

# 西天山东塔尔别克金矿区安山岩 LA-ICP-MS 锆石 U-Pb 年代学、元素地球化学与岩石成因\*

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**Abstract** The eastern Taerbieke gold deposit is geographically located to the south of the Axi gold deposit in the Tulasu basin, western Tianshan, and tectonically occurs in the western section of the Early Paleozoic Borohoro island arc between the Junggar plate to the north and the Yili-central Tianshan micro-plate to the south. The eastern Taerbieke andesites are geochemically similar to high-Mg andesite, e. g., high SiO<sub>2</sub> (58.94% ~ 63.85%) and MgO contents (3.75% ~ 6.59%), Mg<sup>#</sup> values (58 ~ 69), and Cr (94.2 × 10<sup>-6</sup> ~ 241 × 10<sup>-6</sup>) and Ni (54.5 × 10<sup>-6</sup> ~ 126 × 10<sup>-6</sup>) contents, and low FeO<sup>T</sup>/MgO ratios. Except for slight low Sr contents (235 × 10<sup>-6</sup> ~ 696 × 10<sup>-6</sup>), they are also geochemically similar to adakites, e. g., high Al<sub>2</sub>O<sub>3</sub> (15.39% ~ 16.65%), high Sr/Y (23.8 ~ 48.1) ratios, no significant Eu anomalies and clear depletion in Nb, Ta and Ti. LA-ICP-MS U-Pb age data suggest that the eastern Taerbieke andesites were generated in the Early Carboniferous (347.2 ± 1.6 Ma). The enrichment in large ion lithophile elements (e. g. Rb, Th and U) and depletion in high field strength elements (e. g. Nb, Ta and Ti), and higher light rare earth element contents relative to heavy rare earth elements for the andesites, suggest that they are similar to island arc rocks. We suggest that the eastern Taerbieke andesites were possibly generated by the interaction between mantle and melts derived by partial melting of subducting sediments and altered oceanic crust during the subduction of Late Paleozoic northern Tianshan Ocean. The interaction between mantle and melts possibly played an important role in generating Au mineralization.

**Key words** Zircon LA-ICP-MS U-Pb dating; High-Mg adakite; Gold mineralization; Island setting; West Tianshan

**摘要** 东塔尔别克金矿区位于西天山吐拉苏盆地内阿希金矿区南侧, 构造上属于伊犁-中天山微板块与准噶尔板块之间的博罗科努早古生代岛弧带西段。东塔尔别克矿区出露有一些安山岩, 这些岩石具有典型高镁安山岩特征, 如高硅(SiO<sub>2</sub> = 58.94% ~ 63.85%), 具有较高的 MgO 含量(3.75% ~ 6.59%) 和 Mg<sup>#</sup> (58 ~ 69)、高的 Cr (94.2 × 10<sup>-6</sup> ~ 241 × 10<sup>-6</sup>)、和 Ni

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( $54.5 \times 10^{-6} \sim 126 \times 10^{-6}$ )含量,以及低的  $\text{FeO}^T/\text{MgO}$  比值。除了 Sr 含量略微偏低( $235 \times 10^{-6} \sim 696 \times 10^{-6}$ )之外,这些岩石基本显示了埃达克岩的地球化学特征:高的  $\text{Al}_2\text{O}_3$  (15.39% ~ 16.65%),低 Y( $9.86 \times 10^{-6} \sim 14.9 \times 10^{-6}$ )含量,以及高的 Sr/Y 比值(23.8 ~ 48.1),无 Eu 异常,Nb、Ta、Ti 亏损等。LA-ICP-MS 锆石 U-Pb 测年结果显示安山岩的年龄为  $347.2 \pm 1.6\text{Ma}$ ,为早石炭世。安山岩富集 Rb、Th、U 等大离子亲石元素,相对亏损 Nb、Ta、Ti 等高场强元素,轻重稀土元素分馏明显,具有俯冲带岛弧岩浆的特征。东塔尔别克安山岩可能形成于岛弧环境中,并可能由俯冲的晚古生代北天山洋洋壳以及少量上覆沉积物熔融形成,产生的熔体在上升过程中与地幔橄榄岩发生了相互作用。熔体-地幔相互作用对区内金矿的形成具有重要意义。

**关键词** LA-ICP-MS 锆石 U-Pb 定年;高镁埃达克岩;金矿化;岛弧;西天山

**中图法分类号** P588.144;P597.3

## 1 引言

天山造山带是古亚洲洋造山带中的一个重要的造山带。一般认为天山造山带是在古生代由塔里木和西伯利亚板块之间的古亚洲洋的消减闭合,塔里木、准噶尔、哈萨克斯坦等板块的俯冲-碰撞-增生所形成(Coleman, 1989; Windley *et al.*, 1990; Allen *et al.*, 1993; Sengör *et al.*, 1993; Shi *et al.*, 1994; Gao *et al.*, 1998; Xiao *et al.*, 2004; Liu and Fei, 2006)。其中在西天山的伊犁盆地的南北缘发育有大量的石炭纪岩浆岩(Xia *et al.*, 2004b, 2008; Zhu *et al.*, 2005; 王强等, 2006; 朱永峰等, 2006a,b; 朱志新等, 2006; 龙灵利等, 2008)。前人对西天山北部的石炭纪岩浆岩做了大量的工作,但目前对这些石炭纪岩浆岩的形成背景仍存在不同的认识,主要有大陆边缘岛弧(Gao *et al.*, 1998; Zhang *et al.*, 2003; Liu and Fei, 2006; Wang *et al.*, 2007b; Xiao *et al.*, 2008; Qian *et al.*, 2009; 王强等, 2006; 李锦轶等, 2006; 龙灵利等, 2008)还是裂谷(Xia *et al.*, 2004b, 2008; 车自成和刘良, 1996)或地幔柱环境(夏林圻等, 2004)之争。

天山造山带也是一个重要的 Cu-Au 等多金属成矿带,天山北缘出露有许多铜金矿床(点),如阿希、土屋、延东等。其中许多石炭纪岩浆岩与铜金成矿作用十分密切(Qin *et al.*, 2002; Rui *et al.*, 2002; Zhang *et al.*, 2006b, 2008; 王强等, 2006)。通过对岩浆岩的深入研究,有可能对深入了解金属矿床的成因、构造背景和地球动力学提供重要的启示。

本文首次对西天山北缘的阿希金矿区南侧的东塔尔别克金矿区的岩浆岩进行了 LA-ICP-MS 锆石 U-Pb 同位素年代学和元素地球化学研究,发现该区岩浆岩形成于早石炭世,并与阿希金矿的岩浆岩有类似的地球化学特征。本文将重点报道这一成果,并初步探讨其成因和形成的构造背景。

## 2 地质背景与岩相学

塔尔别克金矿位于科古琴山南坡断裂与伊犁盆地北缘断裂之间的吐拉苏盆地内(图 1b),南为伊犁-中天山板块,北为夹于伊犁-中天山板块与准噶尔板块之间的博罗科努早古生代岛弧带西段(图 1a)(Gao *et al.*, 1998)。塔吾尔别克金矿与阿希、京希、伊尔曼得、恰布坎卓它等金矿一起构成了

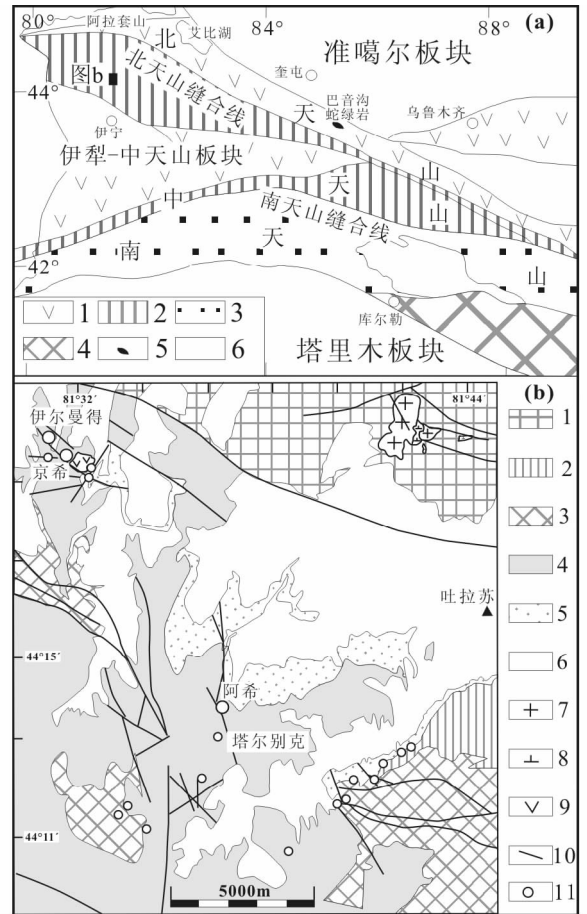


图 1 天山地区大地构造简图(a,据 Gao *et al.*, 1998 修改)和吐拉苏地区地质简图(b,据左学义等, 2005①)

(a): 1-晚古生代火山岩或火山碎屑岩;2-前寒武基底、古生代花岗岩和地层;3-古生代沉积岩;4-前寒武角闪岩相岩石;5-蛇绿岩;6-中新生带地层。(b): 1-前寒武纪;2-泥盆纪;3-早古生代;4-石炭纪大哈拉军组;5-石炭纪阿恰勒组;6-新生代沉积物;7-花岗岩;8-闪长岩;9-安山玢岩;10-断层;11-矿床或矿点

Fig. 1 Simplified geological map of the western Tianshan (a, after Gao *et al.*, 1998) and regional geological sketch map of the Tulasu areas (b)

① 左学义, 漆树基, 伊发源. 2005. 吐拉苏金矿带大型金矿床定位预测. 新疆优势矿产资源勘查评价研究(305 项目, 2003BA12A-06-10)

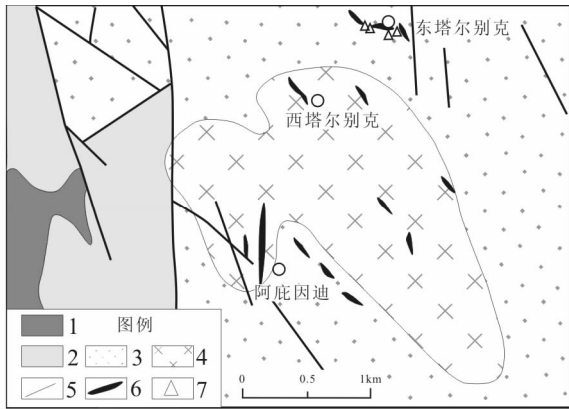


图2 塔尔别克金矿矿区地质简图(据贾斌等,2001 修改)

1-奥陶系奈楞格勒达板组;2-大哈拉军山组第1,2和4岩性段;3-大哈拉军山组第5岩性段;4-二长斑岩;5-断层;6-金矿体;7-采样点

Fig.2 Geological sketch map of the Taerbieke gold deposit (after Jia *et al.*, 2001)

吐拉苏金矿带。该区石炭纪岩浆岩分布广泛,并主要以石炭纪火山岩为主(Zhu *et al.*, 2005; 王强等, 2006; 朱志新等, 2006; 李华芹等, 2006; 龙灵利等, 2008)。该区出露的石炭纪火山岩属于大哈拉军山组。西天山地区的大哈拉军山组火山岩主要由玄武安山岩、安山岩、粗面安山岩、粗面岩、凝灰岩和英安岩组成(Zhu *et al.*, 2005; 朱永峰等, 2005; 左学义等, 2005; 龙灵利等, 2008), 其形成时代为 354 ~ 312Ma (Zhu *et al.*, 2005)。

塔尔别克金矿区包括东西两个部分(图2)。西塔尔别克金矿区以二长花岗斑岩为主,东塔尔别克金矿区以火山岩为主,后者也是本文研究的重点。东塔尔别克金矿区的西侧出露大哈拉军组第1,2和4岩性段火山岩,它们与奥陶系奈楞格勒达板组不整合接触。矿区广泛出露的地层为大哈拉军组第5岩性段火山岩,岩性主要由辉石安山岩、英安岩和凝灰岩组成。其中新鲜安山岩为紫红色,斑状结构,斑晶约占 5% ~ 10%, 以斜长石为主,同时含有少量的角闪石和普通辉石,其中斜长石呈自形板状的居多,最大可达 3mm,斜长石和辉石为半自形,基质以斜长石为主,斜长石有些短柱状,有些呈针状,具有玻晶交织结构,还有少量辉石和 Fe-Ti 氧化物。而蚀变的安山岩为土黄色,也为斑状结构,斑晶以斜长石为主,并大多已经绢云母化,还含有角闪石和石英,其中有些角闪石出现暗化边,有些蚀变为绿泥石或黑云母,基质为微细粒的绢云母化斜长石和钾长石、石英、角闪石和黑云母等,角闪石和黑云母均也都出现暗化边,另外,岩石中有些碳酸盐矿物充填。

### 3 分析方法

主、微量元素的分析测试均在中国科学院广州地球化学

研究所同位素年代学和地球化学重点实验室完成。主量元素分析是用 Rigaku RIX 2000 型荧光光谱仪(XRF)分析,其详细步骤与 Li *et al.* (2005)所述相同。样品的含量由 36 种涵盖硅酸盐样品范围的参考标准物质双变量拟合的工作曲线确定,基体校正根据经验的 Traill-Lachance 程序进行,分析精度优于 1% ~ 5%。微量元素的分析则采用 Perkin-Elmer Sciex ELAN 6000 型电感耦合等离子体质谱仪(ICP-MS),具体的流程见 Li (1997)。使用 USGS 标准 W-2 和 G-2 及国内标准 GSR-1、GSR-2 和 GSR-3 来校正所测样品的元素含量,分析精度一般为 2% ~ 5%。分析数据列于表 1。

为精选锆石样品,先将新鲜的岩石样品粉碎至 120 目以下,用常规的人工淘洗和电磁选方法富集锆石,再在双目镜下用手工方法逐个精选锆石颗粒,未用任何化学试剂。本次锆石定年样品和主元素和微量元素分析的样品相对应。锆石阴极发光图像研究在中国科学院广州地球化学研究所 JXA-8100 电子探针仪上完成。锆石 LA-ICP-MS U-Pb 年龄测定在中国地质大学(武汉)地质过程与矿产资源国家重点实验室完成。ICP-MS 为 Agilent 公司生产的四极杆 ICP-MS Agilent 7500a,激光剥蚀系统为德国 Lamda Physik 公司的 GeoLas 2005 深紫外(DUV) 193 nm ArF 准分子(excimer)激光剥蚀系统。实验中采用 He 作为剥蚀物质的载气,激光斑束直径为 32 $\mu$ m。参考物质为美国国家标准技术协会研制的人工合成硅酸盐玻璃 NIST SRM610,锆石 U-Pb 年龄的测定采用国际标准锆石 91500 作为外标进行校正,每隔 5 个分析点测一次标准,保证标准和样品的仪器条件一致。在样品分析前后以及每隔 20 个测点各测一次 NIST SRM610,以 Si 做内标,测定锆石中的 U, Th, Pb 的含量。详细的分析流程及有关参数见(Yuan *et al.*, 2004; Liu YS *et al.*, 2008)。元素的比率和元素的含量用 GLITTER(4.0 版)来处理,年龄的计算和谐和图用 ISOPLOT(3.00 版)(Ludwig, 2003)来完成。分析数据列于表 2。

## 4 分析结果

### 4.1 LA-ICPMS 锆石 U-Pb 定年

东塔尔别克安山岩样品 06XJ-31 的锆石多呈棱柱状,长约 40 ~ 120 $\mu$ m,长宽比在 1 : 1.5 ~ 3 : 1。阴极发光图像显示发育有清晰的韵律结构(图3),锆石具有比较均一的 U/Th 比(1.27 ~ 3.30),大部分在 1.5 ~ 2.0 之间,为典型的岩浆锆石(U/Th > 1)。25 个分析点中有 23 个<sup>206</sup>Pb/<sup>238</sup>U 表面年龄在 340 ~ 351Ma 之间,在误差范围内比较一致,其加权平均值为 347.2 ± 1.6Ma (MSWD = 0.68) (图3)。分析点 9 落在谐和线上,其<sup>206</sup>Pb/<sup>238</sup>U 表面年龄为 447 ± 6Ma,该锆石成板柱状,无岩浆环带(图3),可能为奥陶纪火山岩的捕获晶。另有一个分析点(点6)也靠近谐和线,其<sup>206</sup>Pb/<sup>238</sup>U 表面年龄为 1009 ± 10Ma,该锆石具有浑圆状的核部,外缘发育有岩浆环带(图3),其可能是岩浆上升过程中捕获的基底岩浆锆石。

表1 东塔尔别克和阿希金矿安山岩主量(wt%)和微量元素( $\times 10^{-6}$ )组成Table 1 Major element (wt%) and trace element ( $\times 10^{-6}$ ) compositions of the andesites in east Taerbieke and Axi gold deposit

样品号	06XJ-30	06XJ-31	06XJ-32	06XJ-33	wxt818	wxt819	wxt821	AX03	AX04	AX14
岩石类型	蚀变安山岩	蚀变安山岩	安山岩	安山岩	安山岩	安山岩	安山岩	安山岩	安山岩	玄武安山岩
地点	东塔尔别克金矿				阿希金矿*					
SiO <sub>2</sub>	63.85	62.02	58.94	61.32	57.93	56.23	60.10	56.59	56.01	53.30
TiO <sub>2</sub>	0.48	0.67	0.81	0.77	0.72	0.74	0.68	0.87	0.88	0.95
Al <sub>2</sub> O <sub>3</sub>	15.90	16.01	15.39	16.65	15.19	15.26	14.28	17.08	17.21	18.22
Fe <sub>2</sub> O <sub>3</sub>	3.77	5.03	5.91	5.28	5.36	7.26	5.38	7.10	7.15	8.06
MnO	0.05	0.07	0.10	0.11	0.06	0.07	0.07	0.09	0.08	0.08
MgO	2.51	3.79	6.59	3.75	7.65	8.47	6.78	4.56	4.04	3.36
CaO	0.70	0.98	2.51	2.18	4.90	4.38	5.88	6.98	7.21	7.21
Na <sub>2</sub> O	2.04	4.87	4.63	6.23	2.54	2.65	2.26	2.64	2.76	3.09
K <sub>2</sub> O	5.92	2.17	1.64	1.04	1.76	1.45	0.66	0.85	0.64	1.08
P <sub>2</sub> O <sub>5</sub>	0.08	0.08	0.11	0.10	0.08	0.06	0.07	0.18	0.18	0.23
LOI	3.99	3.75	3.72	3.03	3.36	3.92	3.20	3.08	3.95	4.11
Total	99.30	99.43	100.35	100.45	99.55	100.49	99.36	103.41	103.32	103.85
Mg <sup>#</sup>	57.00	60.00	69.00	58.00	73.87	69.80	71.40	55.99	52.81	45.23
Sc	8.23	10.9	17.0	10.8	17.0	15.3	17.4	25.2	25.1	20.4
V	75.9	72.4	94.7	76.4	117	118	114	177	253	158
Cr	60.1	70.1	241	94.2	790	768	789	57.4	56.4	10.3
Co	11.4	16.2	22.9	18.6	26.6	27.2	26.4	19.4	20.3	21.4
Ni	35.9	37.1	126	54.5	182	183	186	21.2	21.0	9.20
Ga	17.6	18.7	16.8	17.5	16.1	16.1	15.1	18.5	19.0	20.2
Cs	8.18	5.72	8.11	9.06				2.12	4.12	2.76
Rb	231	86.1	53.6	27.7	59.4	29.5	14.2	24.2	18.5	27.0
Ba	368	345	299	371	295	287	198	214	198	289
Th	4.96	4.76	4.75	5.36	5.50	5.11	4.94	3.75	3.72	4.04
U	1.50	1.36	1.13	1.57	1.58	1.37	1.23	1.06	1.09	1.07
Pb	6.36	7.45	8.20	6.08	12.2	11.4	8.47	8.94	8.83	7.97
Nb	5.83	6.92	9.07	9.53	6.02	5.74	5.35	6.84	6.96	13.3
Ta	0.471	0.559	0.705	0.746	0.536	0.473	0.444	0.510	0.510	0.620
Sr	235	354	686	697	297	294	317	353	387	365
Y	9.86	12.2	14.3	14.8	17.4	17.2	16.6	20.5	21.7	21.3
Zr	149	176	159	168	128	114	109	122	125	171
Hf	3.66	4.19	3.61	3.87	3.58	3.08	2.91	3.64	3.65	3.94
La	13.6	12.7	13.4	14.1	11.3	11.5	11.1	15.8	15.6	19.6
Ce	27.4	28.1	27.6	29.0	24.8	23.4	24.6	33.1	33.1	40.4
Pr	3.19	3.42	3.33	3.48	3.07	2.86	2.95	4.21	4.23	5.06
Nd	11.5	12.8	12.6	13.2	12.2	11.1	11.6	17.1	17.5	19.8
Sm	2.18	2.71	2.74	2.76	2.77	2.52	2.54	3.83	3.98	4.17
Eu	0.601	0.736	0.884	0.878	0.930	0.932	0.869	1.07	1.08	1.23
Gd	1.86	2.34	2.56	2.66	2.85	2.67	2.64	3.84	3.95	4.23
Tb	0.300	0.368	0.428	0.425	0.470	0.456	0.434	0.600	0.610	0.610
Dy	1.70	2.06	2.54	2.59	3.02	2.84	2.75	3.61	3.78	3.74
Ho	0.366	0.445	0.513	0.537	0.602	0.585	0.566	0.770	0.810	0.780
Er	1.07	1.25	1.49	1.55	1.62	1.64	1.55	2.14	2.23	2.15
Tm	0.167	0.183	0.207	0.228	0.268	0.262	0.248	0.330	0.340	0.330
Yb	1.12	1.24	1.53	1.59	1.66	1.69	1.64	2.08	2.12	2.02
Lu	0.175	0.199	0.222	0.232	0.283	0.264	0.265	0.300	0.320	0.310

\* 阿希金矿火山岩数据来自王强等(2006)和龙灵利等(2008)

表 2 东塔尔别克安山岩锆石 LA-ICP-MS 分析结果

Table 2 Zircon LA-ICP-MS analyzing results for the east Taerbieke andesite

分析点号	元素含量 ( $\times 10^{-6}$ )				元素比值				同位素比值				表观年龄 (Ma)			
	Th	U	U/Th	$^{207}\text{Pb}/^{206}\text{Pb}$	$^{207}\text{Pb}/^{235}\text{U}$	$^{206}\text{Pb}/^{238}\text{U}$	$^{207}\text{Pb}/^{206}\text{Pb}$	$^{206}\text{Pb}/^{238}\text{U}$	$^{207}\text{Pb}/^{235}\text{U}$	$^{206}\text{Pb}/^{238}\text{U}$	$^{207}\text{Pb}/^{235}\text{U}$	$^{206}\text{Pb}/^{238}\text{U}$	$^{207}\text{Pb}/^{235}\text{U}$	$^{206}\text{Pb}/^{238}\text{U}$	$^{207}\text{Pb}/^{235}\text{U}$	$^{206}\text{Pb}/^{238}\text{U}$
					1 $\sigma$	1 $\sigma$	1 $\sigma$	1 $\sigma$	1 $\sigma$	1 $\sigma$	1 $\sigma$	1 $\sigma$	1 $\sigma$	1 $\sigma$	1 $\sigma$	1 $\sigma$
06XJ31-1	139	242	1.74	0.05366	0.00089	0.4138	0.00691	0.05593	0.00063	357	37	352	5	351	4	
06XJ31-2	83	136	1.64	0.05639	0.00124	0.43478	0.00949	0.05592	0.00066	467	48	367	7	351	4	
06XJ31-3	84	130	1.55	0.0545	0.00108	0.41928	0.00832	0.0558	0.00064	392	44	356	6	350	4	
06XJ31-4	87	128	1.47	0.05466	0.0012	0.42163	0.00922	0.05595	0.00066	398	48	357	7	351	4	
06XJ31-5	139	255	1.83	0.05299	0.0009	0.40842	0.007	0.05591	0.00063	328	38	348	5	351	4	
06XJ31-6	53	175	3.30	0.07392	0.00104	1.72725	0.02481	0.16948	0.00188	1039	28	1019	9	1009	10	
06XJ31-7	147	201	1.37	0.05493	0.00095	0.41996	0.00731	0.05545	0.00062	409	38	356	5	348	4	
06XJ31-8	111	169	1.52	0.05478	0.00098	0.42285	0.00764	0.05599	0.00063	403	39	358	5	351	4	
06XJ31-9	26	41	1.55	0.05581	0.00252	0.55185	0.02446	0.07172	0.00108	445	97	446	16	447	6	
06XJ31-10	107	156	1.46	0.05135	0.00095	0.39074	0.00727	0.05519	0.00063	256	42	335	5	346	4	
06XJ31-11	145	218	1.51	0.05343	0.00092	0.40663	0.00707	0.0552	0.00062	347	39	346	5	346	4	
06XJ31-12	74	147	2.00	0.05303	0.00112	0.40504	0.00851	0.05539	0.00064	330	47	345	6	348	4	
06XJ31-13	116	201	1.73	0.05358	0.00107	0.40066	0.00799	0.05424	0.00062	353	45	342	6	341	4	
06XJ31-14	56	90	1.61	0.05303	0.00137	0.40129	0.01027	0.05488	0.00066	330	58	343	7	344	4	
06XJ31-15	66	117	1.77	0.05164	0.00129	0.38856	0.00961	0.05457	0.00065	270	56	333	7	343	4	
06XJ31-16	115	179	1.55	0.05407	0.00122	0.40569	0.00913	0.05442	0.00064	374	50	346	7	342	4	
06XJ31-17	68	116	1.71	0.05398	0.00145	0.41253	0.01098	0.05543	0.00068	370	59	351	8	348	4	
06XJ31-18	59	126	2.15	0.05274	0.00131	0.39694	0.00974	0.05459	0.00065	318	55	339	7	343	4	
06XJ31-19	84	153	1.81	0.0566	0.00123	0.43331	0.00939	0.05552	0.00065	475	48	366	7	348	4	
06XJ31-20	31	82	2.61	0.05839	0.00245	0.43571	0.0179	0.05412	0.00079	545	89	367	13	340	5	
06XJ31-21	162	304	1.87	0.05374	0.00098	0.41202	0.00756	0.05561	0.00063	360	40	350	5	349	4	
06XJ31-22	144	286	1.98	0.05411	0.00098	0.41449	0.0075	0.05556	0.00063	375	40	352	5	349	4	
06XJ31-23	52	105	2.03	0.05161	0.0012	0.39316	0.00908	0.05525	0.00065	268	52	337	7	347	4	
06XJ31-24	71	150	2.11	0.05277	0.00108	0.40516	0.00825	0.05568	0.00064	319	46	345	6	349	4	
06XJ31-25	204	320	1.57	0.05363	0.00097	0.40969	0.00744	0.05541	0.00063	355	41	349	5	348	4	

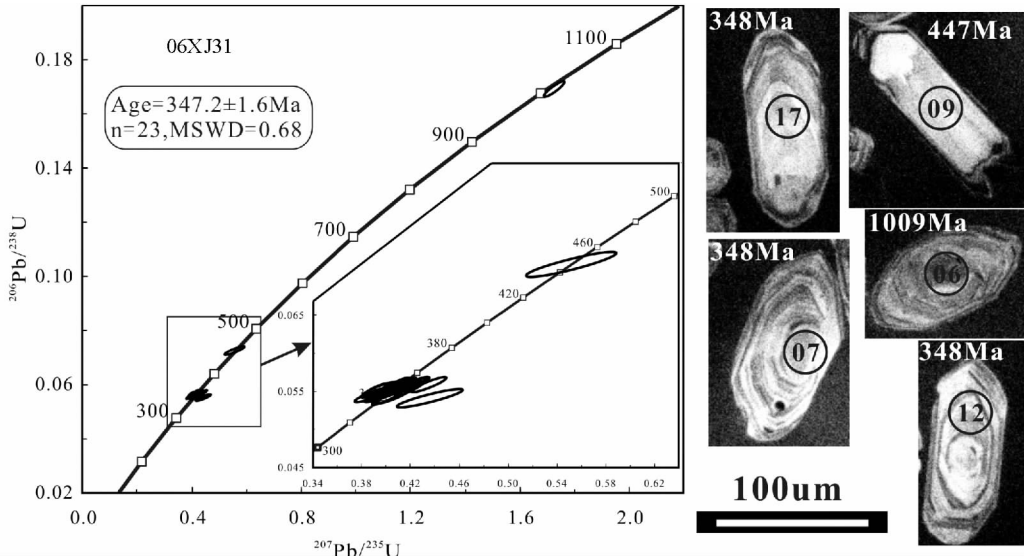


图3 06XJ31 LA-ICP-MS 锆石 U-Pb 谐和图和锆石 CL 图像

Fig. 3 LA-ICP-MS zircon U-Pb concordia diagrams and zircon cathodoluminescence (CL) images

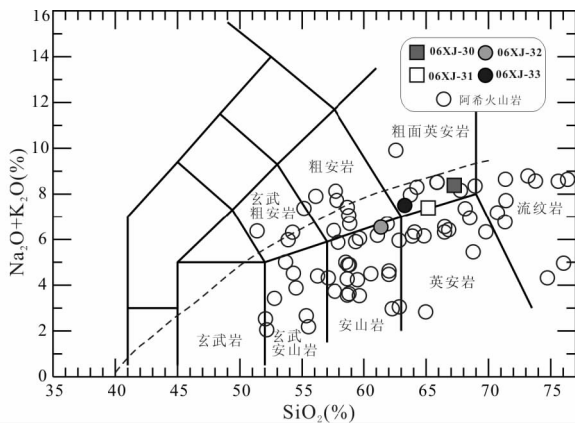


图4 TAS 图解(据 Le Maitre, 2002)

阿希火山岩数据来自左学义等, 2005; 沙德铭等, 2005; 王强等, 2006; 龙灵利等, 2008

Fig. 4 TAS diagram (after Le Maitre, 2002)

#### 4.2 元素地球化学特征

我们对东塔尔别克四件安山岩进行了全岩地球化学分析。其中样品 06XJ-30 和 06XJ-31 两样品取自靠近矿化带处, 岩石有蚀变, 而 06XJ-32 和 06XJ-33 两样品取自远离矿化带的地方, 较新鲜。安山岩的  $\text{SiO}_2$  含量在 58.94% ~ 63.85%, 变化范围较小; 具有较高的  $\text{MgO}$  含量 (2.51% ~ 6.59%) 和高的  $\text{Mg}^\#$  (57 ~ 69); 全碱的含量为 6.27% ~ 7.96%;  $\text{Al}_2\text{O}_3$  含量较高 (15.39% ~ 16.65%)。安山岩在 TAS 图解 (图 4) 中落入亚碱性范围内, 在  $\text{SiO}_2$ - $\text{FeO}^\text{T}/\text{MgO}$  图中 (图 5a) 均落入钙碱性区域。

安山岩稀土元素分布模式为轻稀土富集型, 轻重稀土分

馏明显 (图 6a); 无明显 Eu 异常 ( $\delta\text{Eu}$  为 0.89 ~ 1.02), 但重稀土元素分异不明显, ( $\text{Gd}/\text{Yb}$ )<sub>N</sub> = 1.37 ~ 1.56; 富集大离子亲石元素 (LILE), 如 Sr 具有明显的正异常, 亏损高场强元素 (HFSE) 和重稀土元素, Nb、Ti 有明显的负异常 (图 6b)。以上这些特点显示它们与俯冲带岩浆地球化学特征类似 (Wilson, 1989)。

## 5 讨论

### 5.1 形成时代

前人曾对西天山与铜金相关的岩浆岩的年代学进行了深入研究。但目前对该区出露的岩浆岩的形成时代仍有争议: 李华芹 (1998) 用 Rb-Sr 等时线法测得阿希金矿赋矿围岩大哈拉军山组火山岩的年龄为 346 ~ 321 Ma, 为早石炭世; Zhu *et al.* (2005) 给出分布于西南天山大哈拉军山组火山岩的 SHRIMP 锆石年龄分别为  $353.7 \pm 4.5$  Ma 和  $312.8 \pm 4.2$  Ma 的资料, 分别属于早石炭世和晚石炭世; 翟伟等 (2006) 报道了阿希金矿床赋矿围岩大哈拉军山组火山岩 SHRIMP 锆石年龄为  $363.2 \pm 5.7$  Ma, 但没有对岩石中出现的年轻锆石 (342 ~ 327 Ma) 给予解释。本次对东塔尔别克矿区的花岗闪长斑岩的定年结果显示其结晶年龄为  $347.2 \pm 1.6$  Ma, 这与阿希金矿赋矿安山岩的 SHRIMP 锆石 U-Pb 年龄 (356 ~ 342 Ma, 王强未发表数据) 相一致, 也与该区地层古生物的证据 (王福同, 2006) 一致。因此, 我们认为东塔尔别克岩浆岩形成于早石炭世。

### 5.2 岩石类型与成因

东塔尔别克的安山岩与阿希金矿区的火山岩相似 (图 4 ~ 8), 其具有高的  $\text{MgO}$  含量和  $\text{Mg}^\#$ 、高的 Cr ( $60.1 \times 10^{-6}$  ~

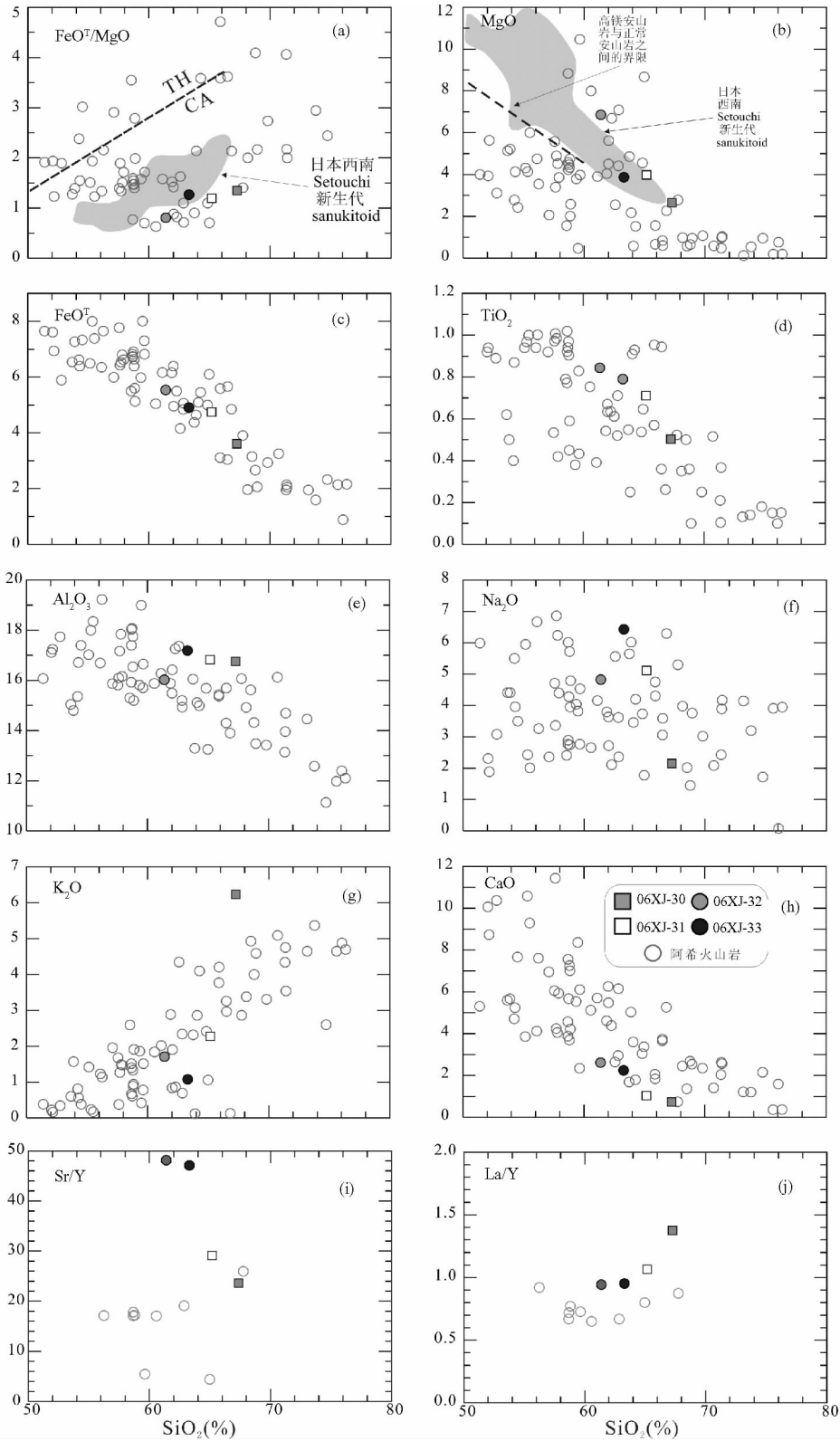


图5  $\text{SiO}_2$ 对  $\text{FeO}^T/\text{MgO}$  (a)、 $\text{MgO}$  (b)、 $\text{FeO}$  (c)、 $\text{TiO}_2$  (d)、 $\text{Al}_2\text{O}_3$  (e)、 $\text{Na}_2\text{O}$  (f)、 $\text{K}_2\text{O}$  (g)、 $\text{CaO}$  (h)、 $\text{Sr}/\text{Y}$  (i) 和  $\text{La}/\text{Y}$  (j) 图 (a) 中 TH-拉斑系列;CA-钙碱系列;(b) 高镁安山岩与正常安山岩分界线来源于 McCarron and Smellie (1998); 日本西南 Setouchi 新生代 Sanukitoid 来源于 Tatsumi (2006); 阿希火山岩数据来自左学义等(2005)、沙德铭等(2005)、王强等(2006)和龙灵利等(2008)

Fig. 5  $\text{SiO}_2$  versus  $\text{FeO}^T/\text{MgO}$  (a),  $\text{MgO}$  (b),  $\text{FeO}$  (c),  $\text{TiO}_2$  (d),  $\text{Al}_2\text{O}_3$  (e),  $\text{Na}_2\text{O}$  (f),  $\text{K}_2\text{O}$  (g),  $\text{CaO}$  (h),  $\text{Sr}/\text{Y}$  (i) and  $\text{La}/\text{Y}$  (j) diagrams

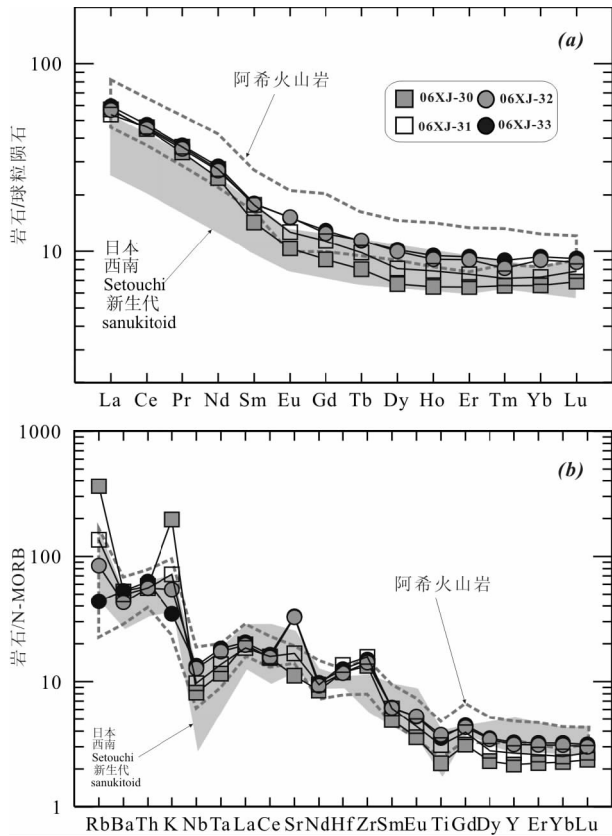


图6 稀土元素球粒陨石标准化配图(a)和微量元素NMORB标准化蛛网图(b) (标准值据 Sun and McDonough, 1989)

阿希火山岩数据来自王强等(2006)和龙灵利等(2008)

Fig. 6 The chondrite-normalized rare earth element (REE) patterns (a) and primitive mantle-normalized multi-element plots (b) (normalized values from Sun and McDonough, 1989)

$241 \times 10^{-6}$ ) 和 Ni ( $35.9 \times 10^{-6} \sim 126 \times 10^{-6}$ ) 含量, 以及低的  $\text{FeO}^T/\text{MgO}$ , 类似于高镁安山岩 (Polat and Kerrich, 2001; Tatsumi, 2001)。但同时与阿希金矿的火山岩相比, 东塔尔别克安山岩具有埃达克岩的特征 (Defant and Drummond, 1990): 无明显 Eu 异常, 其中两新鲜样品具有高的 Sr 含量 ( $685 \times 10^{-6} \sim 696 \times 10^{-6}$ ) 和低 Y 含量 ( $14.3 \times 10^{-6} \sim 14.9 \times 10^{-6}$ ), 以及高的 Sr/Y 比值 (47.1 ~ 48.1), 在 Sr-Y 判别图 (图 7a) (Defant and Drummond, 1993) 中均落入埃达克区。而两蚀变的样品具有相对低的 Sr 含量 ( $235 \times 10^{-6} \sim 354 \times 10^{-6}$ ) 和高的  $\text{K}_2\text{O}$  (2.17% ~ 5.92%) 含量 (表 1), 很可能与蚀变有关。所以东塔尔别克的安山岩为高镁安山岩-埃达克岩组合。

目前, 对埃达克质岩浆的形成提出了不同的机制, 主要包括: (1) 俯冲洋壳熔融 (Defant and Drummond, 1990; Zhang *et al.*, 2004, 2006b; Han *et al.*, 2006); (2) 增厚下地壳熔融 (Atherton and Petford, 1993; Wang *et al.*, 2005; Zhang *et*

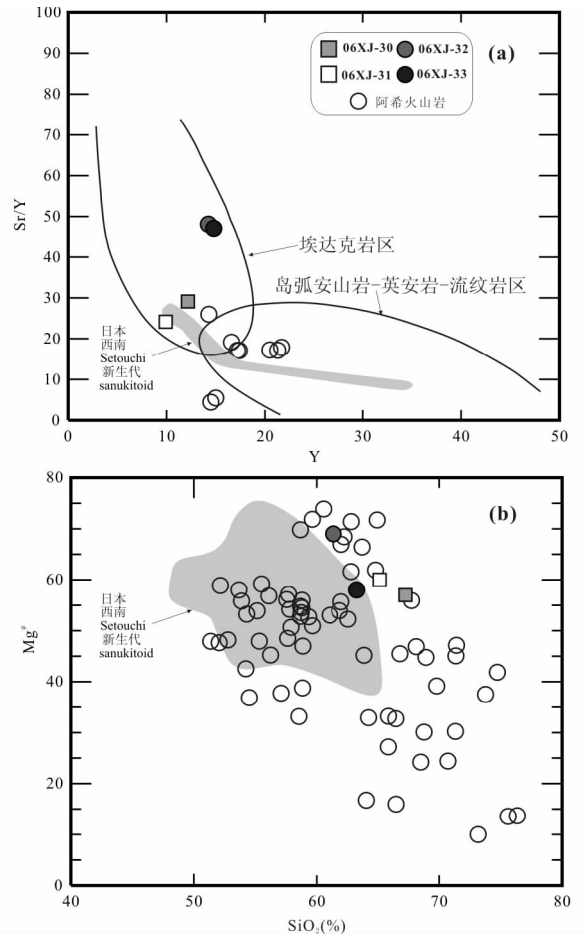


图7 Sr/Y-Y (a, Defant and Drummond, 1993) 和  $\text{SiO}_2$ - $\text{Mg}^\#$  (b)

图解数据来源同图 4

Fig. 7 Sr/Y-Y (a, after Defant and Drummond, 1993) and  $\text{SiO}_2$ - $\text{Mg}^\#$  (b) diagrams

*al.*, 2006a; Lai *et al.*, 2007; Xiao and Clemens, 2007; Zhao *et al.*, 2008; 张旗等, 2001); (3) 拆沉下地壳熔融 (Kay and Kay, 1993; Xu *et al.*, 2002; Gao *et al.*, 2004; Wang *et al.*, 2006a, 2007a; Hou *et al.*, 2007; Huang *et al.*, 2008; Liu S *et al.*, 2008); (4) 玄武质岩浆的结晶分异 (Castillo *et al.*, 1999; Macpherson *et al.*, 2006)。在玄武质岩浆的结晶分异过程中, 石榴石 ± 辉石的高压分离结晶作用导致岩浆中的  $\text{Al}_2\text{O}_3$  随着  $\text{SiO}_2$  的增加而降低, 而 Sr/Y, La/Y 比值则随  $\text{SiO}_2$  的增加而升高, 但东塔尔别克岩浆岩没有这样的规律性变化 (图 5i, j), 所以其不可能由玄武质岩浆的结晶分异所形成。同时东塔尔别克岩浆岩的具有高的  $\text{MgO}$  和  $\text{Mg}^\#$  (57 ~ 69), 所以排除 (2) 的可能性。由于拆沉下地壳熔融往往发生在板内伸展环境 (Xu *et al.*, 2002; Gao *et al.*, 2004; Wang *et al.*, 2006a), 这与西天山北部早石炭世的构造背景不符。最近的研究表明, 在天山北部地区存在石炭纪洋壳的俯冲和熔融作用, 并且形成了典型的岛弧岩浆岩组合: 埃达克岩-高镁安山



岩-富 Nb 玄武岩组合 (Wang *et al.*, 2007b; 王强等, 2006)。因此,我们认为东塔尔别克高镁埃达克岩岩浆的形成更可能与洋壳的俯冲有关,可能是由俯冲洋壳熔融所形成,同时其高的 MgO 含量和  $Mg^\#$  暗示可能受到地幔橄榄岩的混染。

早期含水地幔橄榄岩熔融实验表明,在大多数情况下地幔橄榄岩的熔融一般都是形成玄武质岩石,尤其在高温和高压的情况下更是如此。但含水地幔橄榄岩在低温低压情况下的熔融(约 1000°C, 1GPa)是产生高镁安山质岩浆的一种可能机制(如 Kushiro, 1969; Tatsumi and Ishizaka, 1981, 1982)。这些实验结果使 Crawford *et al.* (1989) 和 Tatsumi and Maruyama (1989) 认为:高镁安山岩可能是由俯冲的洋壳释放的含水流体加入到地幔楔中而引发地幔橄榄岩熔融形成的。Hirose(1997)通过直接熔融含水地幔橄榄岩实验进一步表明,高镁安山岩只能在低于 1300°C、压力为 1GPa 且含水条件下形成。但最近的一系列的研究表明有些高镁安山岩更有可能是由俯冲洋壳(+沉积物)熔融的硅质熔体和上覆的地幔橄榄岩相互反应所形成,尤其是日本西南 Setouchi 火山岩带中的 Sanukitoid (赞岐岩类) (Shimoda *et al.*, 1998; Tatsumi, 2001; Hanyu *et al.*, 2002, 2006; Tatsumi and Hanyu, 2003)。

赞岐岩最早是指在日本四国东北 Sanuki 地区的灰-黑色、有光泽、隐晶质致密的含有针状古铜辉石斑晶的火山岩 (Tatsumi, 2006)。Koto (1916) 进一步将日本西南新生代 Setouchi 火山岩带中与赞岐岩相似且为黑色、斑晶 < 10% 的致密火山岩称为赞岐岩类 (Sanukitoid)。Tatsumi and Ishizaka (1981, 1982) 则将“赞岐岩类”的概念用于 Setouchi 火山岩带中那些斑晶少于 10% (没有或很少有斜长石斑晶) 和无斑隐晶的高镁玄武岩-安山岩,岩石类型包括普通辉石橄榄玄武岩、普通辉石橄榄安山岩、古铜辉石橄榄安山岩和古铜辉石安山岩。后来,Shirey and Hanson (1984)、Stern and Hanson (1991)、Smithies and Champion (2000) 和 Martin *et al.* (2005) 将太古宙具上述赞岐岩特征 (Si 过饱和、 $Mg^\#$  和 Ni、Cr、LILE 含量高) 的深成岩和火山岩 (高镁二长闪长岩、花岗闪长岩和粗安岩) 称为“赞岐岩类”。东塔尔别克的含普通辉石安山岩的地球化学特征有些类似于赞岐岩,在 Th/La-Th 图解中 (图 8) 东塔尔别克安山岩落入 Setouchi 的赞岐岩区域,由于其高的 Th/La 比和 Th 含量,其源区可能包含了更多俯冲洋壳上覆沉积物熔体成分,即由洋壳和沉积物的熔体与地幔橄榄岩相互作用的结果。

### 5.3 动力学与成矿意义

#### 5.3.1 动力学背景

目前,西天山北缘石炭纪火山岩的形成模式主要包括有以下三种:(1) 裂谷 (Xia *et al.*, 2004b; 车自成和刘良, 1996); (2) 地幔柱 (夏林圻等, 2004); (3) 俯冲岛弧环境 (Gao *et al.*, 1998; Zhang *et al.*, 2003; Liu and Fei, 2006; Wang *et al.*, 2007b; Xiao *et al.*, 2008; Qian *et al.*, 2009; 王强等,

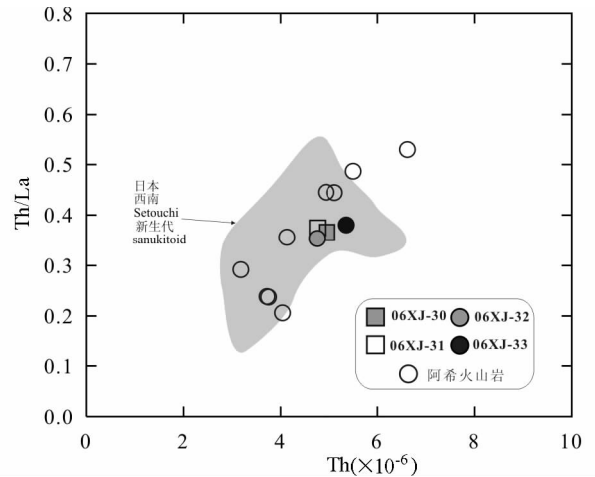


图 8 Th/La-Th 图解

阿希火山岩数据来自王强等(2006)和龙灵利等(2008)

Fig. 8 Th/La-Th diagram

2006; 李锦轶等, 2006; 龙灵利等, 2008)。最近,王强等(2006)在天山北部地区厘定出典型的岛弧岩浆岩组合:埃达克岩-高镁安山岩-富 Nb 玄武岩组合,并认为可能是与石炭纪洋壳的俯冲和熔融有关。龙灵利等(2008)等认为伊犁板块南北缘石炭纪火山岩类似于俯冲带之上大陆边缘火山岛弧岩特征,可能是大陆边缘环境下的产物。随着详细的岩石学和地球化学研究的深入,越来越多的数据显示该区的火山岩以钙碱性为主,碱性岩很少。另外,西天山的中基性熔岩的时代为 354 ~ 313Ma,时限大于 40Ma,该区火山岩并具有典型大陆弧岩浆的地球化学特征 (Zhu *et al.*, 2005)。本文资料表明,东塔尔别克的安山岩也类似于俯冲带岩浆地球化学特征 (Wilson, 1989)。所以,我们趋向认为西天山北缘在早石炭世为岛弧环境。

#### 5.3.2 成矿意义

高镁安山岩-埃达克岩型岩浆岩对于该区的 Au 的成矿作用具有重要意义:(1) Au 是亲硫元素,主要赋存于地幔和铁镁质岩石,所以玄武质洋壳熔融形成的埃达克岩具有良好的成矿物质来源;(2) 板片熔体具有高的氧逸度 (Mungall, 2002), Mungall (2002) 认为板片熔体可以携带大量的  $Fe_2O_3$ , 当携带大量  $Fe_2O_3$  的板片熔体进入到富金属硫化物的地幔楔时,将会导致地幔楔橄榄岩氧逸度 ( $f_{O_2}$ ) 的增高,地幔中的金属硫化物将被氧化, Au 等将顺利进入到板片熔体或岛弧岩浆中;安山岩高的 MgO 含量和  $Mg^\#$  就是板片熔体与地幔橄榄岩相互作用的证据;(3) 俯冲板片富含挥发份  $H_2O$  和 Cl, 挥发份 Cl 在高温高压下是 Au 等金属元素的强烈配合剂,它们将与 Au 形成稳定的配合物随岩浆一起迁移,有利于成矿 (熊小林等, 2005)。东塔尔别克金矿北据阿希金矿仅 2km, 与其共生的岩浆岩与阿希金矿火山岩相似,同时其形成时代也一致。二者的成矿机制可能相同,甚至东塔尔别克金矿很可能是阿希金矿向南延伸的部分。

## 6 结论

(1) 东塔尔别克安山岩的形成时代为  $347.2 \pm 1.6$  Ma, 形成于早石炭世岛弧环境。

(2) 东塔尔别克岩浆岩为高镁安山岩-埃达克岩组合, 其形成可能是与洋壳的俯冲熔融作用所形成, 同时可能包含了更多俯冲洋壳上覆沉积物熔体成分, 即由洋壳和沉积物的熔体与地幔橄辉岩相互作用的结果。

(3) 俯冲带熔体与地幔的作用对于区内金矿的形成可能具有重要的意义。

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