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熊耳山矿集区蒿坪沟 Ag-Au 多金属矿床绿泥石特征及其找矿 意义

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Characteristics of chlorites from the Haopinggou Ag–Au polymetallic deposit in the Xiong'ershan ore concentration area and its exploration implications

Abstract: [**Objective**] The Haopinggou Ag-Au polymetallic deposit is a typical intermediate-sulfidation epithermal deposit in the Xiong'ershan ore concentration area. Ag-Pb-Zn mineralization mainly occurs in steeply dipping veins and breccia matrix. The relationship between large-scale Pb-Zn mineralization and widely developed alteration minerals remains unclear. [**Methods**] In order to discuss chlorite's significance related to Pb-Zn mineralization, chlorite composition in the Haopinggou Ag-Au polymetallic deposit has been analyzed by field geological observation and electron microprobe analysis (EMPA) in this paper. [**Results**] Three types of chlorite were observed in the deposit, occurring in altered wall rocks (Type I), in(with) Pb-Zn sulfides (Type II), and in(with) the breccia matrix (Type III). All three types of chlorite are prochlorites and fall within the compositional range of Fe-rich chlorite, indicating that they could be formed in a partially reducing acidic environment. Fe²⁺ for Mg²⁺ is the primary substitution in chlorite lattice, suggesting a close association between chlorite formation and mafic wall rocks. Based on the corrected chlorite geothermometer, these chlorites formed under aluminum-saturated conditions in the medium to low-temperature range of 196-239 °C. The temperatures of chlorites associated with mineralization (Types II and III) are higher than those in chlorites around quartz veins (Type I). It is believed that during the mineralization process, the hydrothermal fluids evolved from acidic to nearly neutral conditions as the temperature gradually decreased. The initial acidic environment facilitated interaction between water and rocks,

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promoting the dissolution of surrounding rocks and providing space for the further precipitation of metal sulfides. The evolution of ore fluid properties also corresponds to the deposition process of Ag-Pb-Zn. The genesis of chlorite in the deposit is well-correlated with the ore-forming and holds significant prospecting value. (1) Type I chlorites mainly develop on both sides of quartz veins, formed by the dissolution and metasomatism of basic wall rocks by ore-bearing hydrothermal fluids. Type I chlorite's Fe and Mg components are mostly derived from the wall rocks. Although this type does not contain mineralization, it can be used to trace veins. (2) Type III chlorites reflect the migration process of ore-bearing hydrothermal fluids carrying dissolved minerals (biotite/clinopyroxene), which precipitate with changes in the physicochemical environment. Type III chlorite's Fe and Mg components are mainly introduced by ore-bearing hydrothermal fluids. This type of chlorite fills the intergranular pore spaces between minerals and easily replaces minerals such as biotite and hornblende, exhibiting apparent mineral alteration features in hand specimens, which is beneficial for prospecting.(3) The formation mechanism of Type II chlorites includes the possibilities mentioned above. This type of chlorite is formed by complete dissolution and metasomatism of cement in ore-bearing hydrothermal fluids, forming fine-grained cryptocrystalline chlorite fillings in breccia rocks. Ore-bearing hydrothermal fluids and wall rocks both contribute Fe and Mg in the chlorites. Hand specimens of Type II chlorite are dark green, disseminated and filled in the matrix, making it easy to distinguish. The chemical characteristics of Type II chlorites are similar to those of chemical characteristics of granite-related deposits, implying the contribution of magmatic fluids to ore-forming fluids. [Conclusion] The Haopinggou Ag-Au polymetallic deposit contains three types of chlorites. Their chemical characteristics all reflect an acidic and reducing metallogenic environment. In cation exchange, the primary substitution is Fe for Mg, and other substitutions are insignificant. The effect of Fe/(Fe+Mg) must be eliminated in order to calculate the temperature of such deposits using chlorite geothermometers. The formation mechanism of the three chlorites is closely related to mafic wall rock, and their chemical properties suggest the involvement of magmatic fluids in ore-forming fluids. [Significance] These three types of chlorite are well-matched with intense Ag-Pb-Zn mineralization and can serve as key indicators for locating Pb-Zn veins. Keywords: Polymetallic deposit; prospecting significance; Chlorite; EMPA; mineralization; Geochemistry; Xiong'ershan.

摘 要:为理清蒿坪沟 Ag-Au 多金属矿床中多阶段矿化与热液蚀变之间的关系,文章选取与铅锌成矿阶段密切相关的绿泥石进行野外观察及电子探针分析。文章将蒿坪沟 Ag-Au 多金属矿床中的绿泥石分为 3类: Ⅰ型分布在石荚脉两侧的围岩中; Ⅱ型呈细粒、隐晶质填充于隐爆角砾岩基质; Ⅲ型与铅锌硫化物 共生、或以蠕虫状广泛分布在石荚颗粒间隙中。3种类型绿泥石均为斜绿泥石,并落在了铁镁绿泥石的范 围内,指示其形成于偏还原的酸性环境中;在阳离子置换中,主要发生了 Fe²⁺对 Mg²⁺的置换,其余置换作 用均不明显;3种绿泥石形成与镁铁质围岩关系密切。由校正后的绿泥石地质温度计估算出3种类型绿泥 石的形成温度为 196~239 ℃,属于中一低温热液蚀变范围。3类绿泥石与蒿坪沟 Ag-Au 多金属矿床银铅锌 成矿阶段相匹配,对进一步找矿勘查具有重要意义。绿泥石化学特征表明岩浆热液参与了成矿流体的形 成,绿泥石形成于熊耳山矿集区早白垩世大规模岩浆-成矿时期。

关键词:多金属矿床;找矿;绿泥石;电子探针;成矿;地球化学;熊耳山 中图分类号:P614 文献标识码:A 文章编号:1006-6616(2024)01-0129-18 DOI: 10.12090/j.issn.1006-6616.2023121

0 引言

绿泥石属于结构为2:1:1(也记作TOTO)的层 状镁、铁铝硅酸盐矿物,通常是热液蚀变作用的标 志产物,与金、铜和铀等矿床密切相关(Lowell and Guilbert, 1970; Cathelineau and Nieva et al., 1985; Pacey et al., 2020)。绿泥石可以稳定存在于较宽的温度 范围内,且其成因矿物学特征能反映流体性质和 水 -岩反应环境,因此备受矿床学家的重视(Laird, 1988;郑伟等,2013; Bourdelle and Cathelineau, 2015; 戴朝成等,2017; Zhang et al., 2022)。绿泥石的化学成 分和晶体结构在不同的成矿环境中呈现规律性的 变化,常用的通式为($R_x^{2+}R_y^{3+}\square_{6xy}$)^N(Si₂ R_{4-z}^{3+})^N₄O₁₀(OH)₈, 其中 R²⁺代表二价阳离子, R³⁺代表三价阳离子, □代 表八面体空位(Zane and Weiss, 1998); 伴随 3 种主要 的 阳离子 替换:①Fe²⁺⇔Mg²⁺, ② 契尔马克替换 Al^NAl^N⇔Si(Mg²⁺, Fe²⁺), ③二八 -三八面体替换

3(Mg²⁺, Fe²⁺)⇔□ + 2Al^N。结构的多变性表明绿泥 石对成矿时期的温度、压力和全岩成分环境非常敏 感 (Inoue,1995; Inoue et al., 2009; Bourdelle et al., 2013),因此常作为一种重要的标型矿物,用来揭示 金属矿床的成矿作用(刘燚平等,2016;方维萱等, 2017; 张娟等, 2021; Zhang et al., 2022)。针对形成温 度与绿泥石结构转换之间的关系, Cathelineau and Nieva(1985)根据绿泥石 Al^W含量与温度之间的良好 线性关系拟合提出了经验性绿泥石温度计(或铝成 分温度计),后经其他学者校正,衍生出了一系列计 算绿泥石形成温度的线性公式(Cathelineau, 1988; Jowett, 1991; Zane and Fyfe, 1995)。迄今为止, 经验 性绿泥石温度计在矿床研究方面运用广泛(王小雨 等, 2014: Dora and Randive, 2015: 周栋等, 2018: 葛祥 坤等, 2020; Zhang et al., 2022), 但针对不同种类的矿 床其适用性还存在较多争议(刘燚平等,2016)。

蒿坪沟 Ag-Au 多金属矿床位于华北克拉通南 缘熊耳山矿集区西部下峪矿田内,属东秦岭成矿带 中少见的角砾岩-斑岩-浅成低温 Au 多金属矿床 (Tian et al., 2023)。依据原生硫化物组合的硫化状 态的"三分法"方案(Hedenquist et al., 2000), 蒿坪沟 Ag-Au多金属矿床又可归类为中硫型(Sillitoe and Hedenquist, 2005; Tian et al., 2023)。已有研究将该矿 床成矿过程大致划分为3个阶段:金-石英-菱铁矿 阶段、石英-铅锌硫化物阶段、石英-碳酸盐阶段 (陈衍景等, 2003; 程广国, 2013; Li et al., 2013; 徐进 鸿, 2021; Tian et al., 2023)。Li et al(2016)认为金-石 英-菱铁矿阶段与陆-陆碰撞过程中碳酸盐-页岩-燧石建造变质脱水有关,石英-硫化物阶段则以富 集银的铅锌矿化为主,与晚中生代岩浆事件有关。 Tian et al(2023)通过与铅锌同成矿期独居石和磷钇 矿年龄范围 125.7±1.8~123.3±1.7 Ma 对成矿时代进 行了限定,成矿与同期次的花岗斑岩有关。成矿流 体总体被认为属中一低温、中一低盐度流体(陈衍 景等, 2003; Chen et al., 2004; 高建京等, 2010; Li et al., 2013; Han et al., 2014; 徐进鸿, 2021), 主成矿期和 成矿晚期具有大气水混合的特征(Li et al., 2016;徐 进鸿, 2021)。已有的研究成果对蒿坪沟 Ag-Au 多金 属矿床多期多阶段的成矿过程进行了限定,但并未 深入探讨矿化阶段产生的大规模热液蚀变矿物(Li et al., 2013; Tian et al., 2023), 导致矿化过程与热液蚀 变矿物之间的关系尚不明确,一定程度上制约了对 矿床成因认识的深化(吕志成等, 2022; 袁航等,

2022)。作为蒿坪沟 Ag-Au 多金属矿床中重要的蚀 变矿物,绿泥石较为完整地记录了蚀变及矿化过 程,对其开展成分及结构研究可以反演银铅锌成矿 作用过程。基于野外地质调查,显微镜下观察和电 子探针分析,文章对蒿坪沟 Ag-Au 多金属矿床中绿 泥石的类型、组分、形成温度、环境及形成机制进 行了系统研究,以期约束蒿坪沟 Ag-Au 多金属矿床 成矿流体的演化特征及银铅锌沉淀的物理化学环 境,并对成矿期流体性质和绿泥石形成机制进行了 初步探讨,从而为探讨矿床成因及找矿勘查工作提 供新的思路与线索。

1 地质背景

熊耳山矿集区地处华北克拉通南缘,位于华北 地台与秦岭造山带的结合部(图 1a),经历了太古代 陆核形成与增生、元古代大陆裂解与俯冲拼贴、古 生代陆陆碰撞及中生代陆内活动等重要地质演化 过程,形成了多阶段构造-岩浆-成矿格局(陈衍景 等,2003; Chen et al., 2004;李诺等,2008; Mao et al., 2010; Dong et al., 2016; Li et al., 2018; 张国伟等, 2019; Wang et al., 2021)。熊耳山矿集区南部以近东 西走向的马超营断裂为界,北部以洛宁断裂为界, 拆离断层主要位于太华群与熊耳群之间(图 1b)。 整个矿集区东西长约 80 km,南北宽 15~40 km,面 积约 2000 km²。

熊耳山矿集区出露地层单元简单,主体分为上 下两套(Li et al., 2013; 唐克非, 2014)。下部为前寒 武纪的结晶基底,由太华群角闪岩、TTG片麻岩和 富钾片麻岩组成(第五春荣等, 2007; Diwu et al., 2014; Wang et al., 2020); 上部为不整合覆盖于基底 之上的盖层,包括古元古代熊耳群火山岩、中一新 元古代官道口群碳酸盐岩(Zhao et al., 2005; Wang et al., 2010)。洛宁断裂为生长正断层, 控制了新生代 盆地的分布。北倾的马超营断裂在古生代和中生 代活动,伴生一系列次级北东向陡倾左旋断裂系 统,为熊耳山矿集区主干控矿构造(图 1b;张元厚 等, 2006; Han et al., 2009)。燕山期花岗岩在区内广 泛分布,侵入于太华群片麻岩和熊耳群火山岩中, 岩体产状及成因类型多样,与熊耳山矿集区晚中生 代大规模成矿作用具有密切联系(毛景文等,2009; Mao et al., 2010; Wang et al., 2015; Li et al., 2015, 2018; Yang et al., 2019; Zou et al., 2019; Hu et al.,



a一华北克拉通南缘在中国东部的位置; b一熊耳山矿集区大地构造简图及矿产分布图

图1 华北克拉通南缘熊耳山矿集区地质简图(底图据 Tian et al., 2023 修改)

Fig. 1 Simplified geological map of the Xiong'ershan ore concentration area along the southern margin of the North China Craton(Base map modified after Tian et al., 2023)

(a) The inset showing the tectonic location of the southern margin of the North China Craton in eastern China; (b) Geological map of the Xiong' ershan ore concentration area, showing the distribution of the major deposits

2020)。熊耳山矿集区晚中生代发育大规模脉状 金、银铅锌矿床,同时伴有一定规模的斑岩型和隐 爆角砾岩型金矿化。金矿床主要以石英脉型、蚀变 岩型和角砾岩型为主,银铅锌矿床则以陡倾多金属 硫化物脉的形式产出。

2 矿床地质

蒿坪沟 Ag-Au 多金属矿床位于熊耳山矿集区 下峪矿田的西北部(图 1b),与后张沟 Ag(Cu)矿床、 沙沟 Ag-Pb-Zn 矿床、铁炉坪 Ag-Pb(Cu)等矿床相邻 (图 2)。矿区主要发育脉状 Ag-Au 多金属矿体,局 部可见赋存于隐爆角砾岩体中的矿化。总体上,该 矿床累计查明银 270 t、铅锌 80000 t、金 3000 kg。

蒿坪沟 Ag-Au 多金属矿床出露地层主要为太 华群(Arth)片麻岩,矿床区北部被熊耳群(Pt₂x)火山 岩及新生代沉积岩覆盖(图 2)。矿区内断裂构造发 育,以北东向为主,其次为近南北向和北北东向。 矿床区西北部有晚中生代花岗斑岩出露,面积约 0.1 km², 其与蒿坪沟 Ag-Au多金属矿床成矿关系密切。根据已有研究统计,花岗斑岩锆石 U-Pb 年龄共分为3期(~135 Ma,~123 Ma,~110 Ma; 叶会寿,2006; Mao et al., 2011; 梁涛等, 2015; 刘文毅等, 2019; Tian et al., 2023)。北西走向的隐爆角砾岩体位于花岗斑岩的南东部(图2),长约750 m,宽约50~130 m,为蒿坪沟 Ag-Au多金属矿床内的重要赋矿地质体,热液矿物主要赋存于基质中,其矿化阶段和矿物组合与陡倾矿脉一致(Tian et al., 2023)。

根据蒿坪沟 Ag-Au 多金属矿床探矿及采矿成 果(图 3),铅锌矿体主要分布在矿区内的二十多条 断裂带内,矿脉以陡倾斜为主,倾向多为北西向,脉 宽 0.3~5.7 m之间;矿体严格受断裂构造带控制,呈 现明显的分支复合、膨大缩小现象,走向及倾向上 多呈舒缓波状。矿体以脉状、透镜状为主,总体表 现为"伸展脉"的特征(Li et al., 2013)。蒿坪沟 Ag-Au 多金属矿床矿石矿物主要为方铅矿、闪锌矿及 含银硫化物,脉石矿物为石英、铁白云石和菱铁 矿。矿石构造主要以脉状、网脉状及角砾状为主。



图 2 蒿坪沟 Ag-Au 多金属矿床地质简图(底图据梁涛等, 2015 修改)

Fig. 2 Geological map of the Haopinggou Ag-Au polymetallic deposit(Base map modified after Liang et al., 2015)



图 3 蒿坪沟 Ag-Au 多金属矿床勘探线剖面图 (剖面位置见图 2)

Fig. 3 Representative cross-section of the Haopinggou Ag-Au polymetallic deposit (The position of the cross-section is shown in Fig. 2.)

结合已有研究成果, 蒿坪沟 Ag-Au 多金属矿床成矿 过程可以划分为3个阶段; 从早到晚分别为阶段 I: 石英-菱铁矿-黄铁矿阶段, 阶段 II: 石英-方铅矿-闪锌矿阶段, 阶段 III: 石英-碳酸盐阶段(图 4; Li et al., 2013; Tian et al., 2023)。



图 4 蒿坪沟 Ag-Au 多金属矿床矿物共生序列(据 Tian et al., 2023 修改)

Fig. 4 Paragenetic sequence of the Haopinggou Ag-Au polymetallic deposit(modified after Tian et al., 2023)

蒿坪沟 Ag-Au 多金属矿床围岩蚀变普遍发育, 类型多样。通过收集巷道和钻孔中采集的矿化和 蚀变样品,对其矿物组合和结构关系进行了统计和 观察。矿脉与围岩接触界线截然,蚀变矿物常形成 对称的蚀变晕,宽度在 0.2~2m 之间。整体而言,蚀 变带由内向外通常为绢云岩化(最常见)→泥质→青 磐岩化,分别对应的矿物组合为绢云母(伊利石)-石英±碳酸盐,高岭石-蒙脱石以及绿泥石-阳起 石±绿帘石±碳酸盐(Tian et al., 2023)。绿泥石是蒿 坪沟 Ag-Au 多金属矿床中很重要的热液蚀变矿物, 在片麻岩围岩、陡倾矿脉及隐爆角砾岩中都有分 布,尤其是在矿化强烈的部位,绿泥石十分发育 (图 5a)。根据已有成果显示及实地野外观察,绿泥 石的发育与石英铅锌矿脉具有共生/伴生关系(图 5a; Tian et al., 2023),可以用来反映该期热液流体的物 理化学环境。

3 样品及研究方法

通过系统采集含矿石英脉和隐爆角砾岩中的 样品,并开展野外观察和室内镜下鉴定,将蒿坪沟 Ag-Au多金属矿床中的绿泥石划分为3种类型,明 确了不同类型矿物形态、分布位置和组合特征。具 体样品采集位置及矿物组合如下:①样品22H006-1采自740中段H13号脉附近,可见(I型)绿泥石 分布在石英-方铅矿脉体的两侧围岩(图5b);②样 品21H57-04采自钻孔ZK09N20大约孔深7.1m处, 可见(II型)绿泥石以细粒隐晶质形式充填于隐爆 角砾岩基质,与铁白云石共生(图5c);③样品21H60-02取自钻孔ZK11N02孔深242.3m,其中斑岩角砾 发生蚀变,可见基质中(III型)绿泥石与闪锌矿及黄 铁矿共生(图5d)。

根据样品分布及矿物组合特征(图 6a—6d),进 一步开展了电子探针分析。由于绿泥石颗粒细小, 结构复杂,多含有其他矿物的微细包裹体,为提高 数据精确性和可信度,每件样品采取多区域分散取 点,每个区域选取 6~8个点,避免数据偶然性,选 点过程中避开裂隙及矿物包裹体,在 BSE(背散射 光)下尽量选取平整、光滑的绿泥石表面(图 6e—6f)。

绿泥石的化学成分分析在河北省廊坊地质调 查研究院电子探针实验室完成,测试仪器型号为 JEOL EPMA8230,加速电压 15 kV,加速电流 20 nA, 束斑直径 5 μm。标样采用天然矿物或者合成金属 国家标准,分析误差小于 0.01%。

4 绿泥石的化学成分特征

通常认为绿泥石电子探针数据中(NaO+CaO+ K₂O)<0.5%可以排除混染对绿泥石的影响(Hillier, 1993; Zane and Fyfe, 1995)。蒿坪沟 Ag-Au 多金属矿 床绿泥石样品数据中(NaO+CaO+K₂O)均小于 0.5, 可以用来讨论绿泥石的化学特征。Dyar et al. (1992) 认为除了少部分极端氧化环境外,绿泥石 Fe³⁺的含 量都很低,因此可以使用全铁代表 Fe²⁺的含量。手 标本及镜下观察表明 I 型绿泥石为深绿色(图 5, 图 6), 以脉状或叶片状分布于含铅锌矿石英脉两侧。电



Chl一绿泥石; Gn一方铅矿; Sp一闪锌矿; Py一黄铁矿; Ank一铁白云石

a一铅锌矿脉两侧围岩绿泥石蚀变; b一石英脉两侧蚀变绿泥石; c一隐爆角砾岩基质中填充绿泥石与铁白云石; d一隐爆角砾岩中发生绿泥石 蚀变的斑岩角砾

图 5 蒿坪沟 Ag-Au 多金属矿床绿泥石分布情况

Fig. 5 Distribution of chlorites in the Haopinggou Ag-Au polymetallic deposit

(a) Chlorite alteration of surrounding rocks on both sides of Pb-Zn veins; (b) Altered chlorites occurred on both sides of quartz veins; (c) Chlorites and ankerites filled in breccia matrix; (d) Chlorite-altered porphyry breccia

Chl-chlorite; Gn-galena; Sp-sphalerite; Py-pyrite; Ank-ankerite

子探针数据显示 I 型绿泥石中 SiO₂ 含量集中于 25.54%~ 31.84%,均值为 26.71%; Al₂O₃ 含量范围为 13.37%~ 17.61%,均值为 14.41%; FeO 含量集中于 32.93%~ 39.36%,均值为 36.99%; MgO 含量范围为 6.42%~ 9.45%,均值为 7.84%。

II 型绿泥石在手标本下为墨绿色,显微镜下为 浅绿色,呈浸染状填充于隐爆角砾岩基质中。电子 探针数据显示 SiO₂含量集中于 25.52%~26.51%,均 值为 26.03%; Al₂O₃含量范围为 14.09%~15.68%,均 值为 15.27%; FeO 含量集中于 32.61%~34.23%,均值 为 33.30%; MgO 含量范围为 8.82%~9.42%,均值为 9.22%。

Ⅲ型绿泥石在手标本中呈灰绿色,显微镜下为 翠绿色,以蠕虫状分布于石英颗粒间隙;电子探针 数据显示 SiO₂含量集中于 26.65%~29.45%,均值为 27.17%; Al₂O₃含量范围为 13.55%~15.90%,均值为 15.19%; FeO含量集中于 30.86%~33.33%,均值为 32.11%; MgO含量范围为 10.93%~14.31%,均值为

$11.75\%_{\,\circ}$

通过电子探针数据(实验数据在 OSID 码中)可 计算3种类型绿泥石的结构特征,其中I型绿泥石 八 面 体 二 价 阳 离 子 数 为 R²⁺=6.11~7.12, Ⅱ 型 为 R²⁺=6.81~6.94, Ⅲ型为R²⁺=6.45~6.74; Ⅰ型绿泥石 八面体空位为□=0~0.06, Ⅱ型为□=0~0.05, Ⅲ型 为□=0~0.03; I型绿泥石的 Al[™]=0.76~1.02、Al[™]= 0.89~1.35, Ⅱ型为 Al^N=0.97~1.04、Al^N=1.00~1.09, Ⅲ型为Al^N=0.83~1.04、Al^N=0.88~1.04。在绿泥石 特征分析时,首先通过 Fe-Si 图解(图 7a; a.p.f.u 指原 子数,以O=28计算)对绿泥石成分进行分类,然后 使用 Zane and Weiss (1998)图解确定绿泥石结构类 型。绿泥石的3类样品数据多数落在了铁镁绿泥石 区域(图 7b),在Al+□-Mg-Fe 三角图解中表现为富 铁型特征。在 R²⁺-Si 图解中, 蒿坪沟 Ag-Au 多金属 矿床矿区中3种类型绿泥石均落入了三八面体绿泥 石区域,并介于斜绿泥石和须腾石之间,富镁铁而 相对贫铝(图 7c)。



Chl一绿泥石; Ser一绢云母; Gn一方铅矿; Qz一石英; Sp一闪锌矿; Py一黄铁矿; Cal一方解石 a一发育于石英脉两侧的绿泥石(正交偏光); b一隐爆角砾岩基质中填充的绿泥石(正交偏光); c一与黄铁矿、闪锌矿共生的绿泥石(正交偏 光+反射光); d一填充在石英颗粒间隙的蠕虫状绿泥石(正交偏光); e一背散射镜下绿泥石电子探针打点位置分布,可见绿泥石与绢云母共 生; f一背散射镜下绿泥石电子探针打点位置分布

图 6 蒿坪沟 Ag-Au 多金属矿床绿泥石显微形态特征

Fig. 6 Representative photomicrographs of chlorite characteristics of the Haopinggou Ag-Au polymetallic deposit

(a) Chlorite developed on both sides of quartz veins (cross-polarized light);
(b) Chlorite filled in breccia matrix (cross-polarized light);
(c) Chlorite coexisting with pyrite and sphalerite (cross-polarized light+ reflected light);
(d) Worm-like chlorite filling the interstices of quartz grains (cross-polarized light);
(e) Backscattered electron images of the distribution of chlorite EMPA dots, showing coexistence with sericite;
(f) Backscattered electron images of the distribution of chlorite EMPA dots

Chl-chlorite; Ser-sericite; Gn-galena; Qz-quartz; Sp-sphalerite; Py-pyrite; Cal-calcite

绿泥石中的离子间相互置换普遍,可能有 $Fe^{2+} \rightarrow Mg^{2+}$ 之间的任意替换,契尔马克替代 Al^W与 Al^W→Si(Mg²⁺, Fe²⁺)以及二八-三八面体 3(Mg²⁺, Fe²⁺)→□+2Al^W(Inoue et al., 2009)。通常情况下,在 绿泥石结构晶格中,Si和 Al^W占据四面体晶格, Mg²⁺, Fe²⁺和 Al^N占据八面体位置。绿泥石在四面体 和八面体位置上的替换关系通常通过 Al^N、Al^N和(Fe²⁺+ Al^N)-Mg²⁺三者之间的相关关系图(图 8)来说明。 Al^N与 Al^N二者的相关性较差,说明在 Al^N对 Si 原子 的 替 换 过 程 中, Al^N在 八 面 体 位 置 上 对 Mg²⁺或



a-绿泥石 Fe-Si 图解(Deer et al., 1962); b-绿泥石 Al+□-Mg-Fe 图解(Zane and Weiss, 1998); c-绿泥石 Si-R²⁺图解(据刘燚平等, 2016 修改)

图 7 蒿坪沟 Ag-Au 多金属矿床绿泥石化学性质图解

Fig. 7 Chemical diagram of chlorite from the Haopinggou Ag-Au polymetallic deposit

(a) Fe vs. Si diagram of chlorite (Deer et al., 1962); (b) Al+ - Mg-Fe plot (Zane and Weiss, 1998); (c) Si vs. R²⁺ diagram (modified after Liu et al., 2016)

Fe²⁺的置换作用不明显(廖震等, 2010),因此蒿坪沟 Ag-Au多金属矿床中绿泥石 Al 与 Si 置换不属于单 纯的钙镁闪石替代。Fe²⁺/(Fe²⁺+Mg²⁺)、Mg²⁺与 Al^N的 正相关性不明显,也再次印证了 Al^N在八面体位置 上对 Mg²⁺和 Fe²⁺的置换较弱,即不存在 Al^N-Si 置换 的时候同时发生了 Fe²⁺-Mg²⁺置换(Kranidiotis and Maclean, 1987)。

(Fe²⁺+ Al^W)-Mg²⁺呈接近 1:1 的负相关关系,说 明绿泥石八面体位置主要发生了 Mg²⁺被 Fe²⁺+Al^W的 置换,而从 Fe²⁺和 Mg²⁺的相关关系图(图 8)可以看 出, Fe²⁺和 Mg²⁺呈非常好的负相关线性关系,方程为 Fe=-0.928Mg + 0.95 (决定系数 R^2 =0.764),表明 Fe²⁺和 Mg²⁺的置换是绿泥石八面体位置上最重要的 反应, Mg²⁺-Al^W相关性较差,线性方程为 Mg²⁺=0.369 Al^W-0.43(决定系数 R^2 =0.14),所以 Al^W对 Mg²⁺的置换 不明显。蒿坪沟 Ag-Au 多金属矿床中的绿泥石仅 表现 Fe²⁺对 Mg²⁺具有良好的线性关系,而其他阳离 子 Si⁴⁺与 Al^W均与 Mg²⁺相关性较差,暗示了 3 种类型 绿泥石可能具有不同的成因(Xie et al., 1997)。

Al/(Al+Mg+Fe)常被用来判别绿泥石与母岩之 间的关系(Shirozu, 1978; Zane and Fyfe, 1995),通常 由镁铁质岩石形成的绿泥石具有较低的Al/ (Al+Mg+Fe)值(<0.35),与泥质岩相关则比较高 (>0.35)。Ⅰ型、Ⅱ型和Ⅲ型绿泥石Al/(Al+Mg+Fe) 小于0.35,显示均受镁铁质原岩控制,该结果也与蒿 坪沟 Ag-Au 铅锌矿床普遍发育的角闪岩和片麻岩 围岩相符合。因此蒿坪沟 Ag-Au 多金属矿床的绿 泥石中 Fe 元素的来源可能是镁铁质围岩或者含矿 流体。

5 讨论

5.1 绿泥石形成温度

绿泥石成分多变,对环境极其敏感,通过其成 分和结构的规律性变化来反演温度已经得到了广 泛应用(Kranidiotis and Maclean, 1987; Zane and Fyfe,



a一绿泥石 Al^{VI}-Al^{IV}图解; b一绿泥石 Fe²⁺-Mg²⁺图解; c一绿泥石 Fe²⁺/(Fe²⁺+Mg²⁺)-Al^{IV}图解; d一绿泥石 (Fe²⁺+ Al^{VI})-Mg²⁺图解; e一绿泥石 Mg²⁺-Al^{IV}图解; f一绿泥石 Si⁴⁺-Mg²⁺图解

图 8 蒿坪沟 Au-Ag 多金属矿床绿泥石阳离子相关关系图(单位为 a.p.f.u)

Fig. 8 Correlation of cations in chlorites from the Haopinggou Ag-Au polymetallic deposit(unit a.p.f.u)

(a) Al^{VI} vs. Al^{V} diagram of chlorite; (b) Fe^{2+} vs. Mg^{2+} diagram of chlorite; (c) $Fe^{2+}/(Fe^{2+}+Mg^{2+})$ vs. Al^{V} diagram of chlorite; (d) $(Fe^{2+}+Al^{VI})$ vs. Mg diagram of chlorite; (e) Mg^{2+} vs. Al^{V} diagram of chlorite; (f) Si^{4+} vs. Mg^{2+} diagram of chlorite

1995; Inoue et al., 2009; 刘燚平等, 2016; Zhang et al., 2022)。Cathelineau and Nieva(1985)利用绿泥石中的 Al^N与温度之间的正相关性提出了绿泥石固溶体地 质温度计; 之后 Kranidiotis and Maclean(1987)进一步 修正了该温度计, 并强调该地质温度计适用于铝饱 和条件。Jowett(1991)则进一步限定适用的前提为 150~325 ℃的热液条件。蒿坪沟 Ag-Au 多金属矿 床中 3 种类型绿泥石的 Fe/(Fe+Mg)的范围为 0.55~0.74, 其中 Al₂O₃ 含量为 13.23%~17.61%, 在显微镜下可见绿泥石与石英、铁白云石共生, 且各类 绿泥石都基本伴有绢云母化(图 6e-6f), 所以可认为 其处于一个铝饱和的环境, 符合绿泥石地质温度计

的使用范围。由于绿泥石形成环境的复杂性,为探讨地质温度计在蒿坪沟 Ag-Au 多金属矿床中的适用性,文章使用 Cathelineau(1988)提出的温度公式(1)计算出 *T*₁,然后使用 Jowett(1991)提出的修正公式(2)计算出 *T*₂.

$$T_1 = 321.98 \times \text{Al}^{\text{IV}} - 61.92 \tag{1}$$

 $T_2 = 318.5 \times (\text{Al}^{\text{IV}} + 0.10[\text{Fe}/(\text{Fe} + \text{Mg})]) - 68.7$ (2)

3 种类型绿泥石的温度频数直方图如图 9 所示。 根据绿泥石形成温度频数分布直方图可知(图 9a、

9b), *T*₁整体结果较 *T*₂偏小,每个点相差的温度范围 在 9~15 ℃之间。用 *T*₁与 *T*₂的平均值 *T*_{平均}来评估



a一绿泥石形成温度T₁频数分布直方图;b一绿泥石形成温度T₂频数分布直方图;c一绿泥石形成温度T₃频数分布直方图

图 9 蒿坪沟 Ag-Au 多金属矿床绿泥石形成温度频数 (N) 分布直方图

Fig. 9 Hsitogram of formation temperatures of chlorites from the Haopinggou Ag-Au polymetallic deposit

(a) Hsitogram of formation T_1 of chlorites; (b) Hsitogram of formation T_2 of chlorites; Hsitogram of formation T_3 of chlorites

铅锌成矿阶段中热液的流体温度,结果表明 T_{P+h} 相较于已有研究使用流体包裹体等方式获得的温度整体偏高,误差较大。如Zhang et al(2020)通过流体包裹体获得了康山金成矿期的温度(226~305 °C),以及矿田内与蒿坪沟 Ag-Au 多金属矿床临近矿床的铅锌硫化物成矿期的温度(170~270 °C; Li et al., 2013;徐进鸿, 2021)都低于相应的 T_{P+h} 。

绿泥石中 Al^{IV} 与温度有着密切的关系,但也会随着环境中 Fe/(Fe+Mg)的升高而变化(Zane and Fyfe, 1995)。为了消除 Fe/(Fe+Mg)对温度的影响,可以利用公式(3)对绿泥石 Al^{IV} 进行校正(周栋等, 2018):

 $Al_{k\bar{k}_{T}}^{IV} = Al_{t\bar{k}_{T}} + 0.0268 \quad (Fe/Fe + Mg - 0.65) \quad (3)$

 $T_3 = 106.2 \text{Al}_{\text{k} \neq \text{if}}^{\text{IV}} + 17.5 \tag{4}$

然后使用公式(4)计算校正后的温度 T₃,结果 见图 9c。校正后的温度 T₃与已有成果获得对应期 次的成矿流体温度在同一范围内,印证了 T₃的可信 度,因此 T₃可以代表该矿床中绿泥石的形成温度。 结果表明,蒿坪沟 Ag-Au 多金属矿床校正后的Ⅱ型 绿泥石温度范围为224~245℃,平均温度为233℃; Ⅲ型绿泥石比前者稍低,温度范围为(194~237℃),平均为226℃;Ⅰ型绿泥石的温度最低,范围 为179~227℃,平均216℃。Ⅱ、Ⅲ类绿泥石伴生 于石英-铅锌硫化物(图 6b-6d),而Ⅰ类绿泥石伴 成于热液对围岩的改造。形成机制的不同可能造 成了Ⅱ、Ⅲ类绿泥石与Ⅰ类的温度差异。虽然3种 类型绿泥石形成温度存在差异,但总体上相差不 大,也暗示了该期热液活动较为稳定,没有经历多 期次热液-变质活动的叠加改造,与绿泥石的广泛 发育可能指示存在一定规模的铅锌成矿相吻合。

绿泥石中的阳离子个数与温度之间显示出较 好的线性相关关系(图 10)。Cathelineau(1988)认为 在 100~350 ℃ 内,温度与绿泥石 Al^W值呈正相关关 系,而与八面体空位和 Al^W呈负相关关系。蒿坪沟 Ag-Au 多金属矿床中的绿泥石形成温度 $T_{\mp b}$ 与 Al^W及 Si⁴⁺的关系图解指示了良好的线性关系(图 10a、 10b),同样校正后的温度 T_3 与 Al^W及 Si⁴⁺的关系表现 出了明显的线性相关性(图 10c、10d),该结果与其 他学者在地热系统中对绿泥石的研究结果相似,证



a-绿泥石 AI^{N} -T_{平均}关系图解; b-绿泥石 Si^{++} -T_{平均}关系图解; c-绿泥石 AI^{N} - T_{3} 关系图解; d-绿泥石 Si^{++} - T_{3} 关系图解

图 10 绿泥石形成温度 T与 Al^W及 Si⁴⁺相关性图解

Fig. 10 Correlation diagram of T vs. Al^N and Si⁴⁺

(a) Al^N vs. $T_{average}$ diagram of chlorite; (b) Si⁴⁺ vs. $T_{average}$ diagram of chlorite; (c) Al^N vs. T_3 diagram of chlorite; (d)Si⁴⁺ vs. T_3 diagram of chlorite

实了该温度体系的适用性(图 10;周栋等,2018;张 娟等,2021)。

5.2 成矿流体性质

绿泥石的形成过程受水-岩反应控制(廖震, 2010),从热液体系中析出时,其化学组成受宿主岩 性、流体的复杂成分以及物理化学条件影响(Zhang et al., 2022)。在温度条件相近的中一低温热液矿床 的形成过程中,成矿流体的 pH 值对元素的迁移过 程有明显的影响。如pH值较小,则酸性组分更容 易对碱性组分发生取代作用,即发生 Fe 对 Mg 的取 代,反之若 Mg 取代 Fe,则表征相对碱性的环境(张 娟等, 2021)。Ⅰ、Ⅱ、Ⅲ型绿泥石的 Fe/(Fe+Mg)值 较高,分别为0.67~0.76、0.66~0.67、0.59~0.63,说 明3种类型绿泥石均形成于相对酸性的还原环境 中。酸性的环境也有利于水-岩反应的相互作用, 促进了围岩的溶解,为金属硫化物的进一步沉淀提 供空间(Li et al., 2013)。该过程也产生了泥质的蚀 变矿物,如蒿坪沟Ag-Au多金属矿脉周围绿泥石化 和硅化发生的部位常被泥化覆盖,其形成于酸性流 体对围岩中钾质矿物的萃取(Fialips et al., 1998; Reves, 1990)

5.3 绿泥石形成机制及其找矿意义

绿泥石是岩浆热液矿床成矿机制及找矿勘查 的重要指示物。对于斑岩相关的 Cu(Au-Mo)矿床, 可以通过微量元素的变化趋势指示热液中心(Dora and Randive, 2015; Wilkinson et al., 2015; 葛祥坤等, 2020),典型的如中国土屋-延东斑岩型铜矿床(Xiao et al., 2018a, 2018b)通过 V 和 Ti 含量特征来示踪热 液中心。而对于浅成中一低温热液矿床,绿泥石广 泛发育于主成矿期(周栋等, 2018; Zhang et al., 2020; 张娟等, 2021),并为成矿元素的沉淀提供了有利条 件(李占轲等, 2010; 张娟等, 2021)。蒿坪沟 Ag-Au 多金属矿床内共识别出 3 种类型的绿泥石,其形 成机制的厘定对探讨成矿过程及对开展找矿预测 尤为重要。

蒿坪沟 Ag-Au 多金属矿床成矿过程中大量含 铁碳酸盐(菱铁矿、铁白云石)的出现指示了富含 Fe 的成矿热液(Li et al., 2013)。含 Fe 热液与镁铁质 围岩强烈的相互作用导致了矿区内富铁绿泥石的 产生(图 7)。根据其产出部位及矿物组合特征,现 对蒿坪沟 Ag-Au 多金属矿床 3 种类型的富铁绿泥石 形成机制进行分析。

(1) I型绿泥石为溶蚀-结晶。晚中生代热液

顺着北东向陡倾断裂向上运移,同时与两侧的围岩 发生了物理化学反应,顺着片麻理或矿物间裂隙对 片麻岩中的矿物(黑云母/角闪石)进行溶蚀交代(Li et al., 2013),进而在围岩中形成绿泥石(杨献忠等, 2002;李亮, 2011)。母矿物经常出现明显的蚀变特 征,全部/部分发生交代。这种情况形成的绿泥石发 育于脉体两侧,其中Fe、Mg组分多从围岩中获取。

(2)Ⅲ型绿泥石为溶液-迁移-结晶。含矿热液 携带溶蚀矿物(黑云母/角闪石)发生迁移,随着物理 化学条件变化在矿物间裂隙沉淀结晶形成绿泥 石。该类绿泥石在裂隙内部分布均匀致密,而向外 延伸过程中常显示细脉状或蠕虫状分布的特征,该 情况下 Fe、Mg 组分主要为流体带入。

(3) II 型绿泥石的形成过程可能介于前两者之 间。岩浆的浅部就位过程中(Tian et al., 2023), 围岩 由于流体沸腾作用形成隐爆角砾岩, 胶结物为孔隙 度较大的岩粉和岩屑。之后成矿热液进入角砾岩, 充分溶蚀和交代胶结物, 在孔隙度较大、较为宽松 的环境下形成细粒隐晶质绿泥石填充物。该情况 下的 Fe、Mg 组分可能既来自母矿物, 也有流体的贡献。

结合绿泥石的形成机制可以还原银铅锌的成 矿过程,早期成矿流体进入断裂时由于 CO₂ 的释放 导致溶液 pH 酸度的增加(Li et al., 2013),酸性环境 有利于 Ag、Pb、Zn 离子的运移,随后强烈的水-岩 反应则会中和成矿流体中的酸性的 pH 值, pH 值的 减小则会促进铅、锌等硫化物的沉淀(Li et al., 2013)。 该过程发生于整个蒿坪沟 Ag-Au 多金属矿床银铅 锌成矿事件中(高建京等, 2010; Li et al., 2013)。

与蒿坪沟 Ag-Au 多金属矿床相比,熊耳山矿集 区康山金多金属矿床具有类似的成矿环境及流体 特征(表1,图11a、11b),该矿床中Au形成于流体沸 腾, 而流体混合导致 Ag、Pb、Zn 的沉淀(Zhang et al., 2020; 张哲铭等, 2023)。 蒿坪沟 Ag-Au 多金属矿 床中 I 型绿泥石来自大多落在了低温热液矿床区 域外(图 11a、11b), 表明与矿化的关系相对较弱, Ⅱ、Ⅲ型绿泥石数据接近与花岗质侵入体主导的中-低温矿床范围(图 11a, Trumbull et al., 1996), 暗示了 岩浆热液对成矿的重要意义。Tian et al.(2023)提出 蒿坪沟 Ag-Au 多金属矿床中银铅锌成矿与~125 Ma 侵位花岗斑岩具有密切时空联系,表明了成矿 流体中的岩浆来源。矿区中的绿泥石与铅锌矿化 关系密切,与金矿化并无直接关系,结合Lietal. (2016)提出蒿坪沟 Ag-Au 多金属矿床 Au 和 Ag-Pb-Zn来自不同的流体系统,形成于两期独立的成矿事 件,因此推断蒿坪沟 Ag-Au 多金属矿床 Au 可能形 成于更早的一期成矿事件中。

表1 熊耳山矿集区蒿坪沟 Ag-Au 多金属矿床与康山金多金属矿床地质特征

Table 1 Geological characteristics of the Haopinggou Ag-Au polymetallic deposit and the Kangshan Au polymetallic deposit in the Xionger' shan ore concentration area

矿床名称	蒿坪沟Ag-Au多金属矿床	康山金多金属矿床
矿床类型	岩浆热液型	岩浆热液型
大地构造位置	华北克拉通南缘、熊耳山矿集区西北部	华北克拉通南缘、熊耳山矿集区西南部
控矿构造	北东向陡倾断裂和局部隐爆角砾岩	北东向脆性断裂
成矿阶段及矿物组合(Li et al., 2013)	第一阶段: Qz-Sd-Mag-Elc	第一阶段: Qz-Py
	第二阶段: Gn-Sp-Qz-Ank	第二阶段: Qz-Py-Ccp-Au
	第三阶段: Qz-Cal-Fl	第三阶段: Gn-Sp-gold-Qz-Ank-
	_	第四阶段: Qz-Cal-Fl
成矿时代	热液独居石125~123 Ma(Tian et al., 2023)	热液独居石年龄为131 Ma(张哲铭等, 2023)
流体特征及来源	含银硫化物来自蒿坪沟花岗岩体,表现为还 原性的酸性流体(Li et al., 2016)	含金流体来自隐伏花岗岩体,表现为相对弱酸性的还原环境;其成矿机制为 流体沸腾,第三阶段主要成矿机制为流体混合(Zhang et al., 2020)
成矿温度(Li et al., 2013; 徐 进鸿, 2021)	—	Qz-Py: 322~359 °C
	Qz-Sd-Mag-Elc: 217~349 °C	Qz-Py-Ccp-Au: 226~305 °C(256~302 °C)
	Gn-Sp-Qz-Ank: 172~267 ℃(194~237 ℃)	Gn-Sp-Au-Qz-Ank: 185∼246 ℃
	Qz-Cal-Fl: 116~205 ℃	Qz-Cal-Fl: 130∼221 ℃
绿泥石分类	主要为铁镁绿泥石	富铁种属的绿泥石,围岩属于铁镁绿泥石,与矿化相关的绿泥石为铁绿泥石 (周栋等,2018)

注:成矿温度为流体包裹体测温数据(Li et al., 2013;徐进鸿, 2021),括号内为绿泥石EMPA数据所计算出T3温度



不同矿床类型中的绿泥石投图区域:1一活动的地热系统;2一块状硫化物矿床(Kranidiotis and Maclean, 1987);3一热液铜(金)矿化带(Zane and Fyfe, 1995);4一与铜金矿化相关的绿泥石(Dora and Randive, 2015);5一云母石英岩中的绿泥石(Randive et al., 2015);6一与花岗岩相关的矿床(Trumbull et al., 1996);7一热液脉型矿床(Walshe, 1986)

a一绿泥石中 Fe/(Fe+Mg)-(Si/Al)关系图解; b一绿泥石中 Fe/(Fe+Mg)-Al^N关系图解(据周栋等, 2018 修改)

图 11 不同成因类型矿床中的绿泥石特征

Fig. 11 Chlorite characteristics in different genic types of deposits

(a) Fe/(Fe+Mg) vs. (Si/Al) diagram of chlorite; (b) Fe/(Fe+Mg) vs. Al[№] diagram of chlorite (modified after Zhou et al., 2018) Data source: 1–active geothermal system; 2–massive sulfide deposit (Kranidiotis and Maclean, 1987); 3–hydrothermal mineralization zone (Zane and Fyfe, 1995); 4–Chlorites related to Cu-Au mineralization (Dora and Randive, 2015); 5–green-mica quartzites (Randive et al., 2015); 6–deposits associated with granite (Trumbull et al., 1996); 7–hydrothermal vein-type deposit (Walshe, 1986)

蒿坪沟 Ag-Au 多金属矿床从金成矿阶段、铅锌 成矿阶段到碳酸盐阶段,成矿环境和流体性质逐渐 发生改变(Zhang et al., 2020; Tian et al., 2023)。在强 烈的水-岩反应过程中,镁铁质围岩中的斜长石和 黑云母被蚀变为绢云母和绿泥石,热液流体从围岩 中获得了可观的 Fe²⁺、Mg²⁺和 Ca²⁺等阳离子,有利于 碳酸盐矿物的沉淀。流体的 pH 由酸性逐渐变为弱 酸性,最后趋近于中性。低温流体(大气降水)的不 断添加则导致成矿过程中温度逐渐下降(Li et al., 2013)。成矿环境及流体性质的差异也限制了金成 矿与石英-碳酸盐阶段绿泥石的形成。

作为蒿坪沟 Au-Ag 多金属矿床内重要的热液 蚀变产物,绿泥石通常存在于矿化强烈的部位,且 3类绿泥石的野外识别标志都较为明显。Ⅱ型绿泥 石在手标本下为墨绿色,呈浸染状充填于基质中, 极易分辨;Ⅲ型绿泥石呈灰绿色充填于矿物间裂 隙,可以观察到较明显的矿物蚀变特征;Ⅰ型绿泥 石通常发育于脉体两侧围岩,本身不含矿,但该类 绿泥石的广泛发育可能是大规模含矿热液上涌与 太华群片麻岩相互反应的结果,在勘查过程中可以 在蚀变围岩两侧寻找含矿石英脉体。

泥石,其化学特征均反映了酸性、还原性的流体环 境。3种绿泥石发育于铅锌矿化强烈的部位,可以 作为铅锌矿脉的关键识别标志。绿泥石的形成与 镁铁质围岩密切相关,其化学性质暗示了早期成矿 热液中有岩浆流体的参与。

(1) 蒿坪沟 Ag-Au 多金属矿床发育 3 种成因类型的绿泥石: I 型分布在石英脉两侧的围岩中,由 围岩蚀变形成; II 型呈细粒、隐晶质填充质隐爆角 砾岩基质中; II 型与铅锌硫化物共生、以蠕虫状分 布在矿物颗粒间隙。3 种类型的绿泥石均为斜绿泥 石,并落在了铁镁绿泥石的范围内,在结构的阳离 子置换中,主要发生了 Fe²⁺对 Mg²⁺的置换,其余置换 作用均不明显。

(2) 矿区中与矿化相关的绿泥石形成温度略高于石英脉两侧的绿泥石,运用绿泥石经验性公式计算该类矿床的温度时,需要考虑Fe/(Fe+Mg)对温度的影响。

(3)矿区中3种类型绿泥石的形成与镁铁质围 岩密切相关,形成机制可能包含溶蚀-结晶和溶蚀-迁移-结晶两种方式,成矿流体为具有还原性的酸 性富铁流体。

(4) 矿区中绿泥石通常存在于矿化强烈的部位,代表强烈水-岩反应的发生,其产生与大规模铅锌成矿期相匹配,可以作为铅锌矿脉的关键识别标

6 结论

志,其化学性质暗示早期成矿热液中有岩浆流体参与。 致谢:在电子探针实验中得到了河北省区域地质调 查院(河北省地学旅游研究中心)修迪的协助, 在数据处理方面得到了中国地质大学(北京)张 伟博士的帮助;同时感谢两位匿名审稿人以及主 编提出的宝贵意见。

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