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内蒙古阿巴嘎旗阿德拉嘎碱长花岗岩 锆石 LA-ICP-MS U-Pb 年龄、地球 化学特征及构造意义

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内容提要: 阿巴嘎旗阿德拉嘎碱长花岗岩位于贺根山蛇绿岩带北侧。为了确定该碱长花岗岩体的岩石成因类型, 探讨其构造环境, 对该岩体进行了野外地质、岩石学、地球化学和 LA-ICP-MS 锆石 U-Pb 年代学研究。锆石 LA-ICP-MS U-Pb 测年表明, 阿德拉嘎碱长花岗岩体的侵位年龄为 310.7 ± 2.6 Ma, 形成时代为晚石炭世。岩石地球化学研究表明, 阿德拉嘎碱长花岗岩具有较高的 SiO_2 、 $\text{Na}_2\text{O} + \text{K}_2\text{O}$ 含量和较高的 Ga/Al 、 $(\text{Na}_2\text{O} + \text{K}_2\text{O})/\text{CaO}$ 值, 相对贫 CaO/MgO 、 Sr/Ba 、 Eu/Ti 和 P 。稀土元素总量较低, 轻重稀土分馏不明显, 稀土配分曲线呈海鸥性, 负铕异常明显 ($\delta\text{Eu}=0.15 \sim 0.37$)。岩石学和岩石地球化学特征表明, 该碱长花岗岩为 A 型花岗岩, 形成于弧后伸展构造环境, 为晚石炭世古亚洲洋向西伯利亚俯冲作用的产物。

关键词: 碱长花岗岩; A 型花岗岩; 锆石 U-Pb 定年; 晚石炭世; 弧后伸展环境; 内蒙古阿巴嘎旗

A 型花岗岩概念最早由 Loiselle 和 Wones (1979) 提出, 原始定义为碱性 (alkaline)、贫水 (anhydrous) 和非造山 (anorogenic) 的花岗岩。A 型花岗岩概念提出后, 由于其特殊的成因、产出构造背景和重要的地球动力学意义, 引起广大地质工作者的关注 (Whalen et al., 1987; Sylvester, 1989; Bonin, 1990; Eby, 1990, 1992; Whalen et al., 1996; King et al., 1997; Bonin, 2007)。Whalen et al. (1987) 认为 A 型花岗岩具有高 Ga/Al 等地球化学特点, 并建立了相应的 A 型花岗岩判别图解。Eby (1992) 将 A 型花岗岩划分为 A_1 和 A_2 两个亚类, A_1 亚类与洋岛玄武岩 (OIB) 有一定相似性, 产于板内构造环境, A_2 亚类与岛弧玄武岩有一定相似性, 产于后碰撞构造环境。Bonin (2007) 对 A 型花岗岩研究进行了总结, 并将 A 型花岗岩定义扩大为碱性 (alkaline)、贫水 (anhydrous)、非造山 (anorogenic)、铝质 (aluminous) 及模棱两可的 (Ambiguous), 认为 A 型花岗岩可以形成于从陆内到大陆边缘的各种动力学背景或构造环境。

中亚造山带 (CAOB) 是位于西伯利亚板块与塔里木—华北板块之间的巨型复合造山带, 其东段延伸至我国内蒙古及东北地区, 被称为兴蒙造山带

(图 1)。前人在造山带与古亚洲洋演化方面取得较大成果 (Sengor et al., 1993; Dobretsov et al., 1995; Xiao Wenjiao et al., 2003; 张磊等, 2013; Zhu Mingshuai et al., 2014; 邓晋福等, 2015; 刘锐等, 2016; 田树刚等, 2016; 李钢柱等, 2017; 王金芳等, 2017, 2019, 2020, 2021; 李英杰等, 2018; 王树庆等, 2018, 2020; 张晓飞等, 2018a, b, 2019; 程杨等, 2019, 2020; 范玉须等, 2019, 2020; 李锦轶等, 2019a, b; 董培培等, 2021; 李英雷等, 2021; 张夏炜等, 2021; 金松等, 2022; 杨晓平等, 2022)。古亚洲洋于寒武纪之前打开 (Tang Kedong, 1990; Zhu Mingshuai et al., 2014)。寒武纪—志留纪古亚洲洋开始向南北两侧发生双向俯冲 (Xiao Wenjiao et al., 2003), 俯冲持续到石炭纪—早中二叠世 (Xiao Wenjiao et al., 2003; 张磊等, 2013; 李英杰等, 2018; 王树庆等, 2018; 张晓飞等, 2018a, b, 2019; 王金芳等, 2017, 2019, 2020, 2021; 程杨等, 2019, 2020; 范玉须等, 2019, 2020; 董培培等, 2021; 李英雷等, 2021; 张夏炜等, 2021; 金松等, 2022; 杨晓平等, 2022)。古亚洲洋东部最终在晚二叠世末—早三叠世初关闭 (Sengor et al., 1993; Xiao Wenjiao et al., 2003; 邓晋福等, 2015; 刘锐等, 2016; 田树刚等, 2016)。最近, 吕洪波等 (2018) 在

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狼山发现白垩纪蛇绿混杂岩,认为古亚洲洋与华北克拉通是在白垩纪沿儿狼山—阴山一线最终拼合;进而推论,狼山阴山一线以北、白云鄂博—西拉木伦一线以南大片地区是古亚洲洋中的一个地体,而不是华北克拉通的一部分。古亚洲洋形成演化过程中,由北向南形成二连—贺根山(梁日暄,1994;Nozaka and Liu, 2002; Xiao Wenjiao et al., 2003; Zhang Zhicheng et al., 2015)和索伦山—西拉木伦两条重要的蛇绿岩带(梁日暄,1994;王玉净和樊志勇,1997;刘建峰等,2016)。二连—贺根山蛇绿岩带为俯冲带型蛇绿岩带(Xiao Wenjiao et al., 2003),闭合时间晚于早二叠世(张磊等,2013;王金芳等,2017;张晓飞等,2018a,b,2019;程杨等,2019;范玉须等,2019;李英雷等,2021)。索伦山—西拉木伦蛇绿岩带为洋中脊蛇绿岩带(Xiao Wenjiao et al., 2003; Li Jinyi, 2006; Jian Ping et al., 2008)。闭合时间为晚二叠世末—早三叠世(Xiao Wenjiao et al., 2003; Li Jinyi, 2006; 李朋武等,2006; 李锦轶等,2007; 邓胜徽等,2009; 邓晋福等,2015; 刘建峰等,2016)。贺根山蛇绿岩带两侧发育大规模碱性碱长(A型)花岗岩,组成一条近东西向的花岗岩带(洪大卫等,1991; Zhang Xiaohui et al., 2014; Tong Ying et al., 2015)。花岗岩年龄集中于晚石炭世—早二叠世 $271.7 \pm 0.7 \sim 297 \pm 3$ Ma,形成于拉伸构造环境(洪大卫等,1991; 施光海等,2004; 张玉清等,2009; 张磊等,2013; 程银行等,2014; Zhang Xiaohui et al., 2014; Tong Ying et al., 2015; 肖中军等,2015; 贾孝新等,2017; 张晓飞等,2018)。阿巴嘎旗阿德拉嘎碱长花岗岩为首次发现(河北省区域地质矿产调查研究所,2016),本文在岩石学、地球化学和同位素年代学的基础上,探讨碱长花岗岩岩石成因和构造环境为区域碱性(A型)花岗岩带对比研究提供资料。

1 区域地质背景和岩石学特征

研究区位于中亚造山带(CAOB)东南部,贺根山蛇绿岩带北侧,隶属于乌里雅斯太活动陆缘(图1a)。

区域出露地层包括奥陶系、石炭系和第四系(图1b)(河北省区域地质矿产调查研究所,2016)。奥陶系出露地层为下奥陶统铜山组、中奥陶统多宝山组和中—上奥陶统裸河组。铜山组岩性主要为变质中粗粒长石岩屑砂岩、变质细粒岩屑砂岩、变质砾岩夹变质粉砂岩、变质安山岩和结晶灰岩,在变质岩屑长石砂岩见有腕足化石及海百合茎化石。多宝山

组主要为变质玄武安山岩,杏仁状玄武安山岩夹少量变质砂岩、泥质粉砂岩。裸河组主要为变质细粒长石砂岩、长石岩屑砂岩夹变质粉砂岩、变质安山岩,变质粉砂岩中产*Glyptograptus*(矮小雕笔石)化石。石炭系出露地层为上石炭统宝力高庙组,岩性主要为红柱石角岩化粉砂岩、细粒岩屑砂岩、粗粒岩屑砂岩、砾岩、英安质角砾熔结凝灰岩、安山岩、流纹岩等。第四系包括更新统阿巴嘎组和全新统冲洪积物。阿巴嘎组岩性为气孔、杏仁状玄武岩、橄榄玄武岩、玄武质集块角砾岩、玄武质浮岩。冲洪积物由砂砾、砂、黏土等组成。

侵入岩较为发育,岩性主要为二长花岗岩、正长花岗岩、阿德拉嘎碱长花岗岩和花岗斑岩等,岩体均形成于晚石炭世。

阿德拉嘎碱长花岗岩分布于阿德拉嘎附近,呈不规则状产出,长轴方向近东西向,岩石坚硬,节理裂隙发育,一组节理走向北东 20° ,另一组节理走向北西 70° ,倾角 85° ,出露面积约为 13 km^2 。岩体内原生节理发育,局部节理有赤—磁铁矿薄膜充填。该岩体侵入晚石炭世二长花岗岩及正长花岗岩之中,被更新世阿巴嘎组玄武岩角度不整合覆盖。(图2a)。

阿德拉嘎碱长花岗岩新鲜面呈肉红色,具细粒花岗结构,块状构造(图2b)。岩石由钾长石、斜长石、石英组成。钾长石为条纹长石,半自形—近自形板状,粒径 $0.1 \sim 1.5$ mm,具高岭土化,轻微绢云母化,含量 $60\% \sim 70\%$ 。斜长石呈半自形板状,粒径为 $0.1 \sim 0.8$ mm,零散状分布,斜长石牌号 $An = 27$,为更长石,含量 $0\% \sim 5\%$ 。石英呈半自形—它形粒状,粒径为 $0.1 \sim 1.8$ mm,半自形粒状石英多呈堆状聚集分布,它形粒状石英呈填隙状分布,具轻微波状消光,含量 $30\% \sim 35\%$ 。

区内断裂较为发育,以北东向、北西向为主,近东西向及南北向次之。形成时代最早追溯到加里东期,最晚至更新世阿巴嘎期。

2 样品采集和实验方法

系统地质调查的基础上选择新鲜具有代表性样品进行岩石学、地球化学、同位素年代学测试,样品位置及编号见图1b,其中地球化学数据样品6件,编号为P3YQ1、P3YQ2、P3YQ3、P3YQ4、P3YQ5、P3YQ6,同位素样品1件,编号为P3TW,采样位置与P3YQ3位置相同。

样品地球化学分析由河北省区域地质矿产调查

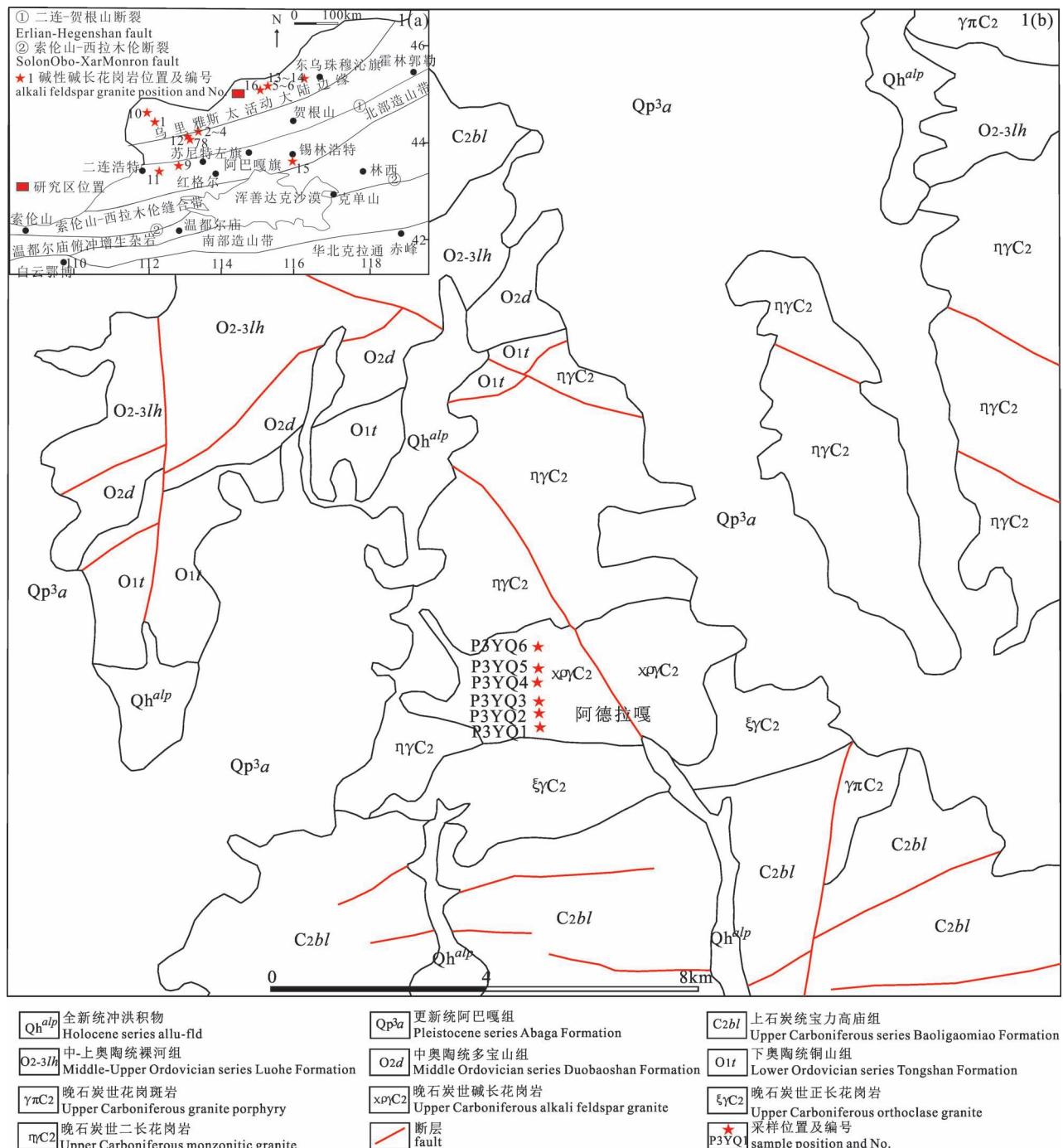


图 1 (a) 阿巴嘎旗阿德拉嘎碱长花岗岩区域构造简图;(b) 阿德拉嘎碱长花岗岩地质简图
(河北省区域地质矿产调查研究所, 2016)

Fig. 1 (a) Sketch tectonic map (a) and geological map (b) of the Adelaga alkali feldspar granite, in Abaga Banner, Inner Mongolia (Hebei Institute of Regional Geology and Mineral Resources, 2016)

图 1a 中数据来源: 1—焦天佳, 2~3—肖中军等, 2015; 4—Zhang Xiaohui et al., 2014; 5—刘建峰等, 2009; 6—张玉清等, 2009; 7—Zhang Xiaohui et al., 2014; 8~11—Tong Ying et al., 2015; 12—贾孝新等, 2017; 13~14—程银行等, 2014; 15—施光海等, 2004; 16—石文杰等, 2019

Data of the Fig. 1a from: 1—Jiao Tianjia, 2015&; 2~3—Xiao Zhongjun et al., 2015&; 4—Zhang Xiaohui et al., 2014; 5—Liu Jianfeng et al., 2009&; 6—Zhang Yuqing et al., 2009&; 7—Zhang Xiaohui et al., 2014; 8~11—Tong Ying et al., 2015; 12—Jia Xiaoxin et al., 2017&; 13~14—Cheng Yinhang et al., 2014&; 15—Shi Guanghai et al., 2004&; 16—Shi Wenjie et al., 2019&



图 2 阿德拉嘎碱长花岗岩野外照片(a)及显微照片(b)

Fig. 2 Filedphotographs (a) and Microphotographs of the Adelaga alkali-feldspar granite

Kfs—钾长石; Qtz—石英

Kfs—K-feldspar; Qtz—quartz

研究所实验室完成。主量元素分析(SiO_2 、 TiO_2 、 Al_2O_3 、 Fe_2O_3 、 MgO 、 MnO 、 Na_2O 、 K_2O 、 CaO 、 P_2O_5)采用X射线荧光光谱(XRF)完成,分析测试仪器为Axios^{max}X射线荧光光谱仪,检测依据为《GB/T14506.28-2010》;灼失量分析采用重量法完成,检测依据《DZG20-1》; H_2O^- 分析采用重量法完成,检测依据《CB/T14506.1-2010》; H_2O^+ 分析采用重量法完成,检测依据《GB/T14506.2-2010》,灼失量、 H_2O^- 、 H_2O^+ 采用P1245电子分析天平测试;FeO采用50mL滴定管滴定,检测依据为《GB/T14506.14-2010》,主量元素、灼失量、 H_2O^- 、 H_2O^+ 分析精度为0.05%。稀土元素及微量元素采用离子体质谱,检测依据为《DZG20-1》,分析测试仪器为X Serise2等离子体质谱仪,稀土元素分析绝对误差为 0.1×10^{-6} ,微量元素 $\leq 5 \times 10^{-6}$ 。实验温度为20℃,湿度为30%。

锆石挑选由河北省区域地质矿产调查研究所实验室完成。样品经过常规的分选后,在双目镜下经人工挑选出纯度在99%以上的锆石,选出的锆石完整、透明度高。将挑选好的锆石粘贴在环氧树脂表面,打磨抛光后露出锆石表面,制成靶样。锆石透、反射光及阴极荧光(CL)照相,由北京锆年领航科技有限公司实验室完成,仪器为JEOL公司生产的JSM6510型扫描电子显微镜。

锆石MC-ICP-MS年龄测试在中国地质科学院矿产资源研究所MC-ICP-MS实验室完成。锆石定年分析所用仪器为Finnigan Neptune型MC-ICP-MS,

激光剥蚀系统为GeoLas 2005。激光剥蚀束斑直径为 $25 \mu\text{m}$,剥蚀深度为 $20\sim40 \mu\text{m}$,频率为10 Hz,能量密度为 2.5 J/cm^2 ,剥蚀过程中采用氦气载气,氩气补偿。信号较小的 ^{207}Pb 、 ^{206}Pb 、 ^{204}Pb (^{+204}Hg)、 ^{202}Hg 用离子计数器(multi-ion-counters)接收, ^{208}Pb 、 ^{232}Th 、 ^{238}U 信号用法拉第杯接收,实现了所有目标同位素信号的同时接收,并且不同质量数的峰基本平坦,进而获得高精度数据。测试过程中在每测定10个锆石样品前后重复测定2个锆石GJ1对样品进行校正,并测量1个锆石Plesovice,用于观测仪器的状态确保测试精确度。详细实验过程见侯可军等(2009)。实验温度20℃,实验湿度55%。对采集的数据采用中国地质大学刘勇胜博士研发的ICP-Ms DataCal程序(Liu Yongsheng et al., 2008)和Ludwig(2012)的Isoplot程序进行处理,并绘制谐和图等图件,置信度为95%。

3 测试结果

3.1 主量元素

阿德拉嘎碱长花岗岩样品烧失量(0.29%~0.69%)较低,总量为99.97%~99.98%,岩石地球化学特征能够有效反应岩石特征(表1)。样品 SiO_2 质量分数为77.12%~77.75%,变化范围较小,含量较高,为酸性岩; TiO_2 为0.06%~0.08%,含量较低; Al_2O_3 为12.36%~12.53%,变化范围较小; Fe_2O_3 为0.88%~1.02%; MgO 为0.02%~0.05%,含量较低; K_2O 为3.87%~4.48%; Na_2O 质量分数为3.96%~

4.36%, 含量较高; $\text{Na}_2\text{O} + \text{K}_2\text{O}$ 为 8.07% ~ 8.61%。阿德拉嘎碱长花岗岩 TAS 图解中, 样品投点于花岗岩区域, 属于亚碱性系列(图 3a)。阿德拉嘎碱长花岗岩 QAP 图解中样品投点于碱长花岗岩区域(图 3b)。 $\text{K}_2\text{O}-\text{SiO}_2$ 图解(图 3c)中样品投点于高钾钙碱性系列区域。岩石分异指数 DI 为 96.75 ~ 97.41, 分异程度较高, 铝饱和指数 A/CNK 为 1.03 ~ 1.09, 为弱过铝质岩石。

3.2 稀土元素

样品稀土总量(ΣREE)为 59.46×10^{-6} ~ 162.29×10^{-6} , 含量较低, 变化范围较大, 轻稀土 LREE 含量为 27.35×10^{-6} ~ 124.23×10^{-6} , 重稀土 HREE 含量为 27.89×10^{-6} ~ 38.06×10^{-6} (表 1)。轻、重稀土 LREE/HREE 比值为 0.85 ~ 3.97, (La/Yb)_N 为 0.26 ~ 2.09, (Ce/Yb)_N 为 3.09 ~ 14.44, 轻重稀土分异程

度较低; (La/Sm)_N 为 1.17 ~ 2.59, 轻稀土分异程度较低; (Gd/Yb)_N 为 0.22 ~ 0.68, 重稀土分异程度较低; δEu 为 0.15 ~ 0.37, 负异常明显; δCe 为 0.84 ~ 8.20, 整体具弱正异常。稀土配分曲线呈海鸥型, 轻重稀土分异程度均较低(图 4a)。

3.3 微量元素

与天山—兴蒙造山系碱长花岗岩(迟清华和鄂明才, 2007)相比, 样品具有高 Hf、Nb, 低 Ba、Sr、U, 中等 Rb、Ta、Th、Ga、Zr 特征。样品微量元素蛛网图配分曲线一致(图 4b), 具有明显 Ba、Sr、P、Ti 负异常。

3.4 锆石 U-Pb 年龄

阿德拉嘎碱长花岗岩用于测试的锆石自形程度较好, 呈长柱状, 发育振荡环带, 具岩浆成因特征(Rubatto and Gebauer, 2000)。阿德拉嘎碱长花岗岩

表 1 阿德拉嘎碱长花岗岩主量元素(%)、稀土元素($\times 10^{-6}$)、微量元素($\times 10^{-6}$)组成

Table 1 Major element(%) ,REE ($\times 10^{-6}$) and trace element($\times 10^{-6}$) compositions of the Adelaga alkali feldspar granite

	P3YQ1	P3YQ2	P3YQ3	P3YQ4	P3YQ5	P3YQ6	*		P3YQ1	P3YQ2	P3YQ3	P3YQ4	P3YQ5	P3YQ6	*
SiO ₂	77.36	77.23	77.31	77.12	77.75	77.15	75.26	Eu	0.36	0.32	0.30	0.32	0.13	0.16	0.40
Al ₂ O ₃	12.42	12.44	12.40	12.47	12.36	12.53	12.98	Gd	5.17	2.89	3.88	5.89	2.35	2.75	5.6
Fe ₂ O ₃	0.99	0.96	0.88	1.02	0.95	0.88	0.90	Tb	1.03	0.84	0.92	1.28	0.86	0.85	0.59
FeO	0.07	0.06	0.17	0.05	0.05	0.05	0.46	Dy	7.40	7.75	7.12	9.62	8.24	7.83	
MnO	0.004	0.008	0.004	0.005	0.005	0.003	0.03	Ho	1.56	1.91	1.61	2.13	2.05	1.90	
TiO ₂	0.07	0.08	0.07	0.06	0.07	0.07	0.16	Er	4.79	6.32	5.12	6.78	6.73	6.23	
CaO	0.16	0.15	0.15	0.30	0.16	0.16	0.47	Tm	0.98	1.31	1.09	1.46	1.41	1.35	
MgO	0.03	0.04	0.02	0.04	0.05	0.05	0.14	Yb	6.24	8.06	6.89	9.23	8.91	8.62	2.9
Na ₂ O	4.13	4.13	4.12	4.36	4.20	3.96	3.71	Lu	1.15	1.46	1.26	1.67	1.56	1.57	0.45
K ₂ O	4.41	4.48	4.43	4.06	3.87	4.41	4.64	Y	41.25	56.46	44.13	63.64	58.83	56.02	23
P ₂ O ₅	0.039	0.018	0.036	0.012	0.012	0.021	0.04	ΣREE	140.89	80.75	103.10	162.29	59.46	109.51	
烧失	0.29	0.38	0.38	0.50	0.50	0.69		LREE	112.57	50.21	75.21	124.23	27.35	78.41	
总量	99.97	99.97	99.98	99.97	99.97	99.97		HREE	28.32	30.54	27.89	38.06	32.11	31.10	
Q	36.1	35.7	36.09	35.52	38.13	36.98		LREE/HREE	3.97	1.64	2.70	3.26	0.85	2.52	
An	0.54	0.63	0.51	1.42	0.72	0.66		(La/Yb) _N	2.09	0.84	1.24	2.03	0.26	0.27	
Ab	35.08	35.11	35.02	37.1	35.74	33.77		(Ce/Yb) _N	14.44	5.34	9.12	9.04	3.09	12.75	
Or	26.16	26.6	26.3	24.13	23	26.26		(La/Sm) _N	1.88	2.59	1.92	2.42	1.36	1.17	
C	0.66	0.57	0.64	0.39	1.01	1.01		(Gd/Yb) _N	0.68	0.30	0.46	0.52	0.22	0.26	
Hy	0.55	0.54	0.52	0.61	0.6	0.54		δEu	0.19	0.37	0.23	0.15	0.21	0.22	
DI	97.33	97.4	97.41	96.75	96.88	97		δCe	1.23	1.34	1.34	0.84	2.19	8.20	
A/CNK	1.06	1.05	1.05	1.03	1.09	1.09		Ba	15.0	23.1	15.5	42.6	51.0	40.1	325
SI	0.31	0.42	0.21	0.42	0.55	0.54		Hf	8.40	9.43	8.12	8.25	8.79	8.36	5.4
AR	4.82	4.82	4.82	4.87	4.63	4.32		Nb	18.26	18.40	18.28	22.08	20.86	25.87	16.0
R1	2635	2613	2633	2624	2770	2696		Rb	164.4	153.5	163.7	145.2	121.4	154.1	160
R2	263	263	261	280	264	267		Sr	7.7	8.9	7.3	14.3	12.9	13.0	70
La	18.19	9.49	11.92	26.16	3.20	3.21	29	Ta	1.58	1.92	1.55	1.49	1.43	1.98	1.56
Ce	55.16	26.34	38.45	51.08	16.83	67.26	63	Th	17.61	17.94	14.54	21.49	17.81	21.56	16.9
Pr	6.65	2.46	4.13	8.40	1.11	1.26		U	1.86	2.01	1.85	2.41	1.99	1.88	2.74
Nd	25.97	9.23	16.41	31.31	4.56	4.75	22	Ga	20.32	19.86	19.95	21.07	17.60	20.37	18
Sm	6.24	2.37	4.00	6.96	1.52	1.77	4.1	Zr	187.5	208.3	181.0	185.9	196.5	183.9	160

注: * 为天山—兴安造山系碱长花岗岩, 数据来源于(迟清华和鄂明才, 2007)。Alkali-feldspar granite of the Ten-zan-Khingan Orogenic System, based on data from (Chiqinghua and Yan Mingcai, 2007)。

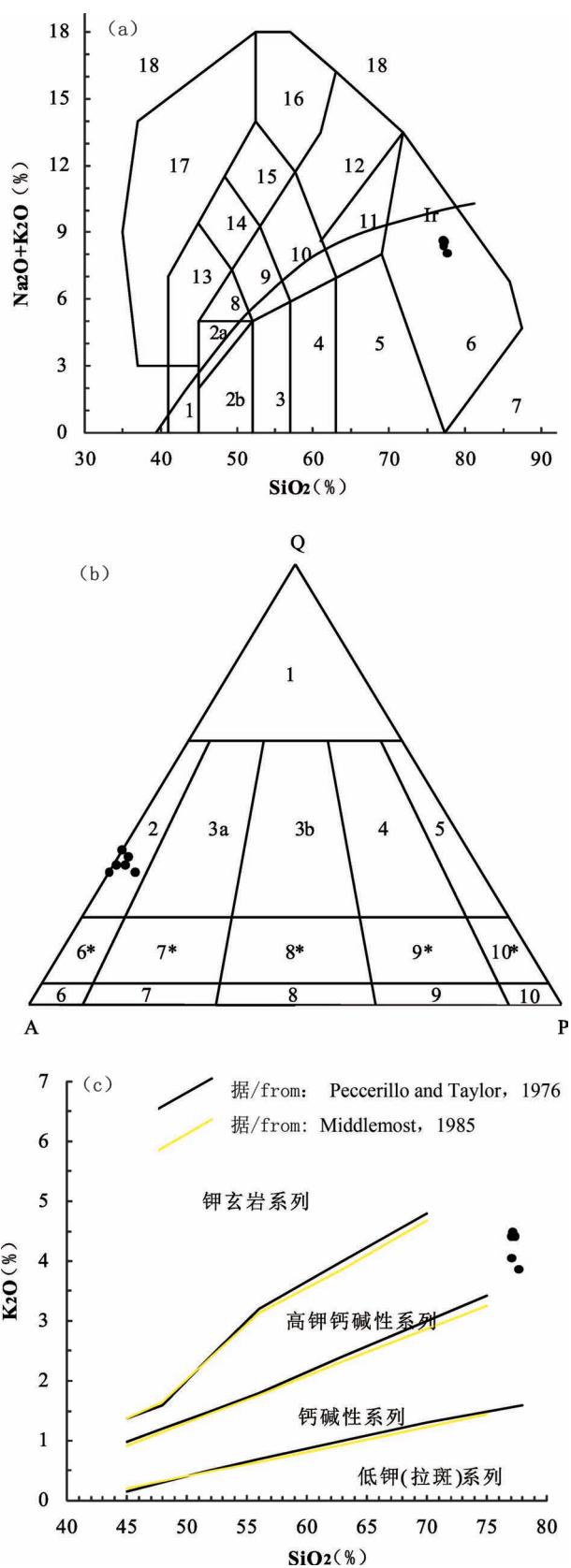


图 3 阿德拉嘎碱长花岗岩 TAS 图解 (a) (底图据 Middlemost, 1994)、QAP 图解 (b) (底图具 Streckeisen, 1976)、K₂O—SiO₂ 图解 (c) (底图据 Middlemost, 1994)

Fig. 3 TAS diagram (a) (after Middlemost, 1994), QAP diagram (b) (after Streckeisen, 1976) and K₂O—SiO₂ diagram (c) (after Middlemost, 1994) of the Adelaga alkali-feldspar granite

(a) Ir—Irvine 分界线, 上方为碱性, 下方为亚碱性 (Irvine and Baragar, 1971)。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—副长石二长闪长岩; 15—副长石二长正长岩; 16—副长正长岩; 17—副长深成岩; 18—霓方钠岩/磷霞岩/粗白榴岩

(b) 1—富石英花岗岩; 2—碱长花岗岩; 3a—花岗岩; 3b—花岗岩(二长花岗岩); 4—花岗闪长岩; 5—英云闪长岩、斜长花岗岩; 6*—碱长石英正长岩; 7*—石英正长岩; 8*—石英二长岩; 9*—石英二长闪长岩; 10*—石英闪长岩、石英辉长岩、石英斜长岩; 6—碱长正长岩; 7—正长岩; 8—二长岩; 9—二长闪长岩、二长辉岩; 10—闪长岩、辉长岩、斜长岩

石进行年龄测试, 在发育振荡环带的位置打点。锆石 Pb 含量 $5.351 \times 10^{-6} \sim 130.3 \times 10^{-6}$, Th 含量 $59.81 \times 10^{-6} \sim 1627 \times 10^{-6}$, U 含量 $89.72 \times 10^{-6} \sim 1525 \times 10^{-6}$ (表 2)。本样品锆石 Th/U 比值 $0.49 \sim 1.07 > 0.4$, 与变质成因锆石 ($\text{Th}/\text{U} \leq 0.1$) 差异明显, 表明锆石为岩浆成因锆石 (Hoskin and Black, 2000; Rubatto and Gebauer, 2000; Hoskin and Schaltegger, 2003; Cleasson et al., 2000; Belousova et al., 2002)。本样品 U-Pb 年龄 <1.0 Ga, 因而采用锆石 $^{206}\text{Pb}/^{238}\text{U}$ 年龄 (Griffin et al., 2004)。样品锆石 $^{206}\text{Pb}/^{238}\text{U}$ 分析点呈点群分布, 分布较为集中, 年龄值为 $294.9 \pm 3.2 \sim 316.2 \pm 3.1$ Ma (图 6a), 加权平均值为 310.7 ± 2.6 Ma, MSWD = 2.5 (图 6b), 表明岩体形成于晚石炭世。

4 讨论

4.1 年代学意义

贺根山蛇绿岩带两侧发育大规模碱性 (A 型) 花岗岩带 (洪大卫等, 2000; 张磊等, 2013; Zhang Xiaohui et al., 2014; 张晓飞等, 2018a)。查干敖包碱长花岗岩 (A 型) 锆石 U-Pb 年龄为 312.2 ± 7.4 Ma (焦天佳, 2015)。阿尔善布拉格细粒碱性花岗岩锆石 U-Pb 年龄为 295.8 ± 1.4 (肖中军等, 2015), 中粗粒碱性花岗岩锆石 U-Pb 年龄 297 ± 3 Ma (肖中军等,

锆石阴极荧光 (CL) 图像显示 (图 5), 锆石图像较暗, Th、U 含量略高。选择不发育裂隙和包裹体的锆

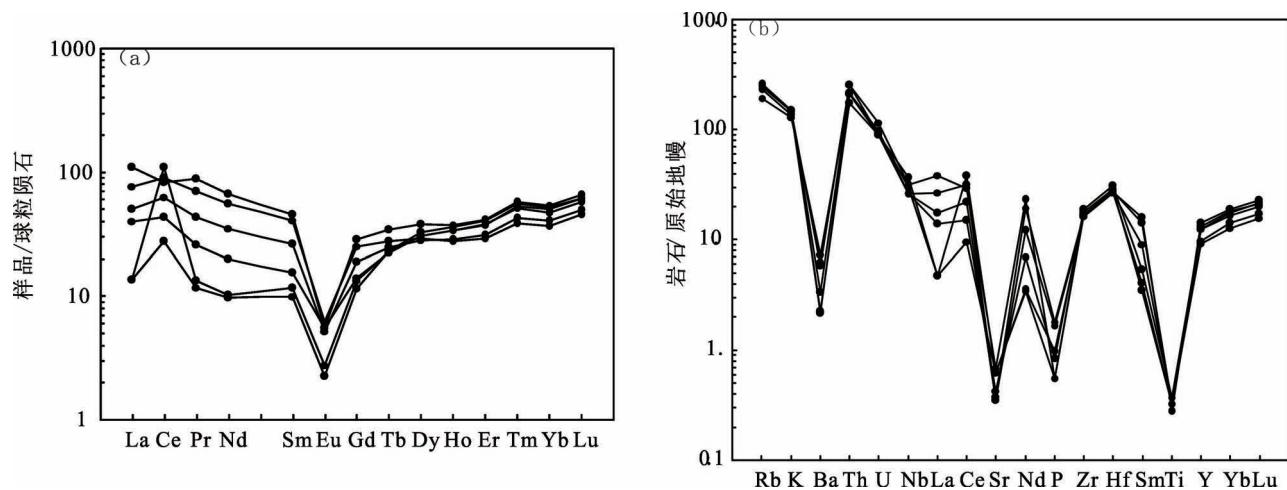


图 4 阿德拉嘎碱长花岗岩稀土元素球粒陨石标准化分布型式图(a)和微量元素原始地幔标准化
蛛网图(b)(标准化数据据 Sun and McDonough, 1989)

Fig. 4 Chondrite normalized REE patterns(a) and primitive mantle normalized trace elements spider diagram of the Adelaga alkali-feldspar granite(modified after Sun and McDonough, 1989)

2015),碱性花岗岩锆石U-Pb年龄为 $288.3\pm2.3\sim290.0\pm2.2$ Ma(Zhang Xiaohui et al., 2014)。京格斯台中粗粒碱长花岗岩(A型)锆石U-Pb年龄为 $295\pm$

2.8 Ma(刘建峰等,2009),碱性花岗岩锆石U-Pb年龄为 284.8 ± 1.1 Ma(张玉清等,2009)。白音乌拉碱性花岗岩锆石U-Pb年龄为 290 ± 2.6 Ma(Zhang

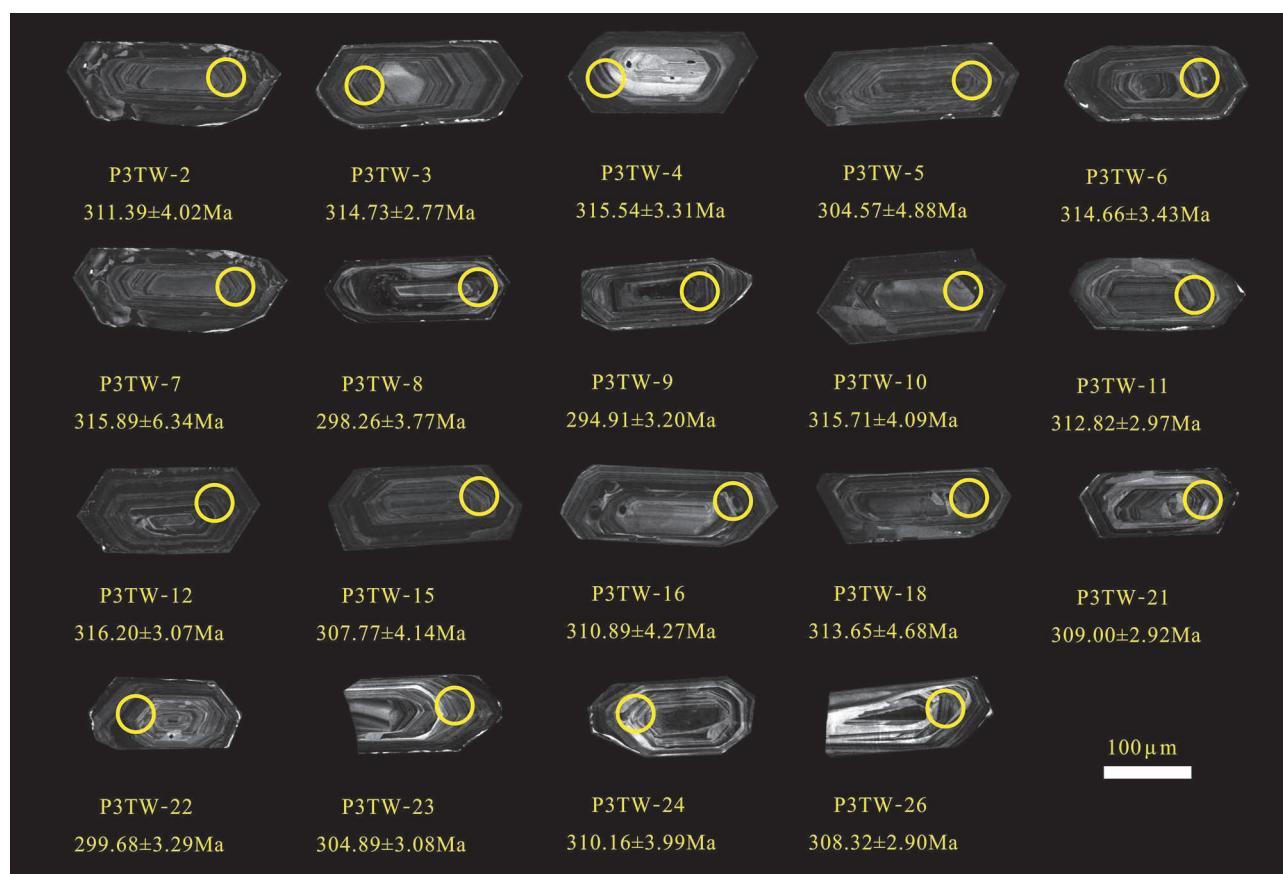


图 5 阿德拉嘎碱长花岗岩锆石阴极荧光(CL)照片

Fig. 5 Zircons cathodoluminescence (CL) images of the Adelaga alkali-feldspar granite

表 2 阿德拉嘎碱长花岗岩(P3TW)LA-ICP-MS 锆石同位素分析结果

Table 2 Zircon LA-ICP-MS analytical results of the Adelaga alkali feldspar granite (P3TW)

测点号	元素含量($\times 10^{-6}$)			同位素比值						同位素年龄(Ma)			$n(^{206}\text{Pb})/n(^{238}\text{U})$				
	Pb	Th	U	Th/U			$n(^{207}\text{Pb})/n(^{206}\text{Pb})$			$n(^{207}\text{Pb})/n(^{235}\text{U})$							
				测值	1σ	测值	测值	1σ	测值	测值	1σ	测值					
P3TW-2	69.53	958.7	1217	0.79	0.05160	0.0009	0.3521	0.0060	0.04949	0.0006	267.7	40.8	306.3	5.2	311.4	4.0	95%
P3TW-3	55.73	650.9	981.9	0.66	0.05326	0.0016	0.3674	0.0064	0.05003	0.0004	339.8	68.8	317.7	5.5	314.7	2.8	98%
P3TW-4	63.75	719.3	1110	0.65	0.05364	0.0023	0.3710	0.0081	0.05017	0.0005	356.0	98.8	320.4	7.0	315.5	3.3	97%
P3TW-5	30.99	390.8	578.4	0.68	0.05781	0.0020	0.3857	0.0065	0.04838	0.0008	522.7	74.2	331.2	5.6	304.6	4.9	98%
P3TW-6	5.351	59.81	89.72	0.67	0.05629	0.0085	0.3882	0.0232	0.05002	0.0005	463.6	335.1	333.1	19.9	314.7	3.4	95%
P3TW-7	83.40	938.3	1386	0.68	0.05340	0.0057	0.3698	0.0523	0.05022	0.0010	345.8	243.2	319.5	45.2	315.9	6.3	97%
P3TW-8	42.77	487.9	762.0	0.64	0.05812	0.0102	0.3795	0.0616	0.04735	0.0006	534.5	384.2	326.7	53.0	298.3	3.8	96%
P3TW-9	60.85	702.0	1130	0.62	0.05902	0.0023	0.3809	0.0075	0.04681	0.0005	567.9	84.4	327.7	6.5	294.9	3.2	98%
P3TW-10	69.28	781.6	1227	0.64	0.05400	0.0021	0.3736	0.0095	0.05019	0.0007	370.1	87.1	322.3	8.2	315.7	4.1	99%
P3TW-11	54.81	654.2	962.5	0.68	0.05566	0.0018	0.3816	0.0074	0.04972	0.0005	438.8	72.8	328.2	6.4	312.8	3.0	95%
P3TW-12	78.71	1045	1315	0.79	0.05530	0.0015	0.3833	0.0078	0.05027	0.0005	424.4	60.4	329.5	6.8	316.2	3.0	98%
P3TW-15	36.25	389.0	679.6	0.57	0.05695	0.0011	0.3840	0.0063	0.04890	0.0007	489.6	42.6	323.0	5.4	307.8	4.1	97%
P3TW-16	43.68	618.9	771.7	0.80	0.05230	0.0013	0.3563	0.0084	0.04941	0.0007	298.7	56.2	309.5	7.3	310.9	4.3	96%
P3TW-18	130.3	1627	1525	1.07	0.05429	0.0048	0.3732	0.0389	0.04986	0.0007	383.2	200.5	322.1	33.6	313.7	4.7	99%
P3TW-21	20.08	306.7	361.1	0.85	0.05541	0.0021	0.3752	0.0069	0.04910	0.0005	429.1	82.6	323.5	6.0	309.0	2.9	99%
P3TW-22	84.54	744.1	1508	0.49	0.05368	0.0032	0.3522	0.0136	0.04759	0.0005	357.6	134.0	306.4	11.8	299.7	3.3	98%
P3TW-23	71.53	983.6	1298	0.77	0.05429	0.0014	0.3626	0.0061	0.04843	0.0005	383.2	58.7	314.1	5.3	304.9	3.1	98%
P3TW-24	55.07	744.5	971.2	0.77	0.05325	0.0022	0.3619	0.0095	0.04929	0.0006	339.6	92.3	313.6	8.3	310.2	4.0	96%
P3TW-26	86.51	1374	1447	0.95	0.05561	0.0025	0.3756	0.0081	0.04899	0.0005	436.8	99.0	323.8	7.0	308.3	2.9	95%

Xiaohui et al., 2014), 285 ± 1 Ma (Tong Ying et al., 2015)。宝拉格碱性花岗岩锆石 U-Pb 年龄为 285 ± 1.1 Ma (Tong Ying et al., 2015)。红格尔碱性花岗岩锆石 U-Pb 年龄为 276 ± 1 Ma (Tong Ying et al., 2015)。塞因乌苏碱性花岗岩锆石 U-Pb 年龄为 279 ± 1 Ma (Tong Ying et al., 2015)。白音乌拉北细粒碱性花岗岩锆石 U-Pb 年龄为 280 ± 3 Ma, 形成于早二叠世(贾孝新等, 2017)。罕乌拉碱长花岗岩(A型)锆石 U-Pb 年龄为 279.8 ± 1.2 Ma (张晓飞等, 2018a)。扎拉嘎碱性花岗岩锆石 U-Pb 年龄为 272.3 ± 0.7 Ma 和 271.7 ± 0.7 Ma (程银行等, 2014)。锡林浩特晶洞花岗岩(A型)锆石 U-Pb 年龄为 276 ± 2 Ma (施光海等, 2004)。白音图嘎碱长花岗岩(A型)锆石 U-Pb 年龄为 294 ± 2 Ma, 形成于早二叠世(石文杰等, 2019)。拉抛碱长花岗岩(A型)锆石 U-Pb 年龄为 325 ± 3 Ma (张磊等, 2013)。区域资料表明碱性碱长(A型)花岗岩带主要形成于早二叠世, 少量形成于晚石炭世(表 3), 本文在阿德拉嘎碱长花岗岩体中获得的 LA-ICP-MS 锆石 U-Pb 年龄为 310.7 ± 2.6 Ma, 表明该岩体形成于晚石炭世, 形成时间相对较早, 为区域晚古生代岩浆活动在本区的响应。

4.2 岩石成因类型

阿德拉嘎碱长花岗岩手标本及镜下鉴定未见角闪石、白云母、堇青石、石榴子石等矿物, 样品中 P_2O_5 含量极低, 与 S型花岗岩特征存在较大差异 (Sylvester, 1998)。花岗岩具有高 Si、富碱 ($\text{Na}_2\text{O} + \text{K}_2\text{O} > 8\%$), 强烈 δEu 负异常 ($0.15 \sim 0.37$), 稀土配分曲线呈海鸥型, $10000\text{Ga}/\text{Al} = 2.68 \sim 3.19 > 2.6$, 表明岩体具 A型花岗

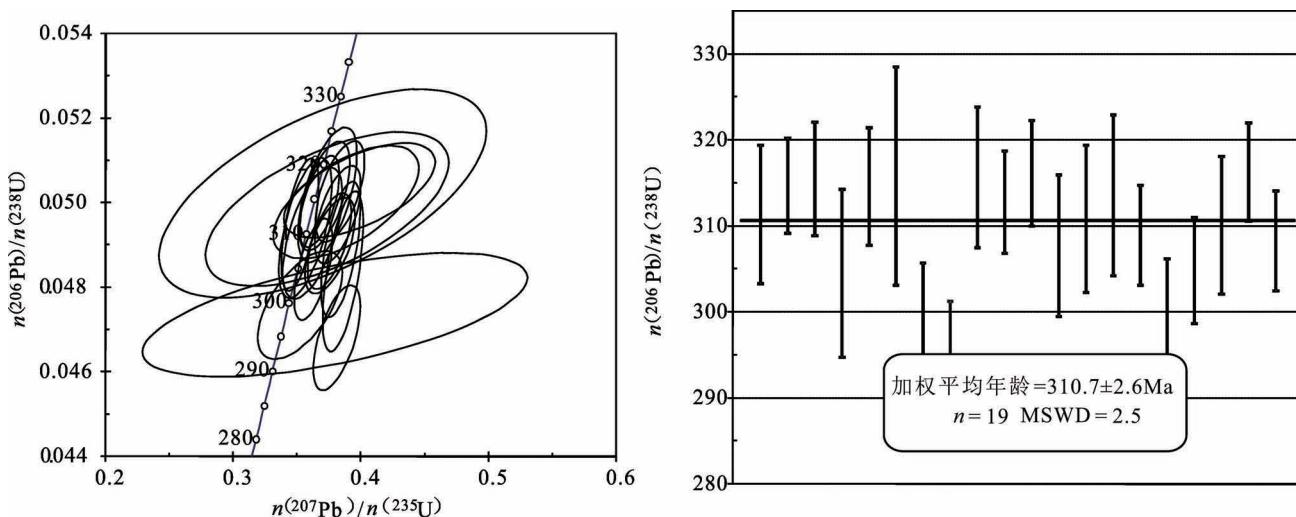


Fig. 6 U-Pb concordia diagram (a) and weighted mean ages diagram (b) of zircons of the Adelaga alkali-feldspar granite

岩的地球化学属性,与拉抛碱长花岗岩和罕乌拉碱长花岗岩等一致(Collins et al., 1982; Whalen et al., 1987; 张磊等,2013; 张晓飞等,2018a)。阿德拉嘎碱长花岗岩($\text{Na}_2\text{O} + \text{K}_2\text{O}$)/CaO—(Zr+Nb+Ce+Y)图解(图11)中样品投点于A型花岗岩区域。阿德拉嘎碱长花岗岩($\text{Na}_2\text{O} + \text{K}_2\text{O}$)/CaO—1000Ga/Al图解(图12)中样品投点于A型花岗岩区域。岩石地球化学特征表明,阿德拉嘎碱长花岗岩的岩石成因类型为A型花岗岩。

花岗岩的源岩部分熔融过程中,HREE中Yb主要富集于石榴子石中,Dy、Ho主要富集于角闪石中(Sisson,1994)。岩浆源区残留相中含有少量石榴子石(10%)时,其部分熔融形成的花岗岩具有高锶低钇(adakitic)特征,并且 Y/Yb 明显大于10(Hollocher et al., 2002), Y/Yb 接近10时,源区以角闪石为主,不含或含极少石榴子石(葛小月等,2002)。阿德拉嘎碱长花岗岩 Y/Yb 比值为6.40~7.00<10,岩浆源区以角闪石为主。

δEu 强烈负异常,表明岩浆源区存在大量斜长石。综合分析认为岩浆源区残留相为角闪石+斜长石。

4.3 构造环境及地质意义

高场强元素(HFSE)受后期作用影响较小,能够有效判别岩石形成的构造环境(Pearce et al., 1984)。Rb、Y(Yb)、Nb(Ta)等能够有效区分大洋脊花岗岩(ORG)、火山弧花岗岩(VAG)、板内花岗岩(WPG)和碰撞带花岗岩(Syn-COLG)等(Pearce et al., 1984)。Eby(1992)将A型花岗岩分为与洋岛玄武岩(OIB)类似,具有很低 Y/Nb , Yb/Ta 值的 A_1 型;与岛弧玄武岩(IAB)类似,具有高 Y/Nb ,

表3 贺根山断裂两侧晚石炭世—早二叠世碱性碱长花岗岩

Table 3 Alkali feldspar granite and aliali granite of Upper Carboniferous epoch and Lower Permian epoch both side of the Hegenshan fault

岩体名称	岩性	年龄(Ma)	测试方法	资料来源
查干敖包	碱长花岗岩	312.2±7.4	LA-ICP-MS	焦天佳,2015
阿尔善布拉格	细粒碱性花岗岩	295.8±1.4	LA-ICP-MS	肖中军等,2015
祖横德楞	中粗粒碱性花岗岩	297±3	LA-ICP-MS	肖中军等,2015
祖横德楞	碱性花岗岩	289±2.2	LA-ICP-MS	Zhang Xiaohui et al., 2014
京格斯台	中粗粒碱长花岗岩	295±2.8	LA-ICP-MS	刘建峰等,2009
京格斯台	碱性花岗岩	285±1	LA-ICP-MS	张玉清,2009
白音乌拉	碱性花岗岩	290±2.6	LA-ICP-MS	Zhang Xiaohui et al., 2014
白音乌拉	碱性花岗岩	285±1	LA-ICP-MS	Tong Ying et al., 2015
宝拉格	碱性花岗岩	285±1.1	LA-ICP-MS	Tong Ying et al., 2015
红格尔	碱性花岗岩	276±1	LA-ICP-MS	Tong Ying et al., 2015
塞因乌苏	碱性花岗岩	279±1	LA-ICP-MS	Tong Ying et al., 2015
白音乌拉北	细粒碱性花岗岩	279±2.7	LA-ICP-MS	贾孝新等,2017
扎拉嘎	碱性花岗岩	272.3±0.7	LA-ICP-MS	程银行等,2014
扎拉嘎	碱性花岗岩	271.7±0.7	LA-ICP-MS	程银行等,2014
罕乌拉	碱长花岗岩	279.8±1.2	LA-ICP-MS	张晓飞等,2018a
锡林浩特	晶洞花岗岩	276±2	LA-ICP-MS	施光海等,2004
白音图嘎	碱长花岗岩	294±2	LA-ICP-MS	石文杰等,2019
拉抛	碱长花岗岩	325±3	LA-ICP-MS	张磊等,2013

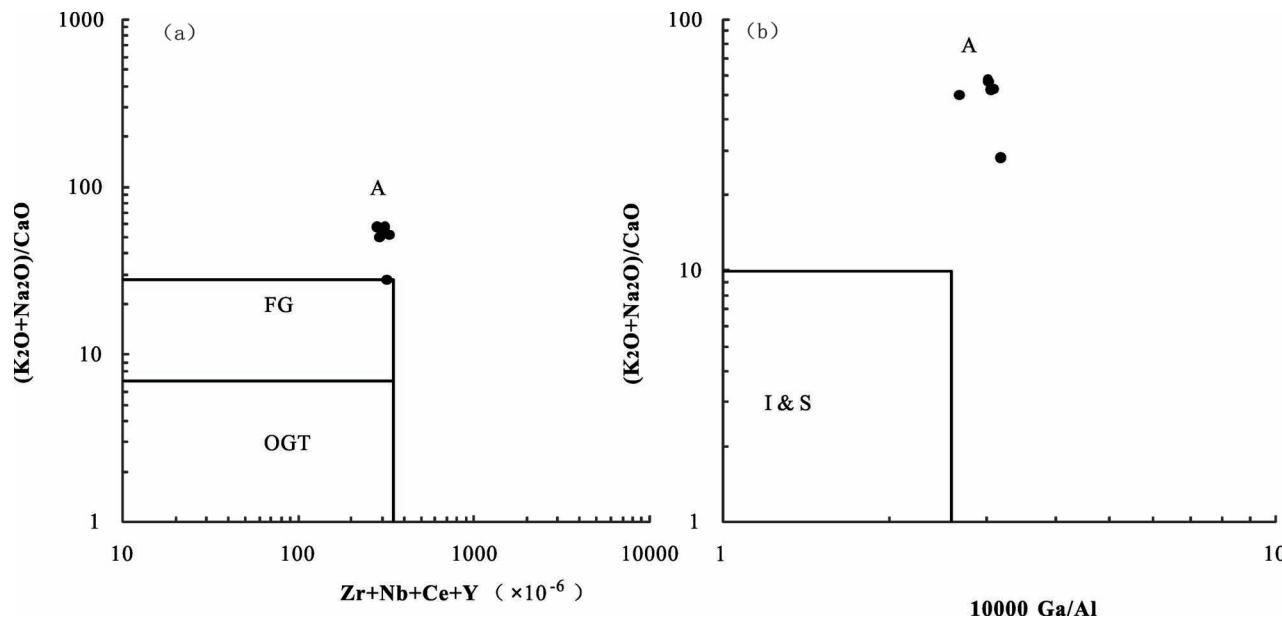


图 7 阿德拉嘎碱长花岗岩 $(\text{Na}_2\text{O}+\text{K}_2\text{O})/\text{CaO}$ — $\text{Zr}+\text{Nb}+\text{Ce}+\text{Y}$ 图解(a)和
 $(\text{Na}_2\text{O}+\text{K}_2\text{O})/\text{CaO}$ — $1000\text{Ga}/\text{Al}$ 图解(b)(底图具(Whalen et al., 1987))

Fig. 7 $\text{Na}_2\text{O}+\text{K}_2\text{O}/\text{CaO}$ — $\text{Zr}+\text{Nb}+\text{Ce}+\text{Y}$ diagram (a) and $\text{Na}_2\text{O}+\text{K}_2\text{O}/\text{CaO}$ — $1000\text{Ga}/\text{Al}$ diagram (b) of the Adelaga alkali feldspar granite(modified after Whalen et al., 1987)

Yb/Ta 值的 A_2 型。阿德拉嘎碱长花岗岩的 Y/Nb 值为 $2.17 \sim 3.07$, 平均 3.12 ; Yb/Ta 值为 $3.95 \sim 6.23$, 平均 5.87 , 都明显 >1 , 显示具岛弧玄武岩亲缘性。在 $\text{Nb}-\text{Y}-3\text{Ga}$ 和 $\text{Nb}-\text{Y}-3\text{Ce}$ 构造环境判别图解上(图 8a,b), 阿德拉嘎碱长花岗岩落在 A_2 型花岗岩区域。阿德拉嘎碱长花岗岩 $\text{Rb}-\text{Y}+\text{Nb}$ 图

解(图 9)样品投点于伸展构造背景(Pearce et al., 1984)。

迪彦庙蛇绿岩枕状玄武岩锆石 U-Pb 年龄为 $333.4 \pm 8.5\text{ Ma}$, 形成于俯冲环境(李英杰等, 2018)。跃进地区花岗岩类锆石 U-Pb 年龄为 $310 \sim 330\text{ Ma}$, 具岛弧侵入岩特征, 形成于俯冲背景下(王树庆等,

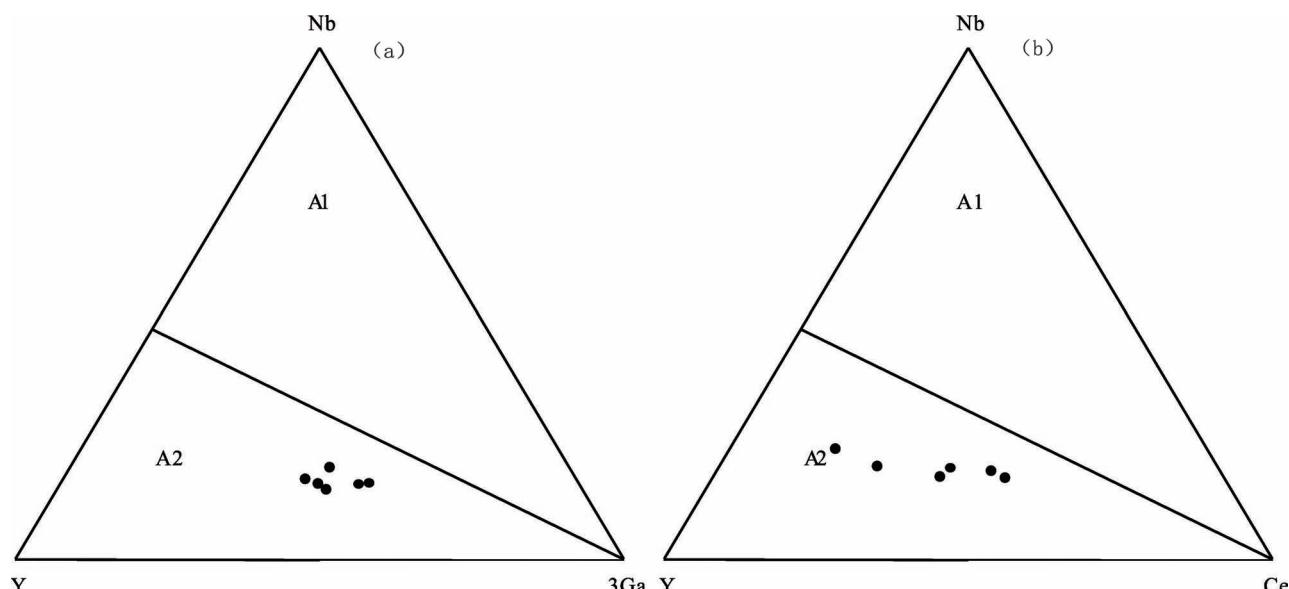


图 8 阿德拉嘎碱长花岗岩 $\text{Nb}-\text{Y}-3\text{Ga}$ 图解(a)和 $\text{Nb}-\text{Y}-\text{Ce}$ 图解(b)(底图具 Eby, 1992)

Fig. 8 $\text{Nb}-\text{Y}-3\text{Ga}$ diagram (a) and $\text{Nb}-\text{Y}-\text{Ce}$ diagram (b) of the Adelaga alkali feldspar granite(modified after Eby, 1992)

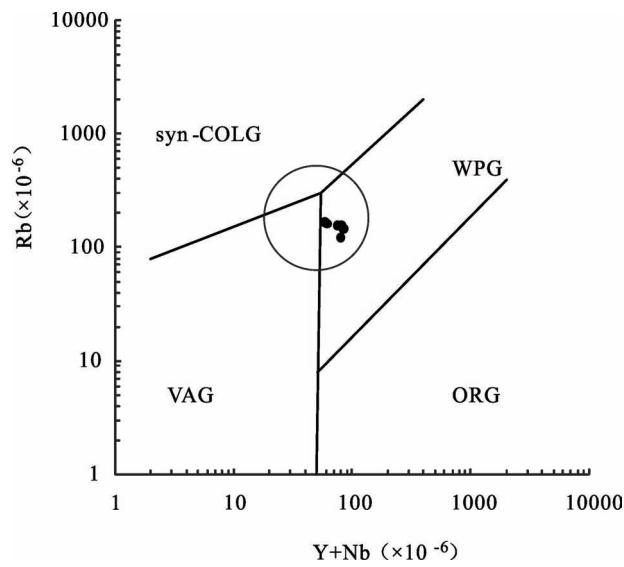


图 9 阿德拉嘎碱长花岗岩 Rb—(Y+Nb) 图解

(Pearce et al., 1984)

Fig. 9 Rb—(Y+Nb) diagram of the Adelaga alkali feldspar granite (modified after Pearce et al., 1984)

2018)。迪彦庙—达青蛇绿岩斜长花岗岩锆石 U-Pb 年龄为 330.1 ± 3.2 Ma、 316.5 ± 2.4 Ma, 辉长岩、高镁闪长岩锆石 U-Pb 年龄分别为 286.1 ± 6.1 Ma、 283.7 ± 4.7 Ma, 结合地球化学和 Hf 同位素数据, 区域石炭纪—二叠纪处于洋盆扩张—洋内俯冲阶段 (程杨等, 2019, 2020)。苏尼特右旗蛇绿岩带中发育英云闪长岩, 锆石 U-Pb 年龄为 322.3 ± 5.2 Ma, 具高锶低钇 (adakitic) 特征, 表明古亚洲洋晚石炭世仍在俯冲 (董培培等, 2021)。迪彦庙奔来可图岩体锆石 U-Pb 年龄为 322.2 ± 1.3 Ma, 形成于古亚洲洋俯冲环境 (范玉须等, 2020)。巴彦都兰角闪石岩锆石 U-Pb 年龄为 310 ± 1 Ma, 地球化学结合电子探针和 Hf 同位素研究认为, 其形成于古亚洲洋俯冲背景下 (张夏炜等, 2021)。梅劳特乌拉 SSZ 型蛇绿岩呼和浩特花岗闪长岩 U-Pb 年龄分别为 294.4 ± 1.7 Ma, 呼都格奥长花岗岩锆石 U-Pb 年龄分别为 306.3 ± 1.9 Ma 和 315.5 ± 1.9 Ma, 白音呼舒奥长花岗岩锆石 U-Pb 年龄为 309.2 ± 1.6 Ma, 均具高锶低钇 (adakitic) 特征; 蛇绿岩中高镁安山岩锆石 U-Pb 年龄 315.0 ± 2.3 Ma, 形成于古亚洲洋俯冲环境 (王金芳等, 2017, 2019, 2020, 2021)。沙巴尔吐蛇绿构造混杂岩超镁铁质岩和碱长花岗岩 279.2 ± 3.3 Ma 和 278.2 ± 1.7 Ma, 结合地球化学和 Hf 同位素研究, 表明古亚洲洋在中二叠世早期尚未闭合 (李英雷等, 2021)。索伦山蛇绿岩带硅质岩中发现早二叠世放射虫动物群, 表明古亚

洲洋俯冲消减闭合应在早二叠世后 (李钢柱等, 2017)。1 : 1000000 地质编图和野外地质调查成果表明大兴安岭地区到中—晚二叠世, 古亚洲洋盆持续俯冲 (杨晓平等, 2022)。大兴安岭海西期花岗岩类研究表明, 晚泥盆世—中二叠世区域广发发育弧岩浆岩, 晚二叠世—早三叠世, 古亚洲洋闭合 (刘锐等, 2016)。九井子蛇绿岩中辉绿岩脉锆石 U-Pb 年龄为 274.7 ± 1.7 Ma, 与蛇绿岩带呈断层接触的粉砂岩碎屑锆石年龄最小值为 249 ± 4.7 Ma, 研究表明, 索伦—西拉木伦蛇绿岩带闭合于晚二叠世末—早三叠世初 (刘建峰等, 2016)。色尔崩岩体和努和亭沙拉岩体锆石 U-Pb 年龄为 255.3 ± 1.4 Ma、 254.4 ± 3.4 Ma, 具高锶低钇 (adakitic) 特征, 形成于晚二叠世俯冲背景下 (范玉须等, 2019)。古地磁数据表明, 西伯利亚和华北板块闭合于二叠纪末期 (~250 Ma) (李朋武等, 2006)。林西地区晚二叠世晚期生物礁和海象化石的发现和马达屯期大规模火山喷发以及巴林右旗和吉林九台县的 P/T 角度不整合表明, 古亚洲洋在该区域晚二叠世末期闭合 (田树刚等, 2016)。

研究表明伸展背景包括大陆裂谷带、大陆减薄区、碰撞后伸展环境、与俯冲有关的洋内岛弧、活动陆缘和弧后盆地等多种环境 (Hochstaedter et al., 1990; 王焰等, 2000)。晚石炭世古亚洲洋俯冲背景下, 贺根山区域晚石炭—早二叠发育洋内弧, 阿德拉嘎地区处于弧后环境, 结合 A 型花岗岩特征, 综合分析认为, 阿德拉嘎碱长花岗岩形成于弧后伸展构造背景。

5 结论

(1) 锆石 LA-ICP-MS U-Pb 测年表明, 阿德拉嘎碱长花岗岩的侵位年龄为 310.7 ± 2.6 Ma, 形成时代为晚石炭世。

(2) 阿德拉嘎碱长花岗岩具有较高的 SiO₂、Na₂O+K₂O 含量和较高的 Ga/Al、(Na₂O+K₂O)/CaO 比值, 相对贫 CaO、MgO、Sr、Ba、Eu、Ti、P, 稀土元素总量较低, 轻重稀土分馏不明显, 稀土配分曲线呈海鸥型, 负铕异常明显 (δ Eu = 0.15~0.37), 为 A 型花岗岩。

(3) 岩石学和岩石地球化学特征表明, 阿德拉嘎碱长花岗岩为 A 型花岗岩, 可能形成于弧后伸展构造环境, 为晚石炭世古亚洲洋向西伯利亚板块俯冲作用的产物。

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注 释 / Notes

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Zircon LA-ICP-MS U-Pb age, geochemical character of Adelaga alkali-feldspar granite, in Abaga Banner, Inner Mongolia, and tectonic significance

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Objectives: This paper provided petrography, geochemical characteristics, LA-ICP-MS zircon U-Pb age of Adelaga alkali feldspar granite, in Abaga Banner, Inner Mongolia. The formation age, genesis and tectonic significance of granites are discussed.

Methods: Based on the field work, through the microscopic observation, the whole rock chemical analysis, the LA-ICP-MS zircon U-Pb isotopic chronology of granites.

Results: The Zircon LA-ICP-MS U-Pb ages indicates that the Adelaga alkali feldspar granite is 310.7 ± 2.6 Ma, and formed in Upper Carboniferous. The granite has the character of Higher content of SiO_2 , $\text{Na}_2\text{O} + \text{K}_2\text{O}$, higher ratio of Ga/Al , $(\text{Na}_2\text{O} + \text{K}_2\text{O})/\text{CaO}$, and relatively poor CaO , MgO , Sr , Ba , Eu , Ti and P . The total amount of rare earth elements is low, the fractionatio of light and heavy rare elements is not aobious, the distribution curve of rare eathr is seagull, and the anomaly of negative europium is obvious ($\delta\text{Eu} = 0.15 \sim 0.37$). The granite belong to A-type granite.

Conclusions: The Adelaga alkali feldspar granite is A-type granite, and formed in back-arc extensional tectonic setting of Upper Carboniferous, which is the product of Paleo-Asian Oceanic subduction to Siberia.

Keywords: Alkali feldspar granite; A-type granite; Zircon U-Pb ages; Upper Carboniferous; Back-arc extensional environment; Abag Banner, Inner Mongolia.

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