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**Constraining the timing of palaeosol development in Iranian arid  
environments using OSL dating**

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## 24 1. Introduction

25 Establishing a chronological framework is of crucial importance to palaeoclimate and  
26 landscape evolution studies. Without any knowledge of the timing of environmental change,  
27 understanding environmental history is not possible. Palaeosols are important archives for  
28 reconstruction of past climates and environments (e.g. Retallack, 2001); they are formed in  
29 response to environmental changes and can be used as records for understanding  
30 palaeoclimatic conditions. Palaeosols in various sediment sequences such as alluvial (e.g.  
31 Kemp et al., 2006), aeolian (e.g. Wang et al., 2016), marginal marine strata and even glacial  
32 deposits (e.g. Mahaney et al., 2013) are reliable indicators for establishing a  
33 chronostratigraphic subdivision and correlation with marine isotope stages (MIS) and glacial-  
34 interglacial cycles (e.g. Gornitz, 2008). Palaeosols developed in alluvial deposits have the  
35 potential of providing information on episodes of landscape instability (sediment deposition),  
36 stability (soil development) and landscape evolution (Kraus, 1999).

37 The palaeoclimate records of Iran have been subject to several studies over the last decade  
38 (e.g. Karimi et al., 2009; Kehl, 2010; Khormali and Kehl, 2011). Quaternary events in Iranian  
39 plateau present important records of palaeoenvironmental changes in the arid environments  
40 of the Middle East. They can be seen as a valuable link for correlating Europe and Central  
41 Asia through the deserts of Saudi Arabia to the West and the deserts of Central Asia to the  
42 East; providing an understanding of palaeoenvironment and landscape evolution resulting in  
43 the genesis of the Old World Dry Belt (Kehl, 2010).

44 Most of the palaeoclimatic studies in Iran have been based on geomorphic and pedogenic  
45 evidence (e.g. Karimi et al., 2009; Khormali and Kehl., 2011; Farpoor et al., 2012;  
46 Ghafarpour et al., 2016). A few studies have applied absolute dating methods to studying  
47 palaeoenvironmental and landscape change in Iran (Kehl et al., 2005; Frechen et al., 2009;  
48 Karimi et al., 2011; Lauer et al., 2016; Büdel et al., 2017), but most focus on the loess-

49 palaeosol sequences of Northern and Northeast of Iran (e.g. Frechen et al., 2009; Karimi et  
50 al., 2011; Lauer et al., 2016). Some of these studies suggest that well-developed palaeosols  
51 correlate with the last interglacial (MIS 5) or MIS 7 and older interglacials (Frechen et al.,  
52 2009; Lauer et al., 2016). Additionally, they suggest loess accumulation during last glacial  
53 period MIS 2 (Karimi et al., 2011).

54 In a recent study in northern Iran, stratigraphic results of Agh Band loess-palaeosol show  
55 poorly- as well as well-developed palaeosol horizons formed around 80 ka and late MIS 7-  
56 MIS 6 with increased humidity and landscape stability (Lauer et al., 2016). In southern Iran,  
57 apart from two studies (i.e. Kehl et al., 2005; Kehl et al., 2009), no information is available  
58 on the various phases of soil formation and sediment deposition. According to Kehl et al.  
59 (2005), based on the stratigraphy and different degrees of soil development, the well-  
60 developed Bw(t) and Btk horizons in the Persepolis basin, reflect probable soil-forming  
61 periods during MIS 5, while the upper loess deposits were likely formed under considerably  
62 cooler and drier climatic conditions during the LGM. The poorly-developed horizons indicate  
63 interstadial periods of MIS 3 and short phases of climate amelioration during the early  
64 Holocene.

65 It appears that there is little information available from the dry zones of Iranian plateau to  
66 allow the reconstruction of palaeoenvironment based on the timing of sediment deposition-  
67 soil formation sequences. This is partly due to the lack of reliable chronological control in the  
68 studies in this area. Alluvial fans and pediments are the most common landforms in the arid  
69 regions of Iran. The most developed soil horizon identified in these landforms is a Btk  
70 horizon in both buried and relict forms. Morphological, micromorphological and  
71 mineralogical characteristics of the palaeosols developed in these landforms have been  
72 previously described (Khademi and Mermurt, 1999; Khormali et al., 2003; Sanjari et al.,  
73 2011; Nejad Zamani., 2014). These studies investigated the pedogenesis of these arid-region

74 soils using various techniques. Unfortunately, none of these studies provides quantitative  
75 information on the timing of palaeosol development. Indeed, in the Middle East in general,  
76 the timing of sedimentation and pedogenesis in alluvial sediment is not well documented.  
77 Given the similarity of palaeosols in the arid areas in Iran, an overall understanding of the  
78 timing of sedimentation and soil genesis episodes is very desirable. Here, we attempt to  
79 establish a first instrumental chronology for pedosedimentary events associated with  
80 previously identified palaeosols; these developed in alluvial fans and pediments in the arid  
81 areas of the Iranian plateau. Luminescence dating is used to set the first chronological  
82 constraints on the palaeosols, by determining the time at which the sediments, in which the  
83 palaeosols later formed, were deposited. Optically stimulated luminescence (OSL) is a well-  
84 established absolute dating method that determines the time elapsed since sediment grains  
85 were last exposed to daylight. In this method, quartz and feldspar grains act as natural  
86 dosimeters and can potentially yield ages of sediment deposition ranging from a few years to  
87 few hundred thousands of years. Many studies have made use of luminescence dating to  
88 establish a chronology for sedimentation and soil formation processes (e.g. Murray and  
89 Clemmensen, 2001; Leopold et al., 2011; Hal and Goble, 2015). In an attempt to establish a  
90 chronology for soil development, we apply luminescence dating to six different soil profiles  
91 in arid alluvial settings in Central Iranian plateau.

## 92 **2. Study area and sampling**

93 The Iranian Plateau is located in southwest Asia, between the Persian Gulf and the Oman Sea  
94 to the south and the Caspian Sea to the north (Darehshouri and Kasraian, 1998) (Fig. 1a). The  
95 study sites are located at six different geographical areas in the centre, east and southeast of  
96 the Iranian plateau (Fig. 1a). The present-day average annual precipitation and temperature  
97 range from 145 to 200 mm and from 14°C to 24.5 °C, respectively, thus the overall climate of  
98 the study area is considered arid.

99 All our sections are located in alluvial and fluvial landforms and are physiographically  
100 similar (Table 1). Each section includes a representative pedon identified and logged in  
101 alluvial or fluvial deposits that originated from adjacent highlands. The location and  
102 environmental characteristics of sections are described in Table 1. These pedons consist of a  
103 sequence of palaeosol-sediment or a sequence of poorly- and well-developed palaeosols (Fig.  
104 1b). The studied pedons and their sediment origins have long geological history, spanning  
105 from Miocene to present and Cretaceous to Pliocene, respectively. Because the most part of  
106 the Iranian plateau is covered by limestone formations, the parent material of most of the  
107 pedons studied here are calcareous; dominated by limestone and marl. The representative  
108 pedons were described and soil horizons identified according to the criteria of the Soil Survey  
109 Staff (2014).

110 In total, seventeen samples were collected for luminescence dating by hammering metal tubes  
111 (5 cm diameter and 15 cm long) into freshly exposed section walls: (i) twelve samples from  
112 well-developed palaeosols, (ii) two samples from less developed palaeosols, and (iii) three  
113 samples from sediment layers without or with very little pedogenic material (Fig. 1b). At  
114 each site (except for Isfahan where the ground surface was reworked by agricultural activity),  
115 an additional sample was collected by brushing the very top surface ( $< 2$  cm) of the ground  
116 into aluminium foil which was then wrapped and sealed in light-tight bags. Such modern  
117 analogues were taken to estimate the degree of bleaching (i.e. residual dose) in currently-  
118 exposed samples. In addition to the luminescence samples, ~250 grams of material from  
119 around each tube was collected for annual dose rate measurements.

### 120 **3. Sample preparation and analytical facilities**

121 Sediment was removed from sampling tubes under subdued orange light. Material from the  
122 exposed ends was saved for water content measurements and sediment from the middle  
123 (unexposed) part of the tubes was wet sieved to 125–250  $\mu\text{m}$ . This fraction was then treated

124 with 10% HCl to remove carbonates. The presence of gypsum in some of the samples  
125 presented a challenge in sample preparation; gypsum dissolves in water very slowly. To  
126 accelerate this dissolution, we treated the samples with a mixture of citric acid and propanol  
127 repeatedly until all or the majority of gypsum was removed. The samples were subsequently  
128 treated by 10% H<sub>2</sub>O<sub>2</sub> to remove any reactive organic material, followed by 10% HF for 40  
129 min. to remove any alpha-irradiated surface layer and weathering products and coatings, and  
130 10% HCl for 20 min. to remove any fluoride precipitation. The K-rich feldspar fractions were  
131 separated by suspension in a water-based heavy liquid solution ( $\rho=2.58 \text{ g.cm}^{-3}$ ; Fastfloat).  
132 The denser (settled) quartz grains ( $\rho > 2.58 \text{ g.cm}^{-3}$ ) were further treated with concentrated  
133 (40%) HF for 60 min. followed by 10% HCl for 40 min..

134 All luminescence measurements were carried out using a Risø TL/OSL reader (model TL-DA  
135 20), with blue light stimulation (470 nm,  $\sim 80 \text{ mW.cm}^{-2}$ ) and photon detection through a 7.5-  
136 mm Hoya U-340 glass filter for quartz, and infrared stimulation (875 nm,  $\sim 135 \text{ mW.cm}^{-2}$ )  
137 and photon detection through a Schott BG39/BG3 filter combination (2 and 3mm,  
138 respectively) for K-feldspar (Bøtter- Jensen et al., 2010). Beta irradiations used a <sup>90</sup>Sr/<sup>90</sup>Y  
139 source mounted on the reader and calibrated for both discs and cups using 180–250  $\mu\text{m}$   
140 calibration quartz grains (Hansen et al., 2015). Grains were mounted as large (9 mm diameter  
141 for quartz) or medium ( $\sim 4 \text{ mm}$  diameter for feldspar) aliquots in a monolayer on 9-mm-  
142 diameter stainless steel discs (quartz) or cups (feldspar) using silicone oil. The heating rate  
143 was  $5 \text{ }^\circ\text{C.s}^{-1}$  throughout. All thermal treatments and stimulations at temperatures higher than  
144  $200 \text{ }^\circ\text{C}$  were carried out in nitrogen atmosphere, and a pause of 5 s was inserted before  
145 stimulation to allow all grains to reach the measurement temperature. Five empty channels  
146 were inserted before and after the stimulation to monitor any isothermal TL signals.

#### 147 **4. Dosimetry**

148 Radionuclide concentrations ( $^{238}\text{U}$ ,  $^{232}\text{Th}$  and  $^{40}\text{K}$ ) were measured using high-resolution  
149 gamma spectrometry. The additional sediment sample collected from around each metal tube  
150 was first dried at 50 °C. A subsample of ~250 g was pulverized and homogenized, and then  
151 heated to 450 °C for 24 h to remove any organic matter. The material was then cast in wax to  
152 prevent radon loss and to provide a reproducible counting geometry. Samples were stored for  
153 at least three weeks to allow  $^{222}\text{Rn}$  to reach equilibrium with its parent  $^{226}\text{Ra}$  before being  
154 measured on a high-purity Germanium detector for at least 24 h. Details of the gamma  
155 spectrometry calibration are given in Murray et al. (1987). The internal beta dose rate from  
156  $^{40}\text{K}$  to feldspar grains was calculated based on an assumed effective potassium content of  
157  $12.5\pm 0.5\%$  (Huntley and Baril, 1997), and the beta contribution from  $^{87}\text{Rb}$  was calculated  
158 assuming a  $^{87}\text{Rb}$  content of  $400\pm 100$  ppm (Huntley and Hancock, 2001). A small internal  
159 alpha contribution of  $0.10\pm 0.05$  Gy ka<sup>-1</sup> from internal  $^{238}\text{U}$  and  $^{232}\text{Th}$  was also included in the  
160 dose rates, derived from  $^{238}\text{U}$  and  $^{232}\text{Th}$  concentration measurements by Mejdahl (1987). For  
161 quartz, an internal dose rate of  $0.010\pm 0.002$  Gy.ka<sup>-1</sup> was assumed (Vandenberghe et al.,  
162 2008). The radionuclide concentrations were converted to dose rate data using the conversion  
163 factors from Guerin et al. (2011). The cosmic-ray dose rates were calculated according to  
164 Prescott and Hutton (1994), assuming an uncertainty of 5%. The long-term water content of  
165 each sample was estimated based on the field water content and saturation water content as  
166 well as the probable position of the water table during the burial time. All radionuclide  
167 concentrations, water contents and dose rates are summarized in Table 2.

## 168 **5. Luminescence characteristics and dose measurements**

### 169 **5.1. Quartz**

170 All the OSL measurements were performed at 125 °C for 40s using a single-aliquot  
171 regenerative (SAR) protocol (Table S1; Murray and Wintle, 2000). A high-temperature blue  
172 light stimulation at 280 °C was applied at the end of each cycle to reduce recuperation (Table

173 S1; Murray and Wintle, 2003). Values for  $L_x$  and  $T_x$  were derived from the initial 0.16 s of  
174 the signal minus an immediate background derived from the following 0.16 s. An early  
175 background subtraction was used to minimize the possible effect of the more difficult to  
176 bleach and more thermally unstable medium and slow components (Jain et al., 2003; Li and  
177 Li, 2006; Cunningham and Wallinga, 2010).

178 For all the samples, the purity of quartz extracts was examined by measuring the OSL signal  
179 from three aliquots from each sample with and without prior IR stimulation at room  
180 temperature for 100 s. The ratio of the two signals, the so-called OSL-IR depletion ratio, was  
181 then calculated for each aliquot (Duller, 2003). The resulting average OSL-IR depletion ratio  
182 was  $0.90 \pm 0.03$  ( $n = 51$ ), implying that any feldspar contamination in our quartz OSL signal is  
183 not significant. Quartz extracts from all the samples were sensitive and the OSL signal was  
184 dominated by the fast component (Fig. 2a, inset). Figure 2a shows a typical dose-response  
185 curve for sample 177118, fitted with a saturating exponential function; the sensitivity-  
186 corrected signals are reproducible (recycling ratio is close to unity), the dose-response curve  
187 passes through the origin (recuperation is small) and the  $D_e$  for this aliquot is  $\sim 111$  Gy.

188 Both natural and dose recovery preheat plateau tests were performed to determine the  
189 appropriate measurement conditions. The natural preheat-plateau test was carried out to  
190 investigate the dependence of equivalent dose ( $D_e$ ) on preheat temperature. Twenty-one  
191 aliquots of quartz from sample 177118 were sorted into groups of three. The  $D_e$  was  
192 measured with different preheat temperatures for each group, in steps of  $20^\circ\text{C}$  ranging from  
193  $180^\circ\text{C}$  to  $300^\circ\text{C}$  (held for 10 s). The temperature of the cut heat was chosen to be  $40^\circ\text{C}$  lower  
194 than that of the first preheat treatment. From Fig. 2b it can be seen that there is no systematic  
195 dependence of  $D_e$  on preheat temperature between  $180$  and  $300^\circ\text{C}$ .

196 Similarly, the dose-recovery preheat-plateau test was carried out on 21 fresh aliquots of  
197 sample 177118. The natural OSL signal was removed by two blue stimulations for 100 s at

198 room temperature. A pause of 10000 s was used between the two stimulations to allow any  
199 charge trapped in shallow refuge traps (especially that associated with the 110 °C TL peak) to  
200 decay and subsequently partly refill the OSL trap before the second stimulation. The aliquots  
201 were then given a laboratory dose of ~75 Gy and measured in a similar manner as in the  
202 natural preheat plateau test. Figure 2c illustrates the results of measured to given dose ratios  
203 at different preheat temperatures. As with the  $D_e$  preheat plateau, no systematic dependence  
204 of dose recovery ratio on preheat temperature is observed. The average dose recovery ratio is  
205  $1.06 \pm 0.04$  ( $n=21$ ), showing that a known laboratory dose given to the sample before any  
206 thermal treatment can be measured accurately. Based on these results, we selected a preheat  
207 temperature of 240 °C, and undertook a further dose recovery tests on all the samples at this  
208 temperature; the overall dose recovery ratio is  $0.88 \pm 0.05$  ( $n = 51$ ; inset to Fig. 2c). There is a  
209 large scatter in these data; the dose recovery ratios vary from 0.26 to 2.7 (inset to Fig. 2c).  
210 Nevertheless, for all quartz OSL measurements described below a preheat/cutheat  
211 temperature of 240/200°C was used, although we further investigate the possible effect of  
212 this very variable dose recovery ratio on our quartz OSL ages in Section 6.

## 213 **5.2. K-rich feldspar**

214 A pIRIR<sub>290</sub> protocol was adopted to measure the K-feldspar fractions (Buylaert et al., 2012).  
215 The  $D_e$  was measured using a SAR protocol with a preheat of 320 °C for 60 s after both  
216 regenerative and test doses. The first IR stimulation at 50 °C was followed by a second IR  
217 stimulation at 290 °C. A high-temperature stimulation at 325 °C was also given at the end of  
218 each SAR cycle to minimise any build-up of charge giving rise to a recuperated signal. All IR  
219 stimulations were carried out for 100 s (Table S1). The pIRIR signals were derived from the  
220 first 1 s of the IR stimulation with a subtracted background based on the last 10 s. Test dose  
221 values were chosen to be ~25% of the pIRIR<sub>290</sub>  $D_e$  estimates for all samples. All dose  
222 response curves were fitted with a single saturating exponential function.

## 223 **6. Quartz OSL ages and bleaching**

224 The quartz OSL ages of all the pedons are summarised in Fig.1b. The ages range from  
225  $5.2\pm 0.6$  ka for the sediment in which a poorly-developed palaeosol (Bk) has formed at the  
226 Jiroft section to  $107\pm 11$  ka for the sediment containing a well-developed Btky2 palaeosol at  
227 Isfahan section. The sediment ages at all sections are in stratigraphic order except at Isfahan  
228 where the age of the bottom Btky3 layer is younger than both the age of the immediately  
229 overlying Btky2 layer, and the unit above, containing Btky1. We do not consider the age of  
230 the bottom unit in the discussion below because it is younger than both the overlying samples  
231 and because the equivalent dose of the immediately overlying sample 177120 is  $\sim 150$  Gy (i.e.  
232 approaching saturation). It is well-known that quartz close to saturation tends to  
233 underestimate age (Chapot et al., 2012) and so it is not surprising that the age of the bottom  
234 sample is underestimated.

235 In the luminescence dating of samples from alluvial settings, one must always consider the  
236 possibility of incomplete resetting or bleaching of the signal before final deposition. The  
237 bleaching rate may be a function of several parameters such as water depth, sediment load,  
238 turbulence, light spectrum, grain size, and transport distance (e.g. Jain et al., 2004). To  
239 investigate the likely importance of incomplete bleaching, we use two approaches: i) we  
240 determine the residual dose in modern analogues, and ii) look at the relative doses recorded  
241 by two signals that bleach at different rates.

### 242 **6.1. Modern analogue**

243 One way to evaluate the significance of incomplete bleaching is to estimate the residual doses  
244 in currently-exposed deposits at the sampling site (e.g. Murray and Olley, 2002; Jain et al.,  
245 2004; Porat et al., 2010). Such modern analogues can provide useful information on the likely  
246 degree of bleaching in the fossil samples prior to deposition. Eighteen aliquots of quartz and  
247 twelve aliquots of K-feldspar were measured on material extracted from samples of

248 identifiably modern sediment from 5 sites. Table 3 summarizes the quartz OSL and K-  
249 feldspar  $IR_{50}$  and  $pIRIR_{290}$   $D_e$  values of modern analogues from these sites. No representative  
250 modern analogue was available at Isfahan section due to significant soil surface disturbance  
251 by agricultural activity. The quartz OSL residual doses range from  $\sim 0.1$  Gy (at Bam, Lar and  
252 Mahan) to  $\sim 1.6$  Gy (at Rayen). The largest  $IR_{50}$  and  $pIRIR_{290}$  doses are  $\sim 2$  Gy (at Mahan) and  
253  $\sim 5$  Gy (at Rayen), respectively. All these doses are small compared to the equivalent doses  
254 measured in the fossil samples from the corresponding site, suggesting that the fossil samples  
255 were likely to have been sufficiently bleached at the time of deposition.

## 256 **6.2. K-feldspar versus quartz**

257 Several studies have shown that the quartz OSL signal resets more rapidly than feldspar  $IR_{50}$   
258 signal (e.g. Godfrey-Smith et al., 1988) and that the  $IR_{50}$  signal, in turn, bleaches more  
259 quickly than the  $pIRIR_{290}$  signal (e.g. Murray et al., 2012). Unfortunately sufficient feldspar  
260 was not available from some samples, but Fig. 3a summarises the dose measurement made on  
261 the 15 samples for which sufficient material could be extracted (all sites). This plot of the  
262  $IR_{50}$  versus  $pIRIR_{290}$  equivalent doses shows that most data points are consistent with a  
263 straight line (Fig. 3a) of slope  $0.47 \pm 0.02$ . Given the heterogeneous nature of the bleaching  
264 process in space and time, the relatively constant relationship over a wide dose range implies  
265 that the feldspar signals in most of our samples are likely to have been well-bleached prior to  
266 deposition (e.g. Sohbati et al., 2016).

267 Based on the differential bleaching of quartz OSL and K-feldspar IRSL signals, Murray et al.  
268 (2012) proposed an approach to identifying well-bleached quartz by comparing quartz OSL  
269 with feldspar  $IR_{50}$  and  $pIRIR_{290}$  ages. Following a similar approach, a comparison between  
270 the quartz OSL and the K-feldspar  $IR_{50}$  ages shows that, except for three samples (at Jiroft  
271 profile), the  $IR_{50}$  ages are comparable to or younger than the corresponding quartz ages (Fig.  
272 3b, Table 2), with an average  $IR_{50}$  to quartz age ratio of  $0.58 \pm 0.04$ , consistent with the degree

273 of fading expected for the IR<sub>50</sub> signal, and suggesting that the quartz OSL signal in these  
274 samples is likely to have been well-bleached prior to deposition (all these samples are  
275 identified as ‘probably well-bleached’ in Table 2). The IR<sub>50</sub> signal from the three samples  
276 from the Jiroft profile has apparently not been as well-bleached as those from the other  
277 profiles, and thus we do not know if the quartz signal in these samples was sufficiently reset.  
278 We also compared the quartz OSL ages with the pIRIR<sub>290</sub> ages. Figure 3c shows that the  
279 pIRIR<sub>290</sub> ages for 6 of the samples are consistent with the corresponding quartz OSL ages  
280 (identified as ‘well-bleached’ in Table 2); the remaining (including the 3 from the Jiroft  
281 profile) overestimate. This implies that despite the negligible pIRIR<sub>290</sub> residual doses in the  
282 modern analogues and the correlation between the IR<sub>50</sub> and pIRIR<sub>290</sub> over a wide dose range  
283 (Fig. 3a), the pIRIR<sub>290</sub> signal in some of our samples may not have been sufficiently bleached  
284 prior to deposition.

### 285 **6.3. Does quartz dose recovery affect the OSL ages**

286 In order to see whether the apparent difficulty in recovering a known dose for some aliquots  
287 (inset to Fig. 2c) is systematically affecting the measurement of dose in quartz, we compare  
288 the OSL to IR<sub>50</sub> age ratio with the quartz OSL dose recovery ratio for all the samples. Figure  
289 3d shows that there is no correlation between the two ratios. Thus, the unsatisfactory dose  
290 recovery ratios observed for some of the samples does not seem to lead to a detectable  
291 systematic underestimation of the quartz age, as might otherwise have been expected. Since  
292 our quartz OSL ages do not seem to be affected either by poor dose recovery or by  
293 incomplete bleaching (with the possible exception of the Jiroft profile) the quartz ages are  
294 used in the geological interpretation of the next section.

## 295 **7. Discussion**

296 The Jirof section provides three OSL ages of between  $5.2 \pm 0.6$  and  $15 \pm 2$  ka. These OSL age  
297 estimates indicate the time of sediment deposition and so suggest that soil development

298 started after 15 ka ago, and was ongoing until at least the middle of the Holocene (although  
299 the youngest unit has only a poorly-developed soil). However, none of these 3 samples are  
300 identified as well- or probably-well bleached by comparison between the quartz OSL age and  
301 the feldspar ages, and so the calculated ages may be overestimates (marked with a (?) in  
302 Table 2).

303 The two samples from Rayen section give OSL age estimates of  $5.5\pm 0.5$  ka for a sediment  
304 with no significant soil development 60 cm below the surface, and  $26\pm 3$  ka for the most  
305 developed Btk horizon at 115 cm; the 2 OSL ages are identified in Table 2 as well bleached  
306 and probably well bleached, respectively. It appears that alluvial deposition of this pedon  
307 took place before or around the LGM, and that soil formation took place later, resulting in a  
308 well-developed Btk horizon in depth of 115 cm. Further alluvial sediments were deposited at  
309  $5.5\pm 0.5$  ka (MIS 1), and no significant soil development took place in this unit, constraining  
310 the period of soil development in the underlying unit to between  $26\pm 3$  and  $5.5\pm 0.5$  ka. The  
311 sharp transition between the lower well-developed soil horizon and the upper underdeveloped  
312 deposits suggests a chronological discontinuity in this pedon, for instance arising from an  
313 erosion event.

314 Finally, the uppermost unit at Mahan contains a well-developed Bt palaeosol in a sediment  
315 with a deposition age of  $18.5\pm 1.5$  ka, and identified in Table 2 as well bleached. Although  
316 less well-constrained than at Jiroft and Rayen, this palaeosol may also have developed during  
317 the Holocene.

318 These results suggest that 5 units (3 at Jiroft, 1 at Rayen and 1 at Mahan) probably all  
319 developed soils to various degrees during the Holocene. The presence of a  $5.5\pm 0.5$  ka  
320 sediment cap at Rayen and only a weakly-developed soil in the youngest unit at Jiroft  
321 probably constrains the active soil development period to the early/mid Holocene.

322 The stratigraphy and soil horizons recorded in the Bam pedon appear similar to those in the  
323 Rayen pedon, but deposition occurred much earlier, at  $61\pm 5$  ka (MIS 4), indicating soil  
324 formation probably after MIS 4. The age of the sample taken from upper coarse-grained  
325 deposits is  $9.1\pm 0.9$  ka, substantially younger than the underlying sediment; this constrains  
326 the formation of this soil most likely to MIS 3, although soil formation in MIS 2 cannot be  
327 ruled out purely on stratigraphic grounds. The time gap between the two horizons may  
328 indicate an erosional event, possibly during MIS 2. One of the two samples appears to have  
329 been well reset, and the other probably well reset (Table 2).

330 The palaeosol-sediment sequence at Mahan gives stratigraphically consistent OSL age  
331 estimates ranging from  $18.5\pm 1.5$  to  $65\pm 4$  ka and all are identified as well or probably well-  
332 bleached (Table 2). The sediment sequence began to be deposited in MIS 4 ( $65\pm 4$  ka) but soil  
333 development did not take place until after  $54\pm 4$  ka, when a well-developed palaeosol (Btk 45-  
334 70 cm) began to form. The upper unit was deposited at  $18\pm 1.1$  ka during MIS 2, and so the  
335 underlying soil probably developed during MIS 3.

336 The Isfahan section consists of a multiple pedocomplex. The OSL age estimates range from  
337  $40\pm 8$  for a sample from the upper horizon at 50 cm below the surface to  $107\pm 11$  ka for the  
338 sedimentary unit at 140 cm (ignoring the unit below this for the reasons discussed above). All  
339 of the samples are judged to have been well-bleached or probably-well bleached before  
340 deposition (Table 2). The sediment sequence began to be deposited at  $107\pm 11$  ka or earlier  
341 and soil formation occurred later, but before  $73\pm 6$  ka, when more sediment was deposited  
342 covering the palaeosol. Further pedogenesis occurred after this time, but before  $40\pm 8$  ka,  
343 when more sediment was deposited. One final period of soil development is recorded at the  
344 top of this section, occurring after  $40\pm 8$  ka. This palaeosol presumably developed at the same  
345 time as that found in the middle unit at Mahan (deposition at  $54\pm 4$  ka), and possibly the  
346 bottom unit at Bam (deposition at  $61\pm 5$  ka). Thus, it appears we have 4 periods of soil

347 development represented at Isfahan, the first (bottom unit) of unknown age but before MIS  
348 5d, one during mid/late MIS 5, one between MIS 5 and mid MIS 3, and the top palaeosol  
349 during late MIS 3.

350 The Lar section is similar to that at Isfahan. The earliest sediment deposition occurs at  
351  $102\pm 10$  ka. This unit is covered by one deposited at  $73\pm 12$  ka, which is in turn covered by  
352 one from  $43\pm 5$  ka. Unfortunately, feldspar was unavailable for these samples, and so we are  
353 unable to comment on the degree of bleaching prior to deposition. Each sedimentary unit  
354 contains a well-developed palaeosol, and thus, as with Isfahan, there are soil formation  
355 periods during mid/late MIS 5, one between MIS 5 and mid MIS 3, and one after mid MIS 3.  
356 In summary, we have firm evidence for soil formation before MIS 5d, during mid/late MIS 5,  
357 between MIS 5 and mid MIS 3, but probably in early MIS 3 (because of the constraining  
358  $54\pm 4$  ka age at Mahan), during late MIS 3, and in the early to mid MIS 1.

## 359 **8. Conclusion**

360 For the first time, a chronology has been developed for the sediments hosting several soil  
361 profiles in the central Iranian plateau. Our OSL ages indicate the time at which the sediment  
362 was deposited; soil formation took place later. The 17 OSL ages constrain at least four broad  
363 phases of sediment deposition and soil formation on the central Iranian plateau: (i) prior to,  
364 and (ii) during, mid/late MIS 5 (at Isfahan and Lar), (iii) MIS 3 (at Bam, Mahan and probably  
365 Isfahan), and (iv) MIS 1 (at Rayen and Jiroft). It appears that, over the last full glacial-  
366 interglacial cycle, there is no convincing evidence for palaeosol formation during MIS 4 and  
367 MIS 2.

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371

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## 500 **Figure captions**

501

502 **Figure 1)** (a) Map of the Middle East and Iran, showing the location of six sampling sites in  
503 the (semi-)arid zones of Iranian plateau. (b) Stratigraphy of palaeosol units and location of  
504 the OSL samples. The dashed lines indicate possible chronological correlations, not coeval  
505 soil development. There are marked discontinuities in all profiles. These may be erosional.

506

507 **Figure 2)** Summary of luminescence characteristics of quartz OSL signal. (a) Typical dose-  
508 response and stimulation (inset) curves, (b) and (c) natural and dose recovery preheat plateau  
509 tests for a sample from Mahan (171118). Each data point represents the average of three  
510 aliquots. The error bars show one standard error. Inset to (c) shows the summary of dose  
511 recovery test for all the samples in this study.

512

513 **Figure 3)** (a) Sample-averaged K-feldspar equivalent doses. Error bars represent one  
514 standard error, (b) quartz OSL and K-feldspar IR<sub>50</sub> ages, (c) quartz OSL and K-feldspar  
515 pIRIR<sub>290</sub> ages, (d) quartz OSL to K-feldspar IR<sub>50</sub> age ratio plotted against quartz dose  
516 recovery ratio.

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ACCEPTED MANUSCRIPT

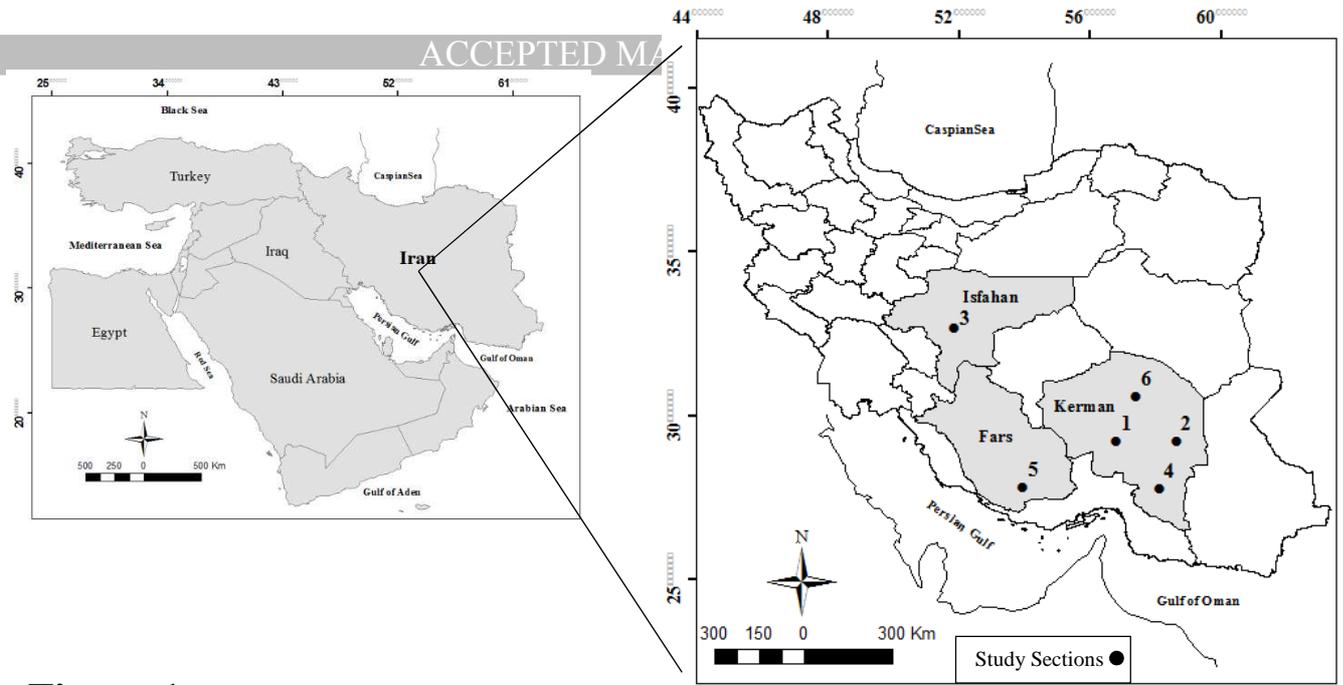


Figure 1a

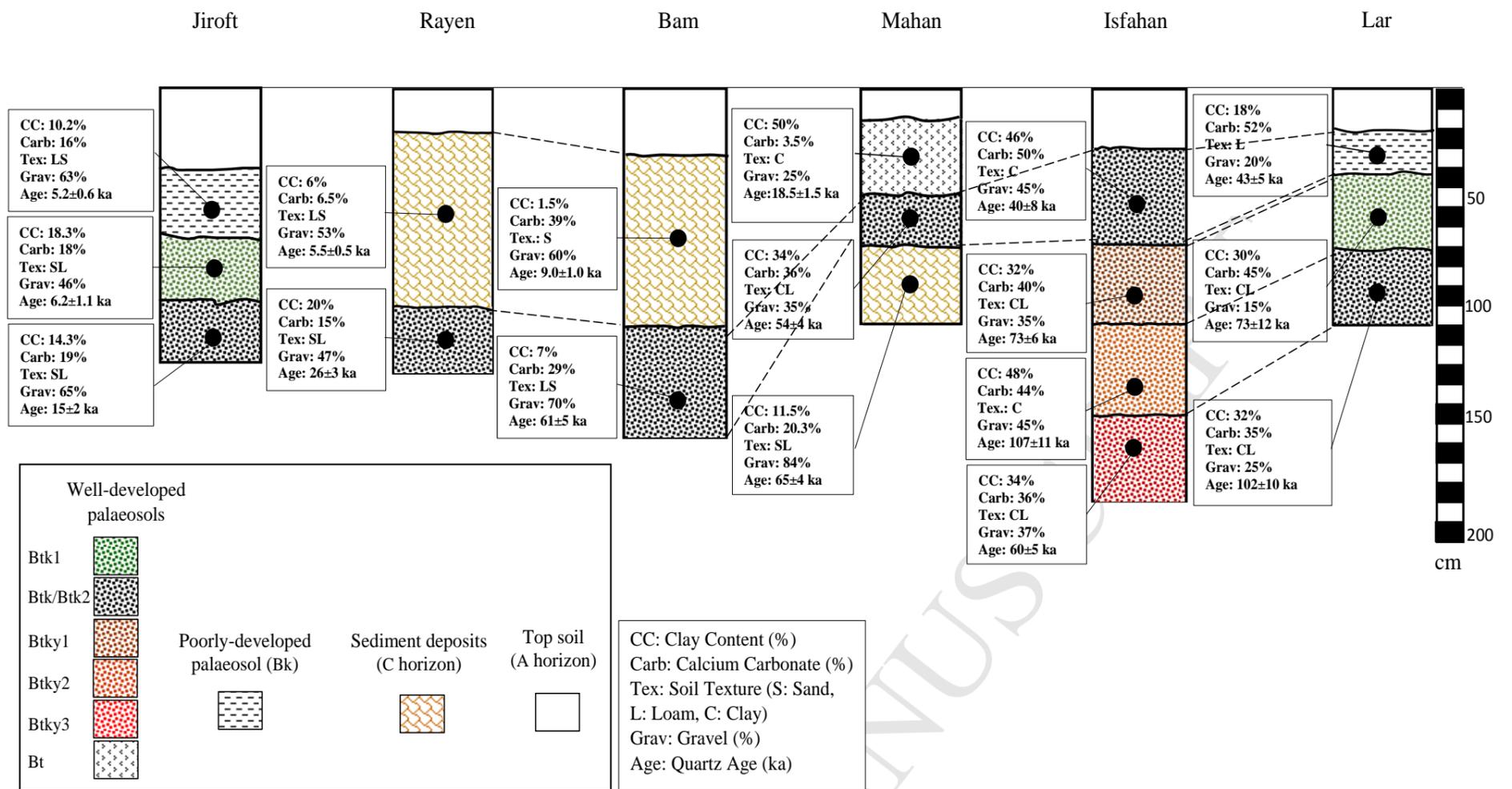


Figure 1b

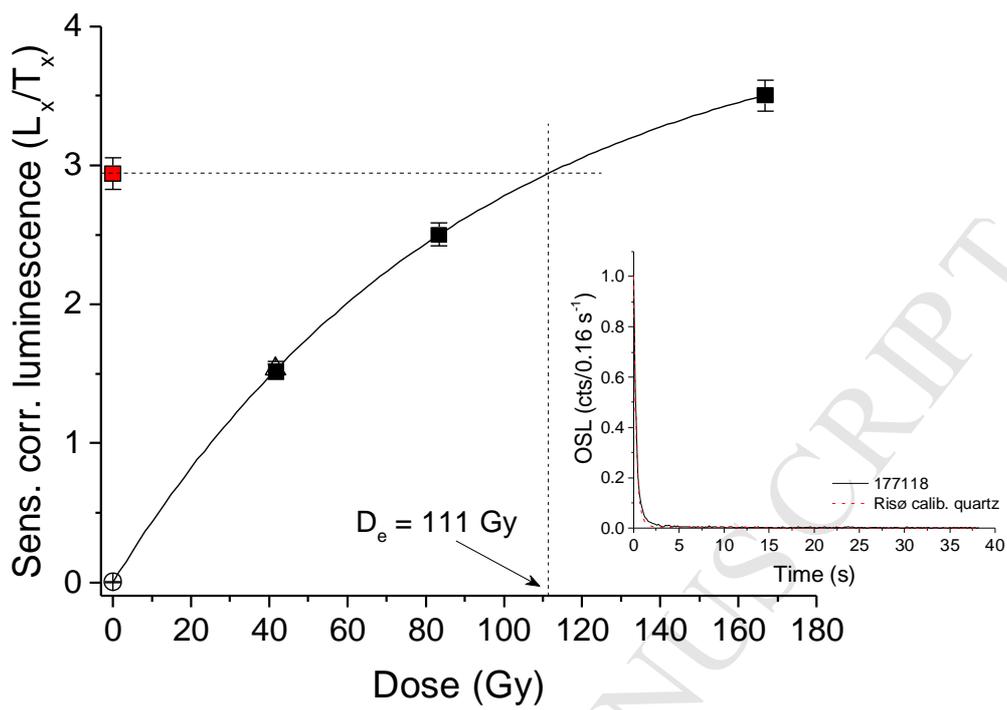
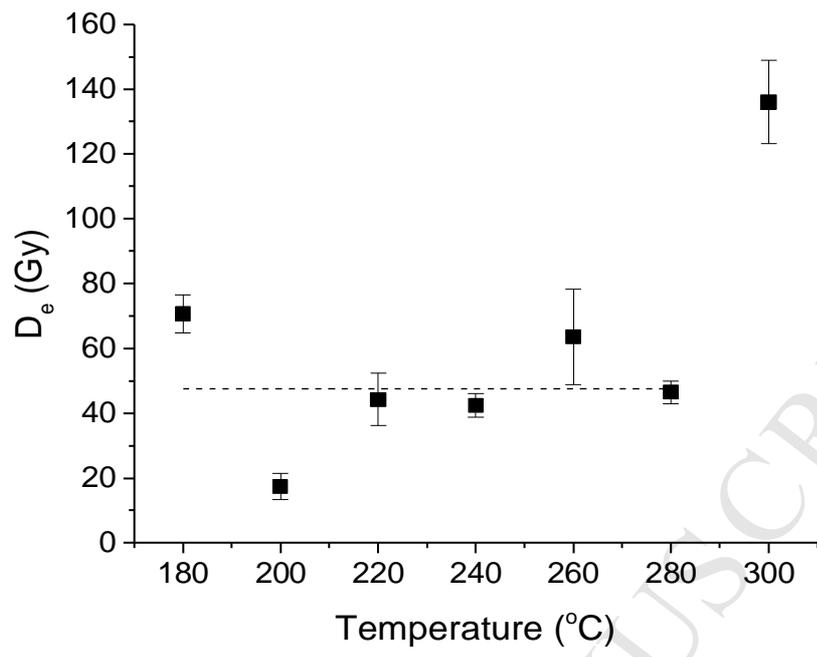


Figure 2a

**Figure 2b**

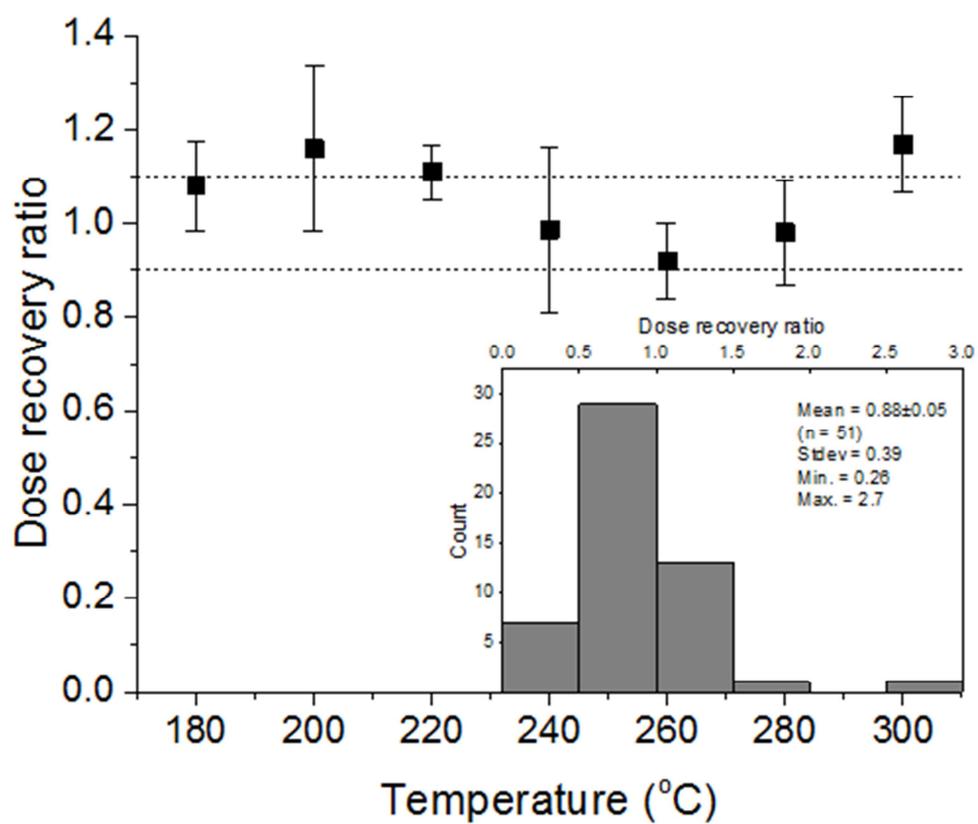


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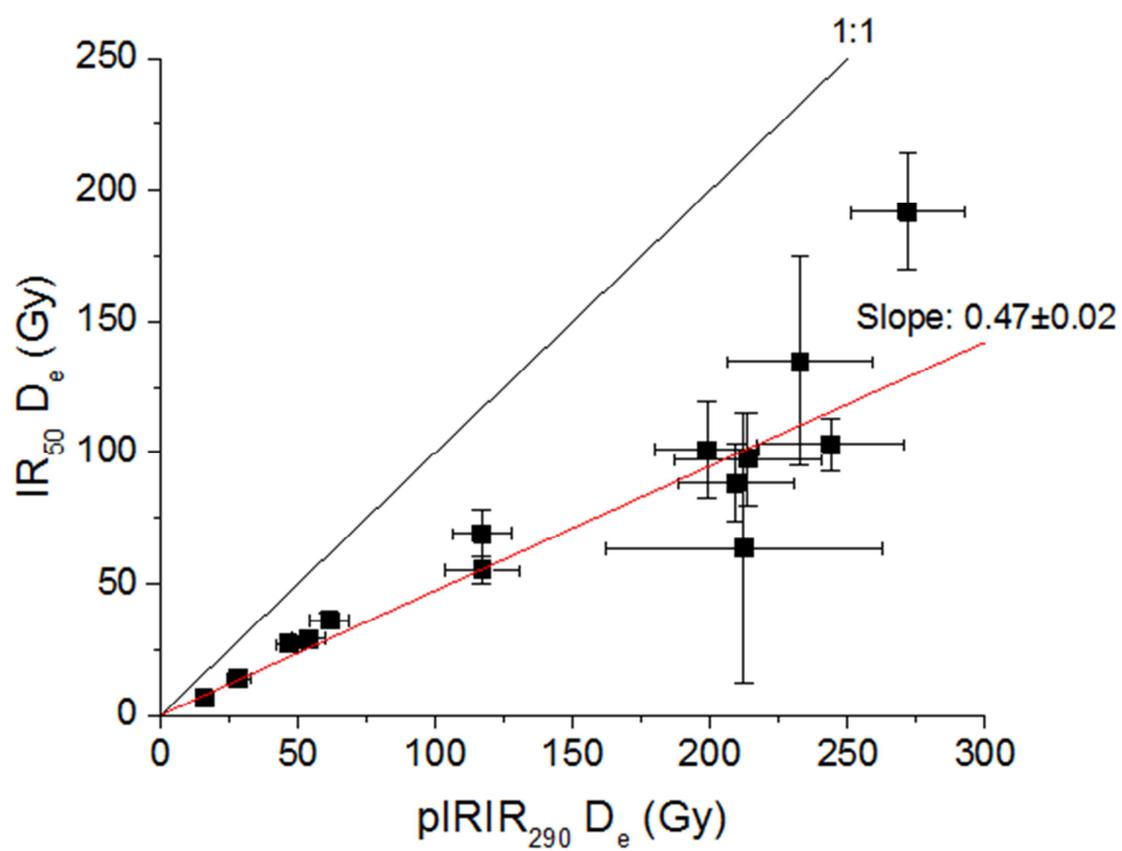


Figure 3a)

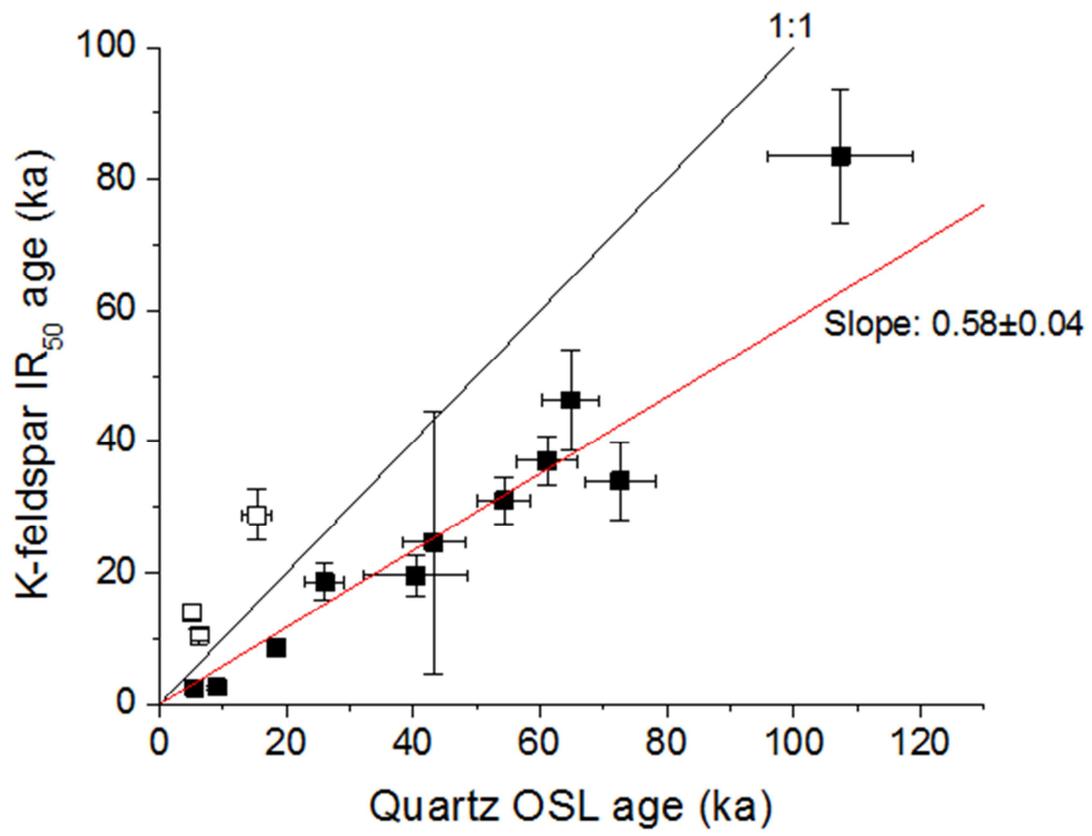


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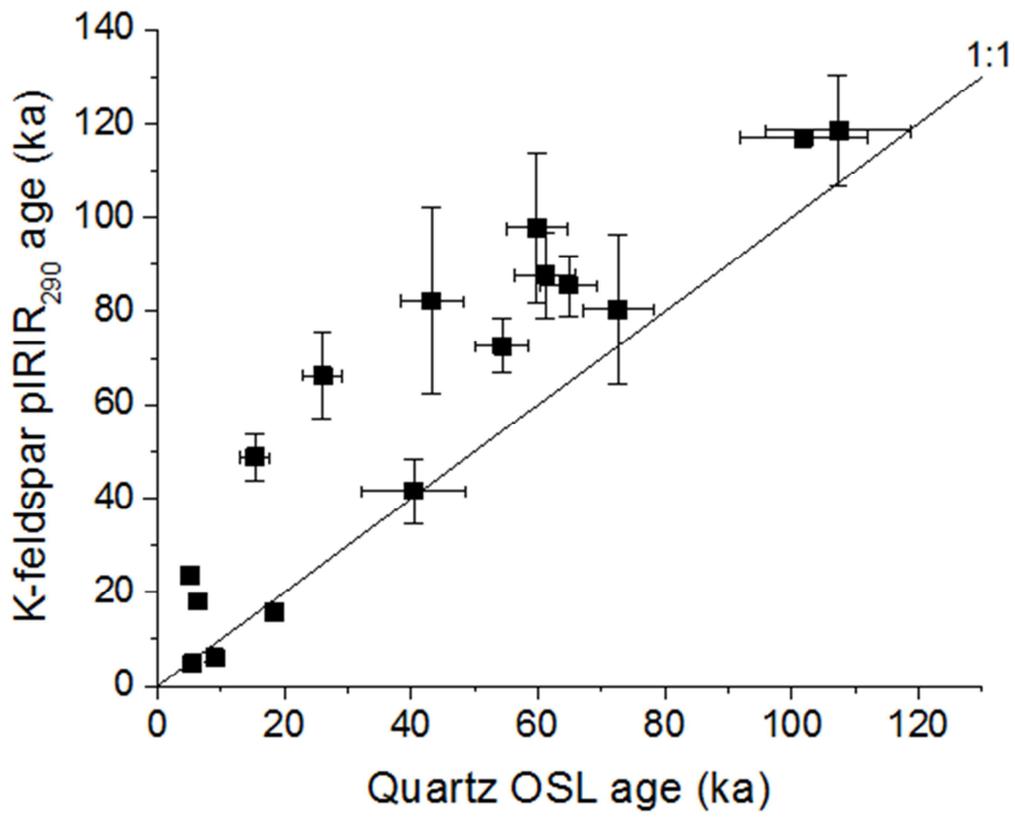


Figure 3c)

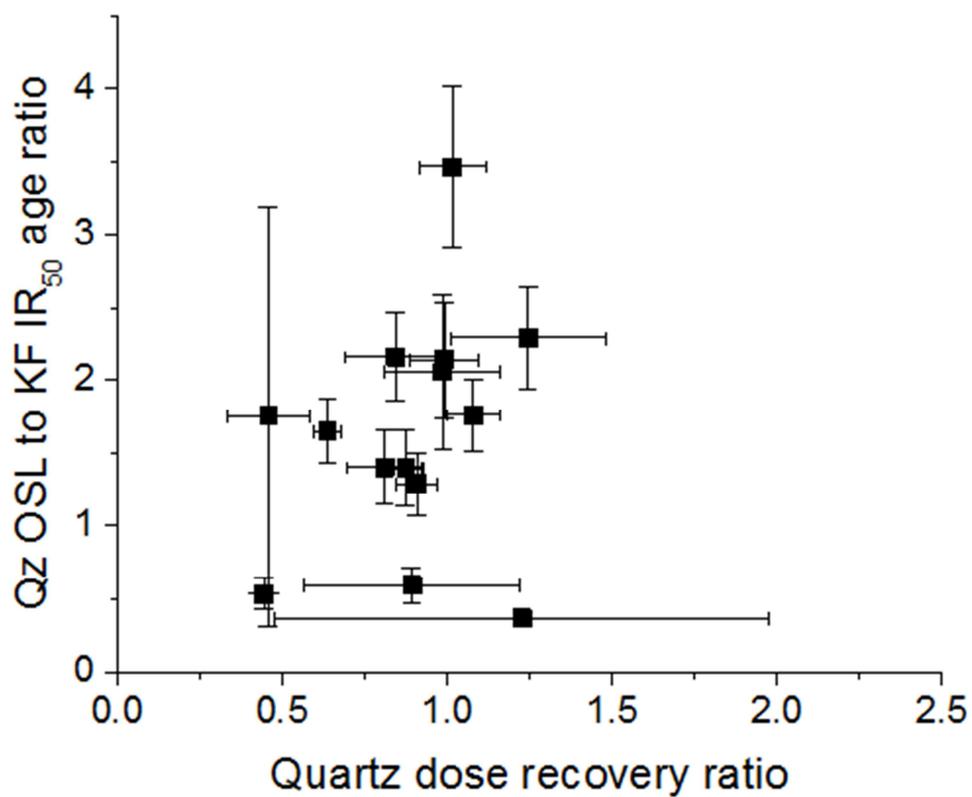


Figure 3d)

**Table 1**  
Sampling locations and pedological description of sections

Section	Longitude (E)	Latitude (N)	Altitud (m)	Mean annual precipitation (mm)	Mean annual temperature (mm)	Landform	local geology	Sediment origin geology
Rayen	57° 39' 21''	29° 27' 25''	2328	200	14	Pediment	Quaternary older gravel fan	Middle Eocene Igneous Rocks and Eocene marl
Bam	57° 39' 39''	29° 39' 55''	2096	145	15.5	Pediment	Upper Neogene to Quaternary conglomerate, sandstone	Cretaceous marl, sandstone and Paleocene to Eocene gypsiferous marl
Isfahan	51° 51' 21''	32° 40' 64''	1598	150	14.7	Alluvial fan	Oligo-Miocene limestone	Cretaceous limestone
Jiroft	57° 42' 50''	28° 37' 53''	660	185	24.5	fluvial plain	Recent alluvium, river terraces	Miocene to Pliocene marl, limestone and conglomerate
Lar	53° 57' 19''	27° 47' 23''	1071	170	23.3	Plateau	Miocene-Pliocene gypseous marl, calcareous sandstone	Eocene limestone and marl
Mahan	57° 17' 23''	30° 08' 38''	1912	160	14.5	Alluvial fan	Quaternary conglomerate and older fan conglomerate deposits	Upper Cretaceous limestone and gypsiferous marl

**Table 2**

Summary of sample code, burial depth, radionuclide concentrations, measured water content, quartz OSL and K-feldspar IR<sub>50</sub> and pIRIR<sub>290</sub> equivalent doses. Feldspar dose rates assume a K concentration of 12.5±0.5% for K-feldspar (Huntley and Baril, 1997). An absolute error of 4% is assumed on the water content values.

section	Sample code	Depth (cm)	Water content (%)	<sup>226</sup> Ra (Bqkg <sup>-1</sup> )±se	<sup>232</sup> Th (Bqkg <sup>-1</sup> )±se	<sup>40</sup> K (Bqkg <sup>-1</sup> )±se	Total dose rate (Gyka <sup>-1</sup> )±se	Quartz OSL De (Gy)±se	n	Quartz OSL age (ka)±se	K-feldspar IR <sub>50</sub> D <sub>e</sub> (Gy)±se	K-feldspar IR <sub>50</sub> age (ka)±se	K-feldspar pIRIR <sub>290</sub> D <sub>e</sub> (Gy)±se	n	K-feldspar pIRIR <sub>290</sub> age (ka)±se	Age ratio IR <sub>50</sub> /OSL	Age ratio pIRIR <sub>290</sub> /OSL	Well bleached	Probably well bleached
Rayen	177107	60	1	37.6±0.64	54.7±0.83	966±19.0	4.79±0.22	27±2	18	5.5±0.5	13.9±1.5	2.4±0.3	28±2.85	9	4.9±0.5	0.44±0.06	0.9±0.12	√	√
	177108	115	2	49.6±1.11	70.0±1.01	815±20.0	4.71±0.22	125±10	17	26±3	105.4±14	19±3	375±49	20	66±9	0.71±0.13	2.5±0.45		√
Bam	177111	70	1	19.5±0.48	14.7±0.37	256±8.0	1.64±0.07	14.7±1.5	17	9.0±1.0	6.6±0.86	2.6±0.3	15.8±2.4	17	6.1±1.0	0.29±0.04	0.68±0.13	√	√
	177112	150	4	22.3±0.45	19.5±0.40	310±8.0	1.84±0.08	112±7	16	61±5	103±9.9	37±4	243.9±22	11	88±9	0.61±0.08	1.4±0.18		√
Isfahan	177118	50	4	26.6±1.00	16.1±0.7	310±14.9	1.87±0.35	76±6	22	40±8	55±5	20±3	117±11.4	12	42±7	0.5±0.12	1.03±0.27	√	√
	177119	95	1	26.2±0.83	13.3±0.6	253±12.0	1.67±0.08	121±7	16	73±6	88.3±14.7	34±6	209±40.7	6	80±16	0.47±0.08	1.1±0.23	√	√
	177120	140	2	24.1±0.39	10.1±0.2	186±5.0	1.36±0.06	148±14	17	107±11	191.7±22	84±10	271.9±24.5	2	119±12	0.8±0.12	1.1±0.16	√	√
	177121	160	2	24.4±0.30	11.8±0.2	201±4.0	1.44±0.06	86±6	15	60±5	134.86±39.7	57±17	232.6±37	17	98±16	0.95±0.3	1.1±0.16	√	√
Jiroft	177125	60	4	17.3±0.81	14.0±0.6	313±14	1.65±0.08	8.5±0.9	16	5.2±0.6	36.1±2.6	14±1.2	61.4±3.9	17	24±1.8	2.7±0.38	4.6±	?	
	177126	85	3	15.2±0.25	13.6±0.2	328±5	1.65±0.07	10.3±1.8	16	6.2±1.1	27±2.2	10.5±0.9	46.6±4	14	18±1.7	1.67±0.33	2.9±	?	
	177127	115	1	11.9±0.26	11.3±0.3	290±6	1.46±0.06	22±3	24	15±2	69.8±8.7	29±4	116.8±10.6	18	49±5	1.9±0.37	3.3±	?	
Lar	177129	30	8	53.9±0.64	9.09±0.3	138±6	1.64±0.08	71±7	20	43±5	63.59±51.5	24.6±20	212±50.2	4	82±20	0.57±0.47	1.9±0.51		√
	177130	60	4	32.3±0.51	5.5±0.4	89±4	1.07±0.05	79±13	9	73±12	-	-	-	0	-	-	-		
	177128	95	5	35.02±0.61	6.18±0.35	99±6	1.2±0.05	122±10	14	102±10	-	-	249.14	1	117	-	1.15±0.11		
Mahan	177131	30	1	26.4±0.83	24.6±0.7	426±14	2.47±0.11	46±3	20	18.5±1.5	29±3.2	8.5±1.0	53.9±3.4	15	15.8±1.2	0.46±0.06	0.86±0.09	√	√
	177132	60	2	21.9±0.51	18.3±0.5	281±9	1.8±0.08	98±6	18	54±4	84.6±9.5	31±4	198.7±13	13	73±6	0.57±0.08	1.34±0.15		√
	177133	90	1	18.2±0.52	15.3±0.5	241±8	1.56±0.07	101±5	21	65±4	115.6±18.7	46±8	213±13.2	11	86±6	0.71±0.13	1.33±0.13		√

**Table 3**

Quartz OSL, feldspar IR<sub>50</sub> and feldspar pIRIR<sub>290</sub> apparent residual doses in modern analogue samples

Sample	Type	Quartz (Gy)	IR <sub>50</sub> (Gy)	pIRIR <sub>290</sub> (Gy)
Rayen	Alluvial	1.64± 0.3	0.7± 0.2	4.7± 1.9
Bam	Alluvial	0.11± 0.02	0.4± 0.2	1.9± 0.5
Jiroft	Fluvial	1.2± 1.01	1.4± 0.2	2.4± 0.5
Lar	Alluvial	0.11± 0.02	0.51± 0.15	2.8± 0.47
Mahan	Alluvial	0.1± 0.01	2.3± 1.9	3.5± 1.02

Supplementary materials

**Fig S1**

The photo shows the typical topography of our sampling sites



**Table S1**

Outline of the SAR OSL and post-IR IRSL protocol (Murray and Wintle, 2003; Buylaert et al. 2012)

step	Quartz		K-feldspar	
	Treatment	Observed	Treatment	Observed
1	Dose		Dose	
2	Preheat (240°C for 10 s)		Preheat (320°C for 60 s)	
3	Blue stimulation (125°C for 40 s)	$L_x$	Infrared stimulation (50°C for 100 s)	$L_x, IR_{50}$
4	-		Infrared stimulation (290°C for 100 s)	$L_x, pIRIR_{290}$
5	Test dose		Test dose	
6	Cut heat (290°C for 0 s)		Preheat (320°C for 60 s)	
7	Blue stimulation (125°C for 40 s)	$T_x$	Infrared stimulation (50°C for 100 s)	$T_x, IR_{50}$
8	-		Infrared stimulation (290°C for 100 s)	$T_x, pIRIR_{290}$
9	Blue stimulation (280°C for 40 s)		Infrared stimulation (325°C for 100 s)	
10	Return to 1		Return to 1	