

1 **Tectonic implications of Palaeoproterozoic anatexis and Late Miocene**
2 **metamorphism in the Lesser Himalayan Sequence, Sutlej Valley, NW India**
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16 Number of words of text 5753

17 Number of references 63

18 Number of tables 4

19 Number of figures 9
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38 TECTONICS OF THE METAMORPHIC CORE, NW INDIA

39 **Abstract:** Unravelling the kinematic evolution of orogenic belts requires that the defining
40 tectono-stratigraphic units, and structural elements that bound them, are properly identified
41 and characterized. In the Sutlej Valley (NW Himalaya), the Muniari and Vaikrita thrusts
42 have both been correlated with the Main Central Thrust. The sequence of amphibolite-grade
43 rocks (the Jutogh Group) bounded by these faults has been variously assigned to the Lesser
44 Himalayan Sequence (based on provenance ages) and to the Greater Himalayan Sequence
45 (from their metamorphic grade). Trace-element and geochronological data from leucogranites
46 in the Jutogh Group (i) indicate crustal melting at *c.* 1810 Ma, before the deposition of the
47 Greater Himalayan Sequence, thus correlating the Jutogh Group with the Lesser Himalayan
48 Sequence, and (ii) record Proterozoic metamorphism overprinted at 10.5 ± 1.1 Ma
49 (established from U–Pb analysis of uraninite) during the Himalayan orogeny. Pressure-
50 temperature-time data affirm that the Jutogh Group and Greater Himalayan Sequence
51 represent distinct tectonic units of the metamorphic core that were decoupled during their
52 extrusion. This precludes extrusion along a single, widening channel, and requires a
53 southward shift of the locus of movement during the Late Miocene, coincident with present-
54 day precipitation patterns.

55

56 **(end of abstract)**

57 Key litho-tectonic units in the Himalaya e.g. the Greater Himalayan Sequence, and the major
58 faults that bound them e.g. the Main Central Thrust, can be traced continuously along the
59 2400 km strike of the orogen (Fig. 1). Understanding the metamorphic evolution of such units
60 together with the recognition of the nature, location and timing of the principle structural
61 elements of an orogenic belt is fundamental both to defining the tectonic architecture of that
62 belt and to understanding the mechanical behaviour of continental crust during its
63 deformation.

64

65 The Greater and Lesser Himalayan Sequences are two key litho-tectonic units in the
66 Himalaya. Using provenance studies based on detrital zircon ages (DeCelles *et al.* 2000;
67 Richards *et al.* 2005) and Nd isotopic compositions (Parrish & Hodges 1996; Ahmad *et al.*
68 2000; Robinson *et al.* 2001; Richards *et al.* 2005) the sequences are stratigraphically defined;
69 the deposition age of the Lesser Himalayan Sequence is Palaeoproterozoic to
70 Mesoproterozoic (*c.* 2500 to 1000 Ma), in contrast with the younger Greater Himalayan
71 Sequence (synonymous with the High Himalayan Crystalline Series), deposited in
72 Neoproterozoic to Cambrian times (*c.* 800 to 500 Ma). The sequences represent two laterally
73 continuous basinal sequences deposited on the Indian passive margin (Le Fort 1975), and
74 prior to continental collision may have been separated by either a ‘proto-Main Central Thrust’
75 lineament or the Himalayan Unconformity (Goscombe *et al.* 2006).

76

77 In general, the crystalline core of the Himalayas (the Greater Himalayan Sequence) was thrust
78 southward over the lower grade Lesser Himalayan Sequence on the Main Central Thrust. This
79 major ductile thrust is a key component in all tectonic reconstructions of the orogen (e.g. Yin
80 2006 and references therein) yet its characteristics, significance and specific location have
81 challenged Himalayan geologists for many decades (Gansser 1964; Le Fort 1975; Hodges
82 2000).

83

84 One reason for this uncertainty is that the crustal architecture of the orogen is complicated in
85 some areas by at least two major thrust faults, one or both of which have been referred to as
86 the Main Central Thrust using various criteria (reviewed in Yin 2006), hindering correlation
87 of the adjacent tectonic units. Such is the case for the Sutlej Valley of NW India (Fig. 2).
88 Although lithologies assigned to the Lesser Himalayan Sequence are typically
89 unmetamorphosed, or at most are metamorphosed to chlorite or biotite metamorphic grade,
90 the Sutlej Valley exposes a more complex transect within which a crystalline zone, the Jutogh
91 Group, is assigned by some authors to the Lesser Himalayan Sequence, and termed the Lesser
92 Himalayan Crystalline Sequence (Vannay *et al.* 1999; Thiede *et al.* 2004). This is supported
93 by a marked contrast in isotope geochemistry, pressure–temperature–time paths, garnet
94 morphologies and monazite ages between these rocks and the Greater Himalayan Sequence
95 (Catlos *et al.* 2001; Kohn *et al.* 2004; Richards *et al.* 2005; Caddick *et al.* 2006). Other
96 authors, using metamorphic grade as the primary criterion for recognising Lesser Himalayan
97 Sequence lithologies and thus the Main Central Thrust (e.g. Sharma 1977; Singh *et al.* 2006),
98 assign the crystalline rocks to the Greater Himalayan Sequence.

99

100 In this study of rocks of the Sutlej Valley we clarify tectono-stratigraphic relationships in this
101 area by a study of leucogranite emplaced into the enigmatic Jutogh Group. We use trace-
102 element data to establish the mode of melting for the leucogranites, and U–Pb dating of
103 accessory phases to a) constrain the ages of both melting and protolith formation, and b)
104 identify a later recrystallization event. In comparing the results with those for the well-
105 documented High Himalayan leucogranites, which intruded high-grade rocks unambiguously
106 assigned to the Greater Himalayan Sequence, we establish the true affinity of the Jutogh
107 Group. Finally, we consider the tectonic evolution of the metamorphic core of the Sutlej
108 Valley with regards to recent thermo-mechanical orogenic modelling, and to the possible
109 feedback mechanism between climate and tectonics.

110

111 **Field relations and petrology**

112 In the Sutlej Valley, a zone of metasediments and orthogneiss separates the high-grade
113 metasediments of the Vaikrita Group (Valdiya 1988) of undisputed Greater Himalayan
114 Sequence affinity from the greenschist-grade sediments of the Rampur window of clear
115 Lesser Himalayan Sequence affinity (Fig. 2, 3). This enigmatic zone is often viewed as part of
116 the Greater Himalayan Sequence (e.g. Singh & Jain 1993) but has also been mapped as the
117 Lesser Himalayan Crystalline Sequence (e.g. Vannay *et al.* 1999). These crystalline rocks,
118 termed the Jutogh Group, include the Wangtu Gneiss Complex and the amphibolite-grade
119 Jutogh (or Jeori) metasediments (Singh & Jain 1993; Vannay *et al.* 2004; Richards *et al.*
120 2005) (Fig. 3).

121

122 There is a gradual transition down-section from the main 1.87 Ga Wangtu orthogneiss
123 (Richards *et al.* 2005), through intercalated orthogneiss, paragneiss and calc-silicate rocks,
124 into the metasedimentary gneisses and mica schists. If the relationship between the gneiss and
125 the metasediments was originally intrusive it is now obscured by subsequent deformation and
126 tectonism, including a 50 to 80 m wide thrust zone which strikes broadly NW–SE between the
127 town of Sarahan and the Sutlej River, here termed the Sarahan Thrust (Figs. 2, 3), and
128 probably equivalent to the Chaura Thrust (Jain *et al.* 2000). This thrust zone is marked by a
129 tectonic *mélange* (locality 55) of sub-rounded clasts (several centimetres to tens of metres
130 across) of fine-grained, mafic amphibolite gneiss in a friable, sheared biotite–chlorite matrix
131 with top-to-the-SE kinematic indicators (Figs. 3, 4). A strongly sheared garnet–amphibolite
132 exposed on the banks of the Sutlej river near Jeori (Fig. 1, locality 83, and Fig. 2 in Singh &
133 Jain 1993) is interpreted to be the SW continuation of the Sarahan Thrust, where the scarcity
134 of pelitic material has inhibited the formation of a *mélange* as described above.

135

136 In the immediate hanging wall of the Sarahan Thrust, kink and chevron-type folds in the
137 Jutogh metasediments verge consistently to the south implying that they are related to the top-
138 to-the-south motion on the Sarahan Thrust. Within the Wangtu Gneiss stretching lineations

139 change orientation progressively from west-plunging in the core of the complex (probably
140 representing a pre-existing lineation orientation) to north- or NNE-plunging in both the
141 hanging wall and footwall of the Sarahan Thrust (Fig. 2), suggesting a genetic relationship
142 with the south-directed thrusting.

143

144 Jain *et al.* (2000) show that the Sarahan (Chaura) Thrust marks a sharp discontinuity in apatite
145 and zircon fission track data, indicating faster exhumation of the Wangtu Gneiss Complex
146 (hanging wall) compared to the Jutogh metasediments (footwall) during the Plio-Pleistocene.
147 However, this contrast in exhumation rates is not clear from other fission track studies based
148 in the Sutlej Valley, nor does it appear to be reflected in muscovite cooling age profiles
149 (Thiede *et al.* 2004; Vannay *et al.* 2004; Thiede *et al.* 2005). Consequently, this suggests that
150 the Sarahan Thrust does not represent a major exhumation structure (i.e. there is no evidence
151 for decoupling of the Jutogh metasediments and the Wangtu Gneiss Complex during their
152 exhumation), and we consider the Jutogh Group as a complete litho-tectonic package.

153

154 Metamorphic grade increases up-section through the Jutogh metasediments from garnet to
155 staurolite (Fig. 3). At the structurally lowest level in the Jutogh Group a quartz-rich kyanite–
156 chlorite–muscovite schist (locality 90, Fig. 2) suggests upper-greenschist conditions of
157 metamorphism, according to the dehydration reaction of pyrophyllite, restricted to high-Al
158 pelites, relative to alkalis (Miyashiro 1994). However, although Pant *et al.* (2006) map
159 kyanite schist directly in the hangingwall to the Munsiri Thrust (above the Rampur
160 Window), we are not so confident about the structural position of the kyanite schists, and
161 herein consider the lowest grade of the Jutogh Group to be garnet. Vannay *et al.* (1999, Fig.
162 1) identified both sillimanite and kyanite in what we here recognize as the hanging wall of the
163 Sarahan Thrust, i.e. in the Wangtu Gneiss Complex. Oxygen isotope studies through this
164 inverted metamorphic field gradient (garnet to sillimanite) suggest a modest increase in
165 recorded temperature, but a *decrease* in recorded pressure up-section, implying that the

166 inverted metamorphic field gradient reflects diachronous mineral growth rather than an
167 inverted geotherm at any one time (Vannay *et al.* 1999).

168

169 Hitherto unrecognised leucogranites in the Jutogh metasediments south of Sarahan have been
170 observed at three localities (63, 70 and 71, Figs. 2 and 3). At localities 63 and 70 the
171 leucogranite bodies form medium to coarse-grained boudins, 1 to 2 m in length, and aligned
172 with the country rock foliation (Fig. 5), whereas at locality 71 deformed, concordant cm-scale
173 fine- to medium-grained leucocratic veins are common. Importantly, there is no evidence for
174 post-tectonic granites intruding the Jutogh Group. The mineralogy of the leucogranites is
175 quartz, alkali-feldspar, plagioclase, muscovite; accessory phases include tourmaline, zircon,
176 uraninite and titanite. Tourmaline forms abundant prisms up to 2 cm across at localities 70
177 and 71. In thin section tourmaline is characterised by a network of colour changes associated
178 with numerous annealed microcracks suggesting alteration by fluid infiltration. High fluid
179 pressure during deformation is indicated by quartz-filled fractures in feldspars whilst
180 deformation lamellae in quartz support a relatively low temperature (*c.* 300 to 400 °C)
181 deformation regime (Passchier & Trouw 1998). Leucogranite margins are undulose but sharp,
182 and interpreted as originally intrusive (rather than formed *in situ*) due to the lack of significant
183 biotite-rich selvages. However, the small size of the Jutogh leucogranites suggests they did
184 not travel far from their source, which is therefore probably the Jutogh metasediments. No
185 evidence for contact metamorphism was observed in the country rocks so either the thermal
186 contrast between intrusion and country rock was low, or contact metamorphic textures have
187 been overprinted by subsequent mineral growth. The paragneiss that encloses the leucogranite
188 lenses contains quartz, alkali feldspar, biotite, muscovite ± plagioclase, with pre-tectonic
189 garnets (1 to 2 mm) at localities 63 and 71.

190

191 Since leucogranites of Early Miocene age are prevalent throughout much of the Greater
192 Himalayan Sequence, determining the age of these leucogranites is critical to the
193 interpretation of host unit affinity (whether to the Lesser or Greater Himalayan Sequence).

194 **Sampling and Methodology**

195 Samples of leucogranite and paragneiss were collected from localities 63 and 70 (Fig. 5) for
196 geochemical and petrological analysis. Locality 71 was sampled for petrological analysis
197 only, as the small volume of leucocratic material available precluded a robust geochemical
198 analysis.

199

200 Whole-rock major and trace element analyses were obtained on an ARL Fisons wavelength-
201 dispersive XRF spectrometer at the Open University. Trace elements were determined from
202 pressed powder pellets.

203

204 Bulk-rock paragneiss samples from localities 57 and 66 were prepared for isotopic analysis
205 following standard techniques as described in Cohen *et al.* (1988) and analysed for Nd isotope
206 data at the Open University using a Triton thermal ionization mass spectrometer (TIMS).
207 Repeat analyses of the La Jolla standard (n=17) gave $^{143}\text{Nd}/^{144}\text{Nd}$ ratios of $0.511849 \pm$
208 0.000004 (2σ) over the analysis period. Total procedural blanks were less than 10 pg.

209

210 For chronometric studies, zircon (ZrSiO_4) and uraninite (UO_2) grains were mounted into a 1
211 inch diameter epoxy resin stub following conventional separation techniques (including
212 diodomethane heavy liquid separation). Most zircons are euhedral (magmatic) and markedly
213 metamict (grey and semi-opaque), between 100 μm and 300 μm in length (Fig. 6a). Back-
214 scatter electron (BSE) imaging of zircons showed no evidence of internal zoning, but did
215 reveal two types of uraninite inclusions, one finely disseminated and the other coarser, <25
216 μm in diameter (Fig. 6c). This characteristic of uraninite in zircon may be the product of
217 exsolution but they are referred to here as inclusions. A small number of zircons separated
218 from sample 70ii were notably different in size (much smaller with an average length of 120
219 μm), shape (rounded to sub-rounded, characteristic of detrital grains) and degree of
220 metamictization (none) (Fig. 6a), and BSE imaging revealed no evidence of internal zoning
221 within them, as well as no inclusions (Fig. 6d). Discrete uraninite grains 250 μm in diameter,

222 also exclusive to sample 70ii, were affixed directly onto a resin block and not polished. One
223 was notably euhedral and lustrous, in comparison to the remainder of uraninite grains which
224 were anhedral and appeared corroded (Fig. 6b). Presumably because of a low modal
225 abundance (e.g. Thorpe *et al.* 1995), no uraninite grains were located in any thin sections
226 precluding further textural analysis.

227

228 Laser ablation multi-collector inductively coupled plasma mass spectrometry (LA–MC–ICP–
229 MS) was used in the analysis of the three phases identified above, which are 1) zircon (both
230 magmatic and detrital grains), 2) coarse uraninite inclusions in magmatic zircon, and 3)
231 discrete uraninite grains. A UP193SS New Wave Research laser ablation system was utilised
232 in conjunction with an Axiom MC–ICP–MS instrument. A 35 μm spot was used statically to
233 ablate samples of zircon using laser fluences of 1 to 2 J/cm^2 , compared to the static ablation of
234 uraninite (both discrete and included crystals) using a 10 μm spot and laser fluences of 2 to 3
235 J/cm^2 .

236

237 The analysis of zircon followed methods similar to those of Horstwood *et al.* (2003) and used
238 the 554 Ma Manangotry monazite standard for Pb/U calibration, coupled with a static ablation
239 pattern. The overall reproducibility of the standard for $^{206}\text{Pb}/^{238}\text{U}$ during the course of these
240 analyses was 4 to 6% (2σ), which has been propagated into the uncertainties for each
241 individual spot analysis. The use of a non-matrix-matched standard for the zircon analyses,
242 coupled to a static ablation protocol, could be expected to introduce a matrix effect and hence
243 Pb/U inaccuracy, on the order of a few percent. However, in this instance, concordant zircon
244 data after normalization to monazite suggest that this effect is either absent or negligible.
245 Also, for the majority of sample zircons analysed here, where small uraninite inclusions are
246 ‘peppered’ throughout, a matrix effect might well be expected even if zircon was used as a
247 standard due to the severe metamictization of the zircon structure. Due to the resulting
248 severely discordant nature of these zircons, any inaccuracy due to non-matrix matched
249 standardisation in this instance is minor and has no significant effect on the interpretation.

250

251 Analyses of uraninite, both discrete grains and coarse inclusions in zircon, were normalized to
252 uraninite crystals separated from a leucogranite near the Rongbuk monastery, South Tibet,
253 and dated by isotope dilution-TIMS, thereby providing an additional calibration control
254 relevant to matrix matching. Data from this 'standard uraninite' are available online at
255 <http://www.geolsoc.org.uk/SUP00000>. A hard copy can be obtained from the Society Library
256 (the number will be allocated by the Staff/Production Editor). Reproducibility of the 'standard
257 uraninite' (18% 2σ) was more heterogeneous than that typically expected for zircon and
258 monazite (2 to 3% 2σ) and indeed for the sample uraninite grains, reflecting a clear difference
259 between the ablation characteristics of both sample and 'standard' uraninites. However,
260 significant (*c.* 19%) difference could be seen in the relative Pb/U normalization values
261 between monazite and uraninite, so despite its poor reproducibility the TIMS-determined
262 uraninite remained the most appropriate standard. As there is no other 'uraninite standard'
263 suitable, propagated measurement uncertainties are therefore larger for dated uraninite,
264 approximately 8% 1σ for individual spot analyses.

265

266 One euhedral uraninite crystal proved concordant at *c.* 11 Ma with no common-Pb; whereas
267 four other anhedral grains had much older components with in part large common-Pb
268 corrections. Precise analysis of uraninite inclusions in zircons proved impossible owing to the
269 small size of the included uraninites leading to uraninite-zircon mixture on a 10 to 20 μm
270 scale. This, coupled with the extreme pulse-to-pulse variations in the Pb and U signals and the
271 need for a large common-Pb correction in many of these grains, resulted in complex results
272 that proved equivocal in their interpretation. Even with time-resolved analysis and a fast
273 washout laser ablation cell, some data were discarded as the results were insufficiently robust.

274

275 **Results**

276 *Granite geochemistry*

277 Major and trace element XRF data for leucogranite samples 63i, 70i and 70ii (Table 1) were
278 compared with two well-characterized generations of leucogranites intruding the Greater
279 Himalayan Sequence; the Miocene High Himalayan leucogranite sheets and plutons
280 (emplaced between 24 and 17 Ma) that have been sampled from across the Himalayan orogen
281 (Inger & Harris 1993; Hodges 2000 and references therein; Singh & Jain 2003), and a less
282 commonly recognized, but probably widespread, suite of deformed leucogranite lenses (*c.* 1
283 m thick) of Eocene (39 ± 3 Ma) age, studied in the Saraswati Valley of the Garhwal
284 Himalaya, 150 km SE of the Sutlej Valley section (Prince *et al.* 2001). Both leucogranite
285 suites provide distinctive trace-element patterns indicative of their differing conditions of
286 formation.

287

288 Major-element compositions define all three leucogranite suites as peraluminous; the Jutogh
289 leucogranites display silica compositions (74 to 76%) intermediate between those of the
290 Miocene (73 to 75%) and Eocene (75 to 77%) leucogranites. The Jutogh leucogranites show
291 depletions in Rb, Ba, Th and Zr relative to an average Miocene High Himalayan leucogranite
292 composition (Fig. 7, Table 1). In contrast the Eocene leucogranites show strong enrichments
293 in Ba and major depletions in all analysed high-field-strength elements (Th, Nb, Zr, Y) (Fig.
294 7). Rb/Sr ratios are significantly lower for the Jutogh leucogranites compared to the Miocene
295 leucogranites (Fig. 8, Table 1).

296

297 *Bulk-rock Nd isotopic data*

298

299 Pelitic Jutogh Group samples in the region around the intruding leucogranites (localities 57,
300 66 and W60, Fig. 1) provide $\epsilon_{Nd}(500)$ values in the range of -16.6 to -20.8 and model Nd ages
301 from 2.52 to 2.82 Ga (Table 2). These data are consistent with an 'Inner' Lesser Himalaya
302 (i.e. Late Archaean) provenance signature (Martin *et al.* 2005; Richards *et al.* 2005).

303

304 *U–Pb zircon and uraninite data*

305 U–Pb analyses of anhedral uraninite grains from sample 70ii define a discordia with an upper
306 intercept at 1810.8 ± 10 Ma (95% confidence, MSWD = 2.3), when regressed on a chord
307 anchored to the 10.5 ± 1.1 Ma (95% confidence, MSWD of concordance = 1.2) concordia age
308 of the three concordant data points from the euhedral uraninite crystal (Fig. 9a). The anhedral
309 uraninite grains evidently suffered modest Pb loss. Analyses of magmatic zircon from the
310 same sample, variably peppered with uraninite inclusions (reflected in the high U content of
311 these zircons), scatter slightly about this chord and probably record multiple periods of Pb
312 loss in some zones, although an inappropriate common-Pb (over)correction may be partly
313 responsible for scatter. Combined with a textural analysis, the more richly ‘peppered’ zones in
314 these zircons suffered the most Pb loss, in contrast to more pristine looking zones that have
315 older ages (Figs. 5e and 5f, Table 3). This suggests that high-U zones of the zircon crystals
316 experienced significant metamictization while zones less riddled with uraninite did not. Two
317 clear, inclusion-free, relatively low-U (*c.* 200 ppm) xenocrystic detrital zircon crystals from
318 sample 70ii (Fig. 6a) gave concordant data at *c.* 1920 Ma (Fig. 9b).

319

320 All analyses from sample 63i, including both zircons variably peppered with uraninite and
321 coarse uraninite inclusions in zircon, are collinear on discordia, reflecting two-component
322 mixing. The lower intercept lies close to 11 Ma as defined by young uraninite from sample
323 70ii. When tied to this value, a regression of the data yields an upper intercept of 1797 ± 20
324 Ma (95% confidence, MSDW = 6.5) (Fig. 9c), which is within error of the upper intercept
325 calculated for sample 70ii.

326

327 A third sample, 70i showed a similar range of high-U zircon data points; however the data are
328 not collinear and do not define the end members described above.

329

330 Despite complicated U–Pb systematics, three leucogranite samples from two localities present
331 low-U detrital *c.* 1920 Ma zircons, *c.* 1810 Ma magmatic high-U zircons and uraninites, and *c.*

332 11 Ma uraninite (as both coarse inclusions in high-U zircons and as discrete euhedral grains).
333 Pb-loss from the high-U zircons was both multi-episodic (data points scattered above chord
334 between 1.8 Ga and 11 Ma) and discrete, as is evident from a fan of all data points
335 terminating at a young (*c.* 11 Ma) age (Fig. 9d). Importantly, the data reflects the formation of
336 high-U zircon and uraninite during igneous crystallization at 1.8 Ga; therefore the Jutogh
337 leucogranites were emplaced during the Palaeoproterozoic.

338

339 **Discussion**

340 Like the Greater Himalayan Sequence throughout the Himalaya, the Jutogh metasediments of
341 the Sutlej Valley display amphibolite-grade metamorphism and are intruded by tourmaline-
342 bearing leucogranites. However, there is persuasive evidence that these intrusives are distinct
343 from the leucogranites that intrude the Greater Himalayan Sequence.

344

345 *Geochemistry*

346 The trace-element geochemistry of the Jutogh leucogranites is quite distinct from that of the
347 widespread Miocene leucogranites of the Greater Himalayan Sequence (Table 1, Figs. 6 and
348 7). Assuming melt saturation of high field strength (HFS) elements during anatexis, lower
349 values of the HFS elements, particularly Zr, are indicative of cooler melt conditions for the
350 Jutogh leucogranites. Applying the data from Table 1 to the equations of Watson and
351 Harrison (1983) for zircon saturation thermometry and assuming the quantity of inherited
352 zircon was negligible, we derive a maximum crystallization temperature of 670 to 690 °C for
353 the Jutogh leucogranites, compared with maximum values of 700 to 750 °C (using the same
354 equations) for the Miocene melts (Ayres *et al.* 1997). Rb, Sr, Ba systematics shed further light
355 on their contrasting petrogeneses. For the Miocene leucogranites, Rb/Sr increases with
356 decreasing Ba, and Rb/Sr ratios are distinctly higher compared with the source pelites (Fig.
357 8). As discussed by Harris and Inger (1992), these trends indicate melting under fluid-absent
358 conditions where low melt fractions and peritectic alkali feldspar contribute to a significant
359 feldspar component in the restite. For the Jutogh leucogranites, Rb/Sr ratios remain constant

360 with varying Ba and are similar to those of coexisting pelites in the Jutogh metasediments,
361 which indicates high H₂O activity during melting. Fluid-present melting for Eocene
362 leucogranites from the Greater Himalayan Sequence of Garhwal was proposed on the basis of
363 increased silica compositions and low HFS-element abundances; Zr thermometry (Watson &
364 Harrison 1983) indicates values of 610 to 670 °C (Prince *et al.* 2001), even lower than for the
365 granites in this study. The extreme enrichment of Ba seen in the Eocene leucogranites (but not
366 in the Jutogh leucogranites, Fig. 7) may be indicative of elemental transfer during Eocene
367 melting, Ba being a particularly mobile element at magmatic temperatures (Harris *et al.*
368 2003). Taken together, the geochemical evidence from the Jutogh leucogranites suggests
369 melting of pelitic compositions under conditions of elevated water activity, but possibly not
370 sufficient for H₂O saturation as in the Eocene crustal melting event. In any case, conditions
371 during melting of the Jutogh metasediments were quite distinct from the widespread fluid-
372 absent melting conditions during the Early to Middle Miocene inferred from the High
373 Himalayan leucogranites.

374

375 *Chronometry*

376 U–Pb ages of crystallization of the Jutogh leucogranites (upper intercept, Fig. 9a) provide
377 robust evidence for Palaeoproterozoic partial melting (*c.* 1810 Ma) followed by crystallization
378 of high-U zircon and uraninite. As crystallization progressed and the residual melt became
379 increasingly saturated in uranium (Guilbert & Park 1986), zircon crystallising out from the
380 melt became increasingly U-rich. This is now evident in zircons which, compared to a
381 relatively inclusion-free/more pristine ‘core’, are peppered with bright uraninite inclusions
382 outside of their cores (also reflected in the degree of metamictization, manifest by the black
383 spots, Figs. 5e and 5f). These melts inherited a few low-U zircons, *c.* 1920 Ma, unaffected by
384 metamictization, Pb loss, or uraninite crystallization. The Jutogh leucogranites thus pre-date
385 the Neoproterozoic to Cambrian (800 to 500 Ma) age of deposition for the Greater Himalayan
386 Sequence in the western and central Himalaya established from detrital zircon ages and the
387 ages of intruding granites (Parrish & Hodges 1996; Ahmad *et al.* 2000). Moreover, granitic

388 gneisses in the Lesser Himalayan Sequence (including the Jutogh Group) are invariably
389 Palaeoproterozoic (1.8 to 1.9 Ga old), as determined by accessory-phase dating (e.g. Miller *et al.*
390 *et al.* 2000; DeCelles *et al.* 2001; Richards *et al.* 2005) and supported by a marked peak in
391 detrital zircons of the same age from the Lesser Himalayan Sequence sediments (DeCelles *et al.*
392 *et al.* 2000; Richards *et al.* 2005; Richards *et al.* 2006). The Wangtu orthogneiss (1866 ± 6 Ma,
393 Richards *et al.* 2005), coeval with orthogneiss intrusions into the Lesser Himalayan Sequence
394 across the Himalaya (e.g. Munsiri granite, Nepal, 1865 ± 60 Ma, Trivedi *et al.* 1984; the
395 Iskere gneiss, Pakistan, *c.* 1850 Ma, Zeitler *et al.* 1989) represents a Palaeoproterozoic period
396 of granite intrusion with which the Jutogh leucogranites in this study (1808 ± 10 Ma) may be
397 associated. However, whereas the Jutogh leucogranites unambiguously intruded Jutogh
398 metasediments, this cannot be confidently said about the Wangtu Gneiss Complex, which is
399 bounded at its base by the Sarahan Thrust (this study). Thus, without further evidence it
400 would be unwise to directly correlate the two intrusive events.

401

402 In summary, a Palaeoproterozoic metamorphic event resulted in crustal melting of the Jutogh
403 Group sediments at temperatures of less than 700 °C, under conditions of high water activity,
404 as determined from the trace-element geochemistry. Thus, despite previous correlations of the
405 Jutogh Group with the Greater Himalayan Sequence (Singh & Jain 1993; Singh *et al.* 2006)
406 we conclude that the Jutogh Group in the Sutlej Valley *is distinct* from the Greater Himalayan
407 Sequence, confirming earlier studies that also recognised a major discontinuity between the
408 two units (Vannay *et al.* 2004; Richards *et al.* 2005).

409

410 We interpret the 10.5 ± 1.1 Ma concordia age to reflect uraninite recrystallization in response
411 to increased fluid activity related to Miocene prograde metamorphism of the Jutogh
412 metasediments, where localized U was mobilized from sites in the metamict crystal lattices of
413 Proterozoic zircon, uraninite and possibly titanite (Webb & Brown 1984). Whether Miocene
414 uraninite nucleated on pre-existing grains or self-nucleated, the analyses of both old and
415 young uraninite from one mineral separate is clear evidence of localized fluid-assisted U

416 dissolution–reprecipitation, as are relatively coarse and euhedral Miocene uraninite inclusions
417 in heavily metamict Proterozoic zircon crystals (e.g. Fig. 6c, Table 3). The absence of a
418 positive Ba anomaly (Fig. 7), in contrast to that seen in the fluid-flushed melts in the Greater
419 Himalayan Sequence (Prince *et al.* 2001), provides further evidence that element mobility has
420 been minimal, or localized, even for the most mobile of elements (Nabelek & Labotka 1993),
421 an inference supported by the localization of tourmaline alteration in the vicinity of
422 microcracks.

423

424 Although uraninite provides a powerful chronometer, the scarcity of published information on
425 its formation and behaviour during metamorphism hinders the assignment of either a pressure
426 (P) or temperature (T) to the proposed mid-Miocene prograde dehydration metamorphic
427 event. However, metamorphic monazite (included and matrix crystals) from a pelitic Jutogh
428 Group sample located within 7 km of the uraninite-bearing leucogranite (Caddick *et al.* 2007),
429 and from Jutogh Group samples in an unpublished study (cited in Vannay *et al.* 2004) yield
430 crystallization ages of 10.6 ± 0.9 Ma and 9.9 ± 0.2 to 6.4 ± 0.5 Ma respectively, i.e. within
431 error of the uraninite concordia age presented in this study. Despite the potential uncertainty
432 concerning the interpretation of some monazite age data (Martin *et al.* 2007), Caddick *et al.*
433 (2007) appropriately justify their monazite dates as primary crystallization ages using
434 systematic textural and trace element analysis. Assuming therefore that the ages of uraninite
435 recrystallization and monazite crystallization are products of the same metamorphic event, the
436 peak P–T conditions experienced by the Jutogh leucogranites and surrounding Jutogh
437 metasediments in the Miocene were 7 to 8 kbars and 600 to 700 °C (Vannay *et al.* 1999;
438 Caddick *et al.* 2007).

439

440 Miocene metamorphism (*c.* 11 Ma, amphibolite grade) related to the Himalayan orogeny
441 overprints Proterozoic metamorphism (*c.* 1.8 Ga, at least upper-amphibolite grade resulting in
442 crustal anatexis). The Jutogh leucogranites are relicts of this Proterozoic metamorphism, as is
443 marked gneissic banding (quartz/feldspar and mica-rich layers segregated on a cm scale)

444 displayed in many Jutogh metasedimentary rocks, which belies their now relatively moderate
445 metamorphic grade. To date, no monazite ages from the Jutogh Group reflect a pre-Miocene
446 metamorphic event, indicating that they have a) not been sampled and analysed, and/or b) pre-
447 existing grains were reset during Miocene metamorphism.

448

449 Polymetamorphism is widely recognised in the lithologies of the Himalayan core, e.g. garnets
450 and monazites from the Greater Himalayan Sequence preserve evidence of a *c.* 500 Ma
451 metamorphic event, now overprinted by Tertiary metamorphism (Argles *et al.* 1999; Martin *et*
452 *al.* 2007). The Lesser Himalayan Sequence also records pre-Tertiary (early Palaeozoic or
453 Precambrian) metamorphism, later overprinted in the Himalayan orogeny, e.g. at Nanga
454 Parbat (western syntaxis of the orogen) and in Nepal (Wheeler *et al.* 1995; Paudel & Arita
455 2000). Such evidence for polymetamorphism cautions strongly against assuming all
456 metamorphic features (mineralogy, textures) in the metamorphic core of the Himalaya reflect
457 the most recent orogenic phase (Gehrels *et al.* 2003). Without chronology it may be
458 impossible to distinguish between pre-Tertiary and Tertiary metamorphism, even for fabric-
459 forming index minerals. Inherited metamorphism has obvious implications for thermo-
460 barometry (e.g. Argles *et al.* 1999) and as a result, for tectonic models (e.g. Gehrels *et al.*
461 2003). Relicts of earlier deformation (e.g. lineations in the core of the Wangtu gneiss) may
462 further obfuscate tectonic interpretations.

463

464

465 *Tectonic evolution of the metamorphic core of the Sutlej Valley*

466 This combined field and geochronological study confirms that the crystalline Jutogh Group is
467 part of the Lesser Himalayan Sequence. Together, the Jutogh Group and the Vaikrita Group
468 (Greater Himalayan Sequence) form the metamorphic core in the Sutlej Valley, juxtaposed by
469 the Vaikrita Thrust which therefore coincides with the Himalayan Unconformity (Goscombe
470 *et al.* 2006) in this transect. The Jutogh and Vaikrita Groups represent discrete litho-tectonic
471 units (summarised in Table 4), with distinct geochemical affinities (Richards *et al.* 2005) and
472 tectonic styles, evident from contrasting P–T–t paths (Caddick *et al.* 2006; Harris 2007) and

473 distinct patterns and timing of cooling and exhumation (Vannay *et al.* 2004; Thiede *et al.*
474 2005). Rapid exhumation of the Vaikrita Group following peak metamorphism at *c.* 23 Ma
475 was facilitated by coeval movement on the Vaikrita Thrust below and the South Tibetan
476 Detachment above, until *c.* 16 Ma when motion ceased on the bounding faults (Vannay *et al.*
477 2004 and references therein). In contrast, the Jutogh Group was (and continues to be)
478 exhumed from *c.* 11 Ma, shortly following peak metamorphism, via concurrent thrust motion
479 on the Munsiri Thrust and extensional movement on the Karcham detachment (Jain *et al.*
480 2000; Janda *et al.* 2001), where semi-brittle top-to-the-east structures are now superimposed
481 on ductile Vaikrita Thrust fabrics (Vannay *et al.* 2004). The modelled cooling history profile
482 across the metamorphic core of the Sutlej Valley published by Vannay *et al.* (2004) reflects
483 comparatively rapid tectonic extrusion of the Jutogh Group compared to the Vaikrita Group,
484 requiring that extrusion of the two crystalline units has been decoupled since the Late
485 Miocene. The combination of cooling age data and other observations suggests that, since *c.*
486 11 Ma, exhumation of the Jutogh Group has been dominated by tectonic extrusion, whereas
487 much slower exhumation of the overlying Vaikrita and Haimanta Groups has been mainly due
488 to erosion.

489

490 This finding is consistent with tectonic models that incorporate foreland thrust propagation
491 (e.g. Dahlen 1990; Bollinger *et al.* 2006). Moreover, the pattern of exhumation has
492 implications for thermo-mechanical orogenic models incorporating the extrusion of the
493 metamorphic core as a ductile channel or wedge (Beaumont *et al.* 2004). In one formulation
494 of this process (Jamieson *et al.* 2004) the extruding channel is predicted to widen with time by
495 drawing in material from the footwall. P–T paths and timing of *peak* metamorphism from the
496 NW Himalaya (Caddick *et al.* 2007) are consistent with predictions from this model, wherein
497 the upper Lesser Himalayan Sequence (Jutogh Group) was exhumed following accretion to
498 the base of the over-thrust, extruding Greater Himalayan Sequence (Vaikrita Group).
499 However the predicted concomitant exhumation of the two units is not consistent with the
500 published Ar isotope data from the Sutlej Valley that suggests they have been decoupled at

501 least during their exhumation. If ductile flow is the mechanism responsible for the
502 exhumation history of these lithologies then the location of focused surface denudation
503 (which partly drives channel flow) has clearly migrated southwards, with early movement
504 along the base of the channel on the Vaikrita Thrust being transferred towards the foreland
505 onto the Munsiri Thrust from *c.* 11 Ma.

506

507 Feedback between tectonics and climate provides a plausible explanation for the tectonic
508 complexity of the metamorphic core of the Sutlej Valley, especially given its high fluvial
509 erosion index (Finlayson *et al.* 2002). Current precipitation is focused on the exposed slopes
510 of the Jutogh Group where the most rapid current exhumation rates are recorded (Thiede *et al.*
511 2004, Fig. 4a). The intensification of the monsoon since the Late Miocene (Vannay *et al.*
512 2004 and references therein) plus southward migration of the precipitation maxima during the
513 growth of the Himalaya may have caused a shift in the locus of exhumation across the orogen
514 towards the foreland. A similar feedback process, active since at least the Pliocene, is
515 suggested for the central Himalaya (Wobus *et al.* 2003; Hodges *et al.* 2004) where the rapidly
516 exhumed upper Lesser Himalayan Sequence coincides with the zone of maximum
517 precipitation.

518

519

520 **Conclusions**

521 This study has established that uraninite, a common accessory in anatectic granites (Bea
522 1996) and pegmatites (Guilbert & Park 1986), may provide a valuable geochronometer as
523 employed by earlier studies (this study, Fraser *et al.* 2001; Santosh *et al.* 2003). It is of
524 particular value in defining Neogene events where the high U content of uraninite will
525 produce appreciable amounts of radiogenic Pb within a few million years.

526

527 The Palaeoproterozoic melting of the Jutogh Group of the Sutlej Valley that has been
528 established by this work confirms that this previously enigmatic unit is part of the Lesser

529 Himalayan Sequence and can not be correlated with the Greater Himalayan Sequence whose
530 deposition post-dates the emplacement of anatectic granites into the Jutogh metasediments.
531 These metasediments exhibit evidence of subsequent Himalayan metamorphism, extending
532 the known range of polymetamorphism preserved in the Himalaya.

533

534 Thus, the metamorphic core exposed in the Sutlej Valley comprises both Lesser and Greater
535 Himalayan Sequence rocks (Jutogh Group and Vaikrita Group respectively) that are both
536 geochemically and tectonically distinct. The discrete exhumation paths of the Jutogh Group
537 and the Vaikrita Group specifically contradict their exhumation as a single unit, as implied by
538 models that require the Himalayan core to be the product of a single, widening ductile
539 channel. Such models thus require modification, and incorporating migrating focused surface
540 denudation may result in more realistic predictions. Further integrated metamorphic and
541 structural studies are required to test and refine tectono-thermal models of India–Asia
542 collision in the Sutlej Valley, to which the effects of climate, including palaeo-climate, can be
543 assessed.

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552 Field work was supported by a NERC Research Studentship field grant (JC) and a NERC
553 Research Grant NE/C513942/1 (TA and NH). Analytical work at the NERC Isotope
554 Geoscience Laboratory in the UK was supported by NIGF grant IP/868/1105. We thank L.
555 Godin and an anonymous reviewer for their comments which greatly improved the
556 manuscript.

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738 **Fig. 1.** Generalized geological map of the Himalaya.

739

740 **Fig. 2. (a)** Geological sketch map of a section of the Sutlej Valley after Caddick *et al.* (2007),
741 based on Vannay *et al.* (1999), with modifications from own field observations. Jutogh Group
742 localities (sample set ‘JC I05’) marked by diamonds (leucogranites), triangles
743 (metasediments) and squares (sheared amphibolite); ‘W60’ from Richards *et al.* (2005).
744 Abbreviations: GHS = Greater Himalayan Sequence; LHS = Lesser Himalayan Sequence;
745 STD = South Tibetan Detachment; VT = Vaikrita Thrust; KD = Karcham Detachment; ST =
746 Sarahan Thrust; MT = Munsiri Thrust; CT = Chail Thrust. **(b)** Cross-section (line A to A’ in
747 (a)). **(c)** Simplified geological sketch map showing lineation trends and metamorphic
748 isograds. Abbreviations: grt = garnet; st = staurolite; ky = kyanite; sill = sillimanite; mig =
749 migmatite.

750

751 **Fig. 3.** Tectono-stratigraphic column across the metamorphic core exposed in the Sutlej
752 Valley. Not to scale. Abbreviations as for Fig. 2, plus chl = chlorite, bi = biotite.

753

754 **Fig. 4.** Tectonic mélangé of the Sarahan Thrust (locality 55, Figs. 2, 3). Sigmoidal clasts of
755 amphibolite gneiss within a friable, sheared biotite–chlorite matrix indicate a top to the SSE
756 sense of shear (see bold arrows) determined from S–C fabrics, σ porphyroclasts, rotated
757 fractures clasts and slickenlines; lineations plunge gently to the NW. JC for scale, top centre.

758

759 **Fig. 5.** Leucogranite boudin (dashed outline) in Jutogh paragneiss (locality 70, Fig. 2):
760 positions of samples i and ii are shown, and lie at the edge of the leucogranite body. Foliation
761 dips moderately to the NE; lineations plunge gently to the north.

762

763 **Fig. 6.** Accessory phases separated from Jutogh leucogranite samples: incident-light images
764 of **(a)** the two different zircon populations, i.e. magmatic (three grey, semi-opaque, euhedral
765 grains) and detrital (one clear, sub-rounded grain, bottom left), and **(b)** of uraninite grains.
766 BSE images of **(c)** a high-U magmatic zircon crystal (analysis 70ii 7.1, Table 3), peppered
767 with bright uraninite inclusions and one relatively coarse-grained euhedral uraninite inclusion
768 (shown by the white arrow, analysis 70ii 7.2, Table 3), **(d)** a detrital (clear, sub-rounded)
769 zircon (analysis 70ii 8.1, Table 3), **(e)** and **(f)** high-U magmatic zircons, 70i 2 and 4
770 respectively (Table 3), with analyses in both richly ‘peppered’ and more pristine zones of the
771 zircon crystal. Scale bar in (a) and (b) is 250 μm . Black circles outline the ablation pits in (c)
772 to (f), where the spot size was 35 μm in all cases except for the smaller one in (c) which was
773 10 μm .

774

775 **Fig. 7.** Element-variation diagram for trace-element compositions of the Jutogh leucogranites,
776 and for Eocene leucogranites in the Greater Himalayan Sequence (Prince *et al.* 2001),
777 normalized against an average composition for High Himalayan (HH) Miocene leucogranites
778 (Table 1).

779

780 **Fig. 8.** Rb/Sr vs. Ba for High Himalayan (HH) Miocene leucogranites and for the Jutogh
781 leucogranites. Plagioclase-bearing metasediments from the Greater Himalayan Sequence
782 (GHS) and the Jutogh Group are also shown and represent possible melt-sources. Fluid-
783 present and fluid-absent melting trends are indicated (Inger & Harris 1993). Data from Table
784 1, Debon *et al.* (1986), Inger & Harris (1993) and references therein.

785

786 **Fig. 9.** U–Pb data from analyses of **(a)** discrete uraninites (from sample 70 ii); **(b)** detrital
787 zircons (from sample 70 ii); **(c)** zircons and uraninites in zircons (from sample 63i); **(d)**
788 zircons and uraninites from all three samples (63i, 70i, 70ii). Concordia ages in (a) and (b)
789 marked by bold dashed grey ellipses. Data point error ellipses are 2σ .

790

791 **Table 1.** *Major and trace element data for leucogranites and pelites from the Jutogh Group*

792 *(sample set 'JC I05')*

793 **Table 2.** *Nd bulk-rock data for selected Jutogh metasediments*

794 **Table 3.** *U–Pb isotopic data of uraninite and zircon grains separated from Jutogh*

795 *leucogranites*

796 **Table 4.** *Characteristics of the metamorphic core of the Sutlej Valley*

797

798

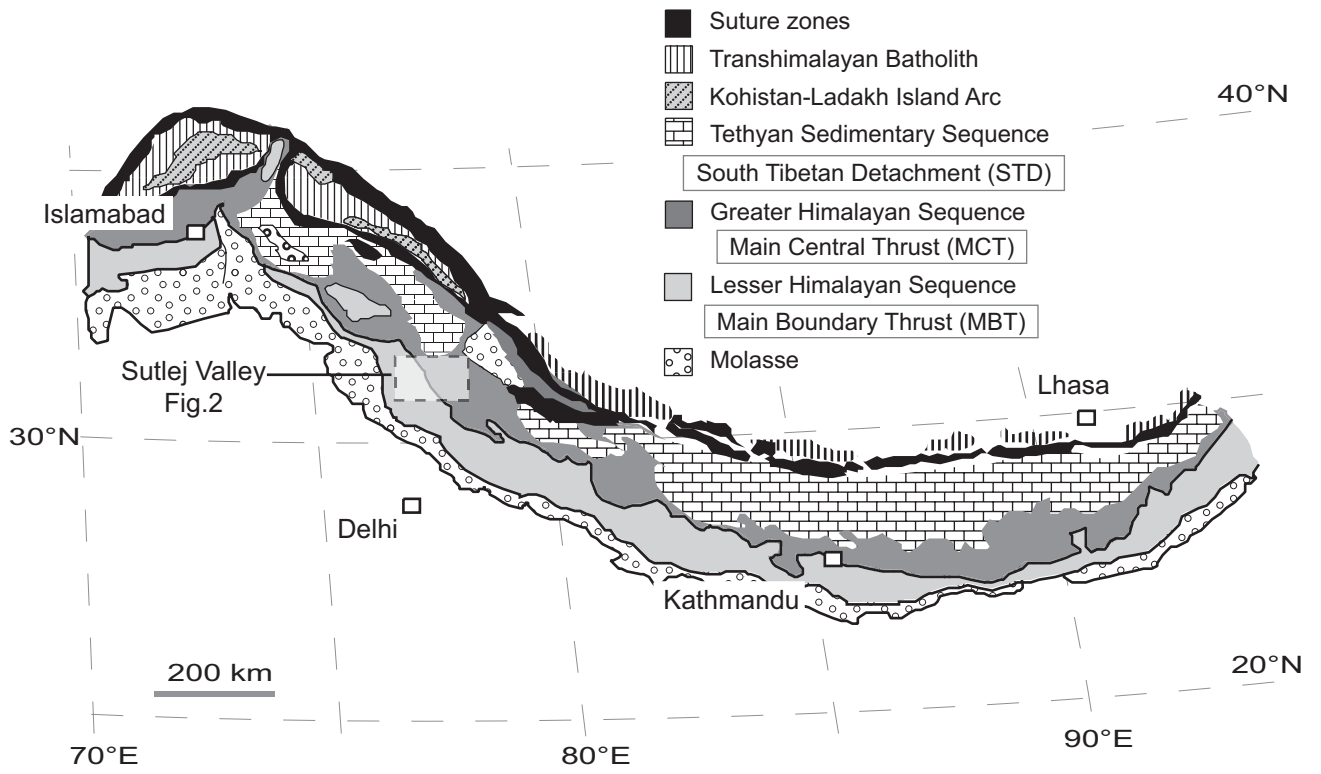


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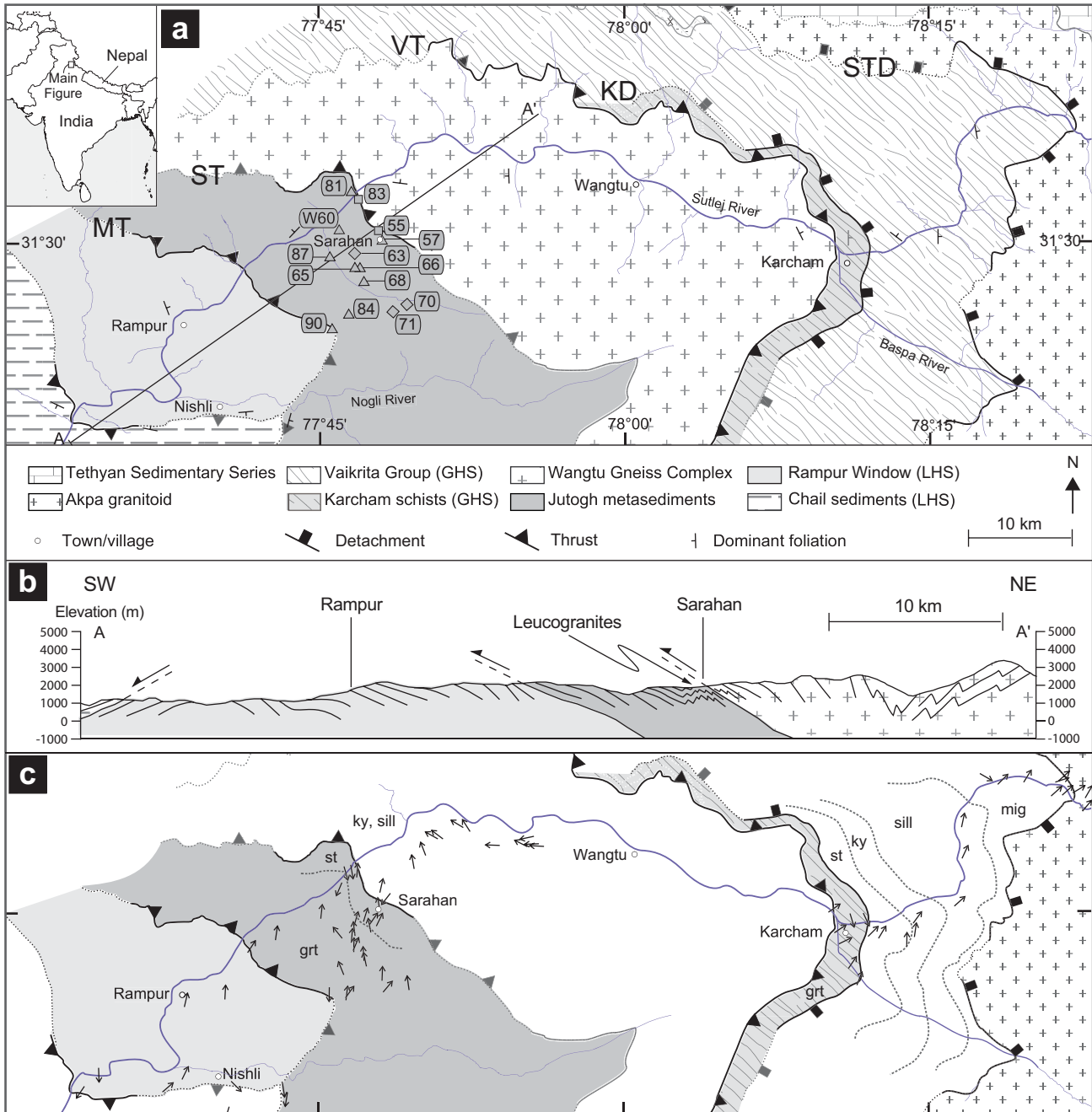


Fig. 2. (a) Geological sketch map of a section of the Sutlej Valley after Caddick *et al.* (2007), based on Vannay *et al.* (1999), with modifications from own field observations. Jutogh Group localities (sample set 'JC 105') marked by diamonds (leucogranites), triangles (metasediments) and squares (sheared amphibolite); 'W60' from Richards *et al.* (2005). Abbreviations: GHS = Greater Himalayan Sequence; LHS = Lesser Himalayan Sequence; STD = South Tibetan Detachment; VT = Vaikrita Thrust; KD = Karcham Detachment; ST = Sarahan Thrust; MT = Muniari Thrust; CT = Chail Thrust. (b) Cross-section (line A to A' in (a)). (c) Simplified geological sketch map showing lineation trends and metamorphic isograds. Abbreviations: grt = garnet; st = staurolite; ky = kyanite; sill = sillimanite; mig = migmatite.

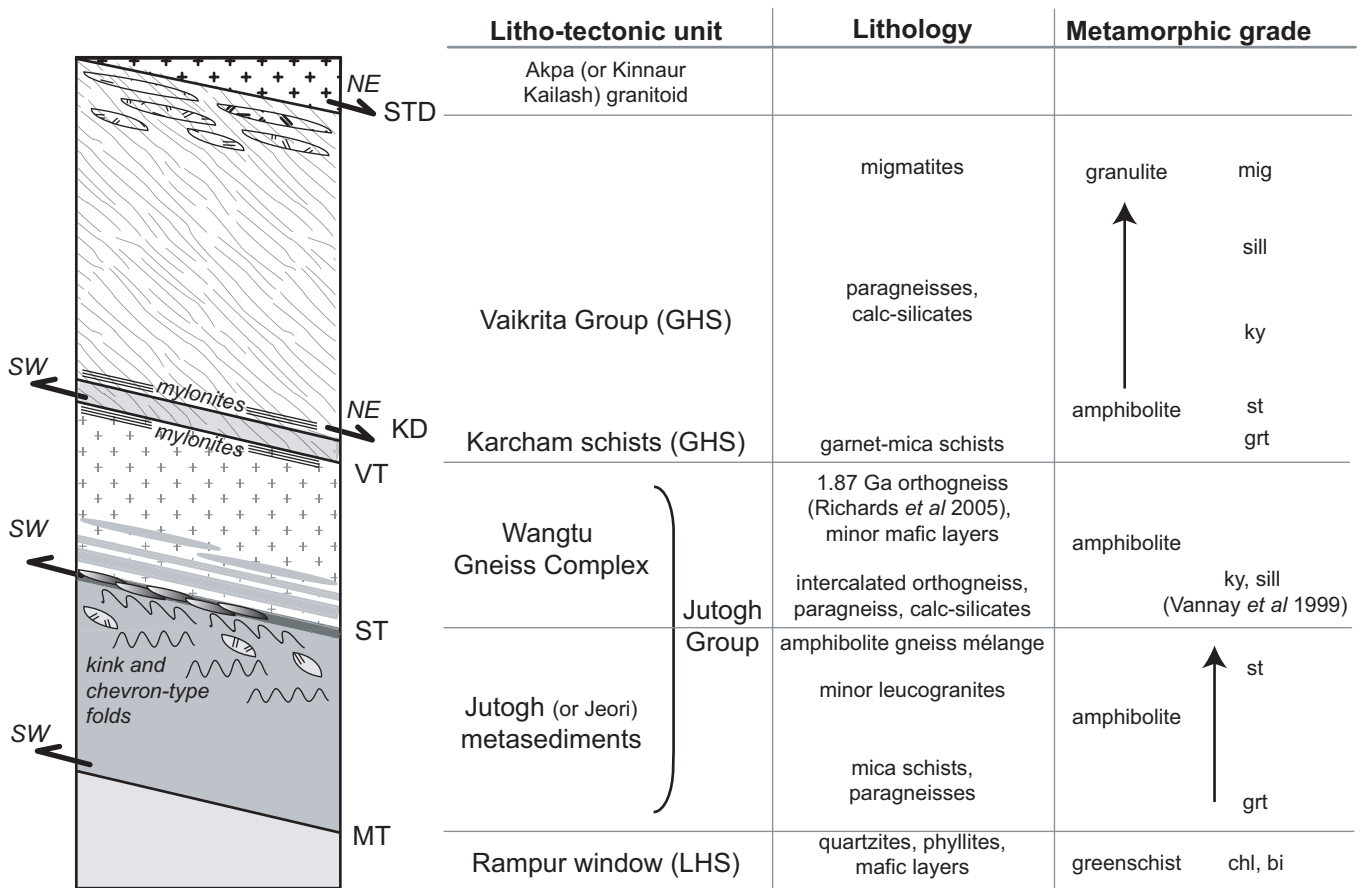


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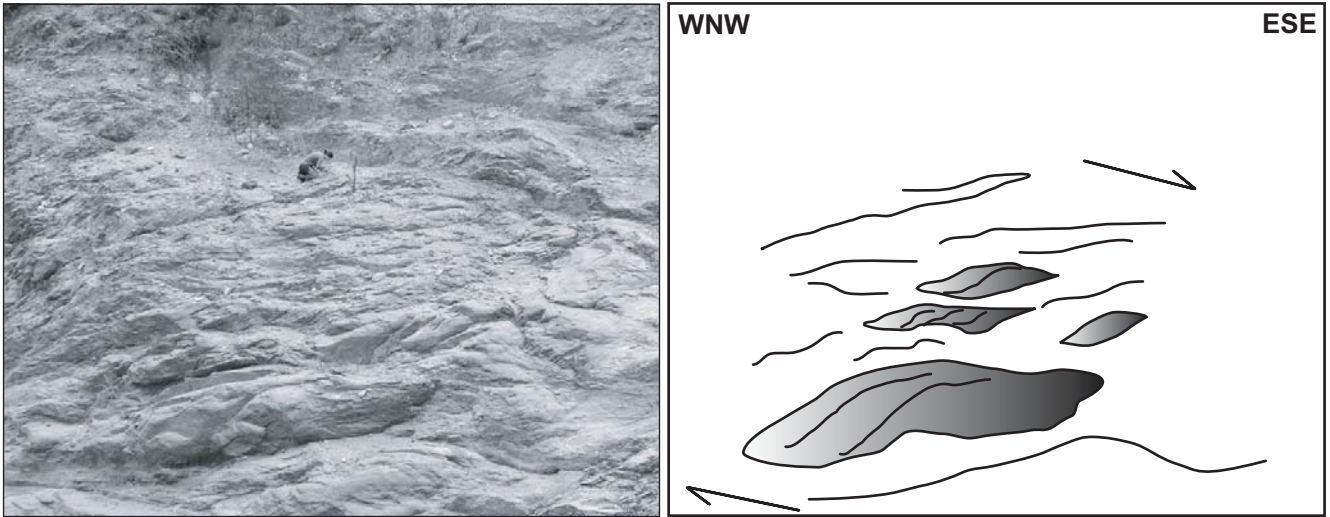


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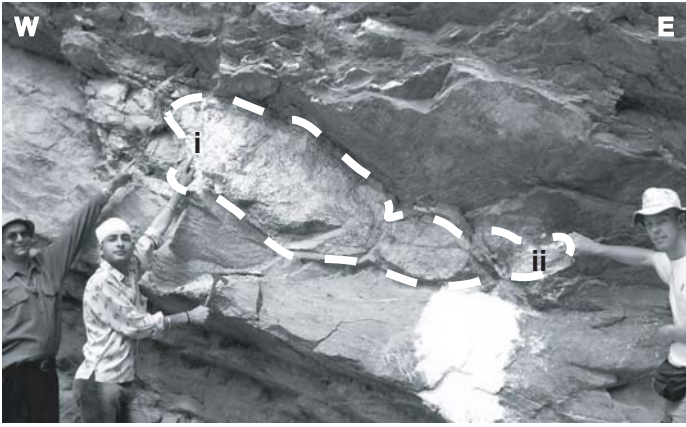


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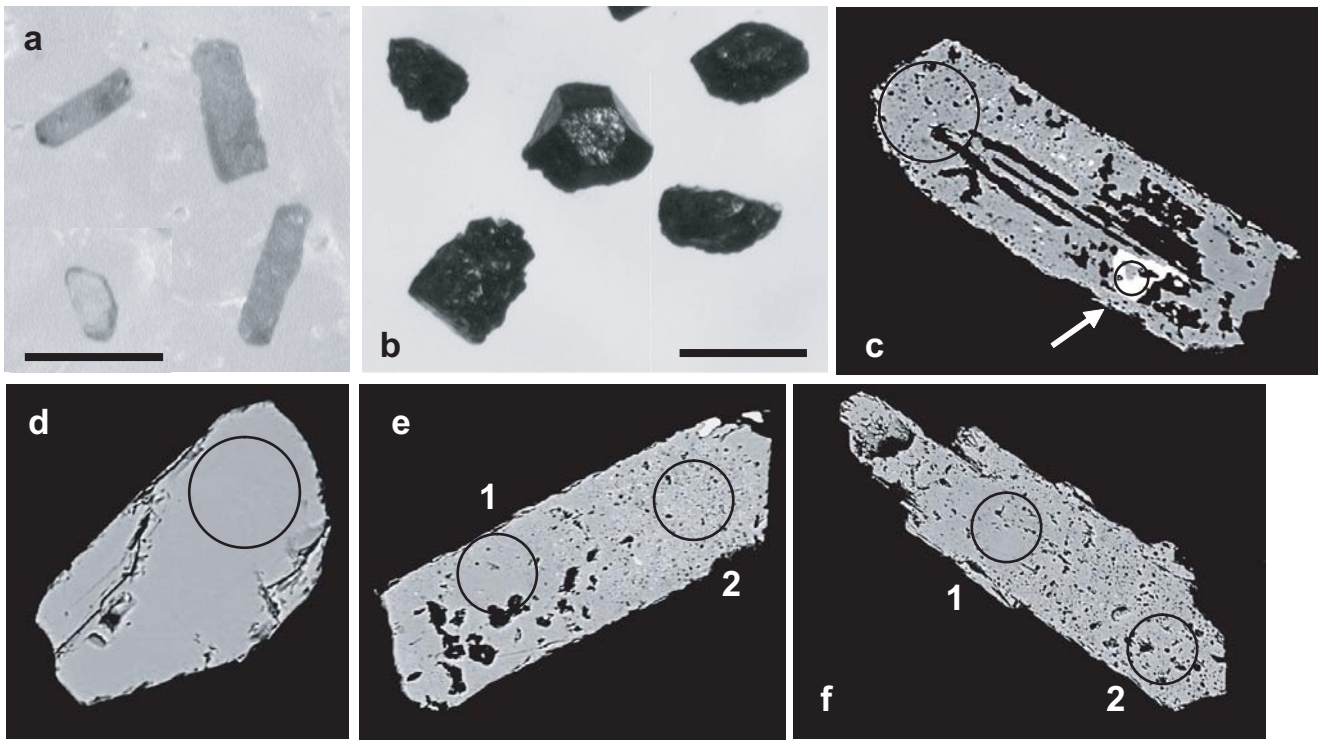


Fig. 6. Accessory phases separated from Jutogh leucogranite samples: incident-light images of (a) the two different zircon populations, i.e. magmatic (three grey, semi-opaque, euhedral grains) and detrital (one clear, sub-rounded grain, bottom left), and (b) of uraninite grains. BSE images of (c) a high-U magmatic zircon crystal (analysis 70ii 7.1, Table 3), peppered with bright uraninite inclusions and one relatively coarse-grained euhedral uraninite inclusion (shown by the white arrow, analysis 70ii 7.2, Table 3), (d) a detrital (clear, sub-rounded) zircon (analysis 70ii 8.1, Table 3), (e) and (f) high-U magmatic zircons, 70i 2 and 4 respectively (Table 3), with analyses in both richly ‘peppered’ and more pristine zones of the zircon crystal. Scale bar in (a) and (b) is 250 μm . Black circles outline the ablation pits in (c) to (f), where the spot size was 35 μm in all cases except for the smaller one in (c) which was 10 μm .

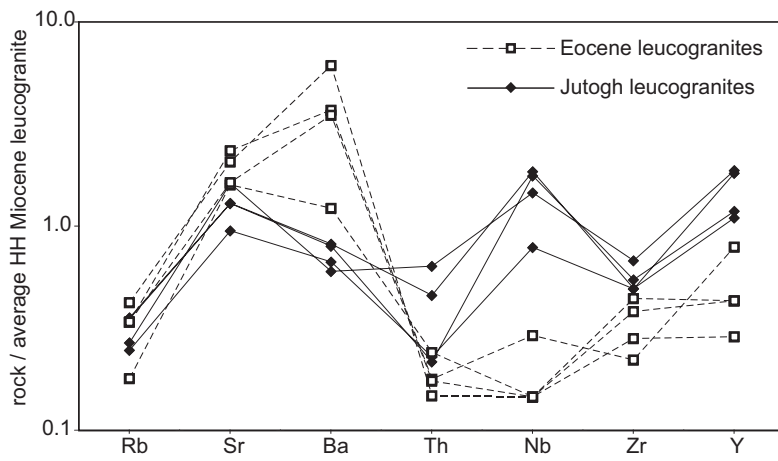


Fig. 7. Element-variation diagram for trace-element compositions of the Jutogh leucogranites, and for Eocene leucogranites in the Greater Himalayan Sequence (Prince *et al.* 2001), normalized against an average composition for High Himalayan (HH) Miocene leucogranites (Table 1).

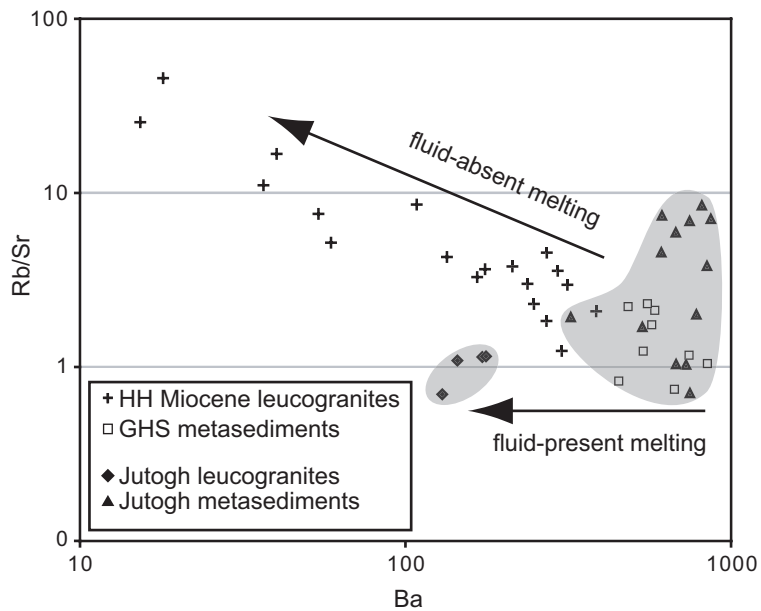


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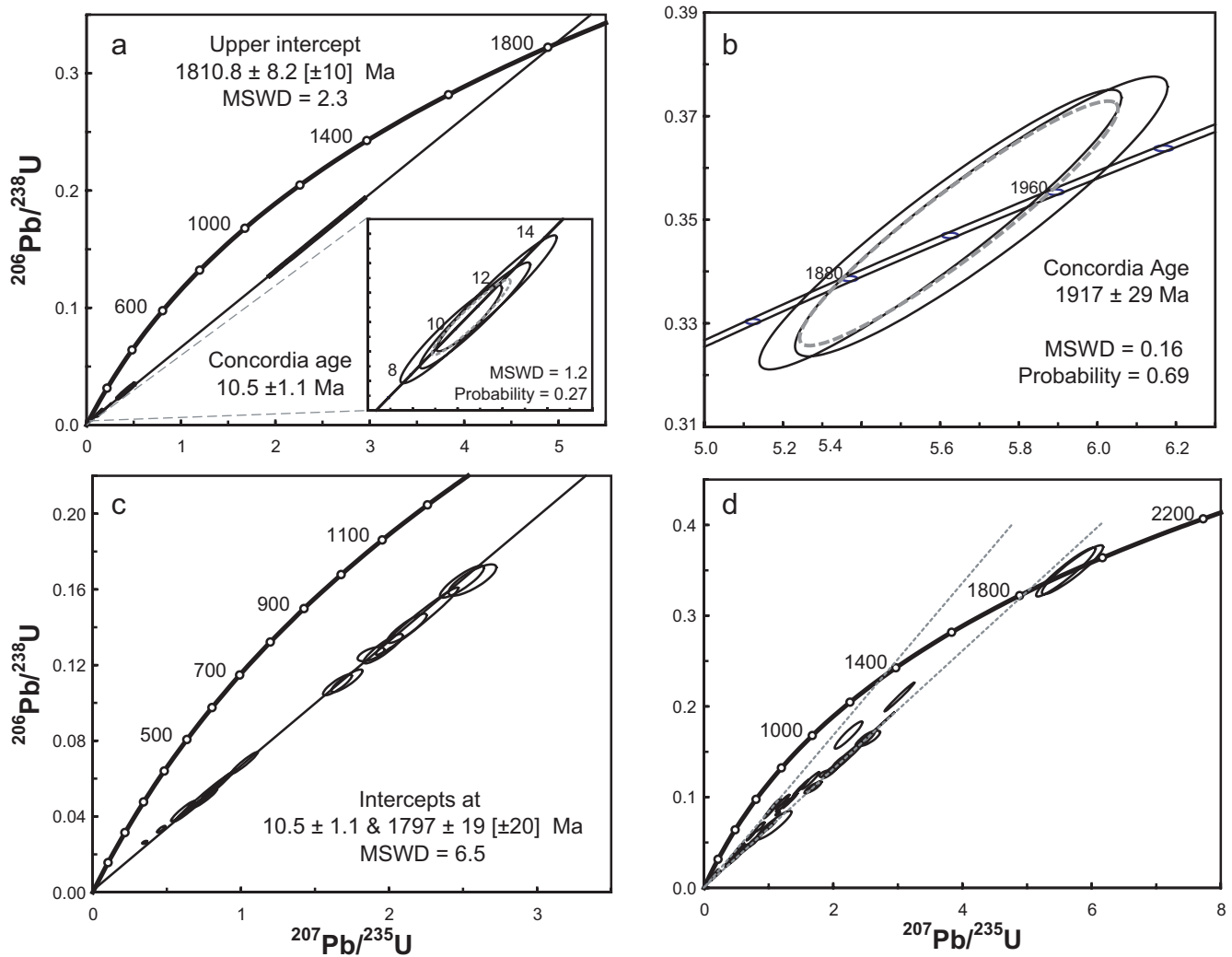


Fig. 9. U–Pb data from analyses of (a) discrete uraninites (from sample 70 ii); (b) detrital zircons (from sample 70 ii); (c) zircons and uraninites in zircons (from sample 63i); (d) zircons and uraninites from all three samples (63i, 70i, 70ii). Concordia ages in (a) and (b) marked by bold dashed grey ellipses. Data point error ellipses are 2σ .

Table 1. Major and trace element data for leucogranites and pelites from the Jutogh Group (sample set 'JC 105')

Sample	Jutogh Group leucogranites				Miocene HH				Jutogh Group pelites												
	63i	63i *	70i	70ii	leucogranite †	57	63ii	65 ‡	66i	68i	68ii	68iii	68iv	70iiix ‡	70iiiy	71i	71ii	81	84 §	87	90
<i>wt. %</i>																					
SiO ₂	75.93	76.33	74.47	76.40		66.35	68.98	61.17	64.81	67.30	70.94	58.42	66.20	65.17	67.29	64.91	67.84	70.70	66.29	59.13	73.65
TiO ₂	0.048	0.050	0.058	0.045		0.574	0.611	0.734	0.768	0.835	0.757	1.065	0.877	0.772	0.705	0.859	0.718	0.494	0.640	0.993	0.895
Al ₂ O ₃	14.93	14.94	15.54	14.29		17.41	15.91	16.63	18.20	15.58	13.41	20.35	16.39	15.88	15.10	15.24	15.39	14.47	16.93	19.19	15.98
Fe ₂ O ₃	0.93	0.91	0.99	1.06		6.15	4.86	6.72	5.19	6.69	6.23	8.06	6.81	6.10	5.66	6.68	5.47	4.74	6.23	7.51	3.30
MnO	0.023	0.021	0.013	0.014		0.052	0.042	0.044	0.055	0.099	0.036	0.054	0.101	0.087	0.081	0.068	0.077	0.254	0.07	0.053	0.005
MgO	0.26	0.27	0.47	0.43		2.51	1.51	6.41	1.99	2.07	1.73	2.52	1.81	3.00	1.29	2.42	1.52	1.32	1.69	2.93	0.25
CaO	1.02	1.02	0.92	0.68		0.24	1.95	0.29	1.89	1.58	0.39	0.61	1.12	0.31	2.21	1.55	2.62	2.92	0.79	0.40	0.33
Na ₂ O	2.76	2.73	4.54	3.80		0.69	1.85	0.31	0.64	0.57	0.35	0.45	0.43	0.45	2.70	1.74	3.12	2.74	1.10	1.38	0.70
K ₂ O	2.98	2.92	1.75	1.98		4.50	3.73	4.02	4.75	3.98	3.99	5.73	4.29	5.76	3.41	4.29	2.54	2.24	4.34	6.37	3.51
P ₂ O ₅	0.161	0.152	0.189	0.199		0.192	0.153	0.196	0.222	0.182	0.162	0.187	0.179	0.179	0.171	0.154	0.167	0.106	0.20	0.199	0.243
LOI	1.43	1.43	1.16	1.26		1.89	1.41	3.05	1.90	2.03	2.23	2.77	1.93	2.24	1.31	1.46	0.96	0.82	2.62	2.26	2.09
Total	100.48	100.77	100.09	100.14		100.55	101.00	99.58	100.42	100.92	100.22	100.22	100.13	99.96	99.92	99.37	100.42	100.81	100.89	100.41	100.96
<i>ppm</i>																					
Rb	110	111	84	77	314	258	163	190	252	240	193	281	223	324	248	223	131	152	250	265	203
Sr	96	97	121	71	75	35	158	20	66	35	33	41	32	13	146	111	185	78	55	31	196
Y	16.5	15.3	26.1	25.3	14.4	30.2	30.6	26.0	30.8	44.1	32.1	47.8	43.6	30.7	39.7	37.0	36.1	31.6	36.4	39.3	41.4
Zr	27	25	34	25	50	166	176	209	213	239	217	303	250	232	217	227	213	149	214	291	260
Nb	12.1	12.7	10.0	5.4	6.9	14.7	12.9	16.0	16.4	14.8	14.3	19.8	16.2	23.1	16.4	17.2	14.9	10.8	16.3	18.4	15.0
Ba	172	177	130	144	217	613	727	367	844	746	676	1004	866	462	534	783	748	322	611	813	678
Pb	36	35	29	22		7	18	7	23	13	22	25	12	7	25	18	30	85	12	14	39
Th	2	4	5	2	7.9	19	18	25	27	25	22	33	25	24	23	23	24	19	26	32	24
U	4	4	9	12		5	7	6	5	7	4	9	5	8	7	4	4	4	6	7	5
Rb/Sr	1.14	1.15	0.70	1.09	4.19	7.42	1.03	9.66	3.81	6.89	5.94	6.82	7.09	24.15	1.70	2.00	0.71	1.93	4.56	8.47	1.04

* duplicate analysis; † average Miocene High Himalayan (HH) leucogranite from Dietrich and Gansser (1981), Le Fort *et al.* (1987), Scaillet *et al.* (1990), Inger and Harris (1993), Ayres and Harris (1997) and Prince *et al.* (2001)

‡ plagioclase-free; § sample not in situ

Table 2. Nd bulk-rock data for selected Jutogh metasediments

Sample	57	66	W59*	W60*
$^{147}\text{Sm}/^{144}\text{Nd}$	0.1189	0.1129	0.1173	0.1328
$^{143}\text{Nd}/^{144}\text{Nd}$	0.51143	0.51130	0.51147	0.51158
Error (2σ)	0.000002	0.000002	0.000008	0.000008
ϵ_{Nd} (500)	-18.7	-20.8	-17.7	-16.6
T_{DM} (Ga)	2.65	2.69	2.52	2.82

* from Richards *et al.* (2005), sample W59 from same exposure as sample 57

Table 3. U–Pb isotopic data of uraninite and zircon grains separated from Jutogh leucogranites

Analysis	Comment/position	^{206}Pb (mV)	^{207}Pb (mV)	^{238}U (mV)	$^{206}\text{Pb}_c$ (%)	U (ppm) †	$^{207}\text{Pb}/^{206}\text{Pb}$ (%)	1 σ (%)	$^{206}\text{Pb}/^{238}\text{U}$ (%)	1 σ (%)	$^{207}\text{Pb}/^{235}\text{U}$ (%)	1 σ (%)	Rho	$^{207}\text{Pb}/^{206}\text{Pb}$ age (Ma)	2 σ abs	$^{206}\text{Pb}/^{238}\text{U}$ age (Ma)	$^{207}\text{Pb}/^{235}\text{U}$ age (Ma)	
Sample 63i																		
<i>magmatic zircon</i>																		
1.1	tip	103.3	10.5	587.2	0.6	5186	0.101	0.6	0.0820	1.0	1.138	1.2	0.857	1638	22	508	772	
1.2	core	44.8	4.9	191.0	0.3	4009	0.111	1.2	0.1117	2.2	1.716	2.5	0.872	1823	44	683	1014	
2.1	relatively inclusion-free	161.5	17.5	600.8	0.0	5306	0.108	1.6	0.1258	1.2	1.878	2.0	0.613	1771	57	764	1073	
2.2	relatively inclusion-free	75.4	8.3	229.0	0.1	4806	0.110	1.2	0.1394	2.3	2.121	2.6	0.880	1805	44	841	1156	
2.3	inclusion-rich	40.3	4.4	249.6	0.1	5258	0.110	1.0	0.0683	3.4	1.023	3.6	0.956	1793	35	426	720	
3.1 a ‡		36.4	3.4	550.0	12.2	4857	0.097	1.6	0.0263	1.2	0.351	2.0	0.603	1559	60	168	305	
3.1 b ‡		44.2	4.5	550.0	bd	4857	0.103	1.7	0.0428	5.0	0.607	5.3	0.948	1676	62	270	482	
3.2		17.9	1.8	269.6	0.2	5657	0.100	1.3	0.0333	2.1	0.461	2.4	0.859	1632	47	211	385	
4.1		276.0	29.6	2268.7	0.5	20035	0.107	0.4	0.0574	5.3	0.851	5.3	0.997	1757	14	360	625	
4.2	rim/edge	70.9	7.8	207.8	0.1	4362	0.110	1.2	0.1640	2.1	2.493	2.4	0.865	1803	44	979	1270	
5.1	tip, relatively inclusion-free	91.0	10.3	265.6	0.1	2345	0.113	1.6	0.1650	2.0	2.564	2.6	0.795	1844	56	984	1290	
5.2 a ‡	edge, relatively inclusion-free	82.7	9.3	807.7	bd	16953	0.111	1.2	0.0498	3.8	0.763	4.0	0.953	1819	44	313	576	
5.2 b ‡	"	48.1	5.4	174.2	0.3	3656	0.111	1.1	0.1432	5.1	2.190	5.2	0.979	1814	39	863	1178	
6.1	fragment	38.2	4.1	178.6	0.3	3748	0.109	1.2	0.1095	2.1	1.651	2.4	0.861	1789	45	670	990	
7.1	fragment	44.7	5.0	185.0	0.0	3883	0.111	1.2	0.1286	2.5	1.960	2.8	0.896	1808	44	780	1102	
<i>uraninite in magmatic zircon</i>																		
2.4		22.5	2.5	172.6	0.0	44976	0.108	1.5	0.0504	10.5	0.748	10.6	0.990	1761	54	317	567	
4.3		369.9	40.5	1890.6	0.0	492747	0.109	0.3	0.0525	9.1	0.788	9.1	1.000	1781	9	330	590	
Sample 70i																		
<i>magmatic zircon</i>																		
1.1		86.7	8.7	213.8	9.3	4504	0.096	1.4	0.1683	3.6	2.236	3.9	0.930	1554	54	1003	1192	
2.1 §	core	57.4	5.9	215.1	0.1	4533	0.103	1.0	0.1183	3.2	1.649	3.3	0.962	1681	36	718	999	
2.2 §	near tip	31.8	3.0	232.0	0.3	4889	0.096	1.0	0.0670	3.2	0.861	3.5	0.909	1551	37	416	643	
3.1		42.7	4.1	454.8	1.1	9582	0.105	1.0	0.0444	3.3	0.588	3.4	0.944	1712	37	285	511	
4.1 §	tip	53.0	4.7	300.4	0.7	6329	0.093	1.0	0.0847	3.9	1.026	4.0	0.965	1494	39	528	752	
4.2 §	core	48.4	4.7	258.3	0.8	5443	0.103	1.0	0.0911	3.2	1.213	3.3	0.951	1672	37	568	847	
5.1	near tip	83.6	7.8	884.9	10.3	18645	0.149	5.7	0.0422	3.4	0.530	3.8	0.905	2336	195	287	671	
<i>uraninite in magmatic zircon</i>																		
5.3		3.4	0.4	9.9	0.6	2573	0.114	4.1	0.0692	9.3	1.084	10.2	0.914	1859	150	431	746	
Sample 70ii																		
<i>magmatic zircon</i>																		
1.1	fragment	17.0	1.5	191.7	0.6	4039	0.092	1.1	0.0401	3.2	0.510	3.4	0.949	1474	41	253	418	
2.1	near tip	26.4	2.5	253.0	0.3	5330	0.095	1.0	0.0419	3.4	0.551	3.5	0.959	1535	38	265	446	

3.1		44.7	4.0	490.0	1.2	10324	0.089	1.2	0.0453	3.9	0.558	4.1	0.958	1414	45	285	451
3.2	core	44.0	4.1	372.9	1.3	7857	0.092	1.1	0.0571	3.4	0.723	3.6	0.955	1465	41	358	553
4.1	near tip	61.0	5.4	337.9	1.7	7120	0.087	1.1	0.0903	3.5	1.088	3.6	0.955	1369	42	558	748
5.1	near tip	155.0	15.3	707.8	0.4	14914	0.098	0.5	0.1105	3.1	1.498	3.1	0.989	1593	17	676	930
5.2	core	151.0	15.9	356.0	0.1	7502	0.104	0.5	0.2103	3.1	3.025	3.1	0.989	1702	17	1231	1414
6.1		94.2	9.5	503.4	1.2	10607	0.101	0.7	0.0961	3.4	1.332	3.5	0.981	1633	25	592	860
6.2	core	51.1	4.8	275.3	0.7	5800	0.093	1.0	0.0947	3.4	1.214	3.5	0.961	1488	37	583	807
7.1 §	tip	40.7	3.9	539.8	0.3	11374	0.095	1.2	0.0420	4.3	0.552	4.5	0.962	1535	46	265	446
<i>uraninite in magmatic zircon</i>																	
4.2		7.6	0.7	81.9	1.7	21350	0.104	3.0	0.0343	10.2	0.490	10.6	0.960	1689	110	218	405
4.3 a ‡		5.0	0.5	40.4	bd	10538	0.097	3.7	0.0370	9.1	0.492	9.8	0.927	1559	138	234	406
4.3 b ‡		11.4	0.9	198.9	0.2	51850	0.078	2.5	0.0167	9.0	0.180	9.3	0.962	1158	100	106	168
7.2 §	largest uraninite inclusion	32.0	2.3	2274.5	0.3	592802	0.074	1.4	0.00418	9.1	0.0426	9.2	0.988	1040	57	27	42
<i>uraninite</i>																	
U1_1	euhedral grain	18.3	0.9	3076.8	0.0	801927	0.049	2.5	0.00178	9.0	0.0122	9.3	0.964	171	116	11.5	12.3
U1_2	on same grain as U1_1	15.2	0.7	2772.4	bd	722568	0.047	2.7	0.00165	8.9	0.0108	9.3	0.956	74	130	10.6	10.9
U1_3	on same grain as U1_1	12.7	0.6	2496.2	0.2	650596	0.046	3.2	0.00152	9.0	0.0097	9.5	0.941	18	155	9.8	9.8
U1_4	on same grain as U1_1	17.9	0.8	2785.3	0.2	725950	0.047	2.6	0.00180	9.0	0.0117	9.3	0.961	69	123	11.6	11.8
U2	anhedral grain	287.4	29.8	2527.8	12.6	658823	0.104	1.2	0.0289	8.7	0.416	8.7	0.991	1702	44	184	353
U3	anhedral grain	1429.1	158.7	2849.6	0.0	742694	0.110	0.1	0.1597	8.6	2.431	8.6	1.000	1806	4	955	1252
U4	anhedral grain	146.9	15.1	2466.8	10.9	642929	0.103	0.8	0.0151	8.7	0.214	8.8	0.996	1676	29	97	197
U5	anhedral grain	73.3	7.0	2241.5	5.1	584201	0.096	0.9	0.0102	8.7	0.136	8.7	0.994	1552	35	66	129
<i>detrital zircon</i>																	
8.1 §		7.9	0.9	11.9	0.3	251	0.117	1.1	0.3481	3.2	5.601	3.4	0.943	1906	40	1925	1916
10.1		5.0	0.6	8.1	0.6	170	0.118	1.3	0.3506	3.1	5.705	3.4	0.927	1926	46	1938	1932

²⁰⁶Pb_c (%) indicates the common portions in total ²⁰⁶Pb; ^{206*}Pb and ^{207*}Pb refers to the radiogenic ²⁰⁶Pb and ²⁰⁷Pb; U (ppm) † indicates that the concentration uncertainty is estimated at ± 25%; ‡ split analysis; § see Fig. 5 (c) to (f)

bd, below detection

Table 4. *Characteristics of the metamorphic core of the Sutlej Valley*

	Vaikrita Group	Jutogh Group
Geochemical affinity *	Greater Himalayan Sequence	Lesser Himalayan Sequence
Age of peak metamorphism (t) † ‡	c. 23 Ma	c. 11 Ma
Pressure (P), bottom to top of unit §	c. 8 kbar	c. 9 to 7 kbar
Temperature (T), bottom to top of unit §	c. 570 to 750 °C	c. 610 to 700 °C
P-T-t path geometry †	clockwise; isothermal decompression after peak P-T before cooling (broad path)	clockwise; peak P coincided with peak T; uplift with immediate cooling (tight path)
Period of exhumation via tectonic extrusion ‡	c. 23 to 16 Ma	c. 11 Ma to present
- thrust motion on	Vaikrita Thrust	Munsiari (Jutogh) Thrust
- extensional motion on	South Tibetan Detachment	Karcham Detachment
Average exhumation rate (Late Miocene to present) ‡	c. 0.7 mm/yr	c. 2.3 mm/yr

* Richards *et al.* (2005); † Caddick *et al.* (2006), see also Harris (2007); ‡ Vannay *et al.* (2004); § Vannay *et al.* (1999)