

Article

Geology and Isotope Systematics of the Jianchaling Au Deposit, Shaanxi Province, China: Implications for Mineral Genesis

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Abstract: The giant Jianchaling Au (52 t Au) deposit is located in the Mian-Lue-Yang Terrane in the southern part of the Qinling Orogen of central China and is hosted by metamorphosed carbonate rocks of the Late Neoproterozoic Duantouya Formation. The deposit consists of multiple generations of mineralised quartz(-carbonate) veins in WNW-trending extensional ductile-brittle shear zones. Based on the mineral assemblages and cross-cutting relationships between the quartz(-carbonate) veins, the paragenesis is characterised by an early coarse-grained pyrite-pyrrhotite-pentlandite-dolomite-quartz assemblage (I), followed by pyrite-sphalerite-galena-carbonate-arsenopyrite-fuchsite-carbonate-quartz containing gold (II), and fine-grained pyrite-dolomite-calcite-quartz-realgar (As₂S₂)-orpiment (As₂S₃) (III). The H-O-C isotope systematics for the three vein sets indicate that the mineralising fluid is probably sourced from the metamorphic dehydration of carbonate rocks in the Duantouya Formation, and gradually mixed with meteoric water during the emplacement of the third vein set. The δ^{34} S values for sulfides (6.3–16.6‰) from the second auriferous vein set are greater than zero, indicating sulfates reduction from the Neoproterozoic metamorphic rocks (Duantouya Fm). The (²⁰⁶Pb/²⁰⁴Pb)_i ratios from pyrite (17.521–18.477) from each of the vein sets overlap those of the ultramafic rocks (18.324–18.717) and the Bikou Group (17.399–18.417), indicating that the units are possible sources for the sulfides in the mineralisation. Both ε Nd(t) and Isr(t) of sulfide overlap with the meta-ultramafic field and Duantouya formation and dominated with mature Sr-Nd character, which indicated that the Duantouya may play an important role during the ore formation and there may exist a minor ultramafic source that is involved in the ore fluid. The S-Pb-Sr-Nd isotopic ratios are closely related to those of the Bikou Group and Duantouya Formation, which indicates that the mineralised fluid has interacted with both units. Combining the previously published data with data from this study on the mineralised area, we surmise that Jianchaling is characteristic of an orogenic-type gold deposit related to the Triassic Qinling Orogeny associated with continental collision.

Keywords: Jianchaling Au deposit; isotope geochemistry; ore genesis; orogenic-type gold deposit; Qinling Orogen; central China



1. Introduction

The different tectonic events are accompanied by different types of mineralisation and associated hydrothermal alteration of different ages [1–3], including orogenic-Au deposits [4–6]. Many of these deposits record multiple deformation events, making then difficult to study and leading to uncertainty in their genesis [7–9]. Nonetheless, some of the deposits have been identified as orogenic-types, such as the Juneau Gold Belt in America [10], the Sukhoi Log gold deposit in Russia [11], the Victorian gold province in SE Australia [12], the Jiaodong gold province in China [13] and the Shanggong, Wenyu and Huachanggou Au deposits, and Tieluping, Weishancheng, Yindonggou, Wangpingxigou, Dahu and Zhifang Ag-Au-Mo deposits [14–26] in China of the Qinling Orogen.

Other deposits present in the Qinling Orogen include porphyry-, skarn-, breccia pipe-, hydrothermal-, Carlin-, and epizonal-types, which have been studied in detail [1,27–38]. Orogenic gold deposits recognised in the orogen include Jianchaling, Lijiagou and Huachanggou [8,25,39], of which Jianchaling is the largest (Figure 1). However, the genesis of the giant Jianchaling Au deposit is still poorly understood.

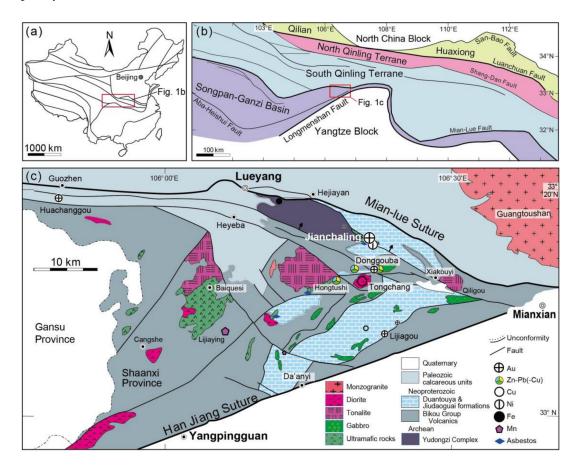


Figure 1. Geological maps showing: (**a**) the tectonic setting of China; (**b**) tectonic subdivision of Qinling Orogen and location of the Mianxian, Lueyang, and Yangpingguan (MLY) area [18]; and (**c**) regional geology and location of gold deposits in the MLY [39].

The Jianchaling Au deposit is located in the triangular area marked by Mianxian, Lueyang, and Yangpingguan towns (MLY), in the South Qinling Terrane of the Qinling Orogen (Figure 1c). The deposit has a measured resource of 52 t Au with an average grade of 5 g/t hosted by quartz-veins in a ductile-brittle shear zone [21,39–43]. Even though the genesis of the mineralisation at the deposit has been studied, the source of the metals and nature of the mineralising fluids have remained unclear.

2. Geological Setting

The Qinling Orogen is located south of the North China Block (NCB) and has experienced a complex tectonic history [44–46]. It is located in the central portion of the east-trending Central China Orogen, and developed during the Mesozoic collision between the North China and Yangtze blocks (Figure 1a,b) [1,47–49]. The Luanchuan Fault is the northern boundary of the orogen, and the southern boundary is the Longmenshan Fault. Deformation associated with the orogen extends into the Huaxiong Domain that is also included in the southern part of the Precambrian North China Block. The subdivisions of the orogen are shown in Figure 1b.

The Jianchaling deposit is a fault-controlled lode-gold deposit hosted by the Neoproterozoic Duantouya Fm in the eastern part of the MLY (Figure 1). The group unconformably underlies Late Paleozoic turbiditic successions in the Songpan-Ganzi Basin situated between the Mian-lue Suture to the north and Han Jiang Suture to the south (Figure 1c). The area is an important part of the Qinling Orogen and hosts many mineral deposits (e.g., Au, Cu, Ni, Zn, Pb, Mn) in various lithological and structural settings (Figure 1c), and is commonly known as the Golden Triangle [42].

The rock-types in the MLY area include metamorphosed Neoarchean granite-greenstones in the Yudongzi Complex, Neoproterozoic volcanic sequences in the Bikou Group, Late Neoproterozoic calcareous units in the Duantouya and Jiudaoguai formations, and Paleozoic carbonate in the Lueyang Formation (Figure 2). The Bikou Group is widespread in the area and consists of greenschist to amphibolite facies felsic and mafic volcanic protoliths. Various generations of metamorphism and ductile to brittle NE- and NW-trending faults (including thrusts and basin-forming extensional faults) have developed in the area, which control the location of felsic, mafic and ultramafic intrusive rocks and mineralisation (Figure 2).

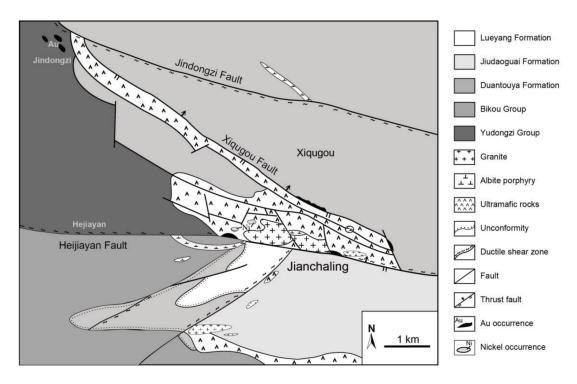


Figure 2. Geological sketch map of the Jianchaling Au deposit [21,39].

3. Geology and Mineralization of the Jianchaling Area

Rocks outcropping in the Jianchaling area Neoarchean Yudongzi Complex, Neoproterozoic Bikou Group, Late Neoproterozoic Duantouya and Jiudaoguai formations, and Paleozoic Lueyang Group (Figure 2). The deposit can be subdivided into northern and southern zones containing five lenticular mineralisation in the WNW-trending Hejiayan and NW-trending Xiqugou faults and their splays. The richest mineralisation is the "No.1 Au orebody" that is 1950 m long, up to 23 m wide, and dips >43° N.

The Hejiayan and Xiqugou faults are brittle-ductile shear zones [50], and host mineralised lenticular quartz veins at the contact between ultramafic rocks and calcareous beds in the Late Neoproterozoic Duantouya and Jiudaoguai formations (Figures 2 and 3). The quartz veins shallowly dip between 20° and 40° WSW is places in steeply dipping sections of the fault.

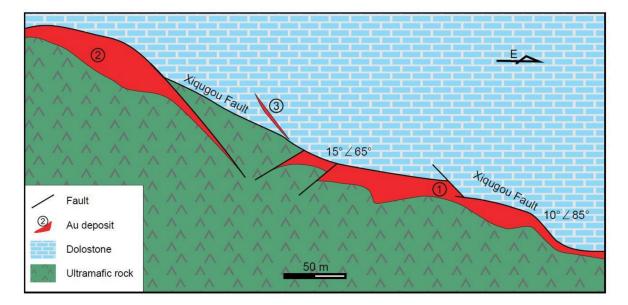


Figure 3. Cross-section of the of the Jianchaling Au deposit at a 870 m elevation showing the relationship of the mineralisation and fault [21,51].

The mineralisation includes native gold and lesser amounts of electrum and native silver associated with pyrite and minor amounts of hematite, magnetite, limonite, chromite, marcasite, realgar, and orpiment. The gangue minerals are dolomite, calcite and quartz, with minor amounts of serpentine, fuchsite, and albite (Figure 4a–c) [39].

Yue [21,39] recognized early deformed and recrystallised quartz—pyrite—carbonate vein crosscut by the quartz—gold—fuchsite—arsenopyrite—pyrite—magnetite second generation vein set, and the third generation of dolomite—calcite—quartz—fine-grained pyrite—realgar—orpiment veins (Figure 4c). The pyrite associated with each of the vein sets are also distinct from each other. The first vein set contains pyrite that is >2 mm across, variably fractured, anhedral to subhedral in shape, and intergrown with euhedral fuchsite and serpentine (Figure 4d,e). The second pyrite generation is not deformed, subhedral, 1–2 mm across, and commonly has cores or inclusions of the first generation pyrite, chalcopyrite, and sphalerite (Figure 4f,g). This pyrite is also commonly altered to hematite with martite pseudomorphing magnetite (Figure 4h). The third generation of pyrite is <10 μ m across, subhedral to euhedral, and present in alteration zones containing the third vein (Figure 4i). The paragenesis is illustrated in Figure 5.

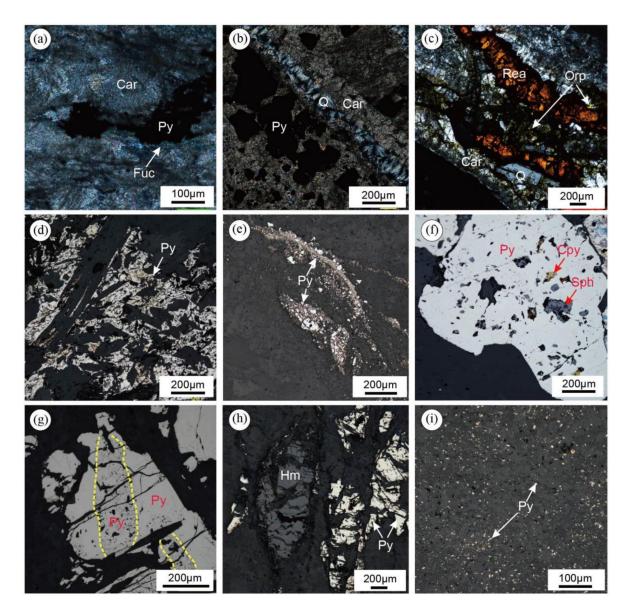


Figure 4. Photomicrographs of samples from the Jianchaling Au eposit showing: (**a**) intergrown pyrite and fuchsite in the mineralised second vein set; (**b**) a second vein set with quartz containing pyrite; (**c**) third vein set with comb-like quartz, calcite and dolomite vein orpiment and realgar; (**d**) pyrite in the first vein set intergrowth; (**e**) fine-grained cataclastic pyrite following schistosity developed during brittle-ductile deformation before emplacement of the mineralised second vein set; (**f**) pyrite wrapping sphalerite and chalcopyrite in the mineralised second vein set; (**g**) pyrite in the first vein set overgrown by pyrite related to the mineralised second vein set; (**h**) hematite formed from pyrite, and martite; (**i**) euhedral fine-grained pyrite in the third vein set. Abbreviations: Cal = calcite; Car = carbonates; Cpy = chalcopyrite; Dol = dolomite; Fuc = fuchsite; Hm = hematite; Orp = orpiment; Py = pyrite; Q = quartz; Rea = realgar; Sph = sphalerite.

Generation Mineral	Vein set 1	Vein set 2	Vein set 3
Pyrite			
Gold			
Pyrrhotite			
Chalcopyrite	-		
Sphalerite	<u> </u>		-
Galena			-
Marcasite			
Arsenopyrite			-
Fuchsite			-
Realgar			
Orpiment			
Dolomite			
Quartz			
Calcite			

Figure 5. Paragenetic relationship of the minerals at the Jianchaling Au deposit [21].

4. Sampling and Analytical Methods

Fresh samples were collected from each of the vein sets at depth in the Jianchaling mine. The samples were examined in thin section to confirm their paragenetic relationship. The sample characteristics are listed in Table 1.

Six samples of pyrite from the second (mineralised) vein set and five samples of the Duantouya Formation and meta-ultramafic from the wall-rock were collected for lead, strontium, and neodymium isotopic studies. The samples were crushed into power.

The samples were then crushed to a minus 10 *mesh* size (420 microns, and sulfide and carbonate fragments were handpicked under a binocular microscope. Between 10 and 50 mg of the powder was leached in acetone and washed in distilled and deionised water to remove contamination, then dried in an oven at 60 °C. The samples were then dissolved in a solution of HF + HNO₃ + HClO₄, dried, and redissolved in 6 N HCl, redried, and redissolved in 0.5 N HCl (for Sr and Nd separation) or 0.5 N HBr (for Pb separation). The Sr and Nd fractions were separated following standard chromatographic techniques using AG50 × 8 and PTFE–HDEHP resins with HCl as eluent. The Pb fraction was separated using a strong alkali anion exchange resin with HBr and HCl as eluents.

The lead isotopes were analysed using a MAT-261 thermal ionization mass spectrometer with the standard NBS 981 in the Analytical Laboratory at the Beijing Research Institute of Uranium Geology, China. Measurements of the common-Pb standard NBS 981 gave average 208 Pb/ 206 Pb values of 2.1681 \pm 0.0008, 207 Pb/ 206 Pb of 0.91464 \pm 0.00033, and 204 Pb/ 206 Pb values of 0.059042 \pm 0.000037; the uncertainty is of <0.1% at the 95% confidence level. Some of the U, Th and Pb values were used to estimate the Pb isotope ratios assuming an age of 198 Ma, which is the 40 Ar/ 39 Ar date determined on fuchsite from the Jianchaling deposit [39]. These Pb isotopic ratios are presented as (208 Pb/ 204 Pb)_i, (207 Pb/ 204 Pb)_i and (206 Pb/ 204 Pb)_i.

A TRITON thermal ionization mass spectrometer was used to measure the Sr and Nd isotopes in the Analytical Laboratory at the Tianjin Institute of Geology and Mineral Resources, China. The ⁸⁷Sr/⁸⁶Sr isotope ratios were normalized against the ⁸⁶Sr/⁸⁸Sr = 0.1194 and ¹⁴³Nd/¹⁴⁴Nd isotope ratios to ¹⁴⁶Nd/¹⁴⁴Nd = 0.7219. The JNdi Nd-Standard yielded ¹⁴³Nd/¹⁴⁴Nd ratios of 0.512118 \pm 0.3 against a reference value of 0.512115 \pm 0.7 [52], and the NBS 987 Sr standard with a reference level of 0.710248 was used yielding ⁸⁷Sr/⁸⁶Sr ratios of 0.710250 \pm 0.7. The Sr and Nd isotopic compositions were measured with a thermal ionization ISOPROBE-T mass spectrometer.

Sample NO.	Mineral	Sample Description	Stage	Analyzed Isotope
800-53-2	Dolomite	Dolomite veins in the Serpentinite with shear deformation	1	С, О
860-52-7	Dolomite	Dolomite veins in the Serpentinite with shear deformation	1	С, О
J-6	Dolomite	Pale green dolomite veins in the cataclastic dolomite	1	С, О
913-32-7	Calcite	Calcite veins in the cataclastic dolomite with a small number of pyrites	2	С, О
913-32-6	Calcite	Calcite veins in the cataclastic dolomite with a small number of pyrites	2	С, О
913-1	Dolomite	Milky white dolomite vein	3	С, О
913-2	Dolomite	Milky white dolomite vein	3	С, О
960-38-1	Dolomite	Milky white dolomite vein	3	С, О
860-51-5	Dolomite	Dolomite veins in the listwanite with a small number of pyrites	3	С, О
860-51-6	Dolomite	Dolomite veins in the listwanite with a small number of pyrites	3	С, О
hjy-8	Pyrite	Grain cluster pyrite with shear deformation	1	S, Pb
J3	Pyrite	Massive sulfide ore	2	S, Pb, Sr, Nd
J3	Pyrrhotite	Massive sulfide ore	2	S
hjy-1	Pyrite	Silicified and fuchsite dolomite ore rocks with disseminated pyrites	2	S, Pb, Sr, Nd
hjy-12	Pyrite	Silicified and fuchsite dolomite ore rocks with disseminated pyrites	2	S, Pb, Sr, Nd
hjy-10	Pyrite	Intensive silicified and fuchsite dolomite ore rocks with disseminated pyrites	2	S, Pb, Sr, Nd
hjy-20	Pyrite	Silicified and fuchsite dolomite ore rocks with disseminated pyrites	2	S, Pb, Sr, Nd
hjy-14	Pyrite	Weak silicified and fuchsite dolomite ore rocks with disseminated pyrites	2	S, Pb, Sr, Nd
H-4	Whole-rock	Shale (Duantouya Formation)		S, Pb, Sr, Nd
960-34-1	Whole-rock	Dolomite		S, Pb, Nd
960-28-1	Whole-rock	Dolomite		S, Pb, Sr, Nd
780-50-3	Whole-rock	Serpentinite		S, Pb, Sr, Nd
800-52-1	Whole-rock	Listwanite		S, Pb, Sr, Nd

Table 1. The geological characteristics of samples from the Jianchaling Au deposit (modified from [21]).

5. Isotope Results

The following are isotopic values determined from samples from the Jianchaling deposit.

5.1. Carbon, Oxygen and Hydrogen Isotopes

The δ^{13} C, δ^{18} O and δ D values from quartz and carbonate from each of the three vein sets and the surroundings rocks are listed in Table 2. These results are published on [21]. The δ^{13} C values for first vein set ranges from -4.4 to 0.6%, the δ^{13} C values for the second (mineralised) vein set ranges from -2.9 to -0.4%, and the δ^{13} C value for the third vein set is between -4.4 to 2.2% (Table 2).

The δ^{18} O values from carbonate and quartz for the first vein set forms two groups of 18.9–23.4‰ and 14.0–17.3‰. The δ^{18} O values from the second vein set are between 13.8 and 19.0‰ (averaging 16.4‰), and for the third vein set are between 11.9 and 16.3‰ (Table 2).

5.2. Sulfur Isotopes

The δ^{34} S isotopic values for the sulfides samples are listed in Table 3 and cited from [21]. The Jianchaling deposit is characterized by highly δ^{34} S positive values with a narrow range. The first vein set has a δ^{34} S value of 14.3‰ (Table 3). The second vein set has δ^{34} S values between 8.2 and 14.3‰. One shale rock sample of the Duantouya Formation have δ^{34} S values of 16.6‰. Two samples of the metamorphosed ultramafic rocks (listwanite and serpentinite) have δ^{34} S values of 6.1 to 8.6‰.

5.3. Lead Isotopes of Sulfides and Wall Rocks

The Pb isotopic analyses completed for this study and previous data are listed in Table 4. In order to compare the contribution of the wall rocks, we also put the lead isotopes of the wall rocks including Yudongzi Fm, Porphyritic granite, Duantouya Fm and Bikou Group in Table 3. Sulfides from gold deposits have ${}^{206}Pb/{}^{204}Pb$ values between 17.257 and 18.477, ${}^{207}Pb/{}^{204}Pb$ values between 15.530 and 15.704, and ${}^{208}Pb/{}^{204}Pb$ values of 36.927 to 38.757 (Table 4). Their calculated (${}^{206}Pb/{}^{204}Pb$)_i, (${}^{207}Pb/{}^{204}Pb$)_i and (${}^{208}Pb/{}^{204}Pb$)_i values are the same as the measured values due to their very low U and Th contents. Therefore, the analytical data is interpreted as being reliable and representative of the source for metals in the mineralising fluid. The ultramafic rocks have ${}^{206}Pb/{}^{204}Pb$ values of 17.952–19.193, ${}^{207}Pb/{}^{204}Pb$ values of 15.520–15.785, and ${}^{208}Pb/{}^{204}Pb$ ratios of 36.029–38.920 (Table 4).

5.4. Strontium and Neodymium Isotopes of Sulfides and Wall Rocks

Four pyrite samples from the gold-bearing second generation vein set have I_{Sr} values in the range 0.706709–0.715929 (average of 0.711973, Table 5), which is similar to that of the meta-ultramafic rocks. The (¹⁴³Nd/¹⁴⁴Nd)_i values of the samples are between 0.511376 and 0.512453 (average of 0.511823, Table 6), and the ε_{Nd} values are between –19.6 and 1.4.

Samples from the Duantouya Formation have $({}^{143}Nd/{}^{144}Nd)_i$ values of 0.511865–0.512114 (average of 0.511953, Table 6), and ϵ_{Nd} values of -10.1 to -5.3. The meta-ultramafic samples have $({}^{143}Nd/{}^{144}Nd)_i$ values of 0.512137–0.512152 (average of 0.512144, Table 6), and ϵ_{Nd} values between -4.8 and -4.5.

Sample No.	Sample	Vein Set	Mineral	δ ¹⁸ O _{mineral} (‰)	Average $\delta^{18}O_{Water}$ (‰)	δD_{Water} (‰)	$\delta^{13} C_{V-PDB}$ (‰)	Average $\delta^{13}C_{CO2}$ (‰)	<i>T</i> [#] (°C)	Reference
MS-57	Au mineralisation	1	quartz	17.3	10.8 (8.3 to 12.1)	-72			310 (250 to 355)	[53]
MS-79	Au mineralisation	1	quartz	17.0	10.5 (8.0 to 11.8)	-85			310 (250 to 355)	[53]
MS-52	Au mineralisation	1	quartz	16.6	10.1 (7.6 to 11.4)	-79			310 (250 to 355)	[53]
MS-77	Au mineralisation	1	quartz	14.0	7.5 (5.0 to 8.8)	-70			310 (250 to 355)	[53]
800-53-2	Au mineralisation	1	dolomite	23.4	17.7 (15.7 to 18.8)		-0.3	1.7 (0.7 to 2.2)	310 (250 to 355)	[53]
J6	Au mineralisation	1	dolomite	18.9	13.2 (11.2 to 14.3)		0.6	2.6 (1.6 to 3.1)	310 (250 to 355)	[53]
860-52-7	Au mineralisation	1	dolomite	19.1	13.4 (11.4 to 14.5)		-4.4	-2.4 (-3.4 to -1.9)	310 (250 to 355)	[53]
	Average			18.0	11.9 (9.6 to 13.1)	-77	-1.4	0.6 (-0.4 to 1.1)		
913-32-6	Au mineralisation	2	calcite	17.0	9.1 (7.2 to 11.2)		-2.9	-1.9 (-3.0 to -0.9)	240 (200 to 300)	[21]
913-32-7	Au mineralisation	2	calcite	16.5	8.6 (6.7 to 10.7)		-1.7	-0.7 (-1.8 to 0.3)	240 (200 to 300)	[21]
P633-61-I1	Au mineralisation	2	calcite	19.0	11.1 (9.2 to 13.2)		-0.6	0.4 (-0.7 to 1.4)	240 (200 to 300)	[41]
J163	Au mineralisation	2	dolomite	16.4	8.3 (6.4 to 10.4)		-1.6	-0.8 (-1.8 to 0.3)	240 (200 to 300)	[41]
J170	Au mineralisation	2	dolomite	13.8	5.7 (3.8 to 7.8)		-0.4	0.4 (-0.6 to 1.5)	240 (200 to 300)	[41]
J180	Au mineralisation	2	dolomite	14.6	6.5 (4.6 to 8.6)		-1.2	-0.4 (-1.4 to 0.7)	240 (200 to 300)	[41]
J14	Au mineralisation	2	ferrocalcite	17.5	9.6 (7.7 to 11.7)		-1.1	-0.1 (-1.2 to 0.9)	240 (200 to 300)	[41]
	Average			16.4	8.4 (6.5 to 10.5)		-1.4	-0.4 (-1.5 to 0.6)		
913-1	Au mineralisation	3	dolomite	13.8	3.2 (0.4 to 4.8)		1.6	1.0 (-0.5 to 1.9)	190 (150 to 220)	[21]
913-2	Au mineralisation	3	dolomite	11.9	1.3 (-1.5 to 2.9)		2.2	1.6 (0.1 to 2.5)	190 (150 to 220)	[21]
960-38-1	Au mineralisation	3	dolomite	13.1	2.5 (-0.3 to 4.1)		0.6	0 (-1.5 to 0.9)	190 (150 to 220)	[21]
860-51-5	Au mineralisation	3	dolomite	14.8	4.2 (1.4 to 5.8)		-4.4	-5.0 (-6.5 to -4.1)	190 (150 to 220)	[21]
860-51-6	Au mineralisation	3	dolomite	14.9	4.3 (1.5 to 5.9)		-4.1	-4.7 (-6.2 to -3.8)	190 (150 to 220)	[21]
MS-69	Au mineralisation	3	quartz	16.3	3.9 (0.8 to 5.8)	-81			190 (150 to 220)	[41]
	Average	3	_	14.1	3.2(0.4 to 4.9)		-0.8	-0.7(-2.9 to -0.5)		
P404-44B-2	Mineralized dolostone		dolomite	22.9			0.0			[41]
P404-43B	Altered dolostone		dolomite	18.0			0.1			[41]
P406-Y-43w	Altered dolostone		dolomite	16.9			0.7			[41]
J142	Dolostone		dolomite	22.5			2.3			[41]
J171	Dolostone		dolomite	23.1			2.3			[41]
J25	Dolostone		dolomite	22.3			-0.4			[41]
2	Average			21.0			0.8			[41]
J167	Serpentinite		dolomite	15.0			-3.8			[41]
J42-1	Listwanite		magnesite	14.3			-2.7			[41]
J59	Listwanite		magnesite	16.3			-3			[41]
Y-Mg	Listwanite		listwanite	13.8			-3.2			[41]
Y-Mg	Listwanite		listwanite	13.8			-3.2			[41]
G-Mg	Listwanite		listwanite	13.4			-2.2			[41]
0	Average			14.5			-3.0			

Table 2. C-H-O isotope composition of mineralisation at the Jianchaling Au deposit (modified from [21]).

Notation: [#] the peak of homogenization temperatures (the range of homogenization temperatures) are from [39]. The $\delta^{18}O_{wtaer}$ values were calculated using equations for quartz–water and carbonate–water [54,55]. The $\delta^{13}C$ in CO₂ equilibrated with dolomite and calculate were calculated using the equations of [56,57].

Sample	Mineral	$\delta^{34} \mathrm{S_{v-CDT}}$ (‰)	Sample	Mineral	$\delta^{34} \mathrm{S_{v-CDT}}$ (‰)
hjy-1 (second vein set) ^c	Py	11.3	Carbonate (Duantouya Fm) ^a	Py	16.4
hjy-12 (second vein set) ^c	Py	10.0	Carbonaceous shale (Duantouya Fm) ^a	Py	5.4
hjy-10 (second vein set) ^c	Py	10.8	Carbonaceous shale (Duantouya Fm) ^a	Py	10.6
J3 (second vein set) ^c	Po	8.2	Carbonate (Duantouya Fm) ^a	Py	12.5
J3 (second vein set) ^c	Py	8.7	Hc-Py1 (serpentinite) ^a	Сру	12.0
hjy-20 (second vein set) ^c	Py	13.2	L-Py-Mt (listwanite) ^a	Po	10.5
hjy-14 (second vein set) ^c	Py	14.3	Ni-bearing ultramafics ^a	Py	9.3
hjy-8 (first vein set) ^c	Py	14.3	Ni-bearing ultramafics ^a	Py	11.3
H-4 Shale (Duantouya Formation) ^c	Whole rock	16.6	Ni-bearing ultramafics ^a	Py	13.2
800-52-1 (listwanite) ^c	Whole rock	6.1	Ni-bearing ultramafics ^a	Py	10.9
780-50-3 (serpentinite) ^c	Whole rock	8.6	Ni-bearing ultramafics ^a	Py	11.1
965-Y-45-46 II	Ру	8.5	Ni-bearing ultramafics ^a	Py	9.5
PD503-Y-Py1-2 II	Py	15.4	Ni-bearing ultramafics ^a	Py	11.7
PD404-cm43-Py ^a	Py	13.9	Ni-bearing ultramafics ^a	Py	10.5
PD406-cm45N-Py ^a	Py	14.2	Ni-bearing ultramafics ^a	Py	12.3
PD404-cm45S-Py ^a	Py	18.5	Ni-bearing ultramafics ^a	Py	14.1
PD503-Ym-Py3 ^a	Py	17.3	Ni-bearing ultramafics ^a	Py	12.1
PD404-cm43B-Py1 ^a	Py	12.8	Ni-bearing ultramafics ^a	Py	11.7
965-47E-Ym ^a	Rea, Orp	10.9	Ni-bearing ultramafics ^a	Py	11.7
PD383-cm58-H1 ^a	Rea, Orp	10.5	Ni-bearing ultramafics ^a	Py	11.4
PD633-cm43B-Py1 ^a	Py	12.6	Ni-bearing ultramafics ^a	Py	10.0
G-E-1 ^a	Py	17.0	Ni-bearing ultramafics ^a	Py	10.9
PD383-Y-Py1 ^a	Py	12.6	Ni-bearing ultramafics ^a	Py	6.1
PD633-Ym-59 ^a	Py	12.0	Ni-bearing ultramafics ^a	Py	12.9
97-ZHE-H1 ^a	Py	11.2	Ni-bearing ultramafics ^a	Po	7.8
Au mineralisation	Py	6.3	Ni-bearing ultramafics ^a	Ро	10.6
Au mineralisation	Py	15.3	Ni-bearing ultramafics ^a	Ро	9.1

Table 3. Sulfur isotope composition of sulfides and whole rocks in the Jianchaling area (modified from [21]).

Table 3. Cont.

Sample	Mineral	$\delta^{34} \mathbf{S_{v-CDT}}$ (‰)	Sample	Mineral	$\delta^{34} \mathrm{S_{v-CDT}}$ (‰)
Sm-P (albite porphyry) ^a	Ру	11.0	Ni-bearing ultramafics ^a	Ро	10.0
N124 (albite porphyry) ^a	Py	14.6	Ni-bearing ultramafics ^a	Ро	12.5
L-Py (albite porphyry) ^a	Py	12.9	Ni-bearing ultramafics ^a	Ро	10.7
Porphyritic granite ^a	Py	11.8	Ni-bearing ultramafics ^a	Ро	12.3
Porphyritic granite ^a	Py	12.8	Ni-bearing ultramafics ^a	Ро	11.4
Porphyritic granite ^a	Py	17.5	Ni-bearing ultramafics ^a	Ро	10.9
Porphyritic granite ^a	Po	11.6	Ni-bearing ultramafics ^a	Ро	11.4
Porphyritic granite ^a	Ро	13.7	Ni-bearing ultramafics ^a	Ро	11.8
Porphyritic granite ^a	Ру	9.0	Ni-bearing ultramafics ^a	Ро	11.1
Porphyritic granite ^a	Py	15.2	Ni-bearing ultramafics ^a	Ро	8.2
Porphyritic granite ^a	Py	15.2	Ni-bearing ultramafics ^a	Ро	10.5
Porphyritic granite ^a	Py	15.2	Ni-bearing ultramafics ^a	Ро	11.6
Porphyritic granite ^a	Py	8.3	Ni-bearing ultramafics ^a	Ро	10.2
Porphyritic granite ^a	Py	10.6	Ni-bearing ultramafics ^a	Ру	10.3
Porphyritic granite ^a	Py	9.5	Ni-bearing ultramafics ^a	Py	12.1
Porphyritic granite ^a	Py	9.9	Ni-bearing ultramafics ^a	Py	9.9
Carbonate (Duantouya Fm) ^a	Py	15.0	Ni-bearing ultramafics ^a	Py	11.4
Carbonate (Duantouya Fm) ^a	Py	18.5	Ni-bearing ultramafics ^a	Py	11.5
Carbonate (Duantouya Fm) ^a	Po	18.6	Ni-bearing ultramafics ^a	Py	13.3
Carbonate (Duantouya Fm) ^a	Ро	16.5	Ni-bearing ultramafics ^a	Py	12.3
Carbonate (Duantouya Fm) ^a	Py	10.3	Porphyritic granite ^b	Py	10.0

Superscripts: ^a [41]; ^b [50]; ^c [21]. Abbreviations: Cyp = chalcopyrite; Orp = orpiment; Py = pyrite, Po = pyrrhotite; Rea = realgar.

Sample No.	Sample	Pb	Th	U	²⁰⁸ Pb/ ²⁰⁴ Pb	²⁰⁷ Pb/ ²⁰⁴ Pb	²⁰⁶ Pb/ ²⁰⁴ Pb	(²⁰⁸ Pb/ ²⁰⁴ Pb) _i	(²⁰⁷ Pb/ ²⁰⁴ Pb) _i	(²⁰⁶ Pb/ ²⁰⁴ Pb) _i	Reference
					Au mi	ineralised-rocks					
hjy-1	pyrite	905.000	0.333	0.216	38.757	15.704	18.477	38.757	15.704	18.477	This study
hjy-12	pyrite	170.000	0.070	0.158	37.996	15.539	18.126	37.996	15.539	18.124	This study
hjy-10	pyrite	50.000	0.035	0.539	37.088	15.550	17.600	37.088	15.549	17.579	This study
hjy-20	pyrite	60.900	0.266	0.047	37.180	15.566	17.522	37.177	15.566	17.521	This study
hjy-14	pyrite	53.800	0.041	0.193	37.135	15.577	17.683	37.135	15.577	17.676	This study
hjy-8	pyrite	352.000	0.019	0.068	37.275	15.549	17.811	37.275	15.549	17.811	This study
J3	pyrite	395.000	0.064	0.226	38.547	15.611	18.381	38.547	15.611	18.380	This study
	pyrrhotite				37.493	15.579	17.931				[41]
	pyrite				37.055	15.553	17.863				[41]
	pyrite				36.927	15.530	17.257				[41]
Average					37.545	15.576	17.865	37.711	15.585	17.938	
					Meta-ı	ultramafic rocks					
	listwanite				36.029	15.520	17.952				[41]
	listwanite				38.591	15.521	18.599				[41]
	listwanite				38.604	15.785	19.193				[41]
800-52-1	listwanite	21.500	0.016	0.062	38.069	15.641	18.723	38.069	15.641	18.717	This study
	listwanite				38.150	15.597	18.260				[41]
	antigorite				38.118	15.632	18.390				[41]
	antigorite				38.489	15.616	18.633				[41]
780-50-3	serpentinite	31.500	0.041	0.052	37.748	15.615	18.327	37.747	15.615	18.324	This study
	serpentinite				38.920	15.657	18.553				[41]
	serpentinite				38.032	15.570	18.833				[41]
	meta-ultrama	fic			38.533	15.536	18.281				[58]
Average					38.117	15.608	18.522	37.908	15.628	18.521	

Table 4. Lead isotope composition of ore sulfide and wallrocks at the Jianchaling deposit and in the Bikou Group.

Sample No.	Sample	Pb	Th	U	²⁰⁸ Pb/ ²⁰⁴ Pb	²⁰⁷ Pb/ ²⁰⁴ Pb	²⁰⁶ Pb/ ²⁰⁴ Pb	(²⁰⁸ Pb/ ²⁰⁴ Pb) _i	(²⁰⁷ Pb/ ²⁰⁴ Pb) _i	(²⁰⁶ Pb/ ²⁰⁴ Pb) _i	Reference
					Yu	dongzi Fm					
	Sericite-quartz-se	chist			38.204	15.663	18.432				[41]
	leptynite				39.161	15.481	17.863				[41]
Average					38.683	15.572	18.148				
					Porpl	hyritic granite					
	magnetite				40.359	15.805	21.881				[41]
	pyrrhotite				39.265	15.771	20.922				[41]
	pyrite				38.338	15.765	21.637				[41]
Average	1.5				39.321	15.780	21.480				
					Du	antouya Fm					
H4	shale	115.000	9.120	4.580	39.156	15.768	19.819	39.103	15.764	19.737	This study
960-34-1	dolostone	15.600	0.064	0.250	37.518	15.667	19.164	37.515	15.665	19.132	This study
960-28-1	dolostone	31.800	0.016	0.217	38.170	15.828	22.364	38.170	15.827	22.350	This study
	limestone				38.172	15.601	18.365				[41]
	dolomite				38.783	15.560	18.931				[58]
Average					38.360	15.685	19.729	38.263	15.752	20.406	
					Bi	kou Group					
	spilite				37.621	15.491	17.661				[59]
	pĥyllite				35.328	15.383	16.308				[59]
	phyllite				36.951	15.447	17.076				[59]
	quartz schist				37.866	15.486	17.677				[59]
	phyllite	3.2	1.3	0.4	38.128	15.471	17.644	37.869	15.459	17.399	[59]
	siltstone	3.2	1.3	0.4	38.394	15.552	18.126	38.132	15.540	17.879	[59]
	phyllite	3.2	1.3	0.4	38.064	15.553	18.016	37.803	15.541	17.770	[59]
	metadolerite	11.8	7.58	1.99	40.069	15.928	18.763	39.639	15.911	18.417	[60]
	metadolerite	1.14	0.34	0.12	39.157	15.833	18.017	38.962	15.822	17.806	[60]
	metadolerite	1.39	0.26	0.09	38.945	15.814	17.910	38.823	15.808	17.781	[60]
Average					38.052	15.596	17.720	38.538	15.680	17.842	

Table 4. Cont.

No.	Material	Vein Set	Rb (ppm)	Sr (ppm)	⁸⁷ Rb/ ⁸⁶ Sr	⁸⁷ Sr/ ⁸⁶ Sr	2σ	I _{sr}	References
				Minerali	ised rock				
hjy-1	Pyrite	2	1.15	2.48	-				This study
hjy-10	Pyrite	2	0.561	2.92	0.556233	0.714179	0.001222	0.712613	This study
hjy-12	Pyrite	2	0.671	3.37	0.576465	0.714265	0.001260	0.712642	This study
hjy-14	Pyrite	2	0.885	6.72	-				This study
hjy-20	Pyrite	2	0.405	2.87	0.408670	0.717080	0.000961	0.715929	This study
]3	Pyrite	2	0.531	1.58	0.972553	0.709447	0.002004	0.706709	This study
Average	N = 4							0.711973	2
PD404-43B	Altered dolostone	2	36.20	89.60	13.164780	0.713400	0.002401	0.710107	[41]
PD404-44B-2	Mineralised rock	2	4.20	7.83	19.593624	0.718400	0.003180	0.714026	[41]
PD633-61-11	Quartz	2				0.720900			[41]
PD633-Y-57	Calcite	3				0.717300			[41]
Average	N = 2							0.712066	
				Wall	rocks				
J-15	Porphyritic granite		77.56	36.67	6.136344	0.735543	0.012494	0.718266	[41]
J-16	Porphyritic granite		83.39	48.79	4.959944	0.738159	0.010187	0.724194	[41]
J-18	Porphyritic granite		83.68	31.21	7.789501	0.749689	0.016065	0.727757	[41]
J-19	Porphyritic granite		64.78	30.82	6.100934	0.740388	0.012508	0.723210	[41]
J-60	Porphyritic granite		58.00	220.70	0.760528	0.709768	0.001597	0.707627	[41]
J-66	Porphyritic granite		75.03	79.03	2.754493	0.735919	0.005704	0.728164	[41]
J-69	Porphyritic granite		86.94	35.30	7.143593	0.732922	0.014431	0.712809	[41]
T D 1	Albite-rich		21.20	F 0.00	1.044(01	0 701000	0.000104	0.718959	[41]
L-Py1	intrusive		21.20	58.80	1.044631	0.721900	0.002184	0.718959	[41]
Average	N = 8							0.720123	
J-17	Meta-ultramafics		0.27	17.16	0.045573	0.718455	0.000498	0.718327	[41]
J-55	Meta-ultramafics		1.59	13.47	0.341431	0.704674	0.000827	0.703713	[41]
J-102	Meta-ultramafics		1.33	5.58	0.689301	0.702761	0.001446	0.700820	[41]
J-118	Meta-ultramafics		0.25	35.27	0.020501	0.703870	0.000472	0.703812	[41]
G-E-Mt	Meta-ultramafics		2.92	64.30	0.131452	0.712300	0.000549	0.711930	[41]
Hc-Fe-1(1)	Meta-ultramafics		0.46	6.59	0.202065	0.712800	0.000631	0.712231	[41]
Hg-N-1	Meta-ultramafics		0.33	3.92	0.243695	0.712800	0.000688	0.712114	[41]
97-Hw-1	Meta-ultramafics		1.11	1.01	5.358452	0.729600	0.006523	0.720628	[41]
J-45	Meta-ultramafics		1.90	4.58	2.842641	0.725862	0.002510	0.722476	[41]
J-50	Meta-ultramafics		0.53	2.11	2.218363	0.717723	0.001553	0.715675	[41]
J-42-3	Meta-ultramafics		0.28	0.69	4.384971	0.726283	0.002460	0.722971	[41]
Y-Mg	Meta-ultramafics		0.33	3.66	1.153334	0.713400	0.000714	0.712665	[41]
G-Mg	Meta-ultramafics		0.29	2.57	1.666748	0.713400	0.000817	0.712480	[41]
780-50-3	Meta-ultramafics		1.57	2.97	1.531251	0.719517	0.003140	0.715206	This study
800-52-1	Meta-ultramafics		0.10	0.80	0.376258	0.711025	0.000895	0.709965	This study
Average	N = 15							0.713001	5

Table 5. The Sr isotope ratios of sulfides and wallrocks at the Jianchaling Au deposit and in the Bikou Group.

No.	Material	Vein Set	Rb (ppm)	Sr (ppm)	⁸⁷ Rb/ ⁸⁶ Sr	⁸⁷ Sr/ ⁸⁶ Sr	2σ	I _{sr}	References
PX406-Y-43W	Duantouya Fm		1.69	108.00	0.447747	0.719000	0.000499	0.718872	[41]
G-E-1	Duantouya Fm		8.89	101.00	2.691745	0.723300	0.000720	0.722582	[41]
H4	Duantouya Fm		105.00	57.20	5.312134	0.734027	0.010856	0.719034	This study
960-28-1	Duantouya Fm		0.03	17.90	0.004370	0.720582	0.000493	0.720570	This study
Average	N = 4							0.720265	5
2000224	Bikou Group		19.55	594.59	1.463281	0.704471	0.000507	0.704203	[61]
2000225	Bikou Group		5.33	118.58	2.088492	0.704899	0.000538	0.704533	[61]
2000226	Bikou Group		3.95	41.58	4.601666	0.703211	0.000720	0.702438	[61]
2000228	Bikou Group		3.87	18.15	10.756975	0.703534	0.001312	0.701797	[61]
2000230	Bikou Group		3.73	39.70	4.931685	0.702650	0.000715	0.701884	[61]
2000231	Bikou Group		23.01	243.96	5.126925	0.713325	0.000732	0.712556	[61]
BK1	Bikou Group		23.40	563.00	2.342130	0.710419	0.000536	0.710080	[60]
BK2	Bikou Group		0.21	111.00	0.110386	0.707353	0.000475	0.707338	[60]
BK3	Bikou Group		0.13	143.00	0.054847	0.706668	0.000474	0.706661	[60]
Average	N = 9							0.705721	

Table 5. Cont.

Table 6. The Nd isotope ratios of sulfides and wallrocks at the Jianchaling deposit.

No.	Sample	Stage	Sm	Nd	¹⁴³ Nd/ ¹⁴⁴ Nd	¹⁴⁷ Sm/ ¹⁴⁴ Nd	(¹⁴³ Nd/ ¹⁴⁴ Nd)i	f _{Sm/Nd}	ε _{Nd} (198)
				Si	ulfides				
hjy-1	Pyrite	2	0.052	0.167	0.512426	0.1883	0.512182	-0.04	-3.9
hjy-10	Pyrite	2		0.018					
hjy-12	Pyrite	2	0.085	0.747	0.513300	0.0688	0.512453	-0.65	1.4
hjy-14	Pyrite	2	0.012	0.041	0.511941	0.1769	0.511712	-0.10	-13.1
hjy-20	Pyrite	2	0.06	0.425	0.511500	0.0853	0.511389	-0.57	-19.4
J3	Pyrite	2	0.026	0.194	0.511481	0.0810	0.511376	-0.59	-19.6
Average	N = 5						0.511823		-10.9
				Wa	allrocks				
H4	Duantouya Fm		3.79	16	0.512051	0.1432	0.511865	-0.27	-10.1
960-34-1	Duantouya Fm		0.216	1.29	0.512012	0.1012	0.511881	-0.49	-9.8
960-28-1	Duantouya Fm		0.039	0.129	0.512350	0.1828	0.512114	-0.07	-5.3
Average	N = 3						0.511953	-0.28	-8.4
780-50-3	Meta-ultramafic rock		0.058	0.242	0.512325	0.1449	0.512137	-0.26	-4.8
800-52-1	Meta-ultramafic rock		0.004	0.013	0.512393	0.1860	0.512152	-0.05	-4.5
Average	<i>N</i> = 2						0.512144	-0.16	-4.7

6. Discussion

6.1. Sources of Metals and Mineralising Fluids

6.1.1. Oxygen and Hydrogen Stable Isotopes

The oxygen and hydrogen stable isotopes are often used to indicate the source of mineralised hydrothermal fluids [62]. However, the overlap in the metamorphic and magmatic fields in δD versus $\delta^{18}O$ diagrams creates uncertainty in deciphering the genesis of mineralisation [63]. This is the case for the mineralising fluid at Jianchaling, which has been interpreted as being magmatic [64,65], meteoric [53], mixed metamorphic and magmatic [42,43,51], and primarily metamorphic [39]. This places a significant doubt on the usefulness of the δD and $\delta^{18}O$ values in directly pinpointing the origin of the mineralising fluid at Jianchaling.

Using fluid-inclusion homogenisation temperatures and detailed paragenetic studies by Yue [39] (Table 2), the $\delta^{18}O_{water}$ values were calculated using the quartz–water equation by [54] and carbonate–water equation by [55]. The calculated quartz-carbonate δ^{18} O values for the hydrothermal fluid associated with the first vein set range from 7.5 to 17.7%, the mineralised second vein set have δ^{18} O values between 5.7 and 11.1‰, and the third vein set range from 1.3 to 4.3‰ (Table 2). Using this data, the δ^{18} O and δ D values for first vein set plot in the magmatic field and close to the metamorphic field (Figure 6). However, assuming that magmatic fluids are generated above the lowest eutectic point at temperatures above 573 °C [66], continuous cooling and water-rock reactions result in reduction of $\delta^{18}O_{water}$ values during the crystallisation of hydrothermal minerals such as quartz and alkali feldspar [66]. Furthermore, magmatic fluids have average $\delta^{18}O_{water}$ values of 18.8% at temperatures of 355 °C [39], which is the average homogenization temperature of fluid inclusions from first vein generation (Table 2). This means that the initial $\delta^{18}O_{water}$ value must have been higher than 18.8% at temperatures above 355 °C, which is higher than the 5.7–11.1% range for the mineralised second vein set. In addition, the Late Triassic to Early Jurassic plutons in the southern part of the Qinling Orogen commonly have δ^{18} O values of <18.8‰. The high δ^{18} O values for the early hydrothermal fluids are consistent with a major contribution from metamorphic sources (rather than a minor contribution as suggested by Figure 6). This interpretation is supported by the low salinity and high CO₂ content of fluid inclusions in the first vein set [7,39]. Furthermore, the δ^{18} O values for quartz in granite porphyry and aplite in the area range between 11.3 and 14.5% [67]. This indicates that the $\delta^{18}O_{water}$ value of 10.0% calculated at a temperature of ~573 °C for the first vein set cannot exceed the $\delta^{18}O_{water}$ value of 13.2% for the second vein set that crystallised at ~300 °C (as determined from the fluid inclusion homogenisation temperatures). In addition, the regional metamorphism in the Jianchaling area is at the greenschist facies [68], further supporting a ~ 300 °C temperature for the crystallisation of the mineralised second vein set.

The 3rd vein $\delta^{18}O_{water}$ values are between 1.3 and 4.3‰, with a δD value of -81%, they are plot in the meteoric field in Figures 6 and 7. The $\delta 18O$ water values of carbonate-quartz from the mineralised vein set vary between 5.7 and 11.1‰, which are values between those for the first and third vein sets. Exactly, on the base of the $\delta^{18}O_{water}$ values (5.7–11.1‰), we think the 2nd vein set is similar to lode gold deposits and add the possible range of the 2nd vein set in Figure 6 that based on the feature of many lode gold deposits worldwide. This is indicative of the mixing of metamorphic and meteoric fluids during mineralisation (Figure 6).

The δ ¹⁸O values for quartz from lode-gold deposits around the world are higher than 10‰ with the values for the mineralising fluids ranging between 5 and 25‰ [6]. The δ ¹⁸O_{water} values of the gold deposits on the Jiaodong Peninsula of eastern China range from 4.9 to 10.9‰, with corresponding δ D values of -78 to -101‰ (Figure 6) [69,70]. The δ ¹⁸O and δ D values for mineralising fluids at Jianchaling plot in the lode-gold deposit field (Figure 6; Table 2; [6]). Figure 6 also shows that the first vein set containing higher δ ¹⁸O and δ D values are sourced from metamorphic fluids. In contrast, the mineralised second vein set are similar lode gold deposits, and the fluids related to the third vein set have (again) a composition close to that of meteoric fluid.

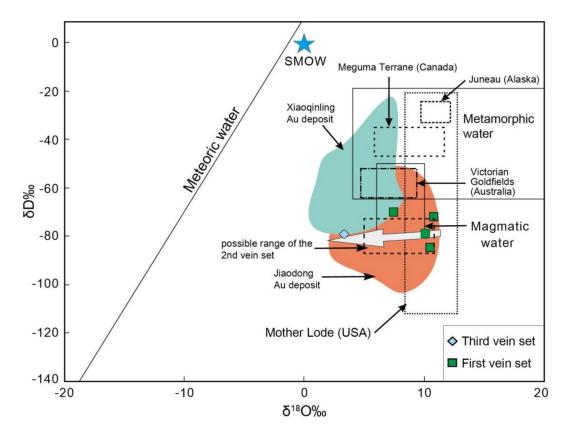


Figure 6. Stable isotope (δD versus $\delta^{18}O$) plot for mineralised fluids at the Jianchaling Au deposit [71]. Included are the fields for the Xiaoqinling and Jiaodong gold deposits [69]; Juneau gold belt, Mother Lode, Victorian Goldfields and Meguma Terrace Au deposits [6].

6.1.2. Carbon and Oxygen Isotopes

The $\delta^{13}C_{CO2}$ values determined for quartz-carbonate from each of the vein sets are essentially the same, hovering around approximated zero (Figure 7). The corresponding $\delta^{18}O_{SMOW}$ values show a significant trend with the first vein set being approximately 15%, the mineralised second vein measuring around 9%, and the third being around 3‰ (Figure 7). The implication for this trend is that all of the vein sets have similar sources for carbon with a gradual decrease in available oxygen in the hydrothermal fluid with time. Another implication is that there was only one hydrothermal fluid that has changed in composition with time.

The $\delta^{13}C_{CO2}$ values for the hydrothermal fluids related to the three vein sets are significantly higher than the values for organic matter (averaging -27%) [62], the content of CO₂ in the atmosphere (-11 to -7% [62]), freshwater carbonate (-20 to -9% [72]), magmatic rocks (-30 to -3% [72]), the continental crustal (-7% [73]), and the mantle (-7 to -5% [72]). However, marine carbonate (i.e., -3 to 2% $\delta^{13}C_{CO2}$ [72]) and carbonate in the wall rock (i.e., -0.4% to 2.3% $\delta^{13}C_{CO2}$; Table 2), are similar in $\delta^{13}C_{CO2}$ composition. This indicates a local source for the hydrothermal fluids related to each of the vein sets. In addition, the close relationship between the ultramafic rocks and the vein sets shown in Figure 7 indicates that the ultramafic rocks are also local sources for the carbon and oxygen in the hydrothermal fluids.

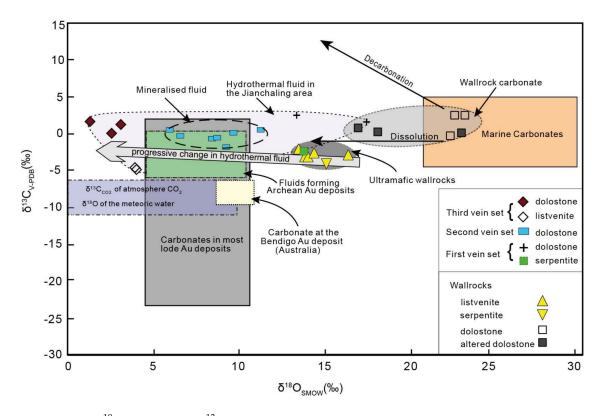


Figure 7. $\delta^{18}O_{SMOW}$ versus $\delta^{13}C_{V-PDB}$ plot for the Jianchaling Au deposit [21]. Data sources: greenschist facies carbonate rocks after [74]; reduced C in sedimentary and metamorphic rocks after [75]; carbonates in most orogenic Au deposits after [76].

On the other hand, the $\delta^{13}C_{CO2}$ and $\delta^{18}O_{SMOW}$ of the mineralised fluids overlap with those of the forming fluid of Archean Au deposits (Figure 7). $\delta^{18}O_{SMOW}$ value of the third vein sets are similar with the meteoric water, indicating meteoric water mixed into the fluid system in late stage.

6.1.3. Sulfur Isotopes

The δ^{34} S signatures for various in the MLY and others from throughout the world are presented in Figure 8. The figure shows that most of the sulfides in the MLY have δ^{34} S signatures greater than zero, are predominantly higher than those for orogenic Au deposits throughout the world, and higher than those for basaltic magmatic rocks, overlap the sulfide values in ultramafic rocks, and are coincide with the high end of sulfides in metamorphic rocks. In fact, the δ^{34} S signatures for the metamorphic rocks are the best fit, which includes the wallrocks at Jianchaling (Table 3). Such heavy δ^{34} S values for the sulfides at Jianchaling is consistent with derivation from sulfates that are characterised by heavy δ^{34} S values [77,78]. It is envisages that organic material derived by the metasedimentary rocks in the study area would have reacted with sulfate resulting in the crystallisation of sulfides [78,79]. Examples of such reducing agents are CH₄, C₂H₆, H₂S and graphite, which have been detected in fluid inclusions using Laser Raman [39]. The conversion of sulfate to sulfide will proceed in the following chemical reaction:

$$CH_4 + SO_4^{2-} \leftarrow \rightarrow H_2S + H_2O + CO_3^{2-} \text{ and } H_2S + Fe^{2+} \leftarrow \rightarrow FeS + 2H^+$$
(1)

Therefore, the Neoproterozoic metamorphic rocks in the MLY are likely sources for sulfur in the Jianchaling area.

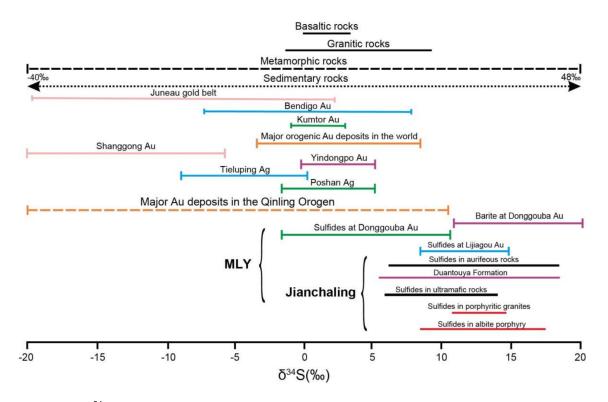


Figure 8. δ^{34} S values for hydrothermal sulfides from the Jianchaling Au deposit. Data sources: Basaltic rocks, Granitic rocks, Metamorphic rocks and Sedimentary rocks after [62,72,80]; Juneau gold belt, Bendigo Au, Kumtor Au and Major orogenic Au deposits in the world after [12,81]; Shanggong Au and Tieluping Ag after [44]; Yindongpo Au after [66]; Poshan Ag after [82]; Major Au deposits in the Qinling Orogen [24].

6.1.4. Lead Isotopes

The country rocks in the Jianchaling area have a wider range in Pb isotopic (²⁰⁶Pb/²⁰⁴Pb, ²⁰⁷Pb/²⁰⁴Pb and ²⁰⁸Pb/²⁰⁴Pb) ratios than the sulfides in the Jianchaling mineralisation (Table 4). The range of the Pb-isotopes is shown in Figure 9 that includes trends defining tectonic settings [83]. The range of Pb-isotopes shown in Figure 9 correlates with defined tectonic setting of orogenic and upper crustal fields, even though there is a limited number of 10 data points. This is in agreement with the observations made in the discussion section above.

Furthermore, these Pb signatures of the sulfides from the mineralisation overlap those of the ultramafic rocks and the Bikou Group, indicating that the units are possible sources for the sulfides in the mineralisation. In contrast, the distinct incongruence of the sulfides from the mineralisation and the granitic rocks indicates that the granitic rocks are not the source for the sulfides in the mineralisation (Figure 9).

6.1.5. Strontium and Neodymium Isotopes

The $I_{Sr}(t)$ values of sulfide samples from the Jianchaling deposit range from 0.706709 to 0.715929 (Table 5; Figure 10a). These values are higher than the mean values (0.703 to 0.705) established for mantle-derived oceanic basalt (MORB, OIB, IAB) and continental basalt, and are suggestive of a crustal source [84,85].

The $I_{Sr}(t)$ values for the sulfide samples from the deposit are also lower than those for the Duantouya Formation (Table 5), and the (¹⁴³Nd/¹⁴⁴Nd)i sulfide ratios for the mineralised second vein set overlap with those of the Duantouya Formation and the meta-ultramafic rocks (Table 6; Figure 10a,b). The ε Nd (t) and Isr (t) of sulfide are indicative of a crustal source and dominated with mature Sr-Nd character (Tables 5 and 6; Figure 10a), and ε Nd (t) overlaps with meta-ultramafic field,

indicated that the Duantouya may play an important role during the ore formation and there may exist minor ultramafic source been involved in the ore fluid.

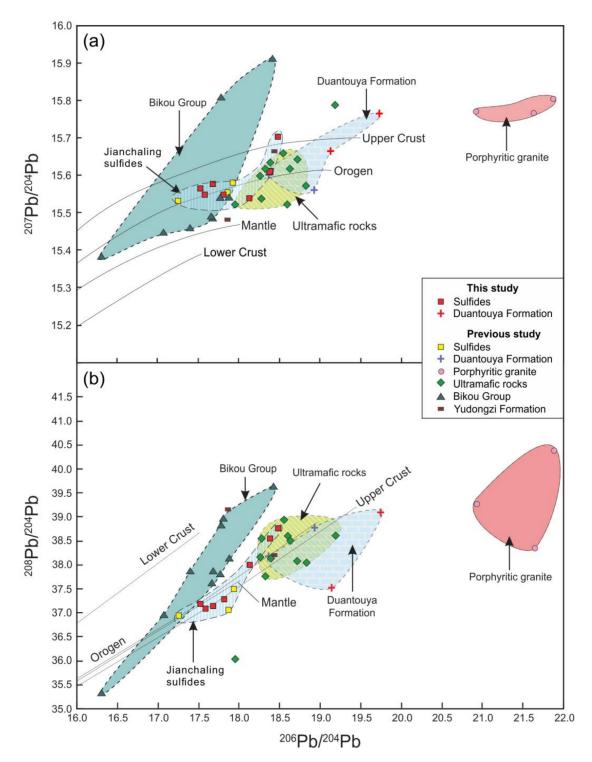


Figure 9. The plumbotectonic model for Pb isotopic systematics for the Jianchaling Au deposit. (a) 206 Pb/ 204 Pb versus 208 Pb/ 204 Pb; (b) 206 Pb/ 204 Pb versus 207 Pb/ 204 Pb.

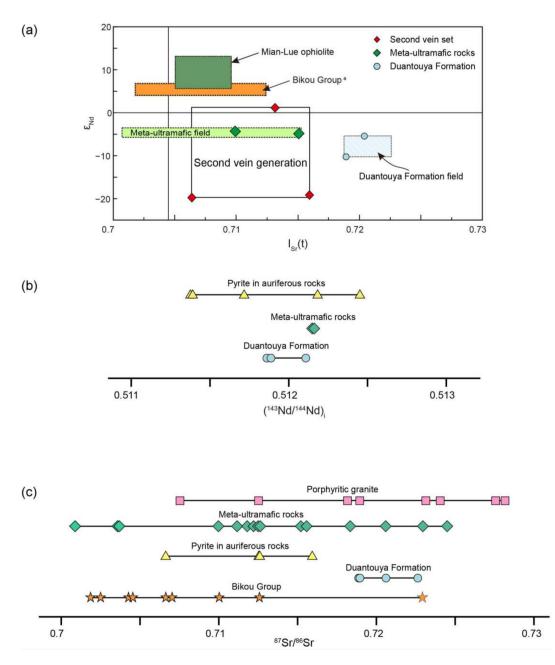


Figure 10. Isotope systematics for the Jianchaling Au deposit showing: (**a**) $I_{Sr}(t)$ versus $\varepsilon_{Nd}(t)$ plot (data for the Bikou Group is from [86]); (**b**) (¹⁴³Nd/¹⁴⁴Nd)i plot; and (**c**) ⁸⁷Sr/⁸⁶Sr plot. The range of the Mian-lue ophiolite is from [23]. T = 198 Ma.

The Bikou Group has $I_{Sr}(t)$ values that overlap with those from the Mian-Lue ophiolite, meta-ultramafic rocks, partly with the second vein set, and are lower than those from the Duantouya Formation (Figure 10a; Tables 5 and 6) [60,61,86]. Although the pyrite in audiferous rocks has $I_{Sr}(t)$ values that overlap with porphyritic granite partly (Figure 10c), it is unlikely to come from porphyritic granite, the evidence is mainly from lead isotopes, as mentioned above.

These relationships can be explained with a mixing model where the mineralised sulfide-bearing fluids interacted with lithological units (Figure 10). This interpretation is supported by the conclusion drawn from the Pb isotope data collected from the Huachanggou gold deposit in the northwestern part of MLY [25,59] (Figure 1c).

6.2. Mineralisation Type

Various mineral-types have been proposed for the Jianchaling Au deposit. These include: (1) a magmatic hydrothermal source associated with Triassic magmatism [65]; (2) Carlin-like affiliations [8]; (3) orogenic affiliations [43]; and (4) sourced from structurally controlled altered rocks [40,53].

Fuchsite separated from the mineralised second vein set yield a well-defined 40 Ar/ 39 Ar isotopic age of 198 ± 2 Ma, which suggests that the mineralisation took place during the Triassic Indosinian Orogeny [39]. Implications of this are that the Neoproterozoic (ca. 927 Ma) magma-hydrothermal event associated with the emplacement of ultramafic intrusions. Triassic (ca. 216 Ma) porphyritic granites in the area are ~18 Ma older than the Jianchaling mineralisation, again indicating that the granites are not related to the gold mineralisation.

The C, H and O stable isotopes indicate that the ore-forming fluids are predominantly metamorphosed in origin and contaminated with meteoric fluid during the deposition of the gold at Jianchaling. On the other words, the mineralising fluid is probably sourced from the metamorphic dehydration of carbonate rocks in the Duantouya Formation, and gradually mixed with meteoric water. The ⁸⁷Sr/⁸⁶Sr, I_{Sr}(t), ε_{Nd} , Pb and S isotopic data (Figure 10) confirm that the sulfides in the mineralised are sourced from the metamorphosed Duantouya Formation, Bikou Group and minor ultramafic rocks.

From the discussion above, combining the comparison of the geological characteristics between Jianchaling and Orogenic gold deposit (Table 7), the Jianchaling deposit is here interpreted as an orogenic-type deposit and summarised in Figure 11.

	Orogenic Gold Deposit ^a	Jianchaling Gold Deposit ^b
Tectonic setting	Subduction hyperplasia, continental collision, intracontinental strike-slip and intracontinental compressional orogenic belt	Continental collision
Ore-bearing terrane	Metamorphic terrane	Metamorphic and sedimentary terrane
Ore-controlling lithology	Ultramafic volcanic rocks, intrusive rocks, miscellaneous sandstones, slate	Ultramafic and dolomite
Metamorphism of the host rock	Greenschist facies (low green schist to low granulite)	Greenschist to low amphibolite facies
Ore-controlling structures	In the secondary or lower faults of the super-lithospheric fracture zone, the ore-forming structures is mainly the high angle oblique slip belt, the reverse overthrust belt, and also the transverse fracture, the ductile-brittle zone	In the brittle fracture of the ductile shear zone (high angle thrust belt in the basement fault zone
Ore and gangue minerals	Mostly pyrite	Arsenic pyrite, marcasite, arsenopyrite, orpiment, realgar, quartz and calcite
Ore type	Quartz vein and altered rock	Altered rock

Table 7. Geological characteristics comparison between the Jianchaling and Orogenic gold deposit.

	Orogenic Gold Deposit ^a	Jianchaling Gold Deposit ^b
Metallogenic hydrothermal fluids characteristics	Aqueous solution with low salinity and low density, containing $CO_2 \pm CH_4 \pm N_2 \pm$ H_2S . The fluids inclusion type has H_2O - CO_2 , rich CO_2 (with an unquantifiable CH_4 and a small amount of H_2O) and gas-liquid two-phase H_2O inclusions	Aqueous solution with low salinity and low density, containing $CO_2 \pm CH_4 \pm H_2S$. The fluids inclusion type has NaCl-H ₂ O, CO ₂ -H ₂ O-NaCl \pm CH ₄ and pure CO ₂ -CH ₄
Metallogenic fluids salinity	3–12 wt % NaCl equiv.	0.4–15.6 wt % NaCl. equiv.
Main metallogenic temperature	$350\pm50~^\circ\mathrm{C}$	200–320 °C
Mineralization pressure	50–400 MPa	117–354 MPa
Initial metallogenic fluids feature	Metamorphic fluids	Metamorphic fluids

Table 7. Cont.

Superscripts: a [63,87,88]; b [21,39].

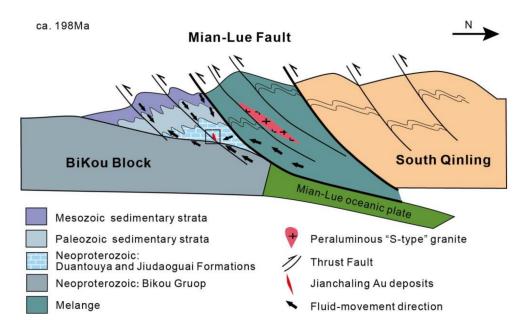


Figure 11. Tectonic model for the Jianchaling Au deposit [39].

7. Conclusions

Hydrogen, oxygen and carbon isotopic systematics indicate that the ore-forming fluids at the Jianchaling Au deposit progressively evolved from an early stage represented by the first vein set associated with deformation and metamorphism of the country rocks. This was succeeded by a late tectonic stage represented by the mineralised second vein set sourced from a mixed metamorphic and meteoric fluid, and progressed to a late stage represented by the third vein set. The S-Pb-Sr-Nd isotopic data indicate that the mineralising fluids were sourced locally around Jianchaling. From these observations, the Jianchaling Au deposit is interpreted as an orogenic Au deposit formed during the Mesozoic Indosinian Orogeny.

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