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4. REFERENCES CITED

1. Analytical Methods

1.1. LA-ICP-MS zircon U-Pb dating

Zircons were separated from pulverized rock by conventional heavy liquid and magnetic techniques at the Lab of Geological Team of Hebei Province, China. Separated zircon grains from each sample were mounted in epoxy resin and polished to expose their interior. Cathodoluminescence (CL) images were obtained at the Institute of Geology, Chinese Academy of Geological Sciences (Beijing) to select suitable positions for in situ analysis. In situ U-Th-Pb isotopes of zircons were measured by LA-ICP-MS using an Agilent 7900 ICP-MS system attached to a NewWave 193 nm excimer ArF laser-ablation system at the Milma Lab of China University of Geosciences (Beijing). All spot analyses were carried out using a beam diameter of 35 μm with a repetition rate of 10 Hz and laser fluence of 4–8 J/cm². Each spot analysis incorporated a background acquisition of ~20 s (gas blank), 50 s data acquisition, and 30 s delay to wash out the previous sample and prepare the next analysis. Zircon 91500 (Wiedenbeck et al., 1995) was used as external standard to calibrate U-Pb dating. The Plešovice zircon (Sláma et al., 2008) and zircon GJ-1 (Jackson et al., 2004) were treated as unknown and used to check the reproducibility and accuracy of the calibration. In each cycle, every six spot analyses were followed by analyzing zircon 91500 twice and one Plešovice and one GJ-1 (i.e., 2 zircon 91500 + 1 Plešovice + 1 GJ-1 + 6 sample spots + 2 zircon 91500 + 1 Plešovice + 1 GJ-1). Off-line data conduction was performed using an Excel-based software ICPMSDataCal (Liu et al., 2010). Concordia diagrams and weighted mean calculations were made using Isoplot/Ex_ver3 (Ludwig, 2003).

1.2. In Situ Zircon Hf Isotope Measurements

In situ zircon Hf isotope measurements were conducted at the Institute of Geology and Geophysics, Chinese Academy of Sciences by using a Neptune multi-collector (MC)-ICP-MS with an attached 193 nm excimer ArF laser-ablation system. Lu-Hf isotope analyses were made on the same zircon grains previously analyzed for U-Pb isotopes. During analysis, a laser repetition rate of 4 Hz was used with spot sizes of 20 or 60 μm depending on zircon grain size. Raw count rates for ¹⁷²Yb, ¹⁷³Yb, ¹⁷⁵Lu, ¹⁷⁶(Hf+Yb+Lu), ¹⁷⁷Hf, ¹⁷⁸Hf, ¹⁷⁹Hf, ¹⁸⁰Hf and ¹⁸²W were collected during analysis. The detailed analytical procedure was described by Wu et al. (2006). Zircon 91500 and Mud Tank were used as external standards and were analyzed once and twice, respectively before and after every 10 analyses. Initial ¹⁷⁶Hf/¹⁷⁷Hf ratios were calculated by using the ¹⁷⁶Lu decay constant of $1.867 \times 10^{-11} \text{ yr}^{-1}$ (Söderlund et al., 2004) and measured ¹⁷⁶Lu/¹⁷⁷Hf. Chondrite Lu-Hf isotopic values (¹⁷⁶Lu/¹⁷⁷Hf = 0.0336 and ¹⁷⁶Hf/¹⁷⁷Hf = 0.282785) reported by Bouvier et al. (2008) are used to calculate $\epsilon_{\text{Hf}}(t)$ values.

1.3. Whole-rock Major and Trace Element Analysis

Major element oxides (wt.%) were measured on fused glass discs, using an X-ray fluorescence (Axios MAX) at the Wuhan SampleSolution Analytical Technology Co., Ltd., Wuhan, China (WSSATCL). Three USGS rock standards (BHVO-2, W-2a, and GSP-2) and two Chinese national rock standards (GBW07103 and GBW07316) were used for calibration. The

analytical uncertainties are generally better than 5% for all elements. Loss on ignition (LOI) was obtained by twice heating.

Whole-rock trace elements were measured by Agilent 7700e ICP-MS at the WSSATCL. Sample powder (200 mesh) were firstly dried for 12 h in an oven at 105 °C. Then 50 mg accurately weighed power was placed in a Teflon bomb followed by slowly adding in 1 ml HNO₃ and 1 ml HF. The Teflon bomb was putted in a stainless steel pressure jacket and heated to 190 °C in an oven for more than 24 h. After cooling, the Teflon bomb was opened and evaporated to incipient dryness on a 140 °C hotplate. Then 1 ml HNO₃ was added in and it was evaporated to dryness again. After evaporation, 1 ml of HNO₃, 1 ml of MQ water, and 1 ml In solution (1 ppm) used as internal standard were added and the Teflon bomb was resealed and placed in the oven for more than 12 h heating at 190 °C. At last, the final solution was diluted to 100 g in a polyethylene bottle by the addition of 2% HNO₃. Four USGS rock standards (AGV-2, BHVO-2, BCR-2, and RGM-2) were used to calibrate the elemental concentrations of samples.

2. LA-ICP-MS ZIRCON U-Pb DATING RESULTS

Nine samples were selected for LA-ICP-MS zircon U-Pb dating. The dating results are reported in Table S1 and illustrated on a concordia and weighted mean age diagram (Fig. S1). All dated samples yielded Late Triassic ages ranging from ~212 Ma to ~220 Ma. Within sample variations of zircon ages are large for most dated samples indicated by large MSWD values. The large MSWD value may indicate that the uncertainty of individual zircon age is underestimated or it may suggest that more than one age population exist, for example, antecrysts recycled from previous batches of magmatism. A thorough work on this issue is beyond the scope of this study and will be covered in a separate study. Currently, a weighted mean ²⁰⁶Pb/²³⁸U age is calculated for each sample from all concordant ages which form a continuous trend in ²⁰⁶Pb/²³⁸U age distribution diagram. However, ages obviously away from the continuous trend are not included in calculation of the weighted mean age.

3. DATA SETS AND FIGURES

Tables S1–S7 are listed in an Excel file

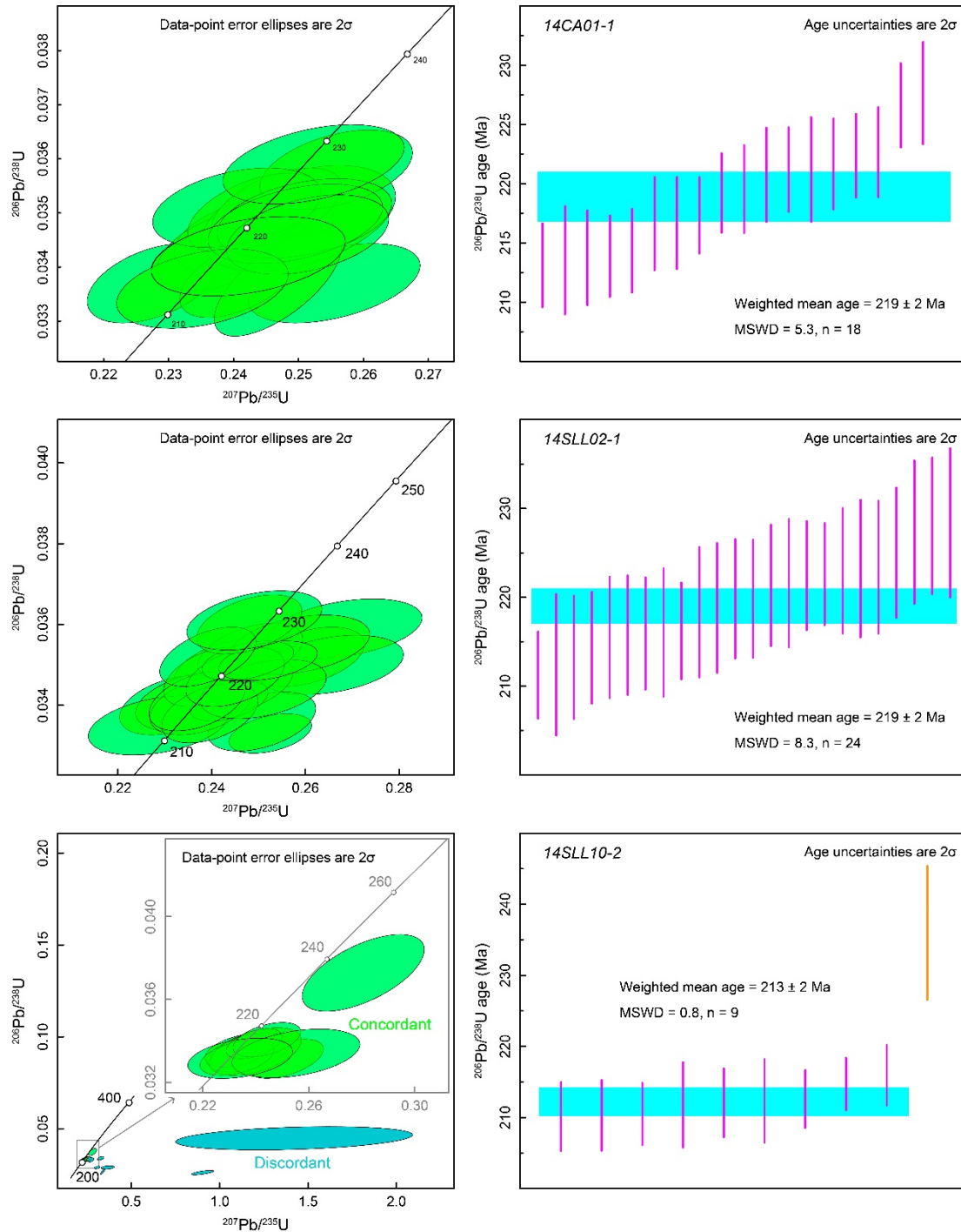


Figure S1. Zircon U-Pb concordia and weighted mean age diagram of nine samples from Late Triassic plutons in the Yidun Terrane, eastern Tibetan Plateau. Green ellipses represent concordant results with 90% to 110% concordance and blue ellipses represent discordant results. Weighted mean age with 2σ uncertainty is represented by light blue box and is calculated from concordant ages which form a continuous trend (pink lines) in the $^{206}\text{Pb}/^{238}\text{U}$ age distribution. Ages away from the continuous trend (brown lines) are not included in the calculation of the mean age.

Figure S1. *Continued.*

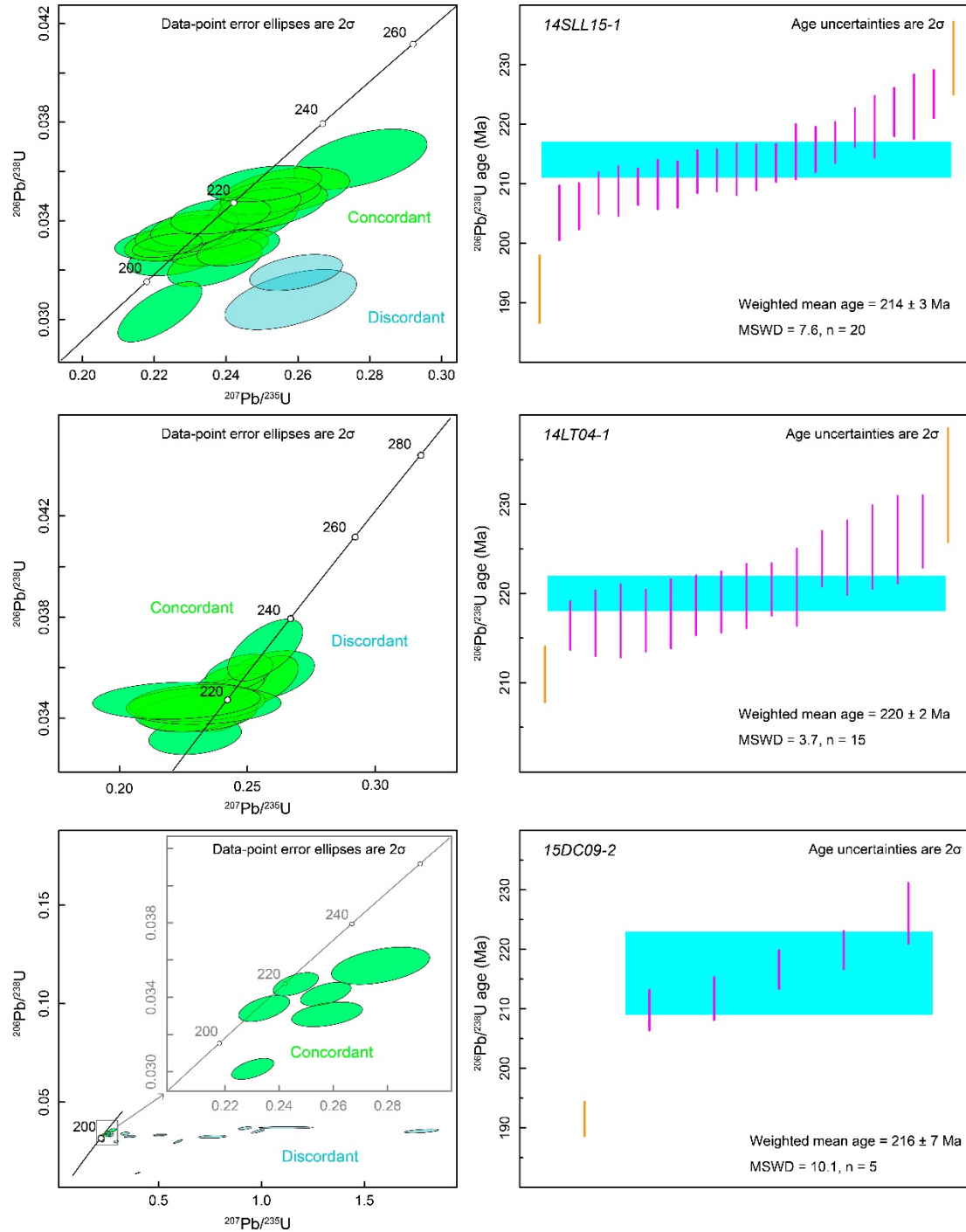


Figure S1. *Continued.*

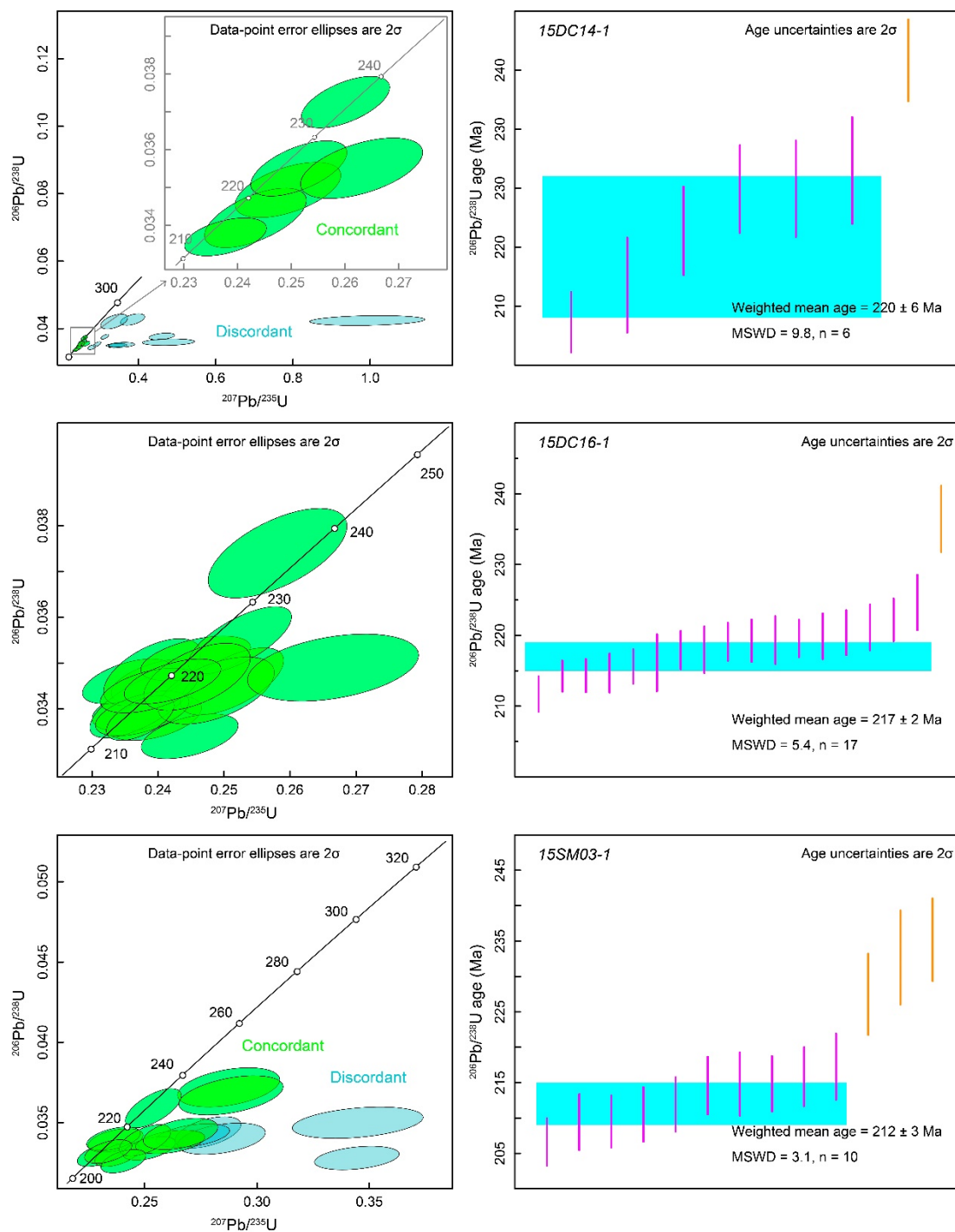


Figure S1. Continued.

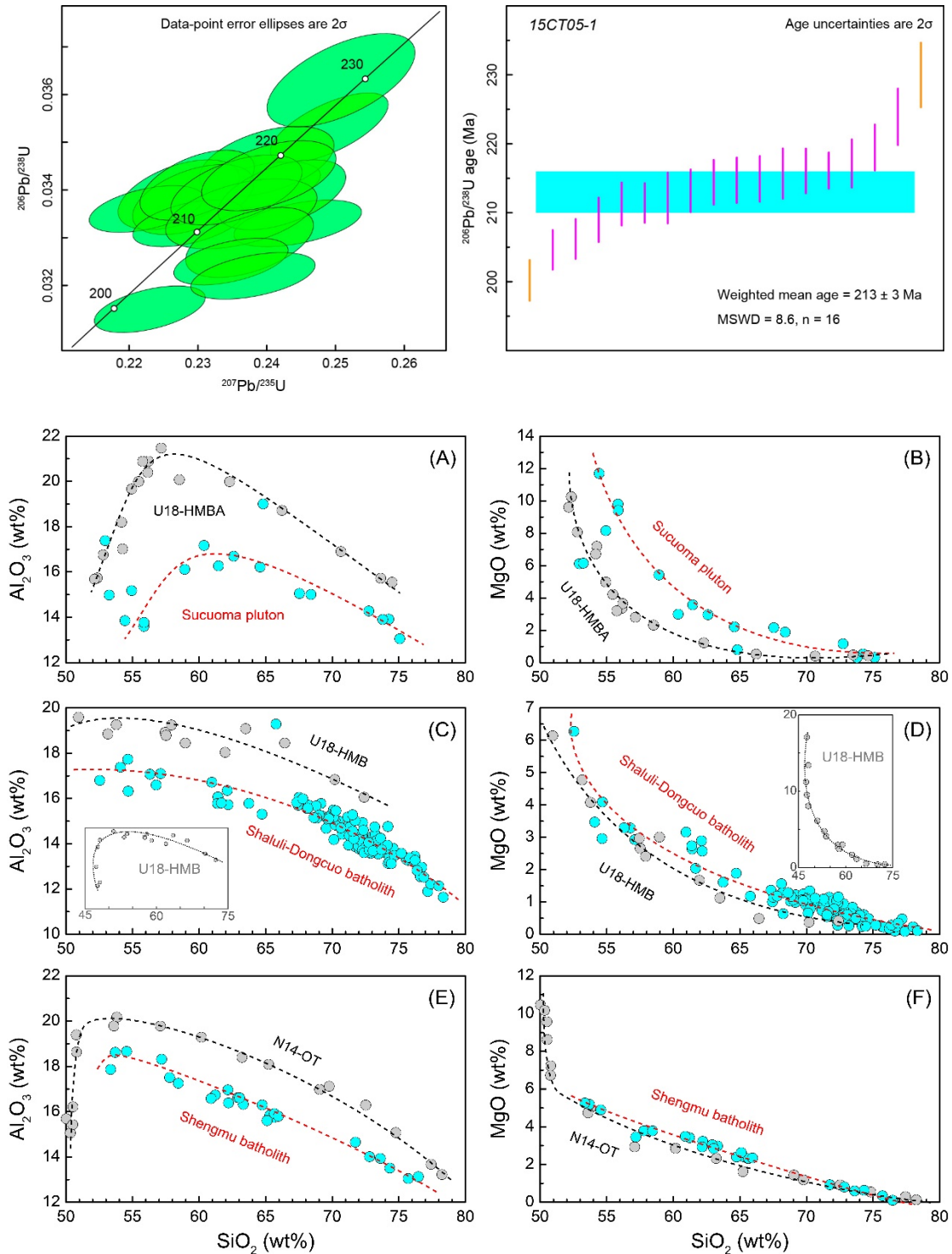


Figure S2. Whole-rock Al_2O_3 and MgO versus SiO_2 contents of samples from the Sucuoma pluton (A, B), Shaluli-Dongcuo batholith (C, D), and Shengmu batholith (E, F) in the Yidun Terrane. Red dashed lines depict the interpreted liquid line of descent (LLD) of the above plutons. Black dashed lines are LLD of hydrous fractional crystallization of a high-Mg basaltic andesite (U18-HMBA) at 1.0 GPa (Ulmer et al., 2018), a high-Mg basalt (U18-HMB) at 1.0 GPa

(Ulmer et al., 2018), and an olivine tholeiite (N14-OT) at 0.7 GPa (Nandedkar et al., 2014). Only part of the LLD of U18-HMB is shown in C and D due to space limitation and the insets in C and d show the intact LLD of U18-HMB. The LLD in these experiments are shown here to reveal that fractional crystallization of mantle derived primitive magmas producing intermediate to silicic magmas will result in curvatures in certain major element systematics. The position of kinks of the LLD is not fixed at given SiO_2 and is related to the composition of parental magmas, crystallization pressure and temperature, and fractionation assemblages. Note that the LLD of N14-OT at the part of $\text{SiO}_2 > 52$ wt% is nearly a straight line. See text for discussion.

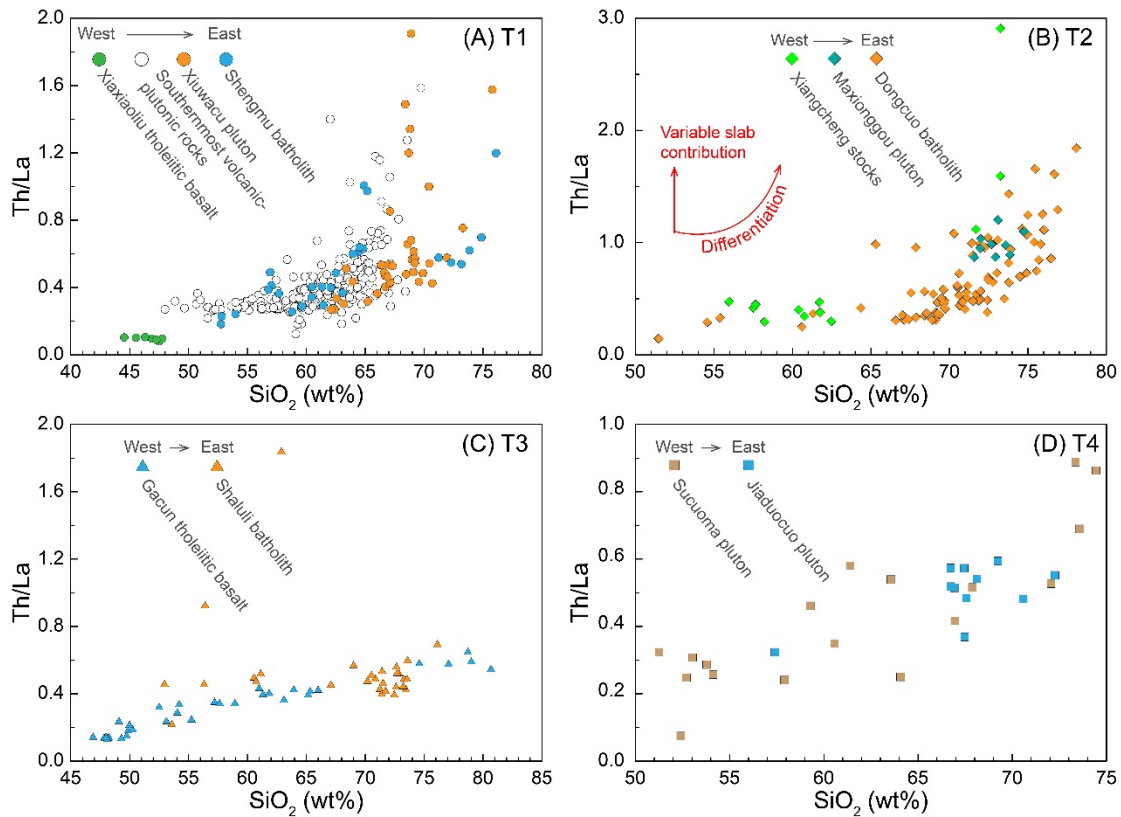


Figure S3. Th/La versus SiO_2 diagram for Late Triassic magmatic rocks along four transects (T1–T4) across the Yidun Terrane. See Figure 1 in the text for locations of the four transects. The Th/La ratio of mafic rocks is a proxy for subduction slab contribution to mantle source (Plank, 2005). Note that mafic rocks derived from variably enriched mantle sources will show different Th/La ratios at given SiO_2 content and magma crystallization differentiation will result in increase of Th/La ratios with increase of SiO_2 (see the different trends depicted in B). From west to east in each transect, mafic rocks show relatively constant rather than increase in Th/La ratios, indicating that west to east enriched in isotopic compositions did not result from higher degree of subduction-related alteration of the lithospheric mantle in the east. See text for discussions.

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