## **Original Article**

## Ages and Sources of Ore-Related Porphyries at Yongping Cu–Mo Deposit in Jiangxi Province, Southeast China

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#### Abstract

Whole-rock geochemistry, zircon U–Pb and molybdenite Re–Os geochronology, and Sr–Nd–Hf isotopes analyses were performed on ore-related dacite porphyry and quartz porphyry at the Yongping Cu–Mo deposit in Southeast China. The geochemical results show that these porphyry stocks have similar REE patterns, and primitive mantle-normalized spectra show LILE-enrichment (Ba, Rb, K) and HFSE (Th, Nb, Ta, Ti) depletion. The zircon SHRIMP U–Pb geochronologic results show that the ore-related porphyries were emplaced at 162–156 Ma. Hydrothermal muscovite of the quartz porphyry yields a plateau age of 162.1  $\pm$  1.4 Ma (2 $\sigma$ ). Two hydrothermal biotite samples of the dacite porphyry show plateau ages of 164  $\pm$  1.3 and 163.8  $\pm$  1.3 Ma. Two molybdenite samples from quartz+molybdenite veins contained in the quartz porphyry yield Re–Os ages of 156.7  $\pm$  2.8 Ma and 155.7  $\pm$  3.6 Ma. The ages of molybdenite coeval to zircon and biotite and muscovite ages of the porphyries within the errors suggest that the Mo mineralization was genetically related to the magmatic emplacement. The whole rocks Nd–Sr isotopic data obtained from both the dacite and quartz porphyries suggest partial melting of the Meso-Proterozoic crust in contribution to the magma process. The zircon Hf isotopic data also indicate the crustal component is the dominated during the magma generation.

**Keywords:** Cu–Mo porphyry, geochronology, magma source, Southeast China, Sr–Nd–Hf isotopes, Yongping deposit.

#### 1. Introduction

The Yongping Cu–Mo deposit is located 200 km east of Nanchang, the capital city of Jiangxi Province, China. The deposit occurs between the Yangtze Cu–Mo metallogenic belt and the south Jiangxi W–Sn–REE metallogenic belt (Jiangxi Geological Exploration Bureau, abbreviated as JGEB, 1996). Its location is close to a bend of the regional Jiangshan-Shaoxing fault, a Neo-Proterozoic suture zone between Yangtze and Cathaysa cratons (Fig. 1). The Cu orebodies at the Yongping deposit were discovered in the 1960s. Openpit mining started in 1984. The Mo orebodies were discovered in 2007 with an average grade of 0.1%. As publically reported, a total of 3.4 Mt ores with a grade of 0.6% Cu and 0.1% Mo were yielded in 2007.

Studies on structural and ore geology, mineralization, geochronology, and hydrothermal alteration have been conducted at the Yongping Cu–Mo deposit (Gu & Xu, 1986; Liu & Huang, 1991; He, 1993; Ding *et al.*, 2005;

Received 12 May 2012. Accepted for publication 18 September 2012.

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**Fig. 1** Simplified maps showing tectonic (a and b) and geologic (c) setting and the location of the Yongping Cu–Mo deposit and adjacent, major Cu-polymetallic deposits (c) in Jiangxi Province, southeast China. SECMY = the southeast continental margin of Yangtze craton (block, plate); NWCMC = the northwest continental margin of Cathaysia craton (block, plate); SCSZ = the South China Suture Zone; YB = Yangtze Block (craton, plate); CB = South China Block (plate) (modified after Chen, 1999).

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Fig. 2 Simplified geological map of Yongping Cu-Mo deposit (modified after Huang et al., 2001).

Du, 2005; Li et al., 2007a). The arguments of the genesis of the deposit have been controversial; the deposit formed by migmatization (He, 1993), metamorphic in origin (Ren, 1994) and mixed origin of sedimentary exhalative (SEDEX) and porphyry mineralization (Gu & Xu, 1986; Liao & Liu, 2003; Du, 2005). However the published geochemical data do not support SEDEX model because the sulfur isotopes of sulfide minerals show considerable discrepancy with that in the host rocks of the Carboniferous Yejiawan Group. The source of magmas, the temporal relationship between the magmatism, hydrothermal alteration, and mineralization at the Yongping deposit remain unclear. This paper reports ion microprobe (SHRIMP) zircon U-Pb ages for the intrusive stocks in the deposit, hydrothermal muscovite and biotite <sup>40</sup>Ar-<sup>39</sup>Ar ages, and molybdenite Re-Os ages. The Sr-Nd-Hf isotopes are also analyzed and presented. Their implications in the source of magmas and the origin of the deposit, as well as in regional metallogeny, are discussed.

### 2. Geology of the Yongping Cu–Mo deposit

The basement rocks in the Yongping deposit and vicinities are the Pre-Cambrian Zhoutan Group, which

consists of metamorphosed shallow marine siltstone interbedded with intermediate-acidic volcanic rocks. The Middle Carboniferous Yejiawan Group is composed of littoral clastic sedimentary rocks and carbonate rocks interbedded with pyroclastic rocks; the Lower Permian Maokou Group is composed of limestone and marble, and silty shale and siltstone (Fig. 2).

The inverted Houjia-Gaoshan anticline and a number of axial faults occur at its eastern limb (such as F1 and F2) in the Yongping district (Liu & Huang, 1991). The axial faults and the anticline axis strike NNW in the north of the district and in NNE in the south of the district. In the middle of the district are several faults that strike N–S. The overall structural framework protrudes to the east (Fig. 2). The axial plane of the Houjia-Gaoshan anticline shows a dip angle of 20–30°. On the east limb of the Houjia-Gaoshan anticline is a second-order inverted syncline, named the Daziping syncline. Its exposed structural nose is 1200 m long, and it strikes N–S at a dip of 30–40°.

The location of orebodies is controlled by both F1 and F2 thrust faults. The NS-trending F1 Fault dips 15–40° east. The Zhoutang migmatite was thrusted westward over the Carboniferous-Permian stratum along the F1 Fault. The orebodies in the deposit occur along the F1 and F2 Faults. The F2 fault is parallel to the F1 fault and exists between the Zhoutang and Yejiawan Groups. It is 5000 m long and dips 60–70° east. The Jurassic quartz porphyry was emplaced along the F2 Fault.

Igneous activities in the district include Silurian granitic porphyry and Jurassic dacite porphyry and quartz porphyry. They were emplaced into the Pre-Cambrian and Cambrian migmatite and the Middle-Carboniferous Yejiawan marine clastic rocks (Liu & Huang, 1991). The Jurassic dacite porphyry with porphyritic texture and/or quartz eye structure was related to the Cu mineralization. Phenocrysts are mainly euhedral plagioclase, platy biotite, and quartz, with minor apatite. The quartz phenocrysts are corroded to produce quartz-eye texture. The groundmass is composed of quartz, K-feldspar, plagioclase, biotite, calcite, fluorite, and magnetite. The Jurassic quartz porphyry with quartz unidirectional solidification textures (USTs) was related to Mo mineralization. Phenocrysts are mainly composed of feldspar and quartz. The groundmass is composed of quartz, feldspar, and fluorite. The volatile-rich minerals, such as fluorite, occur along the boundary between the quartz and K-feldspar (Li et al., 2007a).

# 3. Mineralization at the Yongping deposit

The Yongping Cu–Mo deposit is dominantly hosted by marine clastic rocks of the Carboniferous Yejiawan Group. The N–S-trending orebodies dip east at 20°–50°. They are in stratified or layered-lensoidal forms with massive and disseminated ores. Seven ore zones have been identified. Each of them is composed of multiple orebodies. The II, III, and IV zones are dominated by chalcopyrite; the V and VI zones by pyrite; and the VII zone by Cu-bearing earth and accumulates produced by secondary surface leaching. Lead- and zinc-bearing orebodies occur in the II, III, IV ore zones in the north. The thickness of the orebodies gradually changes with dip direction. The II zone is the largest in the deposit (Huang et al., 2001) (Fig. 2). The ore minerals are chalcopyrite, pyrite, galena, sphalerite, molybdenite, scheelite and magnetite. The gangue minerals are quartz, fluorite, calcite, garnet, anhydrite and barite.

The dacite and quartz porphyries show different ore and alteration mineral assemblages. Illitation, flurotization, biotitization, and chloritization are present in the dacite porphyry. The alteration associated with the

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dacite porphyry occurs along the faults. The faults contain quartz veins with pyrite. Close to the contact between the dacite porphyry and the quartz porphyry, biotitization becomes dominant in the dacite porphyry. The majority of igneous biotite are altered to chlorite and illite, and rutile seggregation along biotite cleavages is observed. The hydrothermal biotite occurs as tiny scales. Feldspar phenocrysts within the igneous rocks are altered to sericite, but their crystal shape is preserved.

The alteration of the quartz porphyry includes illitation, muscovitation, and fluorization; they occur mostly along faults and are pervasive. The feldspars are completely altered to illite and/or muscovite. Some illite recrystallized to become muscovite with diameter of 0.1–5 mm. Muscovite kinking and muscovite fish are common, indicative of a syn-tectonic alteration and mineralization. Muscovite is closely associated with molybdenite in the deposit. Molybdenite-bearing quartz veins and molybdenite veins, as well as quartz+tourmaline veins are documented in the quartz porphyry (Fig. 3). Some molybdenite-bearing quartz veins truncate the quartz porphyry and the dacite porphyry.

The porphyries are associated with skarn along their contact zones with the Yejiawan Group carbonate rocks. The skarn minerals include diopside, tremolite, actinolite, and epidote. The skarn is closely associated with the mineralization. Where garnet co-exists with diopside, tremolite, actinolite, and epidote, chalcopyrite and pyrite are present along with Pb–Zn mineralization. Where the skarn is composed of garnet only, there exist veinlets associated with chalcopyrite and pyrite. Chloritization, sericitization, and silicification overprint on the skarn, followed by carbonate, anhydrite, and fluorite precipitation. Quartz, chalcopyrite, and sphalerite veins are also observed as very late products.

Hydrothermal breccia composed of country rocks is present along the contact between the intrusive rocks and host rocks. They are cemented by fluorite, calcite, kaolinite.

#### 4. Sampling and analytical procedures

Representative samples of the dacite and quartz porphyries, as well as molybdenite-bearing quartz veins were collected from the deposit. The samples were crushed and milled and heavy mineral concentrates were magnetically separated. Inclusion-free zircons,



**Fig. 3** Photomicrographs of the hydrothermally altered igneous rocks of the Yongping Cu–Mo deposit. (a) Muscovite in quartz porphyry; the muscovite shows kinking. (b) Feldspar phenocryst conversion to illite within dacite porphyry. (c) Fluoritization occurs along the boundary of quartz grains and K-feldspar grains in the quartz porphyry. (d) Chloritization and exsolution of rutile from biotite phenocrysts in the dacite porphyry. (e) Biotite flakes in the dacite porphyry. (f) The least altered feldspar; sericitization occurs only along its fractures in the dacite porphyry. (g) Mo mineralization in quartz porphyry with muscovitization. (h) Chalcopyrite was replaced by sphalerite in quartz vein in dacite porphyry. Bt-biotite; Chl-chlorite; Cp-chalcopyrite; Fl-fluorite; ill-illite; Kfs-K-feldspar; Q-quartz; Pl-plagioclase; Mo-molybdenite; Ms-muscovite; Sp-sphalerite; Rt-rutile.

muscovite, biotite, and molybdenite were then handpicked under the binocular microscope.

## 4.1 Descriptions of the samples collected for geochronologic analysis

#### 4.1.1 Sample YP-8

Strongly altered quartz porphyry with unidirectional solidification texture (UST). It is composed of feldspar (40%), quartz (40%), sericite (10%), and fluorite (10%) with minor zircon, topaz and apatite. Feldspar is mostly altered to muscovite at a size of 2–5 mm. Quartz shows mosaic texture. Fine fluorite occurs along quartz grain boundaries.

#### 4.1.2 Sample YP-9

Strongly altered granitic porphyry with UST, and mineralized. It is composed of K-feldspar (40%; with 2V(-) = 40), quartz (40%), calcite (10%), as well as sulfide (5%;, e.g. pyrite). Minor amount of molybdenite (1%), zircon, cassiterite and rutile (1%), and fluorite (2%) found within quartz grains.

#### 4.1.3 Sample YP06-16

Weakly altered dacite porphyry with porphyritic texture. Phenocrysts (30%) are mainly euhedral plagioclase (15%), platy biotite (6%), hornblende (2%), and quartz (6%), with minor apatite (<1%). Most plagioclase phenocrysts are completely altered to sericite, and hornblende is altered to chlorite and rutile. The quartz phenocrysts are corroded to produce quartz-eye texture. The groundmass (70%) is composed of quartz, K-feldspar, plagioclase, biotite, calcite, and fluorite. K-feldspar and plagioclase are considerably altered to sericite and other clay minerals.

#### 4.1.4 Sample YP06-17

Weakly altered dacite porphyry with porphyritic texture. Phenocrysts (20%) consist of euhedral plagioclase (14%), platy biotite (4%), and quartz (2%). The plagioclase phenocrysts are partially altered to sericite, and the quartz is rounded with corroded structure. The groundmass consists of anhedral quartz (40%), euhedral-subhedral plagioclase (24%), anhedral K-feldspar (8%), and biotite (8%), with trace apatite (<1%).

#### 4.1.5 Sample YP06-18

Least altered quartz porphyry with quartz UST structure (Kirwin, 2005, 2006). It is mainly composed of feldspar including large phenocrysts of K-feldspar (with  $2V(-) = 30-60^{\circ}$ , 4–5 mm in diameter), quartz, calcite, and fluorite with trace pyrite and chalcopyrite. Quartz shows mosaic texture. Feldspar crystals are partially altered to clay minerals such as sericite. Fluorite occurs along the boundaries of quartz.

#### 4.1.6 Sample YP06-4

Weakly altered quartz porphyry with quartz UST structure. It is mainly composed of K-feldspar, plagioclase, and quartz. Plagioclase phenocrysts are altered to sericite clay minerals. Opaque minerals include pyrite, radial rutile, molybdenite, as well as fluorite.

#### 4.1.7 Samples YP-10 and YP-2

Collected from two quartz+molybdenite veins within the quartz porphyry.

#### 4.2 Whole-rock geochemical analysis

Two criteria were used for the sample selection from outcrops within the ore-hosting porphyries: (i) homogeneous samples and (ii) samples with the least effects of hydrothermal alteration based on petrographic observations. There were 9 samples collected for whole-rock geochemical analysis. Sample preparation was carried out by jaw-crushing and grinding in a ring mill. Major elements were analyzed by X-ray fluorescence spectroscopy (XRF). Trace elements were analyzed by inductively coupled plasma-mass spectrometry (ICP-MS) and the FeO was analyzed with titration methods, F with ionselective electrode (ISE) method, and S with infrared method. All the wholerock analyses were performed at the Activation Laboratories Ltd. in Canada. Details of the chemical procedures are accessible at http://www.actlabs.com.

#### 4.3 Whole-rock Rb-Sr and Sm-Nd isotopes

Rb–Sr and Sm–Nd isotope compositions of the wholerock samples selected from the ore-hosting porphyries were determined by using a GV IsoProble-T multicollector thermal ionization mass spectrometer at the Geological Institute of Nuclear Industry in Beijing, China, following the method of Zhang *et al.* (1994). <sup>87</sup>Sr/<sup>86</sup>Sr and <sup>143</sup>Nd/<sup>144</sup>Nd ratios are reported as normalized to <sup>86</sup>Sr/<sup>88</sup>Sr of 0.1194 for Sr and to <sup>146</sup>Nd/<sup>144</sup>Nd of 0.7219 for Nd. Measurements for the Johnson Mattey Nd<sub>2</sub>O<sub>3</sub> standard and for NBS-987 Sr standard yielded results of <sup>143</sup>Nd/<sup>144</sup>Nd ratios of 0.511126  $\pm$  9 (2 $\sigma$ , *n* = 27),

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and <sup>87</sup>Sr/<sup>86</sup>Sr ratios of 0.710228  $\pm$  14 (2 $\sigma$ , *n* = 30), respectively. Total analytical blanks were 5 × 10<sup>-11</sup>g for Sm and Nd, and (2–5) × 10<sup>-10</sup>g for Rb and Sr.

Initial <sup>86</sup>Sr/<sup>87</sup>Sr ratios (I<sub>Sr</sub> or (<sup>86</sup>Sr/<sup>87</sup>Sr)<sub>o</sub>) were calculated at time t with  ${}^{87}$ Rb decay constant of  $1.42 \times 10^{-11}$ (Steiger & Jäger, 1977). Initial <sup>143</sup>Nd/<sup>144</sup>Nd ratios with reference to chondritic reservoir (CHUR) at the time t of rock-formation. The  $^{147}$ Sm decay constant of 6.54 imes10<sup>-12</sup> (Lugmair & Marti, 1978) and the present-day chondritic ratios of  ${}^{143}Nd/{}^{144}Nd = 0.512638$  (Goldstein et al., 1984) and <sup>147</sup>Sm/<sup>144</sup>Nd = 0.1967 (Peucat et al., 1989) were adopted in this analysis.  $\varepsilon_{Nd}(t)$  is the measure parameter of the difference between the <sup>143</sup>Nd/<sup>144</sup>Nd ratios of a sample at time of its formation and a reference chondritic reservoir. Single-stage model ages  $(T_{DM1})$  were calculated relative to the depleted mantle (147Sm/144Nd = 0.2137, 143Nd/144Nd = 0.51315, Peucat et al., 1989). Two-stage model ages (T<sub>DM2</sub>) were also calculated assuming a <sup>147</sup>Sm/<sup>144</sup>Nd ratio of 0.118 for the continental crust (Jahn & Condie, 1995).

### 4.4 SHRIMP U-Pb dating

Zircon grains were selected from four samples (YP-9, YP06-16, YP06-17, and YP06-18) of the granite porphyry, dacite porphyry and quartz porphyry in the Yongping deposit. The selected zircons and the G1939 reference zircons were mounted in epoxy before sectioned approximately in half and polished. Reflected, transmitted optic photomicrographs, BSE images, and cathodoluminescence (CL) images were obtained for all zircon samples. The CL images were used to examine internal structure of zircons and to ensure that the 20  $\mu$ m SHRIMP spot was within a single component of a zircon.

The U-Pb isotopic analysis was performed using SHRIMP-II at the SHRIMP Center at the Institute of Geology of the Chinese Academy of Geological Sciences in Beijing, following the procedure described by Jian *et al.* (2003). For each zircon, its  $Zr_2O^+$ , <sup>204</sup>Pb<sup>+</sup> background, <sup>206</sup>Pb<sup>+</sup>, <sup>207</sup>Pb<sup>+</sup>, <sup>208</sup>Pb<sup>+</sup>, U<sup>+</sup>, Th<sup>+</sup>, ThO<sup>+</sup>, and UO<sup>+</sup> were measured on a single electron multiplier by cyclic magnetic field stepping. The mean ion counts were measured and recorded every five scans. A primary ion beam of ~4.5 nA, 10 kV O<sup>2-</sup>, and ~25–30 µm spot diameter was used. Inter-element fraction in the ion emission of zircon was corrected using the RSES reference standard TEM. The software of Ludwig (SQUID 1.0) and accompanying ISOPLOT were used for data processing and plotting (Ludwig, 2001). Ages

were calculated by using the constants recommended by IUGS (Steiger & Jäger, 1977). Unless otherwise stated, analytical uncertainties given in the table and shown in the plots have one standard deviation (1 $\sigma$ ). The ages from polled data are weighted means, with uncertainties given at two standard deviations.

All the analyses were conducted in static mode. Inter-element fractionation was monitored analyzing fragments of 91500, a large concordant zircon crystal with a known age (SHRIMP U–Pb) of 1061 Ma. This reference zircon was analyzed once for every three unknown samples. The reported ages for zircon grains are based on <sup>206</sup>Pb/<sup>238</sup>U ratios due to high errors of the <sup>207</sup>Pb/ <sup>235</sup>U and <sup>207</sup>Pb/<sup>206</sup>Pb ratios. This is due primarily to the low intensity of <sup>207</sup>Pb signal from these young, low-U grains. The <sup>206</sup>Pb/<sup>238</sup>U ratios are corrected for common Pb by the measured <sup>206</sup>Pb/<sup>204</sup>Pb.

For each sample, the  ${}^{206}\text{Pb}/{}^{238}\text{U}$  ages are plotted with  $2\sigma$  error bars that reflect only the error from determined  ${}^{206}\text{Pb}/{}^{238}\text{U}$  and  ${}^{206}\text{Pb}/{}^{204}\text{Pb}$ . For the age of each sample, additional uncertainty from the calibration, decay constant, common lead composition, and variations in the measured  ${}^{206}\text{Pb}/{}^{238}\text{U}$  and  ${}^{206}\text{Pb}/{}^{207}\text{Pb}$  from 91500 standard is considered. These systematic errors were added quadratically to the measurement error. All the reported ages and weighted mean ages have uncertainties at the  $2\sigma$  level. Weighted mean  ${}^{206}\text{Pb}/{}^{238}\text{U}$  ages were calculated, and the uncertainties are reported as 95% confidence limit.

### 4.5 Zircon Hf isotopes

Zircon Hf isotope analysis was carried out in-situ using a Geolas-193 laser-ablation microprobe attached to a Neptune multi-collector ICFMS, at the Institute of Mineral Resources, Chinese Academy of Geological Sciences, Beijing. Instrumental conditions and data acquisition were described by Xu et al. (2004). The laser system is equipped with a 193n UV ArF excimer laser. Typical ablation time for each analysis is about 30 s for 200 cycles of each measurement, with a 10 Hz repetition rate, and a laser power of 100 mJ/pulse. A stationary spot was used for the analysis, with a beam diameter of about 63 µm. Both He and Ar carrier gases were used to transport the ablated materials from the laser-ablation cell via a mixing chamber to the ICP torch. Isobaric interference of 176Lu on 176Hf was corrected measuring the intensity of the interference-free <sup>175</sup>Lu isotope and using the recommended <sup>176</sup>Lu/<sup>175</sup>Lu ratios of 0.02655 to calculate 176Lu/177Hf ratios. Correction for isobaric interference of 176Yb on 176Hf was performed in real time (Wu *et al.*, 2006) as developed by Iizuka and Hirata (2005). Standard zircon 91500 showed similar average <sup>176</sup>Hf/<sup>177</sup>Hf ratios of 0.282307 ± 58, but a bimodal distribution with two peaks at 0.282284 ± 22 and 0.282330 ± 29. The Hf isotope heterogeneity in 91500 brings extra uncertainties to the data, however, our analyses on 91500 in two sessions yielded only one peak at 0.282307 ± 31 (Wu *et al.*, 2006). This means that at least this chip of 91500 analyzed is homogeneous in Hf isotopes. All the Lu–Hf isotope results are reported with a 2 $\sigma$  error (Table 4).

Initial  ${}^{176}\text{Hf}/{}^{177}\text{Hf}$  ratios ( $\epsilon_{\text{Hf}}(t)$ ) were calculated with reference to chondritic reservoir (CHUR) at the time t of zircon growth from magmas, the single grain zircon SHRIMP U-Pb age were used to represent the zircon growth time t from magmas for each point. The <sup>176</sup>Lu decay constant of  $1.865 \times 10^{-11}$  yr<sup>-1</sup> (Nowell *et al.*, 1998) and the present-day chondritic ratios of  ${}^{176}\text{Hf}/{}^{177}\text{Hf} =$ 0.282772 and  ${}^{176}Lu/{}^{177}Hf = 0.0332$  (Blichert-Toft & Albarede, 1997) were adopted in this analysis. Singlestage model ages (T<sub>DM1</sub>) were calculated relative to the depleted mantle with a present-day <sup>176</sup>Hf/<sup>177</sup>Hf ratio of 0.28325, similar to that of the average MORB (Nowell et al., 1998) and a present-day <sup>176</sup>Lu/<sup>177</sup>Hf ratio of 0.0384 (Griffin *et al.*, 2000). Errors in both  $\varepsilon_{Hf}(t)$  and  $T_{DM1}$  values were calculated on the basis of analytical errors, but much bigger and poorly constrained errors could be associated with the Hf model ages due to uncertainties in the reference model (Nowell et al., 1998).

Two-stage model ages ( $T_{DM2}$ ) were also calculated by projecting the initial <sup>176</sup>Hf/<sup>177</sup>Hf of the zircon back to the depleted mantle model growth curve, assuming a <sup>176</sup>Lu/<sup>177</sup>Hf ratio of 0.009 for the upper continental crust of Archean age. A <sup>176</sup>Lu/<sup>177</sup>Hf ratio of 0.015 was assumed for the average continental crust (Griffin *et al.*, 2000), which is consistent with the average <sup>176</sup>Lu/<sup>177</sup>Hf values of Archean sedimentary rocks in Superior Province of North America (Stevenson & Patchett, 1990; Corfu & Noble, 1992).

#### 4.6 Muscovite and biotite Ar-Ar dating

Muscovite samples selected from specimens YP06-4, and the hydrothermal biotite samples selected from specimens YP06-16, and YP06-17 were purified by magnetic separator and then cleaned by ultrasonic treatment with ethanol. The purity of the muscovite grains (at a size of 0.08–0.15 mm) is greater than 99%. The biotite and muscovite samples were irradiated for 51 hours in a nuclear reactor at the Chinese Institute of Atomic Energy in Beijing, and subsequently cooled for

approximately 100 days. <sup>40</sup>Ar/<sup>36</sup>Ar stepwise heating analysis was performed at the Institute of Geology, Chinese Academy of Geological Sciences using a MM-1200B Mass Spectrometer. Measured isotopic ratios were corrected for the mass discrimination, atmospheric Ar component, blanks and irradiationinduced mass interference. The decay constant used is  $\delta = 5.543 \times 10^{-10}$  year<sup>-1</sup> (Steiger & Jäger, 1977). All <sup>37</sup>Ar were corrected for radiogenic decay (half-life 35.1 days). Uncertainty in each apparent age is given at 2 $\sigma$ . The monitor used in this analysis was an internal standard: Fangshan biotite (ZBH-25) of 132.7 Ma and its potassium content is 7.6%. The *J*-values of the monitors are given in Tables 5–7. Detailed analytical techniques are described by Chen *et al.* (2002).

#### 4.7 Molybdenite Re-Os dating

Molybdenite samples were separated from molybdenite-bearing quartz veins for the Re-Os analyses following the method described by Mathur et al. (2002) and Barra et al. (2003). Approximately 0.05-0.10 g of handpicked molybdenite was loaded in a carius tube with 8 mL of reverse aqua regia (1HCl:3HNO<sub>3</sub>). While the reagents, samples, and spikes were frozen, the Carius tube was sealed and left to thaw at room temperature. The tube was placed in an oven and heated to 240°C overnight. The solution was treated in a two-stage distillation process for osmium separation (Nagler & Frei, 1997). Osmium was further purified using a microdistillation technique (Birck et al., 1997) and loaded on platinum filaments with Ba(OH)<sub>2</sub> to enhance ionization. After osmium separation, the remaining acid solution was dried and later dissolved in 0.1 HNO<sub>3</sub>. Rhenium was extracted and purified through a two-stage column using resin and loaded on platinum filaments with BaSO<sub>4</sub>.

Samples were analyzed with ICP-MS at Chinese Academy of Geological Sciences, Beijing. Molybdenite ages were calculated using a <sup>187</sup>Re decay constant of  $1.666 \times 10^{-11}$  per year (Smoliar *et al.*, 1996). Ages were reported with 0.5% error, which is considered a conservative estimate and reflects all sources of errors.

#### 5 Results

#### 5.1 Major and trace elements

Results of major and trace element contents of the ore-hosting quartz and dacite porphyries are listed in Table 1. Some of the analyzed samples contain variable loss on ignition (LOI) values reflecting variable

Samples		Q	uartz porph	yry			Dacite p	oorphyry	
	YP-4	YP-7	YP-8	YP06-18	YP06-26	YP06-16	YP06-17	YP06-20	YP06-21
Wt%									
$SiO_2$	74.76	71.17	74.45	76.21	73.32	63.15	64.53	66.15	63.76
TiO <sub>2</sub>	0.19	0.38	0.06	0.34	0.41	0.57	0.66	0.52	0.55
$Al_2O_3$	13.23	13.57	14.91	10.79	13.15	14.03	15.81	15.53	15.21
Fe <sub>2</sub> O <sub>3</sub>	1.38	1.08	0.28	0.01	0.37	0.81	2.25	0.77	1.58
FeO	1.76	1.84	0.71	0.86	1.13	2.56	1.57	2.38	1.75
MnO	0.003	0.008	0.004	0.015	0.016	0.03	0.02	0.02	0.05
MgO	0.72	1.31	0.95	0.61	1.41	2.35	2.03	1.45	1.75
CaO	0.02	0.98	0.22	1.76	1.07	2.90	3.02	2.21	2.63
Na <sub>2</sub> O	0.09	0.10	0.10	0.13	0.11	0.50	2.74	0.93	1.86
K <sub>2</sub> O	4.51	5.08	5.34	5.74	5.64	6.75	4.29	5.57	5.33
$P_2O_5$	0.02	0.08	0.13	0.10	0.11	0.16	0.20	0.20	0.19
F	0.19	1.84	0.20	0.15	0.17	0.13	0.12	0.13	0.10
LOI	3.43	4.53	3.41	2.71	3.25	5.91	2.71	4.47	4.26
S	2.55	2.19	0.61		_	_	_		_
Total (%)	100.2	100.3	100.3	99.4	100.1	100	100	100.1	99.1
ppm									
Ba	1676	890	865	579	488	2130	912	777	1490
Rb	130	180	170	200	190	207	207	212	187
Sr	14	61	38	90	63	251	251	190	299
Y	6	19	34	9.1	12.5	14.2	14.2	10.1	13.3
Zr	105	168	59	87	114	142	142	149	164
Nb		—	—	9.7	11.9	11.2	11.8	12.2	12.5
Th	6.8	—	7.4	5.45	7.85	7.02	7.54	7.96	8.82
Pb	26	5	16	5	7	5	5	7	9
Ga	_	—	_	14	16	19	21	21	20
Zn	8	12	34	30	30	30	30	30	30
Cu	637	85	27	40	20	220	110	240	50
V	31	67	13	53	69	83	83	67	70
Cr	2.1	23.2	0.5	20	20	30	20	20	20
Hf	2.3	3.8	1.2	2.6	3.2	3.8	4.3	3.9	4.4
Cs	1.1	3.1	3.9	5.2	7.1	7.5	7.7	12.3	9.8
Ta	0.5	0.3	1.1	0.82	1.07	0.84	0.87	0.95	0.99
Co	11.9	10.3	2.8	2	4	8	7	4	5
U	1.2	2	4.8	4.27	3.7	2.38	2.78	2.22	3.16
W	39	50	13	27.4	57.4	30	10.8	27.3	12.3
Mo	3	4	2	19	75	27	46	4	2
La	26.2	31.1	12.5	18.6	24.6	32.6	33.4	38	36.7
Ce	46	61	27	34.9	46.1	62	63.5	68.3	68.4
TT LI	16		12	3.0 12.9	5.00 10.1	6.9	0.90	7.35 26 E	7.44
Sm	10	20 1 6	15	13.0	2.24	4 72	23.8	20.3	20.7
5111	2.36	4.0	0.66	2.40	0.00	4.72	4.4	4.33	4.03
Eu	0.04	1.00	0.00	0.07	0.90	1.20	2.45	1.27	2.51
Gu Th				2.02	2.39	0.5	0.49	0.42	0.54
Du	0.5	0.0	0.7	1.66	2.45	0.5	2.44	1.94	2.75
Dy Ho		—	_	0.3	2.30	2.70	2.44	0.32	2.75
Fr	_	_	_	0.3	1 08	1 22	1 08	0.32	1 2
Tm				0.01	0.15	0.18	0.14	0.01	0.16
Vh	0.53	1.84	2 69	0.11	0.13	1 12	0.14	0.11	1 01
10 I 11	0.00	0.27	2.09 0.40	0.07	0.97	1.13 0.16	0.02	0.7	1.01
Lu	0.00	0.27	0.40	0.090	0.15	0.10	0.12	0.11	0.14

Table 1 Geochemical data of the igneous rocks in the Yongping Cu-Mo deposit

Note: "-" not analyzed.



Fig. 4 Chondrite-normalized REE diagram for both dacite and quartz porphyries, Chondritic normalization is according to Sun and McDonough (1989).

 $H_2O+CO_2$  contents, possibly due to different degree of alteration. In general, the high-field-strength elements (HFSE), rare earth elements (REE), and transition elements are immobile during the hydrothermal alteration (Hawkesworth *et al.*, 1997). However, the Na<sub>2</sub>O and K<sub>2</sub>O contents show a decrease and increase with increasing LOI, respectively, implying their original contents were modified by alteration. Thus, only immobile elements are used in the following discussion.

The total REE concentrations range from 118.64 ppm to 168.16 ppm in the dacite porphyries and from 56.41 ppm to 147.49 ppm in the quartz porphyries. Both porphyries are rich in light REE and poor in heavy REE. The chondrite-normalized REE diagram (Fig. 4) shows a steep curve pattern without Eu depletion. As shown in the primitive mantle-normalized trace element spider diagram (Fig. 5), both types of the porphyries are rich in large ion lithophile elements (Ba, Rb, and K), and depleted in high field strength elements (Th, Nd, Ta, and Ti) and Sr and P. This may suggests subduction-related arc magmatism. The depleted Sr and P may suggest a magma fractional crystallization process.

#### 5.2 Rb–Sr and Sm–Nd isotopes

The Rb–Sr and Sm–Nd isotopic compositions of the dacite and quartz porphyries are listed in Table 2. Table 2 shows the initial Sr isotopic composition for dacite porphyry and quartz porphyry ranging from 0.707 to 0.710, and from 0.704 to 0.712, respectively. The





Fig. 5 Primitive mantle-normalized trace element spider diagram for both dacite and quartz porphyries, Normalizing values are according to Sun and McDonough (1989).

 $\epsilon_{Nd}(t)$  values range from –9.18 to –7.64 and from –10.95 to –6.94, respectively. The Nd model ages of the dacite and quartz porphyries relative to a depleted mantle (T<sub>DM1</sub>) range from 1.26 to 1.38 Ga (with a mean of 1.32 Ga), and from 1.24 to 1.44 Ga (with a mean of 1.30 Ga), respectively. Two-stage Nd model ages of the dacite and quartz porphyries relative to a depleted mantle (T<sub>DM2</sub>) range from 1.57 to 1.69 Ga (with a mean of 1.63 Ga) and from 1.51 to 1.84 Ga (with a mean of 1.62 Ga), respectively.

#### 5.3 Zircon SHRIMP U-Pb ages

Zircons used for SHRIMP U-Pb analysis were examined on the basis of petrographic microscopy and cathodoluminescence (CL) microscopy with a backscattered electron imaging system. Target spots were selected based on CL images (Fig. 6). Zircons are colorless and range from oval, elongate, to long elongate in shape, generally showing 101 type. Most of the zircons were devoid of inclusions, although a few contain minute inclusions.

#### 5.3.1 Sample YP-9

A total of 14 spots were analyzed on 13 zircon grains selected from sample YP-9 which was collected from strongly altered quartz porphyry. All the analyzed zircon grains have relatively uniform texture typical of metamorphic origin. Sample YP-9 yields a  $^{206}$ Pb/ $^{238}$ U weighted average age of 435.2 ± 7.3 Ma (n = 14 zircons, mean square of weighted deviation (MSWD) = 0.45) (Fig. 7a, Table 3).

Table 2 Rb-Sr and Sm-Nd isotope compositions of porphyries in the Yongping Cu-Mo deposit

Samples	Rock Types	<sup>87</sup> Rb/ <sup>86</sup> Sr	<sup>87</sup> Sr/ <sup>86</sup> Sr	( <sup>87</sup> Sr/ <sup>86</sup> Sr) <sub>i</sub>	<sup>147</sup> Sm/ <sup>144</sup> Nd	<sup>143</sup> Nd/ <sup>144</sup> Nd	$^{143}Nd/_{^{144}}Nd_{(t)}$	$Nd_{CHUR(t)} \\$	$\epsilon_{Nd(t)}$	T <sub>DM1</sub> (Ma)	T <sub>DM2</sub> (Ma)
YP06-16	Dacite porphyry	2.5932	0.71608	0.71029	0.0893	0.51206	0.51197	0.51244	-9.18	1340	1693
YP06-17	Dacite porphyry	1.1196	0.71093	0.70844	0.0883	0.51211	0.51202	0.51244	-8.05	1260	1601
YP06-20	Dacite porphyry	3.5494	0.71578	0.70786	0.0973	0.51210	0.51200	0.51244	-8.60	1382	1646
YP06-21	Dacite porphyry	2.0237	0.71363	0.70911	0.093	0.51214	0.51204	0.51244	-7.64	1276	1568
YP-4	Quartz porphyry	30.0169	0.78149	0.71236	0.0883	0.51196	0.51187	0.51243	-10.95	1444	1840
YP06-4	Quartz porphyry	7.1347	0.72241	0.70598	0.09	0.51212	0.51202	0.51243	-7.92	1271	1595
YP06-26	Quartz porphyry	9.8552	0.72671	0.70401	0.0934	0.51217	0.51207	0.51243	-7.02	1244	1521
YP06-18	Quartz porphyry	6.6835	0.72389	0.70849	0.0937	0.51217	0.51207	0.51243	-6.94	1242	1516
YP-9	Granite porphyry	4.4827	0.73155	0.70378	0.0976	0.51168	0.51140	0.51208	-13.23	1929	2246

Notes:

(1) ( ${}^{87}Sr/{}^{86}Sr$ )<sub>i</sub> refer to the initial Sr isotopic ratios at time t;  ${}^{87}Rb$  decay constant  $\lambda = 1.42 \times 10^{-11}$  (Steiger & Jäger, 1977).

(2) <sup>143</sup>Nd/<sup>144</sup>Nd(t) refer to the initial Nd isotopic ratios at time t; <sup>147</sup>Sm decay constant  $\lambda = 6.54 \times 10^{-12}$  (Lugmair & Marti, 1978).

(3)  $Nd_{CHUR(t)}$  is the<sup>143</sup>Nd/<sup>144</sup>Nd isotope ratios of CHUR (chondritic reservoir) at time t; <sup>143</sup>Nd/<sup>144</sup>Nd(CHUR, Today) = 0.512638 (Goldstein *et al.*, 1984), <sup>147</sup>Sm/<sup>144</sup>Nd(CHUR, Today) = 0.1967 (Peucat *et al.*, 1989), (4)  $T_{DM1}$  (Ma) refer to the single stage model age relative to depleted mantle (DM). DM: <sup>147</sup>Sm/<sup>144</sup>Nd = 0.2137, <sup>143</sup>Nd/<sup>144</sup>Nd = 0.51315 (Peucat *et al.*, 1989)

(5) T<sub>DM2</sub> (Ma) refer to the two-stage model age relative to depleted mantle. Continental crustal: <sup>147</sup>Sm/<sup>144</sup>Nd = 0.118 (Jahn & Condie, 1995)

(6) t = 157 Ma is used for dacite porphyry (YP06-16, YP06-17, YP06-20, YP06-21), t = 162Ma is used for quartz porphyry (YP-4, YP06-4, YP06-26, YP06-18), and t = 435 Ma is used for granite porphyry (YP-9).

#### 5.3.2 Sample YP06-16

A total of 15 spots were analyzed on 14 zircon grains selected from sample YP06-16 from the dacite porphyry. This sample yields a  ${}^{206}\text{Pb}/{}^{238}\text{U}$  weighted average age of 157.5 ± 2.9 Ma (n = 14 zircons, MSWD = 0.32). An inherited zircon crystal yields a Silurian age (~430 Ma) (Fig. 7b, Table 3).

#### 5.3.3 Sample YP06-17

A total of 14 spots were analyzed on 14 zircon grains selected from sample YP06-17 from the dacite porphyry. This sample yields a weighted average age of 156.0  $\pm$  3.1 Ma (n = 12 zircons, MSWD = 0.19), although an inherited grain shows a Neo-Proterozoic age (~766 Ma). Another zircon grain gives a Late Cambrian age (~498 Ma) (Fig. 7c, Table 3).

#### 5.3.4 Sample YP06-18

A total of 15 spots were analyzed on 15 zircon grains selected from sample YP06-18 which was collected from the quartz porphyry. This sample yields a weighted average age of  $162.1 \pm 3.4$  Ma (n = 10 zircons, MSWD = 0.35). Three grains (No. 2, No.9, and No.13) yield a weighted average age of  $432 \pm 16$  Ma. A zircon with narrow growth zoning has a Late Triassic age (~208.6 Ma) and another one with narrow growth zoning shows an Early Jurassic age (Fig. 7d, Table 3).

### 5.4 Zircon Hf isotopes

A total of 65 spots of in-situ Hf isotope analysis were analyzed on zircon grains selected from samples dacite porphyry (YP06-16, YP06-17), and quartz porphyry (YP06-18). The results (Table 4) demonstrate negative initial  $\epsilon_{Hf}(t)$  values. The  $\epsilon_{Hf}(t)$  values of sample YP06-16 range from -17 to -5.6, mostly between -10 and -8, except for several spots that have negative initial  $\epsilon_{Hf}(t)$  values ranging from -19 to -13. The  $\epsilon_{Hf}(t)$  values of sample YP06-17 has a wider range between -20.0 and -4.9. The  $\epsilon_{Hf}(t)$  values of sample YP06-18 also has a wider range from -19.8 to -6.3 with two peaks at -18 to -16 and -10 to -8 (Table 4). The negative initial  $\epsilon_{Hf}(t)$  values could imply a crustal magma source, which is in good agreement with the Nd–Sr isotope compositions of the samples.

#### 5.5 Biotite and muscovite Ar-Ar ages

The <sup>40</sup>Ar–<sup>39</sup>Ar ages and  $2\sigma$  uncertainties are plotted against the cumulative released <sup>39</sup>Ar fraction to establish step heating spectra (Fig. 8). The argon isotope ratios are presented graphically on classic isotope correlation plots using program ISOPLOT of Ludwig (2001). The <sup>40</sup>Ar–<sup>39</sup>Ar data of biotite and muscovite are summarized in Tables 5–7, which incorporate the apparent age spectra and the atmospheric argon and Ca:K ratios for each step heating. The muscovite selected from quartz porphyry YP06-4 has a plateau



**Fig. 6** Cathodoluminescence images of zircons selected from samples granite porphyry (YP-9), dacite porphyry (YP06-116, YP06-17) and quartz porphyry (YP06-18). SHRIMP target spots are marked by circles and are approximately 25 μm in diameter. Cathodoluminescence images were acquired at the Institute of Mineral Resources, Chinese Academy of Geological Sciences, using a JOEL JXA 8800R, at 20 kV, 10 nA.

age of 162.1  $\pm$  1.4 Ma, calculated from steps 5–11, representing 62.1% of the total argon release (Fig. 8a). The isochron age of 161  $\pm$  4 Ma (MSWD = 0.106, initial  ${}^{40}$ Ar/ ${}^{36}$ Ar = 318) is concordant with the plateau age (Fig. 8a).

The biotite selected from dacite porphyry (YP06-16, YP06-17) have plateau ages of  $164 \pm 1.3$  Ma and  $163.8 \pm 1.3$  Ma, both of which were calculated from steps 3–10, representing 95% and 95.1% of the total argon release, respectively. Both samples show similar isochron ages: sample YP06-16 of  $164.8 \pm 3.4$  Ma (MSWD = 0.62) and

sample YP06-17 of  $164 \pm 3 \text{ Ma}$  (MSWD = 0.23) (Fig. 8b,c).

#### 5.6 Molybdenite Re-Os results

Two Re–Os molybdenite analyses are reported in Table 8. The concentrations of total Re and <sup>187</sup>Os range in 19.25–79 ppm and in 32–130 ppb, respectively. The samples were analyzed only once because the ages of both samples (155.7  $\pm$  3.6 Ma and 156  $\pm$  2.8 Ma (2 $\sigma$ )) are nearly identical within the allowed error limit.



Fig. 7 Concordia plot showing all SHRIMP data for zircons selected from granite porphyry (YP-9), dacite porphyry (YP06-16, YP06-17) and quartz porphyry (YP06-18).

#### 6. Discussions

## 6.1 Zircon morphology and Th–U geochemistry

#### 6.1.1 Zircon morphology

Cathodoluminescence images of the analyzed samples reveal that all the zircons from the Yongping deposit contain inherited cores surrounded by either magmatic or recrystallized rims (Fig. 6). The ages recorded in the inherited cores can be treated similarly to detrital crystals and used to fingerprint their source (Elburg, 1996). Zircon recrystallized or overgrown during metamorphism could show high or low luminescence in CL images due to different redistribution of REE and U. Hydrothermal zircons may show spongy texture due to the presence of fluid-inclusions (Hoskin, 1998). The apparent systematic distinction between igneous zircons and metamorphic zircons is recongnized in the Th/U ratios, which is very low for the latter (<0.07; Rubatto, 2002). Igneous zircons generally have Th/U ratio equal to or greater than 0.5 (Hoskin & Schaltegger, 2003). Igneous zircons with low Th/U ratios (less than 0.5) may have crystallized in the presence of aqueous fluids (Bacon & Lowenstern, 2005).

		mn 2 - 0	21 T 1 T 1				~~ 9	, 1244										
Spots	Rock types	$\%^{206}\text{Pb}_{c}$	U ppm	Th ppm	$^{232}$ Th /	<sup>206</sup> Pb*	$^{207}\text{Pb*}/$	% <del>+</del>	<sup>207</sup> Pb*/	% +	<sup>206</sup> Pb*/	% +	errcorr	<sup>206</sup> Pb/		<sup>207</sup> Pb/		%
			-		<sup>238</sup> U	mqq	<sup>206</sup> Pb*		<sup>235</sup> U		<sup>238</sup> U			<sup>238</sup> U A	∕ge	<sup>206</sup> Pb .	Age	Discordant
YP-9-1.1	Granite	0.25	1171	28	0.02	72.8	0.05428	1.7	0.540	3.7	0.0722	3.3	.894	449	$\pm 14$	383	+37	-17
YP-9-1.2	porphyry	0.12	892	20	0.02	55.1	0.05667	1.7	0.561	3.7	0.0718	3.3	.895	447	$\pm 14$	479	±37	7
YP-9-2.1	•	0.15	1142	27	0.02	67.1	0.05606	1.6	0.527	3.7	0.0682	3.3	.901	426	$\pm 14$	455	±36	9
YP-9-3.1		0.05	1331	24	0.02	76.9	0.05730	1.5	0.531	3.6	0.0672	3.3	.915	419	+13	503	+32	17
YP-9-4.1		0.33	843	48	0.06	50.4	0.0569	2.2	0.545	4.0	0.0694	3.3	.840	432	$\pm 14$	490	$\pm 48$	12
YP-9-5.1		0.33	839	19	0.02	52.0	0.0561	2.0	0.556	3.9	0.0718	3.3	.853	447	$\pm 14$	458	+45	7
YP-9-6.1		0.20	1164	28	0.02	71.2	0.05620	1.7	0.551	3.7	0.0711	3.3	.893	443	$\pm 14$	461	±37	4
YP-9-7.1		0.14	1213	28	0.02	71.7	0.05697	1.4	0.539	3.6	0.0687	3.3	.924	428	$\pm 14$	491	+30	13
YP-9-8.1		0.25	897	18	0.02	54.2	0.0556	2.0	0.538	3.9	0.0702	3.3	.855	437	$\pm 14$	435	+45	-1
YP-9-8.2		0.13	1867	12	0.01	109	0.05582	1.4	0.521	3.6	0.0677	3.3	.923	422	+13	445	+31	ß
YP-9-9.1		0.14	986	18	0.02	59.6	0.05729	1.4	0.555	3.6	0.0703	3.3	.918	438	$\pm 14$	503	+32	13
YP-9–10.1		0.24	1097	19	0.02	66.3	0.05442	1.8	0.527	3.8	0.0702	3.3	.880	438	$\pm 14$	389	$\pm 40$	-13
YP-9-11.1		0.25	1112	15	0.01	67.4	0.05495	1.8	0.533	3.8	0.0704	3.3	.882	439	$\pm 14$	410	$\pm 40$	-7
YP-9–12.1		0.40	1234	28	0.02	73.9	0.0550	1.9	0.527	3.8	0.0695	3.3	.866	433	$\pm 14$	414	+43	4-
YP06-16-1.1	Dacite	1.71	694	176	0.26	15.1	0.0485	9.0	0.167	9.6	0.02499	3.4	.356	159.1	$\pm 5.4$	128	$\pm 210$	-24
YP06-16-2.1	porphyry	0.52	518	86	0.17	11.3	0.0586	4.7	0.205	5.9	0.02536	3.5	.591	161.5	$\pm 5.5$	551	$\pm 100$	71
YP06-16-3.1	•	1.52	482	101	0.22	10.4	0.0475	7.8	0.162	8.6	0.02466	3.5	.409	157.1	$\pm 5.4$	78	$\pm 190$	-102
YP06-16-4.1		0.86	634	140	0.23	13.3	0.0535	8.0	0.179	8.7	0.02431	3.5	.398	154.9	+5.3	350	$\pm 180$	56
YP06-16-5.1		1.23	767	207	0.28	16.1	0.0484	4.9	0.1613	6.0	0.02415	3.4	.572	153.8	$\pm 5.2$	121	$\pm 120$	-28
YP06-16-6.1		0.86	782	223	0.29	16.9	0.0511	4.8	0.176	5.9	0.02493	3.4	.580	158.7	+5.3	248	$\pm 110$	36
YP06-16-7.1		0.83	713	202	0.29	15.2	0.0502	5.0	0.170	6.1	0.02462	3.4	.562	156.8	+5.3	205	$\pm 120$	24
YP06-16-8.1		1.21	903	344	0.39	19.8	0.0482	6.9	0.168	7.7	0.02522	3.5	.456	160.6	$\pm 5.6$	109	$\pm 160$	-47
YP06-16-9.1		0.73	507	93	0.19	10.7	0.0537	6.5	0.180	7.7	0.02435	4.1	.531	155.1	$\pm 6.2$	361	$\pm 150$	57
YP06-16-10.1		1.78	623	206	0.34	13.8	0.0438	12	0.153	12	0.02529	3.5	.281	161.0	$\pm 5.5$	-120	±290	234
YP06-16-11.1		1.34	575	114	0.21	12.2	0.0471	9.6	0.158	10	0.02435	3.5	.333	155.1	+5.3	56	±230	-178
YP06-16-12.1		1.88	695	253	0.38	15.1	0.0467	11	0.160	11	0.02477	3.5	.301	157.7	$\pm 5.4$	38	$\pm 260$	-318
YP06-16-13.1		0.67	377	38	0.10	22.4	0.0539	5.0	0.513	6.0	0.0689	3.4	.567	430	$\pm 14$	369	$\pm 110$	-17
YP06-16-14.1		3.08	370	76	0.21	8.43	0.0438	19	0.156	19	0.02577	3.6	.192	164.0	$\pm 5.9$	-119	$\pm 460$	238
YP06-16-14.2		1.43	465	100	0.22	9.76	0.0548	9.0	0.183	9.6	0.02413	3.5	.366	153.7	$\pm 5.3$	407	±200	62

Table 3 Zircon SHRIMP U-Pb data for the igneous rocks in the Yongping copper deposit.

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Table 3 Con	tinued																	
Spots	Rock types	$\%^{206} Pb_c$	U ppm	Th ppm	<sup>232</sup> Th/ <sup>238</sup> U	<sup>206</sup> Pb* ppm	<sup>207</sup> Pb*/ <sup>206</sup> Pb*	± %	<sup>207</sup> Pb*/ <sup>235</sup> U	+ %	<sup>206</sup> Pb*/ <sup>238</sup> U	÷ %	errcorr	<sup>206</sup> Pb/ <sup>238</sup> U Ag	e	<sup>207</sup> Pb/ <sup>206</sup> Pb /	\ge	% Discordant
YP06-17-1.1	Dacite	3.67	366	151	0.43	8.23	0.0405	24	0.141	24	0.0253	4.0	.163	161.0	±6.3	-314	$\pm 620$	151
YP06-17-2.1	porphyry	1.75	332	147	0.46	6.91	0.0618	15	0.203	15	0.0238	3.7	.237	151.8	+5.5	668	±320	12
YP06-17-3.1		0.56	424	123	0.30	8.94	0.0594	5.2	0.200	6.3	0.0244	3.5	.557	155.6	$\pm 5.4$	581	$\pm 110$	73
YP06-17-4.1		1.66	406	55	0.14	8.58	0.0455	12	0.152	12	0.0242	3.5	.290	154.2	$\pm 5.4$	-24	±280	736
YP06-17-5.1		0.41	866	255	0.30	18.7	0.0465	5.1	0.1604	6.1	0.0250	3.4	.557	159.4	+5.3	23	$\pm 120$	-584
YP06-17-6.1		0.84	451	112	0.26	9.68	0.0551	4.2	0.188	5.5	0.0247	3.5	.634	157.8	$\pm 5.4$	416	±95	62
YP06-17-7.1		1.58	591	163	0.28	12.6	0.0419	12	0.141	12	0.0244	3.5	.288	155.3	+5.3	-229	±290	168
YP06-17-8.1		2.36	389	85	0.23	8.37	0.0489	17	0.165	17	0.0245	4.1	.241	155.9	±6.3	144	±390	8
YP06-17-9.1		0.31	654	132	0.21	13.9	0.0544	3.7	0.1852	5.0	0.0247	3.4	.682	157.1	±5.3	389	+83	60
YP06-17-10.1		0.48	522	86	0.17	36.2	0.0586	3.0	0.649	4.5	0.0803	3.4	.751	498	$\pm 16$	551	+65	10
YP06-17-11.1		0.31	360	278	0.80	39.1	0.0666	2.5	1.157	4.2	0.1261	3.4	.810	766	±25	824	$\pm 51$	7
YP06-17-12.1		0.69	799	337	0.44	17.1	0.0504	3.4	0.1717	4.8	0.0247	3.4	.702	157.3	±5.3	215	+80	27
YP06-17-13.1		0.62	686	174	0.26	14.3	0.0481	4.8	0.1603	5.9	0.0241	3.4	.581	153.8	±5.2	107	$\pm 110$	-44
YP06-17-14.1		2.21	343	55	0.17	7.37	0.0581	15	0.196	15	0.0245	3.7	.248	156.1	±5.7	534	±320	71
YP06-18-1.1	Quartz	2.56	585	120	0.21	13.0	0.0487	14	0.170	14	0.02526	3.5	.248	160.8	±5.5	136	±320	-18
YP06-18-2.1	porphyry	0.11	1328	~	0.01	81.3	0.05618	1.3	0.551	3.6	0.0712	3.3	.929	443	$\pm 14$	459	±29	4
YP06-18-3.1		1.33	410	94	0.24	9.14	0.0545	11	0.193	12	0.02562	3.5	.297	163.1	+5.6	394	±250	59
YP06-18-4.1		1.39	604	128	0.22	13.3	0.0552	7.4	0.192	8.1	0.02526	3.4	.424	160.8	+5.5	422	$\pm 160$	62
YP06-18-5.1		2.03	643	258	0.41	13.8	0.0428	13	0.145	13	0.02456	3.6	.278	156.4	+5.6	-174	$\pm 310$	190
YP06-18-6.1		1.19	969	138	0.21	19.9	0.0528	6.5	0.240	7.3	0.0329	3.4	.464	208.6	17.0	321	$\pm 150$	35
YP06-18-7.1		0.83	978	416	0.44	21.3	0.0504	5.0	0.175	6.0	0.02515	3.4	.562	160.1	+5.3	215	$\pm 120$	25
YP06-18-8.1		1.44	757	190	0.26	16.8	0.0445	9.2	0.157	9.8	0.02551	3.4	.348	162.4	+5.5	-81	±220	301
YP06-18-9.1		0.28	1322	6	0.01	78.8	0.05457	1.7	0.521	3.7	0.0692	3.3	.888	432	$^{+14}$	395	+38	6-
YP06-18-10.1		0.38	378	170	0.47	8.48	0.0545	5.9	0.195	6.8	0.02599	3.5	.512	165.4	±5.7	393	$\pm 130$	58
YP06-18-11.1		0.33	1520	525	0.36	33.9	0.0494	2.9	0.1764	4.4	0.02590	3.3	.759	164.8	$\pm 5.4$	166	±67	1
YP06-18-12.1		0.18	935	308	0.34	21.2	0.0552	3.3	0.2010	4.7	0.02640	3.4	.722	167.9	±5.7	422	±73	09
YP06-18-13.1		0.25	1257	8	0.01	73.1	0.0539	1.9	0.502	3.8	0.0676	3.3	.862	422	$\pm 14$	367	$\pm 44$	-15
YP06-18-14.1		2.32	455	165	0.38	11.4	0.0483	14	0.190	14	0.0285	3.6	.253	181.1	$\pm 6.4$	115	±320	-57
YP06-18-15.1		1.28	620	275	0.46	13.6	0.0485	9.8	0.169	10	0.02527	3.5	.334	160.9	±5.5	123	±230	-30
Errors are 1-sig	ma; Pb <sub>c</sub> and Pb*	<sup>+</sup> indicate th	te common	n and radio	genic port	ions, rest	ectively.											
Error in standa	rd calibration w	as 1.09% (nu	ot included	l in above	errors but	required	when com	paring	data from	differer	it mounts)							
(1) Common P	b corrected using	g measured <sup>206</sup> ا	<sup>204</sup> .Pb. Ph./ <sup>238</sup> 11_ <sup>207</sup>	Ph / <sup>235</sup> [] ao	e-concord	ance												
(3) Common Pl	b corrected by as	ssuming 2061	Pb/ <sup>238</sup> U- <sup>208</sup>	Pb/ <sup>232</sup> Th ag	ge-concord	lance.												

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Samples	Rock types	<sup>176</sup> Hf/ <sup>177</sup> Hf	<sup>176</sup> Lu/ <sup>177</sup> Hf	t (Ga)	ε <sub>Hf(t)</sub>	<sup>176</sup> Hf/ <sup>177</sup> Hf (t)	T <sub>DM1</sub> (Ga)	T <sub>DM2</sub> (Ga)
YP-9-0-1	Granite	0.28214	0.00032	0.44700	-12.49112	0.28214	1.53395	2.21603
YP-9-0-2	porphyry	0.28212	0.00031	0.42600	-13.84187	0.28212	1.56816	2.28520
YP-9-1-1		0.28218	0.00036	0.42600	-11.79173	0.28217	1.49046	2.15633
YP-9-2-1		0.28214	0.00020	0.44900	-12.55429	0.28214	1.53494	2.22155
YP-9-3-1		0.28217	0.00018	0.41900	-11.96696	0.28217	1.48638	2.16219
YP-9-4-1		0.28214	0.00015	0.43200	-13.00868	0.28214	1.53645	2.23742
YP-9-5-1		0.28216	0.00024	0.44700	-11.95050	0.28216	1.51100	2.18209
YP-9-6-1		0.28215	0.00033	0.43200	-12.65387	0.28215	1.52793	2.21504
YP-9-7-1		0.28216	0.00047	0.43200	-12.23746	0.28216	1.51590	2.18880
YP-9-8-1		0.28215	0.00024	0.44300	-12.22301	0.28215	1.51800	2.19622
YP-9-9-1		0.28214	0.00024	0.42800	-12.85725	0.28214	1.52992	2.22487
YP-9-10-1		0.28215	0.00038	0.43200	-12.78282	0.28214	1.53424	2.22312
YP-9–11-1		0.28215	0.00032	0.43700	-12.52259	0.28215	1.52694	2.21053
YP-9-11-2		0.28207	0.00281	0.42200	-16.33418	0.28205	1.74667	2.43698
YP-9-12-1		0.28210	0.00036	0.43200	-14.36641	0.28210	1.59484	2.32258
YP-9-13-1		0.28216	0.00031	0.43200	-12.23516	0.28216	1.51115	2.18873
YP-9-14-1		0.28216	0.00037	0.43800	-12.27860	0.28215	1.51968	2.19592
YP-9–15-1		0.28213	0.00024	0.43200	-13.12238	0.28213	1.54351	2.24452
YP-9-16-1		0.28215	0.00045	0.43200	-12.60155	0.28215	1.52935	2.21169
YP-9–17-1		0.28212	0.00026	0.43900	-13.35427	0.28212	1.55873	2.26431
YP-9–18-1		0.28218	0.00044	0.43200	-11.68882	0.28217	1.49370	2.15432
YP-9-18-2		0.28208	0.00037	0.43200	-15.20907	0.28207	1.62787	2.37545
YP-9–19-1		0.28215	0.00023	0.43200	-12.61627	0.28215	1.52371	2.21272
YP-9-20-1		0.28212	0.00021	0.43300	-13.43547	0.28212	1.55545	2.26495
YP-9-21-1		0.28218	0.00014	0.43200	-11.61205	0.28218	1.48247	2.14962
YP-9-22-1		0.28217	0.00030	0.43200	-12.05431	0.28216	1.50392	2.17736
YP-9-23-1		0.28213	0.00020	0.43200	-13.09539	0.28213	1.54129	2.24284
YP-9-24-1		0.28213	0.00042	0.43200	-13.15291	0.28213	1.54977	2.24635
YP-9-25-1		0.28216	0.00024	0.43200	-12.37039	0.28215	1.51448	2.19726
YP06-16-1-1	Dacite	0.28245	0.00054	0.15900	-8.05539	0.28245	1.12382	1.72012
YP06-16-1-2	porphyry	0.28245	0.00029	0.16200	-7.84218	0.28245	1.11183	1.70896
YP06-16-3-1		0.28236	0.00035	0.15700	-11.0682	0.28236	1.23459	1.90940
YP06-16-4-1		0.28251	0.00035	0.15700	-6.00981	0.28250	1.03768	1.58897
YP06-16-5-1		0.28245	0.00046	0.15400	-7.94990	0.28245	1.11356	1.70971
YP06-16-6-1		0.28248	0.00040	0.15900	-7.02115	0.28247	1.07992	1.65461
YP06-16-7-1		0.28245	0.00048	0.15700	-7.99673	0.28245	1.11838	1.71492
YP06-16-8-1		0.28240	0.00041	0.16100	-9.78692	0.28240	1.18986	1.83131
YP06-16-9-1		0.28246	0.00040	0.15700	-7.65614	0.28246	1.10303	1.69337
YP06-16-10-1		0.28246	0.00042	0.16100	-7.55635	0.28246	1.10300	1.69004
YP06-16-11-1		0.28247	0.00037	0.15500	-7.21093	0.28247	1.08326	1.66365
YP06-16-12-1		0.28245	0.00026	0.15500	-7.93181	0.28245	1.10871	1.70938
YP06-16-13-1		0.28197	0.00086	0.43000	-19.16091	0.28196	1.79600	2.62115
YP06-16-14-1		0.28241	0.00049	0.15800	-9.28470	0.28241	1.16974	1.79724
YP06-16-15-1		0.28252	0.00041	0.16400	-5.28074	0.28252	1.01644	1.54795
YP06-16-16-1		0.28247	0.00039	0.15700	-7.23301	0.28247	1.08635	1.66654
YP06-16-17-1		0.28247	0.00029	0.15700	-7.10046	0.28247	1.07868	1.65818
YP06-16-18-1		0.28212	0.00006	0.43000	-13.49145	0.28212	1.55083	2.26630
YP06-16-19-1		0.28240	0.00038	0.15700	-9.74901	0.28240	1.18411	1.82593
YP06-16-20-1		0.28242	0.00048	0.15700	-9.20032	0.28241	1.16529	1.79116
YP06-16-21-1		0.28238	0.00104	0.15700	-10.59943	0.28237	1.23596	1.87943
YP06-16-22-1		0.28241	0.00056	0.15700	-9.27837	0.28241	1.17049	1.79607
YP06-16-23-1		0.28252	0.00064	0.15700	-5.55713	0.28252	1.02660	1.56015
YP06-16-24-1		0.28219	0.00138	0.15700	-17.12852	0.28219	1.50602	2.29101
YP06-16-25-1		0.28228	0.00031	0.15700	-14.03855	0.28228	1.34890	2.09702

Table 4 Zircon Hf isotopic compositions of the igneous rocks in the Yongping Cu-Mo deposits

Table 4 Continued

Samples	Rock types	<sup>176</sup> Hf/ <sup>177</sup> Hf	<sup>176</sup> Lu/ <sup>177</sup> Hf	t (Ga)	$\epsilon_{\text{Hf}(t)}$	<sup>176</sup> Hf/ <sup>177</sup> Hf (t)	T <sub>DM1</sub> (Ga)	T <sub>DM2</sub> (Ga)
YP06-17-2-1	Dacite	0.28236	0.00046	0.15200	-11.11120	0.28236	1.23525	1.90832
YP06-17-3-1	porphyry	0.28239	0.00050	0.15700	-10.23911	0.28239	1.20640	1.85689
YP06-17-5-1	1 1 5 5	0.28220	0.00031	0.15700	-16.66223	0.28220	1.45032	2.26241
YP06-17-6-1		0.28232	0.00050	0.15900	-12.52401	0.28232	1.29732	2.00281
YP06-17-7-1		0.28250	0.00046	0.15700	-6.11163	0.28250	1.04414	1.59539
YP06-17-8-1		0.28250	0.00055	0.15400	-6.15630	0.28250	1.04545	1.59594
YP06-17-9-1		0.28223	0.00046	0.15900	-15.63151	0.28223	1.41717	2.19886
YP06-17-10-1		0.28247	0.00041	0.15600	-7.17436	0.28247	1.08358	1.66207
YP06-17-11-1		0.28254	0.00054	0.15700	-4.90944	0.28254	0.99886	1.51905
YP06-17-12-1		0.28213	0.00059	0.76600	-6.05133	0.28212	1.56155	2.05145
YP06-17-12-2		0.28210	0.00034	0.49800	-13.01315	0.28209	1.59798	2.28693
YP06-17-13-1		0.28238	0.00040	0.15300	-10.39933	0.28238	1.20678	1.86407
YP06-17-14-1		0.28209	0.00077	0.49800	-13.51981	0.28208	1.63004	2.31850
YP06-17-15-1		0.28239	0.00057	0.15700	-10.00960	0.28239	1.19956	1.84234
YP06-17-16-1		0.28213	0.00057	0.15700	-19.14853	0.28213	1.55608	2.41864
YP06-17-17-1		0.28246	0.00072	0.15700	-7.78764	0.28245	1.11617	1.70159
YP06-17-18-1		0.28211	0.00102	0.15700	-20.04548	0.28211	1.60792	2.47462
YP06-18-1-1	Quartz	0.28240	0.00034	0.16100	-9.52294	0.28240	1.17765	1.81464
YP06-18-2-1	porphyry	0.28195	0.00101	0.43300	-19.83452	0.28194	1.83000	2.66537
YP06-18-3-1		0.28226	0.00078	0.16200	-14.60124	0.28226	1.38995	2.13596
YP06-18-4-1		0.28220	0.00051	0.16200	-16.77208	0.28220	1.46560	2.27290
YP06-18-5-1		0.28218	0.00111	0.16200	-17.38215	0.28218	1.51048	2.31087
YP06-18-6-1		0.28218	0.00089	0.16200	-17.47156	0.28218	1.50623	2.31665
YP06-18-7-1		0.28219	0.00064	0.16200	-17.22961	0.28218	1.48805	2.30161
YP06-18-8-1		0.28226	0.00092	0.16200	-14.81269	0.28225	1.40270	2.14921
YP06-18-9-1		0.28241	0.00057	0.16200	-9.33715	0.28241	1.17743	1.80353
YP06-18-10-1		0.28243	0.00033	0.16100	-8.66848	0.28243	1.14419	1.76054
YP06-18-11-1		0.28245	0.00051	0.15600	-8.11085	0.28245	1.12266	1.72139
YP06-18-12-1		0.28247	0.00063	0.20900	-6.25648	0.28247	1.09779	1.64373
YP06-18-13-1		0.28219	0.00101	0.16200	-17.11019	0.28219	1.49629	2.29383
YP06-18-14-1		0.28219	0.00094	0.16200	-17.17023	0.28219	1.49618	2.29766
YP06-18-15-1		0.28219	0.00083	0.16200	-17.19565	0.28219	1.49326	2.29934
YP06-18-16-1		0.28223	0.00103	0.43200	-9.96382	0.28222	1.44244	2.04553
YP06-18-17-1		0.28241	0.00078	0.16200	-9.32460	0.28241	1.18263	1.80265
YP06-18-18-1		0.28224	0.00085	0.16200	-15.19702	0.28224	1.41535	2.17348
YP06-18-19-1		0.28214	0.00081	0.16200	-18.90690	0.28214	1.55944	2.40697
YP06-18-20-1		0.28217	0.00076	0.16200	-17.90786	0.28217	1.51865	2.34419
YP06-18-21-1		0.28214	0.00080	0.16200	-18.96232	0.28214	1.56126	2.41046

Note: t (Ma) refers to the time of single zircon grain formation. The single grains zircon SHRIMP U-Pb age was used to represent it.

The inherited cores of the zircon, in the porphyries can be grouped into three types: Type-1 are fragments or cores with well-defined concentric oscillatory zoning (Fig. 6, YP06-17-4); Type-2 are rounded, high luminescent cores with thin faint concentric zoning (Fig. 6, YP-9 and YP06-18-1); and Type-3 are also rounded, high luminescent cores without intra-grain structure (YP06-16-1). The surrounding rims can be grouped into: (i) rims characterized by well-defined concentric and oscillatory zoning, which is interpreted as magmatic overgrowth (YP06-16, YP06-17, and YP06-18); and (ii) nebulous, high-luminescent

grains with structureless dark rims, suggesting a metamorphic recrystallization (YP-9).

#### 6.1.2 Zircon Th–U chemistry

The Th and U concentrations of zircon grains selected from four samples as the result of SHRIMP analysis are shown in Figures 9 and 10.

The Th/U ratios of zircon from dacite porphyry (YP06-17) range from 0.14 to 0.46 expect for the zircon grain No.10. The CL image of the zircon grain No.10 shows typical igneous characteristics in the core and



**Fig. 8** <sup>40</sup>Ar/<sup>39</sup>Ar plateau ages and isochron ages of biotite and muscovite selected from dacite porphyry (YP06-16, YP06-17), and quartz porphyry (YP06-18).

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Table 5 Muscovite <sup>40</sup>Ar/<sup>36</sup>Ar stepwise heating data of the altered quartz porphyry (YP06-4)

T(°C)	( <sup>40</sup> Ar/ <sup>39</sup> Ar) <sub>m</sub>	<sup>(36</sup> Ar/ <sup>39</sup> Ar) <sub>m</sub>	( <sup>37</sup> Ar/ <sup>39</sup> Ar) <sub>m</sub>	( <sup>38</sup> Ar/ <sup>39</sup> Ar) <sub>m</sub>	<sup>40</sup> Ar*/ <sup>39</sup> Ar	<sup>39</sup> Ar (×10 <sup>-14</sup> mol)	<sup>39</sup> Ar (Cum.) (%)	Age (Ma)	Ca/K	± 1σ (Ma)
500	9.7748	0.0092	0.3048	0.0207	7.0725	20.27	0.49	142.0	0.32	8.4
600	8.2012	0.0022	0.0916	0.0145	7.5536	250.36	6.49	151.2	0.2918	1.9
700	7.6441	0.0007	0.0373	0.0128	7.4416	672.72	22.61	149.1	0.1959	1.7
800	7.9930	0.0006	0.0979	0.0127	7.8263	638.83	37.93	156.5	0.0901	2.1
900	8.2396	0.0009	0.1094	0.0128	7.9899	681.60	54.27	159.6	0.0472	1.9
1000	8.2818	0.0007	0.0215	0.0126	8.0793	825.97	74.07	161.3	0.0213	1.9
1050	8.4606	0.0009	0.0374	0.0129	8.1836	302.23	81.31	163.3	0.0661	1.9
1100	8.2864	0.0006	0.0688	0.0130	8.1197	346.39	89.61	162.1	0.0436	1.6
1200	8.4541	0.0012	0.1314	0.0138	8.1029	138.74	92.94	161.7	0.0421	2.2
1300	8.9950	0.0027	0.0648	0.0140	8.1833	123.16	95.89	163.3	0.0469	2.0
1400	10.2800	0.0070	0.1240	0.0141	8.2292	171.36	100.00	164.2	0.0066	2.6

W = 17.24 mg J = 0.011576.Total age = 158.0 Ma.  $F = {}^{40}\text{Ar}{}^*/{}^{39}\text{Ar}$  (radiogenic).

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Table 6 Biotite <sup>40</sup>Ar/<sup>36</sup>Ar stepwise heating data of the altered dacite porphyry (YP06-16)

T(°C)	( <sup>40</sup> Ar/ <sup>39</sup> Ar) <sub>m</sub>	<sup>(36</sup> Ar/ <sup>39</sup> Ar) <sub>m</sub>	( <sup>37</sup> Ar/ <sup>39</sup> Ar) <sub>m</sub>	( <sup>38</sup> Ar/ <sup>39</sup> Ar) <sub>m</sub>	<sup>40</sup> Ar*/ <sup>39</sup> Ar	<sup>39</sup> Ar (×10 <sup>-14</sup> mol)	<sup>39</sup> Ar (Cum.) (%)	Age (Ma)	Ca/K	± 1σ (Ma)
400	12.6635	0.0312	0.1565	0.0232	27.20	144.39	1.68	61.4	61.4	5.0
500	9.8359	0.0157	0.2933	0.0220	53.00	255.72	4.64	92.1	92.1	2.0
600	11.8281	0.0084	0.0420	0.0206	79.12	221.25	7.21	162.1	162.1	2.3
700	9.8851	0.0016	0.0102	0.0158	95.25	1329.45	22.64	163.0	163.0	2.0
800	9.6477	0.0009	0.0144	0.0164	97.17	1086.34	35.25	162.4	162.4	1.7
900	9.7389	0.0014	0.0267	0.0188	95.73	318.46	38.95	161.5	161.5	2.0
1000	10.0394	0.0020	0.0331	0.0183	94.02	283.92	42.24	163.4	163.4	2.0
1100	10.0554	0.0008	0.0073	0.0144	97.71	1046.48	54.39	169.8	169.8	1.9
1200	9.6703	0.0004	0.0073	0.0143	98.77	1893.80	76.37	165.3	165.3	1.7
1300	9.5773	0.0004	0.0180	0.0143	98.78	2006.25	99.66	163.8	163.8	1.8
1400	9.9222	0.0084	0.3187	0.0346	75.05	29.46	100.00	130.2	130.2	8.0

W = 45.19 mg J = 0.010046.

Total age = 160.5 Ma.  $F = {}^{40}Ar^*/{}^{39}Ar$  (radiogenic).

Table 7 Biotite <sup>40</sup>Ar/<sup>36</sup>Ar stepwise heating data of the altered dacite porphyry (YP06-17)

T(°C)	( <sup>40</sup> Ar/ <sup>39</sup> Ar) <sub>m</sub>	<sup>(36</sup> Ar/ <sup>39</sup> Ar) <sub>m</sub>	( <sup>37</sup> Ar/ <sup>39</sup> Ar) <sub>m</sub>	( <sup>38</sup> Ar/ <sup>39</sup> Ar) <sub>m</sub>	<sup>40</sup> Ar*/ <sup>39</sup> Ar	<sup>39</sup> Ar (×10 <sup>-14</sup> mol)	<sup>39</sup> Ar (Cum.) (%)	Age (Ma)	Ca/K	± 1σ (Ma)
400	15.0564	0.0368	0.2068	0.0279	27.82	52.46	0.60	75.5	75.5	5.2
500	13.2957	0.0213	0.1260	0.0235	52.71	359.45	4.68	124.6	124.6	2.7
600	11.2670	0.0069	0.0234	0.0168	81.77	265.07	7.68	162.1	162.1	2.5
700	9.6241	0.0012	0.0075	0.0146	96.20	2183.44	32.47	162.9	162.9	1.7
800	9.4547	0.0007	0.0101	0.0154	98.19	1341.63	47.70	163.3	163.3	1.8
900	9.5649	0.0012	0.0166	0.0160	96.37	455.01	52.86	162.2	162.2	1.8
1000	9.8096	0.0019	0.0300	0.0184	94.28	178.07	54.88	162.7	162.7	2.1
1100	9.9022	0.0010	0.0102	0.0151	96.97	622.38	61.95	168.6	168.6	2.0
1200	9.4998	0.0006	0.0081	0.0146	98.23	1514.17	79.14	164.1	164.1	1.7
1300	9.5381	0.0006	0.0263	0.0152	98.11	1822.16	99.82	164.5	164.5	1.9
1400	9.8520	0.0089	0.2421	0.0349	73.46	15.91	100.00	129	129	12

W = 45 mg J = 0.010205.

Total age = 161.7 Ma.  $F = {}^{40}Ar^*/{}^{39}Ar$  (radiogenic).

Sample	Weight(g)	Re (µg g <sup>-1</sup> )	<sup>187</sup> Re (µg g <sup>-1</sup> )	<sup>187</sup> Os (ng g <sup>-1</sup> )	Age (Ma)
YP-2 YP-10	0.00740 0.01072	$\begin{array}{l} 79.351 \pm 1.649 \\ 19.251 \pm 0.243 \end{array}$	$\begin{array}{l} 49.877  \pm  1.037 \\ 12.101  \pm  0.010 \end{array}$	$\begin{array}{c} 129.57 \pm 0.97 \\ 31.63 \pm 0.32 \end{array}$	$155.7 \pm 3.6$ $156.7 \pm 2.8$

 Table 8
 Molybdenite Re–Os geochronologic data



Fig. 9 Diagram showing U and Th concentrations for granite porphyry (YP-9), dacite porphyry (YP06-116, YP06-17) and quartz porphyry (YP06-18).

metamorphic characteristics in the rim. The Th/U ratio is 0.17 for the rim and 0.80 for the core, which is similar to the results of Bacon & Lowenstern, (2005), and Hoskin and Schaltegger (2003). The Th/U ratio of zircon from dacite porphyry (YP06-16) ranges from 0.17 to 0.39 expect for the spot 13.1 which has a Th/Uratio of 0.1. The spots no. 2.1, no. 9.1, and no. 13.1 in zircon selected from quartz porphyry (YP06-18) show a constant Th/U ratio of 0.01, which is typical of a metamorphic origin, while the other spots in the same zircon show a Th/U ratio ranging from 0.21 to 0.47, which is typical of igneous origin. The spots in the zircon selected from the granite porphyry sample (YP-9) have a constant Th/U ratio of 0.02, except for the spots no. 4.1 (Th/U = 0.06), no. 8.2 (Th/U = 0.01), and no. 11.1 (Th/U = 0.01) (Fig. 9).

In order to discriminate the origin of zircons, this paper compares zircon ages with its Th/U ratios. As shown in Figure 10, the Th/U ratios of the zircons selected from samples YP06-16, YP06-17, and YP06-18 have wide Th/U ranges with constant ages, whereas sample YP-9 yields constant Th/U ratios and a narrow age range. It is proposed that the samples of YP06-16, YP06-17, and YP06-18 were probably derived from a

mixed two sources, one with higher Th/U ratio and the other with lower Th/U ratio. Both sources could be either metamorphic or igneous. Sample YP-9, however, certainly has a metamorphic origin.

## 6.2 Interpretation of U–Pb, Ar–Ar and Re–Os ages

Previously published geochronological data for the Yongping deposit are mainly based on K-Ar and Rb-Sr analyses, and show a wide range of ages with large errors. The new U-Pb, Ar-Ar, and Re-Os molybdenite ages reported in this paper provide a better constraint on the timing of the magmatism and mineralization in the deposit. Zircons in granite porphyry (YP-9) have typical metamorphic characteristics based on their morphology and Th/U ratios. The sample yields a  $^{206}$ Pb/ $^{238}$ U weighted average age of 435.2  $\pm$ 7.3 Ma, suggesting an earlier Silurian metamorphic event. However, zircons in quartz porphyry (YP06-18) have typical igneous characteristics and show a weighted average age of 162.1  $\pm$  3.4 Ma, suggesting a magmatic crystallization age for the quartz porphyry that is associated with the molybdenite mineralization of the Middle-Late Jurassic time. Sample YP06-18 yield two inherited zircons with ages of 432  $\pm$  16 Ma and 208.6  $\pm$  7 Ma, suggesting two metamorphic events prior to the intrusion and mineralization at the Yongping deposit.

Zircons in sample YP06-17 and YP06-16 collected from the dacite porphyries have typical igneous characteristics based on their morphology and Th/U ratios, and yield ages of 157.5  $\pm$  2.9 Ma and 156.0  $\pm$  1.3 Ma, respectively. The near-identical ages represent magmatic crystallization age for the dacite porphyry associated with copper mineralization in the Late Jurassic. Two inherited, metamorphic zircons yield ages of 498  $\pm$  16 Ma and 430  $\pm$  14 Ma, also suggesting two metamorphic events prior to the intrusion and mineralization at the deposit. The inherited age (766  $\pm$  25 Ma) from a typical igneous zircon (with oscillatory zoning and Th/U = 0.8) may indicate a Neo-Proterozoic magmatic activity.

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Fig. 10 Diagram showing 206 Pb/238 U (Ma) and 232 Th/238 U ratios for granite porphyry (YP-9), dacite porphyry (YP06-116, YP06-17) and quartz porphyry (YP06-18).

Ar–Ar ages for hydrothermal biotite from the dacitie porphyry that is associated with copper mineralization and Ar–Ar ages for muscovite from the quartz porphyry that is associated with molybdenum suggest that, the muscovite crystallized later than the hydrothermal biotite. Two near-identical molybdenite Re–Os dating results, 155.7  $\pm$  3.6 Ma and 156.7  $\pm$  2.8 Ma, show a Late-Jurassic molybdenite crystallization subsequent to emplacement of the dacite porphyry.

Based on the zircon U–Pb age data, several episodes of magmatic and metamorphic events occurred in the deposit and its vicinities. The oldest magmatic activity in the deposit occurred in the Neo-Proterozoic time (766  $\pm$  25 Ma), followed by Late Cambrian, Early Silurian, and Late Triassic metamorphic events. The Late Cambrian and Late Triassic events very likely caused migmatization. The Middle–Late Jurassic igneous event produced the dacite and quartz porphyries that were associated with the copper and molybdenum mineralization in the deposit.

#### 6.3 Source of the magmas

Sr-Nd isotopes and zircon Hf isotopes of granitoids are a valuable tracer for magma sources. The quartz porphyry in the deposit has wide  $\varepsilon_{Hf}(t)$  values of -6.2 to -18.9, and wide  $\varepsilon_{Nd}(t)$  values of -10.95 to -6.94. The dacite porphyry has wide  $\varepsilon_{Hf}$  (t) values of -5.3 to -13.5, but narrow  $\varepsilon_{Nd}(t)$  values of -9.18 to -7.64. Their  $\varepsilon_{Nd}(t)$ values are close to that of the granites of southeast China (with  $\varepsilon_{Nd}(t)$  values from -16.4 to -1.1 and  $I_{Sr}$ values from 0.7056 to 0.7404) (Shen *et al.*, 1998). The  $\varepsilon_{Nd}$ (t) values represent the initial values of  $\varepsilon_{Nd}$  in the rock at the time of its crystallization. The negative  $\varepsilon_{Nd}$  (t) value implies a magma source with lower Sm/Nd than CHUR, i.e. enriched mantle sources or crustal source. The Sr-Nd isotopic data at Yongping suggests a dominated crustal source for both the dacite porphyry and the quartz porphyry. In the diagram of  $\varepsilon_{Hf}$  (t)-t (Ma) (Fig. 11), the data plot in the intermediate area between depleted mantle and crust, also indicating a mixed magma source. In the  $\varepsilon_{Nd}$  (t)- $I_{Sr}$  diagram (Fig. 12), all the



Fig. 11 eHf (t)-t(Ma) diagram for granite porphyry (YP-9), dacite porphyry (YP06-116, YP06-17) and quartz porphyry (YP06-18). Depleted mantle and Crustal line are according to Shen *et al.* (1998).

t(Ma)



Fig. 12 εNd(t)-ISr(t) diagram for both dacite and quartz porphyries. Data source: Dexing Cu–Mo deposit (JGEB, 1996); Yinshan Cu–Au–Pb–Zn–Ag deposit (JGEB, 1996); Lengshuikeng Pb–Zn–Ag deposit (Meng *et al.*, 2007); Yulong Cu–Mo deposit (Jiang *et al.*, 2006). Depleted mantle value and pre-cambrian metamorphic rocks are according to Zhu *et al.* (1990) and Shen *et al.* (1998).

data from the dacite porphyry and the quartz porphyry plot in the IV quadrant, again showing a mingling of magma sources between the mantle and the crust. In the diagram of  $\varepsilon_{Nd}$ (t)-t (Ma) (Fig. 13), all the data plot within the Meso-Proterozoic crustal evolution area, suggesting both the dacite porphyry and the quartz porphyry derived from partial melting of the Meso-Proterozoic crust. Two-stage model ages (T<sub>DM2</sub>) of



**Fig. 13** εNd(t)-t(Ma) diagram for both dacite and quartz porphyries (modified after Shen *et al.,* 1998).

dacitie porphyry (1.58Ga–1.69Ga) and quartz porphyry (1.51Ga–1.84Ga) also indicate the contribution of Meso-Proterozoic crust to the magma sources. Compared with other ore-related porphyries, for example, the Dexing granodiorite porphyry and the Yinshan dacite porphyry in northeast Jiangxi Province, the ore-related porphyries at Yongping show characteristics of lower  $\varepsilon_{Nd}$  (t) and older Nd model age (Wang *et al.,* 2006). This may suggest that the source magmas were mixed in different proportions for quartz porphyry and dacite porphyry at the Yongping Cu–Mo–W deposit, Dexing porphyry Cu–Mo–Au, Yinshan polymetallic deposit, and Lengshuikeng Pb–Zn deposit (Fig. 12).

#### 6.4 Implications for regional mineralization

The dacite and quartz porphyries at Yongping deposit are subduction-related high-K calc-alkaline series (Li et al., 2007a). The intrusive rocks at the Dexing porphyry Cu-Mo deposit, located in the northeast Jiangxi Province, about 100 km from the Yongping deposit (Fig. 1), also shows subduction-related magmatic arc characteristics, but the intrusive rocks at both deposits show different geochemical compositions. The intrusive rocks at both deposits are poor in Nb, Ta, and Ti contents, however, the intrusive rocks at the Dexing deposit are rich in Rb, Ba, Zr, Hf, Sr, and P, whereas intrusive rocks at the Yongping deposit is poor in Sr and P. This may reflect differences in their magma sources and in fractional crystallization. The ore-related porphyry in the Dexing Cu-Mo deposit shows adakite affinity, whereas the dacite and quartz porphyries in the Yongping deposit do not (Li & Watanabe, 2006). This suggests a difference in tectonic setting and source

materials of magma between ore-related intrusive rocks at the Dexing and Yongping deposits. Rb/Sr ratios of the intrusive rocks at the Dexing deposit (0.05–0.28) are lower than that of the porphyries in the Yongping deposit (Rb/Sr ratios of 0.35–1.12 for its dacite porphyry and Rb/Sr ratios of 1.59–9.29 for its quartz porphyry).

The zircon SHRIMP U–Pb age analysis of the dacite and quartz porphyries shows that the ore-related porphyries were emplaced at 160 Ma at the Yongping deposit. The molybdenite Re-Os ages show the Mo mineralization at 157 Ma at Yongping. Both the orerelated porphyries and the Cu-Mo mineralization occurred in the Late Jurassic time. The ages of the Dexing porphyry Cu-Mo deposit (zircon SHRIMP U-Pb age of 171 Ma and molybdenite Re-Os age of 170 Ma, Lu et al., 2005; Wang et al., 2006; Li et al., 2013) and the Yinshan polymetal deposit (zircon SHRIMP U-Pb age of 181 Ma and muscovite Ar-Ar age of 179 Ma) (Li et al., 2007b) are older than that of the Yongping deposit, although they are coeval with the largescale W-Sn mineralization (for example, Shizhuyuan W–Sn deposit of 154.7  $\pm$  2.5 Ma and Dangping W deposit of 154.9  $\pm$  2.4 Ma) in southeast China. The Dexing porphyry Cu-Mo-Au and Yinshan polymetallic deposits are considered to be the products of an arc magmatism within a transpressional tectonic setting (Watanabe & Li, 2006; Li & Sasaki, 2007). The orerelated rocks with large-scale W-Sn mineralization in southeast China are mainly biotite granite (Hua et al., 2005). Obviously the differences in magma sources and in tectonic settings were accountable for production of different ore-related igneous rocks and deposits. It is suggested that the Yongping Cu-Mo deposit was produced during the transitional setting from a compressional tectonic regime to an extensional tectonic environment, while the southeast China underwent a widespread crustal extension in Late Jurassic Period (Hua et al., 2005).

## 7. Conclusions

This study reveals that the dacite and quartz porphyries at the Yongping deposit show similar REE patterns with subduction-related features, i.e., LILE-enrichment (Ba, Rb, and K) and HFSE (Th, Nb, Ta, and Ti) depletion. The zircon SHRIMP U-Pb ages indicate that the ore-related porphyries were emplaced at Late Jurassic. The inherited ages (432 Ma and 208.6 Ma) were generated by two metamorphic events. The inherited age of 766 Ma from an igneous zircon suggested a NeoProterozoic magmatic event in the Yongping district. The dating results of hydrothermal muscovite and biotite from the porphyries indicate that the Mo mineralization and the associated hydrothermal activity are coeval with the magmatic intrusions at Yongping. The Sr-Nd isotopic values for the dacite and quartz porphyries indicate a mixture magma source. The  $\varepsilon_{Nd}(t)$ data from the dacite and quartz porphyries suggest a partial melting of the Meso-Proterozoic crust to contribute to the magma process. Comparing with the neighboring Dexing porphyry Cu-Mo and the Yinshan polymetallic deposits, as well as the Shizhuyuan and Dangping W deposits of south China in terms of their geochemical features and tectonic settings, it is proposed that the Yongping deposit was produced during the tectonic transition from a compressional tectonic regime to an extensional tectonic environment at the Late Jurassic.

## Acknowledgments

This study was financially supported by the National Fundamental Research Development Program of China (grant no. 2012CB416705), the National Natural Science Foundation of China (grant no. 41272112, 40872065), and the One Hundred Person Project of the Chinese Academy of Sciences. We thank Huang Zhenjin, Wei Xinglin for their assistance during our field seasons, and Professor Hideo Hirano for his kind help for samples thin sections descriptions. We are indebted to Zhangyan, Dong Huiyan, and Chen Zhenyu for their help with SHRIMP U–Pb analysis and CL imaging.

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