



# Magma generation and sulfide saturation of Permian mafic-ultramafic intrusions from the western part of the Northern Tianshan in NW China: implications for Ni-Cu mineralization

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## Abstract

Compared to Permian magmatic Ni-Cu sulfide deposits in the eastern part of the Northern Tianshan, such as Huangshan and Huangshandong, those in the western part (e.g., Baixintan and Lubei) are poorly documented. Zircon ID-TIMS U-Pb age data ( $279.7 \pm 0.2$  Ma) of ilmenite and LA-ICPMS U-Pb age data ( $277.8 \pm 3.4$  Ma) of a hornblende gabbro suggest the Lubei intrusion was emplaced in the Early Permian. The Baixintan and Lubei mafic-ultramafic intrusive rocks are characterized by enrichments of large ion lithophile elements and Th and U relative to the high field strength elements, as well as depletion of Nb and Ta relative to mid-ocean ridge basalt. The  $(^{87}\text{Sr}/^{86}\text{Sr})_t$  (0.70354–0.70713),  $\epsilon\text{Nd}_{(t)}$  (2.80 to +7.05), and  $\epsilon\text{Hf}_{(t)}$  (2.50–18.4) values for the intrusions indicate that they have assimilated Tianshan basement rocks, whereas the Se/S ratios ( $181 \times 10^{-6}$ – $418 \times 10^{-6}$ ) and  $\delta^{34}\text{S}_{\text{CDT}}$  values ( $-0.13\%$  to  $0.90\%$ ) of the sulfides suggest that crustal sulfur has not been incorporated into the magma. The decrease in Pd/Zr with increasing  $(\text{Th}/\text{Nb})_{\text{PM}}$  indicates that sulfide saturation accompanied crustal contamination. The olivine compositions and whole-rock chalcophile elements of the Baixintan and Lubei intrusions imply that the parental magma of the intrusive rocks in the west contains higher chalcophile elements than those in the east, which suggests that the mafic-ultramafic intrusions in the western part of the Northern Tianshan should be regarded as favorable Ni-Cu sulfide exploration targets.

**Keywords** Magmatic Ni-Cu sulfide deposit · Baixintan mafic-ultramafic intrusion · Lubei mafic-ultramafic intrusion · Western Northern Tianshan · Mantle source · Sulfide saturation

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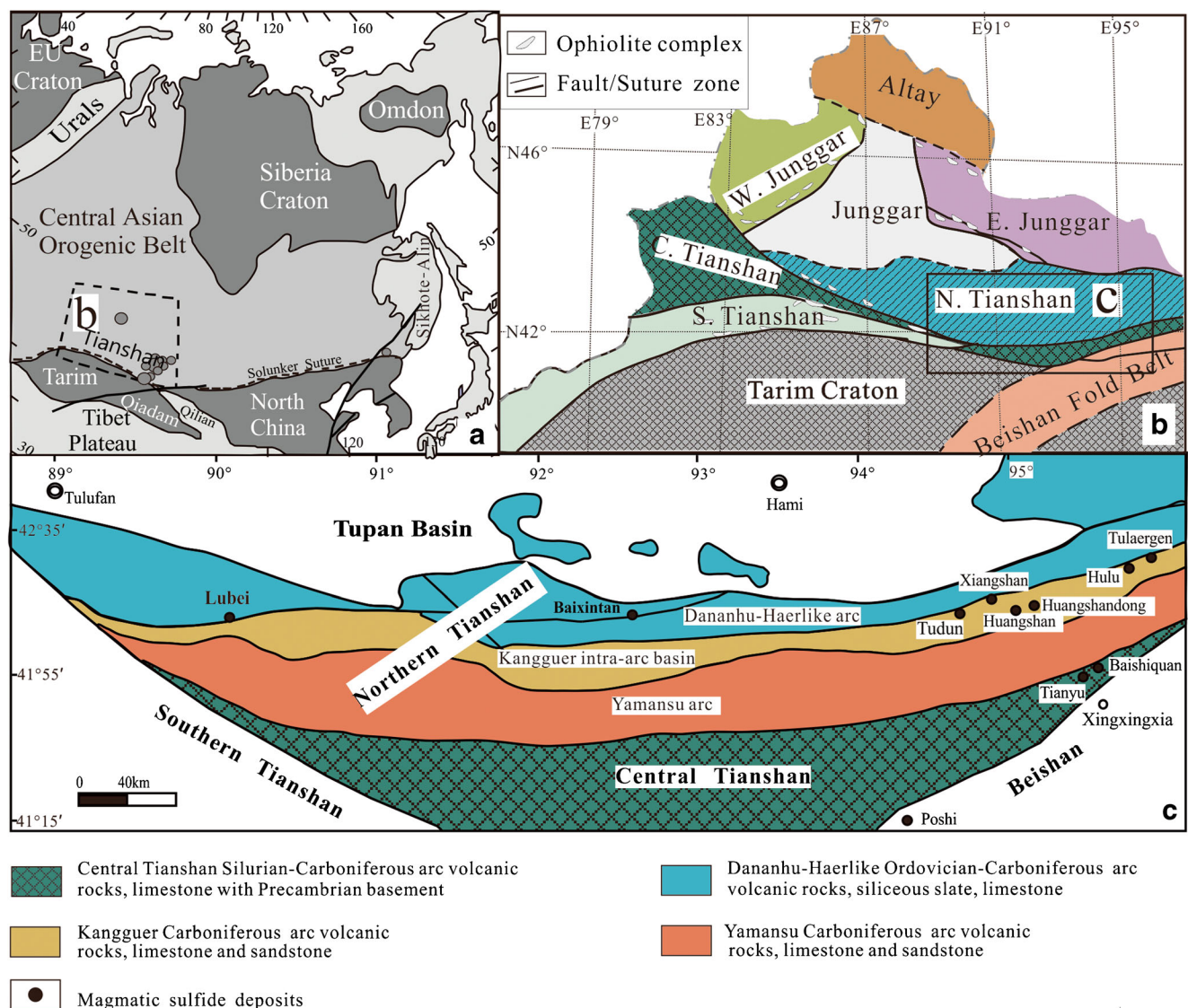
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## Introduction

In the past decades, a growing number of magmatic Ni-Cu sulfide deposits have been found in plate margin settings, such as Xiarihamu (China), Santa Rita (Brazil), Aguablanca (Spain), Quetico (Canada), and Duck Island (USA), indicating significant exploration potential in this tectonic environment (Tornos et al. 2001; Pettigrew and Hattori 2006; Thakurta et al. 2008; Maier et al. 2008; Ripley 2009; Barnes et al. 2011; Xie et al. 2014; Li et al. 2015; Song et al. 2016). Magmatic Ni-Cu sulfide deposits hosted by Permian mafic-ultramafic intrusions in the Northern Tianshan along the southern margin of the Central Asian Orogenic Belt (CAOB) contain roughly 1 million metric tons (Mt) of Ni metal reserves (Fig. 1; Qin et al. 2003; Liu et al. 2005; Su et al. 2013). Magmatic Ni-Cu sulfide deposits in the

eastern part of the Northern Tianshan include Huangshan, Huangshandong, Tulaergen, Huangshannan, Xiangshan, Tudun, and Hulu (Zhang et al. 2011; Sun et al. 2013; Tang et al. 2014; Deng et al. 2014, 2017; Wang et al. 2015; Mao et al. 2015, 2017a). Most of the intrusions that host magmatic Ni-Cu sulfide deposits in the eastern part of the Northern Tianshan formed around 280 Ma (Zhou et al. 2004; Mao et al. 2008; San et al. 2010; Qin et al. 2011; Song et al. 2013; Zhao et al. 2016), due to the interaction between metasomatized lithospheric mantle and ascending asthenospheric mantle, as a result of slab break-off or an upwelling mantle plume (Zhou et al. 2004; Qin et al. 2011; Su et al. 2011; Gao and Zhou 2012; Gao et al. 2013; Song et al. 2013; Deng et al. 2015). Although crustal contamination is generally accepted to have been critical in triggering sulfide segregation to form the deposits, fractional



**Fig. 1** **a** Schematic geological map of the Central Asian Orogenic Belt (modified after Jahn et al. 2000; Xiao et al. 2004). **b** Tectonic blocks of northern Xinjiang (after BGMX, 1993; Song and Li 2009; Song et al.

2013). **c** Simplified geological map of Northern Tianshan (after BGMX, 1993; Xiao et al. 2004; Song et al. 2011; Wang et al. 2015)

crystallization and magma mixing are also thought to have played some role in triggering S saturation during magma evolution in some deposits (Tang et al. 2012; Gao et al. 2013; Sun et al. 2013; Mao et al. 2015; Zhao et al. 2015).

Recent mapping, drilling, and geophysical survey in the western part of the Northern Tianshan have led to the discovery of the Baixintan and Lubei Ni-Cu sulfide occurrences (Li et al. 2014; Yang et al. 2017). The Baixintan intrusion is thought to have formed in a Permian post-orogenic setting, similar to intrusions in the eastern part of the Northern Tianshan (Wang et al. 2015, 2016a, b; Feng et al. 2017). The Permian Lubei mafic-ultramafic intrusions were formed from tholeiitic basaltic magmas derived from decompression partial melting of the metasomatized mantle in a post-collision extensional tectonic setting (Chen et al. 2018). However, it is unclear whether or not the mantle source and the sulfide saturation history of the Baixintan and Lubei intrusions are similar to those in the east. Here, we present new petrological, mineralogical, and geochemical data to constrain magma evolution and sulfide saturation processes of the intrusions in the western part of the Northern Tianshan.

## Geological background

The CAOB is one of the largest accretionary orogens worldwide and comprises a complex collage of continental fragments, island arc assemblages, remnants of oceanic crust, and continental margins, set between the Siberian Craton to the north and the Tarim-North China Craton to the south (Fig. 1a; Sengör et al. 1993; Windley et al. 2007). The southern part of the CAOB in the northern Xinjiang consists, from north to south, of the Chinese Altai, the Junggar and Tianshan terranes, and the Beishan Fold Belt (Fig. 1b). The Tianshan terrane can be further divided into the Northern, Central, and Southern Tianshan (Fig. 1c; BGMX 1993; Xiao et al. 2004).

The Northern Tianshan is separated from the Central Tianshan by the Aqikkuduk fault and comprises, from north to south, the Bogda intra-arc basin, the Dananhu-Harlik island arc, and the Kanggur-Yamansu intra-arc basin (Ma et al. 1993; Jahn et al. 2000; Qin et al. 2002; Xiao et al. 2004, 2008; Fig. 1c). The Bogda intra-arc basin consists of Carboniferous bimodal volcanic rocks, marine carbonates, and epicontinental detrital rocks, as well as Permian bimodal volcanic rocks, conglomerates, and sandstones. The Late Carboniferous pillow basalts and high-alumina basalts are believed to have formed in a back-arc extensional environment (Xie et al. 2016a, 2016b), whereas the Permian bimodal volcanism in the southern Bogda zone may occur in a post-collisional tectonic setting, triggered by slab break-off (Shu et al. 2010; Chen et al. 2011). The Kanggur-Yamansu intra-arc basin contains the Shaerdelan Mesoproterozoic metamorphic core complex and Early Carboniferous submarine lavas, turbidites, pyroclastic rocks,

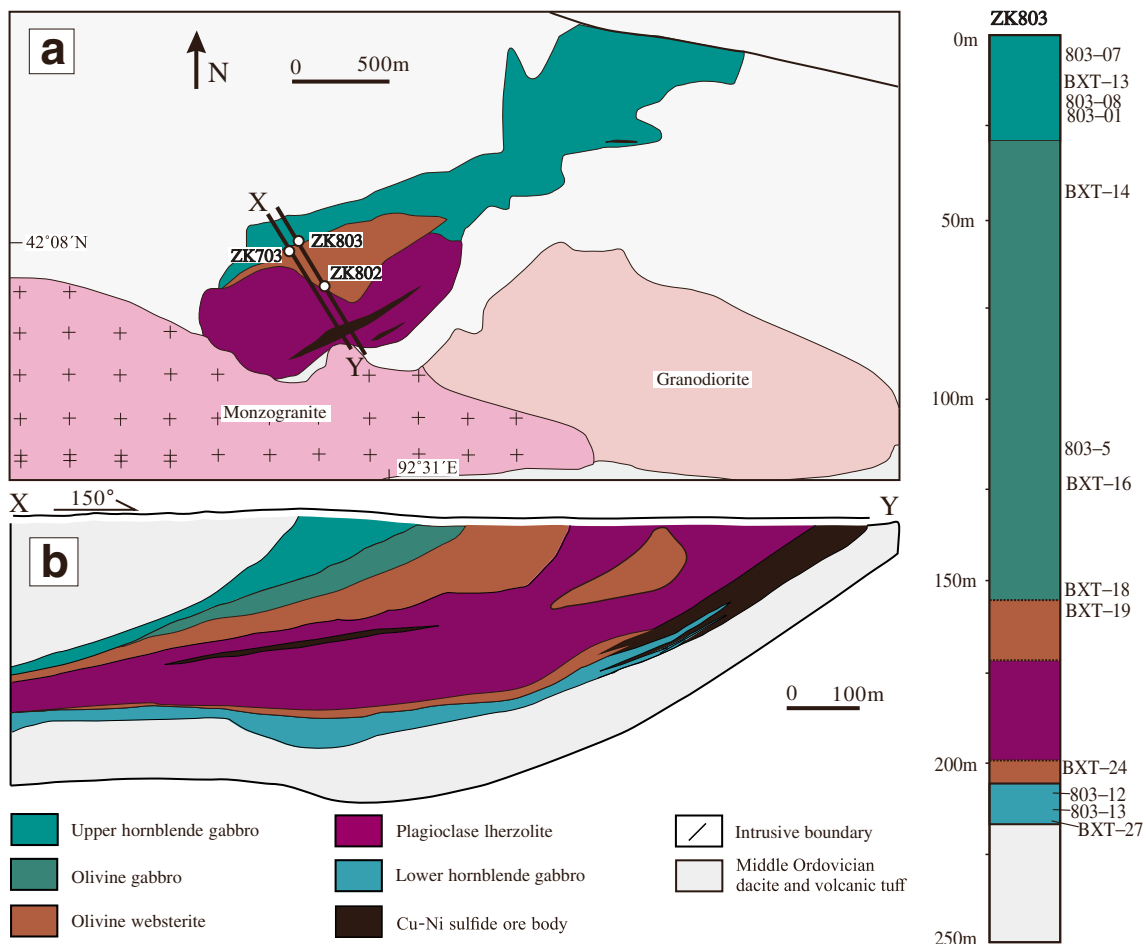
basalt flows, andesitic tuffs, and tuffaceous sandstones (BGMX 1993; Xiao et al. 2004; Su et al. 2008; Zhou et al. 2010; Zhang et al. 2014). Mylonitized granites from the Shaerdelan metamorphic core complex have a zircon LA-ICP-MS U-Pb age of  $921.7 \pm 8.1$  Ma (Su et al. 2008). Native copper-bearing basalts (307–318 Ma) within the Kanggur-Yamansu intra-arc basin have been interpreted to be derived from mafic magmatism generated through lithospheric delamination (Yuan et al. 2007; Zhang et al. 2013). More than 30 Early Permian mafic-ultramafic intrusions have been identified along the east-trending Kanggurtag fault, eight of which are known to host magmatic Ni-Cu sulfide deposits (Fig. 1c; Mao et al. 2008; San et al. 2010; Han et al. 2010; Song et al. 2013; Wang et al. 2015).

The Dananhu-Harlik arc is composed of Ordovician–Carboniferous tholeiitic basalt, calc-alkaline andesite, and pyroclastic rocks (Qin et al. 2002; Li et al. 2006a). Zircon U-Pb dating of volcanic rocks from the Kalatage area in the Dananhu-Harlik arc indicates that these rocks contain Proterozoic-inherited zircons (Long et al. 2017). There are various types of ore deposits in the Kalatage area, including the Silurian Honghai volcanogenic massive sulfide deposit, the Middle Devonian Xierqu skarn deposit, and the Middle Devonian Yudai porphyry deposit (Mao et al. 2010, 2017b; Deng et al. 2016). Adakites in the Early Carboniferous Tuwu-Yandong and Fuxing porphyry Cu deposits in the Dananhu-Harlik arc may have formed by partial melting of subducted oceanic crust (Zhang et al. 2006; Wang et al. 2016a). Several Early Permian mafic-ultramafic intrusions occur along the Kangguer and Dacatan faults, but Ni–Cu mineralization has only been recognized in the Baixintan and Lubei intrusions (Fig. 1c; Li et al. 2006b; Ren et al. 2013; Wang et al. 2015, 2016a, b; Feng et al. 2017; Yang et al. 2017; Chen et al. 2018).

## Petrography and mineralization of the Baixintan and Lubei intrusions

### Baixintan intrusion

The Baixintan intrusion is  $\sim 3.1$  km long, 0.8 km wide, and 0–0.3 km thick, with an areal extent of about 2.1 km<sup>2</sup> (Fig. 2). It comprises an upper hornblende gabbro, a lower hornblende gabbro, and a middle mafic-ultramafic unit with olivine gabbro, lherzolite, and olivine websterite, all emplaced in Middle Ordovician dacite and volcanic tuff of the Qieganbulake Formation (Li et al. 2014; Wang et al. 2015; Feng et al. 2017; Zhao et al. 2018). Zhao et al. (2018) reported a LA-ICP-MS zircon U-Pb Concordia age of  $287 \pm 3$  Ma for the olivine gabbro, which is consistent with a LA-ICP-MS zircon Concordia age of  $285 \text{ Ma} \pm 1$  Ma reported by Feng et al. (2017).



**Fig. 2** a, b Simplified geological map and cross sections of the Baixintan intrusion (after Li et al. 2014; Wang et al. 2015)

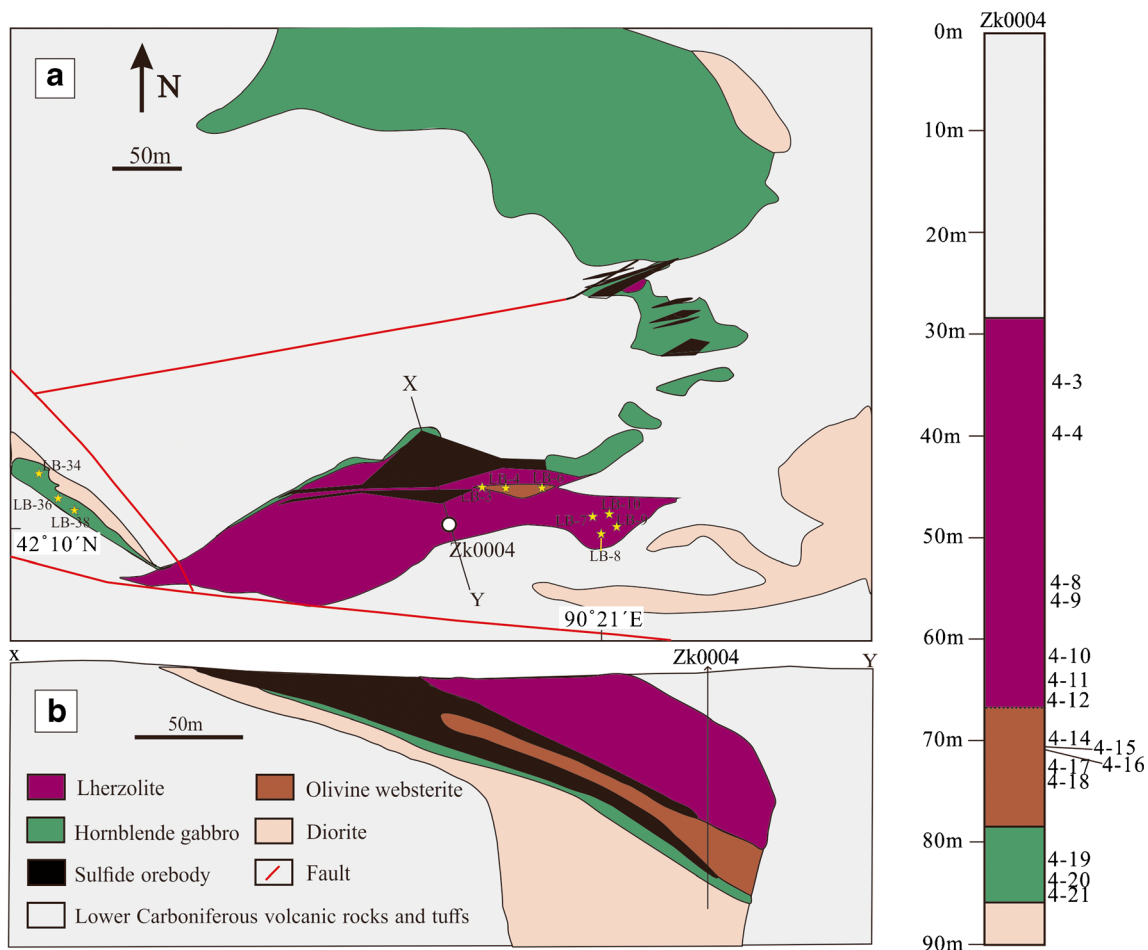
The upper hornblende gabbro is cut by the olivine gabbro of the middle unit and in places is enclosed in olivine gabbro. The contacts between the olivine gabbro, lherzolite, and olivine websterite are generally gradational. The sharp contacts between the lherzolite of the middle unit and the lower hornblende gabbro indicate they formed from different magmatic pulses. The sulfides are mainly hosted in the lherzolite and lower hornblende gabbro with average grades of 0.46 wt% Ni and 1.02 wt% Cu (Li et al. 2014). The mineralization comprises pyrrhotite, pentlandite, chalcopyrite, and violarite.

The upper hornblende gabbro consists of coarse-grained plagioclase (60–65%), clinopyroxene (15–20%), hornblende (10–15%), and minor biotite (0–3%). In the middle unit, the olivine gabbro is composed of 15–20% olivine, 40–50% plagioclase, 5–10% orthopyroxene, 15–20% clinopyroxene, 5–10% hornblende, and 2% sulfide. Some plagioclase crystals are enclosed in olivine and clinopyroxene. The olivine websterite contains 20–35% olivine, 5–10% orthopyroxene, 10–35% clinopyroxene, 5–10% plagioclase, 5–8% hornblende, and 2–20% sulfide. The lherzolite contains 40–55% olivine, 3–5% orthopyroxene, 5–10% clinopyroxene, 5–10%

plagioclase, 5–8% hornblende, and 2% sulfide. The olivine crystals are sub-rounded and enclosed in large orthopyroxene, clinopyroxene, plagioclase, and hornblende. The sulfides are commonly interstitial, but small, rounded sulfide inclusions also occur in some olivine crystals. The lower hornblende gabbro contains fine-grained plagioclase (60–70%), clinopyroxene (15–20%), hornblende (10–20%), and minor biotite (0–2%).

### Lubei intrusion

The Lubei intrusion has an areal extent of about 0.36 km<sup>2</sup> (Fig. 3) and intrudes the Late Carboniferous andesite, dacite, and sandstone of the Xiaorequanzi Formation as well as the Late Carboniferous sandstones and limestones of the Gandun Formation. The intrusion comprises a lower hornblende gabbro overlain by olivine websterite and lherzolite. The contacts between lherzolite and olivine websterite are generally gradational. The lherzolite is crosscut by the lower hornblende gabbro. The sulfides are hosted in the lherzolite and lower hornblende gabbro with average grades of 0.64 wt% Ni and 0.48 wt% Cu (Yang et al. 2017). The mineralization comprises



**Fig. 3** a, b Simplified geological map and cross sections of the Lubei intrusion (after Yang et al. 2017)

pyrrhotite, pentlandite, and chalcopyrite with minor violarite. LA-MC-ICP-MS U-Pb zircon dating yielded a weighted mean age of  $287.9 \pm 1.6$  Ma for the Lubei intrusion (Chen et al. 2018).

The lherzolite contains 50–65% olivine, 3–8% orthopyroxene, 5–10% clinopyroxene, 2–15% plagioclase, 5–8% hornblende, and 2% sulfide. The sulfides are commonly interstitial, but small, rounded sulfide inclusions are also enclosed in olivine crystals. The olivine websterite contains 20–35% olivine, 5–10% orthopyroxene, 10–35% clinopyroxene, 5–10% plagioclase, and 5–8% hornblende. Orthopyroxene is intergrown with clinopyroxene or enclosed in clinopyroxene and hornblende. The lower hornblende gabbro contains fine-grained plagioclase (60–70%), clinopyroxene (15–20%), and hornblende (10–20%).

**Analytical methods**

The samples from the Baixintan and Lubei intrusions were from weakly altered outcrops and drill core. The samples from the Baixintan intrusion included the upper hornblende gabbro,

the olivine gabbro, lherzolite, olivine websterite, and the lower hornblende gabbro, whereas the samples collected from the Lubei intrusion included lower hornblende gabbro, olivine websterite, and lherzolite.

**Zircon U-Pb geochronology**

A gabbro sample (16LB-38) from the Lubei intrusion at  $42^\circ 9' 41''$  N and  $90^\circ 20' 39''$  E was selected for dating. The zircons for LA-ICP-MS U-Pb dating were processed by conventional magnetic and heavy liquid separation methods and then handpicked using a binocular microscope to select zircon grains for analysis. Approximately 30 kg of gabbro was crushed for zircon crystal separation. Sample preparation was carried out by the laboratory of the Regional Geological Mineral Survey Institute of Hebei Province, North China and 23 zircon crystals were recovered from the gabbro. The zircon grains were mounted on an epoxy resin disk and polished. Internal structure of the zircon grains was examined using cathodoluminescence (CL) before U-Pb isotopic analyses. The euhedral to subhedral zircons with striped absorption and oscillatory zoning rims in CL images were selected for

U-Pb LA-ICP-MS dating using an Agilent 7500a quadrupole mass spectrometer and GeoLasPro laser ablation system at the School of Resource and Environmental Engineering, Hefei University of Technology. The laser ablation system used a ComPex102ArF excimer laser (wavelength 193 nm) produced by Coherent Inc. of the USA. The standard zircon 91500 (Wiedenbeck et al. 1995) was used to correct for inter-element fractionation, and U, Th, and Pb concentrations were determined based on standard Mud Tank zircons (732 Ma; Black and Gulson 1978). Data processing was carried out using the Isoplot/Ex 2.49 programs of Ludwig (2001).

Zircon crystals from 16LBD-1 were separated using magnetic and density separation methods. Several clear, euhedral grains were isolated and pre-treated according to the chemical abrasion method of Mattinson (2005), with 60 h of annealing at 850 °C followed by 12 h of leaching in HF and HNO<sub>3</sub>. Five single grains were selected and were cleaned with concentrated distilled HNO<sub>3</sub> and HCl. For ID-TIMS analysis, the samples were spiked with an in-house <sup>205</sup>Pb-<sup>235</sup>U tracer solution, which has been calibrated against SRM 981, SRM 982 (for Pb), and CRM 115 (for U), as well as an externally calibrated U-Pb solution (the JMM solution from the EarthTime consortium). This tracer was regularly checked using “synthetic zircon” solutions that yield U-Pb ages of 500 and 2000 Ma, provided by D. Condon (BGS). Dissolution and equilibration of spiked single crystals were by vapor transfer of HF, using Teflon microcapsules in a Parr pressure vessel placed in a 200 °C oven for 6 days. Due to their small size, no chemical separation was required. The resulting residue was redissolved in HCl and H<sub>3</sub>PO<sub>4</sub> and placed on an outgassed, zone-refined rhenium single filament with 5 µL of silicic acid gel. U-Pb isotope analyses were carried out at Curtin University, Perth, using a Thermo Triton Plus mass spectrometer, in peak-jumping mode using a secondary electron multiplier. Uranium was measured as an oxide (UO<sub>2</sub>). Fractionation and dead time were monitored using SRM 981 and SRM 982. Mass fractionation was 0.02 ± 0.07%/amu. All uncertainties are reported at 2σ. Data were reduced and plotted using the software packages Tripoli (from CIRDLES.org) and Isoplot 4.15 (Ludwig 2012).

### Mineral compositions

Olivines from the Baixintan and Lubei intrusive rocks were analyzed by wavelength-dispersive X-ray analysis using a Cameca SX100 electron microprobe at the Australian National University. The accelerating voltage was 15 kV, the beam current was 20 nA, and the counting time was set at 10 s. Standard Program International (SPI) mineral standards (USA) were used for calibration. Analytical results of natural mineral standards are presented in Appendix 1.

### Major and trace elements

Samples were cut with a diamond-impregnated brass blade, crushed in a steel jaw crusher that was brushed and cleaned with de-ionized water between samples, and pulverized in an agate mortar in order to minimize potential contamination. They were analyzed using a PANalytical Axios X-ray fluorescence spectrometer (XRF) on fused glass beads at ALS Chemex (Guangzhou) Co. Ltd. with an analytical uncertainty less than 5%. Of powder, 0.7 g was mixed completely with Li<sub>2</sub>B<sub>4</sub>O<sub>7</sub>-LiBO<sub>2</sub> flux and then fused to a glass bead at 1050–1100 °C in an automatic melting instrument. Analytical results of standard materials and replicate analyses are presented in Appendix 1.

Selected trace elements were determined using a PerkinElmer Sciex ELAN DRC-e ICP-MS at the Institute of Geochemistry, Chinese Academy of Sciences, with analytical uncertainty better than 10%. Samples were digested with 1 ml of HF and 0.5 ml of HNO<sub>3</sub> in screw top PTFE-lined stainless steel bombs at 190 °C for 12 h. The analytical precision is generally better than 1% for elements with concentrations > 200 ppm and 1–3% when less than 200 ppm. The procedure for the trace elements is described in detail by Qi et al. (2000). Reference standards GBPG-1, OU-6, and AMH-1 were used to monitor the trace element analyses (Appendix 1). The analytical uncertainty of this procedure is better than 5% for most elements.

### Chalcophile elements

PGE were determined by ID-ICP-MS using an improved Carius tube technique (Qi et al. 2007) at the Institute of Geochemistry, Chinese Academy of Sciences, Guiyang. Five grams of powdered sample was digested with 40 ml aqua regia in a 75-ml Carius tube placed in a sealed, custom-made, high-pressure, water-filled autoclave. The solution was used to collect PGE by Te coprecipitation (Qi et al. 2004). The measured results of PGE for the reference standard WGB-1 agree well with recommended values reported by Govindaraju (1994). Analytical precision and accuracy are generally better than 10%, and the duplicate samples match each other very well (Appendix 1). PGEs of the blank are generally less than 1 ppb: Ir < 0.05 ppb, Ru < 0.05 ppb, Rh < 0.05 ppb, Pt < 0.05 ppb, and Pd < 0.5 ppb. Nickel, Cu, and Se of the samples containing 3 to 20 vol% sulfides were measured using Varian ICP735-ES inductively coupled plasma emission spectrometer and PerkinElmer Elan 9000 ICP-MS, respectively, at ALS Chemex (Guangzhou) Co. Ltd. Whole-rock S contents were measured at ALS Chemex (Guangzhou) Co. Ltd. with the detection limits ~ 0.01 wt%. The samples were analyzed for total sulfur using a Leco furnace. The sample (0.01 to 0.1 g) was heated to approximately 1350 °C in an induction furnace while passing a stream of oxygen through the sample. Sulfur

dioxide released from the sample was measured by an IR detection system. The analytical precisions are ~8% for S, ~3% for Ni and Cu, and ~10% for Se.

### Sr-Nd-Hf-S isotopes

For radiogenic isotope analysis, approximately 120 mg of powdered sample was placed in Teflon beakers with a HF + HNO<sub>3</sub> mixture and then heated on a hotplate at about 120 °C for 1 week. Strontium and Nd were then separated and purified by conventional cation-exchange techniques. The isotopic compositions of purified Sr and Nd solutions were measured on a TRITON thermal ionization magnetic sector mass spectrometer (TIMS) at the Institute of Geochemistry, Chinese Academy of Sciences, using the procedure described by Yang et al. (2010). Mass fractionation corrections for Sr and Nd isotopic ratios were based on the values of  $^{86}\text{Sr}/^{88}\text{Sr} = 0.1194$  and  $^{146}\text{Nd}/^{144}\text{Nd} = 0.7219$ . The  $^{87}\text{Sr}/^{86}\text{Sr}$  ratios of the NBS-987 Sr isotopic standards and  $^{143}\text{Nd}/^{144}\text{Nd}$  ratios of the BCR-2 Nd isotopic standards determined during this study were  $0.710255 \pm 10$  ( $2\sigma$ ) and  $0.512550 \pm 8$  ( $2\sigma$ ). Detailed sample preparation and analytical procedures followed Zhang et al. (2001).

Hafnium isotopic compositions were measured over three analytical sessions using a ThermoFinnigan Neptune MC-ICP-MS coupled to an ArF excimer laser (193 nm) and fitted with a fast response two-volume He atmosphere HelEx ablation cell at the Australian National University (Eggins et al. 2005). The analytical conditions followed those described by Hiess et al. (2009).

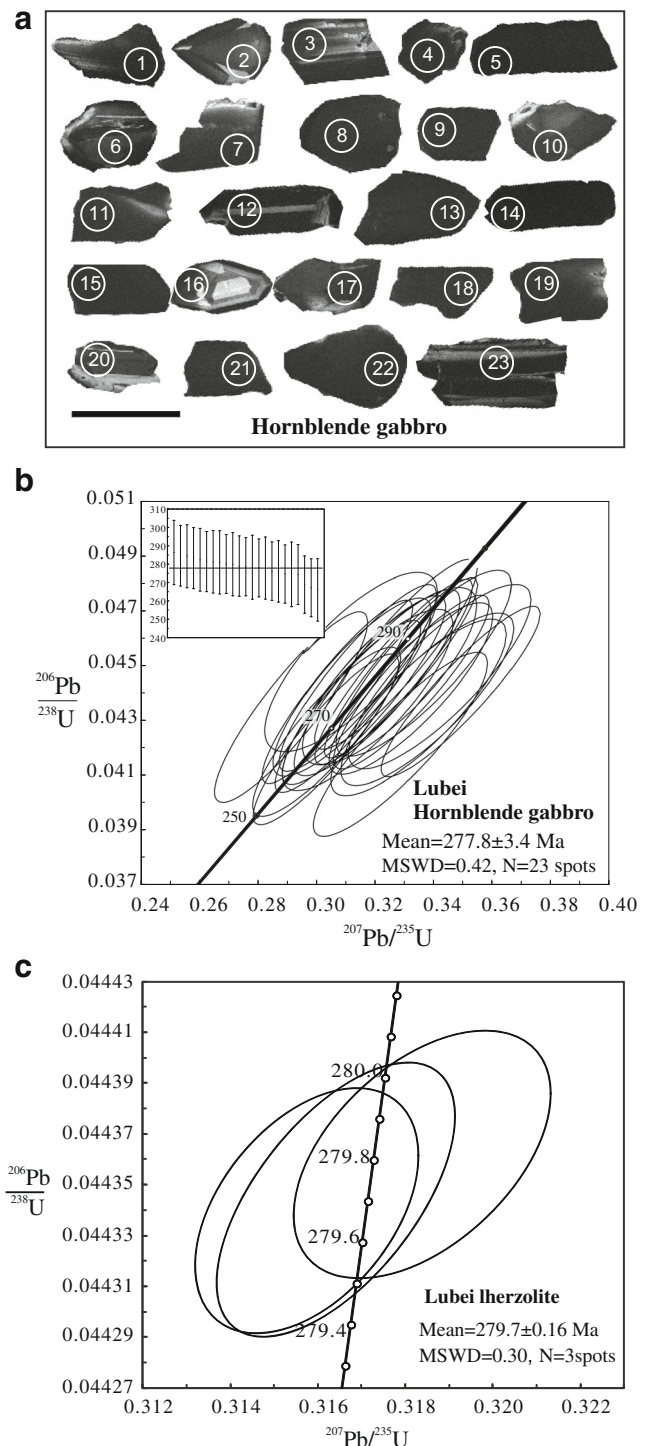
Whole-rock sulfur isotopes of sulfides were measured on a Finnigan MAT 252 continuous flow isotope ratio mass spectrometry at the Institute of Geochemistry, Chinese Academy of Sciences, with an analytical uncertainty less than 0.2‰ (Appendix 1). Analyses of the GBW04414 S standard yielded a value of  $-0.063\text{‰}$  ( $n = 12$ ). All sulfur isotopic data are reported relative to V-CDT in standard  $\delta$  notation.

## Results

### Zircon U-Pb geochronology

The results of LA-ICP-MS U-Pb zircon analyses are listed in Appendix 2 and the U-Pb concordia diagrams are shown in Fig. 4. The zircons have U concentrations of 327–8379 ppm with Th/U ratios of 1.22–2.52. Analyses of the 23 individual zircons yield a concordant age of  $277.8 \pm 3.4$  Ma with a mean square of weighted deviations (MSWD) of 0.42 (Fig. 4) indicating that the gabbro from the Lubei intrusion was formed in the Early Permian.

ID-TIMS U–Pb data of the zircons in the lherzolite (16LBD-1) from the Lubei intrusion are listed in Appendix



**Fig. 4** a, b CL images and Concordia plot of LA-ICP-MS U-Pb analyses of zircons from the hornblende gabbro in the Lubei intrusion. c Concordia plots of ID-TIMS zircon U-Pb geochronological results for the Lubei lherzolite

3. Analyses of three of the five individual zircon crystals form a single concordant group and yield a weighted mean  $^{206}\text{U}/^{238}\text{Pb}$  age of  $279.7 \pm 0.2$  Ma with a MSWD of 0.30 (Fig. 4c). Uranium concentrations were between 100 and 200 ppm, with Th/U ratios between 0.5 and 1.0. Two older

zircons, at 325 and 431 Ma, are considered to be xenocrysts. The crystallization age of the lherzolite is identical to the gabbro of the Lubei intrusion within analytical uncertainties.

### Mineral compositions

Olivine grains were analyzed from the lherzolite and olivine websterite of the Baixintan and Lubei intrusions (Appendices 4 and 5). The olivine crystals in all these samples have low Ca contents (< 1214 ppm; Appendices 4 and 5). The olivines of the lherzolite in the Baixintan intrusion have forsterite (Fo) values varying from 83.0% to 85.2% and Ni from 1486 to 1838 ppm, whereas the olivines of the lherzolite in the Lubei intrusion have higher Fo (84.5–85.8) and Ni contents of 1550 to 2089 ppm.

### Major and trace element geochemistry

Representative major and trace element analyses of the Baixintan and Lubei intrusions are provided in Appendices 6 and 7. The lherzolite of the Baixintan intrusion contains the highest MgO (27.1–27.5 wt%) and  $(\text{Fe}_2\text{O}_3)_T$  (12.1–12.5 wt%) and lowest  $\text{SiO}_2$  (42.3–43.7 wt%) and CaO (3.92–4.40 wt%; Appendix 6). The olivine websterite has lower MgO (20.0–20.3 wt%) and  $(\text{Fe}_2\text{O}_3)_T$  (10.2–10.7 wt%) contents and higher  $\text{SiO}_2$  (44.1–44.8 wt%) and CaO contents (5.79–5.98 wt%) than the lherzolite. The olivine gabbro has higher MgO (13.6–18.9 wt%) but lower  $\text{SiO}_2$  (44.3–45.1 wt%) and CaO contents (6.44–8.87 wt%) than those of the upper and lower hornblende gabbros.

The lherzolite of the Lubei intrusion contains 28.9–33.0 wt% MgO and 38.3–41.2 wt%  $\text{SiO}_2$ . With decreasing MgO,  $(\text{Fe}_2\text{O}_3)_T$  decreases from 14.5 to 12.3 wt% and CaO and  $\text{Al}_2\text{O}_3$  increase from 2.01 and 3.30 to 2.90 and 5.72 wt%, respectively (Appendix 7). The olivine websterite has lower MgO (24.1–28.4 wt%) and higher  $\text{Al}_2\text{O}_3$  and CaO contents (2.09–11.0 wt% and 1.06–7.40 wt%, respectively) than the lherzolite. The hornblende gabbro has the lowest MgO (4.01–15.0 wt%) and highest  $\text{SiO}_2$ ,  $\text{Al}_2\text{O}_3$ , and CaO contents (41.6–49.3, 10.8–17.0, and 2.96–3.42 wt%) of all the samples analyzed.

Normal Mid-Ocean Ridge Basalt (N-MORB) normalized trace element diagrams show that the Baixintan and Lubei intrusive rocks are enriched in large ion lithophile elements (e.g., Rb, Th, U, and La) relative to the high field strength elements (e.g., Zr, Hf, and Yb) which exhibit negative Nb-Ta-Ti anomalies. The Baixintan intrusive rocks contain higher trace element contents than those of the Lubei intrusion (Fig. 5). Except for the olivine websterite of the Lubei intrusion, all the intrusive rocks show positive Sr anomalies. The Baixintan and Lubei rocks are uniformly enriched in LREE relative to HREE ( $(\text{La}/\text{Yb})_N = 1.56\text{--}6.86$ ) and have variable Eu anomalies ( $\delta\text{Eu} = 0.66\text{--}1.29$ ; Fig. 5).

### Chalcophile element geochemistry and S isotopes

The concentrations of PGE, Cu, Ni, S, and Se in the ultramafic rocks and sulfides are presented in Appendix 8. Sulfide contents in the sulfide-mineralized samples were calculated using the procedure of Barnes and Lightfoot (2005):  $C_{(100\%\text{sul})} = C_{\text{wr}} \times 100 / (2.527 \times \text{S} + 0.3408 \times \text{Cu} + 0.4715 \times \text{Ni})$ .  $C_{(100\%\text{sul})}$  is the concentration of Pd or Pt in 100% sulfide;  $C_{\text{wr}}$  is the concentration of the element in the whole rock; and S, Cu, and Ni are the concentrations in weight percent of those elements in the whole rock. The  $\Sigma\text{PGE}$  abundances in the Baixintan intrusive rocks range from 1.87 to 4.41 ppb, which are lower than in the Lubei intrusion (4.06 to 44.3 ppb). The sulfide-bearing intrusive rocks of the Baixintan intrusion have lower  $\Sigma\text{PGE}$  content (6.63–20.6 ppb) than in the Lubei intrusion (9.12–213 ppb; Fig. 6). Copper/Pd ratios of the silicate rocks are between  $4.76 \times 10^4$  and  $2.57 \times 10^5$ , much higher than primitive mantle values (7000–10,000; Barnes and Maier 1999), except one sample (LB-7) which has a Cu/Pd ratio of 8076. The mantle-normalized patterns of Ni, Cu, and PGE, for the intrusive rocks, are PGE depleted relative to Ni and Cu and show PPGE (Pt, Pd, and Rh) enrichments relative to IPGE (Ir, Ru; Fig. 7). The Se/S ratios of the sulfides in the Baixintan and Lubei intrusions range from  $181 \times 10^{-6}$  to  $418 \times 10^{-6}$ , with the majority close to the mantle ratios ( $230\text{--}350 \times 10^{-6}$ ; Eckstrand et al. 1989). The  $\delta^{34}\text{S}_{\text{CDT}}$  values of the sulfides from the Baixintan and Lubei intrusions are low, varying between  $-0.13\text{‰}$  and  $0.90\text{‰}$  (Appendix 8), similar to typical mantle values ( $0 \pm 2\text{‰}$ ; Ohmoto and Rye 1979) but different from those of the country rocks ( $-8.50\text{‰}$  in Baixintan area and  $6.52\text{‰}$  in Lubei area).

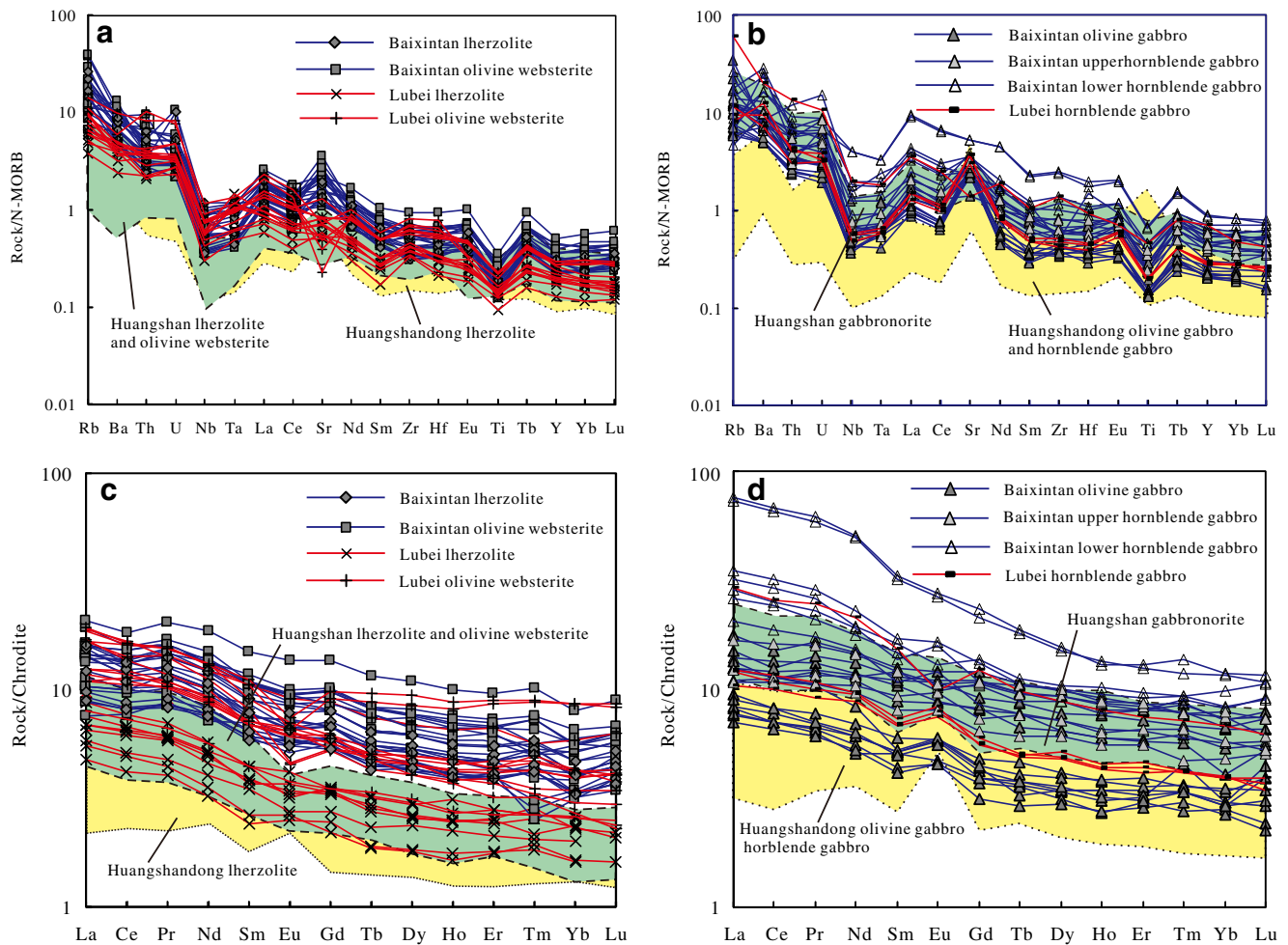
### Sr-Nd-Hf isotopes

Strontium and Nd isotope data for the Baixintan and Lubei intrusions are calculated to have an initial age of 287 and 278 Ma, respectively (Appendix 9; Zhao et al. 2018). The Baixintan intrusive rocks have low initial  $(^{87}\text{Sr}/^{86}\text{Sr})_t$  ratios (0.70354–0.70628) and high  $\epsilon\text{Nd}(t)$  (4.88–7.05), whereas the Lubei intrusive rocks have variable initial  $(^{87}\text{Sr}/^{86}\text{Sr})_t$  ratios (0.70400–0.70713) and  $\epsilon\text{Nd}(t)$  (2.80–6.32; Fig. 8). The Hf isotopes of zircon crystals from Baixintan and Lubei intrusion determined by in situ LA-MC-ICP-MS are listed in Appendix 10. The  $\epsilon\text{Hf}(t = 287 \text{ Ma})$  values of the Baixintan zircons range from 2.5 to 18.4 and the Lubei from 3.18 to 16.1.

### Discussion

Our new zircon U-Pb ages of the lherzolite and hornblende gabbro suggest the Lubei intrusion was emplaced in the Early Permian. The ages of the Baixintan and Lubei intrusions indicate they are contemporary with other known major





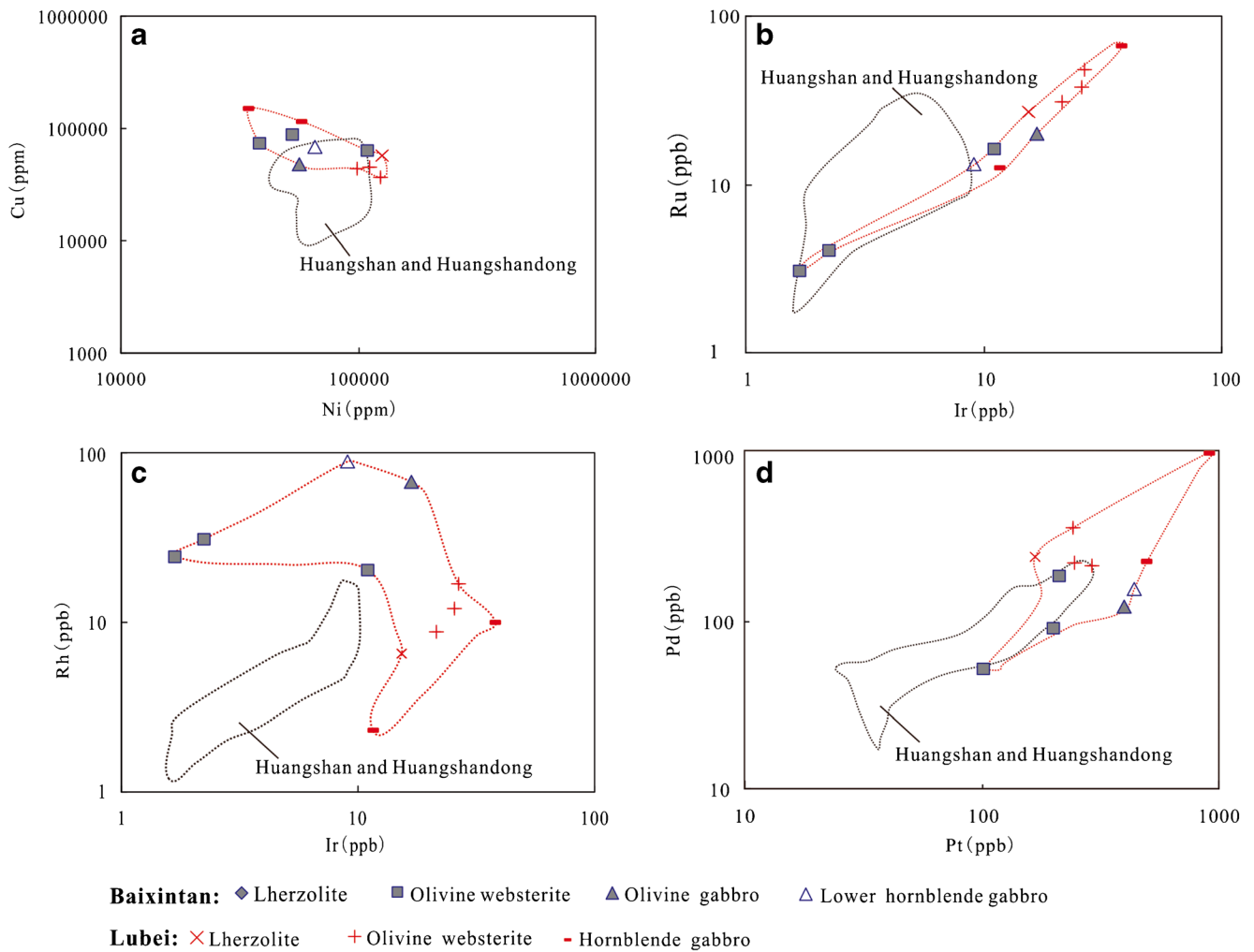
**Fig. 5** a–d N-MORB normalized trace element spider diagrams and chondrite-normalized REE patterns for intrusive rocks in the Baixintan and Lubei intrusions. The data for the Huangshan and Huangshandong intrusions is from Deng et al. (2011a, b, 2014, 2015, 2017). Additional

data for Baixintan intrusion is from Zhao et al. (2018) and Wang et al. (2015). N-MORB normalization and REE chondrite values are from Pearce (1982) and Sun and McDonough (1989), respectively

magmatic Ni-Cu sulfide deposits in the Northern Tianshan (Qin et al. 2011; Zhao et al. 2018; Feng et al. 2018; Chen et al. 2018). The Baixintan and Lubei intrusions are characterized by enrichment of Rb, Ba, U, and Sr, in conjunction with depletions of Nb, Ta, and Ti relative to N-MORB, which is similar to the other Ni-Cu sulfide-bearing intrusions in the Northern Tianshan. However, the  $\epsilon_{Nd(t)}$  values of the Baixintan and Lubei intrusive rocks are lower than those of the Huangshan and Huangshandong intrusive rocks, and the  $\epsilon_{Hf(t)}$  values are more variable (Fig. 8; Su et al. 2012; Sun et al. 2013). The Cu and PGE contents of the sulfides and intrusive rocks in the Baixintan and Lubei intrusions are higher than those of the Huangshan and Huangshandong intrusions (Figs. 6 and 7). The olivine from the lherzolites in the Baixintan intrusion contains higher Ni content than those of the latter. This suggests that the Baixintan and Lubei rocks in the west of the Northern Tianshan underwent a different magma evolution process than the rocks from the Huangshan and Huangshandong intrusions in the east.

### Nature of the mantle source

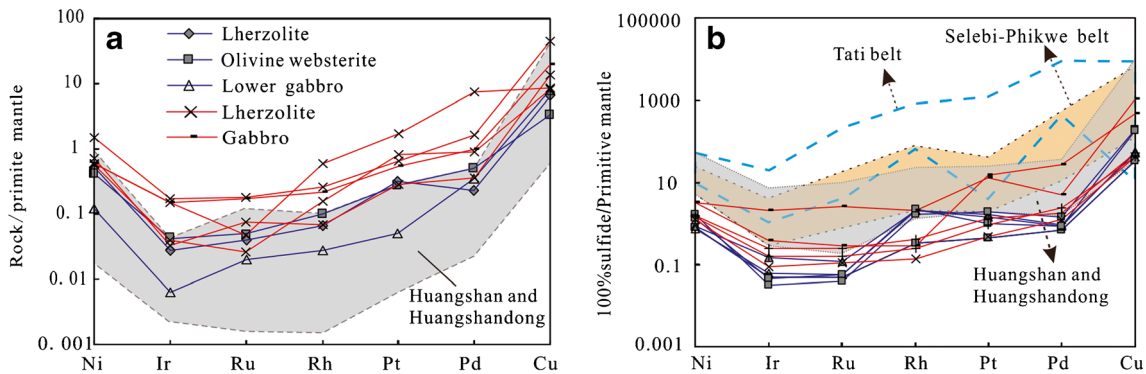
The samples from the Baixintan and Lubei intrusions are enriched in large ion lithophile element and depleted in Nb, Ta, and Ti relative to N-MORB. The depletion in high field strength elements (such as Nb, Ta, and Ti) is normally interpreted to reflect magma generation in a subduction-related environment (Pearce 1983; Hawkesworth et al. 1997; Johnson and Plank 1999) or addition of relatively old continental crust (Wilson 1989). The similar bulk solid/melt partition coefficient of Nb, U, and La (Hofmann 1988; Sun and McDonough 1989) means that they will not be significantly fractionated during magma crystallization processes, and consequently, the Nb/U and La/Nb ratios of the intrusive rocks can be compared with the volcanic rocks to trace the ratios in the mantle source or crustal contamination process. Compared with average crust, N-MORB, ocean island basalt (OIB), and primitive mantle, the Baixintan and Lubei intrusions have lower Nb/U ratio and higher La/Nb ratios, respectively



**Fig. 6** a–d Plots of platinum group elements of the sulfides from the Baixintan and Lubei intrusions on a 100% sulfide basis. The data for the Huangshan and Huangshandong intrusions is from Deng et al. (2014, 2017)

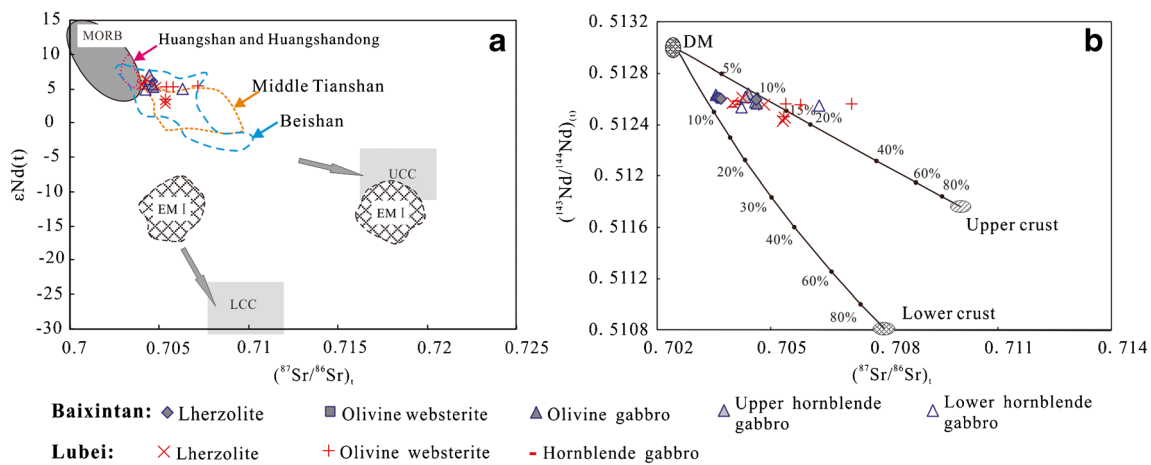
(Appendix 11). Consequently, these trace elements ratios cannot be formed by the contamination of the mantle end-member with the crust but reflect the features of the mantle source of the intrusions. The Nb/U and La/Nb ratios of the

Baixintan and Lubei intrusive rocks are similar to Alaskan-type intrusions (Pettigrew and Hattori 2006), suggesting that the primary magma of the intrusions was derived from a metasomatized mantle source.



**Fig. 7** a, b PGE patterns of intrusive rocks from the Baixintan and Lubei intrusions normalized to primitive mantle (Barnes and Maier 1999). Data for the Huangshan and Huangshandong intrusive rocks is from Deng

et al. (2014, 2017). Also shown are fields for sulfide ores from the Tati and Selebi-Phikwe belts of eastern Botswana (Maier et al. 2008)



**Fig. 8** Plot of **a**  $\epsilon Nd(t)$  versus  $(^{87}Sr/^{86}Sr)_t$ . **b** Modeling calculations of Sr-Nd isotopes of the Baixintan and Lubei intrusions. Additional data from the Baixintan intrusive rocks is from Zhao et al. (2018). Data for the Central Tianshan and Beishan intrusions are from Sun et al. (2013). The

fields for depleted MORB mantle, upper continental crust (UCC), and lower continental crust (LCC) are from Zindler and Hart (1986), Rudnick and Gao (2003), and Windrim and McCulloch (1986), respectively

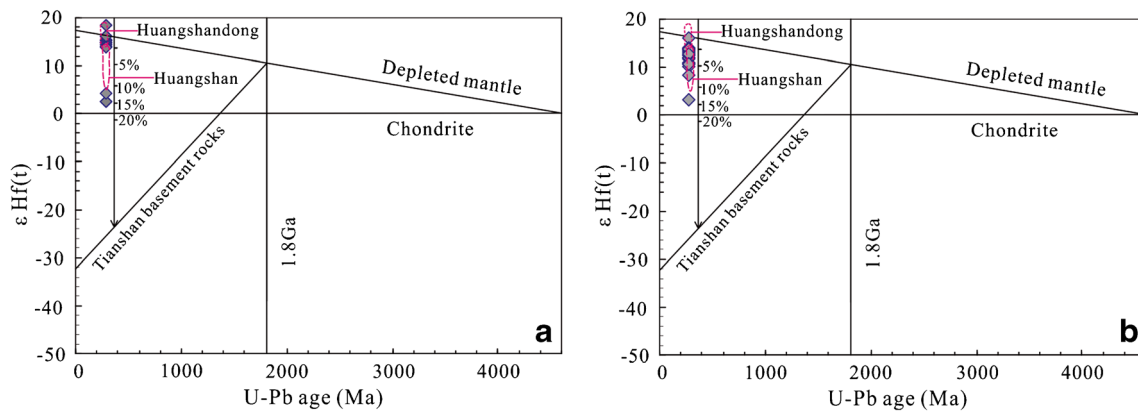
**Sulfur saturation**

Previous studies have proposed that a metasomatized mantle source may experience high degree melting at a relative low temperature and generate the Ni-rich basaltic magmas which will form Ni-Cu sulfide deposits (Song and Li 2009; Ripley 2009). The primary magma of the Permian mafic-ultramafic intrusions from the western and eastern parts of the Northern Tianshan was derived from metasomatized mantle source (Zhang et al. 2011; Deng et al. 2014, 2015). Given the different scale of mineralization in the east and west, we explore what role the sulfide saturation mechanism played in generating the mineralization in the Northern Tianshan.

The Baixintan and Lubei intrusive rocks are characterized by variable initial  $(^{87}Sr/^{86}Sr)_t$  ratios (0.70354–0.70713) and  $\epsilon Nd(t)$  (2.80–7.05) and define a trend towards the upper crust (Fig. 8a). The isotopic variation suggests an upper crustal contamination either in the mantle source or during magma ascent. As the Precambrian crystalline basement of the Central Tianshan has not been subducted beneath the Northern Tianshan, the  $\epsilon Nd(t)$  values of the intrusive rocks cannot be inherited from the mantle source which was metasomatized by subducted crustal materials (Xiao et al. 2004). Modeled  $(^{87}Sr/^{86}Sr)_t$  and  $\epsilon Nd(t)$  values show that < 20% contamination with the upper crust can explain the isotopic variations of the most samples in the Baixintan and Lubei intrusion (Fig. 8b). Previous studies suggest there is no Precambrian basement in the Northern Tianshan. However, zircons from andesites in the Lubei area have yielded concordant U-Pb ages of  $325 \pm 6$  Ma and contain five zircon xenocrysts with an apparent  $^{206}Pb/^{238}U$  age of 1602~1870 Ma (our unpublished data), suggesting that Precambrian crystalline basement has been incorporated in the formation of the volcanic rocks. Furthermore, a Mesoproterozoic metamorphic core complex

has been discovered in the western part of the Northern Tianshan (Su et al. 2008). Thus, we propose that the Mesoproterozoic metamorphic core complex may be the source of the crustal contamination. Additionally, all zircon  $\epsilon Hf(t)$  values of the Baixintan and Lubei intrusions are positive, falling into the range between chondrite and depleted mantle evolution trend lines, consistent with a parental magma derived from a depleted source mantle. The  $\epsilon Hf(t)$  values of the Baixintan and Lubei intrusive rocks indicate that they have assimilated Tianshan basement rocks (Fig. 9).

The role of crustal sulfur in magmatic Ni-Cu sulfides can be constrained using Se/S ratios and  $\delta^{34}S_{CDT}$  (Eckstrand et al. 1989; Ripley et al. 2002; Maier et al. 2008). The Se/S ratios of the sulfides in the Baixintan and Lubei intrusions range from  $181 \times 10^{-6}$  to  $418 \times 10^{-6}$ , with the majority close to the mantle ratios rather than crustal ratios (Eckstrand et al. 1989). These values are higher than those of the Huangshan and Huangshandong sulfide ores, which were segregated from magma that has assimilated crustal sulfur (Fig. 10; Mao et al. 2002; Tang et al. 2012; Deng et al. 2014, 2017). The  $\delta^{34}S_{CDT}$  values of the sulfides from the Baixintan and Lubei intrusions are low, varying between  $-0.13\text{‰}$  and  $0.90\text{‰}$ , similar to typical mantle values ( $0 \pm 2\text{‰}$ ; Ohmoto and Rye 1979) and the sulfide ores from the Huangshan and Huangshandong deposits ( $-0.8\text{‰}$  to  $2.8\text{‰}$ ; Deng et al. 2014, 2017) but different from those of the country rocks ( $-8.50\text{‰}$  in Baixintan area and  $6.52\text{‰}$  in Lubei area). The Se/S ratios and  $\delta^{34}S_{CDT}$  values of the Baixintan and Lubei sulfides resemble those values of mantle-derived rocks but are distinct from the crustal values, suggesting crustal sulfur has not been incorporated in the parent magma (Ohmoto and Rye 1979; Eckstrand et al. 1989). Consequently, sulfide saturation was not caused by assimilation of crustal sulfur in the Baixintan and Lubei intrusions.

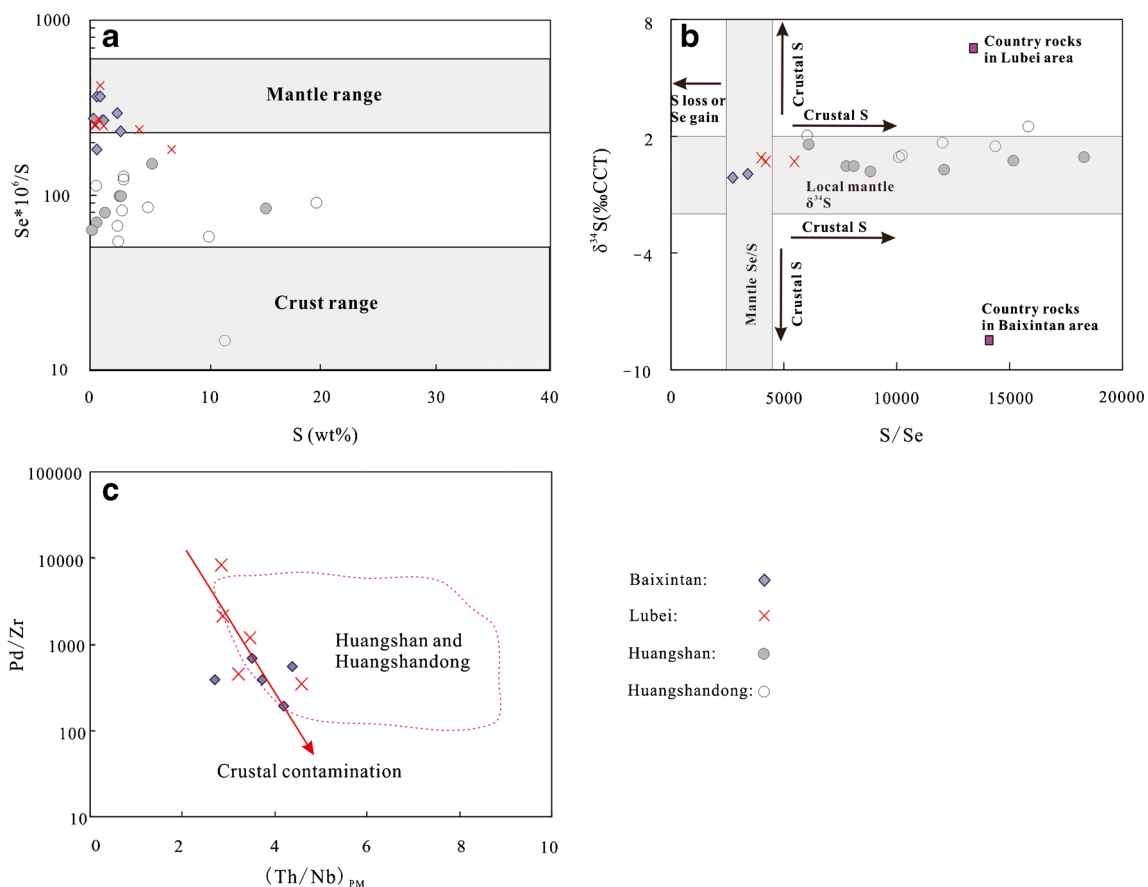


**Fig. 9** a, b Plot of  $\epsilon_{\text{Hf}}(t)$  versus U-Pb age of zircons of the Baixintan and Lubei intrusions. The age of the Tianshan basement rock is based on Hu et al. (2000). The concentration of Hf in N-MORB from Sun and McDonough (1989) is used to represent the composition of melt

derived from depleted mantle. Data of the Huangshan and Huangshandong intrusions are from Su et al. (2013) and Sun et al. (2013). The numbers near the sample points are the degree of crustal contamination

A key question is what has driven the mafic magma to sulfide saturation in the Baixintan and Lubei intrusions. Because Pd/Zr are exceptionally sensitive to sulfide fractionation, they can be used to trace the segregation of even small amounts of magmatic sulfides from mafic magmas. Increasing  $(\text{Th}/\text{Nb})_{\text{PM}}$  (where PM = primitive mantle normalized) ratios

are generally accepted to be the products of increasing degrees of crustal contamination (Lightfoot and Hawkesworth 1988; Keays and Lightfoot 2010). The similar bulk solid/melt partition coefficient of Nb and Th (Sun and McDonough 1989; Aldanmaz et al. 2000) means that they will not be significantly fractionated during magma crystallization processes, so Th/



**Fig. 10** Plots of a  $\text{Se} \cdot 10^6/\text{S}$  versus S, b  $\text{Se} \cdot 10^6/\text{S}$  versus  $\delta^{34}\text{S}_{\text{CCT}}$  and c  $(\text{Th}/\text{Nb})_{\text{PM}}$  versus Pd/Zr for the Baixintan and Lubei mineralized rocks. Additional data for the Baixintan intrusion is from Zhao et al. (2018). The

data for the Huangshan and Huangshandong ores are from Deng et al. (2014, 2017). The range in Se/S ratio of the mantle and crust is taken from Eckstrand et al. (1989)

Nb ratios of both intrusive and volcanic rocks can trace the crustal contamination process. The negative correlation between Pd/Zr and  $(\text{Th}/\text{Nb})_{\text{PM}}$  for the Baixintan and Lubei intrusive rocks suggests crustal contamination and chalcophile metal depletion (Fig. 10c). Crustal contamination would result in addition of  $\text{SiO}_2$  and decreasing temperature which in turn could induce sulfide saturation by decreasing sulfur solubility (Lightfoot and Hawkesworth 1997; Li and Naldrett 2000; Seat et al. 2009). Experimental studies indicated that sulfur solubility of mafic magma decreases with decreasing  $\text{Fe}^{2+}$  content at a constant pressure (Wendlandt 1982; Mavrogenes and O'Neill 1999), so fractionation of mafic silicates may also play an important role in triggering sulfide saturation by sharply lowering the Fe concentration.

## Origin of sulfide mineralization

### Modeling of olivine compositional variations

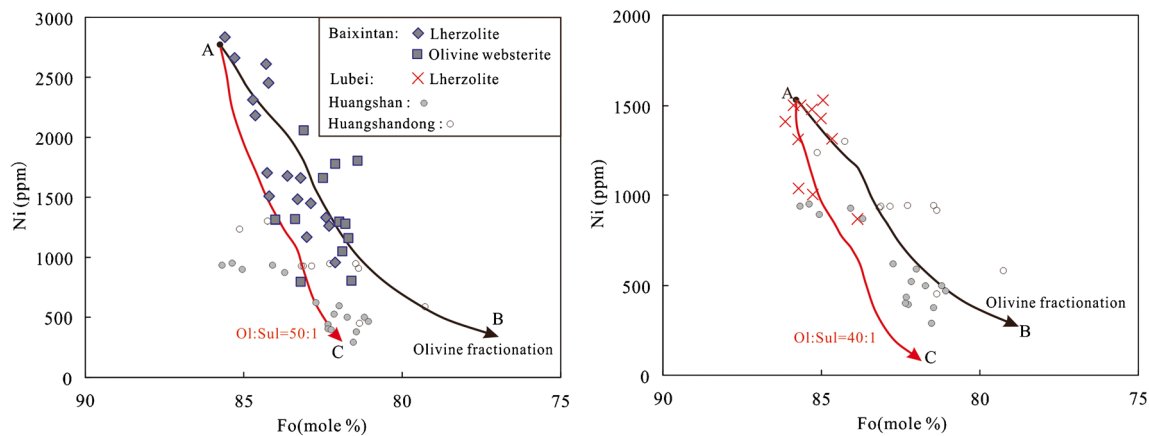
As discussed above, Sr-Nd isotopic compositions indicate the Baixintan and Lubei intrusive rocks assimilated the lower crust in the Northern Tianshan, which would also have resulted in sulfide saturation (Lightfoot and Hawkesworth 1988; Keays and Lightfoot 2010). If crustal contamination has triggered extensive sulfide segregate in the lower crust, the sulfide would accumulate and form Ni-Cu sulfide mineralization whereas the residual metal-depleted magma would not form Ni-Cu sulfide deposits in the upper crust. On the other hand, if the crustal contamination only triggered trace sulfide saturation in the lower crust, the sulfide would extract some PGE from the magma and the residual PGE-depleted magma may form a Ni-Cu sulfide deposit in the shallow crust, as Ni and Cu would have been less depleted in the residual magma due to the lower partition coefficients in sulfide melt than that of PGE. Thus, it is necessary to investigate the sulfide segregation process to evaluate the exploration potential in the Baixintan and Lubei intrusions.

Because most of the lherzolites have poikilitic textures with abundant cumulate olivine and interstitial silicate minerals, we have used the method of Li and Ripley (2011) to estimate the parental magma composition of the Baixintan and Lubei intrusions. In our calculations, we used the average composition of sulfide-barren lherzolite samples containing the most primitive olivine ( $\text{Fo} = 85$ , Baixintan;  $\text{Fo} = 86$ , Lubei), the olivine-liquid exchange coefficient  $(\text{FeO}/\text{MgO})_{\text{Ol}}/(\text{FeO}/\text{MgO})_{\text{Liq}}$  of 0.3 by Roeder and Emslie (1970), and an assumed  $\text{FeO}/(\text{FeO} + \text{Fe}_2\text{O}_3)$  ratio of 0.95 and 0.98 for the liquid in the Baixintan and Lubei intrusions, respectively. The estimated MgO contents in the parental magma of Baixintan and Lubei intrusions are 7.90 and 7.79 wt%, respectively (Appendix 12).

Olivine chemistry is controlled by parental magma composition, fractional crystallization, sulfide segregation,

subsolidus reequilibration with interstitial silicate, and trapped sulfide liquid; consequently, they are used as an indicator of magma evolution and sulfide saturation (Barnes 1986; Barnes et al. 2011, 2013; Li et al. 2001, 2007). We have used the PELE program (Boudreau and Meurer 1999) to investigate the fractional crystallization and sulfide segregation processes of the parental magma for the primitive lherzolite in the Baixintan and Lubei intrusions at 1 kb total pressure and a constant oxidation state equal to that of one log unit above the fayalite-magnetite-quartz buffer. The Rayleigh fractionation equation has been used to describe models for the Ni contents in olivine. In this paper, the partition coefficient of 7 was used for Ni between olivine and magma and 500 between sulfide liquid and magma (Li and Ripley 2005). The olivine Fo value and Ni content correlation curve A–B are the modeled result of normal magmatic evolution (Fig. 11). Curve A–C is the trend of olivine Fo value and Ni content when olivine and sulfide crystallized together. The observed Ni-Fo variations of olivine from the ultramafic rocks can be explained by olivine having crystallized from the silicate liquid that coexisted with the sulfide liquid (Fig. 11).

Although olivines from the lherzolite in the Baixintan and Lubei intrusions have similar Fo values to those of the Huangshan and Huangshandong intrusions, the Ni contents of most olivines in the Baixintan and Lubei (796–2664 ppm) are higher than those of the Huangshan and Huangshandong (282–1295 ppm; Fig. 11). What causes the Ni contents differences in olivines between the intrusions in the west part and east part of Northern Tianshan? Since samples containing  $> 0.5$  wt% S were excluded, the differences in olivine composition did not result from postcrystallization reaction between olivine and sulfide. Similarly, because the olivine compositions of these intrusions do not plot along the modeled path of normal olivine crystallization, fractional crystallization cannot account for the variations of the Ni contents of olivines (Fig. 11). “Trapped silicate liquid shift” generally generates olivine with a composition that is poorer in Mg and Ni contents than the original material (Li and Ripley 2003; Li et al. 2007). The Baixintan and Lubei intrusions have similar Fo values of olivines to those of the Huangshan and Huangshandong intrusions, suggesting that the lower Ni contents in olivine compositions of the Huangshan and Huangshandong intrusions cannot be explained by subsolidus reequilibration with interstitial silicate liquid. The Ni contents of most primitive olivine in the Baixintan intrusion are high (up to 2664 ppm), similar to those of most primitive olivine from in the Huangshannan magmatic Ni-Cu ore deposit ( $\sim 3300$  ppm Ni in olivine, Mao et al. 2017a), and higher than that in Jinchuan and Noril'sk deposits (Li and Ripley 2003; Li et al. 2004). Our modeling yielded estimates of 400 and 220 ppm Ni for the parental magma from which the olivine crystallized in the Baixintan and Lubei intrusion, respectively. As noted above, the high Ni contents of olivine in the Baixintan and Lubei intrusions were not the result of fractional crystallization, sulfide segregation, and subsolidus



**Fig. 11** Model calculation of the variation of nickel content and Fo of the olivine in the Baixintan intrusion (a) and Lubei intrusions (b). For detailed methodology, see Li and Naldrett (1999) and Li et al. (2007). The olivine compositions of the Huangshan and Huangshandong intrusions are from our unpublished data. A is the olivine composition in equilibrium with the parental magma. We use a partition coefficient of

reequilibration; rather, they may have been the result of high Ni contents in the parent magmas.

### Quantitative modeling of sulfide segregation

The parental magma of the Baixintan and Lubei intrusions had higher Ni contents than the Huangshan and Huangshandong intrusions. Consequently, we propose that crustal contamination in the Baixintan and Lubei intrusions would not have triggered extensive sulfide segregation in the lower crust. We investigate the sulfide segregation history of the Baixintan and Lubei intrusions by examining the composition of sulfides. Metal contents of sulfide melts segregated from silicate magma are a function of the  $R$  factors (ratio between silicate melt and sulfide liquid) and the metal partition coefficients (Campbell and Naldrett 1979):

$$C_{sul} i = C_{sil} i \times D_i \times (R + 1) / (R + D_i) \quad (1)$$

where  $C_{sul} i$  and  $C_{sil} i$  represent the content of element  $i$  in the sulfide liquid and in the parental silicate liquid, respectively,  $D_i$  is the partition coefficient of element  $i$  between sulfide liquid and silicate liquid, and  $R$  is the  $R$  factor.

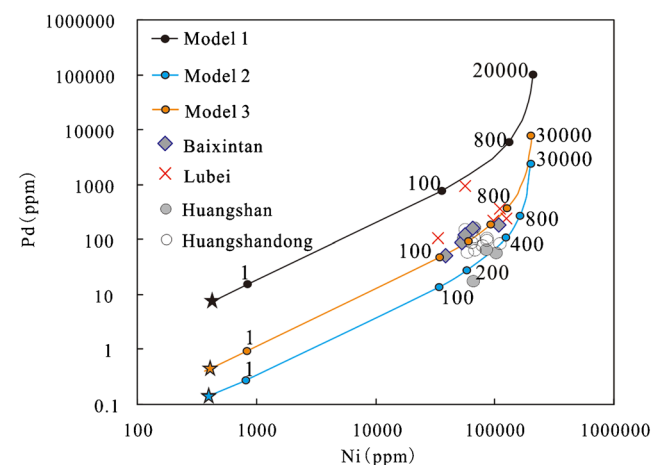
The geochemical signatures of the Baixintan and Lubei intrusions suggest that the parental magmas were derived from metasomatized mantle similar to those of island arc magmas. Consequently, we assume that the parent magma is similar to typical island arc basalt, with 7.5 ppb Pd and 400 ppm Ni (Barnes et al. 1993; Kelemen et al. 2004; Thakurta et al. 2008). The partition coefficients of Pd and Ni between sulfide and silicate magma were assumed to be 40,000 and 500, respectively (Peach et al. 1990; Fleet et al. 1993). The PGE modeling suggests that the source magma for the Baixintan and Lubei intrusions underwent two stages of sulfide

segregation. First, the primary magma underwent 0.007% early sulfide removal at depth and lost much of its chalcophile content (Fig. 12). Then, sulfides in the Baixintan and Lubei intrusions were segregated from the strongly PGE-depleted parent magma under  $R$  factors of 100 to 800 in a shallow magma chamber.

The sulfide-bearing intrusive rocks of the Baixintan intrusion have lower  $\Sigma$ PGE contents (6.63–20.6 ppb) than those of the Lubei intrusion (9.12–213 ppb). The Cu and PGE contents

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The sulfide-bearing intrusive rocks of the Baixintan intrusion have lower  $\Sigma$ PGE contents (6.63–20.6 ppb) than those of the Lubei intrusion (9.12–213 ppb). The Cu and PGE contents



**Fig. 12** Variations of metal tenors in the disseminated sulfide from the Baixintan and Lubei intrusions. The Pd and Ni concentrations of the sulfides are on a 100% sulfide basis. Model 1 presents the sulfide melts segregated from basaltic magma with 7.5 ppb Pd and 430 ppm Ni under variable  $R$  values (labeled in the diagram). Model 2 displays sulfide liquids separated from PGE-weakly depleted magma with 0.46 ppb Pd and 415 ppm Ni (because of 0.007% sulfide removal from magma) under different  $R$  values. Model 3 shows the sulfide liquids produced by sulfide immiscibility in a heavily PGE-depleted magma with 0.14 ppb Pd and 409 ppm Ni (because of 0.01% sulfide removal from magma). The PGE data of the Huangshan and Huangshandong sulfide ores are from Deng et al. (2014, 2017) and Mao et al. (2014, 2015)

of the sulfides in the Baixintan and Lubei intrusions are higher than those of the Huangshan and Huangshandong intrusions (Figs. 6, 7, and 12). The chalcophile element abundance of magmatic sulfides is controlled by some combination of the concentration of these elements in the parental silicate magma, the  $R$  factor, the degree of fractional crystallization of the sulfide liquid, and late hydrothermal alteration (Keays 1995; Barnes et al. 1997; Barnes and Lightfoot 2005). The ratios of  $\text{Ir}/(\text{Ir} + \text{Ru})$  and  $\text{Pt}/(\text{Pt} + \text{Pd})$  (0.36–0.48 and 0.41–0.82, respectively) of the Baixintan and Lubei intrusions are typical of magmatic sulfide (Naldrett et al. 1982), indicating that hydrothermal alteration has a limited effect on the PGE compositions of the sulfides in the Baixintan and Lubei intrusions. Commonly, the fractional crystallization of a sulfide liquid has a greater effect on the chalcophile element compositions of semi-massive and massive sulfides than those of disseminated sulfides (Naldrett 1994). As only disseminated sulfides were investigated, fractional crystallization of sulfides cannot have caused the variation in PGE contents. Moreover, the sulfides in the Baixintan and Lubei intrusions segregated from the magma under  $R$  factors of 100–800, similar to those of the Huangshan and Huangshandong intrusions (Sun et al. 2013; Deng et al. 2014, 2017; Mao et al. 2014, 2015), suggesting that the  $R$  factor did not cause the different PGE contents in sulfides. If more sulfides were removed from the primary magma, the residual magma would be more depleted in PGE. The sulfides segregated from this residual magma would have relatively low PGE contents (Fig. 12). Compared with the Baixintan and Lubei intrusions, the Huangshan and Huangshandong magmas may have experienced more sulfide removal at depth (Fig. 12; Gao et al. 2013; Sun et al. 2013; Deng et al. 2014, 2017). The early removal of sulfide in the Huangshan and Huangshandong intrusions would remove PGE from the magma, resulting in the residual magmas being more depleted in PGE than those in the Baixintan and Lubei intrusions. The more PGE-depleted residual magmas in the Huangshan and Huangshandong intrusions formed sulfide ores with lower PGE contents than those of the Baixintan and Lubei intrusions. In summary, the magmas of the Baixintan and Lubei intrusions were sulfide-saturated in the lower crust and a small amount of sulfides (~0.007%) was removed in staging chambers at depth. A second stage of sulfide saturation was reached in shallow magma chambers and formed the Ni-Cu sulfide mineralization.

### Implications for Ni-Cu sulfide prospectivity

The most important factors required for the formation of a magmatic Ni-Cu sulfide deposit include (1) large amount of mantle-derived magma ascending into the crust, (2) the interaction of the source magma with country rocks to induce sulfide saturation, (3) the concentration of sulfides in conduits through which much magma has flowed, and (4) Ni-Cu upgrading of sulfide

liquid by new, chalcophile-undepleted magma (Naldrett 1997, 1999; Li et al. 2000; Maier et al. 2001; Barnes and Lightfoot 2005). If any mafic-ultramafic intrusive rocks develop these geological features, they would have exploration potential for the magmatic Ni-Cu sulfide deposit.

The mafic-ultramafic intrusions in the western part of the Northern Tianshan formed in the early Permian (278–287 Ma; Fig. 4; Zhao et al. 2018; this study), coeval with the Ni-Cu sulfide-bearing intrusions in the eastern part of the Northern Tianshan. The trace element and isotopic composition of the Baixintan and Lubei intrusions suggest that the primary magmas were derived from metasomatized lithospheric mantle. Combined with the widespread Permian mafic-ultramafic intrusions in the western part of the Northern Tianshan, these intrusions are interpreted to be linked to large volumes of post- or syn-collisional tholeiitic magmatism induced by asthenospheric upwelling (Li et al. 2006b; Song et al. 2011, 2013; Gao et al. 2013; Sun et al. 2013; Mao et al. 2014, 2015; Deng et al. 2015, 2017; Feng et al. 2017). Secondary faults along the Huangshan-Kangguer fault likely provided zones of weakness along which magma ascended. Although no obvious crustal source of sulfur was incorporated into the magma, sulfide liquid saturation was achieved due to crustal contamination and fractional crystallization of mafic silicates and chromites. The modeling of PGE suggests that the sulfide liquids underwent upgrading of PGE and Ni by reaction with successive pulses of magma under  $R$  factors of 100–800 (Fig. 12). Moreover, the olivine compositions and chalcophile elements of mafic-ultramafic intrusions suggest that the parental magma of the intrusive rocks in the western part of the Northern Tianshan contains higher chalcophile elements than those of the eastern part. The Permian mafic-ultramafic intrusions in the west of the Northern Tianshan have a similar mantle source to the Huangshan and Huangshandong intrusions and experienced sulfide saturation and metal upgrading processes. Therefore, they should be regarded as favorable Ni-Cu sulfide exploration targets. All world-class magmatic Ni-Cu deposits have been contaminated by crustal sulfur to achieve sulfide saturation (Naldrett 1999; Keays and Lightfoot 2010; Ripley and Li 2013). Previous studies proposed that more than 50% sulfur in some magmatic Ni-Cu sulfide deposits are from country rocks (Kabanga, Maier and Barnes 2010; Noril'sk, Robertson et al. 2015). Compared to the Huangshan and Huangshandong intrusive rocks, the  $\epsilon\text{Nd}_{(t)}$  and the  $\epsilon\text{Hf}_{(t)}$  values of the Baixintan and Lubei intrusive rocks are relatively low, suggesting that Huangshan and Huangshandong rocks have assimilated more crustal material than the Baixintan and Lubei rocks (Fig. 8a). The Ni contents in the parental magma of the Baixintan and Lubei intrusions are higher than those of the Huangshan and Huangshandong deposits. As the

Baixintan and Lubei intrusions assimilated less crustal sulfur (Fig. 10), even though the volumes of basalt were similar, it is likely that the country rocks did not supply as much sulfur as the Huangshan and Huangshandong deposits. Consequently, the Baixintan and Lubei intrusions do not have the same Ni-Cu potential as the Huangshan and Huangshandong deposits.

## Conclusions

Zircon U-Pb geochronology shows that the Lubei mafic-ultramafic intrusion formed at 278 Ma. The primary magma of the Baixintan and Lubei mafic-ultramafic intrusions was derived from metasomatized mantle. The Baixintan and Lubei intrusive rocks have assimilated Precambrian basement in the western Northern Tianshan. Sulfide saturation was triggered by crustal contamination and fractional crystallization rather than crustal S addition. The parental magma of the intrusive rocks in the Baixintan and Lubei intrusions contains higher chalcophile elements than those of the Huangshan and Huangshandong intrusions.

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