NOTICE: this is the author's version of a work that was accepted for publication in Chemical Geology. Changes resulting from the publishing process, such as peer review, editing, corrections, structural formatting, and other quality control mechanisms may not be reflected in this document. Changes may have been made to this work since it was submitted for publication. A definitive version was subsequently published in Chemical Geology [277, 3-4, 2010] DOI 10.1016/j.chemgeo.2010.08.012 Ridge subduction and crustal growth in the Central Asian Orogenic Belt: Evidence from Late Carboniferous adakites and high-Mg diorites in the western Junggar region, northern Xinjiang (west China) Gongjian Tang^{a, b}, Qiang Wang^{a, c}, Derek A. Wyman^d, Zheng-Xiang Li^c, Zhen-Hua Zhao^a, Xiao-Hui Jia^{a, b}, Zi-Qi Jiang^{a, b} ^a Key Laboratory of Isotope Geochronology and Geochemistry, Guangzhou Institute of Geochemistry, Chinese Academy of Sciences, Guangzhou 510640, P. R. China ^b Graduate School of Chinese Academy of Sciences, Beijing 100049, China ^c The Institute for Geoscience Research (TIGeR), Department of Applied Geology, Curtin University of Technology, GPO Box U1987, Perth, WA 6845, Australia ^d School of Geosciences, Division of Geology and Geophysics, The University of Sydney, NSW 2006, Australia

***Corresponding authors**

25 E-mail address: wqiang@gig.ac.cn (Q. Wang)

26 ABSTRACT: The Central Asian Orogenic Belt (CAOB) is a natural laboratory for the study of accretionary tectonics and crustal growth owing to its massive generation of 27 juvenile crust in the Paleozoic. There is a debate, however, on the mechanism of this 28 29 growth. In the Baogutu area of the western Junggar region, northern Xinjiang (west China), diorite-granodiorite porphyry plutons and dikes are widely associated with Cu-Au 30 31 mineralization. In this study, we present new results of zircon U-Pb geochronology, major and trace elements, and Sr-Nd-Pb-Hf isotope analyses for two diorite-granodiorite 32 porphyry plutons and two dikes from this area. LA-ICPMS zircon U-Pb analyses of four 33 34 plutonic and dike samples yield Late Carboniferous ages of 315 -310 Ma. The Baogutu diorite-granodiorite porphyries exhibit low-Fe and calc-alkaline compositions. They are 35 36 also geochemically characterized by high Sr (346-841 ppm) contents, low Y (9.18-16.5 37 ppm) and Yb (0.95-1.60 ppm) contents, and relatively high Sr/Y (31-67) ratios, which are 38 similar to those of typical adakites. In addition, some samples have relatively high MgO (2.35-8.32 wt.%) and Mg[#] (48-75) values, and Cr (22.7-291 ppm) and Ni (32.0-132 ppm) 39 40 contents, which are similar to those of high-Mg andesites. All rock samples exhibit mid-oceanic ridge basalt (MORB)-like Nd-Sr-Pb-Hf isotope features: high $\varepsilon_{Nd}(t)$ 41 (+5.8+8.3) and $\varepsilon_{Hf}(t)$ (+13.1+15.7) values, and relatively low $({}^{87}Sr)^{86}Sr)_i$ (0.7033 to 42 0.7054) and (²⁰⁶Pb/²⁰⁴Pb)_i (17.842-18.055). The Baogutu adakitic rocks also contain 43 reversely zoned clinopyroxene phenocrysts, which have low MgO cores and relatively 44 45 high MgO rims. Geochemical modeling indicates that the Baogutu adakitic rocks could have been derived by mixing ~95% altered oceanic crust-derived melts with ~5% 46 sediment-derived melts. Taking into account the regional geology, I- and A-type 47 granitoids and Cu-Au mineralization, and the presence of Carboniferous ophiolite 48 mdanges in northern Xinjiang, we suggest that the Baogutu adakitic rocks were most 49 probably generated by partial melting of a slab edge close to a subducting spreading ridge 50

in the Late Carboniferous. Ridge subduction and the resultant slab window probably caused strong extension in the overlying lithosphere, extensive melting of subducting oceanic crust, mantle and juvenile lower crust, and interaction between slab-derived melts and the mantle. Thus, events associated with ridge subduction are likely to have played an important role in crustal growth in the CAOB in addition to previously recognized accretion of subduction and arc complexes and post-collisional crustal melting.

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58 **Key words:** Ridge subduction; slab window; adakite; crustal growth; Central Asian

- 59 Orogenic Belt; western Junggar; Xinjiang
- 60

61 **1. Introduction**

A common tectonic feature in accretionary orogens is ridge subduction accompanied by ridge-trench interaction (Windley et al., 2007). Ridge subduction has been documented at a number of places along the modern Pacific Rim (Sisson et al., 2003; McCrory and Wilson, 2009). However, only a few cases have been well-documented in the pre-Cenozoic geologic record, indicating that this process is grossly underrepresented in tectonic syntheses of plate margins in the ancient geologic record (Sisson et al., 2003).

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Ridge subduction and ridge-trench interaction may impact strongly on magmatic activity, metamorphism and mineralization near convergent plate margins (Cole and Basu, 1992; Haeussler et al., 1995; Sisson et al., 2003; Chadwick et al., 2009; McCrory and Wilson, 2009). When a ridge intersects with the subduction zone, a "slab window" may form between the subducted parts of the diverging oceanic plates (Dickinson and Snyder, 1979, Thorkelson, 1996). The "Blowtorch effect" (Delong et al., 1979) resulting from the upwelling of asthenospheric mantle through the slab window can produce a wide variety 76 of magmas. Typical magmatic rocks resulting from the subduction of slab windows 77 include adakites, tholeiite, high-Mg andesites (e.g., Hole et al., 1991; Abratis and Worner, 2001; Rogers et al., 1985; Guivel et al., 1999; Breitsprecher et al., 2003). The upwelling 78 79 of asthenospheric mantle through the slab window not only provides high heat flow that can induce partial melting of the slab edge, overlying mantle wedge and/or upwelling 80 81 asthenospheric mantle and crustal rocks (e.g., Yogodzinski et al., 2001; Thorkelson and 82 Breitsprecher, 2005), but also causes an extensional stress field above the slab window, 83 leading to the formation of alkaline magmatic rocks or A-type granites (Mortimer et al., 84 2006; Hung et al., 2007; Anma et al., 2009).

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The Central Asian Orogenic Belt (CAOB) (Fig. 1a) is one of the largest orogens in the 86 87 world and comprises island arcs, seamounts, accretionary wedges, oceanic plateaus and, 88 possibly, microcontinents accreted during the closure of the Paleo-Asian Ocean (Sengör et al., 1993; Jahn et al., 2000, 2004; Xiao et al., 2008; Windley et al., 2007). There is 89 90 evidence for possible ridge subduction events resulting from the Paleo-Asian Ocean 91 closure. Windley et al. (2007) suggested that the CAOB contains many key features (e.g., 92 adakites, boninites, near-trench magmatism, Alaskan-type mafic-ultramafic complexes and high-temperature metamorphic belts) that are explicable by ridge-trench interactions 93 94 and that this new perspective may provide a promising approach for resolving many 95 aspects of the orogenic belt's evolution. Based on data for Paleozoic ophiolites, tectonics and magmatism, several possible cases of ridge subduction have been proposed in the 96 CAOB. Windley et al. (2007) focused mainly on the broad diagnostic features of 97 98 ridge-trench interaction in the CAOB and suggested that gold deposits in eastern Tianshan may be related to slab window subduction. Based on geochronological data for ophiolites, 99 100 Jian et al. (2008) suggested that a ca. 430-415 Ma ridge subduction event was recorded in

Inner Mongolia in the southeastern CAOB. Sun et al. (2009) suggested Early Paleozoic ridge subduction in the Altai area, CAOB. Geng et al. (2009) and Yin et al. (2010) proposed that a ridge subduction model could account for the geochemical characteristics of granitoids and coeval mafic rocks from the western Junggar region. Here, we further document such a case in the western Junggar region based on Cu-Au mineralization-related adakites as well as regional geology and the presence of I- and A-type granitoids.

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109 The mechanism of Paleozoic crustal growth in the CAOB has been the subject of dispute (e.g., Seng ör et al., 1993; Gao et al., 1998; Jahn et al., 2000, 2004; Zhou et al., 2004; Liu 110 111 and Fei, 2006). In the western Junggar region of northwestern Xinjiang, there are 112 widespread Late Paleozoic magmatic rocks, typically I or A-type granite batholiths with 113 highly depleted isotopic signatures ($\varepsilon_{Nd}(t)$ of +6.4 to +9.2) (Chen and Arakawa, 2005; Han et al., 2006; Su et al., 2006). They have been attributed to either subduction-related 114 115 sources in an island arc setting (Zhang et al., 2006; Xiao et al., 2008) or to depleted mantle contributions in a post-collisional extensional setting (Chen and Arakawa, 2005; 116 117 Han et al., 2006; Su et al., 2006). These models clearly have significantly different implications for both granite petrogenesis and crustal growth. 118

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In this study, we report on late Carboniferous adakitic and high-Mg diorite-granodiorite porphyry plutons and dikes associated with Cu-Au mineralization in the Baogutu area of the western Junggar region (Fig. 1b and c). On the basis of these results and previous regional studies, we suggest that ridge subduction played an important role in the generation of the large late Carboniferous-Early Permian magmatic suite, contemporary Cu-Au mineralization, and crustal growth in the western Junggar region of the CAOB.

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127 **2. Geological backgroud**

The CAOB, also known as the Altaid Tectonic Collage (Seng ör et al., 1993; Jahn et al., 128 129 2000, 2004; Windley et al., 2007), extends from the Urals in the west, through Kazakhstan, northern China, Mongolia, and southern Siberia to the Okhotsk Sea along the 130 131 eastern Russian coast (Fig. 1a) (Sengör et al., 1993; Xiao et al., 2004). It is located 132 between the Siberian Craton to the north and the North China and Tarim Cratons to the 133 south (Fig. 1a). The CAOB was mainly formed by the progressive subduction of the 134 Paleo-Asian Ocean and the amalgamation of terranes of diverse origins (Coleman, 1989; Xiao et al., 2004; Windley et al., 2007). The most outstanding feature of the CAOB is the 135 136 vast expanse of granitoids and volcanic rocks, which are characterized by positive $\varepsilon_{Nd}(t)$ 137 and young T_{DM} model ages (e.g., Jahn et al., 2000; Wu et al., 2000).

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The western Junggar region is surrounded by the Altai orogen to the north, the Tianshan 139 140 orogen to the south, the Kazakhstan plate to the west, and the Junggar basin to the east (Fig. 1a). No metamorphic basement has been documented in this area and the oldest 141 rocks are ophiolitic mafic-ultramafic types of Cambrian-Ordovician age (Fig. 1b). 142 Post-Cambrian (Ordovician-Quaternary) sedimentary rocks, particularly Devonian and 143 144 Carboniferous volcanic-sedimentary rocks, are abundant in the region (Fig. 1b). The 145 Devonian strata are mainly distributed in the northwest and the Carboniferous strata are mainly in the southeast; both contain extensive volcanic rocks (Fig. 1b). The 146 Carboniferous strata are principally composed of the Tailegula Formation, the Baogutu 147 148 Formation, and Xibeikulasi Formation, in an upward sequence (Fig. 1c) (Shen and Jin, 1993). Both Jin et al. (1987) and Song et al. (1996) reported Late Carboniferous 149 150 deep-water sedimentary rocks close to the Baogutu area in the eastern part of the western 151 Junggar region (Fig.1b), and suggested that a deep-marine basin persisted there until that time. Based on paleogeographic data, Wang (2006) also suggested a late Carboniferous 152 deep-marine environment in that region, in contrast to a contemporary shallow-marine 153 154 environment to the west and northwest (Fig. 1d-f). During the early Carboniferous, however, a deep marine environment is believed to have existed across the entire region 155 (Fig. 1d-f) (Wang, 2006). These observations suggest that the western Junggar might have 156 157 been a growing accretionary prism during the Carboniferous, with a northwest (in present coordinates) directed subduction system (Fig. 1d-f). 158

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Many ophiolitic mafic-ultramafic rocks occur in the western Junggar region, and their ages range from the Cambrian to late Carboniferous (Fig. 1b) (Beijing SHRIMP Unit, 2005; Xu et al., 2006a). Recently discovered ophiolites located to the northeast of Keramay city have a SHRIMP zircon U-Pb age of 332 ± 14 Ma and are the youngest known in the area (Xu et al., 2006a; Xiao et al., 2008).

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166 Carboniferous to Early Permian magmatic rocks occur widely in the western Junggar 167 region and are referred to here as the Keramay arc magmatic rocks (Fig. 1b) (see 168 discussion in part 6). Carboniferous to early Permian granitoid intrusions show systematic 169 changes in both compositions and ages from north to south (Fig. 1b).

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In the north of the western Junggar region (Hebukesaier region) (Fig. 1b), the intrusions mainly consist of I-type granitoids, which have an age range of 338 Ma to 313 Ma (Han et al., 2006; Zhou et al., 2008). They are characterized by the enrichment of light rare earth elements (LREE) and depletion of high field strength elements (HFSE) (e.g., Nb, Ta and Ti) (Zhou et al., 2008).

In the central part of the magmatic province (i.e., the Keramay-Hatu-Tuoli area), large 177 batholiths and relatively small stocks intrude Paleozoic strata (Fig. 1b). These are 178 179 predominantly 315-300 Ma A-type intrusions and minor 300-290 Ma I-type granitoids (Fig. 1) (Chen and Jahn, 2004; Chen and Arakawa, 2005; Han et al., 2006; Su et al., 2006). 180 The I-type granitoids are characterized by enrichment of large ion lithophile elements 181 (LILE) and LREE, depletion of HFSE and relatively high $Mg^{\#}(100 \times Mg^{2+}/(Fe^{2+}+Mg^{2+}))$ 182 (41-63) (Chen and Jahn, 2004; Chen and Arakawa, 2005), i.e., geochemical 183 184 characteristics similar to typical arc magmatic rocks. The more voluminous A-type granites are characterized by high SiO₂ (71-75 wt.%), alkalis and Fe/(Fe+Mg) values, low 185 Al₂O₃ and CaO contents, enrichment in LILEs (Rb, Th and K) and HFSEs (Nb, Ta, Zr 186 187 and Hf), and obvious negative anomalies in Eu, Ba and Sr (Chen and Jahn, 2004; Chen and Arakawa, 2005; Su et al., 2006; Geng et al., 2009), similar to typical A-type granites 188 (Loiselle and Wones, 1979). 189

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191 In the southern part of the region (Tangbale-Mayle area), there are fewer scattered 192 intrusions, which mainly consist of quartz monzonite, biotite granite and diorite, similar to I-type granitoids (Fig. 1b). They are considered to have formed in the late Early 193 Carboniferous (Xinjiang Bureau of Geology and Mineral Resources (XBGMR), 1993). 194 195 Thus, granitoids in the central part of the area are slightly younger than those in the northern and southern areas. Almost all these granitoids exhibit high $\varepsilon_{Nd}(t)$ (+5.4 to +9.2) 196 but the central A-type granites have slightly higher $\varepsilon_{Nd}(t)$ (+6.4 to +8.4) (Chen and Jahn, 197 198 2004; Chen and Arakawa, 2005; Su et al., 2006).

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200 Volcanic rocks and dikes are also abundant in the central part of the magmatic province

(Fig. 1b). The volcanic rocks consist of andesitic basalt, andesite, felsic tuff and minor quartz or olivine tholeiite and occur as part of the Carboniferous Tailegula Formation (Shen and Jin, 1993). A U-Pb zircon SHRIMP age of 328 ± 2 Ma has been reported for felsic tuffs in the Baobei gold deposit from the western Junggar region (Wang and Zhu, 2007). Several hundred intermediate-basic dikes in this area mainly strike NW (280 $^{\circ}$ -300 $^{\circ}$, and cut both granitoid intrusions and Carboniferous strata (Li et al., 2004; Yin et al., 2010).

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209 The western Junggar region is one of the most important copper and gold producing areas in northwestern China (Shen and Jin, 1993; Rui et al., 2002; Zhu et al., 2007). Our study 210 211 area (Baogutu) is located in the centre of the western Junggar region, and is 40 km 212 southeast of the Keramay city (Fig. 1). Porphyry Cu-Au deposits and orogenic (vein) Au 213 deposits in the area (Fig. 1b-c) are hosted in about twenty small plutons and many NW and NE-trending dikes, including the Wudehe No. 2 (outcrop area of $\sim 3 \text{ km}^2$) and No. 5 214 (outcrop area of $\sim 0.84 \text{ km}^2$) plutons in the north, and the Kuogeshaye dikes in the south 215 (Fig. 1c). The plutons intrude the Xibeikulasi and Baogutu formations, which consist 216 217 mainly of tuffaceous breccias and siltstone (Fig. 1c). The Wudehe No. 5 porphyry copper deposit, containing about 1.115×10^6 ton resource averaging 0.28% Cu, >0.01% Mo, and 218 0.25 ppm Au (Shen et al., 2009), is the second largest porphyry copper deposit in 219 220 Xinjiang following the Tuwu-Yandong deposit in eastern Tianshan. The Wudehe Cu (Cu-Mo) mineralization is generally hosted in, or occurs around, small plutons (Fig. 1b). 221 The eastern Kuogeshaye orogenic gold deposit (Rui et al., 2002) is the largest gold 222 223 deposit in the western Junggar area, with an estimated total reserve of 4.2 tons. The eastern and western Kuogeshave gold deposits are located in the southern part of the 224 Baogutu area (Fig. 1b), with lensoid or ribbon-like gold-bearing veins generally hosted in 225

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wall rocks (tuff and tuffaceous siltstone) close to the dioritic porphyrite dikes (Fig. 1b).

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228 **3. Petrography**

The Wudehe plutons mainly consist of quartz diorite and granodiorite or corresponding 229 porphyries whereas the Kuogeshaye dikes are mainly dioritic porphyrite. The intrusive 230 rocks in the Wudehe No. 2 pluton exhibit fine-medium and medium-coarse-grained 231 inequigranular or porphyry texture and mainly contain plagioclase (30-40 vol. %), 232 hornblende (15-20 vol. %), biotite (5-15 vol. %) and quartz (<5 vol. %). Accessory 233 234 minerals include pyroxene, magnetite, titanite, zircon and apatite. Plagioclase, up to 3mm 235 in size, is the most abundant phenocryst phase in all of the Wudehe No.2 intrusive rocks. They are generally euhedral and lath-shaped, and some of them show polysynthetic 236 237 twinning. Hornblende occurs as euhedral phenocrysts with 300-600 μ m in size, but some crystals are altered. Biotite phenocrysts, 100-500 µm in size, are transparent brown and 238 239 subhedral. The intrusive rocks contain minor quartz, which has a round shape caused by resorption. Pyroxene is rare in the intrusive rocks. Fe-Ti oxides appear as 240 241 microphenocrysts and microlites in the matrix, or as inclusions in other phases (e.g., 242 hornblende).

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The porphyries in the Wudehe No.5 pluton contain phenocrysts up to 2 mm. They contain plagioclase (30-45 vol. %), hornblende (15-20 vol. %), biotite (10-15 vol. %), pyroxene (3-6 vol. %), quartz (<5 vol. %), and minor titanite, rutile, apatite, magnetite, and zircon. Plagioclase phenocrysts are generally euhedral with polysynthetic twinning. Horblende commonly occurs as transparent euhedral phenocrysts. Brown biotite occurs as euhedral phenocrysts. Some biotite granules are altered to chlorites and epidote. Pyroxene phenocrysts are sometimes altered to hornblende along grain boundaries.

The Kuogeshaye dioritic porphyrites consist of plagioclase (15-30 vol. %), hornblende (10-20 vol. %), biotite (5-15 vol. %), pyroxene (5-15 vol. %), and accessory minerals similar to those in the Wudehe intrusive rocks. Pyroxene phenocrysts in the Kuogeshaye dioritic porphyries are abundant relative to those of the Wudehe intrusive rocks. They occur as transparent and euhedral phenocrysts that have diameters of 50-300 μm.

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4. Analytical methods

Zircons were separated using conventional heavy liquid and magnetic separation 259 260 techniques. Cathodoluminescence (CL) images were obtained for zircons prior to analysis, using a JEOL JXA-8100 Superprobe at the Guangzhou Institute of Geochemistry, 261 Chinese Academy of Sciences (GIGCAS), in order to characterize internal structures and 262 263 choose potential target sites for U-Pb dating. LA-ICP-MS zircon U-Pb analyses were conducted on an Agilent 7500 ICP-MS equipped with a 193-nm laser, housed at the 264 National Key Laboratory of Geological Processes and Mineral Resources, Faculty of 265 266 Earth Sciences, China University of Geosciences (Wuhan). Zircon 91500 was used as the standard (Wiedenbeck et al., 1995) and the standard silicate glass NIST 610 was used to 267 optimize the machine, with a beam diameter of 30µm. Raw count rates for ²⁹Si, ²⁰⁴Pb, 268 ²⁰⁶Pb, ²⁰⁷Pb, ²⁰⁸Pb, ²³²Th and ²³⁸U were collected and U, Th and Pb concentrations were 269 calibrated using ²⁹Si as the internal calibrant and NIST 610 as the reference material. 270 ²⁰⁷Pb/²⁰⁶Pb and ²⁰⁶Pb/²³⁸U ratios were calculated using the GLITTER program (Jackson et 271 al., 2004). Measured ²⁰⁷Pb/²⁰⁶Pb, ²⁰⁶Pb/²³⁸U and ²⁰⁸Pb/²³²Th ratios in zircon 91500 were 272 averaged over the course of the analytical session and used to calculate correction factors. 273 274 These correction factors were then applied to each sample to correct for both instrumental mass bias and depth-dependent elemental and isotopic fractionation. Common Pb was 275 corrected by ComPbCorr#3 151 (Andersen, 2002) for those with common 206 Pb > 1%. 276

Further detailed descriptions of the instrumentation and analytical procedure for the LA-ICP-MS zircon U-Pb technique can be found in Gao et al. et al. (2002) and Liu et al. (2008, 2009). Uncertainties in the ages listed in Appendix 1 are cited as 1σ , and the weighted mean ages are quoted at the 95% confidence level. The age calculations and concordia plots were made using Isoplot (ver 3.0) (Ludwig, 2003). LA-ICP-MS U–Pb zircon data are presented in Appendix 1.

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Major element analysis and back-scattered-electron imaging for clinopyroxene and plagioclase were carried out at Guangzhou Institute of Geochemistry, Chinese Academy of Sciences (GIG-CAS) using a JXA-8100 electron microprobe. An accelerating voltage of 15 kV, a specimen current of 3.0×10^{-8} A, and a beam size of 1-2 µm were employed. The analytical errors are generally less than 2%. The analytical procedures were described in detail in Huang et al. (2007). The results are listed in Appendix 2.

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Rock samples were examined by optical microscopy and unaltered or least-altered samples were selected for geochemical analysis. Major element oxides were determined using the standard X-ray fluorescence (XRF) method (Li et al., 2006). Trace elements were analyzed by inductively coupled plasma mass spectrometry (ICP-MS), using a Perkin-Elmer Sciex ELAN 6000 instrument at GIGCAS. Analytical procedures are the same as those described by Li et al. (2006). Analytical precision for most elements is better than 3%. The results are listed in Appendix 3.

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Sr and Nd isotopic analyses were performed on a Micromass Isoprobe multi-collector ICPMS at the GIGCAS, using analytical procedures described by Li et al. (2006). Sr and REE were separated using cation columns, and Nd fractions were further separated by

HDEHP-coated Kef columns. Measured ⁸⁷Sr/⁸⁶Sr and ¹⁴³Nd/¹⁴⁴Nd ratios were normalized to ⁸⁶Sr/⁸⁸Sr= 0.1194 and ¹⁴⁶Nd/¹⁴⁴Nd=0.7219, respectively. The reported ⁸⁷Sr/⁸⁶Sr and ¹⁴³Nd/¹⁴⁴Nd ratios were respectively adjusted to the NBS SRM 987 standard ⁸⁷Sr/⁸⁶Sr=0.71025 and the Shin Etsu JNdi-1 standard ¹⁴³Nd/¹⁴⁴Nd=0.512115.

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For Pb isotopic determination, about 100 mg powder was weighed into the Teflon beaker, spiked and dissolved in concentrated HF at 180 °C for 7 h. Lead was separated and purified by conventional cation-exchange technique (AG1×8, 200–400 resin) with diluted HBr as an eluant. Total procedural blanks were less than 50pg Pb. Isotopic ratios were measured using a VG-354 mass-spectrometer at the GIGCAS following procedures described by Zhu et al. (2001). Repeated analyses of SRM 981 yielded average values of 206 Pb/²⁰⁴Pb = 16.9 ±4(2 σ), 207 Pb/²⁰⁴Pb = 15.498 ±4(2 σ) and 208 Pb/²⁰⁴Pb = 36.728 ±9(2 σ).

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Hf isotopic analyses were conducted using a multicollector Thermo Electron Neptune 315 MC-ICP-MS system in the Institute of Geology and Geophysics, Chinese Academy of 316 Sciences (Beijing). The analytical methods are similar to those of Li et al. (2006). 317 ¹⁷⁶Hf/¹⁷⁷Hf measurements were normalized to ¹⁷⁹Hf/¹⁷⁷Hf=0.7325. During the period of 318 data acquisition, standard BCR-2 was also processed for Hf isotopes, which gave a ratio 319 of 0.282881 ± 8 (2 σ m) for 176 Hf/ 177 Hf, in agreement with the recommended value 320 (Bizzarro et al., 2003). The analyzed results, with the calculated initial isotopic values and 321 model ages, are listed in Appendix 4. 322

323

324 **5. Results**

325 **5.1 Zircon geochronology**

326 To determine the emplacement ages of the ore-related porphyries, 4 representative

samples were chosen for LA-ICPMS zircon U-Pb dating, one each from the Wudehe No.
2 and No. 5 plutons, and the eastern and western Kuogeshaye dikes. Zircons have a size
range of 30–150 µm with a length/width ratio of 1:1–3:1. Cathodoluminescence images
of zircon grains used for LA-ICP-MS analysis show micro-scale oscillatory zoning (Fig.
2). These zircon grains also exhibit high Th/U ratios (0.31–1.46), suggesting a magmatic
origin (Belousova et al., 2002). Tera-Wasserburg diagrams and representative CL images
of analyzed zircon are shown in Fig. 2, and the U-Pb age data are given in Appendix 1.

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Twenty-one analyses of zircons from the dominant rock type, quartz dioritic porphyry, were obtained for sample 06XJ145 from the Wudehe No. 2 pluton (84 27 10"N, 45 29 57"E) give a weighted mean 206 Pb/ 238 U age of 315 \pm 4 Ma (2 σ) (Mean square weighted deviation (MSWD) = 0.76) (Appendix 1; Fig. 2a). The remaining five analyzes give 206 Pb/ 238 U ages ranging from 361 Ma to 375 Ma, with a weighted mean 206 Pb/ 238 U age of 367 \pm 9 Ma (2 σ ; MSWD = 1.08) (Fig. 2a), interpreted as the age of inherited zircons.

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The twenty-two analyses of zircons from sample 06XJ147, a quartz dioritic porphyry typical of the Wudehe No. 5 pluton (Fig. 1c, 84 °32'28"N, 45 °28'27"E), result in a single age population with a weighted mean 206 Pb/ 238 U age of 311 ± 4 Ma (2 σ ; MSWD = 0.38) (Fig. 2b). This age is interpreted to be the best estimate of the crystallization age of the Wudehe No. 5 porphyritic intrusion.

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Seventeen of the twenty zircons analyses from the dioritic porphyry dike sample (06XJ-153) in the east Kuogeshaye area (Fig. 1c, 84 $^{2}6'_{25}$ "N, 45 $^{2}3'_{54}$ "E) give a weighted mean 206 Pb/ 238 U age of 314 ± 4 Ma (2 σ ; MSWD = 1.4) (Fig. 2c). This age is

352 interpreted to be the best estimate of the crystallization age of the east Kuogeshaye dike. Analysis number 4 gives the oldest 206 Pb/ 238 U age of ~761 Ma (Appendix 1). Moreover, 353 two other analyses give ²⁰⁶Pb/²³⁸U ages of 344 Ma and 364 Ma, respectively. The old 354 (~761-344 Ma) zircons were likely inherited or entrained from the wall rocks during dike 355 emplacement. Minor 761 Ma zircons were most probably derived from shallow-level 356 sedimentary wall rocks because no Archean-Proterozoic metamorphic rocks have been 357 found in the western Jungger and southern Altay areas (Hu et al., 2000; Sun et al., 2008; 358 2009). 359

360

All 30 analyses zircons from for the west Kuogeshaye dioritic porphyry (06XJ-156, 84 26'14"N, 45 23'24"E) share similar 206 Pb/ 238 U ratios and give a weighted mean 206 Pb/ 238 U age of 310 ± 3 Ma (2 σ ; (MSWD = 0.46) (Appendix 1; Fig. 2d). This age is regarded as the age of crystallization.

365

366 **5.2 Mineral compositions**

Clinopyroxenes from the Baogutu porphyries exhibit variable SiO_2 (49.4 – 53.3 wt.%), 367 Al₂O₃ (0.85-3.42 wt%) and MgO (13.9 -15.5 wt.%) contents and Mg[#] (69 – 85). Most 368 clinopyroxene crystals are augite and diopside (Fig. 3). A prominent feature of the 369 porphyries is the presence of simple reversely zoned clinopyroxene phenocrysts, which 370 have low MgO and Mg[#] cores and relatively high MgO and Mg[#] rims (Fig. 4; Appendix 371 2). The cores also have high FeO and Na₂O contents compared to the rims (Fig. 4e and f). 372 No multiply zoned clinopyroxene has been observed in the Baogutu samples. Plagioclase 373 phenocrysts occur as euhedral weakly zoned crystals, with An (anorthite) contents of 374 46-60 % in the core and An contents of 33-52 % in the rim. Plagioclases in the matrix 375 show a large variation in An contents (3-46 %) (Appendix 5). Large plagioclase crystals 376

377 may occasionally partly enclose or contain some clinopyroxene grains (Appendix 5),

indicating that the crystallization of clinopyroxene occurred prior to that of plagioclase.

379

380 **5.3 Major and trace elements**

All of the studied porphyries share similar major element contents. The samples all plot 381 within the calc-alkaline and low-Fe fields on a SiO₂ vs FeO_{total}/MgO diagram (Fig. 5a), 382 383 and conform to a medium-K calc-alkaline trend on a SiO₂ - K₂O diagram (Fig. 5b). Most Baogutu porphyry samples plot in the field of gabbroic diorites, diorites and granodiorites 384 385 (Fig. 5c) (Middlemost, 1994). Porphyries from the Wudehe plutons show a large range of SiO₂ contents from 50 to 70 wt.% and MgO contents from 0.47 wt.% to 8.70 wt.%, 386 although they are dominantly dioritic (or andesitic) (SiO₂ 53.59-61.49 wt.%; MgO 3.39 387 388 wt%-7.79 wt.%) (see Appendix 3; Fig. 5c). The Kuogeshaye dikes show smaller ranges 389 of SiO₂ (57.08 wt.%-67.88 wt.%) and MgO (1.48 wt%-4.14 w.t%) contents (see Appendix 3; Fig. 5c). The Wudehe plutons and the Kuogeshaye dikes have total alkali 390 391 (Na₂O+K₂O) contents ranging from 4.17 to 8.52 wt.% and Na₂O/K₂O ratios from 1.80 to 9.30 (see Appendix 3), indicating their sodium-rich compositions. 392

The Baogutu samples exhibit geochemical characteristics of typical adakites (Kay, 1978; 394 395 Defant and Drummond, 1990; Martin et al., 2005). They are characterized by fractionated 396 rare earth element (REE) patterns with La/Yb ratios (3.0 -17) higher than those of normal arc magmas (Fig. 6a), and negligible Eu and strongly positive Sr anomalies (Fig. 6b). 397 They also show Nb, Ta and Ti depletions, however, which are similar to the Keramay 398 399 I-type granitoids (Fig. 6b). They have high Al_2O_3 (14.98-18.32 wt.%) and Sr (346–841 ppm) contents (see Appendix 3). Except for sample 06XJ-143, all samples have low Y 400 and Yb contents (9.18–16.5 ppm and 0.95–1.60 ppm, respectively), and high Sr/Y ratios 401

402 ranging from 31 to 67, and plot in the field of typical "adakites" (Fig. 5d).

403

In addition, some gabbroic diorite and diorite samples with SiO₂ contents less than 64 404 wt.% are characterized by elevated MgO (2.35-8.32 wt.%) and Mg[#] (48-75) values (Fig. 405 5e-f), and higher Cr (22.7–291 ppm) and Ni (32.0–132 ppm) contents (Appendix 3). On 406 SiO₂ versus MgO and Mg[#] diagrams (Fig. 5e and f), they partially overlap with high-Mg 407 andesites from the Setouchi Volcanic Belt, Japan (Shimoda et al., 1998; Tatsumi et al., 408 2006; Tatsumi, 2006, and reference therein). The Setouchi high-Mg andesites as a whole, 409 however, are more magnesian and have significantly higher Mg[#] values than the Baogutu 410 411 intrusive rocks.

412

413 **5.4 Nd-Sr-Pb-Hf isotope compositions**

The Baogutu samples have isotopic features broadly similar to those of MORB. Both 414 their initial 87 Sr/ 86 Sr isotopic ratios (0.7033 to 0.7054) and $\epsilon_{Nd}(t)$ values (+5.8 - +8.3, 415 416 average + 6.7) are comparable to those of the East Pacific Rise (EPR) basalts (Fig. 7a; Appendix 4). The Baogutu samples have slightly lower $\varepsilon_{Nd}(t)$ values than the Early 417 418 Carboniferous (345 Ma) volcanic rocks in the Junggar Basin and late Carboniferous-Early Permian (315 - 290 Ma) granitoids in the western Junggar region (Fig. 7a). The Baogutu 419 samples have high $\varepsilon_{\text{Hf}}(t)$ values (+13.5 to +15.7) and positive $\Delta \varepsilon_{\text{Hf}}(t)$ values (+0.9 to+4.3) 420 421 [where $\Delta \varepsilon_{\text{Hf}}(t) = \varepsilon_{\text{Hf}}(t) - (1.59\varepsilon_{\text{Nd}}(t) + 1.28)$] and plot close to or above the mantle array $[\varepsilon_{Hf}(t)=1.59\varepsilon_{Nd}(t) + 1.28]$ of Chauvel et al. (2008) in the $\varepsilon_{Hf}(t)$ versus $\varepsilon_{Nd}(t)$ diagram 422 (Appendix 4; Fig. 7b). In addition, they have variable Pb isotopic compositions 423 $(^{206}\text{Pb}/^{204}\text{Pb}_i = 17.842 - 18.055; \quad {}^{207}\text{Pb}/^{204}\text{Pb}_i = 15.411 - 15.466; \quad {}^{208}\text{Pb}/^{204}\text{Pb}_i = 16.056;$ 424 37.316–37.313) (Appendix 4). 425

427 **6. Discussion**

428 **6.1 Petrogenesis**

Adakites were originally considered to form by melting of subducted young and hot 429 430 oceanic crust (Mechanism A; Defant and Drummond, 1990). Later studies suggested that adakitic magmas could be produced by alternative mechanisms: (a) partial melting of 431 432 thickened basaltic lower crust (Mechanism B; Atherton and Petford, 1993; Rudnick, 1995; Petford and Atherton, 1996; Chung et al., 2003; Condie, 2005; Wang et al., 2005, 2007a); 433 (b) partial melting of delaminated lower crust (Mechanism C; Kay and Kay, 1993; 434 435 Rudnick, 1995; Xu et al., 2002b; Gao et al., 2004; Wang et al., 2006a; Huang et al., 2008); (c) partial melting of subducting continental crust (Mechanism D; Wang et al., 2008); (d) 436 assimilation-low-pressure fractional crystallization from parental basaltic magmas 437 438 (Mechanism E; Castillo et al., 1999; Li et al., 2009); (e) high-pressure crystallization 439 (involving garnet) of typical subduction-related magmas derived from melting of the mantle wedge (Mechanism F; Macpherson et al., 2006); (f) magma mixing, and combined 440 441 assimilation-fractional crystallization of felsic and basaltic magmas (Mechanism G; Streck et al., 2007). We consider these alternative processes below with specific reference 442 443 to the Baogutu adakitic porphyries.

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445 6.1.1 Mechanisms B-G

The geochemical characteristics of the Baogutu adakitic porphyries are inconsistent with partial melting of thickened or delaminated continental lower crust (Mechanisms B and G). Commonly, adakitic rocks derived by melting of thickened lower crust are characterized by relatively low MgO or Mg[#] values (Fig. 5e and f), which are similar to those of experimental melts from metabasalts and eclogites (Sen and Dunn, 1994; Rapp and Watson, 1995; Rapp et al., 1999). In contrast, the Baogutu adakitic rocks possess distinctly higher MgO contents and Mg[#] values than those of experimental melts from
metabasalts and eclogites (Fig. 5e and f) (e.g., Rapp et al., 1999).

454

455 Recently, Wang et al. (2008) proposed that adakitic rocks in some collisional orogenic belts (e.g., Tibet) could have been formed by partial melting of subducted continental 456 crust (Mechanism D). Adakitic rocks formed from subducted continental crust also 457 commonly have higher K_2O contents (> 3 wt.%) than those of the Baogutu adakitic rocks 458 (<3 wt.%) (Fig. 5b). Moreover, adakitic rocks formed in this way have low $\varepsilon_{Nd}(t)$ values 459 460 (generally below -3) (Wang et al., 2008), whereas the Baogutu adakitc rocks have $\varepsilon_{Nd}(t)$ values of between +5.8 and +8.3. In addition, in the case of the Baogutu adakitic rocks, 461 sedimentary data suggest that an oceanic setting persisted into the late Carboniferous in 462 463 the western Junggar region (Jin et al., 1987; Song et al., 1996; Wang, 2006), indicating 464 that no continental collision took place at that time.

465

466 The Baogutu adakitic rocks are also difficult to explain by low-pressure assimilation-fractional crystallization (AFC) from parental basaltic magmas (Mechanism 467 E). If AFC processes could account for the petrogenesis of the Baogutu adakitic rocks, 468 then the most probable candidate for a parental magma would be that of the 345 Ma 469 Kexia basalts with high $\varepsilon_{Nd}(t)$ (+6.8 – +9.6) in the western Junggar Basin (Zheng et al., 470 2007). If olivine and pyroxene fractionated from the Kexia basalts, then the derived 471 magma would show a clear decrease in Mg[#] values as well as MgO contents, but this does 472 473 not occur (Fig. 5e-f). In addition, fractionation of olivine and pyroxene is not consistent 474 with the depletion of HREE (e.g., Yb). These minerals are unable to incorporate HREE elements, which generally leads to concave-upward HREE in the chondrite-normalized 475 REE concentration patterns (Castillo et al., 1999). Low pressure hornblende fractionation 476

477 may drive magmas toward adakitic Sr/Y ratios (Castillo et al., 1999) but even the most Mg-rich samples in the Baogutu suite display this feature (e.g., MgO = 8.7 - 7.4; $SiO_2 =$ 478 51.7 - 54.0; Sr/Y = 23 - 39). The Sr/Y and La/Yb ratios can also not be associated with a 479 480 crystal fractionation assemblage involving plagioclase, given that Eu anomalies are 481 absent and Sr/Y remains high across the entire range of SiO₂ contents (Figs. 5d and 8b, d). Where differentiation of high-Mg adaktic rocks has been observed elsewhere, Sr/Y and 482 483 La/Yb have been reported to increase with higher MgO content (Danyushevsky et al., 484 2008), but these trends are not present for the Baogutu rocks (Fig. 8b). Moreover, most of 485 the Baogutu samples display lower $\varepsilon_{Nd}(t)$ than the Miaoergou and Keramy granite and 486 Kexi basalt samples, which is inconsistent with a crust assimilation model (Fig. 8e). The relatively high MgO rims of reversely zoned clinopyroxene phenocrysts (Fig. 4) also 487 488 argue against crustal contamination.

489

The Baogutu adakitic rocks are also unlikely to be the result of high-pressure 490 491 crystallization (Mechanism F). The high Sr/Y and Dy/Yb ratios can only be achieved by extensive fractionation of amphibole (30-85%) and garnet (15-20%) from the parental 492 493 basalt (Fig. 8a). However, this conflicts with experimental and natural evidence (Müntener et al., 2001), which suggests that amphibole and garnet only occur at higher 494 495 degrees of crystallization, and high-pressure fractionation of hydrous basalts would 496 essentially form pyroxene-rich lithologies. In addition, an important feature of 497 high-pressure crystallization is that some key ratios, such as Sr/Y and Dy/Yb, would increase with increasing SiO₂ contents (Macpherson et al., 2006), because garnet is 498 499 involved in crystallization. However, the data for the Baogutu adakitic rock do not show 500 such trends (Fig. 8b-c).

502 It is also unlikely that magma mixing of felsic and basaltic magmas (Mechanism G) can 503 account for the Baogutu adakitic rocks. Candidates for mantle-derived basaltic and crust-derived felsic end-members in the area are most plausibly represented by the ~ 345 504 505 Ma Kexia basalts (Zheng et al., 2007) and the ~ 290 Ma Miaoergou granites, which have high SiO₂ (71.39–78.81 wt.%) and low MgO contents (0.01–0.85 ppm) (Han et al., 2006; 506 Su et al., 2006). The Baogutu adakitic rocks have an average $\varepsilon_{Nd}(t)$ value of +6.7, which 507 is slightly lower than those of the Kexia basalts (average value = +7.9) and the Miaoergou 508 granites (average value = +7.2) (Fig. 8e), suggesting that the adakitic rocks are not 509 510 products of mixing between the felsic and basaltic end members. In addition, adakitic rocks formed by magma mixing generally have relatively uniform compositions with 511 512 SiO₂ contents that span a narrow range (Streck et al., 2007). In contrast, the Baogutu 513 adakitic rocks display a wide range of SiO₂ contents from 52.46 to 70.65 wt.% (Appendix 514 3). Magma mixing between felsic and basaltic magmas is also inconsistent with the petrographic evidence. Such a process should commonly be accompanied by multiply 515 516 zoned phenocrysts (Troll and Schmincke, 2002), however, the clinopyroxenes from the Baogutu samples only display simple reverse zoning (Fig. 4). The normally zoned 517 518 plagioclase (Appendix 5) also contrasts with the complex zoning caused by magma mixing (Troll and Schmincke, 2002). Although mafic microgranular enclaves (MMEs) 519 520 were found in the Keramay plutons (Chen and Arakawa, 2005), no MMEs were 521 discovered in the Baogutu adakitic rocks, implying that magma mixing between mantle-522 and crust-derived magmas is unlikely to have played an important role in the genesis of the Baogutu samples. 523

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525 6.1.2 Partial melting of subducted oceanic crust (Mechanism A)

526 We suggest that the Baogutu adakitic rocks were generated by partial melting of

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subducted oceanic crust based on both geological and geochemical evidence, as follows:

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(1) The Baogutu adakitic rocks are geochemically similar to slab-derived adakites. They 529 530 are medium-K calc-alkaline and have relatively low K₂O contents (Fig. 5b), similar to slab-derived adakites in Pacific Cenozoic arcs (Stern and Kilian, 1996) and the North 531 Tianshan ranges (Wang et al., 2006b, 2007b) formed by partial melting of young and hot 532 subducted oceanic crust. They also have high MgO, Mg[#], Cr and Ni values, similar to 533 metabasaltic and eclogite melts hybridized by assimilation of 10-20% peridotite (Fig. 534 535 5e-f). Moreover, on the discrimination diagrams for low-SiO₂-adakites (LSA) and high-SiO₂-adakites (HSA) of Martin et al. (2005) (Fig. 9), the Baogutu adakitic rock 536 samples mainly plot in the field of the HSA derived from the interaction between 537 538 slab-derived melts and mantle peridotites. Therefore, the Baogutu adakitic rocks likely 539 resulted from interactions between mantle peridotite and slab melts during their ascent (e.g., Rapp et al., 1999; Martin et al., 2005). 540

541

(2) There is growing evidence for the existence and subduction of a Carboniferous ocean 542 in the Junggar area. Huang et al. (1990) proposed that a "Carboniferous Asian Ocean" or 543 "North Tianshan Ocean" existed in the south Junggar region in the Carboniferous. Xiao et 544 545 al. (1992) also suggested that a Carboniferous ocean existed here, which they named the 546 "North Tianshan Ocean". Recently, several Carboniferous ophiolites have been confirmed in the area, such as the Keramay ophiolites in which gabbro samples yield SHRIMP 547 zircon ages of 332 \pm 14 Ma (Xu et al., 2006a). The Darbut ophiolite formed at 346–347 548 549 Ma as dated by the SHRIMP zircon U-Pb method (Beijing SHRIMP Unit 2005 annual report). The Bayingou ophiolites, from which a gabbro sample yielded a zircon U-Pb age 550 of 344.0 \pm 3.4 Ma (Xu et al., 2006b), developed in response to the opening of the 551

Carboniferous Ocean (Huang et al., 1990). Wang (2006) also argued that there was a Carboniferous shallow to deep ocean setting in the western Junggar region, based on a sedimentary and paleogeographic analysis. In addition, recently indentified Carboniferous adakite, high-Mg andesite and Nb-enriched basalt suites in northern Tianshan (Wang et al., 2006b; 2007b) suggest subduction of Carboniferous oceanic crust along the southern margin of the Junggar Basin.

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(3) The occurrence of the Early Carboniferous Darbut, Keramay and Bayingou ophiolites 559 560 (Fig. 1b) suggests the presence of young oceanic crust, the subduction of which could produce the 315-310 Ma Baogutu adakitic rocks. The Nd-Sr isotopic compositions of the 561 Baogutu adakitic rocks are similar to those of the basaltic rocks from the Bayingou 562 563 ophiolites (Xu et al., 2006c), and also overlap those of the slab-derived adakites resulting from subduction associated with the "Carboniferous North Tianshan ocean" in the North 564 Tianshan ranges (Wang et al., 2006b, 2007b). In addition, the Baogutu adakitic rocks 565 566 display a limited $\varepsilon_{Nd}(t)$ - $\varepsilon_{Hf}(t)$ field overlapping with those of typical MORB (Fig. 7b).

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(4) Reverse zoning in clinopyroxene phenocrysts in the Baogutu adakitic rocks (Fig. 4) reflects a significant MgO or Mg[#] increase and Na₂O content decrease in the melt, which is consistent with reaction between the melt and mantle peridotite (e.g., Yogodzinski and Kelemen, 1998; Gao et al., 2004, 2008; Xiong et al., 2006). Therefore, the Baogutu adakitic rocks are most probably derived by the interaction between ascending subducted oceanic crust-derived adakitic melts and mantle wedge peridotites, based on the petrographic evidence.

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576 (5) It is well documented that subduction zone magmatism is controlled by contributions

from the subducted mafic oceanic crust, the overlying subducted sediments, and the mantle wedge (e.g., Kay, 1978; Defant and Drummond, 1990; Plank and Langmuir, 1993; Hawkesworth et al., 1997; Chauvel et al., 2008). Adakites are generally produced by partial melting of subducted mafic ocean crust, followed by the interaction of this melt with the mantle (Kay, 1978; Defant and Drummond, 1990). If subducted basaltic ocean crust begins to melt, however, then subducted sediments must also undergo partial melting (Kelemen et al., 2003).

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In Fig. 10a, Th/Ce ratios are plotted against initial ¹⁴³Nd/¹⁴⁴Nd ratios for Baogutu adakitic 585 rocks. This diagram is sensitive to the addition of a sediment-derived melt because the 586 subducted sediments have low ¹⁴³Nd/¹⁴⁴Nd and high Th/Ce ratios relative to mantle 587 wedge (Hawkesworth et al., 1997). The Baogutu adakitic rocks show a broad negative 588 589 trend, which is consistent with the addition of a sediment melt. The Baogutu adakitic rocks have high Th/La ratios, although they do not show particularly high Th contents 590 591 (Fig. 10b). They are similar to the high-Mg andesites from the Setouchi Volcanic Belt, Japan (Fig. 10b), which were generated by the partial melting of subducting sediments, 592 593 and subsequent melt-mantle interaction (Shimoda et al., 1998; Tatsumi, 2001; Hanyu et al., 2006). Plots of Th/La versus Sm/La for arc volcanic rocks show mixing toward local 594 595 sediments (Plank, 2005) and it is evident in Fig. 10c that the Baogutu adakitic rock 596 compositions can be produced by a mixture of basalts from the Bayingou ophiolites and subducted sediments. 597

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To model partial melting of subducted oceanic crust and its overlying sediment, we used a Kexia basalt sample (Zheng et al., 2007) and a Bayingou ophiolite gabbro sample (Xu et al., 2006c) as the subducted Carboniferous Junggar Altered Ocean basaltic crust (AOC), 602 and the average Global Subducting Sediment (GLOSS: Plank and Langmuir, 1998) composition as the overlying sediments. Based on these starting materials, the Baogutu 603 samples can be modeled as 99:1 to 90:10 AOC melt:sediment melt mixture on Sr, Nd and 604 605 Pb isotopic variation diagrams (Fig. 11a-c). An average composition of the Baogutu rocks can be derived by mixing at a 95:5 ratio of AOC and sediment melts (large red circles in 606 Fig. 11a-c). Its primitive mantle-normalized trace element pattern can also be reproduced 607 608 by this simple mixing scenario, except for the fluid-mobile elements (Rb, and Ba) (Fig. 11d). This suggests that these highly soluble elements were lost in a fluid phase prior to 609 610 melting. Thus, the Baogutu adakitic rocks were most probably produced by partial 611 melting of subducted oceanic crust and a thin veneer of overlying sediments (Fig. 11), 612 and subsequent melt-mantle interaction (Fig. 5e-f).

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614 **6.2 Geodynamic processes**

Two competing viewpoints for the tectonic evolution of the western Junggar area stress either island arc or post-collisional processes (Chen and Arakawa, 2005; Han et al., 2006; Su et al., 2006; Zhang et al., 2006; Xiao et al., 2008). The most appropriate tectonic model for the western Junggar area must account for the diverse compositional characteristics of the voluminous Carboniferous to Early Permian magmatic rocks, particularly granitoids.

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622 6.2.1 Three types of granitoids in the central western Junggar area

The central western Junggar area hosts many Late Carboniferous to Early Permian granitoids (ca. 315 to 290 Ma) (Fig. 12a). Distinct from the dominantly I-type granitoids of the northern and southern areas (Fig. 1b), the central area contains a diverse range of intrusive rocks, which can be classified into three groups based on their geochemical and

627 geochronological features (Figs. 5-7, 12). From the southeast to the northwest of the central area, these are the Baogutu adakitic rock group (Group 1), the Keramay I-type 628 granitoid group (Group 2), and the Miaoergou A-type granite group (Group 3), 629 respectively (Fig. 1b). Group 3 granitoids include the Miaoergou, Akbastao, Hongshan 630 and Hatu plutons (Fig. 1b). The three groups exhibit systematic variations in terms of 631 ages and geochemical characteristics (Fig. 12). Their ages decrease slightly from Group 1 632 to 3, although Groups 1 and 2 overlap significantly (Fig. 12a). Mg[#] and Sr/Y values 633 clearly decrease from Group 1 to 3 (Fig. 12c-d), whereas zircon saturation temperatures 634 635 increase from southeast (Group 1) to northwest (Group 3; Fig. 12e). If the transition from high to low Sr/Y ratios relates to the presence of garnet and plagioclase in their respective 636 sources, and the variation in Mg[#]s reflects the proportion of mantle component involved 637 638 in their petrogenesis (e.g., Rapp et al., 1999), then the differing compositions suggest that granitoid source depths become shallower from Group 1 in the southeast to Group 3 in 639 the northwest. 640

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642 6.2.2 Ridge subduction and slab window model

During the early Carboniferous, the Paleo-Asia Ocean (the Carboniferous Junggar Ocean) 643 may have been subducting beneath the Keramay arc (Fig. 13a). During this process, 644 upwelling slab-derived fluids would have triggered partial melting of the mantle wedge 645 (Fig. 13a). These 338 - 316 Ma volcanic rocks and I-type granitoids with "island 646 arc-type" geochemical compositions likely originated from partial melting of 647 fluid-metasomatized mantle wedge (Fig. 13a). Based on several lines of evidence, 648 649 discussed below, we suggest that subduction of a spreading centre during the late Carboniferous resulted in a slab window, which led to the formation of 315 - 290 Ma 650 magmatism in the central area of the Keramay arc. 651

Ridge subduction

Ridge subduction can readily provide enough heat for partial melting of a young oceanic 653 crust (≤ 25 Ma) or old oceanic crust (> 25 Ma), generating adakitic rocks (Defant and 654 Drummond, 1990; Aguillon-Robles et al., 2001; Kelemen et al., 2003). Such melting 655 occurs because the resultant slab window permits the upwelling of hot asthenospheric 656 mantle (e.g, Kay et al., 1993; Abratis and Worner, 2001; Breitsprecher et al., 2003; 657 Kelemen et al., 2003; Windley et al., 2007). In this scenario, the Baogutu low Mg (Mg[#] < 658 48) adakitic rocks resulted from partial melting of a slab edge and high-Mg (Mg[#] > 48) 659 660 gabbroic diorites and diorites originated from subsequent interaction between slab melts and the mantle (Fig. 13b). 661

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663 The Baogutu samples generally have slightly lower Sr (346-841 ppm) contents, Sr/Y (31-67 ppm) values and higher HREE and Y contents relative to typical adakites derived 664 from melting of subducted oceanic crust in non-ridge subduction settings (Kay, 1978; 665 666 Defant and Drummond, 1990; Kay et al., 1993; Stern and Kilian, 1996) (Fig. 5d). This is similar to other adakites (e.g., Vizcaino Peninsula, Mexico) formed by ridge subduction 667 (Aguillon-Robles et al., 2001). In addition, the Baogutu samples also show relatively low 668 Nb/Ta ratios (10.7 to 18, with an average of 13.4). This feature, coupled with higher 669 Zr/Sm ratios (16.9 to 75.7 and average 40.2; Appendix 3), is similar to Archean TTGs 670 671 (Condie, 2005). These trace element patterns indicate that slab (AOC + sediments) melting took place at relatively shallow depths, but mainly with a garnet amphibolite 672 residuum (Fig. 11d). These depth and source requirements further suggest that special 673 674 circumstances, such as mantle upwelling by ridge subduction, provided the anomalous heat source for slab melting (e.g., Aguillon-Robles et al., 2001). Interestingly, the zircon 675 saturation temperatures (Watson and Harrison 1983) (642 - 778 ℃) for the Baogutu 676

adakites are distinctly lower than those (729 - 976 °C) of the Miaoergou A-type granites (Fig. 12e), but similar to those (671 - 726 °C) of the southern Costa Rica adakites formed by subduction of the Cocos Ridge (Abratis and Worner, 2001). The temperatures may reflect adakite formation by hydrous partial melting of slab edges saturated with water (Martin, 1999) and generation of A-type granites under H₂O-absent conditions.

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The Keramay I-type granitoids are characterized by relatively high MgO and Sr contents 683 (Fig. 5d-e), high Mg[#], MORB-like Nd-Sr isotopic compositions and young Nd model 684 ages (T_{2DM}) of 300–550 Ma (Figs. 5f, 7a and 12b), indicating that they were derived from 685 a juvenile source (Chen and Arakawa, 2005). Experimental melts of basaltic rocks 686 generally exhibit Mg[#] lower than 40, whereas the Mg[#] of the Keramay series are 687 significantly higher than 40 (Figs. 5f and 11c). Accordingly, the melts were not directly 688 derived by partial melting of basaltic (or equivalent) rocks. The Keramay series are also 689 unlikely to have been generated by direct melting of a mantle source, given that their 690 691 silica contents are as high as 70 wt.% (Fig. 5). Chen and Arakawa (2005) proposed that the parent magma to the Keramay rocks originated from a depleted lithospheric mantle 692 source that had been metasomatized by fluids released from a subducting slab. We 693 suggest that the asthenosphere ascending from the slab window as a result of ridge 694 subduction provided the heat for such partial melting (Fig. 13b). The mantle-derived 695 696 magma may then have evolved into the Keramay I-type granitoids by fractional crystallization (Chen and Arakawa, 2005). Thus, the "island-arc type" characteristics of 697 the Keramay I-type granitoids were probably inherited from their mantle sources (Fig. 698 13a-b). 699

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701 Geochemical and isotopic characteristics of the Miaoergou series A-type granites suggest

702 that they formed above a slab window (Fig. 13b). The granites have low initial Sr isotopic ratios and MgO contents and Mg[#] values, positive $\varepsilon_{Nd}(t)$ values and young Nd model ages 703 (T_{2DM}) (Figs. 5e-f, 7a and 12b-c). Their very high zircon saturation temperatures (729 – 704 705 976 °C) are characteristic of A-type granites (Skjerlie and Johnston 1993). Accordingly, it has been suggested that they were generated by partial melting of juvenile (oceanic?) 706 707 lower crust (Chen and Arakawa, 2005; Geng et al., 2009). We propose that such partial 708 melting occurred because of the upwelling asthenosphere through a slab window (Fig. 13b). In addition, the compositional variations of the Miaoergou series A-type granites 709 710 suggests minor fractional crystallization after the A-type granite magmas were generated. The A-type granites have pronounced negative Eu, Sr and Ba anomalies (Fig. 6), 711 712 indicating the possible fractionation of plagioclase or K-feldspar in the evolution of the 713 A-type granite magmas (Wu et al., 2002). The A-type granites have relatively high Yb (1.96-6.56 ppm) and Y (16.3-80.4 ppm) contents, precluding heavy rare earth 714 715 element-rich garnet in the residue and indicating partial melting under low pressure (< 15) 716 kbar) conditions (Patiño Douce, 2005). In summary, we suggest that they were most likely derived from partial melting of juvenile lower crust above a slab window at low 717 pressure (< 15 kbar), followed by minor fractional crystallization of plagioclase or 718 K-feldspar. 719

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There is additional evidence for ridge subduction in the region. First, charnockites occur as giant enclaves in the Miaoergou batholith, suggesting a hotter than normal geotherm at that time (Fig. 13b). Zhang et al. (2004) proposed that the charnockites were produced by partial melting of juvenile lower crust, heated by mantle upwelling. A slab window would facilitate this upwelling (Fig. 13b). Second, numerous NW-extending basic-intermediate

727 dikes intrude the granite batholiths and the Carboniferous strata. The dikes are only slightly younger than the rocks that they intrude (Fig. 1b). They display typical 728 "subduction zone" characteristics and may have formed by partial melting of 729 730 metasomatized lithosphere mantle in an extensional environment. Intrusion of mafic dikes 731 into the accretionary prism is a typical feature of ridge subduction and slab window 732 scenarios (Sisson et al., 2003; Windley et al., 2007), and the trend of these dikes may be 733 parallel to that of the subducted oceanic ridge (Fig. 13b). Third, other recent studies also 734 document Late Carboniferous adakitic diorites (Geng et al., 2009) and high Mg diorite (or 735 sanukitoid) dikes (Yin et al., 2009) in the west Junggar Basin outside of our study area and also suggest the possible subduction of oceanic ridge (Geng et al., 2009; Yin et al., 736 2009). Fourth, there are Carboniferous primitive tholeiitic basalts in the Hatu area with 737 738 typical N-MORB geochemical characteristics (Shen and Jin, 1993). Such magmas are a 739 common manifestation of spreading ridge subduction (Guivel et al., 1999), and the Hatu basalts were probably derived from partial melting of upwelling mid-ocean-ridge-like 740 741 depleted mantle above a slab window.

742

743 The rock suite in the Keramay arc area is similar to those known to be associated with ridge subduction and slab window formation elsewhere. For example, in the south-central 744 745 Alaska Range, broadly contemporaneous adakites, I-type granitoids, and A-type granite 746 suites are associated with ridge subduction and slab window formation (Cole et al., 2007; Hung et al., 2007). Adakites there (e.g., the Jack River pluton) were generated at $62.7 \pm$ 747 748 0.4 Ma (Cole et al., 2007), I type granitoids (e.g., the Composite plutons) at 67 - 69 Ma, and A-type granites (e.g, the McKinley pluton) at 51 \pm 1 Ma (Hung et al., 2007). 749 Moreover, these A-type granites are considered to have been generated through partial 750 751 melting of lower crust by upwelling asthenosphere above a slab window (Mortimer et al.,

2006; Hung et al., 2007; Anma et al., 2009). In this case, the A-type granites are ca.
10–20 Ma younger than the adakites, which is a similar relationship to that found in the
Keramay arc (Fig. 12a).

755

The short duration (315-290 Ma) of magmatism in the Keramay arc area is also 756 similar to that known to be associated with ridge subduction and slab window formation 757 758 elsewhere. Cole et al., (2006) suggested the Caribou Creek volcanic rocks from the southern Talkeetna Mountains were related to a slab window, which existed between 59 759 760 Ma and 35 Ma. Kinoshita (2002) also documented a similar period of magmatism (~ 20 Ma) above a slab window that formed in response to spreading ridge subduction beneath 761 762 southwest Japan. The Princeton Group in south-central British Columbia formed by 763 remelting of arc basalt above a slab window, and geochronological results indicate that 764 they were erupted between approximately 53 and 47 Ma (Ickert et al., 2009). Anma et al. (2009) demonstrated that the Taitao granites were related to the subduction of a short 765 766 segment of the Chile ridge spreading center that started ~ 6 Ma ago. Therefore, one of the most distinctive features of ridge-subduction is the short duration of magmatism (~20 767 m.y.) above the resulting slab windows, which is consistent with a ridge subduction 768 model for the 315-290 Ma magmatism in the Keramay arc area. 769

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In addition, some plutons in the western Junggar area are rounded to ellipse-shaped, indicating that they are likely to have undergone little or no deformation. Many ridge subduction-related plutons exhibit similar characteristics. For example, the McKinley Sequence and associated Plutons (62-50 Ma) in the Central Alaska Range, which are also related to ridge-subduction (Hung et al., 2007), show no deformation and are mostly rounded to ellipse-shaped. The Taitao granite pluton (southern Chile) formed due to

subduction of the Chile ridge and is undeformed and ellipse-shaped (Anma et. al., 2009).
In fact, in the ridge-subduction model, the upwelling of asthenosphere through a slab
window would cause widespread crustal extension (Wilson et al., 2005; Cole et al., 2006;
Cole and Stewart, 2009), generally resulting in undeformed plutons, in contrast to those
found in a normal (i.e., non-ridge) subduction setting.

782

783 **6.3 Implications for crustal growth in the CAOB**

784 The most remarkable feature of the three groups of magmatic rocks in the central area of the western Junggar region is the fact that they are characterized by positive $\varepsilon_{Nd}(t)$ values 785 (+5.4-+9.2) and very young Nd model ages (T_{2DM}) of 300–600 Ma (Fig. 7a and 12b), 786 787 which is also a common feature of the CAOB. These features indicate that the CAOB 788 crust is made up of young mantle derived material (Jahn et al., 2004). Two competing viewpoints of CAOB crust growth (arc and post-collision related) have frequently caused 789 dispute (Seng ör et al., 1993; Jahn et al., 2000, 2004). Seng ör et al. (1993) hypothesized 790 791 that nearly half of the CAOB was derived from the mantle by successive accretion of arc 792 complexes and subduction accretion during the Palaeozoic. Conversely, many researchers 793 suggest that Phanerozoic CAOB granitoids were generated by basalt underplating in a post-collisional setting (Jahn et al., 2000; Wu et al., 2000). More recently, Jahn et al. 794 (2004) argued that both processes probably played equally important roles. 795

796

We suggest that ridge subduction and slab window formation also played an important role in the growth of CAOB crust. Figure 13 illustrates the most plausible geodynamic scenario: (1) Oceanic crust subduction and fluid release leading to mantle wedge partial melting and resultant 338 – 316 Ma volcanic rocks and I-type granites in an arc setting (Fig. 13a). (2) Direct partial melting of subducted basaltic slab (AOC+sediments), generating adakitic magmas (Process 1: see caption in Fig. 13b). (3) Melting of metasomatised mantle-wedge in the slab window to produce post-315 Ma I-type granitoids (Process 2 of the Fig. 13b). Process 3 in Fig. 13b also illustrates crustal growth via partial melting of the basaltic lower crust by underplating of mantle-derived magmas or direct melting of depleted mantle to produce A-type granites and N-MORB type basalts, respectively. These events were triggered by upwelling of asthenospheric mantle in a ridge subduction setting (Fig. 13b).

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810 Windley et al., (2007) proposed that diagnostic features of ridge subduction are abundant 811 throughout the geological record of the CAOB. Such processes probably accompanied the successive closure of branches of the Paleo-Asian Ocean, given that most subduction 812 813 systems eventually interact with a spreading ridge (Sisson et al., 2003). Although our study was confined to a relatively small area in the western part of the CAOB (Fig. 1), it 814 815 supports the view that ridge subduction was common during formation of the entire belt. If so, then ridge subduction likely made major contributions to crustal growth in the 816 817 CAOB in addition to those made by accretion of subduction and arc complexes (Seng är et 818 al., 1993) and post-collisional crustal melting (Jahn et al., 2000, 2004).

819

820 7. Conclusions

(1) The Baogutu porphyries in the western Junggar region are calc-alkaline quartz
dioritic and granodioritic plutons and dioritic porphyrite dikes. They exhibit
geochemical and petrologic characteristics that are typical of adakites; some of them
also have the geochemical characteristics of high-Mg andesites.

(2) LA-ICPMS zircon U-Pb dating suggests that the Baogutu plutons and dikes have
similar crystallization ages of 315–311 Ma and 314–310 Ma, respectively, and were
generated in the late Carboniferous.

(3) The Baogutu adakitic rocks were most likely generated by partial melting of a slab
edge (containing ca. 95% basaltic oceanic crust and ca. 5% overlying sediments) close
to a subducting spreading ridge in the garnet amphibolite faces as a result of a ridge
subduction, and subsequent interactions between mantle peridotite and slab melts
during their ascent.

(4) Events associated with ridge subduction are likely to have played an important role in
crustal growth in the CAOB in addition to previously recognized accretion of
subduction and arc complexes and post-collisional crustal melting.

836

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1252 Figures



Fig. 1. (a) Simplified tectonic divisions of the CAOB (after Jahn et al., 2000). (b) Geological map of the western Junggar region (modified after XBGMR, 1993). (c) Simplified geological map of the Baogutu deposits (after Cheng and Zhang, 2006). WDH-Wudehe; E-KGSY-Eastern Kuogeshaye; W-KGSY-Western Kuogeshaye. Age

1258 data for mafic-ultramafic or ophiolitic rocks are from Beijing SHRIMP Unit (2005), Xu et al. (2006a) and Xiao et al. (2008). Age data for granite intrusions and volcanic rocks are 1259 from Han et al. (2006), Su et al. (2006), Wang and Zhu (2007), (Zhou et al., 2008), Geng 1260 1261 et al. (2009) and this study. Age data for Baogutu diorite-granodiorite porphyry plutons and dikes are from this study. (d-f) Paleogeographic maps of the western Junggar region 1262 1263 during the Carboniferous to Early Permian (344 – 290 Ma) (from Wang, 2006): 344 - 323 1264 Ma, turbidite facies pyroclastic rocks, tuffaceous sandstones, limestones, radiolarian 1265 cherts and tuff layers were deposited in the Liushugou area. Fusulinid, brachiopod, coral 1266 and gastropods fossils occur in ~ 400 m thick limestones. Marine clastic rocks including a ~ 2500m thick limestone sequence, occur in the Tuoli area and contain brachiopods, coral, 1267 1268 gastropod and plant fossils. 323 - 303 Ma, bathyal facies pyroclastic rocks including ~ 1269 1500 m of andesite, basalts and cherts, are abundant in the central area of the western 1270 Junggar region and radiolarian fossils are widespread. To the north, marine and 1271 continental facies sandstone, mudstone and shales with plant fossils occur and have a 1272 combined thickness of 3000 meters. 303 - 290 Ma, bathyal to deep-marine environments 1273 occurred in the Loushugou area where clastic sedimentary rocks contain cherts with 1274 radiolaria and have a combined thickness of ~ 750 meters. Contemporaneously, however, a continental environment was present in the northwest part of the western Junggar 1275 1276 region.



Fig. 2. LA-ICP-MS U-Pb zircon Tera-Wasserburg diagrams with CL images for (a)
Quartz dioritic porphyry (06XJ-145), (b) Quartz dioritic porphyry (06XJ-147), (c) dioritic

porphyrite dike (06XJ-153) and (d) dioritic porphyrite dike (06XJ-156) from the Baogutu
Cu-Au deposits, western Junggar, CAOB. Circles indicate locations of analyzed sites,
with numbers in the circles representing spot numbers. The age for each spot is given.
Scale bars all represent 100 μm.



Fig. 3. CaSiO₃-MgSiO₃-FeSiO₃ diagram showing the compositions of pyroxene
(Morimoto et al., 1988) from the Baogutu adakitic rocks in the western Junggar region,
NW China.



1288

Fig. 4. (a-b) Two back-scattered electron images of typical cores and rims in reversely zoned clinopyroxene phenocrysts from the Baogutu adakitic rocks. (c-d) Composition variations (MgO) in clinopyroxene phenocrysts along the A-A' and B-B' sections from Fig. a and b, respectively. (e) FeO vs. MgO diagram for clinopyroxene phenocrysts. (f) Na₂O vs. MgO diagram for clinopyroxene phenocrysts.



Fig. 5. (a) SiO₂ versus FeO^T/MgO diagrams (Miyashiro, 1974). Dashed boundaries for the tholeiitic and calc-alkaline fields are after Miyashiro (1974). Discriminate boundaries (grey lines) between low-, medium-, and high-Fe suites are after Arculus (2003). (b) SiO₂ - K₂O plot (Peccerillo and Taylor, 1976). (c) SiO₂ – K₂O + Na₂O plot (Middlemost, 1994). (d) Y versus Sr/Y diagram (after Defant and Drummond, 1993). Crystal fractionation

paths of the primary minerals are from Castillo et al. (1999). (e) SiO₂ versus MgO 1300 diagram. (f) SiO₂ versus Mg[#] diagram. Mantle AFC curves, with proportions of 1301 assimilated peridotite indicated, are after Stern and Kilian (1996) (Curve 1) and Rapp et 1302 1303 al. (1999) (Curve 2), peridotite melts and crust AFC curves from Stern and Kilian (1996). for metabasaltic and eclogite experimental melts (1-4.0 1304 Data GPa), and peridotite-hybridized equivalents, are from Rapp et al. (1999) and references therein. Data 1305 for high-Mg andesites of SW Japan are from the following references: Shimoda et al. 1306 (1998), Tatsumi et al. (2006), Tatsumi et al. (2006), and references therein. Data for the 1307 Keramay I-type granitoids and Miaoergou series A-type granites are from Chen and Jahn 1308 1309 (2004), Chen and Arakawa (2005), Gao et al. (2006) and Su et al. (2006). Data for the 1310 Kexi basalts are from Zheng et al. (2007). Data for the Baogutu samples are from 1311 Appendix 3. The fields of subducted oceanic crust-, delaminated lower crust-, and thickened lower crust-derived adakites are after Wang et al. (2006a). 1312



Fig. 6. Chondrite-normalized REE patterns (a) and primitive mantle normalized trace element diagrams (b) for Baogutu samples from the western Junggar compared with the Keramay I-type granitoids and Miaoergou series A-type granites (Data sources as for Fig. 5). Chondrite and primitive mantle normalized values are from Sun and McDonough (1989).



Fig. 7. (a) $\epsilon_{Nd}(t)$ versus (⁸⁷Sr/⁸⁶Sr)_i, (b) $\epsilon_{Hf}(t)$ versus $\epsilon_{Nd}(t)$, ²⁰⁶Pb/²⁰⁴Pb_i versus ²⁰⁷Pb/²⁰⁴Pb_i 1320 (c) and ${}^{208}\text{Pb}/{}^{204}\text{Pb}_i$ (d) plotes. Data for marine sediments and global subducting sediment 1321 (GLOSS) are from Plank and Langmuir (1998) and Chauvel et al. (2008). Data for 1322 Tethyan Ocean MORB are from Xu et al. (2002a), Xu and Castillo (2004) and Zhang et al. 1323 1324 (2005). Subducted oceanic crust-derived adakites, thickened and delaminated mafic lower crust-derived adakitic rocks are after Wang et al. (2006a, 2007a), Huang et al. (2008) and 1325 references therein. Subducted continental crust-derived adakites are after Wang et al. 1326 1327 (2008). Basalts of the Bayingou ophiolite m danges are after Xu et al. (2006c). Data for the Junggar basin volcanic rocks are from Zheng et al. (2007). The high-Mg andesites of 1328 SW Japan are from Hanyu et al. (2006) and reference therein. EPR: East Pacific Rise 1329 1330 basalts, using the Petdb database (Petrological database of the ocean floor, 1331 www.petdb.org). The mantle array and fields for MORB and OIB are from Chauvel et al.



(2008). Data for the Baogutu samples are from Appendix 4. Other data sources and

1333 symbols are the same as for Fig. 5.

1334

Fig. 8. (a) Nb/La versus Sr/Y; (b) SiO₂ versus Sr/Y; (c) SiO₂ versus Dy/Yb; (d) Yb versus La/Yb, crystal fractionation paths of the primary minerals are from Castillo et al. (1999). (e) SiO₂ versus $\varepsilon_{Nd}(t)$; and (f) SiO₂ versus (${}^{87}Sr/{}^{86}Sr)_i$ plots. HPFC, high-pressure fractional crystallization involving garnet (Macpherson et al., 2006); LPFC: crystal

fractionation of an island arc tholeiite series basalt (Danyushevsky et al., 2008). Fractional crystallization models with different proportions of fractionating amphibole and garnet are shown in (b), using sample 06XJ143 as initial magma (SiO₂ = 50.24 wt.%; MgO = 8.46 wt.%). Partition coefficients are after Mori et al. (2007) and references therein: $D_{Nb} = 0.4945$, $D_{La} = 0.0513$, $D_{Sr} = 0.1095$, $D_Y = 1.0291$. Data sources and symbols are the same as in Fig. 5.



1345

Fig. 9. Comparison between the geochemical characteristices of the Baogutu adakitic rocks and the key geochemical parameters used by Martin et al. (2005) to highlight the differences between high-SiO₂ (HSA) and low-SiO₂ (LSA) adakites., (a) K ppm vs Rb ppm; (b) Sr vs CaO+Na₂O; (c) Cr/Ni vs TiO₂; (d) Sr/Y vs Y.



Fig. 10. (a) Th/Ce versus 143 Nd/ 144 Nd ratios diagram, illustrating a negative correlation for the Baogutu samples. The data set was first filtered to exclude all samples with SiO₂ > 64 wt.%, in order to eliminate late possible AFC processes (Turner et al., 2003). (b) Plot of Th versus Th/La. (c) Sm/La versus Th/La plot after the concept of Plank (2005). Data sources of the Baogutu samples, Bayingou ophiolite m danges, high-Mg andesites of SW Japan, marine sediments and GLOSS are the same as for Figs. 5 and 7.



1358 Fig. 11. Simple mixing modeling results of subducted oceanic crust and overlying sediments for the studied rock suites for (a) Sr-Nd, (b) Sr-Pb and (c) Nd-Pb isotopic 1359 spaces, and model primitive mantle normalized trace element pattern compared with an 1360 average Baogutu adakitc rock (d). The trace element pattern of an average Baogutu 1361 adakitc rock is closely matched by a mixture of 95% AOC melt and 5% sediment melt 1362 1363 (red circles), except for the fluid mobile elements. $AOC_b = Bayingou ophiolite m danges$ in the North Tianshan (Xu et al., 2006c); $AOC_k = Kexi$ basalt in the western Junggar 1364 Basin from Zheng et al. (2007). Bulk solid/melt partition coefficients of andesitic-dacitic 1365 melts in equilibrium with a garnet amphibolite residuum (45% Cpx (clinopyroxene), 5% 1366 Gt (garnet), and 50% Amph (amphibolite)). Individual mineral K_d values are from 1367 Rollinson (1993) and references therein, Barth et al. (2002), Klemme et al. (2005), and 1368 the Geochemical Earth Reference Model (GERM) (http://www.earthref.org). Bulk 1369

1370 solid/melt partition coefficients for sediment melting are from Johnson and Plank (1999).

1371 Composition of AOC and GLOSS melts are based on the assumption of 5% and 10%1372 batch melting, respectively. The Baogutu samples and GLOSS are the same as in Fig. 5.



Fig. 12. Variation of zircon U-Pb ages (a), Nd isotopic model ages (b), Mg[#], (c), Sr/Y (d), and zircon saturation temperatures (e) versus distance along a southeast to northwest traverse in the central zone of the western Junggar magmatic province. Zircon U-Pb ages are from Han et al. (2006), Su et al. (2006), Geng et al. (2009) and this study. Others data sources are the same as for Fig. 5.







Fig. 13. Suggested tectonic model for the Carboniferous to Early Permian magmatic province in the western Junggar region. (a) 338-316 Ma: A Paleo-Asia Ocean plate subducting northwestward beneath the Keramay arc; fluids released from the subducted oceanic crust led to partial melting of mantle wedge and the formation of 338-316 Ma volcanic rocks and I-type granitoids. (b) 315-290 Ma: Formation of a slab window in the

central area of the Keramay due to a spreading ridge subduction. (1) Partial melting of subducted slab edge, and subsequent melt-mantle interaction, producing the Baogutu low Mg (Mg[#] < 48) adakitic rocks and high Mg (Mg[#] > 48) diorites, respectively. (2) Fractional crystallization of a mantle-derived magma, which was metasomatized by fluids released from subducted oceanic slab, to generate the I-type granitoids. (3) Partial melting of the juvenile lower crust triggered by upwelling asthenosphere through a slab window, to form the A-type granites. Other symbols are the same as in Fig. 1b.

Analysis -	Content (ppm)					Isotopic ages (Ma)									
Analysis	Th	U	Th/U	²⁰⁷ Pb/ ²⁰⁶ Pb	±1σ	²⁰⁷ Pb/ ²³⁵ U	±1σ	²⁰⁶ Pb/ ²³⁸ U	±1σ	²⁰⁷ Pb/ ²⁰⁶ Pb	±1σ	²⁰⁷ Pb/ ²³⁵ U	±1σ	²⁰⁶ Pb/ ²³⁸ U	±1σ
06XJ145															
1	31	64	0.48	0.0541	0.0018	0.3763	0.0123	0.0505	0.0007	375	49	324	9	317	4
2	227	331	0.68	0.0565	0.0010	0.3933	0.0071	0.0504	0.0006	474	20	337	5	317	4
3	46	83	0.55	0.0551	0.0019	0.3854	0.0132	0.0507	0.0007	417	52	331	10	319	4
4	57	118	0.48	0.0522	0.0017	0.3660	0.0117	0.0508	0.0007	295	48	317	9	320	4
5	155	199	0.78	0.0506	0.0014	0.3558	0.0097	0.0510	0.0007	224	39	309	7	320	4
6	90	190	0.47	0.0540	0.0022	0.3734	0.0152	0.0502	0.0008	369	65	322	11	316	5
7	54	98	0.55	0.0558	0.0016	0.3894	0.0110	0.0507	0.0007	442	39	334	8	319	4
8	341	259	1.32	0.0542	0.0014	0.3701	0.0099	0.0495	0.0006	380	37	320	7	311	4
9	109	176	0.62	0.0580	0.0023	0.3912	0.0150	0.0489	0.0007	531	59	335	11	308	4
10	36	71	0.51	0.0506	0.0029	0.3483	0.0197	0.0500	0.0009	220	99	303	15	314	5
11	70	125	0.56	0.0544	0.0016	0.3766	0.0108	0.0502	0.0007	386	41	325	8	316	4
12	90	168	0.54	0.0526	0.0012	0.3558	0.0084	0.0491	0.0006	310	31	309	6	309	4
13	61	108	0.56	0.0532	0.0014	0.3674	0.0096	0.0501	0.0007	336	36	318	7	315	4
14	133	353	0.38	0.0584	0.0014	0.4820	0.0119	0.0598	0.0008	546	32	399	8	375	5
15	122	145	0.84	0.0627	0.0014	0.4264	0.0097	0.0493	0.0006	698	28	361	7	310	4
16	292	477	0.61	0.0589	0.0010	0.4695	0.0081	0.0578	0.0007	563	18	391	6	362	4
17	142	144	0.99	0.0568	0.0014	0.3943	0.0094	0.0503	0.0006	484	31	337	7	317	4
18	142	235	0.60	0.0563	0.0012	0.3834	0.0081	0.0494	0.0006	465	26	330	6	311	4
19	297	298	0.99	0.0552	0.0011	0.3851	0.0076	0.0506	0.0006	419	23	331	6	318	4
20	268	552	0.49	0.0586	0.0010	0.4736	0.0083	0.0586	0.0007	554	19	394	6	367	4
21	62	82	0.76	0.0543	0.0026	0.3687	0.0176	0.0492	0.0008	385	78	319	13	310	5
22	119	178	0.67	0.0578	0.0015	0.3991	0.0102	0.0501	0.0007	522	33	341	7	315	4
23	131	193	0.68	0.0578	0.0013	0.4793	0.0112	0.0601	0.0008	522	29	398	8	376	5
24	405	419	0.97	0.0585	0.0019	0.4026	0.0132	0.0500	0.0007	547	47	344	10	314	4
25	165	204	0.81	0.0525	0.0013	0.3677	0.0093	0.0508	0.0007	307	35	318	7	319	4
26	155	328	0.47	0.0563	0.0011	0.4477	0.0088	0.0577	0.0007	465	23	376	6	361	4

Appendix 1 LA-ICP-MS U-Pb isotopic analyses for zircons from the ore-porphyries in the West Junggar

1394 Appendix 1 Continued

Analysis -	Content (ppm)					Isotopic r	atios	Isotopic ages (Ma)							
Analysis	Th	U	Th/U	²⁰⁷ Pb/ ²⁰⁶ Pb	±1σ	²⁰⁷ Pb/ ²³⁵ U	±1σ	²⁰⁶ Pb/ ²³⁸ U	±1σ	²⁰⁷ Pb/ ²⁰⁶ Pb	±1σ	²⁰⁷ Pb/ ²³⁵ U	±1σ	²⁰⁶ Pb/ ²³⁸ U	±1σ
06XJ147															
1	59	84	0.70	0.0566	0.0020	0.3856	0.0134	0.0494	0.0007	477	53	331	10	311	4
2	97	126	0.77	0.0582	0.0037	0.3923	0.0246	0.0488	0.0009	539	104	336	18	307	6
3	57	68	0.84	0.0554	0.0021	0.3820	0.0146	0.0501	0.0007	426	60	329	11	315	4
4	69	80	0.87	0.0511	0.0016	0.3526	0.0108	0.0501	0.0007	244	47	307	8	315	4
5	61	83	0.73	0.0564	0.0016	0.3845	0.0110	0.0494	0.0006	469	41	330	8	311	4
6	103	119	0.86	0.0562	0.0018	0.3819	0.0119	0.0493	0.0007	460	45	328	9	310	4
7	77	93	0.83	0.0530	0.0017	0.3572	0.0114	0.0489	0.0007	328	48	310	9	308	4
8	36	56	0.64	0.0479	0.0028	0.3274	0.0190	0.0495	0.0009	96	97	288	15	312	5
9	51	66	0.78	0.0525	0.0018	0.3571	0.0122	0.0493	0.0007	309	53	310	9	310	4
10	46	61	0.75	0.0622	0.0019	0.4309	0.0133	0.0502	0.0007	682	43	364	9	316	4
11	53	77	0.68	0.0612	0.0037	0.4139	0.0244	0.0490	0.0009	647	95	352	18	309	6
12	70	86	0.81	0.0578	0.0015	0.3915	0.0103	0.0491	0.0006	523	35	335	8	309	4
13	67	90	0.75	0.0585	0.0020	0.4022	0.0136	0.0498	0.0007	550	49	343	10	313	4
14	45	57	0.78	0.0570	0.0034	0.3926	0.0228	0.0499	0.0009	492	96	336	17	314	6
15	50	63	0.79	0.0550	0.0020	0.3733	0.0135	0.0493	0.0007	410	55	322	10	310	4
16	51	75	0.68	0.0597	0.0020	0.4028	0.0132	0.0489	0.0007	592	47	344	10	308	4
17	64	77	0.82	0.0582	0.0019	0.3922	0.0127	0.0489	0.0007	537	47	336	9	308	4
18	47	64	0.72	0.0604	0.0037	0.4130	0.0249	0.0496	0.0009	617	98	351	18	312	6
19	87	104	0.83	0.0521	0.0016	0.3528	0.0106	0.0491	0.0007	289	44	307	8	309	4
20	47	66	0.72	0.0744	0.0022	0.5058	0.0151	0.0493	0.0007	1052	38	416	10	310	4
21	53	72	0.73	0.0520	0.0016	0.3539	0.0105	0.0493	0.0007	286	43	308	8	310	4
22	60	86	0.70	0.0517	0.0031	0.3441	0.0203	0.0482	0.0009	274	102	300	15	304	5
06XJ153															
1	60	79	0.76	0.0554	0.0014	0.3800	0.0098	0.0498	0.0006	429	35	327	7	313	4
2	134	181	0.74	0.0604	0.0017	0.4203	0.0121	0.0505	0.0007	618	39	356	9	317	4
3	108	144	0.75	0.0524	0.0016	0.3518	0.0107	0.0487	0.0007	304	45	306	8	306	4

1396	Appendix 1 Continued

Analysis -	Content (ppm)					Isotopic r	atios	Isotopic ages (Ma)							
Analysis	Th	U	Th/U	²⁰⁷ Pb/ ²⁰⁶ Pb	±1σ	²⁰⁷ Pb/ ²³⁵ U	±1σ	²⁰⁶ Pb/ ²³⁸ U	±1σ	²⁰⁷ Pb/ ²⁰⁶ Pb	±1σ	²⁰⁷ Pb/ ²³⁵ U	±1σ	²⁰⁶ Pb/ ²³⁸ U	±1σ
4	184	600	0.31	0.1522	0.0021	2.6299	0.0382	0.1254	0.0015	2370	11	1309	11	761	9
5	140	378	0.37	0.0646	0.0013	0.5174	0.0103	0.0581	0.0007	761	22	423	7	364	4
6	53	60	0.88	0.0517	0.0041	0.3542	0.0278	0.0497	0.0011	272	140	308	21	313	6
7	55	129	0.43	0.0518	0.0012	0.3592	0.0085	0.0503	0.0006	274	31	312	6	317	4
8	27	44	0.61	0.0668	0.0024	0.4667	0.0168	0.0507	0.0007	830	51	389	12	319	5
9	22	34	0.65	0.0674	0.0030	0.4561	0.0200	0.0491	0.0008	850	65	382	14	309	5
10	37	51	0.72	0.0584	0.0032	0.3931	0.0209	0.0489	0.0008	544	86	337	15	307	5
11	47	95	0.49	0.0500	0.0015	0.3457	0.0105	0.0502	0.0007	193	45	301	8	316	4
12	111	167	0.66	0.0580	0.0017	0.4379	0.0127	0.0548	0.0007	529	40	369	9	344	5
13	128	195	0.66	0.0645	0.0019	0.4442	0.0129	0.0500	0.0007	757	39	373	9	314	4
14	741	506	1.46	0.0554	0.0010	0.3853	0.0071	0.0504	0.0006	429	21	331	5	317	4
15	63	91	0.69	0.0586	0.0024	0.4132	0.0164	0.0512	0.0008	550	60	351	12	322	5
16	87	183	0.47	0.0515	0.0012	0.3489	0.0080	0.0491	0.0006	263	30	304	6	309	4
17	825	1712	0.48	0.0545	0.0014	0.3788	0.0095	0.0504	0.0007	392	33	326	7	317	4
18	329	263	1.25	0.0559	0.0014	0.3772	0.0093	0.0489	0.0006	448	32	325	7	308	4
19	101	100	1.01	0.0618	0.0039	0.4124	0.0256	0.0484	0.0009	668	100	351	18	305	6
20	361	381	0.95	0.0567	0.0015	0.4014	0.0105	0.0513	0.0007	480	35	343	8	323	4
06XJ156															
1	44	85	0.52	0.0542	0.0028	0.3640	0.0181	0.0487	0.0007	380	120	315	13	306	4
2	41	79	0.52	0.0546	0.0016	0.3730	0.0110	0.0495	0.0007	396	42	322	8	312	4
3	36	67	0.54	0.0506	0.0025	0.3410	0.0167	0.0489	0.0008	222	83	298	13	308	5
4	51	98	0.52	0.0694	0.0021	0.4744	0.0144	0.0496	0.0007	909	39	394	10	312	4
5	30	56	0.53	0.0531	0.0029	0.3573	0.0190	0.0488	0.0008	331	89	310	14	307	5
6	37	67	0.55	0.0530	0.0028	0.3578	0.0183	0.0489	0.0007	330	124	311	14	308	4
7	45	84	0.53	0.0534	0.0030	0.3601	0.0199	0.0489	0.0009	348	93	312	15	308	5
8	42	75	0.57	0.1004	0.0036	0.6805	0.0237	0.0492	0.0008	1632	41	527	14	309	5
9	67	115	0.58	0.0614	0.0035	0.4118	0.0227	0.0487	0.0009	652	87	350	16	306	5

1398	Appendix 1 Continued

Analysis -	Content ((ppm)				Isotopic r	atios	Isotopic ages (Ma)							
	Th	U	Th/U	²⁰⁷ Pb/ ²⁰⁶ Pb	±1σ	²⁰⁷ Pb/ ²³⁵ U	±1σ	²⁰⁶ Pb/ ²³⁸ U	±1σ	²⁰⁷ Pb/ ²⁰⁶ Pb	±1σ	²⁰⁷ Pb/ ²³⁵ U	±1σ	²⁰⁶ Pb/ ²³⁸ U	±1σ
10	36	69	0.52	0.0586	0.0033	0.3914	0.0211	0.0484	0.0008	553	125	335	15	305	5
11	51	87	0.58	0.0887	0.0032	0.5818	0.0208	0.0476	0.0008	1398	44	466	13	300	5
12	38	65	0.58	0.0546	0.0039	0.3669	0.0258	0.0488	0.0008	395	167	317	19	307	5
13	35	69	0.51	0.0595	0.0021	0.4056	0.0146	0.0494	0.0007	586	54	346	11	311	4
14	32	63	0.51	0.0631	0.0035	0.4182	0.0226	0.0481	0.0007	712	123	355	16	303	4
15	40	72	0.55	0.0529	0.0029	0.3544	0.0190	0.0486	0.0007	323	129	308	14	306	4
16	51	86	0.60	0.0535	0.0030	0.3567	0.0193	0.0484	0.0007	350	131	310	14	304	4
17	45	84	0.54	0.0533	0.0019	0.3624	0.0126	0.0494	0.0007	339	53	314	9	311	4
18	42	80	0.52	0.0615	0.0029	0.4199	0.0189	0.0496	0.0007	655	104	356	14	312	4
19	49	60	0.81	0.0649	0.0039	0.4327	0.0252	0.0483	0.0008	772	131	365	18	304	5
20	54	68	0.80	0.0679	0.0041	0.4612	0.0266	0.0493	0.0008	864	127	385	18	310	5
21	46	78	0.59	0.0531	0.0028	0.3615	0.0184	0.0494	0.0007	335	123	313	14	311	4
22	54	91	0.59	0.0583	0.0027	0.3932	0.0176	0.0489	0.0007	541	106	337	13	308	4
23	90	91	1.00	0.0884	0.0049	0.5773	0.0305	0.0474	0.0008	1390	109	463	20	298	5
24	51	81	0.63	0.0607	0.0027	0.4068	0.0177	0.0486	0.0008	628	66	347	13	306	5
25	49	84	0.58	0.0609	0.0033	0.4067	0.0209	0.0485	0.0007	635	118	346	15	305	4
26	45	82	0.54	0.0547	0.0032	0.3664	0.0205	0.0486	0.0008	399	134	317	15	306	5
27	51	82	0.62	0.0567	0.0024	0.3808	0.0162	0.0487	0.0008	478	66	328	12	307	5
28	47	82	0.58	0.0584	0.0030	0.3945	0.0191	0.0490	0.0007	544	114	338	14	309	4
29	51	71	0.72	0.0669	0.0045	0.4372	0.0285	0.0474	0.0008	833	144	368	20	299	5
30	26	41	0.65	0.0714	0.0081	0.4833	0.0538	0.0491	0.0012	968	244	400	37	309	8

Spot	Crystal	Zone	SiO2	TiO2	Al2O3	Cr2O3	FeO	MnO	MgO	CaO	Na2O	K2O	Total	Wo	En	Fs	Mg#
1	1	R	51.53	0.27	1.70		8.66	0.27	14.41	22.70	0.37	0.01	100.08	45.1	39.8	13.7	74.8
2	1	R	51.61	0.41	1.57	0.02	8.37	0.26	15.00	22.59	0.39	0.00	100.23	44.5	41.1	13.1	76.2
3	1	R	53.24	0.55	1.67	0.04	8.66	0.25	14.88	20.20	0.44		99.94	41.5	42.6	14.3	75.4
4	1	R	51.20	0.48	1.57		9.27	0.27	14.68	21.67	0.38	0.02	99.54	43.2	40.7	14.7	73.8
5	1	С	51.11	0.49	1.87	0.09	9.78	0.26	14.69	21.16	0.41	0.01	99.87	42.2	40.8	15.5	72.8
6	1	С	51.02	0.38	1.41		9.54	0.31	14.45	21.66	0.32	0.03	99.11	43.4	40.3	15.3	73.0
7	1	С	50.42	0.56	2.32	0.10	9.88	0.23	14.52	21.22	0.41	0.02	99.67	42.5	40.4	15.6	72.4
8	1	С	49.92	0.62	3.18	0.30	9.97	0.22	13.91	21.67	0.49	0.01	100.28	43.5	38.9	15.8	71.3
9	1	С	51.21	0.48	1.54	0.02	9.27	0.24	14.43	21.69	0.40	0.00	99.28	43.5	40.3	14.8	73.5
10	1	С	50.70	0.56	1.88		10.12	0.33	14.31	21.53	0.39	0.01	99.81	42.9	39.7	16.1	71.6
11	1	С	50.13	0.59	2.65	0.31	9.87	0.27	14.16	21.19	0.50		99.68	42.7	39.7	15.8	71.9
12	1	С	50.05	0.58	2.77	0.43	10.12	0.27	14.11	21.00	0.41	0.02	99.74	42.5	39.7	16.2	71.3
13	1	С	50.50	0.49	2.69	0.45	9.58	0.25	14.38	20.91	0.43	0.02	99.71	42.4	40.6	15.4	72.8
14	1	С	50.28	0.54	2.42	0.10	10.22	0.27	14.25	21.18	0.46		99.71	42.4	39.7	16.2	71.3
15	1	R	51.30	0.45	1.59	0.02	9.41	0.27	14.98	21.88	0.33	0.00	100.21	43.1	41.0	14.7	73.9
16	1	R	50.92	0.49	1.80	0.03	9.43	0.22	14.75	21.70	0.43		99.78	43.0	40.7	14.8	73.6
17	1	R	51.45	0.46	1.63		9.61	0.26	14.74	21.55	0.29	0.01	99.99	42.9	40.8	15.2	73.2
18	1	R	51.23	0.34	1.59	0.00	9.21	0.23	14.67	21.90	0.49	0.01	99.67	43.4	40.4	14.4	73.9
19	2	R	51.22	0.32	1.31		8.88	0.28	14.57	22.73	0.33		99.63	44.9	40.0	14.0	74.5
20	2	R	51.66	0.48	1.31	0.01	9.93	0.36	15.23	20.81	0.26	0.01	100.06	41.3	42.0	15.8	73.2
21	2	R	51.16	0.41	1.50		9.60	0.25	14.82	21.72	0.30	0.00	99.77	43.0	40.8	15.1	73.3
22	2	R	51.17	0.30	1.29		9.33	0.29	14.66	22.09	0.41	0.02	99.56	43.6	40.3	14.7	73.7
23	2	С	50.42	0.52	2.56	0.04	9.88	0.21	14.65	21.48	0.30	0.01	100.06	42.8	40.6	15.5	72.5
24	2	С	50.65	0.61	2.07	0.05	9.66	0.30	14.57	21.54	0.36	0.01	99.79	43.0	40.4	15.3	72.9
25	2	С	51.49	0.20	1.28	0.01	8.98	0.28	14.27	22.62	0.38		99.53	44.9	39.5	14.2	73.9
26	2	С	50.94	0.42	1.51	0.02	9.77	0.33	14.29	21.96	0.40	0.01	99.64	43.6	39.5	15.5	72.2
27	2	С	50.63	0.64	2.36	0.01	9.57	0.27	13.99	22.16	0.42		100.05	44.3	38.9	15.2	72.3
28	2	С	51.72	0.27	0.85	0.04	11.38	0.44	14.12	20.99	0.23	0.00	100.03	41.8	39.1	18.3	68.8
29	2	С	50.57	0.54	2.51	0.02	9.43	0.25	14.22	22.29	0.45		100.27	44.3	39.3	14.8	72.9
30	2	С	51.47	0.60	1.87		9.43	0.29	14.24	22.30	0.47		100.67	44.2	39.3	14.9	72.9
31	2	С	50.81	0.56	2.33	0.00	9.41	0.30	13.95	22.05	0.50	0.02	99.92	44.2	38.9	15.1	72.5
32	2	С	50.88	0.64	2.43		9.63	0.30	14.01	22.03	0.47		100.39	44.0	38.9	15.3	72.2
33	2	R	51.59	0.28	0.98		10.15	0.41	14.62	21.15	0.35	0.01	99.55	42.0	40.4	16.3	72.0
34	2	R	51.99	0.22	1.25		9.32	0.27	14.66	21.90	0.41		100.02	43.4	40.4	14.7	73.7
35	2	R	51.43	0.37	1.54		9.66	0.30	14.65	22.02	0.39		100.34	43.4	40.1	15.1	73.0
36	2	R	51.48	0.46	1.65	0.03	9.49	0.30	14.47	22.16	0.36	0.01	100.40	43.9	39.9	15.0	73.1
37	2	R	50.03	0.39	1.24	0.02	9.90	0.33	15.08	21.86	0.35		99.20	42.6	40.9	15.3	73.1
38	3	R	51.28	0.41	1.64	0.02	10.11	0.40	15.14	21.24	0.36	0.01	100.61	41.6	41.2	15.9	72.7

Appendix 2 Major element compositions (wt %) of clinopyroxene and plagioclase

from the Baogutu adakitic ore-porphyries

Appendix 2 Continued

39	3	R	51.33 0	.28 1.4	44 (0.00	9.35	0.30	14.86	22.14	0.41	0.01	100.12	43.4	40.5	14.6	73.9
40	3	R	51.64 0	.34 1.	36 (0.03	9.73	0.35	14.51	22.11	0.41	0.00	100.46	43.5	39.7	15.3	72.7
41	3	R	51.45 0	.35 1.4	41		9.33	0.28	14.51	22.09	0.43	0.02	99.85	43.8	40.0	14.7	73.5
42	3	С	51.53 0	.20 1.	29 (0.01	8.58	0.31	14.27	22.94	0.34		99.47	45.6	39.5	13.7	74.8
43	3	С	51.70 0	.31 1.4	44 (0.04	9.14	0.26	14.25	22.62	0.36	0.00	100.11	44.9	39.4	14.4	73.5
44	3	С	51.67 0	.30 1.	30		9.27	0.29	14.10	22.87	0.37	0.00	100.18	45.3	38.8	14.6	73.0
45	3	С	51.53 0	.48 1.	34 (0.03	9.42	0.30	14.42	22.48	0.39		100.38	44.3	39.5	14.8	73.2
46	3	С	50.87 0	.36 1.	62 (0.02	9.94	0.44	14.26	22.20	0.43	0.01	100.13	43.7	39.0	15.8	71.9
47	3	С	51.85 0	.48 1.4	48 (0.05	9.65	0.33	14.56	22.07	0.41	0.01	100.87	43.5	39.9	15.2	72.9
48	3	С	50.73 0	.48 1.:	58 (0.02	10.04	0.32	14.98	21.34	0.28		99.77	42.1	41.1	15.8	72.7
49	3	R	51.33 0	.40 1.4	48 (0.01	9.80	0.30	14.54	22.00	0.43		100.28	43.3	39.8	15.4	72.6
50	3	R	50.98 0	.43 1.4	43		10.26	0.37	15.23	21.26	0.42	0.01	100.37	41.4	41.2	15.9	72.6
51	3	R	51.23 0	.47 1.:	52 (0.03	10.21	0.33	15.17	21.34	0.40	0.01	100.71	41.6	41.1	15.9	72.6
52	3	R	50.92 0	.20 2.	63 (0.68	6.11	0.14	15.20	23.13	0.44	0.01	99.46	46.4	42.4	9.7	81.6
53	3	С	51.18 0	.39 1.4	44 (0.03	9.37	0.32	14.42	22.07	0.47	0.02	99.70	43.7	39.8	14.8	73.3
54	3	С	50.99 0	.25 1.	65 (0.05	9.37	0.35	14.30	21.88	0.47	0.01	99.33	43.6	39.7	15.0	73.1
55	3	С	50.00 0	.68 3.4	42 (0.33	8.20	0.22	14.17	22.24	0.55	0.01	99.81	45.0	39.9	13.2	75.5
56	3	R	51.04 0	.39 2.	64 (0.76	5.97	0.17	15.25	22.83	0.43	0.01	99.49	46.1	42.8	9.6	82.0
57	4	U	51.01 0	.41 1.:	52		9.06	0.32	15.19	22.06	0.36		99.92	43.2	41.4	14.2	74.9
58	4	U	51.28 0	.29 1.:	50		9.48	0.26	14.22	22.53	0.38		99.93	44.6	39.2	14.9	72.8
59	4	U	51.74 0	.28 1.	39 (0.04	9.53	0.33	14.15	22.67	0.39		100.52	44.7	38.9	15.0	72.6
60	5	U	51.40 0	.35 1.4	42 (0.05	9.51	0.28	14.38	22.14	0.45	0.02	99.98	43.8	39.6	15.0	72.9
61	5	U	51.43 0	.31 1.	31		9.55	0.28	14.40	22.01	0.41	0.02	99.71	43.7	39.8	15.1	72.9
62	5	U	51.68 0	.37 1.	30 (0.00	9.11	0.26	14.41	22.15	0.36	0.01	99.64	44.2	40.0	14.5	73.8
63	6	U	51.14 0	.28 1.	37 (0.03	9.41	0.29	14.53	22.12	0.38		99.55	43.8	40.0	14.8	73.3
64	6	U	51.88 0	.30 1.	07 (0.04	9.10	0.26	14.40	22.69	0.30		100.03	44.9	39.7	14.4	73.8
65	6	U	51.63 0	.39 1.:	50		9.37	0.25	14.68	21.99	0.40	0.00	100.20	43.5	40.4	14.7	73.6
66	6	U	51.50 0	.45 1.:	50 (0.03	9.63	0.30	14.66	22.07	0.31	0.01	100.46	43.5	40.2	15.1	73.1
67	8	U	51.25 0	.54 1.	68 (0.03	9.35	0.27	14.66	21.62	0.41	0.00	99.81	43.1	40.6	14.8	73.6
68	9	R	51.22 0	.36 2.	64 (0.63	5.30	0.14	16.08	22.40	0.35	0.01	99.13	45.2	45.1	8.5	84.4
69	9	С	51.56 0	.42 1.3	88 (0.03	8.69	0.28	14.72	21.99	0.37		99.93	43.9	40.9	13.9	75.1
70	9	С	50.46 0	.47 2.9	94 (0.03	8.40	0.28	14.24	22.25	0.43		99.50	44.9	40.0	13.5	75.1
71	9	С	51.52 0	.21 1.4	47 (0.01	9.35	0.26	14.27	22.19	0.44		99.71	44.1	39.5	14.8	73.1
72	9	R	51.13 0	.43 1.	67 (0.04	8.76	0.29	14.71	21.80	0.31	0.04	99.19	43.7	41.1	14.1	74.9
73	10	U	51.66 0	.36 1.:	53		8.43	0.28	14.92	22.01	0.41		99.59	43.8	41.3	13.4	75.9
74	10	U	51.92 0	.35 1.4	43 (0.05	9.16	0.30	14.47	22.30	0.50		100.46	44.0	39.8	14.4	73.8
75	11	U	51.96 0	.13 1.	30 (0.02	8.42	0.27	14.75	22.57	0.37		99.78	44.7	40.7	13.3	75.7
76	11	U	51.60 0	.27 1.	12 (0.05	10.83	0.30	14.09	21.54	0.41	0.02	100.21	42.7	38.8	17.0	69.9

Appendix 2 Continued

77	12	U	50.70	0.52	2.60	0.06	9.36	0.21	14.23	22.32	0.52		100.52	44.3	39.3	14.6	73.0
78	12	U	50.93	0.43	1.91	0.02	9.22	0.28	14.12	22.30	0.42	0.00	99.63	44.6	39.3	14.7	73.2
79	13	U	50.43	0.58	2.57		8.84	0.29	14.20	22.15	0.39		99.45	44.6	39.8	14.2	74.1
80	13	U	51.51	0.38	1.75		8.53	0.30	14.31	22.31	0.40	0.01	99.49	44.8	40.0	13.8	74.9
81	14	R	51.19	0.40	2.47	0.69	4.88	0.08	16.07	23.04	0.42		99.23	46.1	44.7	7.7	85.4
82	14	R	51.10	0.37	2.68	0.67	5.44	0.11	16.50	22.20	0.36	0.02	99.45	44.3	45.8	8.6	84.4
83	14	R	51.52	0.38	2.42	0.79	5.15	0.14	15.95	22.83	0.42	0.00	99.60	45.8	44.5	8.2	84.7
84	14	R	51.34	0.24	2.40	0.69	4.94	0.12	16.03	22.93	0.38	0.01	99.07	46.0	44.8	7.9	85.3
85	14	R	50.79	0.47	2.82	0.38	7.21	0.19	14.86	22.75	0.48		99.95	45.5	41.3	11.4	78.6
86	14	С	51.39	0.18	1.52		8.84	0.32	14.38	22.24	0.50	0.02	99.38	44.3	39.8	14.1	74.3
87	14	С	51.56	0.19	1.47		9.32	0.26	14.38	22.30	0.46	0.01	99.96	44.1	39.6	14.7	73.3
88	14	R	51.32	0.23	1.51	0.02	7.97	0.24	14.99	22.41	0.55		99.24	44.3	41.2	12.5	77.0
89	15	U	50.84	0.44	1.93	0.04	9.41	0.31	14.53	21.40	0.38		99.27	43.0	40.6	15.1	73.3
90	15	U	51.30	0.48	1.64		9.34	0.21	14.94	21.69	0.29	0.00	99.90	43.0	41.3	14.7	74.0
91	16	U	51.23	0.55	2.09	0.03	9.27	0.29	14.78	21.87	0.44	0.01	100.59	43.2	40.6	14.6	74.0
92	16	U	50.64	0.26	1.75	0.04	8.62	0.21	14.64	22.48	0.46	0.01	99.11	44.5	40.3	13.5	75.1
93	16	U	51.31	0.33	1.46	0.05	9.56	0.24	14.60	21.96	0.36		99.87	43.5	40.2	15.0	73.1
94	16	U	52.02	0.37	1.35	0.02	8.51	0.31	15.06	21.89	0.32		99.84	43.6	41.7	13.6	75.9
95	17	R	51.45	0.42	1.43		8.90	0.28	14.91	22.19	0.36		99.93	43.8	40.9	14.0	74.9
96	17	R	51.21	0.46	1.44		9.90	0.29	14.73	22.00	0.38		100.41	43.1	40.2	15.4	72.6
97	17	С	51.04	0.39	1.49	0.01	9.53	0.31	14.55	21.93	0.43	0.01	99.70	43.4	40.1	15.0	73.1
98	17	С	51.02	0.39	1.40	0.06	9.50	0.32	14.24	22.27	0.42	0.01	99.62	44.2	39.3	15.1	72.7
99	17	R	51.44	0.43	1.40	0.01	9.08	0.30	14.89	22.09	0.28		99.93	43.7	41.0	14.4	74.5
100	17	R	51.72	0.33	1.50	0.06	9.75	0.32	14.28	21.90	0.45		100.31	43.5	39.4	15.5	72.3
101	17	R	50.98	0.34	1.42		9.19	0.33	15.03	22.12	0.28		99.71	43.5	41.1	14.5	74.4
102	17	R	51.23	0.46	1.41		9.90	0.34	14.57	22.16	0.44	0.00	100.50	43.4	39.7	15.4	72.4
103	17	R	51.63	0.57	1.35	0.05	9.22	0.31	14.93	21.84	0.27	0.00	100.18	43.3	41.1	14.6	74.3
104	17	С	51.17	0.47	1.49		9.42	0.27	14.37	21.89	0.42	0.01	99.51	43.7	39.9	14.9	73.1
105	17	С	51.78	0.31	1.16	0.03	9.08	0.21	14.30	22.73	0.36		99.96	45.0	39.4	14.2	73.7
106	17	С	51.29	0.49	1.49	0.01	9.38	0.33	14.34	22.01	0.45		99.78	43.8	39.7	14.9	73.2
107	17	R	51.53	0.51	1.43	0.03	9.30	0.29	14.66	22.21	0.38	0.00	100.33	43.8	40.2	14.6	73.7
108	17	R	51.76	0.35	1.43	0.02	9.10	0.30	14.63	22.40	0.43	0.02	100.43	44.1	40.1	14.3	74.1
109	17	R	51.72	0.37	1.43		9.32	0.26	14.67	21.77	0.34		99.88	43.3	40.7	14.8	73.7
110	18	R	51.29	0.50	1.35	0.03	9.67	0.26	14.85	21.80	0.27	0.01	100.02	43.1	40.8	15.2	73.2
111	18	R	51.50	0.23	1.24	0.01	8.71	0.28	14.86	22.12	0.37	0.00	99.32	43.9	41.0	13.8	75.3
112	18	R	51.68	0.27	1.87	0.18	6.87	0.20	15.33	22.96	0.37		99.73	45.5	42.3	10.9	79.9
113	18	R	51.78	0.22	1.70	0.47	6.88	0.19	16.06	21.77	0.38	0.01	99.45	43.3	44.4	10.9	80.6
114	18	R	51.64	0.48	1.66	0.03	8.74	0.22	14.76	22.13	0.42		100.06	44.0	40.8	13.8	75.1
Appendix 2 Continued

Spot	Crystal	Zone	SiO2	TiO2	A12O3	Cr2O3	FeO	MnO	MgO	CaO	Na2O	K2O	Total	Wo	En	Fs	Mg#
115	18	С	51.17	0.29	1.49		8.91	0.19	14.60	22.21	0.49	0.01	99.37	44.0	40.3	13.9	74.5
116	18	С	51.64	0.26	1.42	0.01	9.58	0.28	14.36	22.07	0.46		100.07	43.7	39.6	15.1	72.8
117	18	С	51.58	0.31	1.34	0.02	9.42	0.34	14.54	22.27	0.51		100.33	43.7	39.7	14.8	73.3
118	18	С	51.33	0.38	1.29		9.95	0.34	14.25	21.83	0.43		99.79	43.3	39.4	15.8	71.8
119	18	С	50.94	0.40	1.90		9.90	0.22	14.18	21.64	0.41	0.00	99.59	43.3	39.5	15.7	71.8
120	18	С	50.94	0.39	1.93	0.00	10.05	0.29	14.42	21.50	0.35	0.01	99.87	42.8	40.0	15.9	71.9
121	18	С	51.43	0.38	1.96	0.03	9.56	0.29	14.36	21.24	0.35	0.03	99.63	42.9	40.4	15.4	72.8
122	18	С	51.99	0.33	1.28		9.03	0.30	14.96	22.27	0.39		100.53	43.7	40.8	14.2	74.7
123	18	R	52.01	0.17	1.68	0.58	6.70	0.20	16.11	21.55	0.45	0.01	99.46	43.0	44.7	10.7	81.1
124	18	R	51.66	0.25	1.73	0.48	6.73	0.20	15.67	21.99	0.41		99.12	44.1	43.7	10.8	80.6
125	18	R	51.68	0.37	1.88	0.01	7.65	0.18	14.97	22.55	0.40	0.01	99.70	45.0	41.5	12.1	77.7

R-Rim; C-Core; U- Unzoned of small pyrocene grains; -below detection limit

1 Appendix 2 Continued

Spot		Zone	SiO2	TiO2	A12O3	Cr2O3	FeO	MnO	MgO	CaO	Na2O	K2O	Total	An
06XJ147 01	Р	R	57.01		26.67	0.06	0.36		0.01	9.43	6.51	0.36	100.42	43.61
06XJ147 02	Р	С	55.71	0.05	27.40		0.35	0.02		10.13	5.82	0.35	99.85	48.06
06XJ147 03	Р	С	53.11	0.02	29.04		0.39	0.02	0.02	12.41	4.48	0.23	99.72	59.68
06XJ147 04	Р	R	58.38	0.05	25.52	0.04	0.36		0.02	8.03	7.02	0.51	99.93	37.64
06XJ147 09	Р	R	58.58	0.03	25.18	0.00	0.30		0.02	7.84	7.37	0.51	99.84	36.00
06XJ147 10	Р	R	57.01	0.02	26.43		0.31	0.04	0.01	9.16	6.62	0.42	100.01	42.35
06XJ147 11	Р	R	58.28	0.02	25.41	0.05	0.24	0.04	0.02	7.96	7.21	0.52	99.74	36.81
06XJ147 12	Р	R	57.69	0.01	26.13	0.02	0.26	0.03	0.02	8.75	6.77	0.46	100.20	40.63
06XJ147 13	Р	R	58.61	0.01	25.35		0.23		0.02	7.72	7.54	0.44	99.93	35.25
06XJ147 14	Р	R	56.32	0.06	27.10	0.00	0.33	0.03	0.03	9.65	6.34	0.39	100.24	44.71
06XJ147 15	Р	R	56.28		26.88		0.30	0.02	0.01	9.71	6.19	0.39	99.80	45.45
06XJ147 16	Р	С	56.12	0.09	27.12		0.29		0.01	9.88	6.08	0.36	99.95	46.37
06XJ147 17	Р	С	54.32	0.09	28.09	0.00	0.27	0.00	0.02	11.26	5.48	0.30	99.86	52.28
06XJ147 18	Р	С	54.80	0.03	27.96	0.01	0.20	0.01	0.01	10.85	5.87	0.21	99.96	49.95
06XJ147 19	Р	С	55.09	0.04	27.90		0.24		0.01	10.89	5.77	0.32	100.26	50.18
06XJ147 20	Р	R	57.65	0.04	26.00	0.02	0.22		0.02	8.63	6.79	0.46	99.88	40.21
06XJ147 22	Р	С	54.75	0.00	28.16		0.28		0.03	11.08	5.68	0.29	100.30	51.05
06XJ147 23	Р	С	54.64		28.11	0.03	0.31	0.00	0.02	11.02	5.46	0.32	99.97	51.77
06XJ147 24	Р	R	59.10	0.05	24.99		0.28	0.05	0.01	7.24	7.73	0.47	99.92	33.22
06XJ153 01	Р	С	56.08	0.02	26.84	0.01	0.47		0.03	10.11	6.12	0.34	100.03	46.85
06XJ153 02	Р	С	54.52	0.02	27.68		0.49	0.04	0.03	10.82	5.70	0.30	99.60	50.35
06XJ153 03	Р	С	53.59	0.03	28.74	0.02	0.35	0.01	0.03	12.04	4.98	0.25	100.06	56.39
06XJ153 04	Р	R	55.86	0.02	26.88	0.01	0.41		0.04	10.24	6.20	0.32	100.02	46.90
06XJ153 05	Р	С	53.74	0.07	28.14		0.55	0.00	0.07	11.72	5.22	0.21	99.72	54.73
06XJ153 06	Р	R	56.01		26.63		0.48	0.03	0.05	9.76	6.44	0.31	99.74	44.80
06XJ153 07	Р	С	52.93	0.10	26.60	0.01	1.21	0.01	0.73	11.92	4.72	0.22	98.51	57.51
06XJ153 08	Р	R	55.03		27.77	0.03	0.50	0.05	0.05	11.04	5.50	0.28	100.26	51.77
06XJ153 09	Μ		58.53	0.03	24.58	0.01	0.39		0.03	7.22	7.46	0.47	98.72	33.92
06XJ153 10	Μ		63.80	0.03	22.31		0.14	0.01	0.01	3.25	9.87	0.64	100.07	14.85
06XJ153 11	Μ		55.47	0.02	26.70	0.04	0.45	0.01	0.04	9.68	6.20	0.26	98.94	45.64
06XJ153 12	Μ		67.39	0.02	20.17		0.25		0.01	0.75	11.57	0.21	100.38	3.40
06XJ153 13	Μ		65.09	0.03	21.67	0.01	0.03			2.42	10.83	0.13	100.22	10.92
06XJ153 14	Μ		62.84	0.04	21.22	0.01	1.26	0.01	0.84	4.08	9.49	0.37	100.17	18.81
06XJ153 15	Μ		65.71		20.79	0.02	0.02			1.69	10.91	0.18	99.37	7.80
06XJ153 18	Μ		57.08	0.05	26.26	0.04	0.30	0.00	0.05	8.57	6.04	1.62	100.01	39.99
06XJ153 19	М		60.54		24.25		0.19	0.02	0.02	6.18	8.46	0.21	99.89	28.42
06XJ153 22	Μ		66.03		20.90	0.05	0.25	0.04	0.00	1.74	10.98	0.14	100.15	7.98
06XJ153 23	Μ		64.67		21.42		0.06	0.01	0.01	2.49	9.80	0.94	99.46	11.67
06XJ153 24	Μ		60.21	0.03	23.70		0.28		0.01	6.66	8.32	0.32	99.63	30.14
06XJ153 25	Μ		62.43		23.00		0.06	0.03	0.03	4.39	9.56	0.17	99.68	20.06
06XJ153 26	Р	R	56.74	0.06	26.20		1.07	0.04	0.15	8.25	6.43	0.56	99.52	40.12

2

R-Rim; C-Core; M-Matrix; -below detection limit

ore-porphyries

Sample	06XJ-137	06XJ-138-1	06XJ-138-2	06XJ-139	06XJ-140	06XJ-141	06XJ-142	06XJ-143	06XJ-144	06XJ-145	06XJ-147	06XJ-148
Position					WDH-N	No.2					WDH	I-No.5
SiO2	56.91	54.02	54.76	57.02	52.46	64.95	52.66	50.24	53.75	60.35	60.55	58.36
TiO2	0.94	0.91	0.80	0.86	0.87	0.54	1.27	1.03	0.90	0.72	0.69	0.73
Al2O3	18.32	17.33	17.62	17.02	16.35	15.96	16.57	15.73	16.26	17.75	16.08	16.03
FeO ^{Total}	6.51	6.75	6.23	6.26	7.44	4.28	8.38	8.56	7.00	3.21	3.44	3.50
MnO	0.08	0.10	0.09	0.09	0.12	0.04	0.15	0.14	0.11	0.06	0.06	0.07
MgO	3.39	5.01	4.72	4.20	7.20	2.23	6.10	8.46	7.28	3.93	4.78	5.84
CaO	5.80	7.83	7.36	6.84	6.83	3.84	6.65	8.01	7.27	6.24	7.13	7.37
Na2O	4.65	3.93	3.89	3.67	3.44	4.31	3.71	3.04	3.79	5.31	4.70	4.48
K2O	1.26	1.24	1.33	1.46	1.91	1.29	1.79	1.02	1.15	0.57	0.60	1.23
P2O5	0.05	0.04	0.04	0.04	0.07	0.05	0.05	0.05	0.06	0.07	0.05	0.04
LOI	1.56	1.55	1.92	1.53	2.72	1.57	1.82	2.53	1.89	1.70	1.05	1.45
Total	100.19	99.48	99.47	99.70	100.23	99.53	100.08	99.77	100.23	100.27	99.51	99.49
$Mg^{\#}$	48.1	56.9	57.4	54.5	63.3	48.2	56.5	63.8	64.9	68.6	71.2	74.9
Sc	8.38	18.4	17.2	18.0	14.7	5.10	12.8	22.2	16.0	12.2	13.2	15.0
V	109	175	151	129	173	93.3	189	187	153	97.2	127	149
Cr	22.7	77.6	87.7	88.2	274	39.6	138	353	291	57.5	152	171
Co	18.8	22.8	21.2	20.7	27.4	11.8	20.7	33.8	27.4	9.28	8.92	8.26
Ni	34.8	40.6	40.4	39.3	133	25.2	61.1	153	132	44.4	62.0	72.0
Ga	17.8	18.0	18.4	17.5	16.9	15.9	15.8	17.4	16.7	17.0	17.0	17.4
Cs	1.58	0.902	1.33	1.13	1.27	1.15	0.979	1.23	1.07	1.14	1.66	1.93
Rb	32.5	31.2	33.7	30.5	42.0	38.5	18.8	30.4	27.2	15.8	38.1	81.2
Ba	337	401	397	499	423	407	243	222	451	241	247	216
Th	1.70	1.35	1.81	2.60	1.07	4.96	0.715	1.03	1.43	2.58	3.21	2.72
U	0.410	0.370	0.481	0.539	0.341	1.15	0.439	0.256	0.348	1.01	1.07	0.822
Pb	3.00	4.18	4.69	4.06	3.13	3.33	1.40	1.60	2.18	2.58	2.16	1.33
Nb	1.91	2.95	2.96	3.32	2.32	3.01	2.50	2.78	2.39	2.71	3.35	2.64
Та	0.172	0.208	0.204	0.251	0.164	0.267	0.187	0.205	0.224	0.228	0.244	0.210
Sr	623	603	614	581	508	487	401	445	575	716	561	540
Y	15.5	16.5	15.3	13.8	13.1	11.8	13.1	19.8	14.5	14.8	15.4	15.9
Zr	144	91.2	96.7	105	102	158	98.5	88.6	110	125	128	101
Hf	3.62	2.46	2.60	2.63	2.59	4.40	2.58	2.51	2.77	3.49	3.62	2.98
La	5.70	7.19	7.22	7.68	7.25	8.33	4.42	5.24	6.18	9.30	11.0	6.44
Ce	14.3	16.6	17.2	17.7	17.3	17.6	12.7	14.5	14.9	24.8	27.8	20.5
Pr	2.13	2.56	2.52	2.42	2.46	2.27	1.95	2.46	2.43	3.54	3.78	3.29
Nd	9.75	11.6	11.0	10.6	10.8	9.38	9.71	12.1	10.9	14.8	15.3	15.1
Sm	2.49	2.88	2.66	2.49	2.59	2.09	2.58	3.21	2.63	3.17	3.27	3.42
Eu	0.887	0.907	0.857	0.881	0.789	0.666	0.777	1.00	0.820	0.877	0.933	0.854
Gd	2.44	2.85	2.68	2.47	2.45	1.91	2.53	3.36	2.53	2.92	3.00	3.27
Tb	0.458	0.504	0.472	0.395	0.428	0.344	0.465	0.615	0.454	0.482	0.507	0.533
Dy	2.73	3.00	2.84	2.41	2.54	2.00	2.80	3.58	2.66	2.75	2.86	2.94
Ho	0.547	0.622	0.580	0.492	0.522	0.412	0.573	0.751	0.542	0.553	0.578	0.592
Er	1.52	1.64	1.63	1.39	1.48	1.22	1.63	2.05	1.49	1.59	1.56	1.62
Tm	0.222	0.235	0.237	0.203	0.203	0.177	0.229	0.284	0.221	0.229	0.226	0.233
Yb	1.48	1.60	1.49	1.42	1.41	1.24	1.49	1.95	1.43	1.55	1.48	1.59
Lu	0.232	0.247	0.235	0.216	0.209	0.207	0.230	0.284	0.209	0.239	0.231	0.233
Sr/Y	40.2	36.5	40.0	42.3	38.8	41.1	30.7	22.5	39.8	48.2	36.5	34.0
Nb/Ta	11.1	14.2	14.5	13.2	14.1	11.3	13.4	13.6	10.7	11.9	13.7	12.6

.

10 Appendix 3 Continued

9

Position WDH-No.5 East KGSY West KGSY SiO2 60.45 58.20 59.78 57.77 59.51 65.66 65.31 58.92 54.87 55 TiO2 0.74 0.67 0.71 0.77 0.68 0.44 0.43 0.67 0.85 Al2O3 18.17 17.24 17.22 17.53 17.33 16.59 16.31 17.57 18.26 14 FeO ^{Total} 2.77 5.70 5.50 6.00 5.08 3.77 3.60 5.50 6.55	56.53 0.66 15.59 7.90 0.13 7.75 6.78 2.91 1.72 0.15
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	56.53 0.66 15.59 7.90 0.13 7.75 6.78 2.91 1.72 0.15
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	0.66 15.59 7.90 0.13 7.75 6.78 2.91 1.72 0.15
Al2O3 18.17 17.24 17.22 17.53 17.33 16.59 16.31 17.57 18.26 1 FeO^{Total} 2.77 5.70 5.50 6.00 5.08 3.77 3.60 5.50 6.55 5.50 MpO 0.04 0.08 0.10 0.11 0.09 0.02 0.07 0.12 0.11	15.59 7.90 0.13 7.75 6.78 2.91 1.72 0.15
FeO ^{Total} 2.77 5.70 5.50 6.00 5.08 3.77 3.60 5.50 6.55 MpO 0.04 0.08 0.10 0.11 0.00 0.02 0.12 0.11	7.90 0.13 7.75 6.78 2.91 1.72 0.15
$M_{\rm PO} = 0.04 = 0.09 = 0.10 = 0.11 = 0.09 = 0.07 = 0.12 = 0.11$	0.13 7.75 6.78 2.91 1.72 0.15
$\frac{1}{100} 0.04 0.06 0.10 0.11 0.09 0.08 0.07 0.12 0.11$	7.75 6.78 2.91 1.72 0.15
MgO 3.96 3.71 3.26 3.66 3.00 1.44 1.44 3.38 3.98	6.78 2.91 1.72 0.15
CaO 7.48 6.23 5.18 2.90 4.57 1.80 2.51 3.85 5.98	2.91 1.72 0.15
Na2O 4.52 3.78 3.82 4.36 4.56 4.61 4.00 4.23 3.46	1.72 0.15
K2O 0.76 1.82 1.74 1.61 1.81 2.07 2.02 1.75 1.16	0.15
P2O5 0.04 0.04 0.08 0.11 0.07 0.09 0.12 0.14 0.19	
LOI 0.98 1.38 2.03 4.09 2.30 2.86 3.45 3.24 4.02	
Total 100.23 99.49 100.03 99.57 99.56 99.83 99.67 99.96 100.16 1	00.57
Mg [#] 71.8 53.7 51.3 52.1 51.3 40.5 41.6 52.3 52.0	68.8
Sc 14.0 14.3 11.7 12.7 10.6 1.91 3.46 10.1 12.9	21.7
V 155 145 134 143 115 65.0 59.5 126 171	
Cr 47.6 67.5 42.5 65.2 23.9 5.84 10.8 63.1 51.7	452
Co 5.87 18.7 17.5 20.4 16.3 4.64 4.30 15.5 21.6	36.1
Ni 41.8 55.8 33.5 42.8 32.0 9.07 6.38 42.4 51.6	176
Ga 17.6 18.5 18.3 18.0 17.7 16.6 16.9 18.1 18.3	
Cs 2.15 4.51 1.07 0.925 0.337 0.917 0.886 0.473 1.44	5.29
Rb 46.7 58.0 25.6 27.9 27.9 34.9 54.0 40.0 23.5	71.3
Ba 230 525 784 759 627 648 363 648 590	293
Th 2.18 2.84 2.47 1.76 2.33 2.31 2.96 1.45 2.24	4.61
U 0.850 0.664 0.755 0.666 0.914 0.679 0.781 0.468 0.575	2.21
Pb 2.37 3.54 5.77 2.46 2.91 7.94 4.21 3.56 2.49	15.2
Nb 2.47 2.41 2.26 1.99 2.27 2.36 2.31 1.50 2.15	4.93
Ta 0.182 0.188 0.171 0.148 0.171 0.209 0.186 0.128 0.141 ().447
Sr 628 577 769 523 624 346 387 637 841	297
Y 16.4 13.4 13.7 12.1 12.9 9.51 10.7 9.56 12.6	15.7
Zr 94.3 62.4 111 95.0 111 125 126 78.0 75.5	97.9
Hf 2.73 2.27 3.39 2.85 3.18 3.49 3.38 2.21 2.19	3.05
La 7.05 8.71 9.94 6.98 9.44 9.68 11.1 8.03 13.4	13.5
Ce 23.4 19.6 21.6 16.6 20.3 18.9 22.8 18.0 27.8	28.8
Pr 3.82 2.76 2.97 2.32 2.73 2.61 2.96 2.43 4.19	3.45
Nd 16.9 11.9 12.3 10.1 11.4 9.97 11.8 10.6 17.9	13.4
Sm 376 2.63 2.62 2.35 2.44 1.89 2.20 2.21 3.53	2.91
Eu 1.05 0.787 0.840 0.783 0.800 0.582 0.635 0.700 1.06 () 891
Gd 3.24 2.46 2.52 2.28 2.34 1.69 1.97 2.08 3.02	2.62
Th 0.539 0.431 0.443 0.390 0.394 0.270 0.317 0.316 0.444 () 454
Dv 319 250 248 230 233 153 178 176 237	2 47
$H_0 = 0.623 = 0.503 = 0.513 = 0.467 = 0.483 = 0.337 = 0.359 = 0.371 = 0.483 = 0.0000000000000000000000000000000000$) 501
Fr = 1.71 = 1.42 = 1.45 = 1.29 = 1.42 = 1.00 = 1.07 = 0.947 = 1.29	1.46
Tm 0.241 0.206 0.211 0.201 0.211 0.156 0.163 0.147 0.179 0.179	1.10
Yh 1.59 1.38 1.46 1.28 1.36 1.08 1.14 0.027 1.15	1 57
$L_{\rm H} = 0.239 = 0.199 = 0.215 = 0.198 = 0.219 = 0.172 = 0.193 = 0.143 = 0.171 = 0.0000000000000000000000000000000000$) 240
Sr/Y = 38.4 = 43.0 = 56.1 = 43.1 = 48.4 = 36.4 = 36.0 = 66.7 = 66.0	19.0
Nb/Ta 13.6 12.8 13.2 13.5 13.3 11.3 12.4 11.7 15.3	11.0

The value for a sanukitoid is the average of the high-Mg andesites from Setouchi

12 Volcanic Belt, Japan (after Tatsumi and Ishizaka, (1981; 1982); Shimoda et al., 2006)

Sample	T(Ma)	⁸⁷ Rb/ ⁸⁶ Sr	⁸⁷ Sr/ ⁸⁶ Sr	2σ	⁸⁷ Sr/ ⁸⁶ Sr i	¹⁴⁷ Sm/ ¹⁴⁴ Nd	¹⁴³ Nd/ ¹⁴⁴ Nd	2σ	εNd(t)	T _{Nd2DM}	f _{Sm/Nd}	¹⁷⁶ Lu/ ¹⁷⁷ Hf
06XJ-137	315	0.1510	0.703976	18	0.7033	0.1555	0.512978	8	8.3	402	-0.21	0.00909
06XJ-138-2	315	0.1590	0.704403	11	0.7037	0.1473	0.512872	8	6.5	546	-0.25	0.01283
06XJ-140	315	0.2394	0.705051	18	0.7040	0.1454	0.512873	10	6.6	538	-0.26	0.01142
06XJ-141	315	0.2290	0.704566	14	0.7035	0.1358	0.512830	16	6.2	574	-0.31	0.00666
06XJ-143	315	0.1973	0.704297	15	0.7034	0.1619	0.512965	15	7.8	445	-0.18	0.01602
06XJ-145	315	0.0639	0.704146	17	0.7039	0.1301	0.512810	7	6.0	587	-0.34	0.00969
06XJ-147	310	0.1964	0.704536	14	0.7037	0.1305	0.512824	9	6.3	566	-0.34	0.00903
06XJ-148	310	0.4346	0.705599	17	0.7037	0.1374	0.512830	11	6.1	580	-0.30	0.01109
06XJ-153	314	0.0961	0.704171	17	0.7037	0.020	0.512798	11	5.8	606	-0.34	0.00897
06XJ-154	314	0.1544	0.704669	14	0.7040	0.032	0.512848	9	6.3	564	-0.28	_
06XJ-156	307	0.2914	0.705185	14	0.7039	0.061	0.512828	8	6.9	512	-0.41	0.00699
06XJ-158	307	0.1817	0.704943	18	0.7041	0.038	0.512862	8	7.1	493	-0.35	0.00918
Sample	¹⁷⁶ Hf/ ¹⁷⁷ Hf	2σ	εHf(t)	f _{Lu/Hf}	T _{HfDM2}	$\Delta \varepsilon Hf(t)$	²⁰⁶ Pb/ ²⁰⁴ Pb	²⁰⁷ Pb/ ²⁰⁴ Pb	²⁰⁸ Pb/ ²⁰⁴ Pb	206Pb/204Pbt	²⁰⁷ Pb/ ²⁰⁴ Pbt	²⁰⁸ Pb/ ²⁰⁴ Pbt
06XJ-137	0.28307	7	15.7	-0.73	325	1.2	18.408	15.446	37.963	17.913	15.420	37.316
06XJ-138-2	0.28305	8	14.0	-0.61	433	2.3	18.278	15.459	37.929	17.907	15.440	37.489
06XJ-140	0.28309	8	15.7	-0.66	325	3.8	18.309	15.485	38.026	17.914	15.464	37.634
06XJ-141	0.28302	7	14.3	-0.80	415	3.1				—	—	_
06XJ-143	0.28310	7	15.0	-0.52	365	1.4	18.518	15.486	38.223	17.964	15.466	37.521
06XJ-145	0.28304	8	14.3	-0.71	409	3.5	19.351	15.534	38.574	17.899	15.457	37.405
06XJ-147	0.28304	7	14.5	-0.73	396	3.3	19.833	15.533	39.378	17.975	15.435	37.608
06XJ-148	0.28304	7	14.1	-0.67	425	3.1				_	—	_
06XJ-153	0.28305	7	14.8	-0.73	378	4.3	18.364	15.461	38.043	17.890	15.436	37.553
06XJ-154	_	—	_		—	—	18.744	15.463	38.331	17.753	15.411	37.503
06XJ-156	0.28304	9	14.8	-0.79	376	2.5	18.365	15.471	38.052	18.055	15.455	37.720
06XJ-158	0.28302	8	13.5	-0.72	456	0.9	18.317	15.456	37.962	17.842	15.431	37.497

Appendix 4 Sr, Nd, Hf and Pb isotopic compositions of the Baogutu intrusive rocks

15 The 147 Sm/ 144 Nd and 143 Nd/ 144 Nd ratios at the present time are 0.1967 and 0.512638 for chondrite, and 0.222 and 0.51315 for depleted mantle,

16 respectively. $\lambda = 6.54 \times 10^{-12}$ a⁻¹; The ¹⁷⁶Hf/¹⁷⁷Hf and ¹⁷⁶Lu/¹⁷⁷Hf ratios at the present time are 0.282772 and 0.0332 for chondrite, and 0.28325 and

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0.0384 for depleted mantle.

Appendix 5. (a) Histogram of plagioclase anorthite contents for the Baogutu adakitic
rocks. (b) Back-scattered electron image of plagioclase. (c-d) Large plagioclase crystals
may occasionally partly or totally enclose some clinopyroxene grains in the Baogutu
adakitic rocks. Plag-Plagioclase; Cpx-clinopyroxene.

