Geochronology, geochemistry and tectonic significance of a Paleoproterozoic diabase at southwestern margin of the NCC

Huan Xu^{1,2}, Jin-hai Luo^{1,*}, Guan-xu Chen^{1,3}, Jia You⁴, and Yi-fei Li¹

¹ State Key Laboratory of Continental Dynamics, Department of Geology, Northwest University, Xi'an, China ² Shaanxi Railway Institute, Department of Surveying and Mapping Engineering, Weinan, Shaanxi, China ³ Yuncheng University, Department of Culture and Tourism, Yuncheng, Shanxi, China ⁴ The Second Oil Production Plant of Changqing Oilfield Company, Qingyang, Gansu, China

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ABSTRACT

The diabase and granite porphyry outcrop in the Baijiagou section within the Meso-Cenozoic Liupanshan fault zone at the southwestern margin of the North China Craton (NCC), and they have been regarded previously as Meso-Cenozoic dykes according to their occurrence. Zircon U-Pb LA-ICP-MS age determining for the diabase and granite porphyry yield respectively 1804 ± 21 and 1792 ± 16 Ma, and they form a bimodal magmatic assemblage of Paleoproterozoic. The diabase can be classified into two major geochemical types, which are low-Ti (LT) and high-Ti (HT) in nature. The former is characterized by relatively high SiO₂ (48.81 - 54.01 wt.%), P₂O₅, and Al₂O₃, and low TiO₂ (1.41 - 1.54 wt.%), FeO, MnO, and CaO, and enriched whole-rock Sr and Nd isotopic composition $[({}^{87}Sr/{}^{86}Sr)_i = 0.703 - 0.708; \epsilon_{Nd} = -3.76$ to -4.37]. The latter has lower SiO₂ (45.9 - 51.19 wt.%), P_2O_5 , and Al_2O_3 , and higher TiO_2 (1.61 - 2.26 wt.%) and FeO than that of the former, and depleted Sr and Nd isotopic composition $[({}^{87}\text{Sr}/{}^{86}\text{Sr})_i = 0.691 - 0.708; \epsilon_{Nd} = 3.27 - 5.03]$. Both types of diabase exhibit light rare earth element enrichment and flat middle-heavy rare earth element patterns. The LT diabase has negative Eu anomalies and higher total REE contents than the HT diabase. Th/Nb ratios of the LT and HT diabases are 0.28 - 0.29 and 0.08 - 0.10, respectively, which are similar to basalts formed in an intraplate spreading setting in association with a mantle plume. The regional geology suggests that intracontinental rifting occurred at the southwestern margin of the NCC in the Paleoproterozoic, and that the rifting was related to a mantle plume. This rifting event was part of the break-up of the Columbia supercontinent.

1. INTRODUCTION

The North China Craton (NCC) includes a history that are up to 4.0 Ga old and most of the rocks in the NCC formed at 3.0 - 2.5 Ga (Zhu et al. 2012), making it one of the oldest cratons in the world. During the late Paleoproterozoic and early Mesoproterozoic orogenesis, uplift, and extension happened to the NCC (Zhai and Santosh 2011, 2013; Zhai et al. 2014). The NCC is divided into eastern and western blocks, separated from each other by the Central Orogenic Belt, which formed prior to 1.85 Ga (Zhao et al. 2005; Jiang et al. 2010). The western block comprises the Yinshan and Ordos blocks (Zhao et al. 2005), which amalgamated at ~1.95 Ga (Li et al. 2013). After 1.8 Ga, a mantle plume caused uplift and extension in the NCC, resulting in rifting and the development of the Helan, Xiong'er, and Yanshan-Taihang aulacogens (Li et al. 2000; Zhao et al. 2007; Gao et al. 2013). Late Paleoproterozoic and Mesoproterozoic rifting was accompanied by extensive magmatism, including extrusion of the Xiong'er Group volcanic rocks (1.80 - 1.75 Ga; Zhao et al. 2004; Xu et al. 2007) and intrusion of coeval granodiorites and porphyries (Zhao et al. 2007; Cui et al. 2010), mafic dike swarms of the Xiong'er aulacogen (Hou et al. 2010), and mafic dike swarms and alkaline intrusive rocks of the Yanshan-Taihang aulacogen (1.77 - 1.78 Ga; Peng et al. 2004; Yang et al. 2005; Dong et al. 2010). The distribution of Paleoproterozoic magmatic rocks in the North China Craton is shown as Fig. 1,

^{*} Corresponding author

E-mail: luojh@nwu.edu.cn

and the related data are in Table 1. The late Paleoproterozoic extension event has been extensively studied, although the mechanisms driving the extension are not well understood. Peng et al. (2008) and Zhai et al. (2014) argued that extension was related to mantle plume activity, and speculated that the southern margin of the NCC (Xiong'er Group volcanic rocks) was the locus of the mantle plume.

The Proterozoic geology of the southern part of the Helanshan and Liupanshan Mountains in this Paleoproterozoic aulacogen is obscured by Meso-Cenozoic metasedimentary strata and the reactivation of the Liupanshan fault system. This has limited our understanding of the Paleoproterozoic and Mesoproterozoic tectonic framework in this region (Che et al. 2011). The Helan aulacogen has been identified at the western margin of the NCC using seismic data, and also from sedimentological observations in the northern part of the Helanshan area. Seismic profiles have revealed a series of extensional faults in the Proterozoic successions, indicating that the 4000-5000-m-thick Mesoproterozoic strata were controlled by syn-sedimentary faulting (Wang 1995; Hua and Qiu 2001). Similar lithological and tectonic characteristics have been identified in the Yanliao aulacogen (Wang 1995; Hua and Qiu 2001; Gao et al. 2010). The limited study of Paleoproterozoic igneous rocks at the southwestern and western margins of the NCC is a barrier to improving our understanding of the evolution of the NCC. Here we present petrological, geochronological, and geochemical data for the Baijiagou diabase and granite porphyry from the southwestern margin of the NCC. We use these data to constrain the petrogenesis and tectonic setting in which the diabase was emplaced, and show that extensional tectonism affected the NCC and formation of the Helan aulacogen in the Paleoproterozoic.

2. REGIONAL GEOLOGY

The Helan aulacogen trends approximately N-S and forms a valley located between the Erdos and Alxa blocks at the western margin of the NCC. During the Proterozoic, the Helan aulacogen extended into the NCC and was filled with littoral clastic sediments and carbonates. The sediments are ~1000 m thick and were overlain by sediments of the south Qilin Trough in the Late Ordovician (Che et al. 2011). Geochronological studies of khondalite-series rocks in the northern Helan Mountains have yielded metamorphic ages of 1.9 - 2.1 Ga (Geng et al. 2010; Zhou et al. 2010; Xiao et al. 2011). The khondalite rocks are the product of Paleoproterozoic crustal development, and intrusive rocks in this area yield emplacement ages of ca. 1.85 Ga (Geng et al. 2010); however, igneous rocks are poorly exposed in the southern part of the Helan aulacogen.

The Baijiagou diabase and associated granite porphyry are exposed in Longxian Country, Shaanxi Province (Fig. 2). The bimodal magmatism potentially provides important insights into the Paleoproterozoic geological evolution of the NCC. Based on contact relationships between the intrusive and country rocks, Che and Wang (1985) suggested that the Baijiagou diabase and granite porphyry are Late Cretaceous to early Paleogene in age, and concluded that their formation was associated with the Liupanshan fault system. The 1:50000 scale geological map of Guguan Town (1996) followed the findings of Che and Wang (1985), and assigned the diabase and granite porphyry a Paleogene age. Our field observations indicate that the sedimentary strata in the study area are Upper Ordovician clastic limestones, Upper Triassic clastic sediments, Lower Cretaceous sandstones, and Quaternary sediments. Faults are widely developed. The contacts between the intrusive and country rocks are generally covered by Quaternary sediments; however, it is clear that metamorphism did not affect the Upper Ordovician limestones. You et al. (2014) obtained an upper intercept U-Pb zircon age of 1846 ± 32 Ma for the Baijiagou granite porphyry, indicating that it was emplaced during the Paleoproterozoic, whereas the diabase has not yet been dated.

3. PETROGRAPHY

There are only a few small outcrops ($< 20 \text{ m}^2$) of diabase and granite porphyry in the Baijiagou area, which have been exposed by river erosion. In almost all outcrops, the relationship between the granite porphyry and diabase is evident (Figs. 3a - c). The diabase is in direct contact with the granite porphyry, and the latter contains diabase xenoliths (Fig. 3b). In addition, granite porphyry veins have intruded into the diabase (Fig. 3c). The diabase samples are greyish green in color (Fig. 3a), massive in structure, and have ophitic textures (Figs. 3d, e). The diabase phenocrysts are predominantly calcic plagioclase and pyroxene, and comprise \sim 70% of the rock. The plagioclase (40%) is idiomorphic and weakly altered. Some plagioclase crystals display simple and polysynthetic twinning. Pyroxenes (30%) are allotriomorphic and infill space between the plagioclase crystals. The pyroxene is locally replaced by mica minerals. Secondary minerals are hornblende and biotite (15%), which are allotriomorphic. The main accessory mineral is magnetite.

The granite porphyry samples are red in color and have a porphyritic texture (Fig. 3f). Phenocrysts are predominantly K-feldspar (25%). The feldspar is altered and occasionally exhibits simple and polysynthetic twinning. Quartz phenocrysts comprise 20% of the rock, and are unaltered with corroded and embayed edges. Plagioclase is weakly altered and comprises 10% of the rock. Secondary biotite is present and the matrix is aphanitic or microcrystalline, comprising feldspar and quartz.

We have collected twelve samples for both of the diabase and granite porphyry, ten of which are for geochemical analysis, one is for chronological analysis, and one is for petrographic analysis.



Fig. 1. Paleoproterozoic magmatic rocks distribution diagram in the North China Craton (after Zhao et al. 2015).

Position	Rock type	Method	Age (Ma)	Reference					
① Baijiagou	granite porphyry	U-Pb LA-ICP-MS	1852 ± 24	You et al. 2014					
② Tiemahe	granite porphyry	U-Pb LA-ICP-MS	1814 ± 12	Xu et al. 2014					
③ Shizuizi	granite porphyry	SHRIMP	1778 ± 14	Wang et al. 2012a					
		U-Pb LA-ICP-MS	1803 ± 15	Gao et al. 2013					
4 Guandishan	granite	SHRIMP	1798 ± 11	Zhao et al. 2008					
Tangershang	granite	SHRIMP	1790 ± 14						
⁽⁵⁾ Hengshan	basic dike	U-Pb LA-ICP-MS	1769 ± 2.5	Li et al. 2001					
		SHRIMP	1778 ± 3	Peng et al. 2005					
⁽⁶⁾ Baiyan obo	basalt	U-Pb LA-ICP-MS	1743	Lu et al. 2002					
⑦ Fengzhen	basic dike	SHRIMP	1769 ± 4	Han et al. 2007					
⑧ Guyang	quartz syenite	U-Pb LA-ICP-MS	1702 ± 31	Wang et al. 2012b					
④ Miyun	rapakivi granite	U-Pb LA-ICP-MS	1681 ± 10	Yang et al. 2005					
10 Miyun	basic dike	SHRIMP	1731 ± 4	Peng et al. 2011					
1 Damiao	monzonite	SHRIMP	1739 ± 14	Zhao et al. 2009					
12 Jianping	clinopyroxene monzonite	U-Pb LA-ICP-MS	1720 ± 4	Wang et al. 2013					
	quartz syenite	U-Pb LA-ICP-MS	1715 ± 4						
① Damiao	anorthosite	U-Pb LA-ICP-MS	1726 ± 9	Zhang et al. 2007					
(1) Beigukou	syengranite	U-Pb LA-ICP-MS	1753 ± 23	Zhang et al. 2007					
15 Chicheng	rapakivi granite	U-Pb LA-ICP-MS	1697 ± 7	Jiang et al. 2011					
16 Datong	basic dike	SHRIMP	1777 ± 3	Peng et al. 2005					
		SHRIMP	1789 ± 28						
17 Luyashan	adamellite	U-Pb LA-ICP-MS	1794 ± 13	Geng et al. 2004					
18 Songshan	basic dike	SHRIMP	1785 ± 18	Hu et al. 2010					
19 Motianzhai	granite	SHRIMP	1797 ± 14	Zhao and Zhou 2009					
2 Xiong'ershan	basic dike	SHRIMP	1773 ± 37	Zhao et al. 2004					
2 Xiong'ershan	subvolcanics	U-Pb LA-ICP-MS	1781 ± 12	Liu 2011					
	rhyolite	U-Pb LA-ICP-MS	1763 ± 15	Wang et al. 2010					
2 Xiong'ershan	syenite	U-Pb LA-ICP-MS	1750 ± 65	Ren et al. 2000					
② Xiong'ershan	rhyolite porphyry	U-Pb LA-ICP-MS	1800 ± 16	Zhao et al. 2004					

Table 1. Magmatic rocks of Paleoproterozoic in the North Chian Craton.



Fig. 2. Geological map of the Baijiagou region in Longxian (modified after the 1:50000 Geological Map of Guguan). 1: Quaternary sediments; 2: Lower Cretaceous sandstone; 3: Upper Triassic sandstone; 4: Upper Ordovician limestone; 5: Middle Ordovician limestone; 6: diabase; 7: granite; 8: fault; 9: inferred fault; 10: sinistral strike-slip fault; 11: sample locality.



Fig. 3. (a) (b) (c) Field contacts between the diabase and granite porphyry; (d) (e) microphotographs of the diabase; (f) microphotograph of the granite porphyry. Af = alkali feldspar; Bt = biotite; Or = orthoclase; Qz = quartz.

4. ANALYTICAL METHODS

All analyses were performed at the State Key Laboratory of Continental Dynamics, Northwest University, Xi'an, China.

4.1 Zircon U-Pb LA-ICP-MS Analysis

Zircons were separated using heavy liquid and magnetic separation techniques, and then handpicked under a binocular microscope. The selected grains were mounted in epoxy resin and polished until their crystal centers were exposed. Cathodoluminescence (CL) images showing the internal structure of the zircon grains were captured using a Quanta 400FEG environmental scanning electron microscope, equipped with an Oxford energy dispersive spectroscopy system and a Gatan CL3+ detector.

U-Pb dating of zircons was conducted with an Agilent 7500a inductively coupled plasma-mass spectrometer (ICP-MS) coupled to a 193 nm ArF excimer laser, flow homogenizer, and optical imaging system. A fixed spot size of 30 µm and laser repetition rate of 6 Hz were used for all analyses. Helium was used as the carrier gas to provide efficient aerosol delivery to the plasma ion source. The National Institute of Standards and Technology (NIST) Standard Reference Material (SRM) 610 was used to optimize the instrument to maximize sensitivity (238U signal intensity of > 460 cps/ μ g·g⁻¹) and minimize oxide production (ThO/ Th < 1%). The U/Th ratio was used to monitor whether zircon vaporization was complete (Günther and Hattendorf 2005). Analyses were made in time-resolved mode using both analog and pulsed detection, and fast peak jumping (20 ms peak-1). GLITTER 4.0 software developed by Macquarie University was used to calculate ²⁰⁷Pb/²⁰⁶Pb, ²⁰⁶Pb/²³⁸U, ²⁰⁷Pb/²³⁵U, and ²⁰⁸Pb/²³²Th ratios, which were corrected for instrumental mass bias and depth-dependent elemental and isotopic fractionation using the 91500 zircon as an external standard (recommended 206Pb/238U age of 1065.4 ±0.6 Ma; Wiedenbeck et al. 2004). Concordia diagrams and weighted mean age calculations were made using Isoplot (v. 2.49; Ludwig 2001). U, Th, and Pb concentrations were calculated using ²⁹Si as an internal standard and NIST SRM610 as an external standard. The zircon standards 91500 and GJ-1 yielded weighted mean 206 Pb/ 238 U ages of 1064.4 ± 3.1 Ma (n = 16; 2σ) and 603.1 ± 3.2 Ma (n = 14; 2σ), respectively, which are in good agreement with the recommended isotope dilution-thermal ionization mass spectrometry (ID-TIMS) ages reported by Wiedenbeck et al. (1995).

4.2 Whole-Rock Major and Trace Element Analyses

Altered surfaces were removed from selected samples prior to sample preparation for geochemical analysis. The samples were then washed, dried, and crushed to 60 mesh size in an alumina jaw crusher. Approximately 60 g of each crushed sample was milled in a tungsten carbide mill (T1-100; CMT) to < 200 mesh (75 μ m) for analysis.

Major element contents were determined by X-ray fluorescence (XRF; Rigaku RIX 2100) spectrometry on Liborate glass discs. Trace elements were measured using an Agilent 7500a ICP-MS after 50 mg samples had been dissolved in sealed high-temperature and high-pressure bombs using equal volumes of ultrapure HF and HNO₃. Analyses of United States Geological Survey (USGS) rock standards BHVO-2 and AGV-2 indicate that the accuracy and precision of the XRF and ICP-MS data were better than 5%.

4.3 Rb-Sr and Sm-Nd Isotope Analyses of Whole Rock

Whole rock Rb-Sr and Sm-Nd isotope ratios were determined using a Micromass MC-ICP-MS Nu Plasma HR. Separation of Sr and Nd was done by using a routine two-columnion ion-exchange technique. Isotopic ratios were measured on Isoprobe mass spectrometer in a static mode. During our analyses, ¹⁴³Nd/¹⁴⁴Nd ratios of 0.512120 \pm 12 (2 σ) were obtained for the international standard Shin Etou JNdi-1and ⁸⁷Sr/⁸⁶Sr ratios of 0.710247 \pm 17 (2 σ) for the international standard NBS-987. Correction of isotopic mass fractionation for Nd and Sr were made with ¹⁴⁶Nd/¹⁴⁴Nd = 0.7219 and ⁸⁶Sr/⁸⁸Sr = 0.1194, respectively (Wei et al. 2002; Liang et al. 2003).

5. RESULTS

5.1 Zircon U-Pb Geochronology

Zircon U-Pb dating was carried out on diabase sample 12LX26 (Table 2) and granite porphyry sample 14LP15 (Table 3). Zircons from the diabase are smaller, euhedral, short columnar in shape, darker in CL images, and have broad zoning that is not oscillatory (Fig. 4). These are interpreted to be the primary zircon from the diabase, and have high U and Th contents, and high Th/U ratios (0.28 - 4.75) indicative of a magmatic origin (Rubatto 2001). Zircons from the granite porphyry are intermediate in size, euhedral-subhedral, columnar in shape, and exhibit oscillatory zon-ing (Fig. 5). The granitic zircon grains are larger than those from the diabase, and have Th/U ratios of 0.49 - 1.08, which are much higher than those of metamorphic zircon.

Twenty zircon grains from the diabase were analyzed, eleven reliable points can form an inconsistency line. From a concordia plot (Fig. 6), an upper intercept ²⁰⁷Pb/²⁰⁶Pb age of 1804 \pm 21 Ma (MSWD = 0.19) was determined, and the weighted mean age is 1803 \pm 15 Ma (n = 8; MSWD = 0.46). The two ages are the same within the error range, so the age of the upper intersection can be used to represent the crystallization age of the diabase. Forty zircon grains from the granite porphyry were analyzed and thirty nine yielded concordant or near-concordant ages (Fig. 7a), and the upper intercept age is 1792 ± 16 Ma (MSWD = 0.99), with a weighted main ²⁰⁷Pb/²⁰⁶Pb age of 1800 ± 8 Ma (MSWD = 1.9) (Fig. 7b). Using the upper intersection age represents the crystallization age of the granite porphyry. Thus, both the diabase and granite porphyry are Paleoproterozoic in age and have identical ages when analytical uncertainties are considered.

5.2 Major Element Geochemistry of the Baijiagou Diabase

Major and trace element data are presented in Supplementary Table 4. Due to the high loss on ignition (LOI) values, which may reflect alteration, the analytical results were recalculated on a LOI-free basis.

The TiO₂, P₂O₅, Al₂O₃, and FeO contents of the diabase samples allow identification of two sample types: low-Ti (LT) and high-Ti (HT). The LT type is characterized by relatively high SiO₂ (48.8 - 54.0 wt.%), P₂O₅ (0.40 - 0.43 wt.%), and Al₂O₃ (14.4 - 16.3 wt.%), and low TiO₂ (1.41 - 1.54 wt.%), FeO^T (10.1 - 11.9 wt.%), MnO (0.14 - 0.15 wt.%), and CaO (3.2 - 5.6 wt.%). The HT type has lower SiO₂ (45.9 - 51.2 wt.%), P₂O₅ (0.14 - 0.20 wt.%), and Al₂O₃ (12.2 - 14.0 wt.%), and higher TiO₂ (1.61 - 2.26 wt.%), FeO^T (12.4 - 16.1 wt.%), MnO (0.11 - 0.32 wt.%), and CaO (5.4 - 13.3 wt.%). The K₂O contents of the two types of diabase are low (average = 1.57 wt.%), but Na₂O contents are relatively high (average = 3.65 wt.%). In a K₂O-SiO₂ diagram, data for the samples plot in the alkaline-subalkaline field (Fig. 8a). The MgO/(MgO + FeO^T) ratios vary from 0.21 -0.31, comparable to those for continental basalts (typically < 0.7; Wilson 1989). An AFM diagram shows that the diabase rocks belong to the tholeiitic series (Fig. 8b).

5.3 Trace Element Geochemistry of the Baijiagou Diabase

Chondrite-normalized rare earth element (REE) patterns for the diabase samples are shown in Fig. 9 (gray line represents E-MORB). Both types of diabase are characterized by light REE enrichment, with this being most pronounced in the LT diabase. Middle to heavy REE patterns are flat for both diabase types. The LT type has a negative Eu anomaly ($\delta Eu = 0.76 - 0.90$) and total REE abundances from 210 - 231 ppm. The HT type only has a slightly negative or no Eu anomaly ($\delta Eu = 0.82 - 1.10$), with total REE abundances of 66 - 97 ppm.

In a mid-ocean ridge basalt (MORB)-normalized multielement (spider) diagram (Fig. 10a), the diabase samples are enriched in K, Rb, Ba, Th, and Ce relative to MORB. In addition, the LT type is characterized by depletions of Nb, Ta, and Ti relative to the HT type. In a chondrite-normalized spider diagram (Fig. 10b), there is a significant negative Sr anomaly in the LT type samples. The Th/Nb ratios of the LT and HT diabases are 0.30 - 0.29 and 0.08 - 0.1, respectively, and their Th/Ta ratios are 5.83 - 5.58 and 1.33 - 1.65, respectively.

5.4 Sr-Nd Isotopic Composition

After the analysis of major and trace element, all the samples were analyzed for Rb-Sr and Sm-Nd isotopes (Table 5). Initial ⁸⁷Sr/⁸⁶Sr and ¹⁴³Nd/¹⁴⁴Nd ratios are expressed as $I_{Sr}(t)$ and $\varepsilon_{Nd}(t)$ values respectively, where t is the age of magma crystallization. The calculated Nd model ages are denoted as $T_{\rm DM1}$ relative to a depleted mantle source for single-stage evolution and as T_{DM2} relative to the average continental crust for two-stage evolution (Jahn and Condie 1995). The HT type diabase shows small variations in initial Nd isotope ratios at t = 1.8 Ga, 3.27 - 5.03 for $\varepsilon_{Nd}(t)$ values, and T_{DM2} of 1.96 - 2.06 Ga. The initial Sr isotope ratios shows large variations, 0.691 - 0.708 for $I_{sr}(t)$ values. Sr isotope system is relatively unstable to Nd and the former is easily modified by later geological actions. The initial Nd isotope ratios at t = 1.8 Ga for the LT type diabase are range from -3.76 to -4.37, 0.703 - 0.708 for $I_{Sr}(t)$ values, and T_{DM2} of 2.61 - 2.65 Ga.

5.5 Major and Trace Element Geochemistry of the Baijiagou Granite Porphyry

The major and trace element data for the granite porphyry are presented in Supplementary Table 6. The average SiO_2 content of the granite porphyry is 70.24 wt.%. SiO_2 contents of a small number of samples are low, and may have been affected by the diabase. The average total alkalis $(K_2O + Na_2O)$ content is 5.6 wt.%, and A/CNK values of 0.46 - 1.19 indicate that the granites are aluminous to weakly peraluminous. The granite samples are characterized by light REE enrichment and negative Eu anomalies $(\delta Eu = 0.44 - 0.51)$, and are relatively depleted in highfield-strength elements (HFSEs; e.g., Nb, Ta, Ti, and Hf) and enriched in large-ion lithophile elements (LILEs; e.g., K, Rb, and Ba). The granites have high Ga/Al ratios, and $10000 \times \text{Ga/Al}$ values ranges from 2.95 - 5.32 (average = 3.96), much higher than the mean values for I- and S-type granite (2.1 and 2.28, respectively) (Whalen et al. 1987). These geochemical features show that the Baijiagou granite porphyry is an A-type granite, which is consistent with the findings of You et al. (2014).

6. DISCUSSION

6.1 Intrusive Age of the Baijiagou Diabase

The age of the Baijiagou diabase was previously estimated to be Late Cretaceous to early Paleogene, as inferred from contact relationships (Che and Wang 1985). We identified direct contacts between the Baijiagou diabase and

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granite porphyry, but the contact relationships with the surrounding strata are generally covered by Quaternary sediments and/or are faulted. It is thus difficult to determine the age of the diabase according to contact relationships. In addition, the country rock (Upper Ordovician calclithite) near the diabase is deformed but not hydrothermally altered, indicating that the diabase pre-dates the Upper Ordovician. The zircon U-Pb isotopic system has a high closure temperature and strong resistance to resetting by late-stage geological processes. Zircons from the diabase yield an upper intercept ²⁰⁷Pb/²⁰⁶Pb age of 1804 ± 21 Ma (MSWD = 0.19), which is within error of the age obtained on the zircons from the granite porphyry [1792 ± 16 Ma (MSWD = 0.99)]. As such, both the diabase and granite porphyry are Paleoproterozoic rocks.

The Tiemahe granite porphyry in Shaanxi province $(1814 \pm 12 \text{ Ma}; \text{Xu et al. 2014})$, the Shizuizi granite in Ningxia province $(1778 \pm 14 \text{ and } 1803 \pm 15 \text{ Ma}; \text{Wang et al.}$ 2012a; Gao et al. 2013) and the Duanjiaxia granite porphyry $(1846 \pm 32 \text{ Ma}; \text{You et al. } 2014)$ are exposed to the north of the present study area. Diabase and basalt associated with granite porphyry are also present in the Tiemahe rocks, but their ages are unknown. These various granites are linearly distributed along the Liupanshan fault system, and the Baijiagou diabase and granite porphyry, Tiemahe granite porphyry, and Shizuizi granite can be considered the products of the same Paleoproterozoic magmatic event, which indicates that there is an important magmatic event during Paleoproterozoic at southwestern margin of the NCC. This Paleoproterozoic igneous belt (i.e., the Helan aulacogen) along the Liupanshan fault system was partially uplifted and exposed by fault movement.

6.2 Petrogenesis of the Baijiagou Diabase

The mineralogical and geochemical characteristics of

mafic igneous rocks are controlled by source temperaturepressure conditions and compositions, lithospheric thickness, and shallow-level processes (e.g., crystal fractionation). The Baijiagou diabase samples show geochemical variations that are further explored in this section using qualitative and quantitative approaches.

6.2.1 Crustal Contamination

The Baijiagou diabase can be classified into LT and HT types. It is generally believed that if the source of mantle basic magma subjected to continental crust material contamination, magma of incompatible elements such as La or Th will be significantly higher, relative to the Zr or Nb. Therefore the ratios of Zr/Nb and Nb/La are used to determine the crust contamination (Xu and Qiu 2010). The Nb/La ratios of the LT type diabase range from 0.22 - 0.23 (i.e., much less than 1), whereas the Nb/La ratios of the HT type diabase range from 0.81 - 1.13, similar to mantle values; the Th/Nb rations of the LT type diabase are about 0.3 (the ration of continental crust is 0.44, Weaver 1991) and the rations of the HT type diabase range from 0.08 - 1 (the ration of mantle is 0.117 - 0.071, Weaver 1991). The trace elements rations suggest that the crustal contamination influenced the LT type diabase, but not the HT type. Such contamination may reflect contributions from enriched lithospheric mantle and/ or lower crust, which resulted in LILE enrichment and HFSE depletion relative to the HT type diabase. Crustal contamination also results in higher Al₂O₃ and P₂O₅ contents.

In the process of emplacement, mantle-derived magma is contaminated by assimilation of crustal materials, and the ratios of Sr and Nd isotopes will evolve toward the crust. The Nd isotopic versue SiO₂ and La/Nb diagrams (Fig. 11) can identify the crustal contamination, easily. From Fig. 11, LT type diabase was obviously contaminated by the crust. According to Fig. 12, in addition to one point of the initial

~ .		222555	²³² Th ²³⁸ U		Istopic ratios						Ages/Ma					
Spot Pb [*]	Pb [*]	²³² Th		Th/U	²⁰⁷ Pb/ ²⁰⁶ Pb		²⁰⁷ Pb	b/ ²³⁵ U ²⁰⁶ Pb		0/ ²³⁸ U ²⁰⁷ Pl		⁰⁷ Pb/ ²⁰⁶ Pb		²⁰⁷ Pb/ ²³⁵ U		²⁰⁶ Pb/ ²³⁸ U
110.		(hhm)	(ppm)		ratios	1σ	ratios	1σ	ratios	1σ	ages	1σ	ages	1σ	ages	1σ
2	2703	1986	7177	0.28	0.11199	0.00382	5.28361	0.15331	0.342	0.00523	1832	31	1866	25	1896	25
3	3286	39524	43497	0.91	0.12402	0.00274	4.29602	0.05493	0.2511	0.00254	1813	18	1693	11	1444	13
4	1840	22824	9019	2.53	0.10967	0.00285	4.92186	0.09209	0.32533	0.0038	1794	18	1806	16	1816	18
6	3802	50630	20325	2.49	0.11078	0.0028	4.18513	0.07408	0.27388	0.0031	1812	17	1671	15	1560	20
7	3646	20737	8390	2.47	0.10985	0.00316	4.72388	0.10563	0.31178	0.00396	1797	23	1771	19	1749	17
8	371	2846	1111	2.56	0.10889	0.00287	4.19913	0.0805	0.2796	0.00328	1781	19	1674	16	1589	15
10	6152	43198	24636	1.75	0.11109	0.00324	4.63937	0.10621	0.30282	0.00393	1817	23	1756	19	1705	19
12	1164	33050	6964	4.75	0.11016	0.00999	3.28629	0.28666	0.21632	0.0077	1802	67	1478	68	1262	18
14	2073	21005	8914	2.36	0.11041	0.00403	4.29201	0.13489	0.28194	0.00442	1806	35	1692	26	1601	18
15	5366	28061	101833	0.28	0.11053	0.0028	3.56664	0.06285	0.23403	0.00264	1808	17	1542	14	1356	18
17	1121	25140	15712	1.60	0.11034	0.00346	4.11476	0.10457	0.27048	0.00369	1805	27	1657	21	1543	19

Table 2. Zircon U-Pb isotope data obtained by LA-ICP-MS for diabase (12LP26) from Baijiagou.

<i>a</i> .		222000	2287.7		Istopic ratios						Ages/Ma					
Spot	Pb [*]	²³² Th (nnm)	²³⁸ U	Th/U	²⁰⁷ Pb	/ ²⁰⁶ Pb	²⁰⁷ Pb	∕ ²³⁵ U	²⁰⁶ Pb	∕ ²³⁸ U	²⁰⁷ Pb/ ²	⁰⁶ Pb	²⁰⁷ Pb/2	²³⁵ U	²⁰⁶ Pb/ ²	²³⁸ U
INO.	(ppm)	(ppm)	(ppm)		ratios	1σ	ratios	1σ	ratios	1σ	ages	1σ	ages	1σ	ages	1σ
1	66.77	167.99	163.46	1.03	0.11128	0.00256	4.83101	0.06473	0.31481	0.00309	1820	12	1790	11	1764	15
2	30.18	55.96	72.46	0.77	0.10897	0.00269	4.7793	0.07766	0.31806	0.0034	1782	15	1781	14	1780	17
3	28.81	51.12	69.53	0.74	0.10898	0.0027	4.78842	0.07902	0.31864	0.00344	1782	15	1783	14	1783	17
4	13.15	24.95	31.6	0.79	0.1081	0.00314	4.71419	0.10626	0.31624	0.00409	1768	23	1770	19	1771	20
5	27.52	33.81	69.6	0.49	0.11039	0.00272	4.88001	0.07968	0.32058	0.00345	1806	15	1799	14	1793	17
6	44.74	109.41	107.98	1.01	0.10921	0.00258	4.58451	0.0672	0.30442	0.0031	1786	13	1746	12	1713	15
7	22.70	43.99	57.26	0.77	0.10935	0.0029	4.6737	0.08779	0.30997	0.00356	1726	59	1730	24	1733	19
8	23.67	47.83	62.24	0.77	0.10961	0.00301	4.58901	0.09242	0.30364	0.00363	1793	20	1747	17	1709	18
9	29.10	48.68	72.58	0.67	0.11215	0.00278	4.92364	0.08008	0.31838	0.0034	1797	49	1786	20	1777	18
10	19.24	43.06	46.21	0.93	0.10824	0.00297	4.66774	0.0941	0.31274	0.00373	1770	20	1761	17	1754	18
11	18.66	39.26	46.86	0.84	0.11355	0.00314	4.80976	0.09849	0.30718	0.00374	1784	64	1748	27	1717	20
12	26.05	47.61	67.84	0.70	0.10689	0.00277	4.48064	0.08058	0.30399	0.00338	1747	17	1727	15	1711	17
13	26.77	50.59	65.83	0.77	0.10948	0.00281	4.7334	0.08364	0.31354	0.0035	1791	17	1773	15	1758	17
14	32.93	56.38	83.79	0.67	0.11035	0.00271	4.74798	0.0759	0.31203	0.0033	1805	15	1776	13	1751	16
15	18.71	31.73	47.74	0.66	0.1115	0.00303	4.78594	0.09491	0.31127	0.00371	1778	58	1758	24	1741	19
16	38.78	88.51	95.75	0.92	0.11174	0.00272	4.71436	0.0733	0.30596	0.0032	1828	14	1770	13	1721	16
17	14.76	26.6	37.05	0.72	0.11134	0.00322	4.69946	0.10482	0.30609	0.00393	1768	66	1739	27	1715	21
18	19.55	36.57	50.06	0.73	0.11264	0.00307	4.72404	0.09452	0.30413	0.00365	1777	60	1737	25	1703	19
19	41.84	102.12	100.2	1.02	0.11071	0.00266	4.67126	0.07073	0.30597	0.00316	1811	14	1762	13	1721	16
20	28.45	71.2	71.83	0.99	0.11219	0.00311	4.79842	0.09946	0.31016	0.00383	1835	20	1785	17	1741	19
21	25.06	46.19	63.29	0.73	0.11331	0.00295	4.85338	0.08899	0.31061	0.00357	1853	17	1794	15	1744	18
22	34.00	59.17	82.06	0.72	0.11051	0.00269	4.86001	0.07718	0.31893	0.00339	1808	15	1795	13	1785	17
23	23.30	39.51	58.5	0.68	0.10776	0.00279	4.6317	0.08402	0.3117	0.00353	1762	17	1755	15	1749	17
24	41.64	89.28	100.09	0.89	0.10967	0.00261	4.67409	0.07008	0.30907	0.00319	1794	14	1763	13	1736	16
25	18.72	39.01	45.59	0.86	0.11021	0.00302	4.7157	0.09605	0.31032	0.00378	1803	20	1770	17	1742	19
27	25.69	48.46	64.57	0.75	0.10861	0.00279	4.55987	0.08192	0.30448	0.00344	1776	17	1742	15	1713	17
28	16.14	28.82	41.15	0.70	0.10948	0.00319	4.55897	0.10371	0.30201	0.00392	1791	23	1742	19	1701	19
29	18.50	32.77	47.56	0.69	0.10878	0.00299	4.57358	0.09424	0.30492	0.00373	1779	20	1744	17	1716	18
30	34.51	69.41	86.88	0.80	0.11152	0.00292	4.64793	0.08722	0.30228	0.00352	1824	18	1758	16	1703	17
31	29.92	71.54	73.97	0.97	0.11363	0.00309	4.77757	0.09666	0.30494	0.00374	1858	20	1781	17	1716	18
32	31.29	64.23	72.46	0.89	0.10967	0.00273	4.75268	0.08015	0.31431	0.00346	1794	16	1777	14	1762	17
33	28.79	52.1	67.99	0.77	0.10951	0.00275	4.7726	0.08214	0.31609	0.00352	1791	16	1780	14	1771	17
34	24.67	42.02	61.89	0.68	0.10916	0.00284	4.62872	0.08588	0.30756	0.00356	1785	18	1754	15	1729	18
35	37.71	77.88	96.3	0.81	0.11296	0.00289	4.70504	0.0842	0.30212	0.00345	1848	17	1768	15	1702	17
36	39.90	106.54	98.66	1.08	0.10862	0.00278	4.50009	0.08095	0.30052	0.00341	1776	17	1731	15	1694	17
37	26.13	53.38	60.44	0.88	0.12102	0.00306	5.08814	0.08941	0.30498	0.00349	1828	60	1756	25	1697	19
38	19.68	36.03	48.63	0.74	0.11083	0.00302	4.6695	0.09495	0.30563	0.00375	1813	20	1762	17	1719	19
39	21.28	36.41	52.95	0.69	0.1101	0.00292	4.64473	0.08983	0.30604	0.00364	1801	19	1757	16	1721	18
40	34.60	62.76	82.04	0.76	0.11838	0.00286	5.03398	0.08014	0.3085	0.00335	1795	53	1752	22	1715	18

Table 3. Zircon U-Pb isotope data obtained by LA-ICP-MS for k-feldspar granite porphyry (14LP15) from Baijiagou.

Note: Pb* represents radiogenic lead in the table.



Fig. 4. Cathodoluminescence (CL) images of zircons from the Baijiagou diabase (12LP26) [the numbers in the figure stand for age (Ma) of the zircons].



Fig. 5. Cathodoluminescence (CL) images of zircons from the Baijiagou granite porphyry (14LP15) (the numbers in the figure stand for age (Ma) of the zircons).



Fig. 6. U-Pb concordia diagram for zircons from the Baijiagou diabase (the blue circle in the figure is the concordia point and the red circle is inconcordia point).



Fig. 7. U-Pb concordia diagram for zircons from the Baijiagou granite porphyry.

1 .			-								
sample	14LP41	14LP43	14LP44	14LP46	14LP42	14LP45	14LP47	14LP52	14LP53	14LP54	
SiO_2	54.01	50.95	49.49	48.81	48.76	46.29	42.41	48.39	48.47	41.77	
TiO_2	1.47	1.41	1.54	1.47	2.12	2.26	2.04	1.93	1.88	1.61	
Al_2O_3	14.76	14.37	16.29	15.08	13.51	13.31	13.99	13.91	12.86	12.23	
$Fe_2O_3^T$	11.43	10.10	11.86	11.48	15.89	16.14	14.89	14.82	13.86	12.42	
MnO	0.14	0.15	0.14	0.15	0.32	0.19	0.11	0.22	0.23	0.28	
MgO	5.20	4.03	5.39	4.97	5.38	6.07	4.05	5.33	4.61	4.59	
CaO	3.51	5.56	5.05	5.58	5.37	7.24	7.92	6.64	7.48	13.34	
Na ₂ O	3.19	3.97	3.64	3.62	4.79	2.91	4.35	3.92	3.21	2.91	
K_2O	1.47	2.07	1.66	1.63	0.88	1.85	0.93	1.63	1.90	1.68	
P_2O_5	0.41	0.40	0.43	0.42	0.20	0.14	0.20	0.18	0.18	0.15	
LOI	4.40	6.73	4.86	6.62	2.80	3.40	8.97	2.71	5.61	9.02	
TOTAL	99.99	99.74	100.35	99.83	100.02	99.80	99.86	99.68	100.29	100.00	
M ^g #	51.5	48.2	51.4	50.2	44.1	46.7	38.8	45.6	43.7	46.3	
Li	50.1	27.1	40.3	36.6	28.1	34.1	37.1	30.2	38.0	31.1	
Be	1.86	1.13	1.79	1.88	2.23	0.73	1.89	1.37	2.65	1.83	
Sc	25.3	23.0	26.5	25.8	38.5	40.4	36.4	36.0	33.7	31.5	
V	154	131	156	159	375	542	355	360	324	343	
Cr	88.0	83.9	92.2	90.0	65.5	70.7	83.1	61.6	63.3	68.3	
Co	33.8	30.4	31.9	26.7	47.3	54.6	49.4	45.3	43.6	41.8	
Ni	42.7	38.9	43.9	42.7	38.0	49.3	49.0	36.5	33.9	41.0	

Table 4. Major-element (%) and trace-element (ppm) compositions of diabase from Baijiagou.

HT

59.5

117

17.1

2.32

78.8

260

25.4

88.2

9.55

LT

Cu

Zn

Ga

Ge

Rb

Sr

Y

Zr

Nb

23.0

217

19.7

1.50

33.0

363

42.3

218

9.36

59.8

131

17.7

1.08

35.9

266

38.3

213

8.96

39.6

139

22.7

1.55

32.6

515

42.7

229

9.98

14.5 $w(Eu)_{N} \{ (1/2)[w(Sm)_{N}+w(Gd)_{N}) \}, w(Eu)_{N}, w(Sm)_{N} and w(Gd)_{N} are the standard values of chondrite.$

58.1

179

21.7

2.03

35.3

207

28.2

134

81.2

130

18.8

2.03

98.3

254

21.3

92.2

10.1

66.5

206

23.1

1.30

36.8

238

25.7

122

13.2

61.3

130

19.9

1.80

79.0

206

25.6

110

12.5

53.9

123

19.4

1.69

86.4

214

29.4

145

13.5

15.0

135

22.7

1.35

30.3

341

41.0

222

9.57

		L	Т		НТ						
sample	14LP41	14LP43	14LP44	14LP46	14LP42	14LP45	14LP47	14LP52	14LP53	14LP54	
Cs	1.23	0.51	0.71	0.43	0.26	0.62	0.21	0.87	0.50	0.99	
Ba	917	1289	1064	1201	226	926	357	418	601	843	
La	42.5	39.3	43.3	40.9	13.0	8.93	14.1	11.4	13.8	11.8	
Ce	89.8	83.9	92.4	84.9	32.2	21.6	33.1	27.0	33.4	26.0	
Pr	11.1	10.3	11.3	10.5	4.40	3.09	4.32	3.83	4.44	3.61	
Nd	44.6	41.2	45.1	42.5	19.6	13.9	18.8	17.2	19.7	16.1	
Sm	8.80	8.06	8.88	8.39	4.94	3.66	4.49	4.41	4.90	4.07	
Eu	2.23	1.91	2.33	2.37	1.45	1.30	1.22	1.46	1.58	1.51	
Gd	8.07	7.37	8.07	7.72	5.01	3.79	4.59	4.56	5.01	4.31	
Tb	1.23	1.10	1.20	1.15	0.82	0.62	0.74	0.74	0.83	0.70	
Dy	7.53	6.68	7.26	7.06	5.11	3.92	4.58	4.64	5.16	4.32	
Но	1.51	1.35	1.51	1.44	1.02	0.79	0.92	0.94	1.04	0.87	
Er	4.32	3.94	4.32	4.14	2.84	2.19	2.59	2.61	2.97	2.40	
Tm	0.61	0.58	0.62	0.60	0.41	0.32	0.37	0.37	0.43	0.33	
Yb	4.06	3.84	4.12	4.03	2.76	2.04	2.38	2.45	2.81	2.14	
Lu	0.60	0.58	0.62	0.60	0.41	0.31	0.36	0.36	0.42	0.32	
Hf	5.22	5.01	5.50	5.25	3.40	2.32	3.04	2.78	3.59	2.21	
Та	0.49	0.47	0.52	0.49	0.88	0.61	0.82	0.75	0.87	0.59	
Pb	42.7	130	19.6	7.38	8.57	6.54	59.0	11.7	10.3	7.06	
Th	2.78	2.66	2.89	2.85	1.27	0.84	1.31	0.99	1.44	0.84	
U	0.47	0.43	0.49	0.55	0.46	0.35	0.69	0.39	0.57	0.34	
δEu	0.81	0.76	0.84	0.90	0.89	1.06	0.82	1.00	0.98	1.10	
∑REE	227.1	210.0	230.9	216.3	93.9	66.4	92.6	81.9	96.5	78.6	

Table 4. (Continued)



Fig. 8. (a) K₂O-SiO₂ and (b) AFM discriminant diagrams (modified after Irvine and Baragar 1971) for the Baijiagou diabase.



Fig. 9. Chondrite-normalized REE diagram for the Baijiagou diabase (chondrite data are from Boynton 1984).



Fig. 10. Multi-element patterns for the Baijiagou diabase (MORB data are from Pearce 1982; chondrite data are from Boynton 1984).

1 J											
Sample	⁸⁷ Rb/ ⁸⁶ Sr	87 Sr/ 86 Sr ± 2 σ	147Sm/144Nd	143 Nd/ 144 Nd ± 2 σ	$\epsilon_{\rm Nd}(t)$	$({}^{87}{ m Sr}/{}^{86}{ m Sr})_i$	t _{DM1} (Ga)	t _{DM2} (Ga)			
14LP41	0.263	0.714994 ± 11	0.1193	0.511412 ± 4	-4.04	0.705094898	2.79	2.63			
14LP42	0.494	0.720542 ± 14	0.1524	0.512271 ± 5	4.24	0.701962536	2.18	2.01			
14LP43	0.391	0.717897 ± 15	0.1182	0.511413 ± 5	-3.76	0.703196637	2.76	2.61			
14LP44	0.183	0.713436 ± 12	0.1190	0.511392 ± 6	-4.37	0.706543927	2.81	2.65			
14LP45	1.122	0.731517 ± 10	0.1592	0.512311 ± 5	3.27	0.689306833	2.34	2.06			
14LP46	0.257	0.718045 ± 10	0.1193	0.511417 ± 8	-3.96	0.70836653	2.78	2.62			
14LP47	0.448	0.724754 ± 19	0.1444	0.512201 ± 10	4.94	0.707900517	2.08	1.97			
14LP52	1.112	0.732503 ± 26	0.1550	0.512316 ± 7	4.45	0.690671501	2.16	1.99			
14LP53	1.171	0.734906 ± 14	0.1504	0.512284 ± 7	5.03	0.690856575	2.08	1.96			
14LP54	0.879	0.728720 ± 12	0.1528	0.512304 ± 5	4.78	0.695672588	2.11	1.97			

Table 5. Sm-Nd isotopic composition of Baijiagou diabase.

Note: $\varepsilon_{Nd}(t) = [({}^{143}Nd/{}^{144}Nd)_{sample}/({}^{143}Nd/{}^{144}Nd)_{CHUR}-1] \times 10000.$

	5			11	/ 1		1 0	1 1	5 5	3.0
sample	14LP17	14LP18	14LP19	14LP20	14LP21	14LP22	14LP23	14LP24	14LP25	14LP26
SiO_2	73.66	78.24	73.80	68.41	71.26	68.07	69.18	67.61	62.02	70.16
TiO_2	0.56	0.55	0.59	0.55	0.52	0.59	0.67	0.56	0.71	0.60
Al_2O_3	7.59	6.96	8.86	13.48	7.41	12.03	12.54	11.37	11.47	11.87
Fe ₂ O ₃ ^T	4.72	6.32	4.29	5.59	3.76	5.94	4.98	6.31	8.01	6.13
MnO	0.11	0.11	0.09	0.09	0.09	0.10	0.09	0.15	0.10	0.13
MgO	1.01	1.16	1.11	1.51	0.86	1.58	1.20	1.75	0.95	1.61
CaO	4.10	1.53	2.55	0.33	6.11	1.21	0.48	1.60	4.34	1.05
Na ₂ O	2.67	2.06	1.72	2.38	1.88	2.05	2.15	2.72	2.56	3.82
K ₂ O	0.56	0.74	3.21	5.29	1.90	5.42	5.65	2.80	4.52	1.86
P_2O_5	0.11	0.10	0.11	0.11	0.10	0.11	0.13	0.10	0.15	0.11
LOI	4.67	2.66	3.47	2.04	6.18	2.52	3.06	4.87	4.71	2.54
TOTAL	99.76	100.43	99.80	99.78	100.07	99.62	100.13	99.84	99.54	99.88
A/CNK	0.61	1.00	0.81	1.32	0.46	1.05	1.19	1.09	0.67	1.16
Mg#	33.3	30.0	37.6	38.6	34.8	38.3	36.0	39.3	21.7	38.0
Li	13.0	20.7	13.4	18.6	19.0	17.7	18.3	21.3	11.5	20.7
Be	1.87	1.95	1.57	2.64	1.76	2.60	1.80	2.03	1.64	1.85
Sc	9.88	10.6	8.45	10.1	11.0	10.4	11.2	9.62	11.0	10.7
V	8.06	9.50	7.58	10.3	9.30	13.1	10.7	6.60	10.3	9.39
Cr	7.89	7.35	6.78	20.4	6.68	8.15	7.50	16.3	7.99	7.07
Co	16.1	15.4	12.8	11.7	11.9	12.3	11.7	14.6	11.7	12.2
Ni	5.08	6.82	3.76	16.9	3.99	5.07	3.42	10.2	3.82	3.77
Cu	8.82	5.66	2.45	5.62	2.77	3.81	2.72	2.50	2.97	2.52
Zn	121	133	126	156	98.8	127	144	215	112	196
Ga	20.0	19.6	17.2	25.8	16.7	24.4	23.2	26.8	17.9	25.1
Ge	1.66	1.85	1.24	1.39	1.26	1.13	1.16	1.48	1.32	1.24
Rb	37.2	38.6	74.0	148	55.9	141	129	91.9	99.2	58.0
Sr	51.9	44.2	49.3	64.9	63.6	48.8	34.2	45.7	57.3	45.1
Y	88.2	80.7	79.9	81.8	88.1	79.6	84.3	78.8	122	78.1
Zr	535	520	580	388	525	611	678	558	763	616
Nb	25.2	24.5	25.9	27.1	23.2	28.2	31.1	25.6	33.8	27.3
Cs	0.38	0.58	0.29	0.69	0.34	0.41	0.33	0.57	0.32	0.34
Ba	114	132	1044	2738	857	1631	1061	510	1535	672
La	211	251	82.9	101	91.8	24.7	63.2	120	133	91.3
Ce	401	482	171	207	189	59.2	132	242	276	193
Pr	49.6	59.3	21.1	25.0	23.3	7.88	16.4	30.4	34.3	23.3
Nd	193	229	82.2	95.6	88.2	34.2	65.2	113	124	87.7
Sm	31.3	36.3	17.4	17.6	16.7	9.19	13.5	21.3	22.9	15.9
Eu	4.21	5.20	2.76	2.50	2.27	1.45	1.90	2.83	3.42	2.05
Gd	22.9	27.2	15.6	14.9	14.7	9.98	11.9	17.0	21.8	13.0
Tb	2.95	3.44	2.36	2.19	2.30	1.95	1.99	2.46	3.42	2.05
Dy	16.6	17.8	14.1	13.7	14.8	13.4	13.5	14.5	21.5	13.0
Но	3.18	3.02	2.90	2.89	3.00	2.95	3.02	2.88	4.42	2.77
Er	9.13	8.29	8.48	8.58	8.73	8.70	9.43	8.33	12.3	8.43
Tm	1.31	1.19	1.27	1.27	1.31	1.31	1.47	1.23	1.75	1.28
Yb	8.72	8.05	8.60	8.58	8.58	8.83	10.1	8.19	11.5	8.80
Lu	1.28	1.18	1.28	1.28	1.24	1.31	1.50	1.20	1.68	1.33
Hf	13.1	12.8	14.4	10.0	12.6	14.8	16.6	13.4	18.5	15.0
Та	1.34	1.31	1.48	1.43	1.24	1.51	1.70	1.36	1.82	1.43
Pb	5.71	8.97	4.06	5.67	5.15	5.81	4.52	6.07	5.10	3.50
Th	18.5	18.7	19.9	18.8	19.8	17.7	21.8	19.2	23.3	19.3
U	4.83	3.14	4.25	5.60	3.17	3.16	3.81	3.82	5.84	4.61
δΕυ	0.48	0.51	0.51	0.47	0.44	0.46	0.46	0.45	0.47	0.44

Table 6. Major-element (%) and trace-element (ppm) compositions of k-feldspar granite porphyry from Baijiagou.

 $Note: A/CNK = (Al_2O_3/101.96)/[(CaO/56.08) + (Na_2O/61.98) + (K_2O/94.20)].$



Fig. 11. $\varepsilon_{Nd}(t)$ -SiO₂ (a) and $\varepsilon_{Nd}(t)$ -La/Nb (b) diagram for Baijiagou diabase (after Wei and Xu 2013).



Fig. 12. $({}^{87}Sr/{}^{86}Sr)_i$ -t and ε_{Nd} -t diagram for Baijiagou diabase (after Han and Ma 2003).

Sr isotopes, the initial Sr, Nb isotopes of the HT type diabase basically follow the mantle evolution line; the evolution of initial Sr, Nb isotopes for LT type diabase is between the mantle and the crust, which is the result of crustal contamination.

6.2.2 Crystal Fractionation

Fractional crystallization of different minerals from parental magmas of the same composition will produce rocks with contrasting geochemistry and mineralogy. The crystallization of different minerals from parental magmas of the same composition can result from differences in pressure-temperature conditions. Figure 13a show that MgO is rough correlated with $Fe_2O_3^T$ for the LT type diabase, which is indicative of olivine fractionation. But the correlations are insignificant for the HT type diabase. No correlation or negative correlation between CaO and MgO for both LT and HT type diabase (Fig. 13b) suggests non fractionation of calcic pyroxene (e.g., diopside). The LT type shows positive correlations between Sr and Al₂O₃ (Fig. 13c), reflecting the crystallization of calcic plagioclase during the evolution of the LT diabase magma. The LT diabase also has significant negative Eu and Sr anomalies, consistent with plagioclase crystallization; no correlation between Sr and Al₂O₃ for the HT type indicates that fractionation of calcic plagioclase did not occur, consistent with the lack of Eu anomalies. Figure 13d shows that the LT type diabase mainly formed by crystal fractionation.

6.2.3 Source of the Parental Magma

In a mid-ocean ridge basalt (MORB)-normalized multielement (spider) diagram (Fig. 9a), the diabase samples are enriched in K, Rb, Ba, Th, and Ce relative to MORB, which is similar to continental basalts produced in an extensional setting (Pearce 1982; Dong et al. 1998; Kalsbeek et al. 2001). The LT type is characterized by depletions of Nb, Ta, and Ti relative to the HT type, and the Th/Nb ratios of the LT and HT diabases are 0.30 - 0.29 and 0.08 - 0.1, respectively,



Fig. 13. Harker diagrams showing fractional crystallization trends for the Baijiagou diabase.

and their Th/Ta ratios are 5.83 - 5.58 and 1.33 - 1.65, respectively. These ratios are similar to those of continental rift basalt (LT) and mantle plume basalt (HT), respectively (Wang et al. 2001; Sun et al. 2003). The chondrite-normalized REE patterns of the HT type are distinct from MORB and similar to ocean island basalt (OIB). The main difference between the HT type diabase and OIB is that the former is LILE enriched and has some continental characteristics. The HT type diabase is geochemically similar to HT basalts of the Emei Mounto igneous province (Xu and Chung 2001; Xiao et al. 2003; Wei and Xu 2013). These characteristics suggest that the HT type diabase may come from convective mantle. And the LT type diabase maybe derived from convective mantle and crust.

The Sr and Nd isotopic systems are commonly used as magma source region tracers. On a $({}^{87}\text{Sr}/{}^{86}\text{Sr})_{i^-}$ t and ε_{Nd} -t diagram (Fig. 12) the HT type diabase plot on the mantle evolution line, and the T_{DM2} of the HT type (1.96 - 2.06 Ga) is roughly near crystallizing age. But the LT type diabase plot between the mantle and the crust, which suggest that the rich end-member came from the ancient crust material. The T_{DM2} of the LT type (2.61 - 2.65 Ga) can also confirm this.

The flat middle-heavy REE patterns of the LT diabase

indicate derivation from a garnet-free source. The minimum pressure of garnet stability is 1.6 - 1.7 GPa (Deng et al. 2009), meaning that the source of the LT diabase magma experienced a pressure of < 1.6 GPa (i.e., < 56 km), within the lithospheric mantle. In contrast, the HT diabase did not experience plagioclase crystallization (no negative Eu-Sr anomalies), and its REE pattern (Figs. 9 and 10) indicates that its source was garnet-bearing. The garnet stability zone is generally at a depth of > 80 km, meaning that the source of the HT diabase magma was at depths of > 80 km (Deng et al. 2009). Hence, the HT diabase magma may have been derived from a mantle plume or the asthenosphere.

Petrographic and geochemical data show that the granite porphyry located near the diabase is an A-type granite. The formation of A-type granite generally requires low pressures and water contents, and high temperatures. Partial melting experiments have demonstrated that A-type granites can be produced from the shallow continental crust at P < 0.8 GPa and T > 800°C (Zhang et al. 2010). It is thought that zircon saturation temperatures represent the near-liquidus temperatures of granitic rocks (Watson and Harrison 1983). Miller et al. (2003) calculated a series of zircon saturation temperatures and showed that there are perhaps two types of granites: "Hot granites" have $T_{Zr} > 800^{\circ}$ C, whereas "cold granites" have $T_{Zr} < 800^{\circ}$ C. The difference in the temperature estimates is attributed to the amount of inherited zircon, with the "hot granites" containing few inherited zircons. We calculated T_{Zr} for the Baijiagou granite porphyry using the following equation:

$$T_{\rm Zr}(^{\circ}{\rm C}) =$$
12900/[2.95 + 0.85M + ln(496000/Zr)] - 273.15
(1)

In this equation, M is the atomic fraction normalizing parameter, where Si + Al + Fe + Mg + Ca + Na + K + P = 1 and M = $(2Ca + K + Na)/(Si \times Al)$. This equation yielded crystallization temperatures for the Baijiagou granite porphyry of ca. 880°C. There are also no inherited zircons in the granite porphyry. Zircon saturation temperatures typically represent the lowest temperature of magmatic crystallization, and so the formation temperature of the Baigaogou granite was > 880°C. According to the geothermal gradient and at 0.8 GPa pressure, the temperature would be $< 700^{\circ}$ C; however, with input of mantle heat the temperature at 30 km depth can reach $\sim 900^{\circ}$ C. Therefore, a thermal anomaly was involved in the formation of granite porphyry, perhaps provided by upwelling mantle.

The above discussion leads to the conclusion that the Paleoproterozoic diabase and granite porphyry formed in a continental rift environment, possibly an active rift caused by mantle plume activity. This conclusion is supported by Th/Hf-Ta/Hf and 2Nb-Zr/4-Y tectonic discrimination diagrams (Fig. 14).

Figure 15 shows a schematic depiction of the magmatic evolution of the Baijiagou region. The ascending mantle plume caused regional crustal uplift, extension, and thinning. At the same time, the high-temperature magmas from the mantle plume caused partial melting of the lower crust and formation of A-type granitic magma. The early mantle plume material mixed with crustal material during upwelling and the LT diabase magma formed. The LT diabase and A-type granite were formed at the same time. Later magmas



Fig. 14. Th/Hf-Ta/Hf and 2Nb-Zr/4-Y tectonic setting discrimination diagrams for basalts [(a) modified after Wang et al. 2001; (b) after Meschede 1986]. I: N-MORB; II: convergent plate margin basalt (III: oceanic arc basalt; II2: continental island arc basalt); III: T-MORB, E-MORB; IV: continental intraplate basalt (IV1: continental rift valley or continental margin rift basalt; IV2: continental rift valley alkali basalt; IV3: continental extensional zone and incipient rift valley basalt); V: mantle plume basalt; A: intraplate alkali basalt; B: mantle plume oceanic ridge basalt; C: intraplate tholeiitic basalt; D: normal oceanic ridge basalt.



Fig. 15. Schematic diagram of the magma sources of the Baijiagou rocks (modified after Wilson 1989).

were derived directly from the mantle plume and formed the HT diabase. Subsequently, the crust and lithosphere were further extended and formed a fully developed continental rift. Later tectonism, sedimentation, and erosion mean that the sedimentary strata of this Paleoproterozoic are now rarely exposed; however, the bimodal magmatism of the Baijiagou region provides direct evidence for the southward extension of the Helan aulacogen.

6.3 Tectonic Implications

Previous studies have examined the extensional tectonic events of the NCC and the formation of its aulacogens in the late Paleoproterozoic (Li et al. 2000; Zhai and Peng 2007; Zhao et al. 2007; Zhai et al. 2014; Gao et al. 2013). These extensional events produced a rift system in the NCC. E-W-trending rifts evolved into the Qinling-Qilian trough and the NNE- to NE-trending rifts (Helan, Xiong'er, and Yanliao rifts) evolved into aulacogens (Che et al. 2011). Previous studies on the Xiong'er and Yanliao aulacogens have suggested that extension was associated with breakaway of the NCC from the Columbia Supercontinent (Yang et al. 2005; Zhao et al. 2007; Cui et al. 2010), and may have been related to a mantle plume (Peng et al. 2005; Xu et al. 2007; Yang et al. 2019). Zhai (2004) and Zhai et al. (2014) provided evidence for mantle plume involvement, including: (1) large-scale mantle upwelling and uplift of the lower crust to the surface, accompanied by strong migmatization and meso-ductile deformation that exposed granulite-amphibolite-facies metamorphic rocks; (2) medium-pressure granulite- and amphibolite-facies retrograde metamorphism of high-pressure granulite dikes in the NCC at 1820 - 1800 and 1790 - 1760 Ma, with isothermal or decreasing-T decompression P-T-t paths; (3) basement uplift following the rift events; and (4) rapid enrichment of ancient depleted mantle in the period from 2000 to 1900 - 1760 Ma. It can thus be inferred that mantle upwelling caused uplift of the NCC, which resulted in lithospheric thinning, partial melting, and the development of a rift system. Other studies have reached similar conclusions, and consider that the Xiong'er volcanic rocks and mafic dikes, anorthosites, alkaline granites, and rapakivi granites with ages of 1750 - 1680 Ma represent the mantle plume or break-up of the Columbia supercontinent (Peng et al. 2005, 2007; Hou et al. 2008; Hou 2012; Zhang et al. 2012). Xu et al. (2007) and Yang et al. (2019) argued that the Xiaoliangling volcanic rocks in the Luliang area are similar to the Xiong'er volcanic rocks in terms of lithology, geochemistry, and age, and speculated that their formation was also related to mantle plume activity.

Mantle plumes can produce significant, short-term environmental effects due to large volcanic eruptions, and longerterm effects on global tectonism, including: (1) regional crustal uplift and subsequent continent rifting; (2) the evolution and extinction of biota; and (3) regional and global changes in climate; however, such effects from an ancient Proterozoic mantle plume are difficult to identify (Xu and Qiu 2010). Therefore, identification of ancient mantle plumes relies on petrological and geochemical studies of igneous rocks. Our study area is located at the southwestern margin of the NCC, and connects the Qinling-Qilian and Helan troughs. The two types of diabase and granite porphyry present in this region formed at 1.8 Ga, and have geochemical features consistent with formation in a continental rift environment associated with a mantle plume. These rocks were produced during a period of rifting in the NCC that was linked to the break-up of the Columbia supercontinent.

7. CONCLUSIONS

- The crystallization age of the Baijiagou diabase is 1804 ± 21 Ma (MSWD = 0.19), which is therefore Paleoproterozoic in age.
- (2) The diabase can be divided into LT and HT types.
- (3) The diabase is associated with a coeval granite porphyry, and these collectively represent a Paleoproterozoic bimodal magmatic assemblage related to continental rifting and a mantle plume.
- (4) The major 1.8 Ga rifting event (Helan aulacogen) at the southwestern margin of the NCC, which may have been caused by a mantle plume, was associated with the break-up of the Columbia supercontinent.

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