  $^{87}\text{Sr}/^{86}\text{Sr} = 0.7062$  to  $0.7063$ ;  $\varepsilon\text{Nd}(t) = -1.6$  to  $-1.7$ ). The Sr–Nd isotopic compositions of the MMEs are significantly different from those of the Bayankhongor ophiolites that are thought to derive from a depleted asthenospheric mantle (Fig. 9a) (Jian et al., 2010). Furthermore, MME zircons with positive  $\varepsilon\text{Hf}(t)$  values show single-stage Hf model ages (1.05–1.12 Ga) that are much older than the zircon U–Pb ages of 547 Ma, implying a late Mesoproterozoic lithospheric mantle source. Recently, a Mesoproterozoic lithospheric mantle of the Hangay zone was proposed based on the Mesoproterozoic Nd model ages (1.4–1.1 Ga) of granitoids (Kovalenko et al., 2004) and  $T_{\text{RD}}$  model ages (1.5–1.3 Ga and 1.0 Ga) for the sulfides from spinel peridotite xenoliths (Wang et al., 2013). The above evidence indicates that the MMEs were likely derived from partial melting of Mesoproterozoic lithospheric mantle that reworked at the early Cambrian.

Rocks from the host granite exhibit high  $\text{SiO}_2$ , low MgO, Cr and Ni contents, and enriched Sr–Nd–Hf isotopic compositions, reflecting a crustal origin. Compositional diversity of crustally derived melts strongly depends on source compositions and variable melting conditions (Beard and Lofgren, 1991; Rapp et al., 1991; Patiño Douce and Beard, 1995; Thompson and Connolly, 1995). To identify the possible source, it is necessary to compare the chemical and isotopic compositions of the host granite with those of crustal rocks in the surrounding areas. The Archean to Paleoproterozoic basement rocks in the Baydrag block are characterized by far more negative  $\varepsilon\text{Nd}(t)$  values ( $-14.52$  to  $-33.24$ , calculated at 547 Ma) (Kozakov et al., 1997) than those of the host granite. Compared

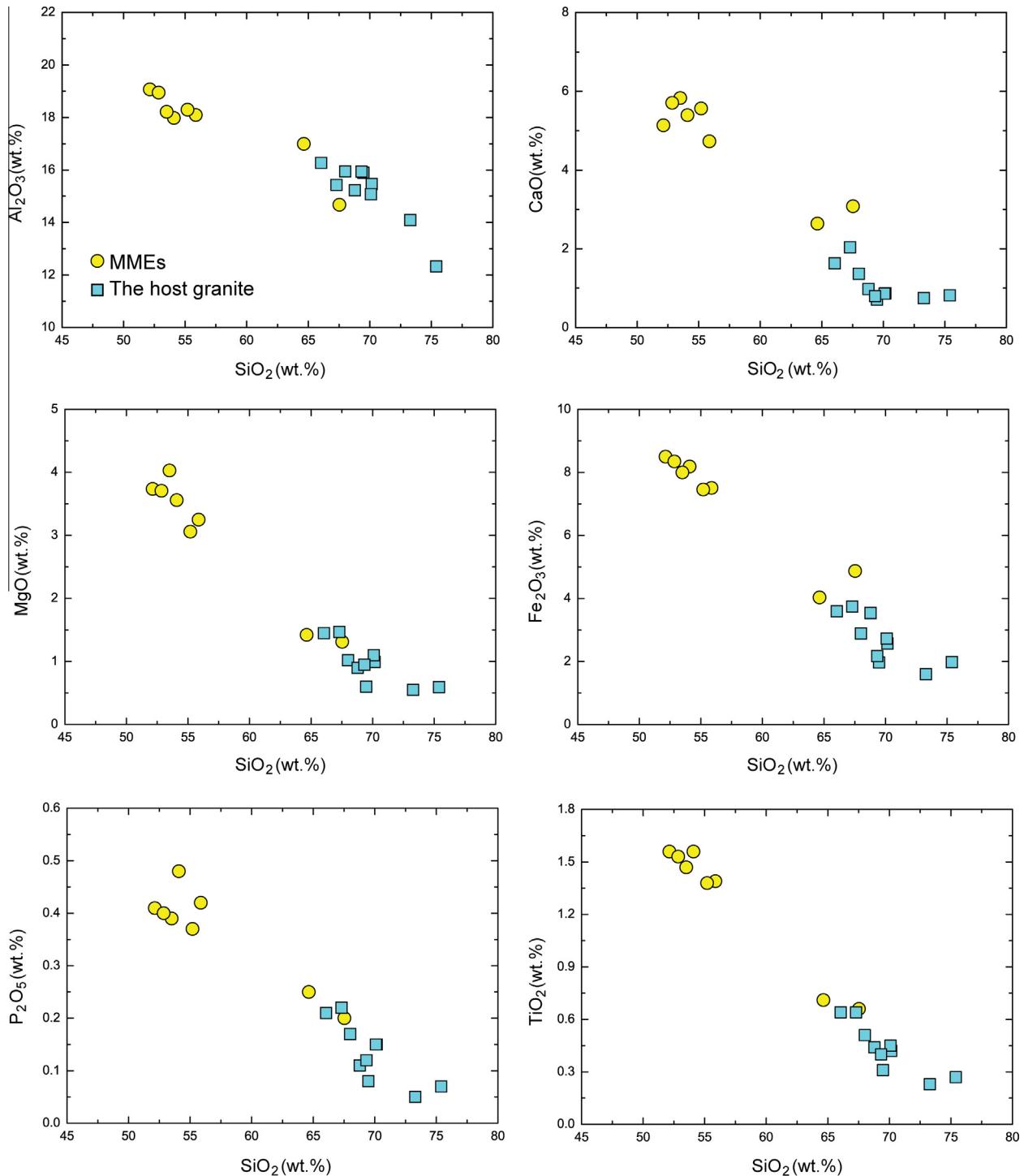


Fig. 7. Binary diagrams of oxides vs.  $\text{SiO}_2$  for the host granite and MMEs.

with the basement rocks, the Neoproterozoic Burd Gol accretionary complex that derived from the Baydrag block displays less negative initial  $\epsilon\text{Nd}(t)$  values ( $-11.5$  to  $-16.1$ ) (Kovalenko et al., 2005). However, both the Archean to Paleoproterozoic basement rocks and the Neoproterozoic Burd Gol accretionary complex cannot be the possible protolith due to their strongly negative  $\epsilon\text{Nd}(t)$  (Fig. 9a and b). In contrast, sedimentary rocks from the Bayankhongor zone and the Dzag zone exhibit slightly negative  $\epsilon\text{Nd}(t)$  and young Nd model ages, which are similar to those of the host granite. For instance, phyllites from the Huluut Bulag mélangé gave initial  $\epsilon\text{Nd}(t)$  values of  $-4.2$  to  $-3.2$  with Nd modal ages of 1.5–

1.7 Ga (Fig. 7b) (Kovalenko et al., 2005). Pelitic schists of the Dzag zone yielded initial  $\epsilon\text{Nd}(t)$  values of  $-3.1$  to  $-3.7$  with Nd model ages of 1.6–1.5 Ga (Jahn et al., 2004; Kovalenko et al., 2005). Therefore, we suggest that the felsic pelite is a potential candidate for the source of the host granite. This conclusion is further supported by whole-rock geochemistry. Jung and Pfänder (2007) proposed that  $\text{CaO}/\text{Na}_2\text{O}$  ratios of crust-derived melts strongly depend on the source compositions and can be used to distinguish between pelite-derived melts ( $\text{CaO}/\text{Na}_2\text{O} < 0.5$ ) and melts derived from greywackes or igneous rocks ( $\text{CaO}/\text{Na}_2\text{O}: 0.3\text{--}1.5$ ). Samples from the host granite have low  $\text{CaO}/\text{Na}_2\text{O}$  ratios (0.19–0.49, mostly

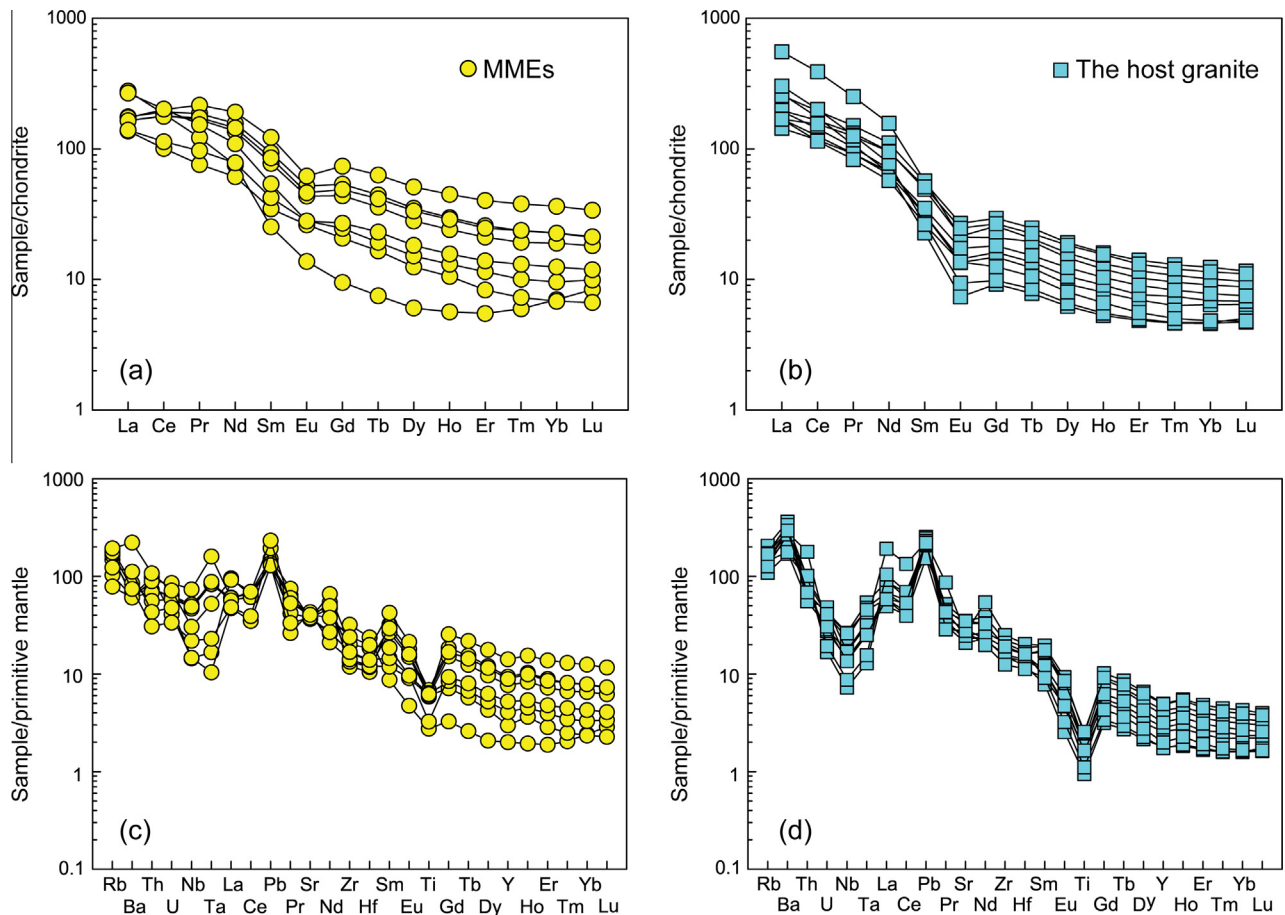


Fig. 8. Chondrite-normalized REE patterns (a and b) and primitive mantle-normalized multi-element patterns (c and d) for the host granite and MMEs. Normalizing values from Sun and McDonough (1989).

Table 2

Sm–Nd and Rb–Sr isotopic analytical results of the Ulaan Uul batholith and associated MMEs in the Bayankhongor area, central Mongolia.

Sample	Rb (ppm)	Sr (ppm)	$^{87}\text{Rb}/^{86}\text{Sr}$	$^{87}\text{Sr}/^{86}\text{Sr}$	$(^{87}\text{Sr}/^{86}\text{Sr})_i$	Sm (ppm)	Nd (ppm)	$^{147}\text{Sm}/^{144}\text{Nd}$	$^{143}\text{Nd}/^{144}\text{Nd}$	$(^{143}\text{Nd}/^{144}\text{Nd})_i$	$T_{\text{DM1}}(\text{Ga})$	$T_{\text{DM2}}(\text{Ga})$	$\epsilon_{\text{Nd}}(t)$
MG11-89-2	104	791	0.382	0.709188	0.706213	11.9	63.8	0.1124	0.512256	0.511854	1.35	1.41	–1.6
MG11-89-3	65.2	791	0.239	0.708483	0.706621	3.87	35.1	0.0667	0.512070	0.511831	1.12	1.44	–2.0
MG11-89-5	119	854	0.402	0.709466	0.706335	13.0	67.4	0.1168	0.512264	0.511846	1.39	1.42	–1.7
MG11-89-8	123	854	0.416	0.709255	0.706009	8.22	51.1	0.0972	0.512174	0.511826	1.28	1.45	–2.1
MG11-89-9	106	614	0.497	0.709769	0.705894	3.47	30.6	0.0687	0.512074	0.511827	1.13	1.45	–2.1
MG11-89-14	92.0	554	0.480	0.710208	0.706463	8.70	73.2	0.0718	0.512097	0.511840	1.13	1.43	–1.8
MG11-89-15	91.3	441	0.600	0.711120	0.706443	5.37	31.3	0.1038	0.512212	0.511840	1.30	1.43	–1.8
MG11-89-18	108	736	0.423	0.709353	0.706052	7.81	44.4	0.1062	0.512216	0.511835	1.33	1.44	–1.9

<0.3), suggesting a pelite-dominated origin. In addition, their high  $\text{Al}_2\text{O}_3/(\text{MgO} + \text{FeO}^{\text{T}})$  and low  $\text{CaO}/(\text{MgO} + \text{FeO}^{\text{T}})$  ratios also suggest a felsic pelite source (Fig. 13), because melts derived from metapelites possess higher  $\text{Al}_2\text{O}_3/(\text{MgO} + \text{FeO}^{\text{T}})$  but lower  $\text{CaO}/(\text{MgO} + \text{FeO}^{\text{T}})$  ratios than those from metabasalts and metagreywackes (Gerdes et al., 2002). Partial melting of felsic pelites needs a heat source, which can be provided by the underplating of a basaltic magma (represented by the MMEs). Thus, intrusion of Mesoproterozoic SCLM-derived basaltic magma created a thermal anomaly to trigger partial melting of the felsic pelites and subsequent mixing between the mantle- and pelite-derived melts resulted in the formation of the host granite.

### 5.3. Implication for the petrogenesis of high Ba–Sr granitoids

The Ulaan Uul batholith displays high Ba (1189–2544 ppm), Sr (440–745 ppm), K/Rb (354–418), but low Rb/Ba (<0.07), compara-

ble to those of typical high Ba–Sr granitoids (Fig. 11). High Ba–Sr granitoids share some common geochemical features: (1) high contents LILEs such as Ba, Sr and  $\text{K}_2\text{O}$ ; (2) crust-like Sr–Nd–Hf–Pb–O isotopic compositions (Tarney and Jones, 1994; Fowler and Rollinson, 2012; Peng et al., 2013). To explain these features, an origin from partial melting of an enriched SCLM metasomatized by sedimentary melts and/or fluids via subduction was proposed (Qian et al., 2003; Fowler et al., 2001, 2008; Fowler and Rollinson, 2012; Jiang et al., 2012; Peng et al., 2013). Terrigenous sediments-derived melts or aqueous fluids could carry voluminous Ba and Sr (Plank and Langmuir, 1998; Kilian and Behrmann, 2003; Portnyagin et al., 2007), and transfer of these elements would result in enrichment of associated lithospheric mantle. Large volume of sediments eroded from the continents will deposit on ocean floor and significantly recycle into the mantle through subduction. Moreover, almost all of the high Ba–Sr granitic intrusions in the world are located on continental margins. Therefore, it is

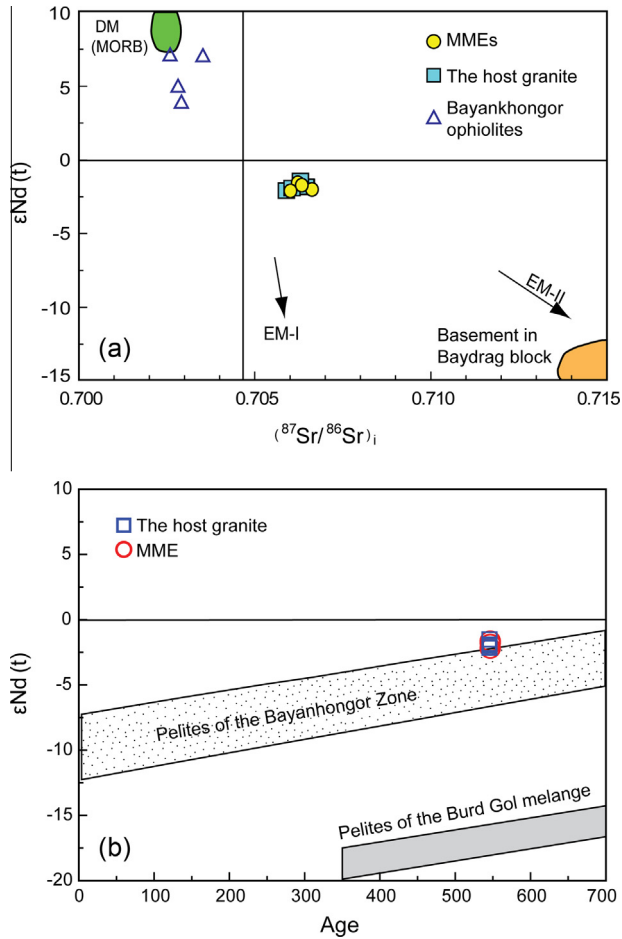


Fig. 9. (a) Diagram of  $\epsilon Nd(t)$  vs.  $(^{87}Sr/^{86}Sr)_i$  for the host granite and associated MMEs (modified after Jahn et al., 2009); (b)  $\epsilon Nd(t)$  vs. U–Pb ages for zircons from the host granite and associated MMEs (modified after Kovalenko et al., 2005).

reasonable to infer that terrestrial materials can significantly recycle into the mantle source in the form of sediment-derived fluids and/or melts via subduction along the continental margins, and that interaction of terrestrial sediment-derived fluid or melt can give rise to an enriched lithospheric mantle. Subsequent low-degree partial melting of the metasomatized lithospheric mantle will produce melts with high Ba–Sr signatures.

Our petrogenetic model for the Ulaan Uul batholith, i.e., melts of sedimentary materials and subsequent mixing with mantle-derived magma may shed new lights on the origin of high Ba–Sr granitoids. Our geochemical data indicate that the high Ba–Sr batholith was not entirely derived from an enriched lithospheric mantle as other high Ba–Sr granitoids in the world. In fact, rocks from the high Ba–Sr Ulaan Uul batholith display more remarkable sedimentary signatures than those of the high Ba–Sr appinites and granitoids from the Scottish Caledonides, as suggested by their higher A/CNK and A/NK ratios (Fig. 6c). As discussed above, the newly accreted sediments in the Haluut Bulag mélangé played an important role in the formation of the high Ba–Sr batholith. This model suggests that the high Ba–Sr signatures can be achieved not only by the recycle of terrigenous sediments in the mantle, but also by crust–mantle magma mixing process.

#### 5.4. Geodynamic implication

Generally, high Ba–Sr granitoid rocks have been considered to be emplaced in a post-collisional extensional setting caused by

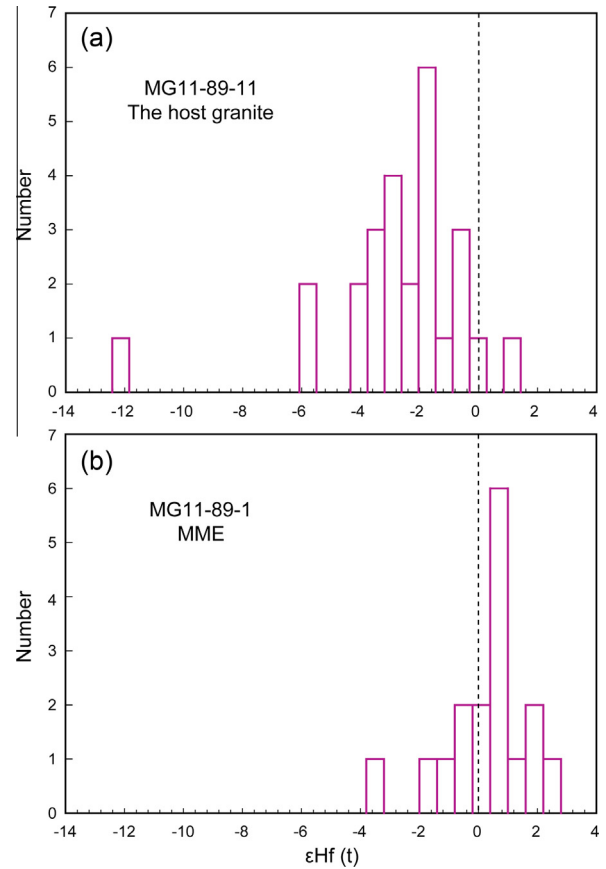


Fig. 10. Relative probability diagrams for  $\epsilon Hf(t)$  values of zircons from the host granite (a) and associated MMEs (b).

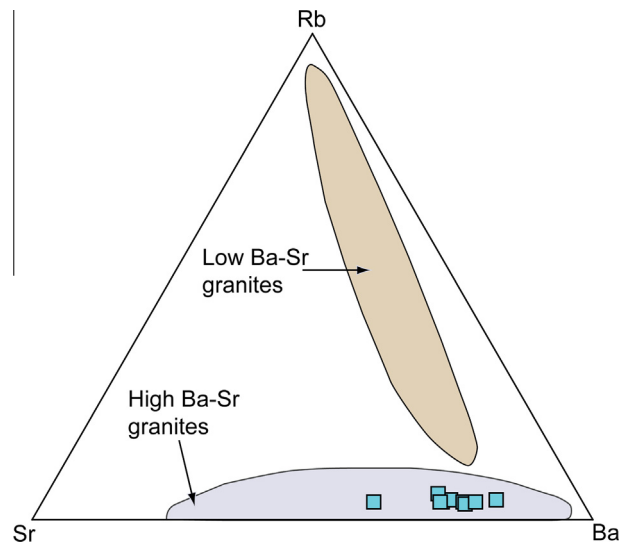
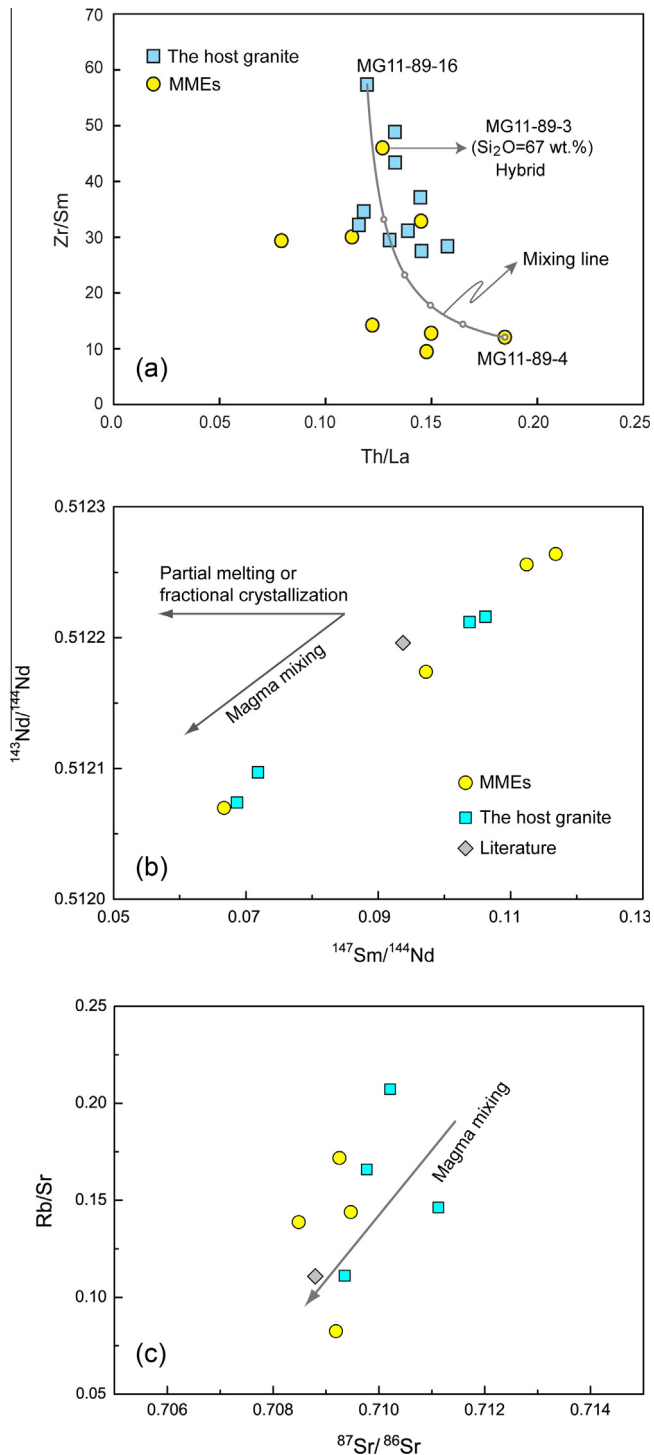


Fig. 11. Sr–Rb–Ba plot (modified after Tarney and Jones, 1994) for the host granite.

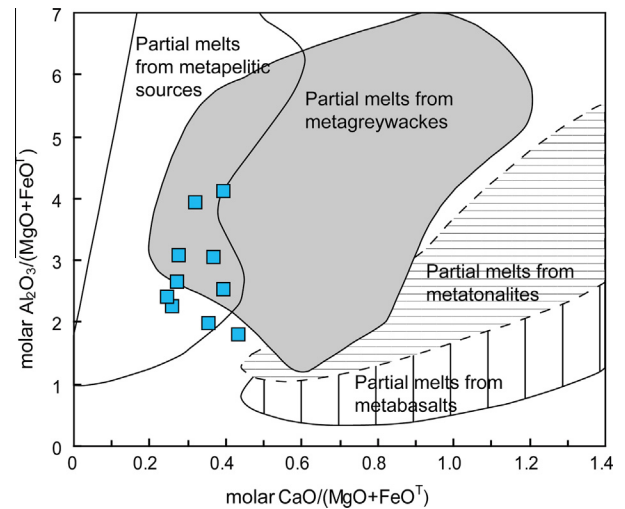
slab break-off or gravitational collapse (Qian et al., 2003; Fowler et al., 2001, 2008; Fowler and Rollinson, 2012; Ye et al., 2008; Heilimo et al., 2010). Recently, a subduction-related setting has been proposed to account for the formation of high Ba–Sr granitoids (González-Guillot et al., 2012; Peng et al., 2013; Cardona



**Fig. 12.** Selected geochemical variation diagrams for the host granite and MMEs. These variations strongly show that crust–mantle magma mixing takes place. (a) Th/La vs. Zr/Sm covariant diagram; (b) <sup>143</sup>Nd/<sup>144</sup>Nd vs. <sup>147</sup>Sm/<sup>144</sup>Nd and (c) Rb/Sr vs. <sup>87</sup>Sr/<sup>86</sup>Sr diagrams.

et al., 2014). In conjunction with regional geology and magmatic evolutionary history, we suggest that formation of the Ulaan Uul batholith with high Ba–Sr signatures was not related to a post-collisional setting, but rather formed in a subduction-related setting.

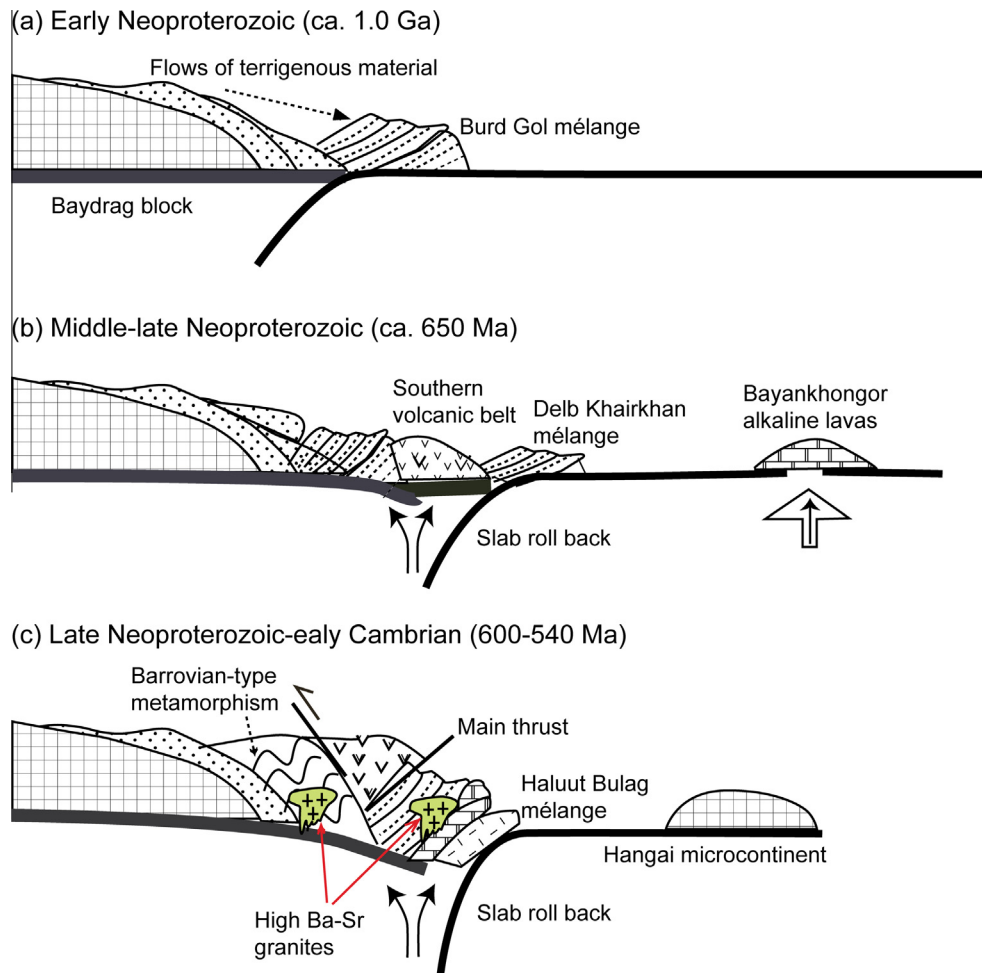
As described in Section 2.1, several tectonic terranes have been recognized in the Bayankhongor region. A striking feature is that the intervening mélanges between the Baydrag block and the



**Fig. 13.** Molar Al<sub>2</sub>O<sub>3</sub>/(MgO + FeO) vs. molar CaO/(MgO + FeO) diagram for the host granite (after Gerdes et al., 2002).

Hangai microcontinent decrease in age from south to north (Buchan et al., 2002; Demoux et al., 2009; Kröner et al., 2011), suggesting that the subduction zone migrated northwards. In the early Neoproterozoic, the Burd Gol mélangé developed on the northern margin of the Baydrag block, due to the subduction of the Bayankhongor oceanic lithosphere beneath the Baydrag block (Fig. 14a) (Demoux et al., 2009). Along with the sweep-back of the subducting Bayankhongor oceanic slab, the South Volcanic Belt and the Delb Khairkhan mélangé formed on a southward-dipping subduction zone in the middle to late Neoproterozoic (Fig. 14b) (Dergunov et al., 1997). Meanwhile, the Bayankhongor alkaline lavas were built on the oceanic crust, and subsequently accreted to the Delb Khairkhan mélangé (Fig. 14b and c).

Regional metamorphic and magmatic events also recorded the accretionary process of the intervening mélanges onto the Baydrag microcontinent. The Burd Gol mélangé underwent Barrovian-type metamorphism with metamorphic grade increasing northwards (Buchan et al., 2001), and metamorphic white micas from a pelitic schist gave a K–Ar age of  $699 \pm 35$  Ma reflecting an early-stage accretion of the Burd Gol mélangé onto the Baydrag block (Teraoka et al., 1996). In addition, a biotite–garnet schist from the Burd Gol mélangé yielded a biotite <sup>40</sup>Ar/<sup>39</sup>Ar plateau age of  $533 \pm 3$  Ma (Höck et al., 2000). This prolonged metamorphism probably reflects the punctuated southward accretion–collision. Recent zircon U–Pb dating results demonstrate that late Neoproterozoic to early Cambrian granitic magmatism was widespread in the Bayankhongor region (Buchan et al., 2002; Jahn et al., 2004; Kozakov et al., 2006, 2008; Demoux et al., 2009; Jian et al., 2010). Furthermore, it is found that the late Neoproterozoic (580–540 Ma) granitic intrusions are mainly distributed in the Baydrag block and Burd Gol mélangé, whereas those of early Cambrian (540–510) are widespread in the whole region. This trend clearly reflects the northward migration of a subduction zone. For instance, two gneissic granites that intruded the Burd Gol mélangé yielded zircon U–Pb ages of 579 and 560 Ma, respectively (Demoux et al., 2009). The Ulaan Uul batholith formed at  $546 \pm 3$  Ma and was probably related to the roll-back of the subducted Bayankhongor oceanic lithosphere. In this scenario, a descending slab migrating backwards would induce upwelling of the deep asthenosphere and cause a thermal anomaly in the mantle wedge. Subsequent partial melting of the Mesoproterozoic SCLM and underplating basaltic magma gave rise to partial melting of overlying sedimentary materials (Fig. 14c).



**Fig. 14.** Schematic diagrams for the Neoproterozoic to early Cambrian tectonic evolution of the Bayankhongor area, central Mongolia. (a) Southward subduction of the Bayankhongor oceanic lithosphere and the formation of the Burd Gol mélangé; (b) accretion of the Burd Gol mélangé onto the Baydrag block and a new subduction zone developed on the northern margin of the Burd Gol mélangé which account for the formation of the southern volcanic belt and the Delb Khairkhan mélangé; (c) due to the roll-back of the Bayankhongor oceanic lithosphere, the Bayankhongor ophiolite was accreted onto the Burd Gol mélangé by incorporation of two accretionary wedges (the Delb Khairkhan and Haluut Bulag mélangé), and the high Ba–Sr granites were emplaced into the mélangés.

## 6. Conclusions

- (1) LA–ICP–MS zircon U–Pb dating has illustrated that the ages of the host granite and MMEs are  $546 \pm 3$  Ma and  $547 \pm 3$  Ma, respectively, indicating that the mafic and felsic magmas were coeval.
- (2) The petrographic observations and geochemical data indicate that the parental magma of the MMEs was derived from an enriched Mesoproterozoic SCLM. The host granite, classified as high Ba–Sr granitoids, was produced by partial melting of felsic pelites triggered by upwelling of lithospheric mantle-derived magma (represented by MME) and subsequent mixing of the mantle- and sediment-derived melts.
- (3) Considering available regional tectonic and magmatic data, it can be concluded that the formation of the Ulaan Uul batholith and associated MMEs was related to an extensional setting, as a tectonic response to the roll-back of the subducted Bayankhongor oceanic lithosphere.

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## Appendix A. Supplementary material

Supplementary data associated with this article can be found, in the online version, at <http://dx.doi.org/10.1016/j.jseas.2014.11.029>.

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