

钟福平, 钟建华, 艾合买提江·阿不都热合曼, 等. 华北克拉通破坏时间与破坏范围分布特征——来自银根—额济纳旗盆地苏红图坳陷早白垩世火山岩的启示[J]. 中国地质, 2015, 42(2): 435–456.

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华北克拉通破坏时间与破坏范围分布特征 ——来自银根—额济纳旗盆地苏红图坳陷早白垩世 火山岩的启示

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提要:克拉通破坏的时间和范围是华北克拉通破坏研究的重要基础问题,但是在华北克拉通破坏时间与破坏范围的问题上存在着不同观点。本文通过对位于华北克拉通西北部银根—额济纳旗盆地苏红图坳陷内采集的火山岩进行年代学及地球化学研究,认为苏红图火山岩年龄为 105~113 Ma,为一套钾质碱性系列玄武岩,其形成机制是由于岩石圈发生减薄,软流圈地幔岩浆上涌,经分离结晶而形成的,而动力学机制主要是由于西伯利亚板块、内蒙古褶皱带和华北板块在晚侏罗世发生的碰撞拼合。此外,本文还在前人对华北克拉通破坏研究基础上,依据作者对苏红图坳陷火山岩做的一些工作,粗浅地探讨了华北克拉通破坏的时间与范围的问题,认为华北克拉通周缘均为构造薄弱带,北缘为兴—蒙造山带,南侧为大别—秦岭造山带,东侧为苏鲁带和太平洋俯冲带,河套裂陷、汾渭裂陷分别与古元古代高温变质孔兹岩带及约 18.5 亿年前华北克拉通东、西部块体拼合时形成的中部造山带内位置大致重合,而苏红图坳陷位于中亚造山带南缘,同时也处于两板块拼合交汇处。这些构造薄弱带处在不同时期发生的俯冲与碰撞的结合部位,它们可能是岩石圈减薄的起始位置,并且它们的俯冲与碰撞时间是华北克拉通破坏的起始时间。克拉通破坏范围主要发生在太行山以东地区,太行山以西的河套裂陷、汾渭裂陷发生了减薄,而苏红图坳陷在早白垩世也发生减薄,所以,破坏范围分布在地理上呈不连续分布特征,造成这种分布特征的主要原因是由于不同区域的破坏时间与破坏的动力学机制不同。

关 键 词:华北克拉通;银根—额济纳旗盆地苏红图坳陷;破坏范围;动力学机制

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Timing and scale of the destruction of the North China craton: Revelation from the Early Cretaceous volcanic rocks in Suhongtu Depression of Inggen–Ejin Banner Basin

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Abstract: Suhongtu Depression in Inggen–Ejin Banner basin is located on the northwestern margin of the North China Craton. The periphery of the North China Craton is a weak tectonic belt, with Hing – Mongolian orogenic zone on the northern margin, Dabie – Qinling orogenic belt on the southern margin, Tan–Lu fracture zone and Pacific subduction zone on the eastern margin. Hetao rift and Fenwei rift are roughly coincident with positions of Paleoproterozoic high-temperature metamorphic rocks and the central orogenic belts, which were formed during the breakup period of the North China Craton and western blocks approximately 1.85 billion years ago. However, Suhongtu Depression is located on the southern edge of the Central Asia Orogenic Belt, and also lies at the junction of two plates. These structural weakness belts experienced subduction in different periods, and might have served as the starting locations of lithospheric thinning, and the time of their subduction and collision was the beginning of the destruction of the North China Craton. Craton destruction regions occurred mainly to the east of the Taihang Mountains, whereas thinning happened in Fenwei rift, Hetao rift and Suhongtu Depression west of the Taihang Mountains. Therefore, damages to the Suhongtu Depression lithosphere occurred on the northwest margin of the North China Craton in the late Early Cretaceous (~ 110Ma), and hence the damaged regions in the Craton were distributed discontinuously in geography, which resulted from different destruction periods and dynamic mechanisms of the lithosphere in different regions.

Key words: Suhongtu depression of Inggen–Ejin Banner Basin; North China Craton; destruction time; destruction range; dynamic mechanism

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银根—额济纳旗盆地(以下简称银—额盆地)苏红图坳陷位于华北板块西北部,同时又处于中亚造山带南缘,塔里木地块与华北板块的交汇处(图1),构造背景复杂。盆地内火山岩发育3期,其中早白垩世晚期火山岩活动最为强烈,波及面较广,由中基性火山熔岩组成,产于下白垩统苏红图组中。银—额盆地是中国具有一定勘探潜力的含油气盆地^[1],而火山岩对油气的生成、运移和储存具有重要影响^[2–3]。已有学者对该区早白垩世火山岩做过研究,并进行过一些有益的探索^[4–7]。

华北克拉通以发育古老岩石和中生代岩石圈巨量减薄闻名于世,是世界上少有的遭受破坏及岩石圈巨量减薄的克拉通。20世纪初翁文灏先生^[8]提出了燕山运动概念,随后陈国达先生^[9]提出“地台活化”观点,在此基础上,20世纪90年代中外科学家提

出的“岩石圈减薄”^[10–11]或“去根”^[12]的概念。随着研究的不断深入,人们逐步意识到克拉通东部不仅发生了巨厚岩石圈地幔的丢失,而且岩石圈地幔物质成分、热状态和流变学性质等方面发生了根本性的改变^[13]。破坏和减薄伴随着一系列强烈的构造作用、岩浆作用、成矿大爆发及大型含油气盆地的形成,在全球极为少见,因此,华北克拉通成了近年来国内外研究的最热门课题之一。

克拉通破坏的时间和范围是华北克拉通破坏研究的重要基础问题,它的确定是探索克拉通破坏机制及其动力学控制因素的关键^[14]。目前,在华北克拉通破坏发生的时间上存在着在石炭纪^[15,16]、三叠纪^[17,18]、侏罗纪^[19–21]、白垩纪^[22]和新生代^[23]等不同的认识,而其破坏范围主要发生在华北克拉通东部,即太行山断裂带以东地区,但在太行山以西的汾渭

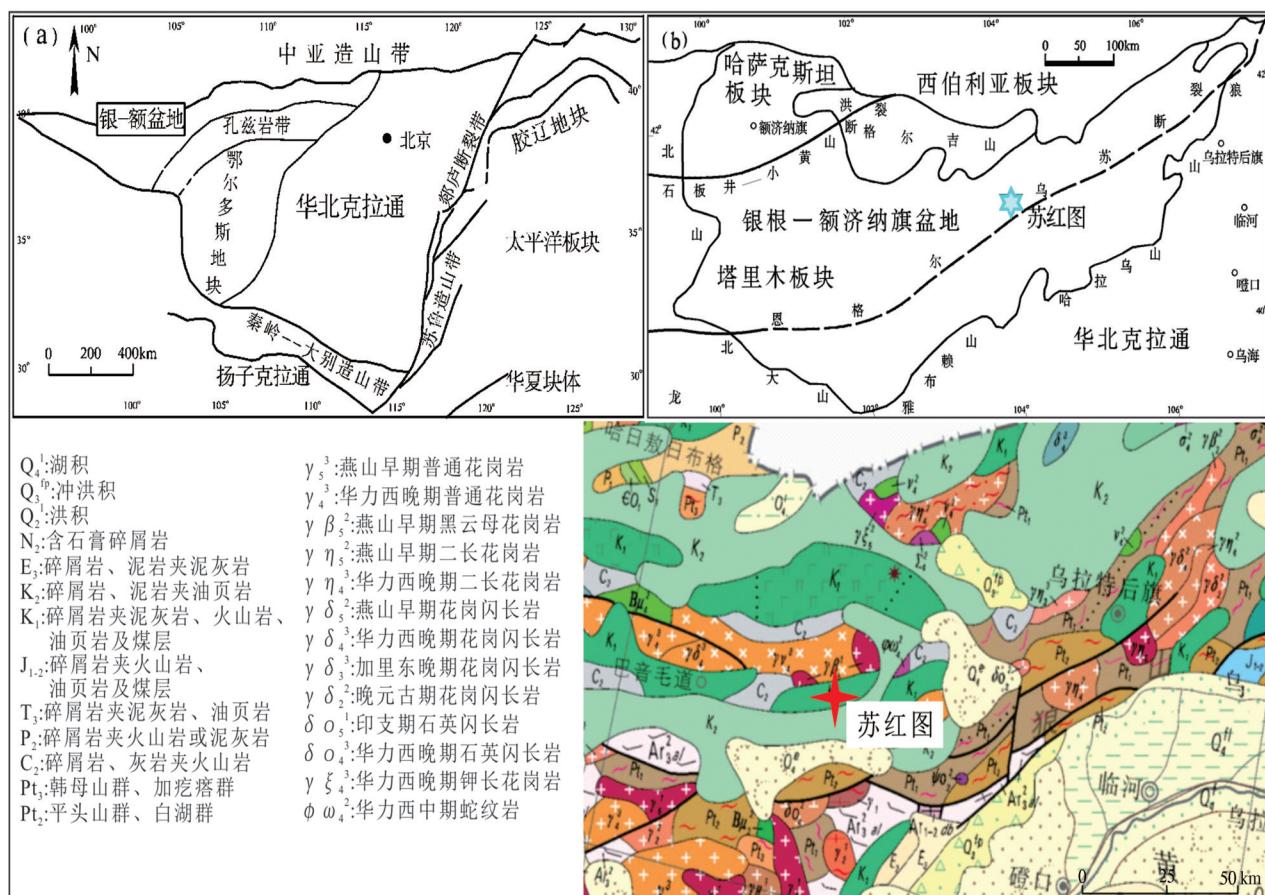


图1 样品采样位置图
Fig.1 Geographic map showing sampling sites

地堑、河套地堑下伏岩石圈也发生了减薄^[24]。

通过对华北地区岩浆岩的研究,可以探讨华北克拉通破坏时间和范围。本文通过对银—额盆地苏红图坳陷早白垩世火山岩研究,探讨火山岩的地球化学特征及喷发动力学机制,依据苏红图坳陷大地构造位置特殊性,通过对华北克拉通发生破坏的时间和范围的探讨,结合对研究区火山岩年代学及动力学研究,探讨华北克拉通破坏范围分布特征,对于华北克拉通破坏研究具有重要意义。

1 苏红图火山岩年代学及地球化学特征

1.1 火山岩年代学

笔者在银—额盆地苏红图坳陷进行野外踏勘(图1~3)时采集大量火山岩样品,选送其中的6件样品(编号分别为Sht08、Sht11、Sht16、Sht18、Sht23、Sht28)进行⁴⁰Ar/³⁹Ar定年,得到年龄值为105~113

Ma(表1,图4)。

⁴⁰Ar/³⁹Ar同位素定年法是利用原子反应堆的快中子,先将矿物或岩石中的稳定态同位素³⁹K活化为³⁹Ar。经中子活化后的样品,利用质谱分析方法,一次完成⁴⁰Ar、³⁹Ar、³⁷Ar及³⁶Ar等氩同位素的分析测量工作,经由⁴⁰Ar/³⁹Ar比值推算出⁴⁰Ar*/⁴⁰K(⁴⁰Ar*为放射性产生的氩的同位素)比值。然而在中子照射过程中,³⁹Ar与³⁹K含量的关系常数并非为定值,会随中子通量大小而改变,因此在实际操作过程中⁴⁰Ar/³⁹Ar定年法系利用已知的K-Ar同位素年龄的矿物,即标准矿物与欲测年龄的样品一起放入原子反应堆中接受中子活化,求得一些与快中子通量有关的常数,进而计算得到样品同位素年代。

1.2 火山岩地球化学特征

1.2.1 火山岩地化数据测试方法

将采集的样品选送部分至中国科学院地质与地球物理研究所进行测试,数据见表2,测试项目为

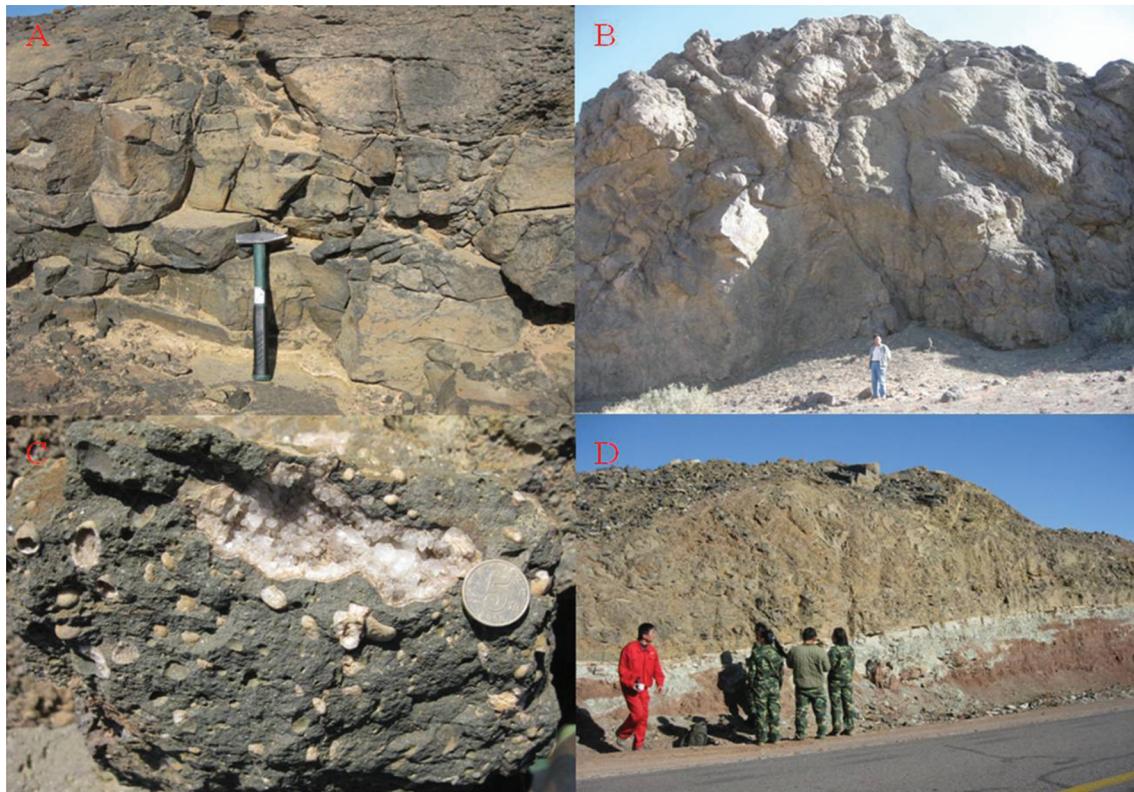


图2 银—额盆地苏红图坳陷火山岩野外照片

A—块状玄武岩(41°12'067N, "104°12'929"E); B—块状玄武岩(41°11'406 N, "104°13'333"E); C—杏仁状玄武安山岩(41°12'014 "N, 104°12'741"E); D—块状橄榄玄武岩(41°13'213 "N, 104°11'006"E);括号“()”内经纬度表示火山岩的地理位置

Fig.2 Field photographs of volcanic rocks

A—Massive basalt(41°12'067N, "104°12'929"E); B—Massive basalt(41°11'406 N, "104°13'333"E); C—Amygdaloidal basaltic andesite (41°12'014 "N, 104°12'741"E); D—Massive olivine basalt(41°13'213 "N, 104°11'006"E); The latitude and longitude in brackets denote the geographic location of volcanic rocks

常量元素、微量元素、Sr—Nd—Pb同位素,其中常量、微量元素测试各24件,同位素6件。

常量元素分析方法:用0.6 g样品和6 g四硼酸锂制成的玻璃片在Shimadzu XRF-1500上测定氧化物的质量分数,精度优于2%~3%。

微量元素采用酸溶法:样品溶液制备好后,在Element II型ICP-MS上测试,按照GSR-1和GSR-2国家标准,元素质量分数大于 10×10^{-6} 的元素精度优于5%,小于 10×10^{-6} 的元素优于10%。

Sr、Nd同位素分析方法:称取约100 mg全岩粉末样品,加入适量的 $^{87}\text{Rb}-^{84}\text{Sr}$ 和 $^{149}\text{Sm}-^{150}\text{Nd}$ 混合稀释剂和纯化的HF-HClO₄混合试剂后,在高温下完全溶解。Rb-Sr分离和纯化在装有5 mL AG50W-X12交换树脂(200~400目)的石英交换柱进行,Sm和Nd的分离和纯化是在石英交换柱用1.7 mL Teflon粉末为交换介质完成。采用 $^{146}\text{Nd}/^{144}\text{Nd}=$

0.7219 和 $^{86}\text{Sr}/^{88}\text{Sr}=0.1194$ 分别校正测得 Nd 和 Sr 同位素比值。Nd 标样 Jndi-1 测定结果为 $^{143}\text{Nd}-^{144}\text{Nd}=0.512107$, Sr 标样 NBS987 测定结果为 $^{87}\text{Sr}-^{86}\text{Sr}=0.710276$ 。Rb-Sr 和 Sm-Nd 的全流程本底分别为 100 pg 和 50 pg 左右。

Pb 同位素的分析方法:采用 HF 溶解样品,溶样后蒸干样品溶液,用 6 mol/L HCl 将氟化物样品转为氯化物,蒸干后,用 0.6 mol/L HBr 提取样品。在装有 80 μL AGLx(100~200 目)交换树脂的 Teflon 交换柱上,采用 0.6 mol/L HBr 和 6 mol/L HCl 流程分离纯化 Pb 样品。同位素比值测量采用德国 Finnigan 公司 MAT262 固体源质谱仪, $^{147}\text{Sm}/^{144}\text{Nd}$ 和 $^{87}\text{Rb}/^{86}\text{Sr}$ 比值误差小于 0.5%。

1.2.2 火山岩地球化学特征

苏红图组火山岩主、微量元素分析结果(表2)显示, SiO_2 含量为 47.58%~56.28%, 为中基性火山

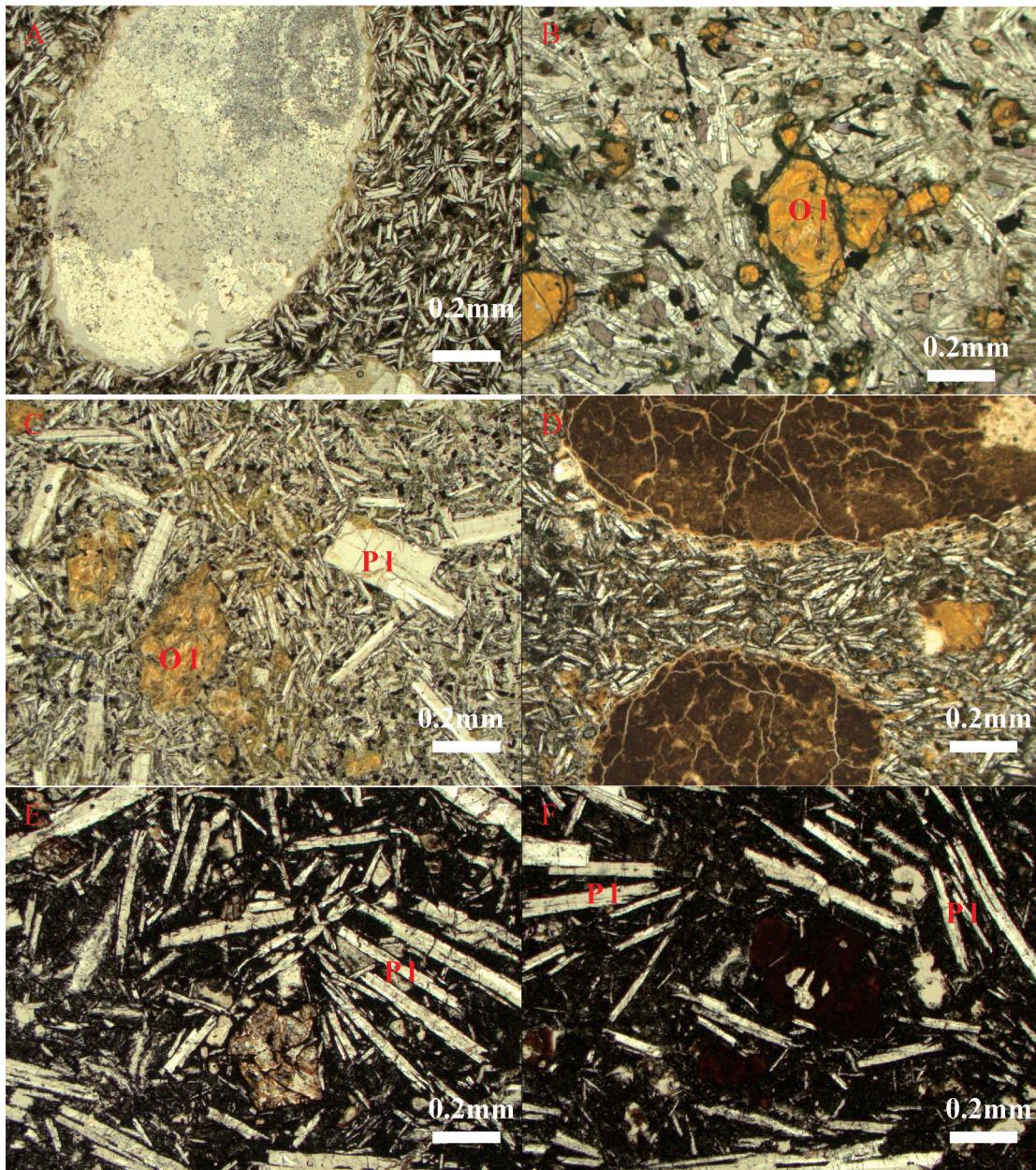


图3 银—额盆地苏红图坳陷火山岩镜下照片

A—杏仁状辉石安山岩;B—橄榄玄武岩;C—玄武岩;D—杏仁状辉石安山岩;E—杏仁状铁质玄武岩;
F—杏仁状橄榄铁质玄武岩;PI—斜长石;OI—橄榄石

Fig. 3 Microscope photographs of volcanic rocks

A—Amygdaloidal pyroxene andesite; B—Olivine basalt; C—Basalt; D—Amygdaloidal pyroxene andesite; E—Amygdaloidal iron basalt;
F—Amygdaloidal olivine iron basalt; PI—Plagioclase; OI—Olivine

岩, 平均含量49.63%;里特曼指数为 $4.7 < \sigma < 7.9$, 平均值为6.3, 属碱性系列火山岩; $(K_2O + Na_2O)$ 为4.95%~7.51%, 平均含量为6.37%;在 $SiO_2 - (K_2O + Na_2O)$ 图解中24个样品均落在碱性玄武岩系列区域中(图5)。在 $SiO_2 - K_2O$ 图解中, 绝大多数样品落

高钾玄武岩区域中(图6)。从微量元素蛛网图解(图7)中可以看出Nb、Ta元素具有富集的特点。

通过计算, 苏红图火山岩的 ε_{Nd} 为-6.24~-3.08。 $^{87}Sr/^{86}Sr$ 和 $^{143}Nd/^{144}Nd$ 比值分别为0.705477~0.707098 和 0.512318~0.512480(表3)。 $^{143}Nd/^{144}Nd$

表1 火山岩(Sht8、Sht11、Sht16、Sht18、Sht23、Sht28)⁴⁰Ar/³⁹Ar阶段升温分析结果Table 1 Analytical results of ⁴⁰Ar/³⁹Ar stage warming

火山岩(Sht8)								
温度/℃	⁴⁰ Ar/ ³⁹ Ar	³⁷ Ar/ ³⁹ Ar	³⁶ Ar/ ³⁹ Ar	⁴⁰ Ar*/ ³⁹ Ar _k	⁴⁰ Ar*/%	³⁹ Ar _k /%	Age / Ma	± 2s /Ma
800	400.53596	1.89396	1.29445	18.204491	4.54	1.07	159.6	±83.3
850	600.56064	1.53985	1.91936	33.556077	5.58	3.56	284.1	±107.3
890	199.55504	1.25822	0.60574	20.681794	10.35	6.57	180.3	±36.2
930	42.46466	1.33096	0.10237	12.333764	29.01	13.84	109.7	± 6.6
960	33.16382	1.39697	0.06999	12.608255	37.97	11.62	112.0	± 4.5
990	33.04161	1.80906	0.07016	12.471141	37.69	6.09	110.8	± 4.6
1020	34.15268	1.44533	0.07368	12.509556	36.58	4.89	111.2	± 5.4
1050	31.62098	0.85513	0.06583	12.245327	38.70	4.01	108.9	± 5.0
1080	29.31229	0.70512	0.05814	12.193512	41.57	3.64	108.5	± 4.0
1110	24.09522	0.69613	0.03871	12.720275	52.76	5.07	113.0	± 2.8
1140	17.52755	1.26822	0.01754	12.458789	71.00	11.95	110.7	± 1.5
1170	19.37254	2.55239	0.02450	12.361393	63.67	7.50	109.9	± 2.1
1200	21.68269	4.25215	0.03216	12.563433	57.73	3.63	111.6	± 3.3
1260	21.02537	7.18903	0.03174	12.292286	58.11	4.41	109.3	± 2.9
1340	16.64307	6.99289	0.02219	10.704157	63.93	4.93	95.5	± 2.5
1450	17.09333	6.80533	0.01953	11.932095	69.40	7.23	106.2	± 1.8
火山岩(Sht11)								
800	701.20117	2.83541	2.37229	0.414464	0.06	0.34	3.6	±150.4
850	1347.6337	1.51256	4.54096	5.908866	0.44	1.83	50.7	±275.7
900	377.25698	1.54868	1.22754	14.659909	3.88	4.67	123.2	±70.7
950	94.59198	1.71983	0.26732	15.759120	16.64	13.26	132.2	±15.6
990	44.86243	1.25496	0.10511	13.915625	30.99	13.21	117.2	± 6.4
1030	33.75397	1.08670	0.06894	13.479338	39.90	10.35	113.6	± 4.4
1070	35.48988	0.81590	0.07659	12.932506	36.41	5.71	109.1	± 4.9
1110	27.21932	0.76648	0.04611	13.662610	50.16	6.75	115.1	± 3.3
1150	20.10580	1.47922	0.02323	13.376534	66.45	8.36	112.8	± 1.9
1180	21.19297	2.36783	0.02721	13.368360	62.95	5.71	112.7	± 2.7
1210	20.25695	2.63317	0.02429	13.319324	65.60	6.26	112.3	± 2.4
1260	19.66864	4.47057	0.02273	13.358679	67.66	10.48	112.6	± 1.9
1330	21.43974	9.43589	0.03002	13.427904	62.13	4.98	113.2	± 2.7
1450	20.24739	7.92726	0.02506	13.563974	66.54	8.11	114.3	± 2.1
火山岩(Sht16)								
800	40.90162	0.46527	0.13673	0.534630	1.31	2.08	4.7	±150.4
850	90.03898	0.97592	0.28733	5.214394	5.79	4.43	45.6	±275.7
900	45.48567	1.43442	0.11843	10.617135	23.31	6.73	91.7	±70.7
940	17.64187	0.59178	0.01755	12.509980	70.88	10.28	107.5	±15.6
980	16.97458	0.36660	0.01484	12.620800	74.33	9.48	108.4	± 6.4
1020	17.96807	0.34655	0.01868	12.478521	69.43	5.35	107.3	± 4.4
1050	17.62515	0.38095	0.01844	12.208998	69.25	6.23	105.0	± 4.9
1080	16.99371	0.40724	0.01694	12.023626	70.73	6.46	103.5	± 3.3
1110	16.83544	0.46998	0.01576	12.221395	72.56	7.12	105.1	± 1.9
1140	17.32462	0.49255	0.01683	12.394906	71.51	8.02	106.6	± 2.7
1170	18.15213	0.66033	0.01937	12.486624	68.75	7.63	107.3	± 2.4
1200	18.69208	1.03487	0.02142	12.456461	66.58	6.43	107.1	± 1.9
1250	18.87149	1.61694	0.02162	12.627883	66.82	9.50	108.5	± 2.7
1330	22.92370	2.06242	0.03613	12.433196	54.14	4.52	106.9	± 2.1
1450	22.39469	1.88448	0.03420	12.459035	55.54	5.74	107.1	± 2.3
火山岩(Sht18)								
800	78.74380	0.56554	0.26173	1.448856	1.84	1.97	12.9	±18.2
850	189.24489	0.96147	0.62531	4.544667	2.40	4.38	40.3	±40.6
900	94.08039	1.18663	0.28533	9.870298	10.48	7.77	86.3	±17.8
940	21.27121	0.43903	0.03040	12.328428	57.94	9.48	107.2	± 2.0
980	20.05984	0.38474	0.02585	12.455932	62.07	8.75	108.3	± 2.0
1020	21.02690	0.36767	0.03005	12.179294	57.90	6.40	105.9	± 2.1
1050	21.58730	0.37745	0.03226	12.088074	55.98	6.96	105.2	± 2.3
1080	23.09013	0.48025	0.03766	12.002888	51.96	9.80	104.5	± 2.5
1110	24.75671	0.56895	0.04324	12.029227	48.57	7.43	104.7	± 2.8
1140	25.41226	0.60758	0.04502	12.163771	47.84	7.48	105.8	± 3.0
1170	27.03045	0.67962	0.04993	12.335917	45.61	8.17	107.3	± 3.2
1200	25.33075	0.86710	0.04448	12.263561	48.38	6.83	106.7	± 3.4
1250	26.59450	1.34128	0.04871	12.319973	46.27	6.23	107.1	± 3.3
1330	29.33296	1.72995	0.05849	12.205260	41.55	3.00	106.2	± 4.0
1450	28.11178	1.56163	0.05306	12.571883	44.66	5.36	109.3	± 3.5
火山岩(Sh23)								
800	243.87498	1.07126	0.79933	7.763698	3.18	1.08	68.7	±54.4
850	452.17901	0.85577	1.49424	10.707317	2.37	2.97	94.1	±90.7
900	123.49801	0.79431	0.38590	9.532681	7.71	5.21	84.0	±23.7
940	30.29007	0.88810	0.06454	11.297737	37.27	10.89	99.2	± 4.4
980	26.58034	0.85804	0.05034	11.781364	44.29	10.92	103.3	± 3.4
1020	32.87398	0.89845	0.07028	12.185219	37.04	7.03	106.7	± 4.6
1050	33.23477	0.66023	0.07069	12.405373	37.31	5.72	108.6	± 4.7
1080	20.65185	0.50446	0.02798	12.429463	60.16	6.00	108.8	± 2.0
1110	17.31100	0.67712	0.01705	12.332348	71.20	6.45	108.0	± 1.7
1140	17.65415	0.89421	0.01825	12.340914	69.85	5.85	108.1	± 1.7
1170	19.93714	0.90509	0.02608	12.312473	61.71	7.45	107.8	± 1.9
1200	17.47566	1.38969	0.01738	12.465453	71.25	11.03	109.1	± 1.4
1250	16.89189	2.52097	0.01591	12.416105	73.35	9.15	108.7	± 1.4
1330	16.93102	5.20982	0.01683	12.428147	73.08	4.81	108.8	± 1.7
1450	17.41517	8.93531	0.01960	12.429052	70.83	5.44	108.8	± 1.6

续表1

温度/ °C	$^{40}\text{Ar}/^{39}\text{Ar}$	$^{37}\text{Ar}/^{39}\text{Ar}$	$^{36}\text{Ar}/^{39}\text{Ar}$	$^{40}\text{Ar}*/^{39}\text{Ar}_k$	$^{40}\text{Ar}*/\%$	$^{39}\text{Ar}_k/\%$	Age / Ma	$\pm 2s / \text{Ma}$
火山岩 (Sht28)								
800	296.49593	0.88497	0.95605	14.063408	4.74	2.07	123.4	± 74.2
880	249.96868	0.61497	0.76178	24.925178	9.97	8.72	213.3	± 47.0
960	24.55579	0.65110	0.04308	11.882813	48.36	29.39	104.8	± 3.0
990	25.76912	0.71244	0.04589	12.271538	47.59	7.92	108.2	± 2.9
1020	31.25041	0.63736	0.06364	12.502820	39.99	4.21	110.1	± 4.5
1050	30.65379	0.48972	0.06100	12.672812	41.32	3.22	111.6	± 4.6
1100	25.25518	0.45594	0.04399	12.296685	48.67	4.20	108.4	± 3.3
1150	21.61958	0.56800	0.03143	12.381785	57.24	5.04	109.1	± 2.5
1200	20.56691	0.75113	0.02708	12.632402	61.38	9.43	111.2	± 2.0
1230	15.97513	1.11264	0.01219	12.472792	78.00	6.89	109.9	± 1.7
1260	15.32703	1.94431	0.01041	12.426233	80.94	7.74	109.5	± 1.4
1290	15.40786	2.75785	0.01136	12.298386	79.63	4.05	108.4	± 1.9
1340	15.38935	7.21215	0.01230	12.404024	80.11	4.97	109.3	± 1.6
1390	19.09551	12.69582	0.02723	12.190053	63.15	1.73	107.5	± 4.5
1450	26.42477	8.59044	0.05675	10.412478	39.12	0.43	92.2	± 10.5

注:由中国科学院地质与地球物理研究所测试。

比值均小于未分异球粒陨石地幔值 0.512638。在 $^{87}\text{Sr}/^{86}\text{Sr}$ 、 $^{143}\text{Nd}/^{144}\text{Nd}$ 及 ε_{Nd} 相关性图解上(图 8),火山岩的 Nd、Sr 同位素组成靠近第一类富集地幔(EMI)端元,并有向 EM II 延伸的趋势,显示火山岩源区主要受 EMI 控制,并受 EM II 的微弱影响。通过火山岩 $^{206}\text{Pb}/^{204}\text{Pb}$ 与 $^{207}\text{Pb}/^{204}\text{Pb}$ 、 $^{206}\text{Pb}/^{204}\text{Pb}$ 与 $^{208}\text{Pb}/^{204}\text{Pb}$ 关系图解(图 9)也可以看出,该区火山岩主要受 EMI 源区的影响。

2 苏红图火山岩形成及动力学机制

2.1 苏红图火山岩形成机制

火山岩样品中 Nb、Ta 元素具有富集的特点,显示出了具有洋岛玄武岩(OIB)特征,说明了苏红图火山岩岩浆主要来自于对流的软流圈地幔;而从同位素特点来看,该区 6 个火山岩 ε_{Nd} 均小于 0,为 $-6.24\sim-3.08$,很显然岩浆源区具有富集的特征,且在 $^{87}\text{Sr}/^{86}\text{Sr}$ 、 $^{143}\text{Nd}/^{144}\text{Nd}$ 及 ε_{Nd} 相关性图解上,火山岩的 Nd、Sr 同位素组成靠近第一类富集地幔(EMI)端元,并有向 EM II 延伸的趋势,显示本区火山岩的源区受 EMI 影响很大,受 EM II 的微弱影响,对该区火山岩 $^{206}\text{Pb}/^{204}\text{Pb}$ 与 $^{207}\text{Pb}/^{204}\text{Pb}$ 、 $^{206}\text{Pb}/^{204}\text{Pb}$ 与 $^{208}\text{Pb}/^{204}\text{Pb}$ 关系图解,也可以看出,该区火山岩主要受 EMI 源区的影响。

从图 10 中可以看出,随着 ε_{Nd} 的减少,Zr、 P_2O_5 、Rb、Sr、Ba 及 Nb 逐渐富集,暗示着苏红图火山岩没有经历明显的地壳物质混染。而 Na_2O 、 CaO 等主量元素随着 $\varepsilon_{\text{Nd}}(\varepsilon_{\text{Nd}}=-6.24\sim-3.08)$ 的增加而增加,是因为随着部分熔融程度增加,地幔中的角闪石、金红石和金云母开始部分熔融,富集了 Na 和 Ca 元素。此外,如果火山岩岩浆被上地壳物质混染, $^{87}\text{Sr}/^{86}\text{Sr}$

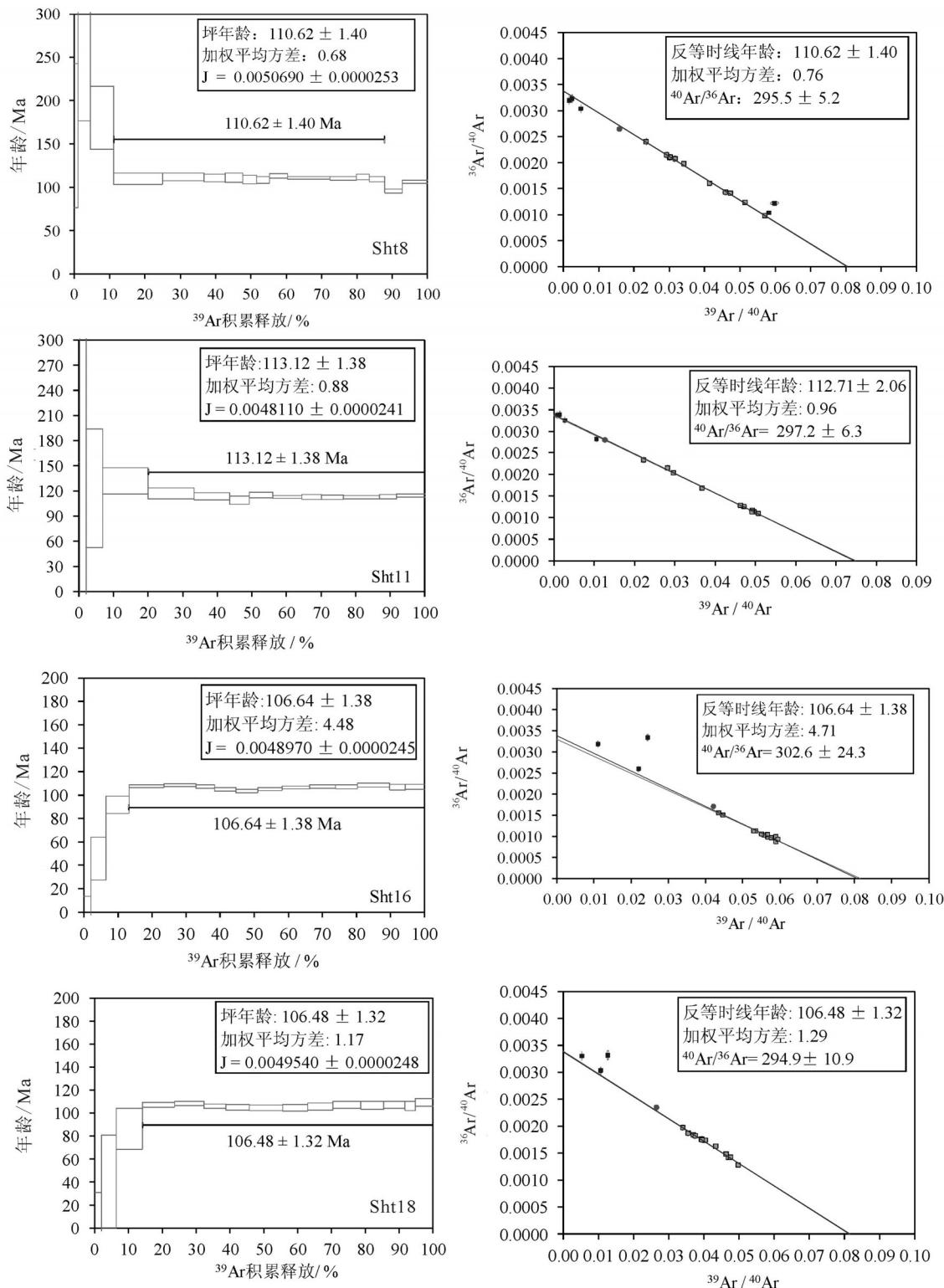
比值应该更大些,变化范围更宽些。苏红图火山岩 $^{87}\text{Sr}/^{86}\text{Sr}$ 值远小于现今大陆地壳 $^{87}\text{Sr}/^{86}\text{Sr}$ 值(表 3 中平均为 0.719),也可以说明它们没有受到明显的地壳物质混染,此外,低 Rb/Ba、Rb/Sr 值和高 Ti/Y 值排除了岩浆被地壳混染的可能。

从 La/Sm-La 图解(图 11)显示苏红图火山岩岩浆既有部分熔融特征,又带有分离结晶的特征。因此,苏红图火山岩可能为平衡部分熔融及随后的分离结晶共同作用的产物。

2.2 苏红图火山岩喷发动力学机制

目前为止,人们对西伯利亚、华北和塔里木等稳定块体之间的碰撞拼合已有了较为一致的认识^[4,27~31]。Zhao Xixi 等^[27]根据古地磁研究结果,提出西伯利亚板块、内蒙古褶皱带和华北板块的碰撞拼合最后结束时间为晚侏罗世;任收麦等^[4]通过对苏红图火山岩古地磁研究,提出在早白垩世,西伯利亚板块、内蒙古褶皱带和华北块体在动力学上已成为整体;研究认为苏红图坳陷火山岩属于钾质碱性玄武岩,而钾质碱性玄武岩通常只形成于大陆环境^[32],通过对火山岩微量元素分析也得到相同结论(图 12~13)。银—额盆地内的地震资料表明^[33],当时盆地沉降和沉积充填受张性断裂构造控制,且岩浆活动并不仅仅局限于银—额盆地,在其周缘、南蒙古地区、甚至在整个中亚地区都普遍出现,说明了早白垩世的岩石圈在较大范围内处于一种拉伸状态^[33],岩石圈厚度为 90 km 左右^[34]。

所以,由于西伯利亚板块、内蒙古褶皱带和华北地块在晚侏罗世碰撞拼合,导致华北克拉通西北部苏红图坳陷下伏岩石圈减薄,随着岩石圈地幔的



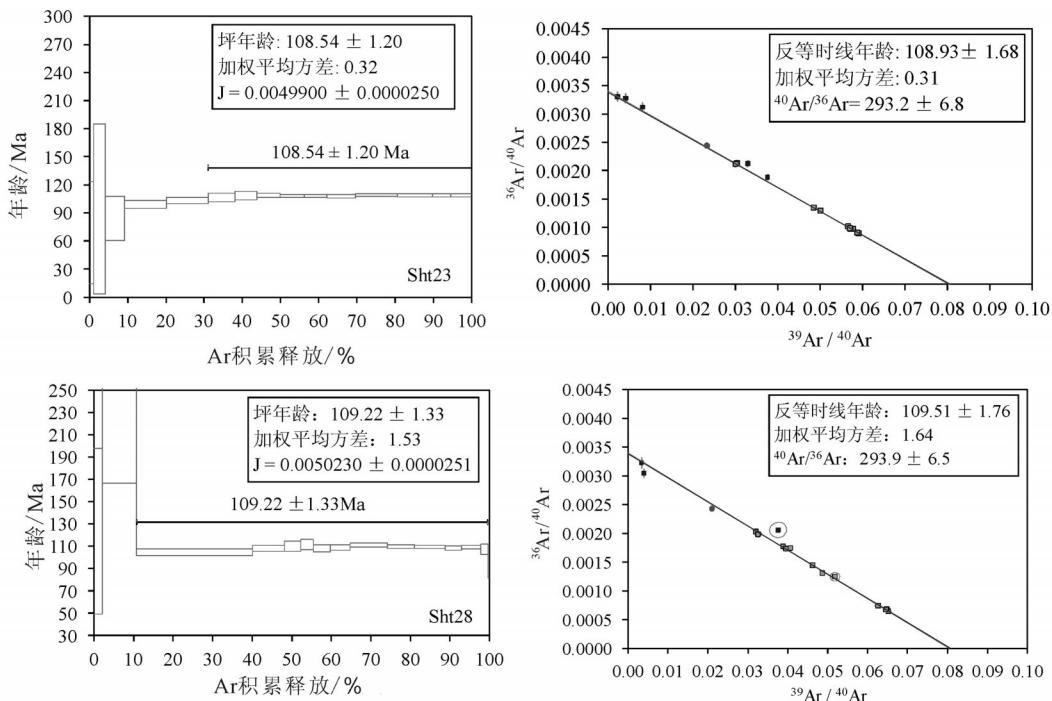


图4 火山岩(Sht8、Sht11、Sht16、Sht18、Sht23、Sht28)坪年龄与反等时线年龄

Fig.4 Plateau age and reverse isochron age of Suhongtu volcanic rocks

减薄,在早白垩世(105~113 Ma)该区下伏软流圈地幔岩浆上涌,在上涌的过程中没有经历地壳物质的明显混染,经平衡部分熔融及随后的分离结晶共同作用形成陆内苏红图高钾玄武岩。

3 克拉通破坏时间与范围的探讨

华北克拉通破坏时间与范围是克拉通破坏研究的重要科学问题。许多学者利用华北克拉通破坏在地壳浅部产生的响应对这些问题展开了一系列研究,但并没有达成统一共识。文章前面部分对位于华北克拉通西北缘的苏红图火山岩年代学与地球化学进行了研究,下面主要针对前人对华北克拉通研究成果,结合本文作者在苏红图火山岩方面所做的一些工作,对华北克拉通在破坏时间与范围上进行一些探讨。

岩浆活动是华北克拉通破坏在地壳浅部产生的重要响应之一。华北地区岩浆活动从晚古生代到新生代均有发育(图14)。华北地区最早的岩浆活动出现在华北北缘的内蒙古隆起(图14-a),为一套钙碱性、I型花岗岩,年龄主要分布在324~300 Ma^[35]。晚石炭世岩浆活动说明了当时克拉通边缘

岩石圈结构和热状态发生了改变,因此无论是何种岩浆动力学机制,正是自晚石炭世开始华北克拉通周边发生了多次构造拼贴和相关岩浆活动,拉开了华北岩石圈破坏的序幕。所以,由此认为华北克拉通有可能是从晚石炭世开始发生破坏的^[15,35]。

晚三叠世岩浆作用的分布较为局限,主要分布在苏鲁超高压变质带及其北延部分(图14-b)。根据Chen et al.^[36]和Xie et al.^[37]的研究,发育在甲子山的晚三叠世正长岩是由于扬子板块向华北板块俯冲过程中发生板片断离,软流圈地幔沿板片窗上涌导致上覆华北岩石圈地幔发生熔融所致。研究显示在徐淮弧形构造区内早侏罗世(191 Ma)同造山侵入岩中发现的高压榴辉岩包体^[38]及在徐淮弧形构造带中发育的早侏罗世和早白垩世侵入杂岩体中发现了含220 Ma的榴辉岩捕虏体^[39],与大别—苏鲁造山带超高压变质作用时间一致。这一结果暗示扬子地块与华北板块碰撞导致了华北克拉通的破坏,即碰撞导致了华北下地壳的加厚,形成了大量榴辉岩,继而发生拆沉作用^[19,40]。所以,在晚三叠世,克拉通边缘的岩石圈结构和热状态发生了改变。

华北地区大约从中侏罗世开始出现较大范围

表2 银—额盆地苏红图坳陷火山岩岩石地球化学分析结果

Table 2 Analytical results of volcanic rocks

	Sht-04	Sht-05	Sht-06	Sht-07	Sht-08	Sht-09	Sht-10	Sht-11	Sht-12	Sht-13	Sht-14	Sht-15
SiO ₂ / %	49.96	48.13	47.90	49.40	47.80	48.60	47.85	47.90	49.03	47.58	49.45	52.02
TiO ₂	2.03	1.73	1.77	2.04	1.79	1.94	1.76	1.75	2.01	1.69	2.13	1.98
Al ₂ O ₃	17.94	15.84	15.97	16.55	15.69	16.34	15.66	15.54	16.16	15.75	16.80	17.74
TFe ₂ O ₃	7.17	11.10	10.79	9.82	11.38	10.74	11.28	11.32	10.31	10.99	9.84	7.60
MnO	0.07	0.16	0.13	0.13	0.15	0.13	0.15	0.16	0.14	0.16	0.17	0.09
MgO	4.63	7.31	6.55	5.38	7.06	5.93	7.61	7.82	5.84	7.60	5.02	4.56
CaO	6.33	6.20	6.85	6.19	6.88	7.61	7.33	6.52	6.66	6.41	7.34	4.85
Na ₂ O	3.75	5.29	4.39	5.15	4.09	3.34	3.41	4.51	4.70	4.59	3.62	4.03
K ₂ O	3.02	1.07	1.03	1.77	1.47	1.77	1.55	1.17	1.86	1.16	2.18	2.99
P ₂ O ₅	1.05	0.43	0.44	0.59	0.43	0.48	0.43	0.44	0.57	0.43	0.60	0.85
LOI	3.88	3.02	4.24	2.96	3.16	3.10	3.14	3.06	2.86	3.42	2.26	2.70
Total	99.83	100.26	100.06	99.98	99.91	99.98	100.16	100.18	100.14	99.77	99.42	99.41
Li / 10 ⁻⁶	56.03	22.40	31.40	21.82	33.15	30.86	27.51	26.35	20.58	29.70	41.19	41.01
Be	0.97	1.13	1.15	1.57	1.11	1.27	1.10	1.11	1.51	1.12	1.10	1.10
Sc	16.82	26.27	26.66	24.72	26.00	26.54	26.70	26.96	24.70	26.41	25.00	18.12
V	111.10	169.61	167.80	156.65	170.66	170.98	172.59	171.010	157.68	168.85	165.84	116.66
Cr	49.17	237.14	221.13	181.90	234.46	222.06	274.12	230.59	188.94	233.57	178.54	69.22
Co	25.47	41.56	39.49	32.76	41.20	37.61	40.86	43.52	35.09	42.20	34.54	25.35
Ni	42.07	112.34	109.12	78.92	114.76	102.33	114.31	119.61	93.34	111.24	96.49	33.23
Cu	23.64	42.28	46.19	42.76	48.22	50.18	44.57	38.77	37.85	44.72	57.75	33.29
Zn	121.00	98.73	95.83	97.52	98.33	99.42	95.31	99.44	95.39	100.12	121.60	149.72
Ga	23.29	18.60	18.71	19.28	18.98	19.99	17.76	17.84	19.21	18.44	20.21	23.81
Rb	26.86	7.60	6.87	45.63	25.27	31.09	25.23	9.54	47.53	10.72	37.72	33.77
Sr	827.17	338.17	406.24	473.09	447.89	549.36	473.00	324.93	509.25	314.08	555.06	747.77
Y	41.13	25.97	25.78	30.72	26.25	28.17	25.96	27.33	30.96	25.89	30.99	36.64
Zr	331.88	172.19	172.11	236.21	172.73	190.96	170.12	173.99	224.93	165.93	217.00	367.20
Nb	57.28	29.24	29.22	42.17	29.40	32.34	29.50	30.30	41.94	28.92	40.05	56.91
Cs	0.29	8.59	6.35	8.36	5.54	1.26	1.99	7.23	6.21	7.21	0.39	0.15
Ba	1296.51	543.63	446.16	731.17	482.08	625.00	519.31	542.94	708.23	514.73	684.56	1124.50
La	62.32	23.29	23.10	33.46	23.58	26.11	22.98	24.27	33.45	22.75	32.89	56.15
Ce	126.34	45.43	45.30	64.43	45.97	50.51	45.46	47.40	64.27	44.17	64.04	115.74
Pr	15.99	5.88	5.68	8.18	5.90	6.55	5.88	6.22	7.96	5.66	8.10	14.15
Nd	62.94	23.77	23.04	33.42	24.29	26.14	23.03	24.44	32.00	23.00	33.76	54.55
Sm	11.21	5.22	5.06	6.57	5.40	5.67	4.91	5.39	6.52	5.05	6.76	10.37
Eu	2.71	1.75	1.67	2.03	1.74	1.88	1.73	1.73	2.15	1.69	2.18	2.60
Gd	10.40	5.35	5.23	6.55	5.32	5.78	4.94	5.02	5.97	4.93	6.29	9.35
Tb	1.53	0.88	0.84	1.02	0.86	0.94	0.81	0.82	0.97	0.81	0.99	1.34
Dy	8.37	5.25	5.04	6.07	5.29	5.57	4.94	5.27	5.90	4.79	5.86	7.13
Ho	1.62	1.05	1.03	1.25	1.06	1.17	1.02	1.06	1.18	0.97	1.18	1.42
Er	4.19	2.84	2.81	3.26	2.88	3.07	2.74	2.83	3.12	2.63	3.10	3.69
Tm	0.59	0.40	0.40	0.47	0.42	0.44	0.39	0.42	0.47	0.39	0.46	0.53
Yb	3.65	2.51	2.50	3.02	2.58	2.75	2.50	2.61	3.02	2.47	2.85	3.32
Lu	0.54	0.38	0.38	0.46	0.39	0.41	0.38	0.38	0.46	0.37	0.43	0.50
Hf	8.99	4.27	4.23	5.78	4.28	4.78	4.21	4.33	5.47	3.99	5.38	8.83
Ta	3.51	1.86	1.85	2.72	1.90	2.11	1.82	1.91	2.55	1.81	2.53	3.58
Tl	0.13	0.05	0.04	0.07	0.06	0.07	0.06	0.05	0.07	0.05	0.06	0.11
Pb	10.12	3.56	4.33	6.00	3.29	3.44	3.08	3.10	4.17	3.07	4.46	7.32
Bi	0.06	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.02	0.02
Th	5.45	2.34	2.31	3.59	2.40	2.66	2.18	2.34	3.48	2.20	3.26	5.49
U	2.23	0.57	0.59	0.86	0.54	0.62	0.53	0.60	0.91	0.58	0.47	1.02
REE	312.39	123.98	122.08	170.18	125.68	136.99	121.69	127.85	167.42	119.68	168.88	280.82
LREE	281.51	105.34	103.85	148.09	106.88	116.86	103.98	109.46	146.35	102.32	147.73	253.55
HREE	30.89	18.65	18.23	22.09	18.80	20.13	17.71	18.39	21.07	17.36	21.14	27.27
LREE/HREE	9.11	5.65	5.70	6.70	5.69	5.81	5.87	5.95	6.94	5.89	6.99	9.30
δCe	0.94	0.91	0.93	0.91	0.91	0.91	0.92	0.91	0.92	0.91	0.92	0.96
δEu	0.75	1.00	0.98	0.94	0.98	1.00	1.06	1.00	1.03	1.02	1.01	0.79

续表2

	Sht-16	Sht-17	Sht-18	Sht-19	Sht-20	Sht-21	Sht-22	Sht-23	Sht-25	Sht-26	Sht-27	Sht-28
SiO ₂ / %	54.63	51.14	56.28	49.78	51.51	48.77	48.68	49.00	48.88	49.24	48.82	48.88
TiO ₂	1.83	2.09	1.89	2.12	2.00	2.12	2.04	2.04	2.01	2.04	1.95	2.02
Al ₂ O ₃	15.44	16.67	14.67	16.67	15.69	16.93	16.21	16.42	16.59	16.48	16.33	16.31
TFe ₂ O ₃	8.88	9.05	10.96	9.42	9.34	10.33	10.42	9.24	9.08	9.19	9.38	9.30
MnO	0.10	0.12	0.06	0.13	0.15	0.19	0.17	0.14	0.13	0.14	0.14	0.14
MgO	0.51	0.94	0.78	4.94	1.48	5.64	6.29	5.95	6.03	5.91	6.49	6.31
CaO	7.34	8.36	5.52	6.77	7.76	6.32	6.85	6.36	6.62	6.11	6.87	6.57
Na ₂ O	3.88	4.48	4.00	4.23	4.13	3.62	3.53	5.19	5.03	5.39	4.68	4.87
K ₂ O	2.90	3.03	2.95	2.28	3.09	2.24	2.08	1.87	2.00	1.89	1.84	1.74
P ₂ O ₅	0.72	0.81	0.72	0.60	0.72	0.60	0.58	0.61	0.60	0.62	0.59	0.58
LOI	3.18	3.59	2.04	2.66	4.00	2.70	2.94	2.68	2.38	2.82	2.60	2.74
Total	99.41	100.29	99.89	99.61	99.88	99.45	99.80	99.50	99.35	99.84	99.69	99.46
Li / 10 ⁻⁶	51.27	33.06	43.04	37.02	30.72	55.37	52.80	22.98	11.90	25.98	29.19	22.17
Be	1.56	1.49	1.87	1.77	3.18	1.20	1.15	1.89	1.65	1.68	1.58	1.65
Sc	16.89	18.52	15.04	24.81	18.15	25.36	24.03	22.19	23.00	21.79	22.99	23.24
V	110.17	127.71	101.98	168.64	134.03	172.95	156.32	141.32	142.06	134.99	149.07	148.43
Cr	137.28	133.09	124.20	188.84	128.91	190.88	177.89	130.34	153.00	125.47	225.73	149.68
Co	10.38	16.14	13.62	34.50	15.18	38.26	34.86	30.77	30.98	30.07	33.68	32.53
Ni	15.93	33.58	27.84	85.25	27.69	92.24	93.17	78.89	79.88	77.56	88.85	85.09
Cu	22.43	47.30	26.60	41.63	33.17	37.41	27.83	32.47	31.57	32.43	30.84	31.67
Zn	48.99	62.79	50.23	108.78	63.35	130.00	117.27	98.94	85.21	85.94	86.53	89.86
Ga	18.49	19.77	18.34	18.89	19.42	20.91	19.21	19.04	19.01	19.22	18.61	18.65
Rb	58.56	59.02	58.25	45.28	64.20	40.99	36.13	12.03	18.74	13.73	13.00	15.88
Sr	643.49	634.63	642.72	509.84	609.82	542.11	489.81	554.08	600.25	539.63	564.15	574.73
Y	29.67	30.21	28.17	29.53	28.75	29.33	28.53	26.94	27.71	27.95	26.40	27.65
Zr	262.42	264.97	262.25	222.85	282.83	211.46	205.21	223.30	229.76	233.31	223.98	229.86
Nb	46.63	49.15	47.02	43.77	47.30	40.90	38.31	48.04	48.98	50.79	47.41	49.40
Cs	1.57	1.26	3.19	14.48	1.81	0.51	1.46	1.86	1.26	2.12	1.52	1.62
Ba	1659.75	816.17	782.62	877.45	1029.91	706.57	605.90	810.74	809.16	792.00	818.67	846.99
La	37.04	40.77	38.03	33.07	37.33	32.15	31.53	32.43	34.10	34.66	33.41	34.20
Ce	73.69	80.76	75.49	66.92	74.31	61.69	61.16	62.00	64.78	65.16	62.90	63.77
Pr	9.47	10.56	9.73	8.41	9.73	8.24	7.79	7.75	7.93	8.14	7.97	8.07
Nd	37.21	40.36	38.48	33.88	37.84	32.39	30.45	30.27	31.32	30.95	30.60	31.63
Sm	6.95	7.51	7.25	6.82	7.48	6.82	6.29	6.01	6.50	6.41	6.19	6.27
Eu	2.09	2.27	2.04	2.12	2.13	2.18	2.11	2.07	2.12	2.16	1.98	2.00
Gd	6.84	7.20	6.88	6.57	7.40	6.88	5.77	5.74	5.63	5.93	5.64	5.74
Tb	0.99	1.06	0.97	0.99	1.03	1.01	0.89	0.86	0.91	0.94	0.91	0.89
Dy	5.60	5.98	5.53	5.79	5.57	5.78	5.42	5.09	5.29	5.51	5.11	5.33
Ho	1.13	1.15	1.13	1.16	1.09	1.16	1.08	0.99	1.04	1.08	1.01	1.04
Er	3.04	3.07	2.99	3.10	2.98	3.12	2.88	2.64	2.65	2.80	2.69	2.79
Tm	0.44	0.45	0.44	0.46	0.44	0.46	0.42	0.38	0.39	0.40	0.39	0.40
Yb	2.90	2.95	2.77	2.95	2.82	2.91	2.51	2.41	2.46	2.48	2.49	2.54
Lu	0.44	0.44	0.42	0.45	0.43	0.44	0.37	0.36	0.37	0.39	0.38	0.39
Hf	6.04	6.23	6.18	5.37	5.78	5.28	5.06	5.62	5.61	5.85	5.48	5.87
Ta	2.77	2.91	2.87	2.67	2.69	2.53	2.36	3.04	3.09	3.12	3.02	3.23
Tl	0.140	0.12	0.17	0.09	0.14	0.06	0.05	0.045	0.07	0.05	0.15	0.06
Pb	7.000	6.15	8.29	4.34	7.12	4.08	3.89	3.94	4.15	4.12	4.17	4.20
Bi	0.014	0.01	0.02	0.01	0.03	0.02	0.01	0.011	0.01	0.01	0.01	0.01
Th	3.710	3.81	3.84	3.87	3.55	3.30	2.94	3.35	3.49	3.61	3.72	3.65
U	0.98	1.44	1.13	0.93	2.50	0.43	0.53	0.94	1.00	0.96	0.97	1.03
REE	187.81	204.53	192.13	172.68	190.57	165.22	158.68	158.99	165.48	167.01	161.65	165.05
LREE	166.43	182.22	171.02	151.21	168.81	143.47	139.33	140.52	146.75	147.48	143.03	145.93
HREE	21.38	22.30	21.12	21.46	21.76	21.76	19.35	18.48	18.73	19.53	18.62	19.12
LREE/HREE	7.78	8.17	8.10	7.05	7.76	6.59	7.20	7.61	7.83	7.55	7.68	7.63
δCe	0.93	0.92	0.92	0.94	0.92	0.89	0.91	0.91	0.92	0.90	0.90	0.89
δEu	0.92	0.93	0.87	0.96	0.87	0.96	1.05	1.06	1.05	1.06	1.00	1.00

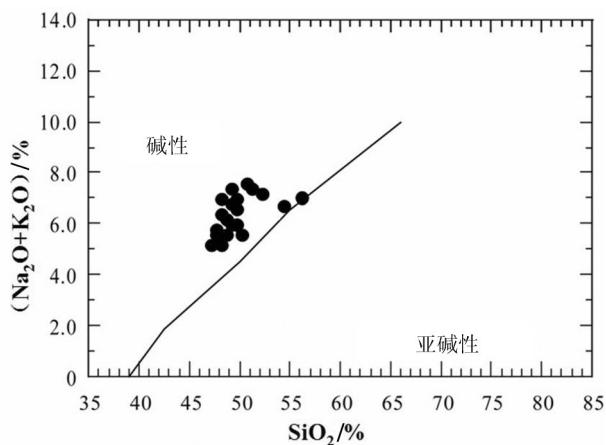


图 5 SiO_2 - $(\text{K}_2\text{O}+\text{Na}_2\text{O})$ 图解
Fig. 5 SiO_2 - $(\text{K}_2\text{O}+\text{Na}_2\text{O})$ diagram

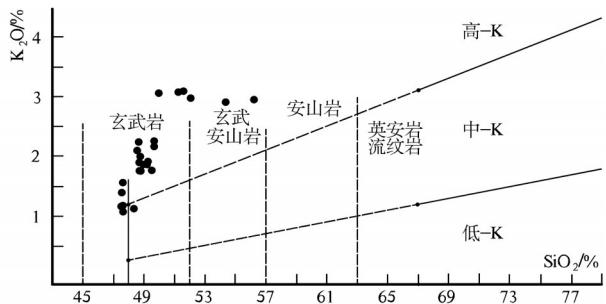


图 6 SiO_2 - K_2O 图解
Fig. 6 SiO_2 - K_2O diagram

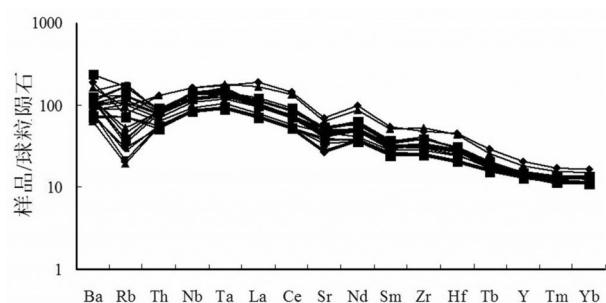


图 7 苏红图火山岩微量元素/球粒陨石标准化模式图
Fig. 7 Chondrite-normalized trace element diagram

的岩浆活动(图 14-C),其活动时期主要在 190~155 Ma^[22,41,42]。Gao et al.^[20]通过对辽西地区中生代高镁 adakite 研究,结合该岩石高 Ni 和 Cr 含量及其在矿物中的分布特征,认为这些岩石是拆沉进入地幔的

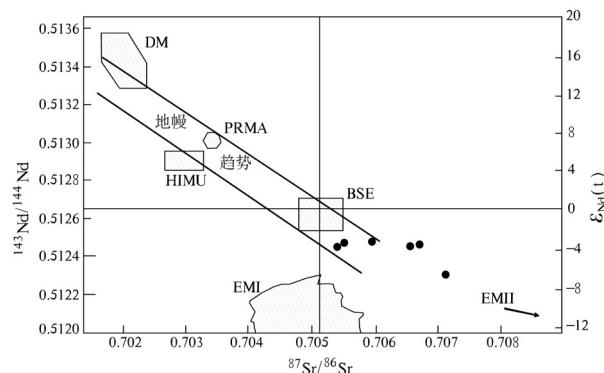


图 8 $^{87}\text{Sr}/^{86}\text{Sr}$ 、 $^{143}\text{Nd}/^{144}\text{Nd}$ 及 ε_{Nd} 关系图解^[25]
DM—亏损地幔; BSE—全硅酸盐地球; EM I、EM II—富集地幔 I、II;
HIMU—具有高 U/Pb 比值的地幔; PREMA—普通地幔
Fig. 8 Diagram showing relationships between $^{87}\text{Sr}/^{86}\text{Sr}$, $^{143}\text{Nd}/^{144}\text{Nd}$ and $\varepsilon_{\text{Nd}}^{[25]}$
DM—Depleted mantle; BSE—Whole silicate earth; EM I, EM II—Type I and II enriched mantle; HIMU—Range of high U/Pb ratio mantle;
PREMA—Ordinary mantle

地壳岩石部分熔融而成,从而暗示华北克拉通的破坏发生在中侏罗世以后。Charles et al.^[43]认为晚侏罗世时华北克拉通东部南、北缘局部开始了伸展活动,但是在其内部还很少发育岩浆活动与伸展构造,反映了在该时期华北克拉通东部岩石圈的改造只发生在其南、北缘^[24]。在克拉通东部南缘胶北地区与合肥盆地北缘发现了晚侏罗世的玲珑岩基和荆山—涂山花岗岩体,研究表明这些晚侏罗世岩浆活动是发生在岩石圈减薄的环境中^[44~46]。

在晚侏罗世—早白垩世时期,岩浆活动的范围和强度达到了顶峰,除了分布在燕山、太行山和苏鲁大别带外,也出现在华北地台腹地及郯庐断裂两侧,且岩浆活动呈现出由克拉通边缘向克拉通腹地迁移的趋势(图 14-d)^[47]。此外,在克拉通西北缘的苏红图坳陷在早白垩世时期也发生了强烈的岩浆活动。克拉通破坏在地壳浅部响应除了岩浆作用外,还有变质核杂岩、成矿作用及断陷盆地的产生。变质核杂岩在 120~130 Ma 最为发育^[41],且大部分分布在克拉通北缘。在华北克拉通发育了两条近南北向的岩浆—矿床带,岩浆—成矿作用年龄分别集中在 140~102 Ma 和 135~100 Ma。在中国东部的大型金矿成矿作用出现在 125~120 Ma^[48,49],其与吴福元等^[41]研究认为金属金矿床形成在 125 Ma 左右的一个狭窄的时间范围内,以及翟明国^[50]研究认为

表3 苏红图火山岩同位素
Table 3 Isotopes of Suhongtu volcanic rocks

样品号	$^{206}\text{Pb}/^{204}\text{Pb}$	2SE(M)%	$^{207}\text{Pb}/^{204}\text{Pb}$	2SE(M)%	$^{207}\text{Pb}/^{204}\text{Pb}$	2SE(M)%	$^{87}\text{Sr}/^{86}\text{Sr}$	(2 σ)	$^{143}\text{Nd}/^{144}\text{Nd}$	(2 σ)
sht-06	17.9368	0.015	15.4832	0.016	38.1678	0.018	0.706611	0.000012	0.512457	0.000015
sht-08	17.8987	0.009	15.4818	0.010	38.1503	0.010	0.706782	0.000008	0.512475	0.000015
sht-11	17.9089	0.011	15.4831	0.012	38.1483	0.013	0.705867	0.000012	0.512480	0.000015
sht-18	17.8307	0.011	15.4916	0.012	37.4447	0.013	0.707098	0.000014	0.512318	0.000014
sht-23	17.8386	0.011	15.4689	0.012	37.9882	0.011	0.705498	0.000010	0.512471	0.000015
sht-28	17.8309	0.020	15.4745	0.023	38.0358	0.030	0.705477	0.000012	0.512445	0.000016

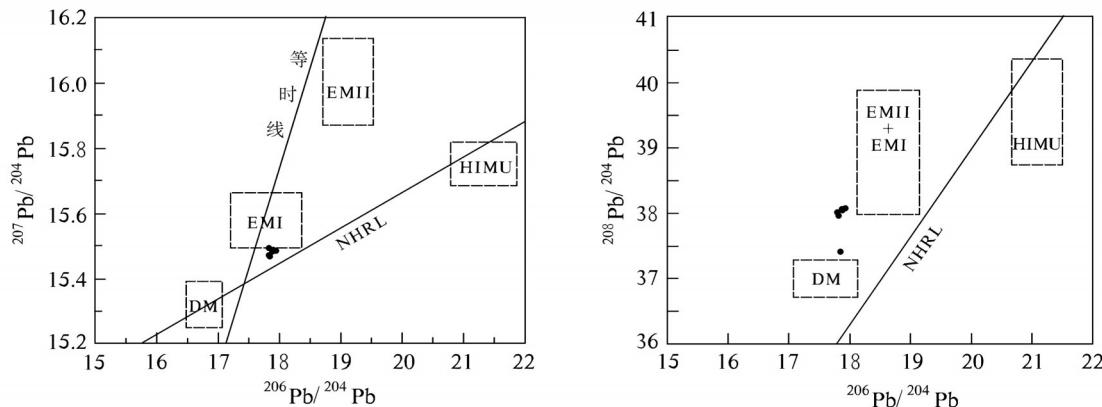


图9 火山岩 $\frac{^{207}\text{Pb}}{^{204}\text{Pb}}$ - $\frac{^{206}\text{Pb}}{^{204}\text{Pb}}$ 与 $\frac{^{208}\text{Pb}}{^{204}\text{Pb}}$ - $\frac{^{206}\text{Pb}}{^{204}\text{Pb}}$ 关系图解^[26]

EMI、EMII—富集地幔端元; DM—亏损地幔同位素组成范围; HIMU—高 U/Pb 地幔组成范围; NHRL—北半球参考线

Fig. 9 Pb isotopic compositions of Suhongtu volcanic rocks $^{206}\text{Pb}/^{204}\text{Pb}$ versus $^{207}\text{Pb}/^{204}\text{Pb}$ and $^{206}\text{Pb}/^{204}\text{Pb}$ versus $^{208}\text{Pb}/^{204}\text{Pb}$ ^[26]
EMI—Type I enriched mantle, EMII—Type II enriched mantle; DM—Range of depleted mantle isotopic composition; HIMU—Range of high U/Pb ratio mantle; NHHRL—Northern hemisphere reference line

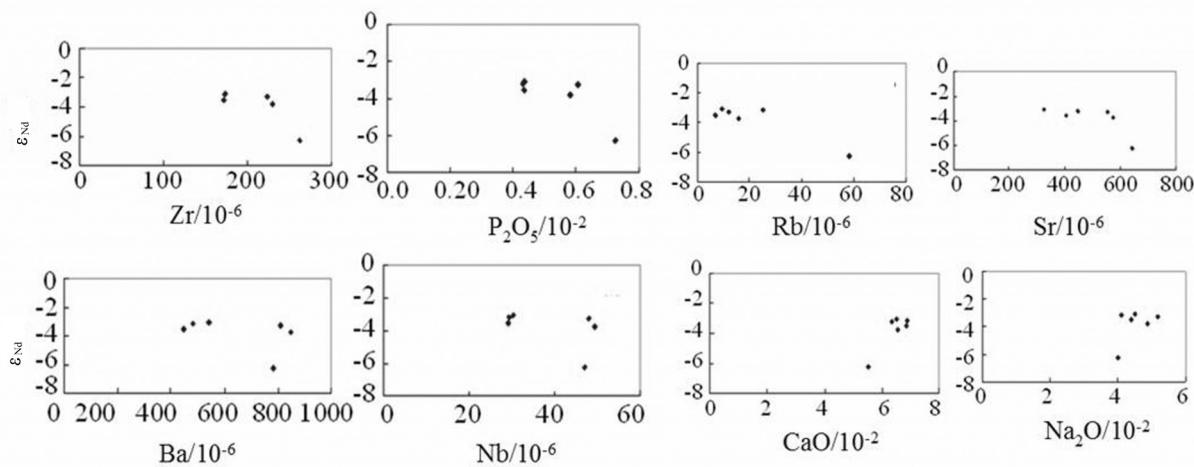


图10 苏红图火山岩 $\epsilon_{\text{Nd}}(t)$ 与Zr、P₂O₅、Rb、Sr、Ba、Nb、CaO、Na₂O关系图

Fig.10 Diagram showing relationships between $\epsilon_{\text{Nd}}(t)$ and Zr, P₂O₅, Rb, Sr, Ba and Nb for volcanic rocks of Suhongtu depression

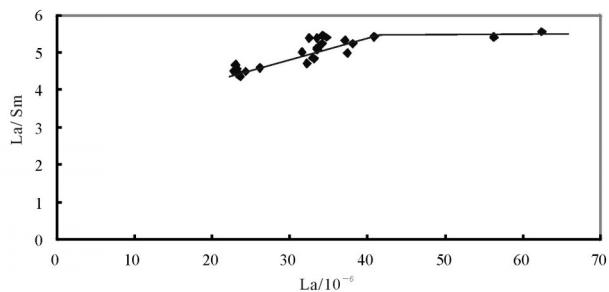


图11 La-La/Sm关系图

Fig.11 Diagram showing relationship between $w(\text{La})/w(\text{Sm})$ and La

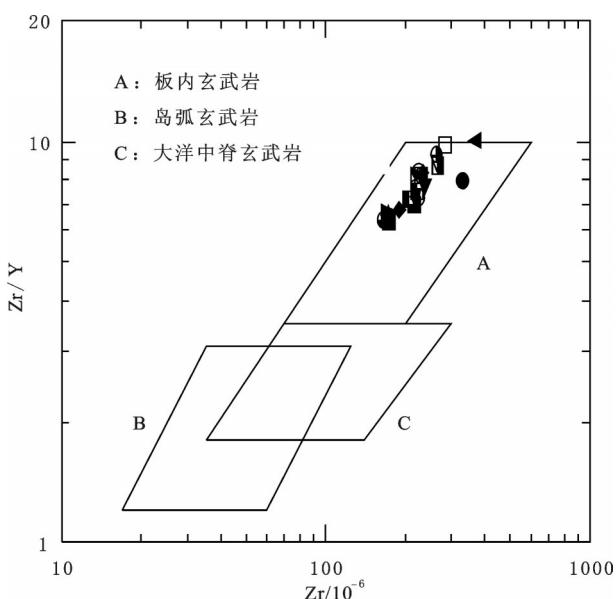


图12 Zr/Y与Zr关系图

Fig. 12 Diagram showing relationship between Zr/Y and Zr

遍布在华北东部的克拉通边缘及克拉通内部的爆发式大规模金成矿作用主期在 (120 ± 10) Ma 的结论相一致。120~140 Ma 是中国东部及邻区裂陷盆地主要形成和发育时期^[51], 盆地形成时代和强烈的岩浆活动、大范围的变质核杂岩和大规模成矿作用时间都相一致^[52]。所以, 在晚侏罗世—早白垩世时期克拉通破坏产生的浅部响应在范围上和强度上达到高峰, 说明了岩石圈在此时期内遭受强烈破坏, 甚至破坏达到最高峰^[22]。

晚中生代—新生代, 火山作用仍有分布^[53~56], 但强度相对减弱(图 14-e), 玄武岩以碱性和强碱性为主, 且主要出现在裂谷两侧^[47]。如在汾渭裂谷系北

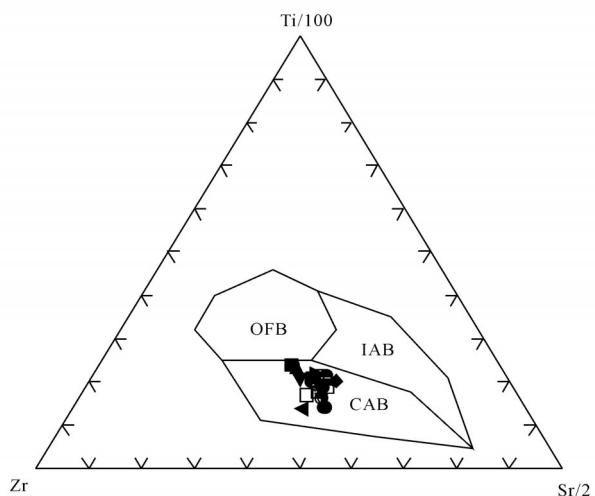


图13 Ti/100、Zr、Sr/2之间的关系图

Fig.13 Diagram showing relationships between $\text{Ti}/100$, Zr and $\text{Sr}/2$

部发现新生代玄武岩喷发, 范围主要在恒山—大同及其以北地域^[57]。

上述发育地壳浅部响应的区域可能也是克拉通发生破坏的区域, 这些破坏区域与地震学成像结果显示的华北克拉通现今岩石圈厚度存在明显的区域差异^[58~62]是相吻合的。克拉通东部广泛分布的较薄的岩石圈^[58~62]以及新生代相对饱满的地幔包体地球化学特征^[63~67], 反映该地区岩石圈地幔的物理和化学性质在显生宙发生了根本改变, 即克拉通岩石圈已经被普遍减薄甚至发生了破坏。克拉通东部岩石圈厚度为 60~100 km, 向西北内部逐渐增厚至 90~100 km, 向克拉通内部逐渐过渡为 100~150 km。同时, 较薄的地壳^[68~72]、较高的地表热流^[73]和活跃的地震活动^[74]表明, 东部地壳岩石圈改造不仅横向范围广泛, 而且在深度上涉及整个岩石圈。中、西部地区薄于 100 km 的岩石圈在河套和汾渭两个狭长的新生代局部裂陷区^[24], 在银—额盆地苏红图坳陷下伏岩石圈也发生减薄, 厚度仅为 90 km 左右^[34], 其他区域岩石圈普遍厚于 120 km, 而在鄂尔多斯盆地之下的岩石圈厚度达 200 km, 且岩浆活动微弱、地热梯度低, 表明了克拉通中、西部大部分地区可能保留着太古宙克拉通的核心部分。

通过克拉通破坏时在地壳浅部产生的响应对破坏时间与范围的探讨发现华北克拉通破坏的时间在空间上具有不均一性^[75~77]。在石炭纪、三叠纪、

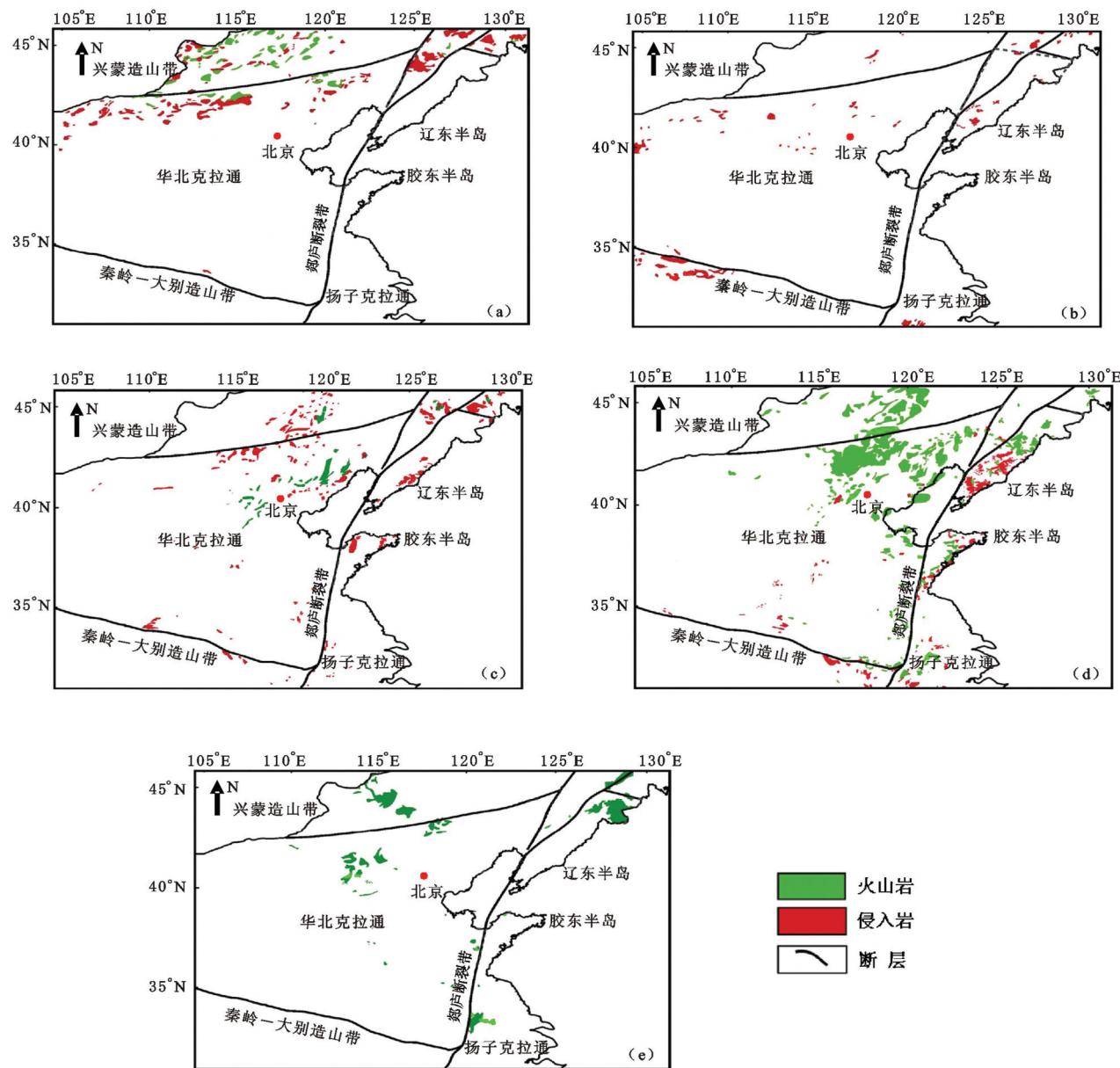


图14 晚古生代—新生代华北克拉通岩浆岩分布图^[47]
a—石炭纪—二叠纪; b—三叠纪; c—侏罗纪; d—白垩纪; e—新生代
Fig. 14 Distribution of NCC late Paleozoic – Cenozoic igneous rocks^[47]
a—Carboniferous–Permian; b—Triassic; c—Jurassic; d—Cretaceous; e—Cenozoic

侏罗纪时发生的破坏主要集中在克拉通周缘地区,而在白垩纪则影响到了克拉通腹地,上述通过对苏红图坳陷火山岩研究得出克拉通西北缘的苏红图坳陷在早白垩世也发生了破坏。总体来说,破坏主要集中在太行山以东地区,以西的汾渭裂陷、河套裂陷与苏红图坳陷等地区发生了减薄,在地理分布上呈现出不连续的特征。

4 克拉通破坏的动力学背景与破坏机制的探讨

从华北克拉通构造纲要图中我们可以发现华北克拉通周边均为显生宙造山带所隔(图1)。克拉通北缘为古亚洲洋闭合后形成的兴—蒙造山带,南侧为著名的大别—秦岭造山带,东侧为苏鲁带和太

平洋俯冲带,克拉通中西部的河套裂陷、汾渭裂陷分别与古元古代高温变质孔兹岩带^[60,61]及约18.5亿年前华北克拉通东、西部块体拼合时形成的中部造山带内^[60,62]位置大致重合,而苏红图坳陷位于中亚造山带南缘,也位于两板块拼合交汇处。这些构造带在不同时期经历了重大地质事件而发生了改造。如:在晚古生代期间,克拉通北部古亚洲洋向华北克拉通俯冲,最终在二叠纪末发生了华北克拉通与蒙古地块碰撞^[78],随后,蒙古—鄂霍次克洋在晚侏罗纪中期关闭,使得华北—蒙古陆块与西伯利亚克拉通发生拼合^[79,80];晚三叠世扬子与华北板块发生了强烈碰撞^[81~85];中生代发生了太平洋板块对欧亚板块俯冲,在新生代发生了印度板块对欧亚板块^[86~89]的俯冲等重大构造事件。

针对晚侏罗世—早白垩世时期,从全球来看,在早白垩世发生了大规模的海底火山活动、太平洋超级地幔柱的上升^[90]与特提斯洋关闭的相关的地幔雪崩事件^[91],以及冈瓦纳大陆的裂解事件^[92]。从区域来看,在中侏罗世 Izanagi 板块开始以 4.7 cm/a 的速度向东亚大陆正向俯冲,而在早白垩世初期 Izanagi 板块运动方向和速度突然发生改变,以 30 cm/a 的速度向正北方向俯冲于东亚大陆之下^[93]。Kopper et al.^[94,95]研究也认为自从 140 Ma 以来,太平洋板块向欧亚板块俯冲方向改变了多次,而最重要的一次改变是发生在 120~125 Ma^[94,96],方向大致由南转变为北西方向,转幅达 80°^[97]。在晚侏罗世—早白垩世克拉通破坏除了可能达到峰期外,还有学者认为中国东部在该时期构造体制发生了根本的转变,侏罗纪对应于古太平洋板块相关的挤压构造环境,而白垩纪是伸展构造背景^[98~101]。如:郑亚东^[102]等和 Davis et al.^[103]对河北承德一带的构造研究表明区域内的挤压构造主要发生在 127~180 Ma,而伸展作用主要发生在 127 Ma 以后;Tang et al.^[104]通过对宁芜盆地火山岩岩浆认为,火山岩在短时间内(133~130 Ma)快速喷发而形成,暗示了中国东部早白垩世构造体制的快速转变。

研究发现克拉通破坏在石炭纪、三叠纪、侏罗纪时主要集中在克拉通周缘地区,而白垩纪则影响到了克拉通腹地。华北克拉通周缘板块碰撞拼合的时间、位置与克拉通破坏的时间、范围相吻合。那么克拉通的破坏与当时的区域构造背景甚至是

全球构造背景及克拉通周缘的存在的构造带之间是否存在联系?

通过矿物物理实验和理论模拟结果显示,先前存在于岩石圈中的古老构造带在后期的热-构造事件过程中,可以引起上地幔顶部热扩散^[105]和机械变形^[106]的各向异性,并进一步导致岩石圈的不均匀加热和应变集中,有利于古老构造带的活化,从而使得古老岩石圈发生改造^[105,107]。所以,根据克拉通破坏的时间、范围及华北地区在早白垩世发生构造体制重大转折的客观事实,结合当时全球及区域构造背景,认为二叠纪末、晚三叠世甚至新生代等时期发生在克拉通周缘的板块碰撞拼合事件使得先前存在的古老构造带的活化,造成岩石圈发生破坏。而在晚侏罗世—早白垩世初期,处于全球构造背景中的太平洋板块在对亚欧板块俯冲过程中,全球构造事件导致太平洋板块俯冲速度与方向发生重大变化,致使华北克拉通构造体制发生重大转折,从而造成了华北克拉通东部在早白垩世发生了强烈破坏,甚至达到了破坏高峰期。

所以,克拉通周缘的构造薄弱带的俯冲、碰撞与华北克拉通破坏关系极为密切,它们可能对华北克拉通破坏产生了重要影响^[108~110],构造薄弱带的位置可能就是岩石圈减薄的起始位置,并且它们的俯冲与碰撞的时间是华北克拉通破坏的起始时间,从而最终导致了克拉通的破坏在时空上存在不一致性^[75~77]。

5 结 论

西伯利亚板块、内蒙古褶皱带和华北地块在晚侏罗世碰撞拼合,导致华北克拉通西北部苏红图坳陷下伏岩石圈减薄,随着岩石圈地幔的减薄,在早白垩世(105~113 Ma)该区下伏软流圈地幔岩浆上涌,在上涌的过程中没有经历地壳物质的明显混染,经平衡部分熔融及随后的分离结晶共同作用形成了陆内苏红图高钾火山岩。

华北克拉通破坏在石炭纪、三叠纪、侏罗纪时主要集中在克拉通周缘地区,而白垩纪则影响到了克拉通腹地。克拉通破坏范围主要发生在太行山东部地区,太行山以西的河套裂陷、汾渭裂陷发生了减薄,而苏红图坳陷在早白垩世也发生减薄,破坏范围分布在地理上呈不连续分布特征,造成这种分布特征的主要原因是由于在克拉通周缘先前存

在的构造薄弱带受到了后期构造活动的改造,改造导致了岩石圈的破坏,从而导致了克拉通的破坏在时空上存在不一致性。

致谢: 在野外工作的还有浙江油田的张磊、胜利油田的禚元杰、中国石油大学(华东)研究生王安东、大庆油田王金华、孟玮等,在成文过程中得到了中科院广州地化所郭锋研究员的无私帮助,广西地质调查局的覃晓峰博士在火山岩分析方面给予了无私帮助,中国科学院地质与地球物理研究所的老师对火山岩进行了精心的分析测试,在此一并表示感谢!

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