Coordination and competition between magnetic particles driven by opposite climate transitions

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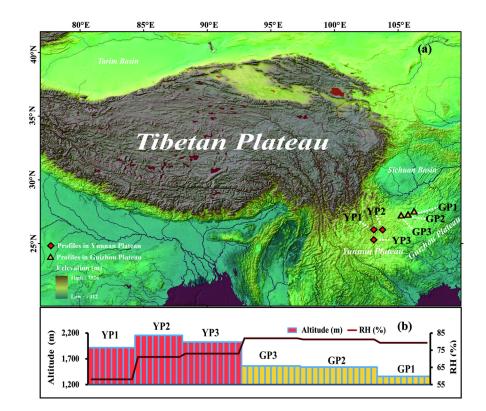
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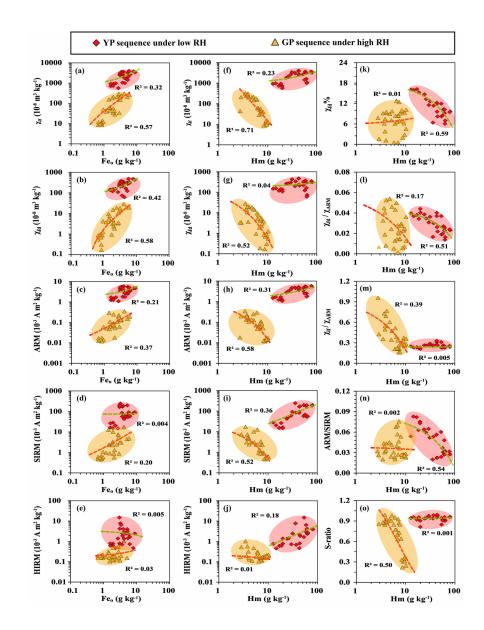
Abstract

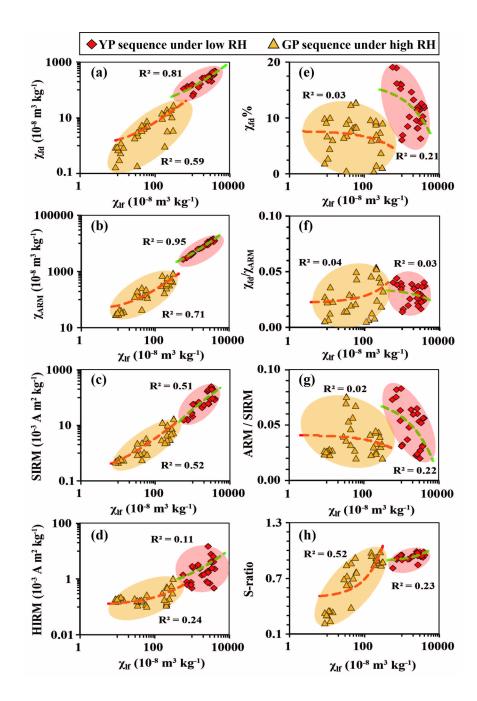
The ferrimagnetic (FM) and antiferromagnetic (AFM) particles of iron oxides are considered to be pedogenic and climatic indicators in soil taxonomy and paleoclimate reconstruction due to their enrichment trends as a function of increasing rainfall and temperature. However, opposite climate can retard chemical weathering but promote significant transformation between iron oxides, which could account for a nonlinear response of magnetism and color to extreme climate. We examined two soil sequences undergone opposite climate on the eastern edge of the Tibetan Plateau. The dry and warm climate transition favors the dehydration of amorphous iron oxides to form AFM hematite and FM particles, while the wet and cool climate transition impedes the formation but leads to their competition. The outcome well interprets the synchronous and asynchronous changes in color and magnetism under extreme opposite climate, and suggests that evaporation is as important as precipitation in extreme paleoclimate reconstructions based on iron oxides.

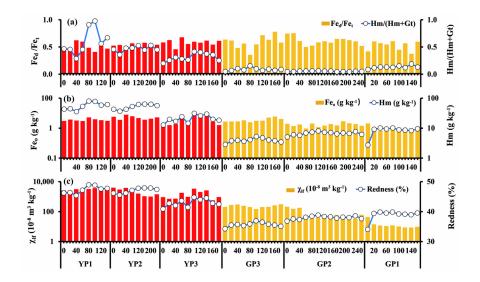
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3	driven by opposite climate transitions
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14	Key Points:
15	• Dry and warm climates favor the dehydration of amorphous iron oxides to
16	form antiferromagnetic hematite and ferrimagnetic particles
17	• Wet and cool climates retard their formation but lead to competition
18	• Evaporation is as important as precipitation in extreme climate cycles and
19	patterns reconstruction

20 Abstract

21 The ferrimagnetic (FM) and antiferromagnetic (AFM) particles of iron oxides are 22 considered to be pedogenic and climatic indicators in soil taxonomy and paleoclimate 23 reconstruction due to their enrichment trends as a function of increasing rainfall and temperature. However, opposite climate can retard chemical weathering but promote 24 25 significant transformation between iron oxides, which could account for a nonlinear response of magnetism and color to extreme climate. We examined two soil sequences 26 undergone opposite climate on the eastern edge of the Tibetan Plateau. The dry and 27 28 warm climate transition favors the dehydration of amorphous iron oxides to form 29 AFM hematite and FM particles, while the wet and cool climate transition impedes the formation but leads to their competition. The outcome well interprets the 30 synchronous and asynchronous changes in color and magnetism under extreme 31 32 opposite climate, and suggests that evaporation is as important as precipitation in 33 extreme paleoclimate reconstructions based on iron oxides.

Key words: Magnetism; Color; Iron oxides; Paleoclimate reconstruction 34

35

Plain Language Summary

Iron oxides are ubiquitous on the surface of Earth and Mars. They can be divided 36 into antiferromagnetic (AFM) and ferrimagnetic (FM) phases according to physical 37 properties. The former is found in colors ranging from red to yellow, while the latter is 38 39 the dominant form of magnetism in soils and sediments. Color and magnetism are considered sensitive pedogenic and climatic indicators in soil taxonomy and 40 paleoclimate reconstruction because iron oxides are commonly enriched as a 41

42	result of increasing regional rainfall and temperature. However, inverse changes in
43	temperature with rainfall can retard chemical weathering but promote significant
44	transformation between FM and AFM particles, which could result in a nonlinear
45	climatic response of soil color and magnetism to extreme climate cycles and patterns.
46	The uplift of the Tibetan Plateau (TP) has led to different orographic lifts in the
47	Yunnan Plateau (YP) and Guizhou Plateau (GP) on the eastern edge, which has
48	resulted in contrasting climate transitions on both plateaus. We found that the dry and
49	warm climate transition present in the YP is favorable to the dehydration of
50	amorphous iron oxides that then synchronously form FM particles and AFM hematite,
51	while the wet and cool climate transition present in the GP retards the formation but
52	leads to their competition. It well interprets synchronous and asynchronous change of
53	magnetism and color in dry and wet climate stages. Additionally, it also suggested that
54	evaporation is as important as precipitation when performing extreme paleoclimate
55	reconstruction based on iron oxides, especially during extreme climate cycles and
56	patterns.

57 1 Introduction

Iron oxides are ubiquitous on the surface of Earth and Mars [*Cornell and Schwtermann, 2003; Christensen et al., 2001*] and can be divided into chromogenic and magnetogenic groups according to their physical properties [*Long et al., 2011*]. The former includes antiferromagnetic (AFM) hematite (Hm, α -Fe₂O₃) and goethite (Gt, α -FeOOH) dominate optical properties, while the latter includes ferrimagnetic

63	(FM) magnetite (Mgt, Fe_3O_4) and maghemite (Mgh, γ -Fe ₂ O ₃), which dominate
64	magnetism in soils and sediments [Cornell and Schwartzman, 2003; Liu et al., 2012].
65	These particles are often synchronously enriched as immobile weathering products
66	under aerobic conditions with comparable increase in rainfall and temperature [Long
67	et al., 2011, 2016; Torrent et al., 2006]. Consequently, color and magnetism are
68	considered reasonable pedogenic and climatic indicators in soil taxonomy and
69	paleoclimate reconstruction if iron contents in parent materials are comparable
70	[Cornell and Schwartzman, 2003; Maher, 1998]. Over the past few decades, magnetic
71	properties have been successfully incorporated into paleorainfall reconstruction,
72	especially with aeolian sediments in the Chinese Loess Plateau (CLP) [Heller et al,
73	1991, 2010; Liu et al., 1995, 2003; Liu et al., 2007; Nie et al., 2008, 2013; Maher;
74	2016] and other temperate regions [Liu et al., 2001,2012; Chlachula, 2003].
75	Meanwhile, the color indices of soils have also been employed to reflect changes in
76	temperature [Yang et al., 2001; Yang and Ding, 2003].

However, growing evidences have been accumulated on asynchronous of color 77 78 indices and magnetism properties in soils [Han et al., 1996; Gao et al., 2018; Maher, 79 1998; Yang et al., 2001] and sediments at different scales, especially in loesses and paleosols layers of CLP, such as S1 [Liu and Ding, 1998], S5 [Guo et al., 2013], and 80 81 S9 [Xie et al., 2003], driven by extreme climate cycles, or deposits ranging from the 82 Tertiary red clay to the Quaternary loess driven by the dramatic climate pattern shifts of the CLP [Ji et al., 2004; Nie et al., 2014; Balsam et al., 2004; Hao et al., 2009]. 83 84 Therefore, the other soil chemical and mineral parameters have been introduced to

85	understand these shifting correlations. The Fe _d parameter indicating the total amount
86	of iron oxides has been applied to trace the changes in pedogenic intensity [Ding et
87	al., 2001]. The Fe _o parameter reflecting amorphous iron oxides was also introduced to
88	interpret the formation and transformation of FM particles [Hu et al., 2009].
89	Moreover, the ratio of Hm and Gt determined by diffuse reflectance spectra method
90	[Ji et al., 2001] is used to reconstruct changes in the relative humidity (RH) rather
91	than individual changes in rainfall and temperature [Ji et al., 2001, 2004; Balsam,
92	2004; Hao et al., 2009]. It was found that these extreme stages are often characterized
93	by high pedogenic intensity [Ding et al., 2002], lower iron oxide crystallinity [Hu et
94	al., 2009] and significant changes in Hm/(Hm+Gt) [Ji et al., 2004; Hao et al., 2009].
95	Theoretically, AFM Hm forms under warm, dry and seasonal climates, while Gt
96	forms under cool, wet and less seasonal climates [Schwertmann, 1985]. The change in
97	Hm/(Hm+Gt) can be promoted by inverse changes in rainfall and temperature [Long
98	et al., 2011, 2016]. Moreover, FM Mgh particles with differing sizes compete with
99	AFM Hm as an intermediate product from amorphous iron oxides under aerobic
100	conditions [Barrón and Torrent, 2002; Torrent, 2015], which also depends on the
101	formation efficiency of the Hm estimated by Hm/(Hm+Gt) [Hu et al., 2013; Long et
102	al., 2015]. However, the pedogenesis derived from aeolian sediments is disturbed by
103	the dust provenances [Li et al., 2009], deposition rates [Kukla, 1987], physical erosion
104	processes [Lu et al., 2006] besides chemical weathering controlled by specific
105	climates. Therefore, it is difficult to discern the independent contributions of climate
106	to pedogenic iron oxides and related changes in color and magnetism.

107 The Yunnan Plateau (YP) and Guizhou Plateau (GP) on the eastern edge of the 108 Tibetan Plateau (TP) have undergone differential uplifts and opposite climate 109 transitions at least since the Quaternary [Yang et al., 2010; Yan et al., 2011] due to the 110 uplift of the TP [*Pan et al., 2004*]. Moreover, the change in rainfall is accompanied by 111 the inverse pattern of temperature, which enhance the difference in the relative 112 humidity (RH) of the two plateaus. As a result, marked soil reddening in the YP and 113 yellowing in the GP has been observed and indicates that the iron oxide 114 transformations are driven by opposite climate transitions. These conditions provide a 115 good opportunity to understand the correlation between chromogenic and 116 magnetogenic iron oxides and their climatic implications in extreme wet-dry cycles 117 and patterns.

118 2 Materials and Methods

119 2.1 Geographical settings and soil sampling

120 The YP and GP belong to the tectonic extrusion zone of the TP [Bao et al., 121 2015]. Compared to the flatter TP with that has experienced rapid uplifting to an 122 average altitude of approximately 4000 m [Li and Fang, 1999], the surfaces of the 123 YP and GP are rugged with an average altitude of approximately 2000 m and 1100 m, respectively [Liu and Dong, 2013; Yang et al., 2010]. (Appendix A). The climate of 124 125 the YP is characterized by a higher mean annual temperature (MAT) of 13 $^{\circ}C \sim 20 ^{\circ}C$ 126 and a lower mean annual precipitation (MAP) of 600 ~ 900 mm/yr [Tong et al., 1994] than the GP characterized by a MAT of 12 $^{\circ}C \sim 16 ^{\circ}C$ and a MAP of 800 $\sim 1200 \text{ mm/}$ 127

128 yr [Liu and Xiong et al., 2015]. The opposite trends in MAT and MAP enlarge the 129 difference in the RH between the YP and GP from 50% to 85% [Xu, 1991]. Moreover, 130 it is near the climatic inflection point that controls soil reddening and yellowing, as proposed in our previous studies [Long et al., 2016]. As a result, the saprolitic soils 131 132 derived from the widespread Triassic carbonate rocks [Feng, 2005] have 133 demonstrated a common reddening trend $(3.2 \text{ YR} \sim 6.3 \text{ YR})$ in the YP due to its lower 134 RH from 58% to 73% while the soils have demonstrated a significant yellowing trend 135 $(6.2 \text{ YR} \sim 9.6 \text{ YR})$ in the GP due to its high RH from 79% to 82%.

136 We collected two saprolitic soil profile sequences from the YP and GP that are 137 separated by the boundary of the Wumeng Mountains. The profiles of YP1, YP2 and YP3 were collected from the YP under increasing MAT from 13.4 °C to 18.2 °C and 138 decreasing MAP from 924 mm/yr to 762 mm/yr. Similarly, the profiles of GP1, GP2 139 140 and GP3 were collected under slowly increasing MAT from 12.9 °C to 14.4 °C and a 141 decreasing MAP from 937 mm/yr to 899 mm/yr. These profiles were collected on the 142 local highland and were covered by natural vegetation ranging from herbaceous plants 143 in YP to evergreen forests in GP. The soil type in the YP can be categorized as an Acrisol, while that in the GP can be categorized as an Alisol [IUSS Working Group] 144 WRB, 2015]. The soil samples were collected from the surface to the bottom of 145 146 outcrops at intervals of 20 cm or 40 cm covering the main horizons depending on the thickness of the outcrops. 147

148 2.2 Chemical and physical measurement

149 The air-dried samples were sieved by a 2-mm sieve and ground into powder for 150 chemical analysis. The chemical compositions were determined with an ARL9800XP 151 + X-ray spectrophotometer and have been expressed as oxides. The chemical 152 weathering index (CIA) was calculated as the molar percentage of $Al_2O_3 / (Al_2O_3 +$ $CaO + Na_2O + K_2O$ [Nesbitt and Young, 1982], and the Sa index was calculated as 153 154 the molar ratio of SiO₂/Al₂O₃ [Bayan and Ruxton, 1968]. Free iron (Fe_d) and amorphous iron (Fe_o) were extracted by the citrate-bicarbonate-dithionate (CBD) 155 156 method [Mehra and Jackson, 1960] and ammonium oxalate method [Schwertmann, 157 1964], respectively.

158 The diffuse reflectance spectra (DRS) were measured with a Perkin Elmer 159 Lambda 900 UV/VIS/NIR spectrometer at 2 nm intervals. The standard Hm and Gt 160 minerals used in the experiment were the Pfizer R1599 pure red from Pfizer Company 161 and Synox HY610 pure yellow nanoscale iron oxides from the Hoover Color Corporation. The redness was calculated according to the ratio of mean reflectance 162 163 between the red-light band ($630 \sim 700 \text{ nm}$) and the visual light band ($400 \sim 700 \text{ nm}$) 164 [Judd and Wyszecki, 1975]. The Hm content was estimated using a working curve 165 established by the sample substrate after CBD treatment mixed with a series of standard Hm and Gt samples in different ratios [Long et al., 2011, 2016]. Finally, we 166 167 assigned Fe_d to be the combination of Fe in stoichiometric Hm, Gt [Torrent et al., 168 2007] and Fe_o, and the contents of Hm and Gt were calculated by the following 169 equation:

170 Hm
$$(g kg^{-1}) = 0.0012 \times e^{0.227 * Redness}$$

171 Gt
$$(g kg^{-1}) = 1.59 \times (Fe_d - Fe_o - Hm/1.43)$$

172 The magnetic susceptibility of all samples was measured in the laboratory at 0.47 173 kHz (χ_{lf}) and 4.7 kHz (χ_{hf}) with a Bartington MS2B susceptibility meter. The 174 frequency-dependent susceptibilities χ_{fd} and χ_{fd} % representing the absolute and relative contributions of SP particles were calculated as $\chi_{lf}-\chi_{hf}$ and $(\chi_{lf}-\chi_{hf})/\chi_{lf}\times$ 175 176 100%, respectively [Dearing et al., 1996; Worm, 1998]. Meanwhile, the anhysteretic 177 remanent magnetization (ARM) was imparted using a peak of the 100 mT alternating 178 field and a 0.05 mT biasing field with a Molspin demagnetizer [Dunlop and 179 *Özdemir*, 1997]. The χ_{ARM} parameter was calculated by the ARM and normalized by 180 the biasing field. The saturation isothermal remanent magnetization (SIRM) was 181 attained at 1 T with the ASC-10 impulse magnetizer, and all the remanence 182 magnetizations were measured in the AGICO JR6 spinner magnetometer. HIRM was 183 calculated by $(IRM_{-300mT} + SIRM)/2$, which mainly reflects the content changes of 184 high-coercivity minerals, such as Hm [Nie et al., 2010], and the S-ratio is calculated 185 by -IRM._{300mT}/SIRM [King et al., 1991], which indicates the relative abundance of the 186 FM to AFM minerals [Thompson and Oldfield, 1986; Liu et al., 2007].

187 **3. Results**

As illustrated in **Figure 1**, the change of Fe_d/Fe_t is comparable in both sequences. In contrast, the Hm/(Hm+Gt) demonstrates a significant shift from 0.20 to 0.98 in the YP sequence but remains low from 0.05 to 0.19 in the GP sequence (**Figure 1a**). In

191	addition, the Fe_o content is also little higher in the YP sequence than those in the GP
192	sequence. However, the Hm content in the YP sequence is much higher than that in
193	the GP sequence (Figure 1b). Correspondingly, the χ_{1f} ranges from 573.9×10 ⁻⁸ m ³ kg ⁻¹
194	to 4005.1×10^{-8} m ³ kg ⁻¹ in the YP sequence, which is much higher than that ranging
195	from 8.7×10^{-8} m ⁻³ kg ⁻¹ to 310.5×10^{-8} m ³ kg ⁻¹ in the GP sequence (Figure 1c). More
196	importantly, χ_{lf} changes in phase with redness control by Hm in the YP sequence but
197	out of phase in the GP sequence. The χ_{fd},χ_{ARM} and SIRM, also exhibit synchronous
198	changes with χ_{ff} (Figure 2a-2c), although the χ_{fd} %, χ_{fd}/χ_{ARM} , ARM/SIRM are more
199	comparable across both sequences (Figure 2e-2g). $\chi_{\rm fd}\%$ and ARM/SIRM are only
200	slightly higher in the YP sequence, while χ_{fd}/χ_{ARM} is slightly higher in the GP
201	sequence. In addition, the HIRM is higher in the YP sequence than that in the GP
202	sequence (Figure 2d), but it has a less correlation with the increasing χ_{lf} . The <i>S-ratio</i>
203	exhibits a rapid increase with increasing $\chi_{\rm lf}$ in the GP sequence but remains close to 1
204	in the YP sequence (Figure 2h).

205 4. Discussion

4.1 Comparable chemical weathering and significant iron oxides transformation driven by opposite climate transitions

Theoretically, the increasing of rainfall and temperature favors chemical weathering and the enrichment of iron oxides because primary iron-bearing minerals are often preferentially weathered to form secondary iron oxides [*Kump et al., 2000*]. However, in the GP under the wet and cool climate, the higher rainfall is

superimposed by lower evaporation. Although the changed leaching could be 212 213 enhanced by the more effective rainfall, the chemical reaction rates should be retarded by the lowering temperature [Kump et al., 2000; White and Blum, 1995]. The 214 215 phenomena can be widely observed in mountainous, where the rainfall also 216 accompanied by the inverse change of the temperature [Long et al., 2016]. 217 Nevertheless, the effective rainfall would favor the formation of iron oxyhydroxides 218 such as Gt [Schwertmann, 1971]. However, in the YP under the dry and warm climate, 219 the lower rainfall superimposed by the increased evaporation retards chemical 220 leaching but promotes the dehydration of amorphous iron oxides to form iron oxides 221 like Hm and Mgh [Schwertmann, 1971; Barrón and Torrent, 2002; Grogan et al., 222 2003]. Generally, these inverse changes of temperature with rainfall lead to less 223 variability in chemical weathering intensity and more significant differentiation of 224 iron oxides indicated by Hm/(Hm+Gt) in both sequences.

225 4.2 Coordination and competition between FM and AFM particles driven by 226 opposite climate transitions

The magnetic properties revealed a common positive correlation with Fe_o in both sequences except that the SIRM and HIRM reveals more consistent correlation with Fe_o (Figures 3a-3e). Although the contents of Hm and FM particles are much higher in the YP sequence than these in the GP sequence (Figures 3f-3j). However, the Hm and FM particles change in phase in the YP sequence but out of phase in the GP sequence (Figures 3f-3i) except that the HIRM reveals more consistent correlation with Hm (Figure 3j). Moreover, the χ_{fd}/χ_{ARM} , χ_{Jf}/χ_{ARM} in the GP soils (Figures 3l-3m), as well as the χ_{fd} %, χ_{fd}/χ_{ARM} and ARM/SIRM in the YP soils, which indicate the ratio of fine FM particles to coarser FM particles both decrease with Hm in both sequences (Figures 3k-3n). However, the *S-ratio* reveals significant decreasing with Hm in the GP yellowing soils but remains constant in the YP reddening soils (Figure 3o).

238 These outcomes verify the FM particles with growing size as the intermediate 239 products of AFM Hm aging from amorphous iron oxides [Barrón and Torrent, 2002] 240 but the correlation between FM particles and Hm depends on Hm/(Hm+Gt) controlled 241 by RH [Torrent et al., 2006; Long et al., 2015]. The positive correlation between FM 242 particles and Hm, as revealed in the YP reddening soils, occurs under the condition 243 with a high formation efficiency of Hm, indicate by Hm/(Hm+Gt) from 0.20 to 0.98 under the low RH. It is consistent with the result revealed in aerobic soils with high 244 Hm/(Hm+Gt) [Torrent et al., 2006], especially in the red Ferralsols derived from 245 246 basalt with the Hm/(Hm+Gt) above 0.6 [Long et al., 2015]. However, the negative 247 correlation between Hm and FM particles, as revealed in the GP yellowing soils, 248 occurs under the conditions with a low Hm/(Hm+Gt) from 0.05 to 0.19 under high RH. These negative correlations can be observed in each profile of the GP sequence 249 250 (Figure 1c). This result apparently accords with the yellow soils derived from the 251 downslope of a subtropical granitic toposequence, with Hm/(Hm+Gt) < 0.2 and Hm%<1% controlled by the downward increasing of water activity [*Guo et al., 2021*]. 252 253

However, in contrast to the topsequence affected by the dynamic migration of magnetic particles [*Guo et al., 2021*]. The negative correlation between FM particles

and Hm in the climosequence of the GP under high RH indicates there would be a 255 256 competition between FM particles and Hm since high soil water activity retards the 257 formation of iron oxides but promote the formation of iron oxyhydroxides [Tardy and 258 Nahon, 1985; Trolard and Tardy, 1987]. Moreover, with the slow increasing of RH 259 from GP1 to GP3, a little amount of FM particles has accumulated at the cost of Hm. 260 The outcome confirms the FM particles as rainfall indicator while the Hm as temperature or evaporation indicator at a large scale [Gao et al., 2018]. If the different 261 262 aging processes under aerobic conditions in natural systems of Hm and Gt from Fh

are combined as Gt 1 Fh 2 SP Mgh3 SD Mgh4 Hm [Schwertmann, 1971; Barrón and

Torrent, 2002], in the wet and cool climate transition, the step 1 and step 2 are favored, which results in the accumulation of a large amount of Gt and a limited number of FM particles at the cost of previously formed Hm. However, in the dry and warm climate transition with low RH, the step 3 and step 4 are favored, which results in the accumulation of a large number of FM particles, as well as their significant grain growth and transformation into Hm.

270 4.3 Significance in paleoclimate cycle and pattern reconstruction

In both sequences under high and low RH, an increase in rainfall is often accompanied by decreasing temperature. It could help us understand the response of iron oxide to extreme dry-wet cycles and patterns. In paleoclimate reconstruction, changes in rainfall are often considered to be accompanied by synchronous changes in temperature [*Liu et al., 2001, 2012; Kukla, 1987; Maher, 1998, 2016*]. This climate 276 pattern could lead to remarkable changes in the Fe_d content controlled by chemical 277 weathering that constrains the changes in the Hm/(Hm+Gt) controlled by RH [Ji et al., 2004; Balsam et al., 2004; Ding et al., 2001]. However, if the temperature and 278 279 rainfall changes in opposing directions, a significant change in Hm/(Hm+Gt) can be 280 observed, and the formation efficiency of Hm and FM particles and their correlation 281 could change, resulting in the mismatching of color indices and magnetic properties. 282 Actually, the magnetism and redness are often coupled in loess deposits under dry and 283 cool climates [Ji et al., 2004], but they are frequently observed as decoupled in 284 paleosols under warm and wet climate, with Hm/(Hm+Gt) commonly decreasing 285 below 0.2 [Ji et al., 2004; Hao et al., 2009]. This phenomenon agrees with the 286 competition occurring between Hm and FM particles under the high RH present in the 287 GP sequence with low Hm/(Hm+Gt). In addition, the change content of FM particles

estimated by
$$\chi_{\rm lf}$$
 (approximately 30×10⁻⁸ m³ kg⁻¹ ~ 190×10⁻⁸ m³ kg⁻¹) [*Ji et al., 2004*]

well matches the content changes in the hematite content (approximately 0.1% ~ 0.2%) [*Guo et al.*, 2021; *Ji et al.*, 2004] if the $\chi_{\rm lf}$ of pure FM particles are estimated around 110, 000×10⁻⁸ m³ kg⁻¹ [*Worm and Jackson, 1999*].

However, in some warm stages, such as red clays with Hm/(Hm+Gt) above 0.6 [*Hao et al., 2009*], uncertain and even opposite correlation between redness and magnetism were also found [*Ding et al., 2001; Nie et al., 2008; Hu et al., 2009*]. In the YP sequence as well as red Ferrosols with high Hm/(Hm+Gt) under low RH and 296 high evaporation, the positive correlation between FM particles and Hm still remains 297 although the formation of fine FM particles are observed to deaccelerate with Hm 298 [Long et al., 2015]. Therefore, the opposite correlation between redness and 299 magnetism in Tertiary Red Clay could also correlate with the increased iron 300 crystallinity with longer aging time [Hu et al., 2009; Jiang et al., 2018], less ligand protection of highly weathering soils [Ren et al., 2020] in addition to high 301 Hm/(Hm+Gt). Nevertheless, it verifies the magnetic parameters indicating the ratios 302 303 of different FM particles could be more reasonable than the magnetic parameters 304 indicating the contents of magnetic particles in paleoclimate reconstruction under 305 widely climate scale [Nie et al., 2014].

306 In addition, it should be noted that the soil sequences exhibit comparable changes in the Fe_d contents but significant changes in the Hm/(Hm+Gt). The higher 307 308 redness and degree of magnetism are observed in the YP sequence under dry and 309 warm climate. In contrast, under comparable changes in rainfall and temperature on 310 the CLP, the Fe_d of the paleosols is often two times that of loesses [*Ding et al., 2001*], 311 while the Hm/(Hm+Gt) is slightly lower in paleosols [Ji et al., 2004]. The higher 312 redness and degree of magnetism are observed in paleosols formed under a wet and 313 warm climate. Since the soil reddening and magnetic enhancement could be achieved 314 by the dehydration of iron oxides associated with strong evaporation [Barrón and 315 Torrent, 2002; Long et al., 2015] even when chemical weathering is restrained by low 316 rainfall. The accompanying change in temperature should be considered as important 317 as rainfall in paleoclimate reconstruction, especially under extreme climate cycles and

318 patterns shifts.

319 **5.** Conclusion

320 To unravel the changing relationship between AFM and FM particles and their 321 climatic implications under extremely climate, we examined two soil sequences in the 322 YP and GP on the eastern edge of TP. The YP and GP sequences have undergone 323 significant reddening and yellowing, respectively, as a result of dry and wet climate 324 transitions accompanied by inverse changes in temperature and rainfall. The AFM 325 Hm and FM particles are much more enriched in the YP reddening soils than these in 326 the GP yellowing soils although the change in total amount of iron oxides controlled 327 by chemical weathering are comparable. The dry and warm climate favors the 328 dehydration of amorphous iron oxides to form higher contents of AFM Hm and FM 329 particles, while the wet and cool climate impedes their formation and leads to 330 competition. The little amount of FM particles could form at the cost of previously 331 formed Hm under high RH. The model well interprets the synchronous and 332 asynchronous changes in color and magnetism under the dry and wet cycles, and 333 suggests that evaporation is as important as precipitation in extreme paleoclimate reconstruction based on iron oxides. 334

335

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- 341 paper can be accessed through the public domain repository Zenodo at
- 342 http://doi.org/10.5281/zenodo.4495880

Appendix A: (a) Location of sampling points on the YP and GP; (b) The YP is
characterized by higher elevation and lower RH than the GP. The profiles of YP1,
YP2, YP3 and GP1, GP2, GP3, with increasing relative humidity in the sequences
were sampled in the YP and GP, respectively.

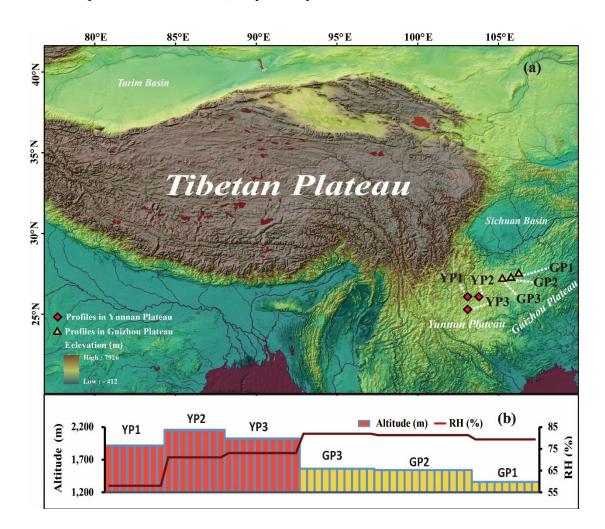
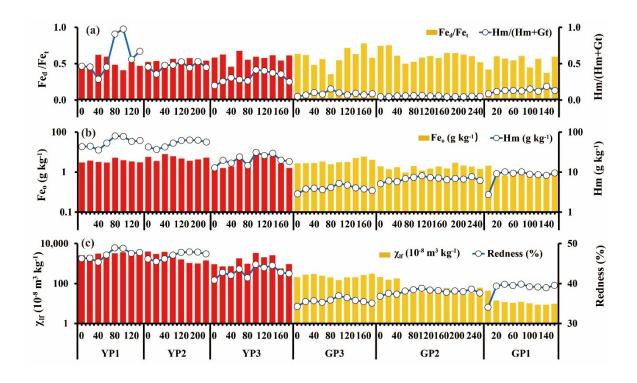
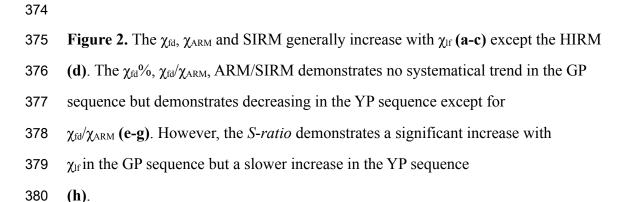


Figure 1. The Fe_d/Fe_t keeps comparable in both sequences but the Hm/(Hm+Gt) is

- 358 much higher in the YP sequence than that in the GP sequence
- **359** (a). The Fe_0 in the YP sequence is a little higher than that in the GP sequence while
- 360 the Hm is significantly higher in the YP than that in the GP
- 361 (b). The magnetic susceptibility changes in phase with the redness in the YP sequence
- but out of phase in the GP sequence (c). Note that the Fe_0 , Hm and χ_{lf} are shown in
- 363 logarithmic form because of the significant difference between profiles.

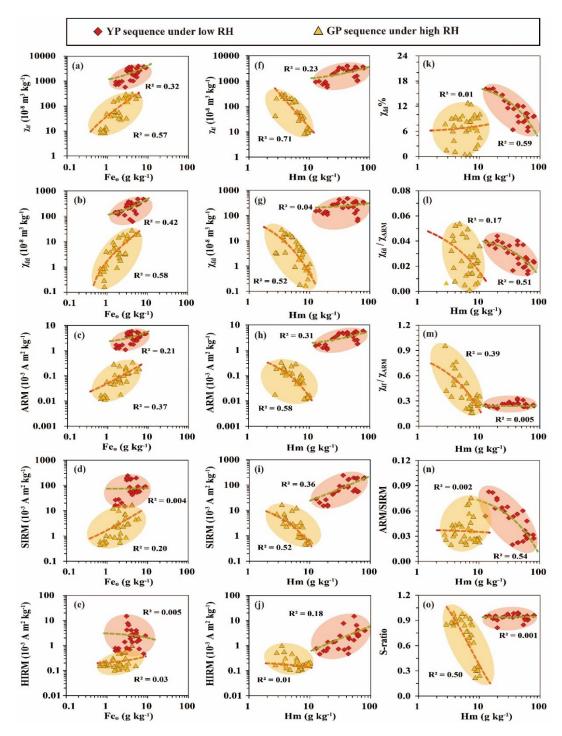






381 YP sequence under low RH ▲ GP sequence under high RH 1000 20 382 (a) (e) $R^2 = 0.81$ $\chi_{fd} (10^{-8} \text{ m}^3 \text{ kg}^{-1})$ 383 $R^2 = 0.03$ χ_{fd}% 10 10 384 385 $\mathbf{R}^2 =$ 0.2 $R^2 = 0.59$ 0.1 0 386 100 10000 100 10000 1 1 χ_{lf} (10⁻⁸ m³ kg⁻¹) χ_{lf} (10⁻⁸ m³ kg⁻¹) 387 100000 0.10 (f) **(b)** $\chi_{ARM} (10^{-8} \text{ m}^3 \text{ kg}^{-1})$ 388 $R^2 = 0.95$ Xfd/Xarm 0.05 389 $R^2 = 0.04$ = 0.03 1000 390 = 0.71 R² 391 0.00 10 10000 100 1 1 100 10000 392 χ_{lf} (10⁻⁸ m³ kg⁻¹) χ_{lf} (10⁻⁸ m³ kg⁻¹) 0.10 1000 393 SIRM (10⁻³ A m² kg⁻¹) (c) (g) $R^2 = 0.51$ $R^2 = 0.02$ **ARM / SIRM** 394 0.05 10 395 396 $R^2 = 0.52$ R² = 0.220.1 0.00 397 100 10000 100 1 1 10000 χ_{lf} (10⁻⁸ m³ kg⁻¹) χ_{lf} (10⁻⁸ m³ kg⁻¹) 398 1.3 100 HIRM (10⁻³ A m² kg⁻¹) (d) (h) $R^2 = 0.11$ 399 $R^2 = 0.52$ S-ratio 400 0.7 1 $R^2 = 0.23$ 401 $R^2 = 0.24$ 402 0.01 0.1 10000 100 10000 100 1 1 403 χ_{lf} (10⁻⁸ m³ kg⁻¹) χ_{lf} (10⁻⁸ m³ kg⁻¹)

Figure 3. The magnetic parameters generally increase with Fe_o (**a-e**). Both the FM particles and Hm are more enriched in the YP than that in the GP but they change in phase in the YP and out of phase in GP except for the HIRM (**f-j**). The χ_{fd}/χ_{ARM} , χ_{If}/χ_{ARM} decreases with Hm in the GP while χ_{fd} %, χ_{fd}/χ_{ARM} and ARM/SIRM decreases with Hm in the YP and the *S-ratio* reveals significant decreasing with Hm in the GP but remains constant in the YP (**k-o**).



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