https://doi.org/10.3799/dqkx.2021.209



浙江郎村钨钼矿床花岗斑岩岩石地球化学、年代学及 岩石成因

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摘 要:即村钨钼矿床位于浙西北安吉县境内,是钦杭成矿带东段新发现的中型斑岩型矿床.矿体主要产于花岗斑岩岩体中 以及岩体与围岩的内外接触带中.为探讨成矿岩浆来源及动力学背景,在野外地质工作基础上对矿区与成矿相关的花岗斑岩 进行了详细的岩相学、锆石年代学和岩石地球化学研究.结果表明,花岗斑岩LA-ICP-MS 锆石U-Pb年龄为129.7±1.1 Ma.岩 石地球化学结果显示高硅、富碱和弱过铝质特征,属于钾玄岩系列花岗岩类,其稀土配分曲线显示轻稀土富集型特征,并具有 明显的Eu负异常(δEu=0.36~0.40).高硅、富碱、高的10 000 Ga/A1比值(2.57~2.90)和高Zr+Nb+Ce+Y值(379.5×10⁻⁶~ 462.0×10⁻⁶),显示了A型花岗岩的特征.花岗斑岩全岩的(⁸⁷Sr/⁸⁶Sr),为0.707 17~0.709 08, ε_{Nd}(t)为-6.1~-4.0,二阶段模式 年龄(T_{DM2})为1.25~1.42 Ga;锆石的 ε_{HI}(t)为-5.94~-0.87,二阶段模式年龄(T_{MD2})介于1.23~1.56 Ga,表明花岗斑岩来源于 中元古代地壳部分熔融,并有少量地幔物质的参与.花岗斑岩可能形成于中国东部岩石圈伸展—减薄的构造背景下,是华南大 规模岩浆-构造事件的产物.

关键词:郎村钨钼矿床;花岗斑岩;地球化学;年代学;岩石学;矿床学.
中图分类号: P581;P597
文章编号: 1000-2383(2023)10-3725-18
收稿日期: 2021-09-03

Geochemistry, Geochronology and Petrogenesis of Granite Porphyry in Langcun W-Mo Deposit, Zhejiang Province

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Abstract: The Langcun tungsten-molybdenum deposit is a medium porphyry-skarn type deposit in the Northwest Zhejiang

Citation: Cui Kai, Xie Yuling, Qu Yunwei, Tang Yanwen, Li Lamei, Chen Qi, 2023. Geochemistry, Geochronology and Petrogenesis of Granite Porphyry in Langcun W-Mo Deposit, Zhejiang Province. *Earth Science*, 48(10): 3725-3742.

基金项目:国家"十二五"科技计划项目(No. 2011BAB04B00);国土资源部公益性行业基金项目(No. 201011011).

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引用格式:崔凯,谢玉玲,曲云伟,唐燕文,李腊梅,陈琦,2023.浙江郎村钨钼矿床花岗斑岩岩石地球化学、年代学及岩石成因. 地球科学,48(10):3725-3742.

Province, western Qinzhou-Hangzhou metallogenic belt (QHMB). In this paper, it analyzed the petrological, chronological and geochemical characteristics of the ore-forming related granite porphyry in this deposit to constrain its genesis and interpret its tectonic significance. Granite porphyry formed at 129.7 \pm 1.1 Ma (zircon U-Pb, LA-ICP-MS) and belongs to shoshonite series with high SiO₂, alkali contents, and weakly peraluminous. The characteristics, i.e., right-dipping chondrite normalized REE pattern with negative Eu anomalies (δ Eu=0.36-0.40) and the 10 000 Ga/Al and Zr+Nb+Ce+Y values range from 2.57 to 2.90 and 379.5×10^{-6} to 462.0×10^{-6} , respectively, suggest that the granite porphyry belongs to A-type granite. Whole rock Sr-Nd compositions are characterized by $I_{\rm Sr}$ and $\varepsilon_{\rm Nd}(t)$ range from 0.707 17 to 0.709 08 and -6.1 to -4.0, respectively, with Nd model ages of 1.25 to 1.42 Ga. Meanwhile, zircon Hf compositions are characterized by $\varepsilon_{\rm HI}(t) = -5.94$ to -0.87 with two stage model ages of 1.23 to 1.56 Ga. All these evidences suggest that granite porphyry is formed by an ancient continental crust mixed with mantle material source under the tectonic of lithosphere extension and thinning in South China.

Key words: Langcun W-Mo deposit; granite porphyry; geochemistry; geochronology; petrology; mineralogy.

0 引言

中国是世界第一大钨钼资源国和生产国,斑岩 钼矿是中国钼资源的主要来源.中国的斑岩钼矿主 要沿古板块和现代板块结合带分布,形成了多条时 代各异的钼成矿带.钼矿常与钨矿共生,形成了一 些重要的钨钼矿床,如阳储岭钨钼矿床、南泥湖钨 钼矿床、新田岭钨钼矿床等.

传统观点认为,斑岩钼矿、钨锡矿一般与壳源 岩浆岩有关(Chen et al., 2000; Hua et al., 2003; 侯增谦和杨志明,2009).但越来越多的研究表明,斑 岩钼矿和云英岩型钨锡矿的成矿岩浆也显示了幔 源岩浆的贡献(毛景文等,2020;谢玉玲等,2020;蒋 少涌等,2020).因此,与钼、钨、锡成矿有关的岩浆岩 的源区仍存在争议.华南是我国钨、钼、锡等金属的 重要矿集区,且成矿与该区广泛分布的中生代花岗 质岩石有密切的时空或成因联系(陈毓川,1983;华 仁民等,2003,2010;毛景文等,2008;赵葵东和蒋少 涌,2022).前人通过岩石学、岩石化学和同位素地球 化学研究,在该区划分出一系列与钨一锡一钼成矿 有关的A型花岗岩带(朱金初等,2008;蒋少涌等, 2008).朱金初等(2008)通过岩石学和岩石地球化学 厘定出了南岭地区燕山期含钨锡的A型花岗岩带, 也有学者称之为十杭带湘南-桂北段A型花岗岩 带,并认为A型花岗岩为W、Sn等成矿元素的富集 提供了有利条件(蒋少涌等,2008);南岭成矿带东 缘的白石嶂钼矿、锡田钨锡多金属矿床(周云等, 2017)和皖南桂林郑钼矿,与成矿相关的岩体也显 示了A型花岗岩的特征(陈雪锋等,2017;Ren et al., 2018;谢昊等, 2020).A型花岗岩与钨、钼、锡 成矿有关的认识得到越来越多学者的认同,且这些 A型花岗岩的岩石化学均显示了壳幔混源的特征,

幔源岩浆作用在成矿岩浆形成过程中可能具有重 要作用.

郎村钨钼矿床位于浙江省安吉县,大地构造位 置处于华夏板块与扬子板块结合带(简称钦杭结合 带)东段.该结合带是我国重要的钼、钨、铜、多金属 成矿带,除已知的德兴超大型斑岩铜矿、朱溪钨金矿 外,近年还发现有竹溪岭、逍遥、郎村、安吉等多个大 型、中型钼(钨)矿和热液型铅锌矿,具有重要的找矿 潜力.郎村钨钼矿床是近年发现的以黑钨矿、白钨 矿、辉钼矿为主要矿石矿物的中型钨钼矿床,估算钨 333资源量(WO₃)1.85万t,平均品位1.25%(陈琦, 2015). 朱高伶俐(2016)和 Tang et al. (2020)分别对 该矿床的成矿流体演化和成矿年代学方面进行了研 究,而对与成矿相关的岩体尚未有深入探讨.本文选 择郎村矿区花岗斑岩为研究对象,在详细的野外地 质调查和室内研究基础上,开展了花岗斑岩的岩石 学、年代学、岩石地球化学及全岩 Sr-Nd、锆石 Hf同 位素等的研究工作,旨在对花岗斑岩的岩石成因,尤 其是岩浆源区进行约束,同时探讨其与钨钼成矿作 用的关系及其指示的构造环境.

1 区域地质概况

郎村钨钼矿床处于扬子板块东南缘,大地构造 上属于钦杭成矿带东段.杨明桂和梅勇文(1997)首 次提出钦杭结合带的概念,即位于华南地区扬子板 块和华夏板块在晋宁期碰撞拼贴形成的巨型古板 块结合带(图1),其南西起自钦州湾,北东至杭州 湾,全长约2000 km,呈反S状弧形展布(杨明桂和 梅勇文,1997).对钦杭结合带的构造演化目前尚存 在不同认识.其先后经历了晋宁期、加里东期、印支 期和燕山期等复杂的构造岩浆活动,造成其多期次

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图 1



钦杭成矿带及周围晚中生代W矿床分布 Fig.1 Distribution of the W deposits in the Qinzhou-Hangzhou metallogenic belt (QHMB) and surrounding area 据参考文献:杨明桂和梅勇文(1997);Tang et al.(2017)

的岩浆事件(杨明桂和梅勇文,1997;毛景文等, 2011;舒良树,2012).有学者认为,新元古代时期,华 南处于多岛弧古洋盆体系,在晋宁期经历了洋一陆 (弧)俯冲、陆陆碰撞,并导致华夏板块和扬子板块 碰撞对接(胡开明,2001;杨明桂等,2009;周永章 等,2017).加里东期后,华夏板块和扬子板块经历又 一轮陆内裂解和陆陆碰撞,最终完全拼贴形成统一 的华南陆块(胡肇荣和邓国辉,2009).印支期,扬子 板块和华夏板块发生了强烈陆内褶皱造山,华南进 入板内构造发展阶段(Li and Li, 2007).进入燕山 期,钦杭结合带东段主要受太平洋板块斜向俯冲的 影响,中一晚侏罗世,由于俯冲角度的转变,构造体 制由挤压碰撞转换为伸展拉张,造成岩石圈减薄, 并引起广泛的岩浆作用和大规模的成矿作用(Zhou et al., 2006; 毛建仁等, 2013). 钦杭结合带作为一 条古板块结合带和构造薄弱带,其内和旁侧形成了 我国华南地区重要的燕山期花岗岩带和W多金属 成矿带(毛景文等,2007;杨明桂等,2009),其中包 括典型的香花岭、瑶岗仙、柿竹园、香炉山、阳储岭、 东源、竹溪岭等大型、超大型钨、钨锡、钨钼等多金 属矿床.

区内扬子和华夏地块具有不同的结晶基底.钦 杭成矿带北东侧华夏地块由太古代一古元古代结 晶基底组成(胡开明,2001;胡肇荣和邓国辉, 2009), 基底岩石 Nd 模式年龄大多分布在 1.8~ 2.2 Ga(陈江峰等,1999);南西侧扬子地块主要由元 古代结晶基底组成,扬子地块西南部基底岩石 Nd 模式年龄主要在1.5~1.8 Ga之间(陈江峰等, 1999).研究区内地层发育较为齐全,自新元古界至 中生界均有出露.区内发育多条北东向和东西向深 大断裂,主要形成于晋宁期和燕山期,如江山一绍 兴深断裂(NW)、学川-湖州大断裂(NW)和湖 州-嘉善大断裂(EW)等(朱安庆等,2009),这些断 裂构造控制了区内中生代花岗岩及其矿床的侵位. 区内燕山期中酸性岩浆岩分布较为广泛,吕劲松等 (2017)将区内燕山期岩浆活动分为两期:早期 (180~145 Ma) 以 I 型 和 S 型 花 岗 岩 为 主;晚期 (145~100 Ma)为S型和A型花岗岩,特别是

136 Ma之后,主要以A型花岗岩为主.

2 矿床地质

郎村钨钼矿区位于浙江省安吉县报福镇.矿区 出露的地层主要为南华系休宁组和南沱组,震旦系 蓝天组、寒武系荷塘组、大陈岭组和杨柳岗组、志留 系霞乡组、白垩系劳村组和黄尖组、第四系(图2;浙 江省第一地质大队内部资料).其中,南华系休宁组 以角岩为主,是主要的含矿地层.矿区内褶皱和断 裂构造发育:以南华系为核部发育复式背斜,两翼 由震旦系-寒武系组成;受区域性北东向北川-湖 州大断裂(F₁)构造影响,矿区内发育一系列以北东 向为主的次级断裂,控制着矿体的产状和分布

矿区内燕山期岩浆活动强烈,具有多期侵位的 特点,以酸性侵入岩为主,可见少量的基性岩脉,岩 性从老到新主要为黑云母二长斑岩、细粒花岗岩、 花岗斑岩和煌斑岩.其中,花岗斑岩呈岩脉状侵入 于南华系休宁组中,受构造控制明显.

浙江省第一地质大队通过在矿区地表和深部 工程共控制3条矿化带:郎村I号钨钼矿带、丁丁石 II号铜多金属矿带、阮村西侧II号钼矿带.本次研 究集中在I号钨钼矿体,矿体产于细粒花岗岩、花 岗斑岩中及与南华系休宁组角岩内外接触带中.岩 体及围岩内外接触带中发育的北西向和东西向节 理裂隙控制着含矿石英脉的产状,与成矿关系较为 密切.

I号矿体中的矿石矿物主要为黑钨矿、白钨矿和辉钼矿(图3),矿化类型以细脉浸染型和细脉一网脉状为主,后者为主要的矿化类型.此外,矿区矿石矿物还包括黄铁矿和磁铁矿,以及少量钛铁矿、黄铜矿、磁黄铁矿、方铅矿和闪锌矿等.矿区内蚀变以钾化、硅化、绢英岩化、绿泥石化及碳酸盐化为主,其中,硅化、绢英岩化与成矿密切相关;围岩局部发育砂卡岩化,并可见与砂卡岩化有关的白钨矿化.





据浙江省第一地质大队内部资料修绘.1.第四系;2.劳村组;3.乡霞组;4.杨柳岗组;5.荷塘组;6.大陈岭组;7.灯影组;8.陡山沱组;9.南沱 组;10.休宁组;11.细粒花岗岩;12.花岗斑岩;13.黑云母二长斑岩;14.花岗闪长斑岩;15.硅化破碎带;16.采样位置;17.断裂;18.探矿区范 围;19.矿化带范围





3 样品及测试方法

本次测试样品采自矿区地表新鲜的花岗斑岩 岩体.岩石呈肉红色(图4a),斑状结构,块状构造. 斑晶主要为钾长石、斜长石和石英(图4b~4d),基 质主要为长英质,其中石英约25%~30%.金属矿物 主要为含钛磁铁矿,少量的钛铁矿和黄铁矿,副矿 物主要有金红石、锆石、磷灰石、独居石等.弱蚀变 样品中,可见绿泥石化和长石绢云母化(图4e,4f).

选取5件新鲜的花岗斑岩样品粉碎至200目, 在核工业北京地质研究院分析测试中心进行全岩 主微量元素分析.测试方法依据《GB/T 14506.28-93硅酸盐岩石化学分析方法》和《DZ/T 0223-2001 电感耦合等离子体质谱(ICP-MS)方法》通则进行. 主量元素分析使用Philip PW2404型X荧光光谱仪 (XRF)完成,分析精度优于1%;微量元素和稀土元 素分析使用Finnigan MAT Element I型电感耦合 等离子体质谱仪(ICP-MS)完成,分析精度多优于 3%.具体分析流程见Wang et al. (2013).

锆石单矿物挑选均在河北廊坊区调研究所实 验室利用标准技术完成.锆石制靶以及阴极发光 (CL)观察和照相工作由北京锆年领航科技有限公 司完成.本次锆石 U-Pb 年龄测定在澳大利亚塔斯 马尼亚大学优秀矿床研究中心(CODES)完成,使 用仪器为 Agilent 7500cs 激光剥蚀四极杆电感耦合 等离子质谱(LA-ICP-MS).测试采用 Ar 作为剥蚀 物质的载气,激光波长 193 nm,样品测试采用的激 光束斑直径为 32~33 μm.同位素分馏校正采用 91500 为标准锆石(参见 Wiedenbeck *et al.*, 1995), 元素含量校正以 Zr 作为内标元素,以美国国家标准 技术研究院人工合成硅酸盐玻璃 NIST 610 为标准 参考物质.

本研究全岩 Sr-Nd 同位素分析在核工业北京 地质研究院分析测试中心利用热电离质谱仪 (TIMS)进行比值测定.Rb的误差为2%,Sr的误差 为 $0.5\%\sim0.1\%$,Sm和Nd的误差<0.5%,同位素比 值采用⁸⁶Sr/⁸⁸Sr = 0.1194和¹⁴⁶Nd/¹⁴⁴Nd = 0.7219 进行标准化校正.

锆石 Hf 同位素分析是在中国科学院地球化学研究所矿床地球化学国家重点实验室完成.测试采用 Nu Plasma III型 MC-ICP-MS 和 RESOlution S-155型激光剥蚀系统,实验方法见 Yuan *et al.* (2008).



图 4 郎村矿区花岗斑岩手标本及镜下照片 Fig.4 Photographs and micrographs of the Langcun granite porphyry a.花岗斑岩中见暗色包体; b~c. 钾长石和斜长石斑晶(正交偏光); d. 和石英斑晶(单偏光); e. 长石斑晶绢云母化(正交偏光); f. 绿泥石化 (单偏光); Kfs. 钾长石; Pl. 斜长石; Qz. 石英; Ser. 绢云母; Chl. 绿泥石

4 分析结果

4.1 岩石地球化学

4.1.1 主量元素特征 郎村花岗斑岩的主量元素 分析结果见表 1. 通过分析,花岗斑岩具有以下特 征:(1)花岗斑岩为 SiO₂过饱和岩石, SiO₂含量为 70.58%~71.58%,平均 71.18%; Al₂O₃含量为 14.07%~14.46%,平均 14.27%,.其分异指数 DI为 88.42~90.48,说明花岗斑岩分异演化程度高;(2) 岩石全碱 Alk含量(K₂O+Na₂O)较高,为9.17%~ 9.56%,平均9.37%,相对富碱,在TAS图解中(图 5a),花岗斑岩投点落入花岗岩区域,且属亚碱性岩 石;(3)K₂O/Na₂O变化于1.30~1.47,平均1.36,相 对富钾;在SiO₂-K₂O图解上样品也落入钾玄岩系 列(图5b).铝饱和指数A/CNK介于1.04~1.12,A/ NK介于1.12~1.20,在A/CNK-A/NK图解上,样 品投点落入过铝质岩石区域(图5c),判定花岗斑岩 为弱过铝质岩石;(4)CaO含量为0.52%~0.61%, MgO含量为0.38%~0.40%,P₂O₅含量为0.08%~ 0.10%,相对贫钙、镁和磷.





a. TAS图解(底图据 Middlemost, 1994); Ir-Irvine 分界线, 上方为碱性, 下方为亚碱性; 1. 橄榄辉长岩; 2a. 碱性辉长岩; 2b. 亚碱性辉长岩; 3. 辉 长闪长岩; 4. 闪长岩; 5. 花岗闪长岩; 6. 花岗岩; 7. 硅英岩; 8. 二长辉长岩; 9. 二长闪长岩; 10. 二长岩; 11. 石英二长岩; 12. 正长岩; 13. 副长石辉长 岩; 14. 副长石二长岩; 15. 副长石二长正长岩; 16. 副长正长岩; 17. 副长深成岩; 18. 宽方钠岩/磷霞岩/白榴岩; b. SiO₂-K₂O岩石序列图解(底图 据 Rickwood, 1989); c. A/CNK-A/NK 图解(底图据 Maniar and Piccoli, 1989) 第10期

Table 1	Major ele	ment comp	ositions (%) and rare e	arth and tra	ice element con	npositions	(10^{-6}) of g	ranite porp	ohyry from	Langcu
样品号	LC1089-1	LC1089-2	LC1089-3	LC1089-4	LC1089-5	样品号	LC1089-1	LC1089-2	LC1089-3	LC1089-4	LC1089-
SiO_2	71.53	71.46	71.58	70.58	70.77	Sb	0.74	0.83	0.72	0.64	0.56
Al_2O_3	14.25	14.07	14.19	14.46	14.40	Cs	3.95	4.24	4.23	5.33	5.00
$\mathrm{Fe}_{2}\mathrm{O}_{3}$	2.92	3.25	2.79	3.57	3.29	Ва	674.00	666.00	751.00	736.00	701.00
FeO	2.61	2.34	1.39	1.43	1.92	La	58.70	62.40	61.50	56.00	60.90
MgO	0.38	0.40	0.40	0.39	0.40	Ce	99.50	93.00	108.00	87.00	99.20
CaO	0.53	0.55	0.57	0.52	0.61	Pr	11.70	12.10	11.60	11.60	12.60
Na_2O	4.09	4.10	4.07	3.71	3.81	Nd	45.30	46.70	50.20	45.60	50.80
K_2O	5.44	5.33	5.39	5.44	5.34	Sm	8.81	8.42	8.84	8.96	9.71
MnO	0.05	0.05	0.06	0.06	0.05	Eu	1.00	0.97	1.04	1.09	1.14
TiO_2	0.29	0.29	0.28	0.33	0.33	Gd	7.88	7.83	8.06	7.13	8.57
P_2O_5	0.08	0.08	0.08	0.10	0.10	Tb	1.36	1.33	1.41	1.34	1.38
LOI	0.41	0.43	0.53	0.80	0.87	Dy	7.01	6.85	6.83	6.84	7.22
Total	99.67	99.75	99.78	99.79	99.75	Но	1.42	1.22	1.40	1.39	1.34
A/CNK	1.05	1.04	1.04	1.12	1.09	Er	4.11	3.87	3.99	4.05	3.92
A/NK	1.13	1.12	1.13	1.20	1.19	Tm	0.67	0.67	0.69	0.61	0.61
DI	89.15	89.03	90.48	88.86	88.42	Yb	4.79	4.23	4.54	4.18	4.27
Li	11.50	7.84	10.70	14.60	15.20	Lu	0.60	0.64	0.63	0.59	0.61
Be	3.71	3.58	3.78	3.72	3.60	Та	1.69	1.84	1.99	1.90	2.01
Sc	6.09	5.83	6.12	6.58	7.06	W	3.40	3.77	2.98	4.44	4.94
V	11.80	11.70	12.10	15.40	14.60	Re	0.00	0.01	0.01	0.01	0.01
Cr	11.70	10.30	10.70	10.30	7.27	T1	0.95	1.01	0.98	1.03	0.94
Со	2.59	2.75	2.66	2.68	2.56	Pb	17.70	19.10	20.90	17.90	16.80
Ni	6.07	5.70	5.10	5.18	3.82	Bi	0.14	0.13	0.17	0.25	0.15
Cu	7.69	6.95	7.28	7.79	9.38	Th	21.20	22.30	21.90	21.70	21.30
Zn	52.70	53.40	53.70	54.00	54.30	U	8.55	4.88	5.45	4.88	4.96
Ga	18.40	19.30	20.40	20.30	20.90	Zr	215.00	269.00	270.00	291.00	298.00
Rb	183.00	193.00	195.00	214.00	216.00	Hf	6.77	8.55	8.54	8.12	8.47
Sr	87.30	89.40	102.00	133.00	136.00	∑REE	252.84	250.22	268.73	236.38	262.27
Υ	38.20	38.40	41.90	34.20	39.90	LREE/HREE	2.99	3.14	3.12	3.02	3.19
Nb	26.80	29.10	27.60	27.60	24.90	(La/Yb) _N	8.26	9.95	9.13	9.03	9.62
Mo	4.74	5.38	4.53	5.06	3.15	(La/Sm) _N	4.19	4.66	4.38	3.93	3.95
Cd	0.35	0.29	0.27	0.37	0.40	(Gd/Yb) _N	1.33	1.49	1.43	1.38	1.62
In	0.08	0.08	0.08	0.07	0.06	δEu	0.36	0.36	0.37	0.40	0.37

表1 郎村花岗斑岩主量元素(%)和微量、稀土元素(10-6)分析结果

注:A/CNK代表Al₂O₃/(CaO+Na₂O+K₂O)摩尔比;A/NK代表Al₂O₃/(Na₂O+K₂O)摩尔比;分异指数DI=Qz+Or+Ab+Ne+Le+Kp. 主量元素质量分数单位为%;微量和稀土元素质量分数单位为10⁻⁶.

4.1.2 微量及稀土元素特征 花岗斑岩微量元素 和稀土元素分析结果见表1.微量元素蛛网图和稀 土元素球粒陨石标准化曲线分布见图6.微量元素 和稀土元素地球化学特征如下:

(1) 稀土元素总量较高($\sum REE=250.22 \times 10^{-6} \sim 268.73 \times 10^{-6}$), $\sum LREE/\sum HREE = 8.05 \sim 8.75$, $(La/Yb)_{N}=8.26 \sim 9.95$, 显示相对富集轻稀土元素. $(La/Sm)_{N}=0.41 \sim 0.88$, $(Gd/Yb)_{N}=0.70 \sim$

0.79,轻稀土和重稀土内部未发生明显分馏.球粒陨 石标准化稀土元素配分模式图上(图 6a),Eu呈明显 的负异常(δEu=0.36~0.40),稀土元素总体表现为 海鸥式配分模式.

(2)原始地幔标准化微量元素蛛网图(图 6b)显示,花岗斑岩富集大离子亲石元素K、Rb、Th、Pb等,亏损高场强元素Nb、Ta、Zr、P、Ti等的特征,与大陆地壳特征相似.



Fig.6 Chondrite-normalized REE patterns and primitive mantle-normalized trace element patterns of the granite porphyry from Langcun

a. 标准化值据 McDonough and Sun(1995); b. 标准化值据 Boynton(1984);数据来源:骑田岭花岗岩据柏道远等(2005);白菊花尖花岗岩据 Wong *et al.*(2009)





4.2 锆石 U-Pb 年代学

郎村花岗斑岩样品中锆石颗粒晶型完整,多呈 柱状,大小在70~280 μm. 阴极发光图像显示,锆石 大多具有明显的岩浆振荡环带(图7a),且锆石中的 Th和U含量较高,Th含量为77×10⁻⁶~1703× 10⁻⁶,U含量为104×10⁻⁶~2237×10⁻⁶,Th/U比值 范围为0.64~1.24(表2),均大于0.4,表明其为岩浆 结晶的产物(Hoskin and Black, 2000; Crofu *et al.*, 2003; 吴元保和郑永飞,2004).

花岗斑岩中锆石共分析了15个点,其中有14 个点年龄范围较集中,均落在谐和线附近(图7b), 为有效点,另有一点年龄为142±1.4 Ma,可能为岩 浆上升时捕获的老锆石,不参与年龄计算.14个数 据点的U-Pb同位素下交点年龄为129.7±1.1 Ma (MSWD=1.3),笔者认为其可代表花岗斑岩的成 岩年龄.

4.3 Sr-Nd 同位素

郎村花岗斑岩样品 Sm-Nd 和 Rb-Sr 同位素分析及计算结果见表 3.花岗斑岩的⁸⁷Sr/⁸⁶Sr 和¹⁴³Nd/¹⁴⁴Nd 值分别为 0.717 349~0.719 853 和 0.512 264~0.512 368,初始⁸⁷Sr/⁸⁶Sr(即 I_{sr})和 ε_{sd}(t)

		Table 2	Zircon U-Pb	dating data of the grani	te porphyry from Lang	gcun		
	Th ²³²	U^{238}	T1 /II	同位素	素比值	年龄(Ma)		
分析息	(10) ⁻⁶)	- Ih/U	²⁰⁷ Pb/ ²⁰⁶ Pb	²⁰⁶ Pb/ ²³⁸ Th	²⁰⁷ Pb/ ²⁰⁶ Pb(Ma)	$^{206}{\rm Pb}/^{238}{\rm U(Ma)}$	
LC1089-1	237	191	1.242 1	0.0834 ± 0.0506	$0.020\ 1 {\pm} 0.022\ 0$	1278 ± 99	123 ± 3	
LC1089-2	114	173	0.658 6	$0.066\ 3 {\pm} 0.058\ 3$	$0.020\ 3 {\pm} 0.018\ 0$	816 ± 122	127 ± 2	
LC1089-3	100	148	0.677 3	$0.074.8 \pm 0.061.8$	$0.020\;6\!\pm\!0.016\;5$	$1062\!\pm\!124$	127 ± 2	
LC1089-4	177	237	0.747 9	$0.061.6 \pm 0.037.0$	$0.020\ 3 {\pm} 0.017\ 0$	$659\!\pm\!79$	128 ± 2	
LC1089-5	152	203	0.751 0	$0.050\ 9 \pm 0.033\ 2$	$0.020\ 1 {\pm} 0.016\ 8$	$239\!\pm\!77$	128 ± 2	
LC1089-6	129	161	0.804 1	$0.048\ 5{\pm}0.033\ 5$	$0.020\ 2 {\pm}\ 0.015\ 2$	124 ± 79	129 ± 2	
LC1089-7	121	158	$0.768\ 4$	$0.046\ 6 \pm 0.038\ 5$	$0.020\ 2 {\pm} 0.015\ 9$	$27\!\pm\!92$	$129\!\pm\!2$	
LC1089-8	80	113	0.706 5	$0.057~8 \pm 0.039~2$	$0.020\;5\!\pm\!0.018\;5$	521 ± 86	129 ± 2	
LC1089-9	79	122	0.646 4	$0.046~8 \pm 0.045~1$	$0.020\ 4 \pm 0.016\ 4$	$38\!\pm\!108$	130 ± 2	
LC1089-10	179	206	0.866 0	$0.045\ 9 \pm 0.049\ 4$	$0.020\;5\!\pm\!0.017\;7$	-9 ± 119	131 ± 2	
LC1089-11	78	106	0.735 9	$0.049\ 0 \pm 0.057\ 2$	$0.020\;6\!\pm\!0.022\;9$	148 ± 134	131 ± 3	
LC1089-12	108	165	0.653 5	$0.078~8 \pm 0.027~2$	$0.021\ 4\!\pm\!0.013\ 3$	$1168\!\pm\!54$	131 ± 2	
LC1089-13	149	215	0.694 0	$0.053\ 1{\pm}0.038\ 6$	$0.020~8 \pm 0.014~3$	332 ± 88	132 ± 2	
LC1089-14	1 731	2 275	0.760 8	$0.052\ 1 {\pm} 0.020\ 2$	$0.020~8 \pm 0.011~6$	291 ± 46	132 ± 2	
LC1089-15	180	176	1.020 3	$0.0532\!\pm\!0.0462$	$0.0224\!\pm\!0.0261$	338 ± 105	142 ± 4	

表2 郎村花岗斑岩锆石 U-Pb 定年结果

表3 郎村花岗斑岩 Sr-Nd 同位素组成

Table 3 Sr-Nd isotopic compositions of the granite porphyry from Langcun

样品号	⁸⁷ Rb/ ⁸⁶ Sr	⁸⁷ Sr/ ⁸⁶ Sr	$I_{ m sr}$	147Sm/144Nd	143Nd/144Nd	$I_{ m Nd}$	$\epsilon_{\rm Nd}(t)$	$T_{\rm DM}({\rm Ma})$	$T_{\rm DM2}({\rm Ma})$
LC1089-1	6.835 5	0.719 853	0.707 21	0.120 9	0.512 289	0.512 186	-5.6	1 415	1 376
LC1089-2	6.818 6	0.719 735	0.707 13	0.120 7	0.512 280	0.512 177	-5.7	1 427	1 390
LC1089-3	6.545 9	0.719 489	0.707 38	0.121 6	0.512 264	0.512 161	-6.0	1 467	1 417
LC1089-4	4.487 0	0.717 349	0.709 05	0.122 3	0.512 368	0.512 264	-4.0	1 306	1 253

表4 郎村花岗斑岩Hf同位素组成

Table 4 Hf isotopic compositions of zircon from granite porphyry from Langcun

测点 号	t(Ma)	¹⁷⁶ Yb/ ¹⁷⁷ Hf	2σ	¹⁷⁶ Lu/ ¹⁷⁷ Hf	2σ	¹⁷⁶ Hf/ ¹⁷⁷ Hf	2σ	$I_{ m Hf}$	$\varepsilon_{\rm Hf}(t)$	$T_{\rm DM2}$ (Ma)	$f_{\rm Lu/Hf}$
1	142	0.064 848	0.001 100	0.001 571	0.001 100	0.282 534	0.000 030	0.282 530	-5.71	1 548	-0.95
2	127	0.033 649	0.000 140	0.000 843	0.000 140	0.282 637	0.000 026	0.282 635	-2.00	1 313	-0.97
3	131	0.057 155	0.000 310	0.001 427	0.000 310	0.282 587	0.000 026	0.282 583	-3.84	1 429	-0.96
4	129	0.053 719	0.000 160	0.001 306	0.000 160	0.282 528	0.000 026	0.282 524	-5.93	1 562	-0.96
5	130	0.029 816	0.000 084	0.000 749	0.000 084	0.282 558	0.000 028	0.282 556	-4.79	1 490	-0.98
6	129	0.052 278	0.000 240	0.001 292	0.000 240	0.282 555	0.000 025	0.282 552	-4.94	$1\ 500$	-0.96
7	123	0.068 835	0.002 400	0.001 681	0.002 400	0.282 675	0.000 041	0.282 670	-0.76	1 233	-0.95
8	128	0.078 133	0.000 270	0.002 026	0.000 270	0.282 550	0.000 026	0.282 545	-5.18	$1\ 517$	-0.94
9	131	0.041 398	0.000 034	0.001 057	0.000 034	0.282 594	0.000 022	0.282 591	-3.56	1 412	-0.97

以花岗斑岩成岩年龄 129.7 Ma计算,分别为 0.707 13~0.709 05和-6.0~-4.0.该花岗斑岩的 一阶段模式年龄(T_{DM})为1.31~1.47 Ga,两阶段模 式年龄(T_{DM2})为1.25~1.42 Ga.

4.4 锆石 Hf 同位素

花岗斑岩的9颗锆石 Hf 同位素分析结果见表 4.Hf 同位素值均根据对应的上述锆石 U-Pb 年龄计 算.花岗斑岩 Hf 同位素初始¹⁷⁶Hf/¹⁷⁷Hf 值为 0.282 524~0.282 671,平均为 0.282 576; ε_{Hf}(*t*)为 -5.94~-0.87,均为负值,两阶段模式年龄(T_{MD2}) 介于1.23~1.56 Ga,平均1.44 Ga.

5 讨论

5.1 岩石成因类型

传统的花岗岩成因分类方案是将花岗岩分为S型、I型、M型和A型(Chappell and White, 1974; Collins *et al.*, 1982).S型花岗岩一般为过铝质,铝 饱和指数(A/CNK)>1.1,含有白云母、堇青石、石 榴子石、红柱石等特征矿物;I型花岗岩一般为偏铝 质或弱过铝质,A/CNK<1.05,含有黑云母、普通角 闪石等特征矿物;A型花岗岩一般含有霓(辉)石、钠 (铁)闪石、铁云母等碱性暗色矿物.但高分异的I型 和S型花岗岩在矿物学及地球化学特征上与A型花 岗岩相似,因此,区分岩石成因类型需从多方面综 合考虑.

研究表明, P_2O_5 和 SiO₂的相关关系可以有效判 别 S型、I型花岗岩(Chappell and White, 1992; Wu *et al.*, 2003; Li *et al.*, 2007). 郎村花岗斑岩 P_2O_5 含 量较低且随着 SiO₂含量的升高而降低, 与A/I型花 岗岩的演化趋势相同(图 8). 花岗斑岩为弱过铝质 岩石(图 5c), 不具有 S型花岗岩的强过铝质特征, 结 合岩石化学上表现为高硅(SiO₂含量 70.46%~ 71.58%)、富碱(全碱 Alk含量 9.17%~9.56%)的特 点, 认为郎村花岗斑岩不属于 S型花岗岩, 为 I型或 者 A型花岗岩.

郎村花岗斑岩的锆石 U-Pb 年龄为 129.7 Ma, 形成于早白垩世早期,前人对钦杭成矿带东段花岗 岩进行研究,识别了多处同一时期的 A 型花岗岩岩 体,如大桥坞花岗斑岩和杨梅湾花岗岩(136~ 133 Ma, Yang et al., 2012)、大茅山和桐山花岗岩 (129~21 Ma, Jiang et al., 2011)、密坑山花岗岩 (124 Ma,邱检生等, 2005)、白菊花尖花岗岩 (123 Ma, Wong et al., 2009)等.本文研究的郎村 花岗斑岩与这些 A 型花岗岩体具有相似的地球化 学特征.

Whalen *et al.*(1987)通过对A型花岗岩数据的 总结发现,A型花岗岩具有高硅、富碱、高Ga、Zr、 Nb、Ce和Y含量的特点,据此提出以10000Ga/Al> 2.6和Zr+Nb+Ce+Y>350×10⁻⁶作为判别A型 花岗岩的标准.在10000Ga/Al和Zr+Nb+Ce+ Y-(Na₂O+K₂O)/CaO图解上,花岗斑岩大多落在 A型花岗岩区域内,与区域内A型花岗岩范围一致



Fig.8 $SiO_2 - P_2O_5$ diagram of granite porphyry from Langcun

(图 9),表明花岗斑岩为A型花岗岩.这一结论也得 到以下证据的支持:(1)花岗斑岩具有高硅(平均 71.18)、富碱(平均 9.37%)、高铝饱和指数(A/CNK 为 1.04~1.12)、贫 CaO和 MgO的特点(邱检生等, 2000;吴锁平等,2007);(2)花岗斑岩 10 000 Ga/Al 比值为 2.57~2.90(>2.60),Zr+Nb+Ce+Y为 379.5×10⁻⁶~462.0×10⁻⁶(>350×10⁻⁶);其稀土含 量较高,亏损Ba、P、Ti和Eu的特征与A型花岗岩一 致(Wong et al., 2009);(3)花岗斑岩稀土配分模式 与区域上骑田岭和白菊花尖等A型花岗岩一致,为 典型的右倾海鸥型(图 6a);(4)花岗斑岩的锆饱和 温度为 808~844 ℃,由于缺乏继承锆石,锆饱和温 度代表了岩浆熔体的最低温度(赵振华,2016),形 成花岗斑岩的岩浆温度明显高于I型花岗岩的平均 值 781℃(King et al., 1997).

5.2 岩浆源区

关于A型花岗岩的岩浆起源,目前有几种不同的认识:(1)富F和Cl的麻粒岩相下地壳部分熔融(Whalen et al., 1987; Creaser et al., 1991);(2)幔源的碱性基性岩一中性岩分异演化(Anderson et al., 2003; Bonin, 2007);(3)地壳浅部钙碱性花岗岩脱水熔融(Skjerlie and Dana Johnston, 1993; Patiño Douce, 1997);(4)壳幔岩浆混合(Min-gram et al., 2000; Yang et al., 2006).Wong et al. (2009)对白菊花尖A型花岗岩的研究认为,其源区以地幔物质为主,有少量古老地壳物质的参与; Ji-ang et al.(2011)研究的铜山和大茅山A型花岗岩来自下地壳的部分熔融;邱检生等(2005)认为密坑岩



底图据 Whalen et al.(1987);数据来源:白菊花尖花岗岩据 Wong et al.(2009);大茅山和桐山花岗岩据 Jiang et al.(2011);大桥坞花岗斑岩 和杨梅湾花岗岩据 Yang et al.(2012);密坑山花岗岩据邱检生等(2005);FG.分异型的长英质花岗岩;OGT.未分异的 I、S和 M 型花岗岩

体为壳幔物质在源区混合后经历高度分异演化 形成.

花岗斑岩为弱过铝质和钾玄岩一高钾钙碱性系列;稀土元素球粒陨石标准化配分图表现为相对富 集轻稀土的特征,轻、重稀土内部分馏不明显,Eu负 异常;微量元素蛛网图显示富集大离子亲石元素K、 Rb、Th、Pb等,亏损高场强元素Nb、Ta、Zr、P、Ti等 的特征,与大陆地壳特征相似.据前人研究,这些特 征指示了陆壳参与岩浆过程,即岩浆源区以壳源物 质为主(张宏飞和高山,2012;黄兰椿和蒋少涌, 2012).

花岗斑岩的⁸⁷Sr/⁸⁶Sr初始值为0.717349~ 0.719853和 $\varepsilon_{Nd}(t)$ 值为一6.0~一4.0,在 $I_{sr}-\varepsilon_{Nd}(t)$ 图解上,花岗斑岩投点落入地幔组分和华南壳源花岗 岩之间(图10a),表明岩浆主要来源于地壳,但受到 地幔物质的混染(White and Chappell, 1983; Faure, 1986). 花岗斑岩的 $\varepsilon_{Hf}(t)$ 比值均为负值 (-5.94~-0.87),在 $\varepsilon_{Hf}(t)-t$ 图解上,花岗斑岩投 点在亏损地幔之下,地壳1.0 Ga和1.5 Ga演化线之 间(图10b),前人研究表明这种特征的岩浆源区物 质主要来源于古老地壳(吴福元等,2007).花岗斑岩 的全岩 Nd两阶段模式年龄(1.25~1.42 Ga)与Hf两 阶段模式年龄(1.23~1.56 Ga)相近,稍晚于扬子板 块基底岩石的 Nd同位素模式年龄(1.5~1.8 Ga),也 暗示了其源区物质主要为中元古代地壳,但形成时 有明显的地幔物质或新生地壳物质加入.由于花岗 斑岩的 T_{DM} 为1.31~1.47 Ga,不倾向于新生下地壳 的存在(陈江峰等,1999),因此认为郎村花岗斑岩 源区以中元古代地壳为主,有少量地幔物质的混合. 并且在矿区出露有同期的煌斑岩脉,也印证了区内 存在同期幔源岩浆活动.



图 10 郎村花岗斑岩 $I_{Sr} - \epsilon_{Nd}(t)$ 图解(a)和 $\epsilon_{Hf}(t) - t$ 图解(b)





图11 郎村花岗斑岩构造环境判别图

Fig.11 Tectonic environment diagrams of granite porphyry from Langcun a. 底图据 Batchelor and Bowden(1985); ①地幔斜长花岗岩;②破坏性活动板块边缘(板块碰撞前)花岗岩;③板块碰撞后隆起期花岗岩; ④晚造山期花岗岩;⑤非造山期A型花岗岩;⑥同碰撞(S型)花岗岩;⑦造山期后A型花岗岩;b. 底图据 Pearce *et al.*(1984)

5.3 成岩成矿时代和构造环境

前人对区域内晚中生代花岗岩的成岩时间做 过大量研究.Tang et al.(2017)测得铜村钼矿区二长 花岗岩年龄为148.1±1.7 Ma; 厉子龙等(2013)获得 木瓜铜钨多金属矿花岗斑岩年龄为142.2±1.2 Ma; 唐增才等(2017)测得浙西淳安双溪口锡矿花岗岩 年龄为130.4±1.1 Ma; 黄国成等(2012a, 2012b)对 千亩田钨铍矿区顺溪岩体和夏色岭钨矿细粒花岗 岩进行锆石定年,获得其成岩年龄分别为123.5~ 125.5 Ma和126.9±1.7 Ma,距离郎村钨钼矿床最近 的安吉铅锌矿床的细粒花岗岩成岩年龄为133.9± 1.3 Ma和134.5±1.6 Ma(谢玉玲等, 2012),可见浙 西地区成岩成矿时间主要集中在晚侏罗世至早白 垩世.

本文采用LA-ICP-MS 锆石 U-Pb 定年测得郎 村花岗斑岩的侵位时间为129.7 ± 1.1 Ma,与 Tang et al.(2020)获得的郎村黑钨矿 U-Pb 年龄127.4 Ma 误差范围内一致.在空间上,郎村钨钼矿体赋存在 花岗斑岩岩体中及岩体与围岩的内外接触带.因 此,郎村钨钼矿床钨钼矿化与花岗斑岩具有密切的 时空联系,其侵位时代同时或略晚,与区域内的大 规模成岩时间一致,是浙西地区乃至华南地区大规 第10期

模构造一岩浆活动的产物.

华南地区白垩纪侵入岩主要为高钾 I 型和 A 型 花岗岩,分布在浙江、福建、粤北等东南沿海地区, 江西和湖南等地也有少量分布(Li et al., 2000;华 仁民等,2003).早期研究认为,华南与钨锡成矿有关 的花岗岩均为壳源改造型或 S 型花岗岩类,但后来 研究发现部分岩体可归为 A 型花岗岩类,但后来 研究发现部分岩体可归为 A 型花岗岩(华仁民等, 2010).A 型花岗岩形成于伸展环境的观点已得到了 广泛认可,研究 A 型花岗岩对区域构造演化有重要 的指示意义.矿区花岗岩构造环境判别图解(图 11) 显示,郎村花岗斑岩在 R₁-R₂关系图解上落入晚造 山期 - 非造山期 A 型花岗岩区域,Yb+Nb-Rb关 系图解上落入后碰撞花岗岩区域,均指示郎村花岗 斑岩形成于伸展环境.

早中生代华南大陆动力体制经历了从古亚洲 洋一古特提斯洋向滨太平洋构造域的转换(Li et al., 2000; Wang et al., 2005, 2007; Shu et al., 2008;舒良树,2012;毛建仁等,2013,2014),华南大 陆的构造体制主要受古太平洋板块向华南大陆之下 俯冲作用影响.中晚侏罗世一早白垩世,受古太平洋 板块向华南大陆之下俯冲角度变化的影响,俯冲板 块回撤,华南地区构造环境由陆内挤压造山向造山 后伸展作用转换(Li et al., 2000; Zhou and Li, 2000; Zhou et al., 2007;张岳桥等, 2012). 目前, 对于 挤压向伸展转换的时间尚有不同认识, Li et al. (2007)研究认为构造转换时间为145 Ma, 而张岳桥 等(2012)和毛建仁等(2014)认为转变的时间大致在 135 Ma. 在这一大的构造背景影响下, 浙西北在145 Ma也由活动大陆边缘向造山后伸展构造转换,直至 123 Ma岩浆成矿活动基本结束(厉子龙等,2013).郎 村花岗斑岩形成于早白垩世伸展环境,也证明了浙 西地区在130 Ma之前构造环境转换已经结束.

6 结论

(1)花岗斑岩为具有高硅、富碱的特点,稀土含量较高,强烈的Eu负异常,其稀土元素配分曲线具有A型花岗岩的特点,总体表明,花岗斑岩属于高分异的A型花岗岩.

(2)Sr-Nd-Hf同位素研究结果表明,花岗斑岩 成岩物质主要来源于中元古代地壳部分熔融,并有 少量地幔物质的加入.

(3)花岗斑岩 LA-ICP-MS 锆石 U-Pb 年龄为 129.7±1.1 Ma,与区域大规模的岩浆活动时限一 致,形成于伸展一减薄的构造环境.

致谢:两位匿名审稿专家审阅了全文,对文章 内容提出了建设性修改意见,在此致以诚挚的 感谢!

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