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### Abstract

A paleosol from the Middle Pleistocene lacustrine-fluvial succession of Sulmona Basin, central Italy, was analysed for the land snail shell content, and the stable isotope composition of the shells and associated pedogenic carbonates. The paleosol – known as Fiorata Paleosol – is covered by a thick tephra layer dated to ca. 527 ka allowing the pedogenetic horizons to be correlated to the marine isotope stage (MIS) 14-early MIS 13 interval. The terrestrial mollusc assemblage contained few individuals and was characterized by a low number of species which predominantly indicate open and dry habitats, thus suggesting that Fiorata Paleosol likely developed during glacial conditions of the MIS 14. The  $\delta^{13}\text{C}$  values of pedogenic carbonates and terrestrial shells indicate prevailing C3-type vegetation, probably marked by some degree of water stress. Calculation of the  $\delta^{18}\text{O}$  precipitation values, derived from pedogenic carbonates and shell  $\delta^{18}\text{O}$  values, indicate that the average temperature was 3-5°C lower than present day. This study highlights how paleosols, despite offering only snapshots of past climate and environments, provide valuable complementary information to paleoclimatic data obtained in the adjacent lacustrine intervals, specifically for the Sulmona successions.

<b>Keywords</b>	tephra layers; paleosol; C3-type vegetation; Glacial
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Dear Editor

We would like to submit the manuscript entitled: “Middle Pleistocene (MIS 14) environmental conditions in the central Mediterranean derived from terrestrial molluscs and carbonate stable isotopes from Sulmona Basin (Italy)” by Zanchetta et al.

The manuscript deals with paleontological and geochemical investigation on a paleosols from the Sulmona basin, developed during the late part of MIS14, as constrained by volcanic layers.

Sincerely

Giovanni Zanchetta


# Middle Pleistocene (MIS 14) environmental conditions in the central Mediterranean derived from terrestrial molluscs and carbonate stable isotopes from Sulmona Basin (Italy)

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## Abstract

A paleosol from the Middle Pleistocene lacustrine-fluvial succession of Sulmona Basin, central Italy, was analysed for the land snail shell content, and the stable isotope composition of the shells and associated pedogenic carbonates. The paleosol – known as Fiorata Paleosol – is covered by a thick tephra layer dated to ca. 527 ka allowing the pedogenetic horizons to be correlated to the marine isotope stage (MIS) 14-early MIS 13 interval. The terrestrial mollusc assemblage contained few individuals and was characterized by a low number of species which predominantly indicate open and dry habitats, thus suggesting that Fiorata Paleosol likely developed during glacial conditions of the MIS 14. The  $\delta^{13}\text{C}$  values of pedogenic carbonates and terrestrial shells indicate prevailing  $\text{C}_3$ -type vegetation, probably marked by some degree of water stress. Calculation of the  $\delta^{18}\text{O}$  precipitation values, derived from pedogenic carbonates and shell  $\delta^{18}\text{O}$  values, indicate that the average temperature was 3-5°C lower than present day. This study highlights how paleosols, despite offering only snapshots of past climate and environments, provide valuable complementary information to paleoclimatic data obtained in the adjacent lacustrine intervals, specifically for the Sulmona successions.

*Keywords:* tephra layers; paleosol;  $\text{C}_3$ -type vegetation; Glacial;

## 1. Introduction

Stable isotopes (e.g.  $^{13}\text{C}/^{12}\text{C}$  and  $^{18}\text{O}/^{16}\text{O}$ ) of pedogenetic carbonate (Cerling, 1984; Jiamao et al., 1991; Cerling and Quade, 1993; Zanchetta et al., 2000) and terrestrial mollusc shells (e.g. Balakrishnan et al., 2005a; Colonese et al., 2007, 2010; Yanes et al., 2011) can provide valuable snapshots of past environments, notably vegetation type (e.g.  $\text{C}_3/\text{C}_4$  ratios, Cerling and Quade, 1993; Zanchetta et al., 2006), and past precipitation regimes (Lecolle, 1985; Zanchetta et al., 2005). Despite representing important sources of complementary paleoclimatic information, combined studies on molluscan association and stable isotope analysis of pedogenic and shell carbonates are scarce (e.g. Zanchetta et al., 2006; Leone et al., 2000; Balakrishnan et al., 2005a).

Furthermore, whilst the Holocene and Late Pleistocene terrestrial mollusc assemblages of the Mediterranean and continental Europe are rather well known and studied (e.g. Kerney, 1976; Esu, 1981; Rousseau et al., 1992; Limondin-Lozouet and Antoine, 2006; Limondin-Lozouet et al., 2016), the Middle Pleistocene successions are relatively rare and often chronologically poorly constrained (e.g. Rousseau and Keen, 1989; Limondin-Lozouet and Preece, 2004). In particular, terrestrial molluscs of glacial periods have been previously described for the Last Glacial in Europe from loess successions (e.g. Ložek, 1990; Rousseau et al., 1990; Moine, 2008), but for older glacial intervals our knowledge is essentially fragmentary and incomplete.

In this paper we discuss the terrestrial mollusc assemblages and stable isotope geochemistry ( $^{13}\text{C}/^{12}\text{C}$  and  $^{18}\text{O}/^{16}\text{O}$  ratios) of their shells and of associated pedogenic carbonates from a Middle Pleistocene paleosol developed within a fluvial to lacustrine succession at the Sulmona Basin (Abruzzo, central Italy, Fig. 1). The lacustrine successions have been extensively investigated and yielded important insights into past climate conditions in the Central Mediterranean between the late Early to the Late Pleistocene (Giaccio et al., 2015; Regattieri et al., 2015,2016,2017). However, stratigraphic evidence indicates that the lake level was substantially lower during some glacial intervals, and subaerial processes (fluvial-colluvial deposition, erosion and/or pedogenesis) dominated with respect to the lacustrine sedimentation (Giaccio et al., 2015; Regattieri et al., 2015,2016,2017). Pedogenic horizons represent alternative sources of information to complement paleoclimatic inferences derived from lacustrine sediments at Sulmona Basin.

## 2. Site description

The Sulmona Basin (Fig. 1) is an intramontane depression formed during the Plio-Quaternary extensional tectonic phase that dissected the earlier orogenic, fold-and-thrust-belt system of the Apennine chain (e.g. Patacca and Scandone, 2007). The progressive formation of the basin was

68 driven by the Sulmona or Morrone NW–SE-trending fault system (Galli et al., 2015),  
69 accommodating the volume for the accumulation of a thick Quaternary succession (e.g. Cavinato  
70 and Miccadei, 2000; Giaccio et al., 2012, 2013b). The Pleistocene succession is subdivided in three  
71 main unconformity-bounded alluvial–fluvial–lacustrine units; SUL-6, SUL-5 and SUL 4-3 (Figs. 1,  
72 2). Each unit is constrained by  $^{40}\text{Ar}/^{39}\text{Ar}$  dating of the tephra layers and magnetostratigraphy  
73 (Giaccio et al., 2012, 2013b; Sagnotti et al., 2014; Figs. 1, 2). The paleosol discussed in this paper  
74 (hereafter referred to as Fiorata Paleosol, from the toponymal of the type section; Fig. 1) occurs  
75 immediately above the lower boundary of SUL-5 and formed on the gravel-sand succession which  
76 fills a deep fluvial incision carved into the underlying SUL-6 unit (Figs. 1, 2). The pedogenic  
77 horizon consists of ca. 40-50 cm of gray-dark-grayish brown (10YR 4/2 -4/1 dark) silty to coarse  
78 massive sands A horizon (Fig. 2). The lower boundary fades in the lower fluvial interval through a  
79 massive, bioturbated C horizon. Thin root traces are preserved in the upper horizon, sometime with  
80 walls impregnated by oxides. Small carbonate concretions are also visible. The Fiorata Paleosol is  
81 directly capped by a syn-depositionally reworked tephra layer (SUL5b-12 in Fig. 2) up to 0.5 m-  
82 thick of fine lapilli to coarse ash made up of green, porphyritic and finely grained micro-scoria (Fig.  
83 2). Some fine volcanic ash fills small burrows and/or root traces within the upper soil horizon.  
84 Based on its peculiar foiditic composition of the glass from the layer, SUL5b-12 (Fig. 2) was  
85 correlated to the Tufo di Bagno Albule eruption by Giaccio et al. (2013b), from Colli Albani  
86 volcanic district dated by  $^{40}\text{Ar}/^{39}\text{Ar}$  to  $527\pm 2$  ka (Marra et al., 2009).

87 A detailed description of climatic and hydrological settings of the Sulmona Basin can be found  
88 in Regattieri et al. (2015,2016,2017) and is only briefly summarized here. The Sulmona  
89 meteorological station (ca. 400 m a.s.l.) records a mean annual temperature of  $13.7^\circ\text{C}$ , and an  
90 average rainfall of ca. 700 mm. Precipitation is strongly influenced by local topography and by the  
91 rising margin of the tectonic basin, reaching values of about 1200 mm at mountain summits. Winter  
92 precipitation is largely regulated by conditions in the North Atlantic and the North Atlantic  
93 Oscillation (López-Moreno et al., 2011). The meteoric precipitation at the nearby L'Aquila Station  
94 (ca. 710 m a.s.l.) has an average  $\delta^{18}\text{O}$  value of  $\text{‰}7.13\text{‰}$  (Longinelli and Selmo, 2003). The  
95 measured isotopic altitudinal gradient ( $\delta^{18}\text{O}/100$  m) ranged from  $\text{‰}0.23$  to  $\text{‰}0.13$  ‰/100 m (e.g.  
96 Barbieri et al., 2005; Giustini et al., 2016).

97  
98

### 3. Material and Methods

99 Two distinct sedimentary samples (SUL-16/01 and SUL-16/02) of ca. 10 kg each were collected  
100 from two different positions along the section exposing the Fiorata Paleosol (Figs. 1 and 2).  
101 Samples were first dried at room temperature for several days and then disaggregated using a very

102 dilute solution of H<sub>2</sub>O<sub>2</sub> (ca. 5%) and deionized water. The material was gently washed and sieved  
103 using 2000, 1000, 500 and 250 µm mesh screens, and all the identifiable shells and fragments were  
104 picked out under a binocular microscope and counted using the convention of Sparks (1961), where  
105 every gastropod apex is recorded to give a minimum number of individuals. The higher systematics  
106 followed Bouchet and Rocroi (2005) except for the helicoideans, for which the revision by Razkin  
107 et al. (2015) was adopted. The taxonomy and nomenclature of the extant species followed Welter-  
108 Schultes (2012).

109 During mollusc picking, small carbonate concretions (mm-sized) were also found and selected  
110 for isotopic analysis. Well preserved shells of a helicelline geomitrid species and small carbonate  
111 concretions were soaked in a solution of distilled water and H<sub>2</sub>O<sub>2</sub> (30%) and sonicated to remove  
112 contaminants. Samples were then dried, powdered and homogenized for stable isotope analyses.  
113 Samples of helicelline shells were also checked for mineralogical composition using X-ray  
114 diffraction (XRD). XRD was performed using a Bruker D2 Phaser diffractometer (30 kV, 10 mA)  
115 operating in Bragg-Brentano geometry (Θ-Θ scan mode) and equipped with a 1-dimensional  
116 Lynxeye detector. Ni-filtered Cu Kα radiation was used. Data were collected in the scan range 4-  
117 65° in 2Θ, with scan step of 0.02° and counting time of 0.1 s/step. Data were processed through the  
118 software Diffrac.Eva (Bruker AXS Inc., 2015). Similarly, carbonate concretions were checked for  
119 mineralogical composition using XRD and inspected using a SEM-EDS for microscopical  
120 observation (Philips SEM 515 coupled with an EDS EDAX-DX micro-analyser).

121 Stable oxygen (δ<sup>18</sup>O) and carbon (δ<sup>13</sup>C) isotope ratios were determined using a Gas Bench II  
122 (Thermo Scientific) coupled with an IRMS Delta XP (Finnigan Mat) at the IGG-CNR in Pisa  
123 (Italy). Each carbonate sample of ca. 0.15 mg was dissolved in H<sub>3</sub>PO<sub>4</sub> (100%), for 1 h at 70 °C in a  
124 sealed vial flushed with helium. The headspace gas (CO<sub>2</sub>) was entrained in a helium stream,  
125 automatically dried and purified and then injected into the continuous flow isotope ratio mass  
126 spectrometer via an active open split. Sample results were corrected using the international standard  
127 NBS-19 and a set of internal standards (two marbles, MOM and MS, and a carbonatite NEW12,  
128 previously calibrated using the international standards NBS-18 and NBS-19, e.g. Negri et al., 2015)  
129 and normalized to the V-PDB international standard and expressed in the well-known δ-notation.  
130 All samples were analysed in duplicate and analytical uncertainties for replicated analyses of δ<sup>18</sup>O  
131 and δ<sup>13</sup>C were ±0.15 ‰ or better. The δ<sup>18</sup>O values of water are reported as V-SMOW.

132

## 133 **4. Results**

### 134 *4.1 The terrestrial mollusc assemblage*

135 A very low number of terrestrial mollusc shells were recovered from the sampled sediments.

136 These are all gastropods, no bivalves were found. Sample SUL-16/01 was virtually devoid of fossil  
137 remains, whereas sample SUL-16/02 contained less than 20 shells/kg. Despite the relatively low  
138 number of individuals, the counts can be considered representative for ecological analyses due to  
139 the amount of material sieved. All the shell remains belonged to terrestrial molluscs (Table 1), and  
140 only scarce fragments of unidentifiable micromammal bones were recovered.  
141 Six species of terrestrial molluscs were recovered (Tab. 1; Fig. 3), two of which could not be  
142 identified to species level. The enid *Jaminia* sp. is represented by two shell apices only. As the  
143 extant species of this genus can be distinguished from the extinct Quaternary *Jaminia malatestae*  
144 (Esu, 1988) only based on the shell aperture it was not possible to identify these specimens to  
145 species level. The helicelline geomitrid, the most represented species in the paleosol, belongs to a  
146 group whose genus level taxonomy is entirely based on anatomical characters of the soft-parts.  
147 Therefore it was also not possible to make a more detailed taxonomic identification for this taxon.  
148 Shell features of the Sulmona geomitrid matched those of species in genera *Candidula*, *Cernuella*,  
149 *Helicella*, *Helicopsis*, *Xerocrassa* and *Xerosecta* (see the thorough iconographic survey on  
150 European species by Welter-Schultes, 2012). It could be hypothesized that the Fiorata Paleosol  
151 species corresponded to *Helicopsis striata* (Müller, 1774), a small European xerophilous geomitrid  
152 often associated with *Pupilla muscorum*.

153 From an ecological perspective, the identified species can be associated with dry to  
154 mesophilous open habitats (Tab. 1). *Pomatias elegans* is a medium-sized European prosobranch  
155 snail living among litter, humus and plant debris in many different dry to mesophilous habitats with  
156 some plant cover and preferably on calcareous substrates. *Truncatellina cylindrica* is a very minute  
157 European-Mediterranean pulmonate snail living among humus and leaf litter in dry to mesophilous  
158 habitats with some plant cover. The extant *Jaminia* species are small to medium-sized European  
159 calciphile pulmonates living on the soil surface or among rocks in dry, open grasslands and  
160 limestone reliefs. The Middle and Late Pleistocene *Jaminia malatestae* has been reported in open-  
161 dry paleoenvironments from the central-southern Italian peninsula during cold climatic periods (Di  
162 Vito et al., 1998; Marcolini et al., 2003; D'Amico and Esu, 2011; Limondine-Louzet et al., 2016).  
163 *Pupilla muscorum* is a very small Holarctic pulmonate snail usually living among humus, leaf litter  
164 and rock debris in dry, cool, open habitats, preferably on calcareous substrates. *Vallonia costata* is a  
165 very small Holarctic snail, living among humus, litter, moss and plant debris in mesophilous open  
166 or sparsely vegetated habitats. This is also valid for the helicelline geomitrid, which is typical of  
167 dry, open, sunny habitats (indeed its shell is a typical “chaliconcha” following the shell  
168 classification by Sacchi, 1952).

169 *4.2 Mineralogical and isotopic analyses*

170 The XRD analysis confirmed that the terrestrial shells preserved their primary aragonite  
171 mineralogy, as also suggested by their well-preserved aspect. Only one sample had a considerable  
172 amount of calcite (ca. ¼) and thus it was excluded from further isotopic analysis. However, we  
173 suspect that this was due to the presence of thin and superficial encrustations of pedogenic  
174 carbonates. Oxygen isotope composition of terrestrial shells ranged from  $\delta 4.26$  ‰ to  $\delta 2.55$  ‰,  
175 whereas their carbon isotope composition ranged from  $\delta 9.42$  ‰ to  $\delta 8.34$  ‰ (Table 2).

176 Pedogenic carbonates were mm-sized, often elongated fine-grained concretions, preserving  
177 small cylindrical holes, sometimes ramified. Some were consistent with “hypocoatings-type”  
178 carbonate concretions (e.g. Barta, 2011). The XRD analysis indicated that the small pedogenic  
179 concretions were mostly formed of calcite, along with minor quartz inclusions and traces of  
180 feldspars and micas, as further confirmed by SEM-EDS. Oxygen isotope composition of pedogenic  
181 carbonates ranged from  $\delta 7.60$  ‰ to  $\delta 6.54$  ‰, whereas their carbon isotope composition ranged  
182 from  $\delta 10.02$  ‰ to  $\delta 9.52$  ‰ (Table 2).

183

184 **5. Discussion**

185 *5.1 Chronology and paleoenvironmental significance of the mollusc assemblage*

186 The deep unconformity at the base of SUL-5, which cuts up to ~50 m of the underlying unit  
187 SUL-6 (Figs. 1 and 2), filled by fluvial gravels and sands, and capped by the Fiorata Paleosol,  
188 indicates a pronounced and long phase of lake low-stand probably associated with the desiccation of  
189 the lacustrine system. Based on the available chronological and stratigraphic constraints (Figs. 1  
190 and 2), this subaerial phase can be roughly dated between ~650 ka and ~530 ka, or between MIS 16  
191 and MIS 14. However, the Fiorata Paleosol documents only the final stages of this long subaerial  
192 phase, and the timing of the end of the soil-forming phase can be precisely constrained by the  
193 deposition of the thick tephra layer correlated to the Tufo di Bagno Albule, dated to  $527 \pm 2$  ka (Fig.  
194 2). Recently, a tephra layer dated to  $531 \pm 5$  ka, and tentatively correlated to the Tufo di Bagno  
195 Albule, has also been identified in the archaeological succession of Valle Giumentina (Villa et al.,  
196 2016), located ~15 km NE of Sulmona Basin. The Fiorata Paleosol can therefore be correlated with  
197 the period corresponding to MIS 14 (Fig. 2) and/or to the transition to MIS 13 (Railsback et al.,  
198 2015). However, the non-marine fauna is suggestive of an open-dry environment and more  
199 indicative of a glacial phase, better corresponding to a later phase of the glacial MIS 14.



200 The fauna is characterized by a low number of species and shares some general similarities with  
201 other terrestrial mollusc assemblages from the central Italian Peninsula considered typical of colder  
202 and drier conditions of glacial periods (e.g. Esu, 1981; Di Vito et al., 1998; Limondin-Lozouet et  
203 al., 2016; Boretto et al., 2017). Similarly, the terrestrial molluscs from Sulmona Basin indicate open  
204 and moderately dry habitats, as inferred from the presence of mesophilous and slightly  
205 thermophilous components (Table 1, i.e. *Pomatias elegans*).

206

## 207 5.2 Stable isotope geochemistry

208 The shells and pedogenic carbonates have very distinctive  $\delta^{18}\text{O}$  and  $\delta^{13}\text{C}$  values (Fig. 4),  
209 which also differ from the values obtained from lacustrine carbonates dated to MIS 11 and MIS 12  
210 in the same basin (Regattieri et al., 2016), as well from clastic marine carbonates from the  
211 substratum (Regattieri et al., 2016; Villa et al., 2016). Along with the mineralogical evidence for the  
212 preservation of the shell mineralogy, the isotopic distribution provides robust evidence that  
213 terrestrial carbonates were not isotopically altered by diagenesis. This also indicates that pedogenic  
214 carbonates were not significantly contaminated by clastic carbonate.

215 Aragonite is usually enriched in  $^{18}\text{O}$  and  $^{13}\text{C}$  compared to calcite (Tarutani et al., 1969;  
216 Grossman and Ku, 1986; Romanek et al., 1992; Kim and O'Neil, 1997), but the difference observed  
217 between pedogenic carbonates and shells is not simply related to different isotopic fractionation  
218 factors. Terrestrial shells have  $\delta^{13}\text{C}$  values higher than pedogenic carbonate by ca. 1 ‰. This value  
219 is slightly lower than those expected for calcite and aragonite precipitating close to isotopic  
220 equilibrium from the same solution (ca. 1.7 ‰, Romanek et al. 1992). We noticed, however, that  
221 analyses of aragonite and calcite in biogenic carbonates yielded differences in values closer to ca. 1  
222 ‰ (Lécuyer et al., 2012). The  $\Delta$  ( $\delta^{18}\text{O}_{\text{aragonite}} - \delta^{18}\text{O}_{\text{calcite}}$ ) values show some differences according to  
223 the equations used (e.g. usually lower than 1 ‰, Tarutani et al., 1969; Grossman and Ku, 1986;  
224 Patterson et al., 1993; Kim and O'Neil, 1997; Lécuyer et al., 2012), but these differences are always  
225 lower than those observed between the  $\delta^{18}\text{O}$  of aragonitic shells and pedogenetic carbonates in our  
226 record (ca. 4 ‰, Fig. 4). This difference can be explained by kinetic (vital) offset compared to  
227 equilibrium conditions for terrestrial mollusc shells, and to different environmental water sources  
228 from which the two carbonate polymorphs precipitate (i.e. the shell and the pedogenic carbonate).

229 The  $\delta^{18}\text{O}$  values of terrestrial gastropod shells is proven to be related to the  $\delta^{18}\text{O}$  values of  
230 environmental waters absorbed/ingested by the snails (e.g. water vapour, dew, local meteoric  
231 precipitation, e.g. Lécuyer, 1985; Goodfriend et al., 1989; Zanchetta et al., 2005; Prendergast et al.,

232 2015), and to isotopic effects linked to the exchange of fluid between the external environment  
233 (through the body of the snails) and internal fluid (Balakrishnan and Yapp, 2004), which are  
234 influenced by relative humidity (Balakrishnan and Yapp, 2004) and temperature. Therefore, no  
235 simple isotopic equilibrium with meteoric water could be assumed. However, empirical relations  
236 between the  $\delta^{18}\text{O}$  values of meteoric water and the shells have been found within living populations  
237 (e.g. Lécolle, 1985; Goodfriend and Ellis, 2002; Zanchetta et al., 2005; Yanes et al., 2008;  
238 Prendergast et al., 2015), although in very arid lands a direct correlation is often not particularly  
239 robust (Goodfriend et al., 1989). Considering the data available from different living populations,  
240 there is no conclusive evidence that oxygen isotopic composition of shells is species-dependent  
241 (e.g. Lécolle, 1985; Goodfriend and Ellis, 2002; Zanchetta et al., 2005; Baldini et al., 2007; Yanes  
242 et al., 2008, 2009; Colonese et al., 2013ab, 2014). Physiological and ecological factors, along with  
243 environmental conditions of different populations, can additionally influence the final oxygen  
244 isotope composition of the shells (Goodfriend et al., 1989; Balakrishnan et al., 2005b; Yanes et al.,  
245 2008, 2009; Colonese et al., 2013b, 2014). The most complete model to interpret oxygen isotope  
246 composition of terrestrial gastropod shells is that proposed by Balakrishnan and Yapp (2004). The  
247 model indicates that the steady-state  $\delta^{18}\text{O}$  value of shell carbonate depends upon temperature,  
248 relative humidity,  $\delta^{18}\text{O}$  of the input liquid water and  $\delta^{18}\text{O}$  of ambient water vapour. However,  
249 quantitative prediction using this model involves several assumptions, which complicate its  
250 applicability to past samples (Balakrishnan et al., 2005a; Colonese et al., 2013a).

251 For living populations in Italy, Zanchetta et al. (2005) found an empirical relation between isotopic  
252 composition of precipitation ( $\delta^{18}\text{O}_p$ ) and isotopic composition of shell ( $\delta^{18}\text{O}_s$ ):

$$254 \delta^{18}\text{O}_p = 0.65 \times \delta^{18}\text{O}_s + 5.44 \quad (r^2 = 0.79) \quad (1)$$

255  
256 If relation (1) is assumed valid also for the past and for Sulmona settings, the average values of the  
257 oxygen isotope composition of meteoric water during the period of shell calcification can be  
258 calculated to be  $+7.8 \pm 0.5$  ‰.

259 Oxygen isotope composition of pedogenic carbonate is mostly related to local rainfall (Cerling,  
260 1984), with additional evaporative effects in the soil and the effect of temperature-related isotopic  
261 fractionation during calcite precipitation. Using Cerling's (1984) data on modern soils, Jiamao et al.  
262 (1997) proposed the following relationship between  $\delta^{18}\text{O}$  values in water and soil carbonate, which  
263 incorporates the evaporative effect in soils (Zanchetta et al., 2000) and the effect of the temperature  
264 in the fractionation factor:

265

266  $\delta^{18}\text{O}_p = \text{‰}1.361 + 0.955 \times \delta^{18}\text{O}_{\text{CaCO}_3}$  ( $r^2 = 0.98$ ) (2)

267

268 Boretto et al. (2017) found that equation (2) is a good predictor of isotopic composition of current  
269  $\delta^{18}\text{O}_p$  along the Tuscan coast. Assuming that equation (2) can also be applied to older pedogenic  
270 carbonates, it provides an average  $\delta^{18}\text{O}$  precipitation value of  $\text{‰}8.3\pm0.4$  ‰. This  $\delta^{18}\text{O}$  value is  
271 similar, though lower, to those obtained from the Fiorata Paleosol shells, a fact that mutually  
272 supports the two estimations. We note that the equations (1) and (2) have been obtained using the  
273 annual average  $\delta^{18}\text{O}$  value of local rainfall, so in principle the calculated values should be  
274 interpreted accordingly. However, terrestrial molluscs form their shells predominantly during wetter  
275 and/or warmer conditions (Balakrishnan and Yapp, 2004), therefore, the isotopic signal would be  
276 skewed toward the growth period (Kehrwald et al., 2010). If the warmer part of the year (e.g.  
277 spring/summer) was the principal season for the shell growth, higher  $\delta^{18}\text{O}$  values of meteoric  
278 precipitation would be expected (Rozanski et al., 1993; Fricke and O'Neil, 1999) as, indeed,  
279 observed.

280 While pedogenic carbonates may precipitate from soil water which is more representative of  
281 the annual average recharge conditions (Cerling, 1984), Breecker et al. (2009) observed that  
282 pedogenic carbonates in very dry environments form during warmer, drier periods and from soil  
283 solution mostly recharged during wetter periods. If the soil water solution from which carbonate  
284 precipitates represents the colder months of recharge, this would explain the lower estimated  $\delta^{18}\text{O}$   
285 for precipitation compared to that obtained for shells. On the other hand, pedogenic carbonates can  
286 be the result of repeated events of carbonate deposition (and eventually re-dissolution), and thus  
287 their isotopic composition represents the weighted average of multiple events over a certain period  
288 of time. Similarly, shells dispersed within soil horizons are not necessarily coeval, but may  
289 represent a different period of burial. All these factors can complicate the proposed interpretation,  
290 however, the relatively narrow  $\delta^{18}\text{O}$  values measured also suggest that conditions did not change  
291 significantly during the soil formation.

292 The most striking feature is that the estimated  $\delta^{18}\text{O}$  values of precipitation are, on average,  
293 lower than the values measured today at L'Aquila station ( $\text{‰}7.13$  ‰), also considering a small  
294 altitudinal correction for the Sulmona basin (ca.  $\text{‰}0.15$  ‰). Lower  $\delta^{18}\text{O}$  of precipitation during  
295 glacial/stadial conditions is expected at latitudes where isotopic composition of meteoric  
296 precipitation is strongly related to temperature (Rozanski et al., 1993). In the central Mediterranean  
297 a dependence between precipitation  $\delta^{18}\text{O}$  and surface temperature ( $\delta^{18}\text{O}/T$ ) has been found to be ca.

298 +0.2 ‰/°C, and this relation can be assumed also for the last two glacial periods (Bard et al., 2002).  
299 Therefore, the differences between present day  $\delta^{18}\text{O}$  values of precipitation and the predicted  
300 average  $\delta^{18}\text{O}$  value from pedogenic carbonates would account for ca. 5°C lower temperature at the  
301 time of soil carbonate formation, perhaps representing winter recharge; whereas the Fiorata  
302 Paleosol shells indicate less than 3°C lower temperature, probably due to expression of the warmer  
303 season.

304 It has been suggested that in the central Mediterranean most of the  $\delta^{18}\text{O}$  signal in Quaternary  
305 continental carbonates is dominated by the “amount effect” of rainfall (ca.  $\approx 2$  ‰/100 mm; Bard et  
306 al., 2002). This assumption has been used specifically for lakes for which additional evaporative-  
307 enrichment effects have been suggested during drier periods (Zanchetta et al., 1999,2007a; Roberts  
308 et al., 2008; Giaccio et al., 2015; Regattieri et al., 2015,2016), and for speleothem carbonate (Bar-  
309 Matthews et al., 2000; Regattieri et al., 2014; Zanchetta et al., 2007a,2016), for which the  
310 evaporative effect could be considered minor. Oxygen isotope composition of authigenic, bio-  
311 mediated calcite from lacustrine intervals from the Sulmona Basin unequivocally indicate that  
312 carbonates tend to have higher  $\delta^{18}\text{O}$  values during colder and drier periods (Regattieri et al.,  
313 2015,2016,2017; Giaccio et al., 2015). Higher  $\delta^{18}\text{O}$  values of lacustrine calcite during colder and  
314 drier periods at Sulmona probably result from the combination of several factors. During a glacial  
315 period  $^{16}\text{O}$ -enriched water is stored in continental ice (the “ice volume effect”, e.g. Mix and  
316 Ruddiman, 1984) leading to  $^{18}\text{O}$ -enriched ocean waters; this enrichment is propagated into the  
317 hydrological cycle. The most obvious local effect is the lower temperature of carbonate  
318 precipitation, with related changes in the fractionation factor (Kim and O’Neil, 1997 and references  
319 therein), even though the occurrence of algal blooms responsible for calcite precipitations cannot  
320 occur for too lower temperatures. Moreover, drier conditions during glacial periods could enhance  
321 evaporation, causing enrichment in  $^{18}\text{O}$  in residual water (Gonfiantini, 1986).

322 While environmental conditions derived from the terrestrial mollusc assemblages and from  
323 the general stratigraphic features are consistent with a general reduction in the amount of  
324 precipitation, both shell and pedogenic carbonates predict considerably lower  $\delta^{18}\text{O}$  values in  
325 meteoric precipitation. This clearly challenges the assumption that the rainfall amount exerts a  
326 dominant effect on the isotopic composition of continental carbonates in the Mediterranean area, at  
327 least for the interval considered.

328 A possible explanation to reconcile these discordant interpretations is that terrestrial  
329 carbonates show distinct responses to precipitation regimes. Lakes and speleothems tend to have  
330 recharge systems that average and mix rainfall over the catchment area, and thus are more sensitive

331 to the total amount of precipitation. Terrestrial gastropod shells, instead, are more susceptible to  
332 local precipitation and humidity during the period of growth. Pedogenic carbonates would be more  
333 sensitive to local rainfall and specific periods of recharge of soil interstitial water. For instance, the  
334 pedogenic carbonates of the Fiorata Paleosol would have been influenced by a shift in large-scale  
335 atmospheric circulation. This may have taken the form of frequent incursions of cold air masses,  
336 depleted in  $^{18}\text{O}$ , deriving from northern latitudes of continental Europe, producing mostly snow  
337 precipitation (Enzi et al., 2014). Melting snow can have a different soil infiltration pattern  
338 compared to rainfall. This may have led to carbonates mostly recharged by  $^{18}\text{O}$ -depleted waters.  
339 Shells may have formed during warmer parts of the year but during wetter precipitation events  
340 characterised by particularly lower-than-average  $\delta^{18}\text{O}$  values (e.g. Colonese et al., 2007,2013).

341 It is important to emphasize that the empirical equations discussed herein may not be widely  
342 applicable to past climates, in particular for glacial periods for which different synoptical climate  
343 conditions would have existed (Kuhleemann et al., 2008; Kehrwald et al., 2010).

344 On the other hand, the MIS 14 is a particularly weak glacial in many records that one could its  
345 designation as glacial (Lang and Wolff, 2011), as shown by the global benthic stack of Lisieki and  
346 Raymo (2005) or ice core temperatures (Jouzel et al., 2007). The pollen record at Thenaghi  
347 Philippon in Greece did not show a very prominent decrease in arboreal vegetation for MIS 14,  
348 even though phases of increased grasses due to drier and colder conditions were recognized  
349 (Tzedakis et al., 2006). This is also evident in Lake Ohrid where MIS 14 seems one of the less  
350 expressed glacial periods of the record (Franke et al., 2016).

351 Carbon isotope composition of both shells and pedogenic carbonates is indirectly related to  
352 vegetation cover, but with different and complex relationships. Terrestrial molluscs form their shells  
353 mainly from respired  $\text{CO}_2$ , and shell  $\delta^{13}\text{C}$  values mostly reflect the stable carbon isotope  
354 composition of ingested vegetation (e.g. Goodfriend et al., 1989; Stott, 2002; Metref et al., 2003;  
355 Balakrishnan et al., 2005b; Liu et al., 2007). However, depending on the species and environmental  
356 settings (calcareous areas) shell  $\delta^{13}\text{C}$  values may also be affected by the ingestion of soil carbonates  
357 (Yates et al., 2002; Romaniello et al., 2008; Yanes et al., 2008; Colonese et al., 2014). Moreover,  
358 different feeding behavior and food preferences may variably affect shell  $\delta^{13}\text{C}$  values (Colonese et  
359 al., 2014). Using vegetation as the unique  $\text{CO}_2$  source for shell carbon isotopes, Stott (2002) found a  
360 strong positive linear relationship between plant and shell  $\delta^{13}\text{C}$  ( $\delta^{13}\text{C}_{\text{shell}}$ ):

361  
362 
$$\delta^{13}\text{C}_{\text{diet}} = 1.35 \times \delta^{13}\text{C}_{\text{shell}} \boxtimes 11.73 \quad (3)$$
  
363

364 The equation (3) has been obtained for *Cornu aspersum* and the applicability to other species might  
365 be questionable. However, applied to our shell  $\delta^{13}\text{C}$  values, it provides an average value for the  
366 ingested food of  $\approx 23.7 \pm 0.6$  ‰. The average value for  $\text{C}_3$  plants is ca.  $\approx 27$  ‰ (e.g. Deines, 1980),  
367 and the calculated values of ingested vegetation are consistent with carbon isotope values obtained  
368 from  $\text{C}_3$  vegetation from moderately dry conditions. Carbon isotope composition of  $\text{C}_3$  vegetation in  
369 drier environments can be significantly higher than plants from wet environments (e.g. Kohn,  
370 2010). Specifically in the Mediterranean, remarkable differences in  $\delta^{13}\text{C}$  of  $\text{C}_3$  plants are observed  
371 related to changes in water-use efficiency, which also varies largely between species, with higher  
372  $^{13}\text{C}/^{12}\text{C}$  ratio measured in drier areas (Hartman and Danin, 2010). Moreover, in the Mediterranean  
373 area similar shell  $\delta^{13}\text{C}$  values have been reported in ecosystems dominated by  $\text{C}_3$  plants  
374 (Goodfriend et al., 1989; Colonese et al., 2014; Prendergast et al., 2015).

375 For pedogenic carbonates there are theoretical equations which can be applied for  
376 calculating the  $\delta^{13}\text{C}$  values of vegetation from which it is precipitated and the relative amount of  $\text{C}_3$   
377 and  $\text{C}_4$  vegetation. According to Wang and Zheng (1989) the  $\delta^{13}\text{C}$  values of vegetation over a soil  
378 can be estimated from the carbon isotope composition of pedogenic carbonate, with the following  
379 equation:

$$380 \quad x = (11.9 + \delta^{13}\text{C}_{\text{pedogenic}}) / 14 \quad (4)$$

383 Applying the equation (4) to our carbonate  $\delta^{13}\text{C}$  values, a proportion of  $\text{C}_4$  plants ranging from 30  
384 to 38 % is obtained.  $\text{C}_4$  vegetation is relatively rare in southwestern Europe and mostly belonging to  
385 herb and shrubs (Pyankov et al., 2010), as such these estimations could seem particularly high.  
386 Indeed, these estimations are based on the assumption that  $\text{C}_3$  plants have a mean carbon isotopic  
387 value of ca.  $\approx 27$  ‰, which is only a first-order estimation, whereas higher values of prevailing  $\text{C}_3$   
388 vegetation can be obtained by water stress, as previously discussed.

389 Considering the  $\text{C}_4$  estimation from the isotopic composition of pedogenic carbonate, Breecker et  
390 al. (2009) observed that in dry environments pedogenic carbonates form predominantly during  
391 warm and dry conditions, and during periods of low soil respired- $\text{CO}_2$ , thus overestimating the  
392 presence of  $\text{C}_4$  vegetation. This supports the observation obtained from shell  $\delta^{13}\text{C}$  values that the  
393 relatively high  $\delta^{13}\text{C}$  values in pedogenic carbonates at Sulmona derive from  $\text{C}_3$  vegetation enriched  
394 in  $^{13}\text{C}$  during dry seasons. The integration of these two sources of paleoenvironmental information  
395 offers stronger arguments for interpreting past vegetation cover during soil formation, notably in  
396 Mediterranean areas, where the isotope ecology of modern and fossil shells is relatively well  
397 known.

398

## 6. Conclusion

Chronological, stratigraphic, and paleontological data indicate that the paleosol at the base of SUL5 in the Sulmona Basin sedimentary succession was formed during drier and probably colder conditions at the time of the MIS 14 glacial phase. The local mollusc assemblage indicates an open, dry environment. Carbon isotope compositions of pedogenic carbonates and shells consistently suggest prevailing C<sub>3</sub> vegetation adapted to dry environments. Inferred oxygen isotope composition of past rainfall from shells and pedogenic carbonates indicates that precipitation was generally <sup>18</sup>O-depleted over the region compared to present-day. While this could imply a decrease in the atmospheric temperature of ca. 3-5°C compared to present day, this also conflicts with the current interpretation of speleothems and lake δ<sup>18</sup>O values in the central Mediterranean. For example, lower δ<sup>18</sup>O values in carbonates (arising from lower precipitation δ<sup>18</sup>O values) should reflect increased rainfall, due to the amount effect. But increased rainfall is not supported in our record according to the paleontological data and/or the carbon isotope composition of carbonates. We propose that more frequent incursions of <sup>18</sup>O-depleted cold air masses deriving from northern latitudes of continental Europe, along with a general context of reduced precipitation, would have influenced the isotopic composition of pedogenic carbonates and terrestrial shells. This work emphasizes the importance of integrating isotopic approaches on terrestrial carbonates (molluscs and pedogenic carbonates) to derive more robust interpretative frameworks on past climate and environments in the Mediterranean region.

418

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791 Figure and table captions

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800 Figure 3. The most represented land snail species in Fiorata paleosol: *Pupilla muscorum* (top),  
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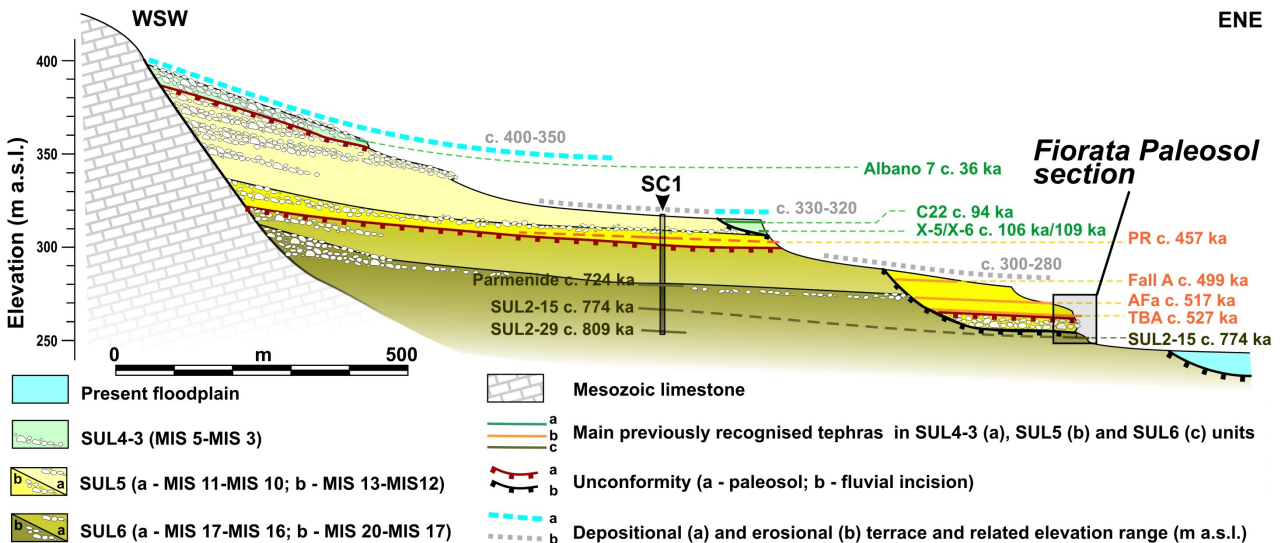
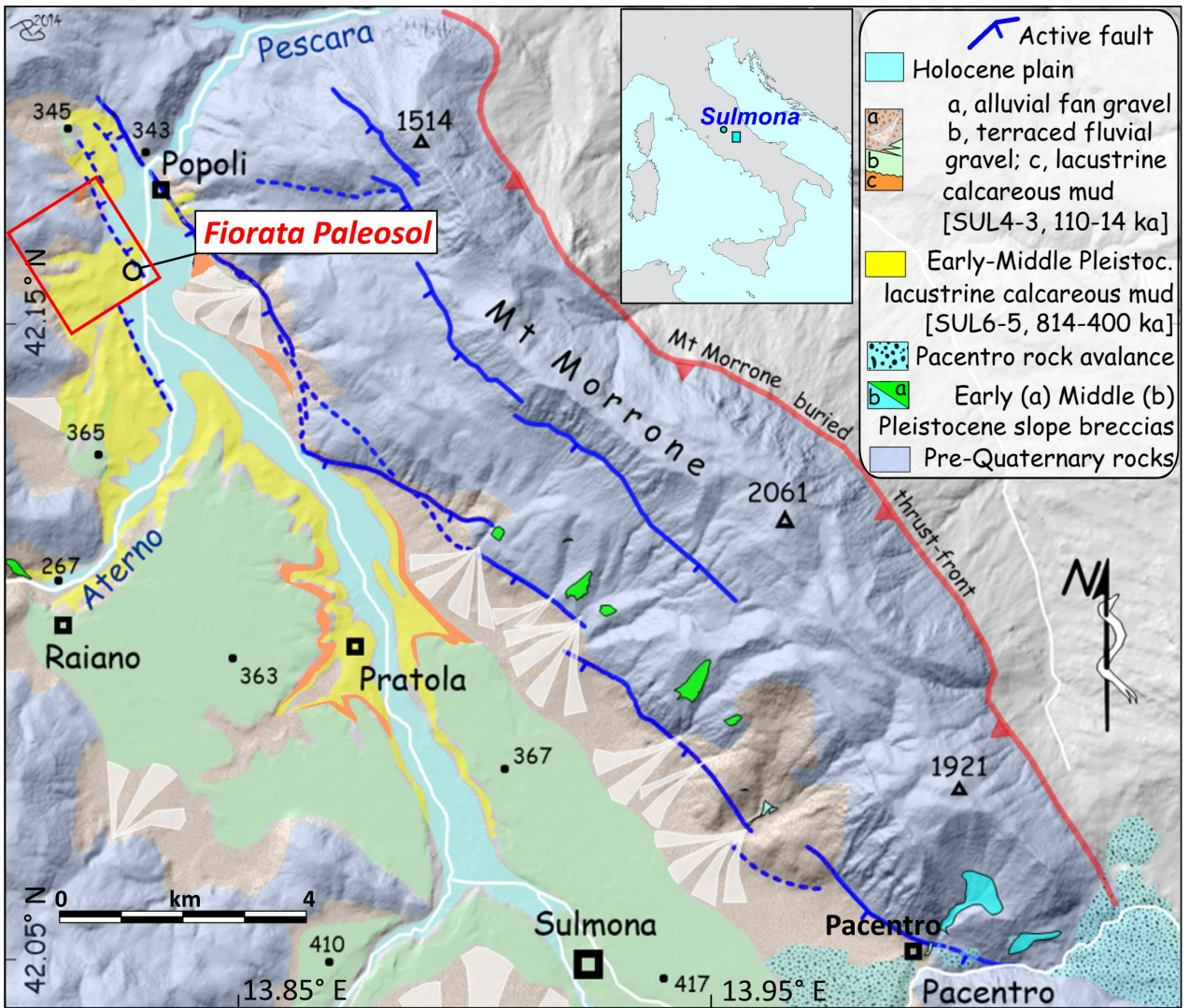
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803 Table 1. Landsnails species and their ecological requirement.

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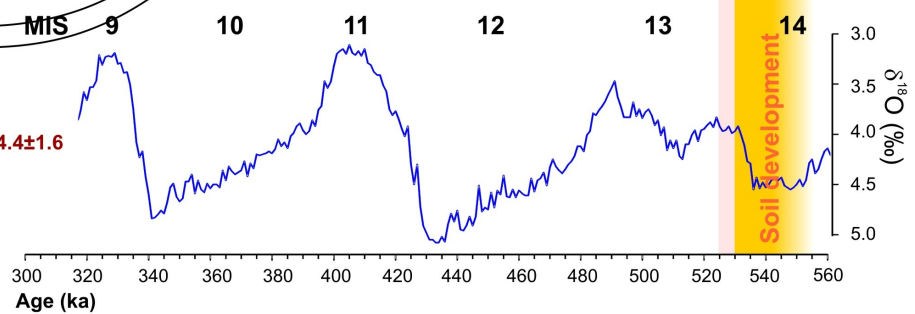
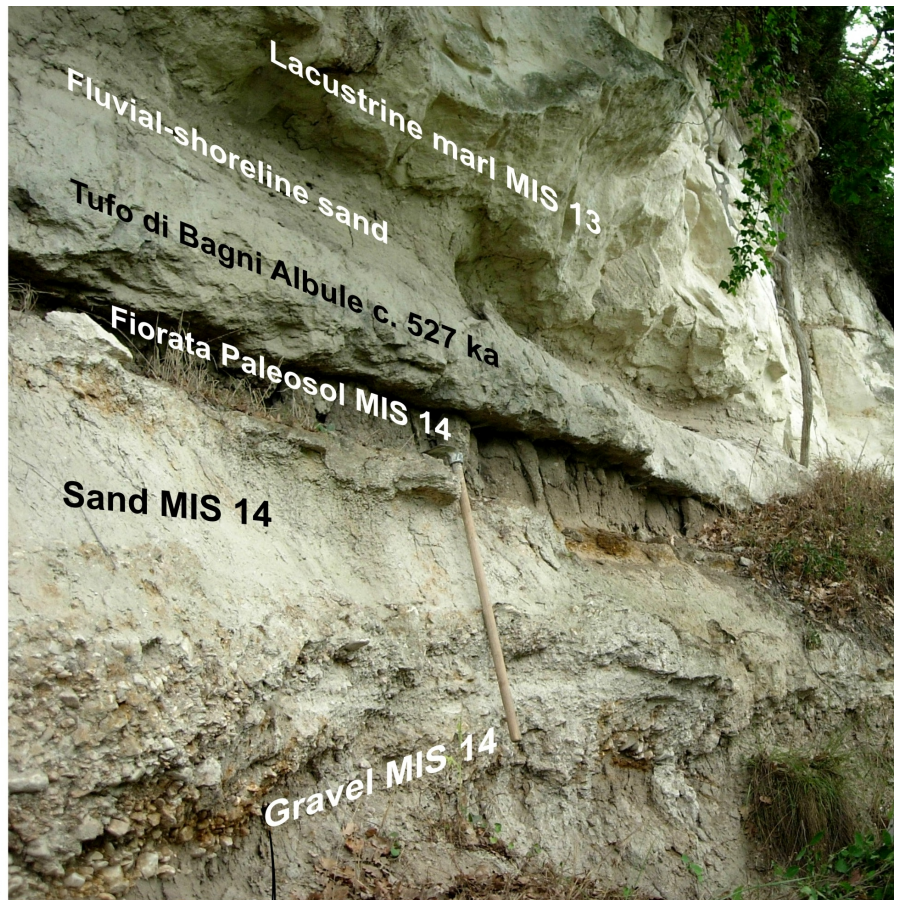
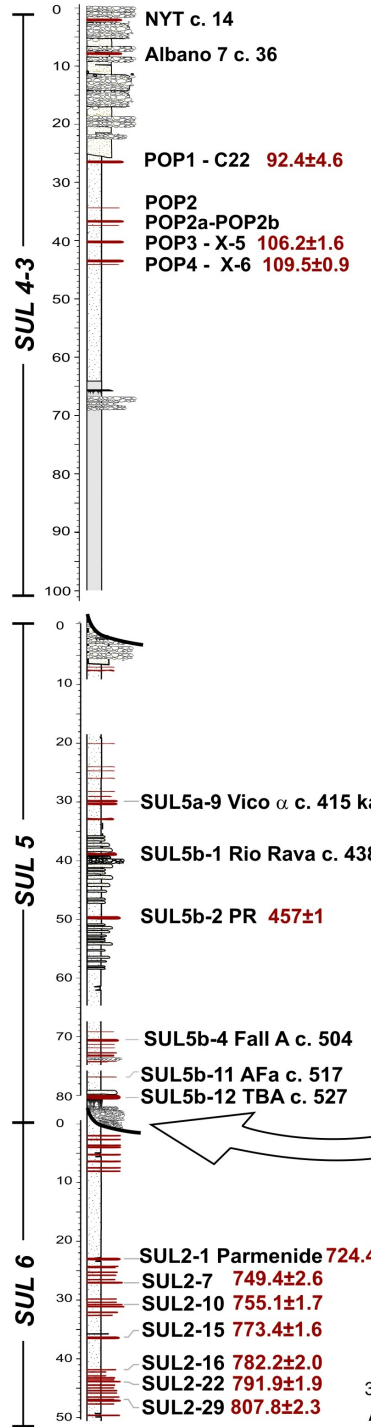
805 Table 2. Stable isotope composition of land snail shells and pedogenic carbonate from the Fiorata  
806 paleosol.

807



# Sulmona composite section

Depth (m) Tephra and related age (ka)



- |  |                                   |  |                         |  |              |
|--|-----------------------------------|--|-------------------------|--|--------------|
|  | Lacustrine whitish marls          |  | Fluvial/lacustrine sand |  | Paleosol     |
|  | Lacustrine-palustrine greysh clay |  | Fluvial/alluvial gravel |  | Tephra layer |
|  | Peat                              |  |                         |  |              |
- 106.2±1.6** <sup>40</sup>Ar/<sup>39</sup>Ar age (ka) of Sulmona tephra  
**c. 36** <sup>40</sup>Ar/<sup>39</sup>Ar age (ka) of correlated tephra



2 mm



2 mm



5 mm

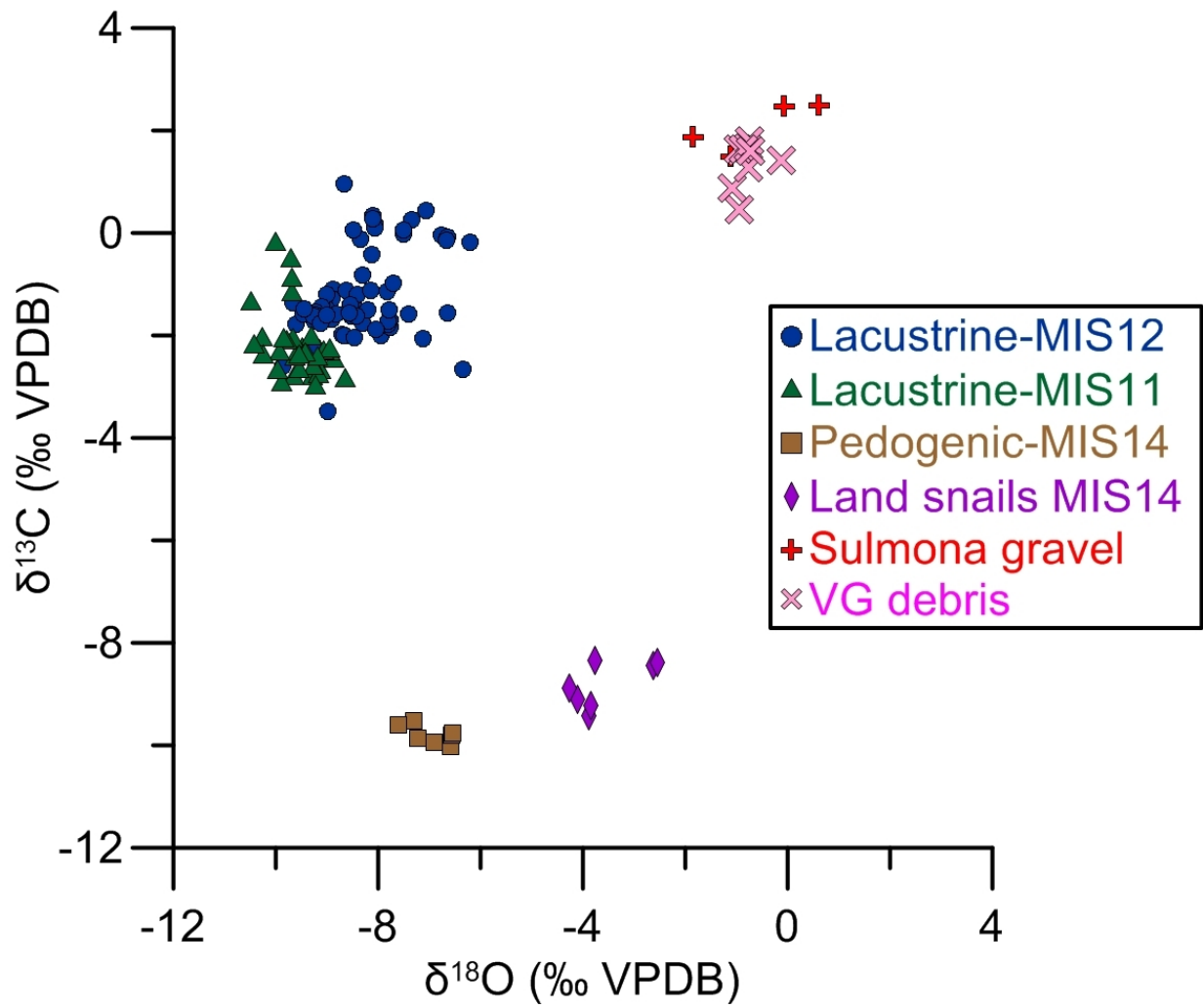


Fig. 4

## Figure and table captions

Figure 1. Location map, geological sketch map and general stratigraphy of the area. SC1 hole is discussed in Sagnotti et al., 2014 and Regattieri et al., 2015.

Figure 2. General stratigraphy of the lacustrine succession of the Sulmona Basin, with details of Fiorata paleosol succession and age constraints for the formation of the soil. Main tephra layers are indicated along with  $^{40}\text{Ar}/^{39}\text{Ar}$  dating.

Figure 3. The most represented land snail species in Fiorata paleosol: *Pupilla muscorum* (top), *Vallonia costata* (middle) and an unidentified geometrid helicelline (bottom).

Figure 4. Carbon and oxygen isotopic data from lacustrine deposits of Sulmona Basin (Regattieri et al., 2016), shell and pedogenic carbonate from Fiorata paleosol (this work); and clastic carbonate (Villa et al. 2016; Regattieri et al., 2016). VM: Valle Giumentina.

Table 1. Landsnails species and their ecological requirement.

Table 2. Stable isotope composition of land snail shells and pedogenic carbonate from the Fiorata paleosol.



Species and higher systematics	Material	Main habitats
<b>Caenogastropoda, Hypsogastropoda, Pomatiidae</b>		
<i>Pomatias elegans</i> (Müller, 1774)	2 sps: one operculum of a juvenile and one last whorl fragment of an adult/subadult	dry to mesophilous woodland edges and glades
<b>Heterobranchia, Pulmonata, Orthurethra, Vertiginidae</b>		
<i>Truncatellina cylindrica</i> (Férussac, 1807)	2 sps: two incomplete shells	mesophilous open habitats
<b>Heterobranchia, Pulmonata, Orthurethra, Pupillidae</b>		
<i>Pupilla muscorum</i> (Linnaeus, 1758)	> 16 sps: many well preserved shells	dry to mesophilous open habitats
<b>Heterobranchia, Pulmonata, Orthurethra, Valloniidae</b>		
<i>Vallonia costata</i> (Müller, 1774)	32 sps many bad preserved shells	mesophilous open habitats and woodlands
<b>Heterobranchia, Pulmonata, Orthurethra, Enidae</b>		
<i>Jaminia</i> sp.	2 sps: two shell apices	dry open habitats
<b>Heterobranchia, Pulmonata, Sigmurethra, Geomitridae, Helicellinae</b>		
Unidentified helicelline geomitrid	> 70 sps: many fragmentary shells	dry open habitats

Table 1. Landsnails species and their ecological requirement.

Sample label	$\delta^{13}\text{C}$ ‰ (V-PDB)	$\delta^{18}\text{O}$ ‰ (V-PDB)
Pedogenic carbonate		
SUL14-1	-10.02	-6.58
SUL14-2	-9.80	-6.57
SUL14-3	-9.95	-6.90
SUL14-4	-9.52	-7.22
SUL14-14a	-9.52	-7.31
SUL14-12	-9.77	-6.54
SUL14-13	-9.60	-7.60
<b><i>Average (<math>\pm 1</math> st dev)</i></b>	<b><i>-9.79 <math>\pm</math> 0.18</i></b>	<b><i>-6.96 <math>\pm</math> 0.42</i></b>
<i>Land snail shell</i>		
SUL14-6	-9.42	-3.88
SUL14-7	-9.11	-4.10
SUL14-8	-8.45	-2.63
SUL14-9	-8.34	-3.77
SUL14-10	-8.89	-4.26
SUL14-11	-9.23	-3.85
SUL14-11a	-8.39	-2.55
<b><i>Average (<math>\pm 1</math> st dev)</i></b>	<b><i>-8.83 <math>\pm</math> 0.44</i></b>	<b><i>-3.10 <math>\pm</math> 0.69</i></b>

Table 2 Stable isotope composition of land snail shells and pedogenic carbonate from the Fiorata paleosol

## Highlights

Mollusks association from Fiorata Paleosol indicates dry climate during late MIS14.

Carbon isotopes of pedogenic carbonates and land shells support the notion of dry climate.

Oxygen isotope composition of pedogenic carbonate and shells suggest lower 3-5 °C temperature than present.