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From leaves to seeds? The dietary shift in late Miocene colobine monkeys of southeastern Europe

Running title: Dietary shift in late Miocene colobines

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The authors have no conflict of interest to declare.

Data accessibility:

The R code is available in its entirety as online supplementary material, along with all data used for the preparation of this manuscript [<https://dx.doi.org/10.13140/RG.2.2.13861.42722>]. When it was possible, surface meshes of the specimens used in the topographic analysis have been uploaded on Morphosource [<https://www.morphosource.org>].

39 **From leaves to seeds? The dietary shift in late Miocene colobine monkeys of southeastern**
40 **Europe**

41 **Abstract**

42 Extant colobine monkeys are specialized leaf eaters. But during the late Miocene, western Eurasia
43 was home to colobines which were less efficient at chewing leaves than they were at breaking
44 seed shells. To understand the link between folivory and granivory in this lineage, the dietary
45 niche of *Mesopithecus delsoni* and *M. pentelicus* was investigated in southeastern Europe, where a
46 major environmental change occurred during the late Miocene. We combined dental topographic
47 estimates of chewing efficiency with dental microwear texture analysis of enamel wear facets. *M.*
48 *delsoni* was more efficient at chewing leaves than *M. pentelicus*, the dental topography of which
49 matches an opportunistic seed eater. Concurrently, microwear complexity increases in *M.*
50 *pentelicus*, especially in the northernmost localities corresponding to present-day Bulgaria. This is
51 interpreted as a dietary shift towards hard foods such as seeds or tubers, which is consistent with
52 the savanna and open mixed forest biomes that covered Bulgaria during the Tortonian. The fact
53 that *M. delsoni* was better adapted to folivory and consumed a lower amount of hard foods than *M.*
54 *pentelicus* suggests that colobines either adapted to folivory before their dispersal to Europe, or
55 evolved adaptations to leaf consumption in multiple occurrences.

56 **Keywords:** Balkans; Colobinae; Dental microwear texture analysis; Dental topography;
57 *Mesopithecus*; Paleoecology.

58 **Background**

59 *Climate change, plant communities and the adaptive answer of plant eaters*

60 Climate change can have drastic and long-term effects on the structure and the diversity of
61 terrestrial plant communities. Plenty of examples can be found in the fossil record, such as the
62 collapse of the ‘Coal Forest’ in Europe and North America following the greenhouse-gas forced
63 aridification at the end of the Carboniferous (Montañez et al. 2007), or the replacement of C₃
64 carbon fixation plants by C₄ carbon fixation plants in the intertropical zone at the end of the
65 Miocene, triggered among other factors by global aridification in a cooling context (Osborne and
66 Beerling 2006; Osborne 2008). Such major changes in plant communities can dramatically affect
67 the ability of contemporaneous fauna to feed on their preferred food resources, hence impacting
68 both taxonomic diversity and ecological disparity of animals. For instance, the collapse of the
69 Carboniferous rainforest resulted in a large faunal turnover in tetrapods as well as the emergence
70 of new dietary strategies, such as carnivory or tree fern browsing (Sahney et al. 2010). Similarly,
71 the expansion of C₄ plants during the late Miocene resulted in the diversification of grazing
72 mammals in North America (Janis et al. 2000; Osborne 2008), Asia (Behrensmeyer et al. 2007)
73 and Africa (Cerling et al. 2005).

74 Still, the consequences of the global climatic events can have different outcomes, depending on
75 the regional context. For instance, the global aridification which took place at the end of the
76 Miocene did not result in the expansion of C₄ plants in Western Eurasia, even though they could
77 have been locally abundant (Böhme et al. 2017); instead, it induced the shrinking of the Paratethys

78 and then, the disappearance of the Central European Wet Zone (van Dam 2006; Utescher et al.
79 2011; Pound et al. 2012). At the same time, a quasi-continuous biome of woodland/savanna and
80 open mixed forest arose from western China to eastern Europe (Böhme et al. 2017; Kaya et al.
81 2018; but see Denk et al. 2018), associated with the Pikermian and Baodean chronofaunas adapted
82 to these new environments (Kostopoulos 2009; Ataabadi et al. 2013). Concurrently, one likely
83 consequence of this aridification trend was the rise and fall of successive primate families in
84 Europe (Fig. 1). The Miocene epoch is indeed characterized by a radiation of hominoid primates
85 in Eurasia, with more than 10 genera ranging from Spain to south-eastern Asia (Moyà-Solà et al.
86 2009; Casanovas-Vilar et al. 2011; Begun 2015; DeMiguel et al. 2021). At the end of the Middle-
87 Miocene Climatic Transition (MMCT), the taxonomic diversity of European primates reached a
88 peak for hominoids and the now extinct pliopithecoids (Fig. 1). This high specific diversity
89 mirrors a wide ecological disparity, with species foraging on both soft and hard foods (Ungar and
90 Kay 1995; King et al. 1999; Merceron et al. 2005; DeMiguel et al. 2013, 2014), occupying
91 different types of habitats (Eronen and Rook 2004; Merceron et al. 2007, 2012, 2013, 2016;
92 Sukselainen et al. 2015; DeMiguel et al. 2021). Afterwards, the fossil record progressively
93 decreased for both pliopithecoids (Harrison 2005; Begun 2016) and hominoids (Casanovas-Vilar
94 et al. 2011), and the cercopithecids started occurring in Eurasia during the early Turolian (de
95 Bonis et al. 1990; Koufos 2009), where they persisted until Pleistocene times (Elton and O'Regan
96 2014).

97 The fossil record can translate how biodiversity is impacted by a global and/or regional
98 environmental changes, but it can also help exploring the adaptive answer of species to the local
99 effects of these environmental changes (Davis and Pineda-Munoz 2016). When faced with climate

100 change and its influence on primary production, species can avoid extinction either (i) by tracking
101 their preferred environment or (ii) by adapting to their new environmental conditions. Comparing
102 the dietary niches of extinct species at different localities, but within a constrained spatial and
103 temporal framework, could give important insights into the mechanisms underlying adaptation and
104 how they ultimately result in macroevolution.

105 *When colobine monkeys roamed the Balkans*

106 Colobine monkeys are a subfamily of catarrhines feeding principally on leaves. But back in the
107 late Miocene, the oldest colobine, *Microcolobus tugenensis* from Kenya, had low-cusped molars
108 compared to extant colobines (Benefit and Pickford 1986) and was thus unlikely to have been as
109 folivorous as extant species. This raises the issue of the evolutionary origins of folivory in
110 colobines. Following the terminology of Gould and Vrba (1982), it has been suggested that the
111 aptitudes of extant colobines for leaf consumption, such as their multi-chambered stomach or their
112 dental morphology, could in fact be exaptations co-opted from adaptations for seed consumption
113 (Chivers 1994; Lucas and Teaford 1994). For instance, the multi-chambered stomachs of
114 colobines are interpreted by Chivers (1994) as exaptations to folivory, which would have been co-
115 opted from an adaptation to digest high concentration of toxic carbon chains in seeds. . Similarly,
116 dental tools used during leaf chewing (i.e. lophs) could have been co-opted from an adaptation to
117 process tough seed shells (Happel 1988; Lucas and Teaford 1994). Still, it is not known when such
118 dietary shift to folivory would have occurred, and what could have caused it.

119 Today, colobines are restricted to inter-tropical biomes of Asia and Africa, with the notable
120 exceptions of three species of snub-nosed monkeys living in the temperate forests of China

121 (Nüchel et al. 2018) and at least two species of gray langurs living in the Himalayas (Groves
122 2001). But from the late Miocene to the Pliocene, they also thrived in western Eurasia. The earliest
123 reliable occurrence of a Eurasian colobine is *Mesopithecus delsoni* found in the Greek locality of
124 Ravin des Zouaves-5 (de Bonis et al. 1990; Koufos 2009); magnetostratigraphic estimations date
125 the site to ~ 8.2 Ma, while relative datation, based on the faunal assemblage, have assigned the site
126 to the Mammalian Neogene zone (MN; see Guérin 1980, 1982; Van Dam et al. 2001) MN11 (~
127 8.7 to 7.75 Ma; de Bonis et al. 1990; Koufos 2009). The status of *M. delsoni* is debated and
128 regarded either as a bona fide species (Koufos et al. 2004; Koufos 2009) or as a subspecies of the
129 later species *M. pentelicus* (Alba et al. 2015), which spans the second half of the MN12 (the
130 Pikermian event ~ 7.43 to 7.27 Ma; De Bonis and Koufos 1999; Böhme et al. 2018; Spassov et al.
131 2018) and possibly later at the beginning of the Messinian (Spassov et al. 2012). *Mesopithecus*
132 *delsoni* differs from *M. pentelicus* in its less inclined mandible planum alveolare, larger teeth, a
133 large honing facet on the lower third premolar, and a large hypoconulid on the lower third molar
134 with a deep groove on the distal face (de Bonis et al. 1990; Koufos 2009; Spassov et al. 2018).
135 Given these differences, we consider *M. delsoni* as a separate species.

136 From the analysis of its dental microwear texture (Merceron et al. 2009b), *M. pentelicus* has been
137 described as an opportunistic forager feeding occasionally on stress-limited food such as hard
138 seeds, having a high microwear complexity (Asfc) and a low anisotropy (EpLsar), as expected
139 from crushing wear facets with a high count of pits (Merceron et al. 2009a). A similar pattern was
140 observed in *M. delsoni* fossils from Ravin des Zouaves-5 and Hadjidimovo (Merceron et al.
141 2009b). The upper second molars of *M. pentelicus* have thin enamel combined with a low relief,
142 which indicates a moderate ability to crush hard foods, such as seeds, tubers or the exocarp of

143 some fruit, as well as a low ability to shear tough and pliant foods, such as leaves (Thiery et al.
144 2017). Compared to their extant relatives, these monkeys may have ingested a larger proportion of
145 sturdy, seed-like foods and a smaller proportion of tough, leaf-like foods.

146 The present study aims to explore how the dietary niche of the genus *Mesopithecus* changed
147 during the late Miocene in the Balkans (Fig. 1). This time and geographic location are especially
148 interesting because of the environmental change that occurs there at the end of the late Miocene.
149 Indeed, by the end of the MN11, the dominating evergreen broadleaf forest which occurred in
150 southeastern Europe had lost ground to forest savanna and open woodlands, particularly east of
151 longitude 28°E (Eastern border of Romania/Bulgaria; Pound et al. 2012). This aridification and
152 landscape opening then continued during the Pikermian event (MN12; Böhme et al. 2017).

153 ***Multi-scale analysis of the dietary niche***

154 Dietary estimates can only track the diet at a given temporal scale (Davis and Pineda-Munoz
155 2016). For instance, the dietary niche of a species can be inferred from cranial and dental
156 morphology, which reflect adaptations to ingest and process a range of foods at a large organismal
157 extent i.e., metapopulations. Since cranio-dental evolution may take thousands of years,
158 morphological estimates operate at a large temporal scale. At the same time, the food eaten by
159 individuals can leave a trace in the isotopic composition of the enamel (Ben-David and Flaherty
160 2012) or at the surface of the enamel wear facets, in the form of microscopic wear marks (Teaford
161 and Oyen 1989; Merceron et al. 2016; Ramdarshan et al. 2016; Teaford et al. 2017); but in
162 contrast with cranio-dental morphology, stable isotopes and dental microwear work to small
163 organismal extents i.e., the individuals, and operate at a small time scale (from weeks in dental

164 microwear to years in stable isotopes of the enamel; Davis and Pineda-Munoz 2016 and references
165 therein).

166 This work combines dental estimates from (i) dental topography and (ii), dental microwear texture
167 analysis. Dental topography measures the three-dimensional geometry of the tooth surface (Ungar
168 and Williamson 2000; Guy et al. 2013). Like cranio-dental morphology, the topography of the
169 teeth (before any significant modification by dental wear) is the result of selection over multiple
170 generations (Davis and Pineda-Munoz 2016). Dental topographic variables measure several
171 aspects of the tooth surface geometry. First, dental relief i.e., height variation over the tooth
172 surface can be estimated using slope (Ungar and Williamson 2000) and relief indices such as RFI
173 (Ungar and Williamson 2000; Boyer 2008; Thiery et al. 2019). The sharpness of the dental tools
174 can be estimated using Dirichlet normal energy (DNE: Bunn et al. 2011) and other curvature-
175 based indices, such as the area-relative curvature (ARC) introduced therein (see also Guy et al.
176 2017). Lastly, dental complexity is estimated using variables based on the heterogeneity of
177 exposure, such as orientation patch count rotated (OPCR; Wilson et al. 2012; Guy et al. 2013).
178 Leaf-eating cercopithecids tend to have higher relief and sharper teeth compared to more
179 frugivorous species (Bunn and Ungar 2009). Based on Thiery et al. (2017), the topography of *M.*
180 *pentelicus* and *M. delsoni* is expected to match with cercopithecids consuming a large proportion
181 of hard food i.e., we expect their molars to have a low-to-medium relief and a low sharpness.

182 In contrast, dental microwear texture analysis (DMTA) investigates texture changes of enamel
183 wear facets caused by daily contacts with food and other ingested particles (Scott et al. 2006,
184 2012). Experimental data and modern datasets confirm that hard food eaters have higher texture
185 complexity (Asfc) values (Scott et al. 2012; Ramdarshan et al. 2016). The heterogeneity of texture

186 complexity quantifies the variation of texture complexity observed within a scan (Asfc) and is
187 expected to be higher for animals with a versatile diet compared for instance to species with a
188 specialized diet (Scott et al. 2012; Ramdarshan et al. 2016; Merceron et al. 2018). The anisotropy
189 of the microwear texture (epLsar) measures the orientation concentration of surface roughness,
190 which is expected to be more directional (higher epLsar) for leaf eating species, notably when
191 their diet includes tough foliages and silica-bearing plants (Scott et al. 2012; Merceron et al.
192 2021). Finally, the textural fill volume (Tfv) is the result of an algorithm that fills a surface with
193 square cuboids of different volumes. Tfv does not depend on the surface shape but on its finer
194 texture. We expect a higher Tfv to be associated with higher Asfc values (Scott et al. 2012). Based
195 on earlier works on 2D microwear analysis (Merceron et al. 2009a) and DMTA (Merceron et al.
196 2009b), the texture complexity of *M. pentelicus* and *M. delsoni* is expected to reflect the
197 opportunistic inclusion of hard foods, with regional and possibly temporal variations in response
198 to local differences in environmental conditions (Kostopoulos 2009).

199 Results

200 *Dental topography*

201 Average slope is significantly higher in colobine monkeys than in the other catarrhines examined
202 here (Kruskall-Wallis' chi squared test: $K = 13.601$, $p < 0.001$). However, both *Trachypithecus*
203 *auratus* and *T. cristatus* have a noticeably lower slope than other colobines. In addition,
204 *Erythrocebus patas* has a higher slope than other non-colobine monkeys (Fig. 2A). Similarly,
205 ARC is significantly higher in colobines (Kruskall-Wallis' chi squared test: $K = 18.652$, $p <$
206 0.001). Again, *T. auratus* shows noticeably lower ARC values than other colobines, while *E. patas*
207 shows higher ARC values than other non-colobines (Fig. 2B).

208 Both *M. aff. delsoni* from Hadjidimovo and *M. pentelicus* from Pikermi have low slope values
209 grouping them with the most granivorous catarrhines, although *M. pentelicus* has an even lower
210 slope than the granivorous *Lophocebus albigena*. Similarly, the ARC of *M. pentelicus* is as low as
211 the ARC of *Chlorocebus aethiops* and *L. albigena*. In *M. aff. delsoni* however, ARC is visibly
212 higher, being similar to that of *T. auratus*. In contrast, *M. cf. pentelicus* (noted *M. sp.* on Fig. 2)
213 from Azmaka shows high slope values for a colobine, comparable to *Ch. Aethiops* or *T. auratus*,
214 and intermediate ARC values between *M. aff. delsoni* and *M. pentelicus*. Regarding other
215 estimates, the higher relief in *M. cf. pentelicus* is confirmed by relief index (RFI, Fig. 3B) sensu
216 Ungar and Williamson (2000) but not by relief index sensu Boyer (2008; LRFI, Fig. 3C). Both
217 variables do not separate *M. delsoni* from *M. pentelicus*. Concerning dental sharpness, DNE values
218 confirm that *M. delsoni* had sharper molars than *M. pentelicus* (Fig 3E). Finally, *M. delsoni* had

219 lower OPCR values indicating that its molars were less complex than the molars of *M. pentelicus*
220 (Fig. 3F).

221 ***Dental microwear texture analysis***

222 Complexity Asfc on phase II (crushing/grinding molar facets) is significantly higher in the
223 omnivorous *Chlorocebus aethiops* and *Erythrocebus patas*, and in the seed-eating mangabeys *C.*
224 *atys* (*Cercocebus* sp. on Fig. 2) and *L. albigena* than in the African colobines and in *T. auratus*
225 (post-hoc LSD-test, p-value < 0.05). The mangabeys also have significantly higher Asfc values
226 than *S. entellus*. In contrast, *Nasalis larvatus* shows a broad range of Asfc values overlapping with
227 mangabeys, and *T. cristatus* has Asfc values exceeding mangabeys' (Fig. 2A, B).

228 The Asfc of *Mesopithecus* ranges over most of the spectrum observed in extant species (Fig. 2A-
229 B). However, *Mesopithecus delsoni* from Ravin des Zouaves-5 and *M. pentelicus* from Pikermi
230 have low Asfc values (Fig. 2C). The highest Asfc values are observed in the northernmost
231 population of *M. pentelicus*, from Gorna Sushitsa (N = 4). They are significantly higher than what
232 is measured for Ravin des Zouaves-5 and Pikermi (post-hoc LSD-test, p-value < 0.05) and range
233 as high as the values observed for *T. cristatus* (Fig. 2A-B). In addition, Asfc values are
234 significantly higher in *M. pentelicus* from the Bulgarian locality of Kalimantsi than in Pikermi
235 (post-hoc LSD-test, p-value < 0.05). Lastly, specimens from Vathylakkos-2 (N = 1) and Perivolaki
236 (N = 4) attributed to *M. delsoni/pentelicus* (noted M. sp on Fig. 2) only differ significantly from
237 Gorna Sushitsa (post-hoc LSD-test, p-value < 0.05).

238 The heterogeneity of Asfc (Hasfc) shows a contrasting picture in extant species and does not
239 reflect dietary categories and phylogeny (Table 1). In extinct species however, it is worth noting

240 that the dispersion of data is much larger in Pikermi compared to the other sites (Fig. 4B).
241 Regarding epLsar, African colobines and *T. auratus* all have a significantly higher anisotropy than
242 the frugivorous and seed-eating cercopithecids, together with *T. cristatus* (Table 1). The
243 anisotropy is also significantly lower in the MN12 locality of Kalimantsi (*M. pentelicus*) than in
244 the MN11 locality of Hadjidimovo (*M. delsoni*), denoting a lower reliance on leaves (Fig. 4C).
245 Still, other MN12 localities do not mirror this difference. Again, the epLsar variance is the largest
246 at the MN12 locality of Pikermi, indicating a wide range of behaviors. Finally, the textural fill
247 volume (Tfv) does not separate extinct species or their localities (Fig. 4D; see also Table S8).

248 **Discussion**

249 *Fundamental niche predicted from dental topography*

250 Caution is needed when interpreting dental topography, as morphology is strongly constrained by
251 phylogeny, and what seems like an adaptation may be the remnants of an ancestral morphology
252 (Davis and Pineda-Munoz 2016). Still, a large number of studies have shown that folivorous
253 primates are characterized by high-relief molars which are on average sharper (Boyer 2008; Bunn
254 and Ungar 2009; Bunn et al. 2011; Winchester et al. 2014; see also Kay and Hiiemae 1974; Kay
255 1975, 1977). This is most likely related to the selective pressure exerted by flat, tough and pliant
256 foods such as leaves, which are displacement-limited foods and usually take more chewing
257 motions to be comminuted (Lucas 2000). The higher the molar relief, the sharper the tooth, and
258 the more displacement-limited food can be comminuted in a single chewing motion.

259 Apart from *T. auratus*, all colobine monkeys have high slope and ARC values, which indicates a
260 high leaf-chewing efficiency (Fig. 2A, 1B). The three species of African colobines forages
261 primarily on leaves even though *Co. polykomos* has been reported to fall back on seeds (protected
262 by pods) in several locations (Dasilva 1994; Davies et al. 1999). It is worth noting that colobine
263 monkeys usually remove the pods' shell using their anterior teeth i.e., incisors and canines and do
264 not use their molars for that purpose (Oates 1994; McGraw et al. 2016). Moreover, Fashing (2001)
265 reports on two populations of *Co. guereza* in western Kenya spending 40% to 60% of their
266 foraging time on fruits for 6 month a year. This degree of frugivory represents exceptions driven
267 by local environmental conditions and is most likely undetectable by dental topography due to the
268 scale of evolutionary biological changes through time. That being said, the consumption of seed-
269 like foods would have probably affected the dental topography of worn teeth even on a short-term,
270 local scale, as hinted by several studies (Ungar and Williamson 2000; Yamashita et al. 2012; Li et
271 al. 2020).

272 The four species of Asian colobines are also known to favor leaves. However, *T. auratus* has been
273 reported to target seeds as well (Kool 1993). This is consistent with their average slope, which is
274 the lowest among colobines. In addition, some populations of *N. larvatus* devote 15-20% of their
275 feeding time to seeds (Bennett and Sebastian 1988; Yeager 1989), and the most terrestrial colobine
276 monkey *S. entellus* also relies on fruits, seeds and even underground storage organs (USOs),
277 depending on resource availability (Hladik 1977; Sayers and Norconk 2008). This is consistent
278 with their lower ARC scores compared to other colobines. Both *Ch. aethiops* and *E. patas* have
279 intermediate slope scores, but the former has very low ARC scores, denoting a low leaf-chewing
280 ability, whereas *E. patas* has ARC scores comparable to *N. larvatus* or *T. auratus*. Again, this is

281 consistent with their diet, as vervets feed frequently on fruits and seeds (Wrangham and Waterman
282 1981; Whitten 1983), whereas patas feed more on tough but soft foods such as arthropods (Isbell
283 1998; Isbell et al. 1998). In contrast, the mangabeys have low slope and ARC values, pointing to a
284 poor leaf-chewing efficiency. Their molars better suit the grinding and crushing of sturdy hard
285 food such as seeds (Lambert et al. 2004; McGraw et al. 2014).

286 Given its low-relief and relatively dull molars, the theoretical dietary niche of *Mesopithecus*
287 *pentelicus* can be inferred as more versatile than extant colobines (Fig. 2A-B). It was probably
288 able to masticate or crack sturdy, hard foods such as seeds, seed shells or tubers without damaging
289 its molars, as suggested in previous studies (Merceron et al. 2009a,b; Thiery et al. 2017). The low
290 leaf-chewing ability of *M. pentelicus*, similar to that of *Ch. aethiops*, also indicates that it would
291 have experienced difficulties to thrive from leaves alone (because of longer chewing times, lower
292 digestibility, etc.). In addition, the thin enamel of *M. pentelicus*, which differs from the thick
293 enamel of both mangabey species, would have mitigated its ability to crack the hardest foods,
294 while improving the natural sharpening of enamel crests as tooth wears down (Thiery et al. 2017).

295 Having sharper molars, *M. delsoni* appears to have been slightly more efficient at chewing
296 leathery, leaf-like foods. However, the topography of only one specimen of *M. aff. delsoni* was
297 measured, so this interpretation remains highly hypothetical. Lastly, *M. cf. pentelicus* from
298 Azmaka (AZM-6) have intermediate ARC values between *M. delsoni* and *M. pentelicus*, but a
299 higher slope which reminds the molar topography of *Cercocebus*. This suggests a decent aptitude
300 to crush hard foods such as seeds, but also some underground storage organs e.g., tubers (Dominy
301 et al. 2008).

302 ***Realized niche inferred from dental microwear texture***

303 High Asfc on phase II molar facets is correlated with the consumption of USOs and any other food
304 which may carry grit and dust on their surface. However, and despite the fact that quartz grains
305 present on food do contribute to tooth wear, experiments which emulated natural conditions
306 concluded that dust loads do not obliterate the dietary signal (Merceron et al. 2016; Schulz-Kornas
307 et al. 2019). The complexity (Asfc) of crushing/grinding molar facets is also higher for primates
308 consuming hard, sturdy food items, such as fruits with hard exocarp or hard seeds (Scott et al.
309 2012; this study). This relationship has been shown experimentally on captive primates (Teaford
310 and Oyen 1989; Teaford et al. 2017) and on captive ewes (Ramdarshan et al. 2016). The two
311 mangabeys *C. atys* and *L. albigena* have high Asfc values (Fig. 2), while all colobines except *N.*
312 *larvatus* and *T. cristatus* have low to moderate Asfc values. Because they occupy one of the
313 wettest dense forest around the world, dusty sandstorms are unlikely to have affected dental
314 microwear of *N. larvatus* and *T. cristatus*, and their higher Asfc is more likely to result from the
315 seasonal inclusion of small seeds, as observed in *N. larvatus* (Yeager 1989; Matsuda et al. 2013).
316 During MN11, *M. delsoni* and *M. aff. delsoni* present Asfc scores within the range of *S. entellus*
317 and *N. larvatus*. Like these species, the dietary niche of *M. delsoni* was probably restricted to a
318 relatively narrow range of foods, likely fruit and leaves – albeit wider than other colobine
319 monkeys such as African colobines, the dietary niche of which is even narrower. In contrast, there
320 is a significant increase in the range of complexity (Asfc) between *M. delsoni* during MN11 and
321 *M. pentelicus* during MN12. While the complexity (Asfc) of *M. pentelicus* at the southern Greece
322 locality of Pikermi and to some extent in Vathylakkos-2 and Perivolaki) remains comparable to
323 the complexity of *M. delsoni*, it is similar to *C. atys* in the Bulgarian locality of Kalimantsi (N =

324 16) and it is even higher in Gorna Sushitsa (N = 4), pointing to the consumption of harder foods
325 such as seeds or tubers.

326 It is unlikely that this dietary shift would have resulted from interspecific competition with other
327 primates. The extinction of Pliopithecidae after the Middle Miocene Climatic Transition (Fig. 1),
328 and the progressive extinction of European hominoids during the late Miocene (Merceron et al.
329 2010; Casanovas-Vilar et al. 2011) before the genus *Macaca* disperses to Europe from Africa
330 (Alba et al. 2014), could have instead left dietary niches open to colobines, although hominoids
331 might have co-existed with *Mesopithecus pentelicus* in a different site of Azmaka (respectively
332 found in AZM-4 and AZM-6, Spassov et al. 2012; Böhme et al. 2017). More likely, the dietary
333 shift that we observe (Fig. 2C; Fig. 4) is the consequence of the environmental changes which
334 occurred in southeastern Europe during the late Miocene. The opening of landscapes, especially
335 east of longitude 23°E (Western border of Romania/Bulgaria; Pound et al. 2012) after MN11
336 benefited species with broader dietary niches. This was probably the case for *M. pentelicus* which
337 displays a broad geographic range, from the Iberian peninsula (Alba et al. 2015) to China
338 (Jablonski et al. 2011; Dong and Qi 2013), and was likely able to disperse and inhabit a wide
339 range of habitats, including woodland/savanna environments (Clavel et al. 2012; Pound et al.
340 2012; Kaya et al. 2018).

341 Periods of environmental change are prone to select more generalist species at the expense of
342 specialized ones. *Mesopithecus delsoni* joins the cohort of species affected by the great
343 environmental change which occurs in southeastern Europe between the early/mid-Miocene and
344 the late Miocene, coinciding with the extinction of Moschidae and the decrease in richness of
345 Cervidae as well as forest-affiliated small taxa (Casanovas-Vilar et al. 2005). At the same time,

346 the richness of Bovidae, Equidae, Giraffidae and rodents adapted to open environments increase as
347 the global climate cools and Eurasia aridifies (Merceron et al. 2005, 2010; Fortelius et al. 2014;
348 Cano et al. 2015). It will not be until MN13 and the decline of the evergreen forest in southern
349 Europe (Casanovas-Vilar et al. 2005; Cano et al. 2015) that *M. pentelicus* disperses from eastern
350 Europe to Italy (Pradella and Rook 2007) and then to the Iberian basin (Alba et al. 2015), denoting
351 its high dispersal capacity in a landscape opening context.

352 *Implications for the evolutionary origins of folivory in colobines*

353 While our results do not contradict the general assumption that adaptation to folivory would have
354 emerged from granivory in colobine monkeys (Chivers 1994; Lucas and Teaford 1994), they show
355 that *M. delsoni* during MN11 was less prone to the consumption of hard foods than *M. pentelicus*
356 during MN12 (Fig. 2C; Fig. 3A). Regarding dental topography, much caution is needed, as only a
357 single unworn molar of *M. delsoni* could be examined. Still, its higher average ARC indicates a
358 higher leaf-shearing ability and most likely a lower ability to process hard foods (Fig. 2B).
359 Together, these data indicate that *M. delsoni* was less likely to consume seeds than the later
360 colobine *M. pentelicus*.

361 If a shift from a seed-rich diet towards folivory did actually occur in colobines, two evolutionary
362 scenarios come to mind. On the one hand, folivory could have already emerged in colobines
363 before the dispersal of *Mesopithecus* to Eurasia; *M. pentelicus* would have then adapted to a
364 broader dietary niche, including the consumption of hard foods (Thiery et al. 2017). In fact, the
365 oldest colobine monkeys could have been folivorous to some extent, and colobine monkeys might
366 have evolved sharper molars later as their body size increased, thus resulting in higher nutritional

367 needs (Rossie et al. 2013). A thorough analysis of both dental topography and microwear of
368 *Microcolobus tugenensis* and other early African colobine monkeys is required to test whether
369 folivory was the ancestral dietary niche of European colobines or not.

370 On the other hand, folivorous colobines might have emerged several times, each time when a
371 favorable environmental context occurred. While it seems like a costly hypothesis, the early
372 presence of key physiological adaptations to seed consumption (Happel 1988) might have been
373 co-opted as exaptations to folivory and facilitated the multiple rises of convergent folivorous
374 lineages. As a matter of fact, existing differences in physiology and gut anatomy within extant
375 colobines are not connected to phylogeny but rather reflect the importance of seasonal fruit and
376 seed consumption (Matsuda et al. 2019) which supports the hypothesis of a multiple emergence of
377 the folivory. In addition, the whole-genome sequencing of the golden snub-nosed monkey
378 (*Rhinopithecus roxellana*) revealed that Asian colobines have evolved early on an enhanced
379 ability to derive energy from fatty acids and to degrade xenobiotics (Zhou et al. 2014), which
380 would have directly improved their ability to digest seeds and at the same time facilitated the
381 digestion of low-energy food such as lichens or leaves. Lastly, folivory is hypothesized to have
382 evolved from seed consumption in many mammalian orders, including primates, rodents and
383 marsupials (Cork 1996). Rumination (chewing after microbial pre-digestion into a forestomach)
384 could have also evolved in artiodactyls in order to digest hard seeds too hard to break with the
385 teeth, by directly swallowing them and softening them into the forestomach before re-chewing
386 them, as proposed by Bodmer (1989) while studying the feeding behavior of the brocket deer
387 (*Mazama americana*) in Amazonia. Further comparison is required in order to identify key

388 adaptations which would have enabled the convergent evolution of folivory in early colobines and
389 other potential seed-eating mammals.

390 **Conclusions**

391 This work finds that dental microwear complexity within the genus *Mesopithecus* increased
392 between MN11 and MN12 in southeastern Europe. This observed increase was more important in
393 northern regions corresponding to present Bulgaria, which were then covered by savanna and open
394 mixed forests, whereas in southern regions (notably Pikermi) corresponding to present Greece,
395 there was no increase despite a more open biome. We interpret this result as a dietary niche shift
396 towards a more challenging diet, with the likely inclusion of hard foods such as seeds or
397 underground storage organs, especially in the northernmost populations of *M. pentelicus*. This
398 ecological shift is concurrent with a change in dental topography, the molars of *M. pentelicus*
399 showing lower slopes and sharpness scores compared to *M. delsoni*, which attests of its large scale
400 occurrence, at least at the specific level.

401 While our results do not contradict the assumption that adaptation to folivory would have emerged
402 from granivory in colobine monkeys, they show that *M. delsoni* was more likely to eat leaves
403 during MN11 than *M. pentelicus* during MN12, albeit to a lesser extent than extant colobines. This
404 indicates that colobine monkeys either evolved folivory before their dispersal to Eurasia, or re-
405 evolved folivory in multiple occurrences, likely because of key adaptations to seed predation
406 which would have been co-opted for the consumption of leaf-like foods. This may have included
407 bilophodont molars, but also multi-chambered stomach, rumination strategy or physiological

408 adaptations such as enzymes improving the digestion of fatty acids and the degradation of
409 xenobiotics.

410 This work illustrates how the analysis of the fossil record within a constrained temporal and
411 geographic scale can give insights into the mechanisms underlying adaptation and
412 macroevolution. Here, combining dietary estimates operating on populations at a large timescale,
413 such as dental topography, with estimates operating on single individuals at a small timescale,
414 such as dental microwear textures, allowed us to characterize change in behavior driven by
415 environmental change, and how it ultimately resulted in the emergence of dietary adaptations. This
416 multi-scale approach could be applied to multiple evolutionary discussions related to
417 environmental change and how it can drive dietary evolution in terrestrial tetrapods.

418

419 **Material and methods**

420 ***Material***

421 Dental microwear texture analysis: One hundred sixty nine extant specimens belonging to four
422 African species of cercopithecines and seven of Asian and African colobines were considered. The
423 colobines are represented by *Colobus guereza*, *C. polykomos*, *Ptilocolobus badius* and
424 *Semnopithecus entellus*, *Nasalis larvatus*, *Trachypithecus cristatus*, *T. auratus*. The
425 cercopithecines are represented by *Lophocebus albigena*, *Cercocebus atys*, *Chlorocebus aethiops*

426 and *Erythrocebus patas*. The complete list of the specimens can be found in supplementary
427 materials (Table S9).

428 Sixty six specimens of the extinct genus *Mesopithecus*, coming from Bulgarian and Greek late
429 Miocene sites, were considered. Fifteen specimens are attributed to *Mesopithecus delsoni* from
430 Ravin des Zouaves-5 (type-locality, Greece, N = 3) and to *Mesopithecus aff. delsoni* from
431 Hadjidimovo (Bulgaria, N = 12). Ravin des Zouaves-5 is dated from MN11 (de Bonis et al. 1990)
432 while Hadjidimovo is located at the end/second half of MN11 (Li and Spassov 2017; Spassov et
433 al. 2018). Forty-six specimens are attributed to *M. pentelicus* from Pikermi (type-locality, Greece,
434 N = 26), Kalimantsi (Bulgaria, N = 16), and Gorna Sushitsa (Bulgaria, N = 4; Spassov et al. 2019).
435 Koufos (2009) considered specimens from Perivolaki (Greece, N= 4) and Vathylakkos-2 (Greece,
436 N = 1) as intermediate forms between *M. delsoni* and *M. pentelicus*.

437 Dental topographic analysis: The skulls of twenty seven specimens were selected from the same
438 species as the microwear sample. The list of the specimens can be found in supplementary
439 materials (Table S2). Dental topography and especially dental relief and complexity can be greatly
440 affected by dental wear (Ungar and Williamson 2000; Bunn and Ungar 2009; Pampush et al.
441 2018; Li et al. 2020). In order to mitigate this effect, only the relatively unworn second molars of
442 subadults were selected following previous works (Guy et al. 2013, 2015; Thiery et al. 2017)
443 hence the limited sample size. Scans of extant specimens were combined with scans of molar row
444 casts. In addition, nine specimens of the genus *Mesopithecus* coming from Bulgarian and Greek
445 late Miocene sites were considered. Five previously investigated specimens (Thiery et al. 2017)
446 are attributed to *M. pentelicus* from Pikermi (type-locality, Greece), two specimens from Azmaka
447 (Bulgaria) which are under study will be considered here as *M. cf. pentelicus*, one specimen is

448 attributed to *M. pentelicus* from Gorna Sushitsa (Bulgaria; Spassov et al. 2019) and one to *M. aff.*
449 *delsoni* from Hadjidimovo (Bulgaria).

450 **Methods**

451 Dental Microwear Texture Analysis: Methods for dental microwear texture analysis have been
452 discussed in detail by Scott et al., (2006, 2012; see also Calandra and Merceron 2016). Molars
453 were cleaned with cotton swabs soaked in alcohol in order to remove grit, dust, and glue residues.
454 After cleaning, dental impressions were made with a silicone material (polyvinyl siloxane ISO
455 4823, President Regular Body, Coltène-Whaledent Corp.). Scans were made directly from the
456 silicon molds with the "TRIDENT" Leica DCM8 white-light scanning confocal microscope (Leica
457 Microsystems) with a $\times 100$ long-distance lens (Numerical Aperture=0.90; working
458 distance=0.9mm), housed at the PALEVOPRIM laboratory. For each specimen, one Phase II facet
459 (Facets 9 to 13) was scanned on either the upper or lower molars (M1, M2, M3). The scanning
460 process generates $350 \times 264 \mu\text{m}$ point clouds with a vertical sampling lower than $0.002 \mu\text{m}$ and a
461 lateral sampling (x, y) of $0.129 \mu\text{m}$. The resulting surfaces were then imported in the LeicaMap
462 software (Mountain technology, Leica Microsystems) to remove aberrant peaks with automatic
463 operators including a morphological filter. The surfaces were also leveled and a $200 \times 200 \mu\text{m}$ area
464 was selected and saved as a Digital Elevation Model (Merceron et al. 2016).

465 Complexity (Asfc), heterogeneity of complexity (HASfc) and anisotropy (epLsar) were calculated
466 with Toothfrax software, while textural fill volume (Tfv) was calculated with Sfrax software
467 (www.surfract.com). Hasfc is obtained by subdividing the surface to a 9×9 grid, hence 81 cells,
468 in which texture complexity is calculated. Textural fill volume is the result of an algorithm that

469 fills a surface with square cuboids of different volumes (10 and 2 μm - side cuboids). All DMTA
470 data can be found in supplementary materials (Table S6-9). Inter-group differences were tested
471 after Box-Cox transformation using an ANOVA (Table S7), followed by post-hoc LSD-test (Table
472 S8). Only the most discriminant variable was retained to be compared with topographic variables
473 (Asfc, Table. 2; see supplementary material for details on the evaluation of the discriminant power
474 of variables).

475 Dental topography: Outer enamel surfaces were scanned using high-resolution microtomography
476 at the Centre de Microtomographie de Poitiers. Dental surfaces were segmented and exported as
477 three-dimensional polygonal meshes using Avizo. The meshes were then oriented, smoothed and
478 re-tesselated using Geomagic Studio as in Guy et al., (2015) except that surfaces were re-
479 tessellated to an amount of 10,000 triangles instead of 50,000. Afterwards, the surfaces have been
480 cropped above the lowermost point of the occlusal basin (Ungar and Williamson 2000; Bunn and
481 Ungar 2009).

482 Topographic variables were measured in R 4.0.0 (R Core Team 2021) using the beta version of the
483 package doolkit 0.9.9. Among the variables currently supported by the package, we compared
484 three-dimensional area, occlusal patch count rotated 'OPCR' (Wilson et al. 2012; Guy et al. 2013),
485 Dirichlet normal energy 'DNE' (Bunn et al. 2011), as well as all relief estimates presented in
486 Thiery et al., (2019), notably the average slope (Ungar and Williamson 2000; Bunn and Ungar
487 2009), the relief index 'RFI' (Ungar and Williamson 2000) and the linearized relief index (Boyer
488 2008). We also use an original estimate of tooth sharpness, the area-relative curvature 'ARC' (Guy
489 et al., 2017).

490 In contrast with dental microwear data, the sample size for the topographic analysis was too small
491 for performing a parametric test to compare species. Therefore, we compared the ability of
492 topographic variables to separate colobine monkeys from other cercopithecids using Wilcoxon's
493 rank-sum test (Table 2). Only the most discriminant variables were retained for comparisons with
494 microwear variables (average slope and average ARC, Fig. 2A-B; see supplementary materials for
495 details on the evaluation of the discriminant power of variables). Average ARC and average slope
496 are correlated (Pearson's $r = 0.8$, $p\text{-value} < 0.001$; see supplementary materials: Table S4). Other
497 variables were correlated, such as RFI, LRFI and DNE, but since they measure the same aspect as
498 slope and ARC, they were only represented for fossil taxa (Fig. 3B, 3C, 3E). In contrast, variables
499 which were not correlated to ARC nor slope, such as OPCR (see supplementary materials: Fig. S2;
500 Table S3-S4), were not able to discriminate colobines from other cercopithecids. Nevertheless,
501 OPCR was represented for fossil taxa because it is a widely used variable (Fig. 3F).

502 **References**

- Alba, D. M., E. Delson, G. Carnevale, S. Colombero, M. Delfino, P. Giuntelli, M. Pavia, and G. Pavia. 2014. First joint record of *Mesopithecus* and cf. *Macaca* in the Miocene of Europe. *J. Hum. Evol.* 67:1–18.
- Alba, D. M., P. Montoya, M. Pina, L. Rook, J. Abella, J. Morales, and E. Delson. 2015. First record of *Mesopithecus* (Cercopithecidae, Colobinae) from the Miocene of the Iberian Peninsula. *J. Hum. Evol.* 88:1–14.
- Ataabadi, M. M., L.-P. Liu, J. T. Eronen, R. L. Bernor, and M. Fortelius. 2013. Continental-Scale Patterns in Neogene Mammal Community Evolution and Biogeography. Pp. 629–655 *in* M. Fortelius, X. Wang, and L. Flynn, eds. *Fossil Mammals of Asia*. Columbia University Press.
- Begun, D. R. 2015. Fossil Record of Miocene Hominoids. Pp. 1261–1332 *in* W. Henke and I. Tattersall, eds. *Handbook of Paleoanthropology*. Springer, Berlin, Heidelberg.
- Begun, D. R. 2017. Evolution of the Pliopithecoidea. Pp. 1–4 *in* A. Fuentes, ed. *The international encyclopedia of primatology*. Wiley, Hoboken, NJ.
- Behrensmeyer, A. K., J. Quade, T. E. Cerling, J. Kappelman, I. A. Khan, P. Copeland, L. Roe, J. Hicks, P. Stubblefield, B. J. Willis, and C. Latorre. 2007. The structure and rate of late Miocene expansion of C4 plants: Evidence from lateral variation in stable isotopes in paleosols of the Siwalik Group, northern Pakistan. *GSA Bull.* 119:1486–1505.
- Ben-David, M., and E. A. Flaherty. 2012. Stable isotopes in mammalian research: a beginner's guide. *J. Mammal.* 93:312–328.
- Benefit, B. R., and M. Pickford. 1986. Miocene fossil cercopithecoids from Kenya. *Am. J. Phys. Anthropol.* 69:441–464.

- Bennett, E. L., and A. C. Sebastian. 1988. Social organization and ecology of proboscis monkeys (*Nasalis larvatus*) in mixed coastal forest in Sarawak. *Int. J. Primatol.* 9:233–255.
- Bodmer, R. E. 1989. Frugivory in Amazonian Artiodactyla: evidence for the evolution of the ruminant stomach. *J. Zool.* 219:457–467.
- Böhme, M., N. Spassov, M. Ebner, D. Geraads, L. Hristova, U. Kirscher, S. Kötter, U. Linnemann, J. Prieto, S. Roussiakis, G. Theodorou, G. Uhlig, and M. Winklhofer. 2017. Messinian age and savannah environment of the possible hominin *Graecopithecus* from Europe. *PLOS ONE* 12:e0177347.
- Böhme, M., C. G. C. Van Baak, J. Prieto, M. Winklhofer, and N. Spassov. 2018. Late Miocene stratigraphy, palaeoclimate and evolution of the Sandanski Basin (Bulgaria) and the chronology of the Pikermian faunal changes. *Glob. Planet. Change* 170:1–19.
- Boyer, D. M. 2008. Relief index of second mandibular molars is a correlate of diet among prosimian primates and other euarchontan mammals. *J. Hum. Evol.* 55:1118–1137.
- Bunn, J. M., D. M. Boyer, Y. Lipman, E. M. St. Clair, J. Jernvall, and I. Daubechies. 2011. Comparing Dirichlet normal surface energy of tooth crowns, a new technique of molar shape quantification for dietary inference, with previous methods in isolation and in combination. *Am. J. Phys. Anthropol.* 145:247–261.
- Bunn, J. M., and P. S. Ungar. 2009. Dental topography and diets of four old world monkey species. *Am. J. Primatol.* 71:466–477.
- Calandra, I., and G. Merceron. 2016. Dental microwear texture analysis in mammalian ecology: DMTA in ecology. *Mammal Rev.* 46:215–228.
- Cano, A. R. G., J. L. Cantalapiedra, M. Á. Álvarez-Sierra, and M. H. Fernández. 2015. A macroecological glance at the structure of late Miocene rodent assemblages from Southwest Europe. *Sci. Rep.* 4:6557.

- Casanovas-Vilar, I., D. M. Alba, M. Garcés, J. M. Robles, and S. Moyà-Solà. 2011. Updated chronology for the Miocene hominoid radiation in Western Eurasia. *Proc. Natl. Acad. Sci. U. S. A.* 108:5554–5559.
- Casanovas-Vilar, I., S. Moyà-Solà, J. Agustí, and M. Köhler. 2005. The geography of a faunal turnover: tracking the vallesian crisis. Pp. 247–300 *in* A. M. T. Elewa, ed. *Migration of Organisms*. Springer-Verlag, Berlin/Heidelberg.
- Cerling, T. E., J. M. Harris, and M. G. Leakey. 2005. Environmentally Driven Dietary Adaptations in African Mammals. Pp. 258–272 *in* I. T. Baldwin, M. M. Caldwell, G. Heldmaier, R. B. Jackson, O. L. Lange, H. A. Mooney, E.-D. Schulze, U. Sommer, J. R. Ehleringer, M. Denise Dearing, and T. E. Cerling, eds. *A History of Atmospheric CO₂ and Its Effects on Plants, Animals, and Ecosystems*. Springer, New York, NY.
- Chivers, D. J. 1994. Functional anatomy of the gastrointestinal tract. Pp. 205–228 *in* A. G. Davies and J. F. Oates, eds. *Colobine Monkeys: Their Ecology, Behaviour and Evolution*. Cambridge University Press.
- Clavel, J., G. Merceron, L. Hristova, S. Nikolay, D. Kovachev, and G. Escarguel. 2012. On *Mesopithecus* habitat: Insights from late Miocene fossil vertebrate localities of Bulgaria. *J. Hum. Evol.* 63:162–79.
- Cork, S. J. 1996. Optimal digestive strategies for arboreal herbivorous mammals in contrasting forest types: Why Koalas and Colobines are different. *Aust. J. Ecol.* 21:10–20.
- Dasilva, G. L. 1994. Diet of *Colobus polykomos* on Tiwai Island: Selection of food in relation to its seasonal abundance and nutritional quality. *Int. J. Primatol.* 15:655–680.
- Davies, A. G., J. F. Oates, and G. L. Dasilva. 1999. Patterns of Frugivory in Three West African Colobine Monkeys. *Int. J. Primatol.* 20:327–357.
- Davis, M., and S. Pineda-Munoz. 2016. The temporal scale of diet and dietary proxies. *Ecol. Evol.* 6:1883–1897.

- de Bonis, L., G. Bouvrain, D. Geraads, and G. Koufos. 1990. New remains of *Mesopithecus* (Primates, Cercopithecoidea) from the late Miocene of Macedonia (Greece), with the description of a new species. *J. Vertebr. Paleontol.* 10:473–483.
- De Bonis, L., and G. Koufos. 1999. The Miocene large mammal succession in Greece. Pp. 205–237 *in*.
- DeMiguel, D., D. M. Alba, and S. Moyà-Solà. 2014. Dietary Specialization during the Evolution of Western Eurasian Hominoids and the Extinction of European Great Apes. *PLOS ONE* 9:e97442.
- DeMiguel, D., D. M. Alba, and S. Moyà-Solà. 2013. European pliopithecoid diets revised in the light of dental microwear in *Pliopithecus canmatensis* and *Barberapithecus huerzeleri*. *Am. J. Phys. Anthropol.* 151:573–582.
- DeMiguel, D., L. Domingo, I. M. Sánchez, I. Casanovas-Vilar, J. M. Robles, and D. M. Alba. 2021. Palaeoecological differences underlie rare co-occurrence of Miocene European primates. *BMC Biol.* 19:6.
- Denk, T., C. M. Zohner, G. W. Grimm, and S. S. Renner. 2018. Plant fossils reveal major biomes occupied by the late Miocene Old-World Pikermian fauna. *Nat. Ecol. Evol.* 2:1864–1870.
- Dominy, N. J., E. R. Vogel, J. D. Yeakel, P. Constantino, and P. W. Lucas. 2008. Mechanical Properties of Plant Underground Storage Organs and Implications for Dietary Models of Early Hominins. *Evol. Biol.* 35:159–175.
- Dong, W., and G. Qi. 2013. Hominoid-Producing Localities and Biostratigraphy in Yunnan. Pp. 293–313 *in* *Fossil Mammals of Asia: Neogene Biostratigraphy and Chronology*.
- Elton, S., and H. J. O'Regan. 2014. Macaques at the margins: the biogeography and extinction of *Macaca sylvanus* in Europe. *Quat. Sci. Rev.* 96:117–130.
- Eronen, J. T., and L. Rook. 2004. The Mio-Pliocene European primate fossil record: dynamics and habitat tracking. *J. Hum. Evol.* 47:323–341.

- Fashing, P. J. 2001. Feeding Ecology of Guerezas in the Kakamega Forest, Kenya: The Importance of Moraceae Fruit in Their Diet. *Int. J. Primatol.* 22:579–609.
- Fortelius, M., J. T. Eronen, F. Kaya, H. Tang, P. Raia, and K. Puolamäki. 2014. Evolution of Neogene Mammals in Eurasia: Environmental Forcing and Biotic Interactions. *Annu. Rev. Earth Planet. Sci.* 42:579–604.
- Gould, S. J., and E. S. Vrba. 1982. Exaptation-A Missing Term in the Science of Form. *Paleobiology* 8:4–15.
- Groves, C. 2001. *Primate taxonomy*. Smithsonian Institution Print. Washington D.C.
- Guérin, C. 1980. Les rhinocéros (Mammalia, Perissodactyla) du Miocène terminal au Pléistocène supérieur en Europe occidentale. Comparaison avec les espèces actuelles (fascicule 3). *Trav. Doc. Lab. Géologie Lyon* 79:787–1185. Persée - Portail des revues scientifiques en SHS.
- Guérin, C. 1982. Première biozonation du Pléistocène Européen, principal résultat biostratigraphique de l'étude des Rhinocerotidae (Mammalia, Perissodactyla) du Miocène terminal au Pléistocène supérieur d'Europe Occidentale. *Geobios* 15:593–598.
- Guy, F., F. Gouvard, R. Boistel, A. Euriat, and V. Lazzari. 2013. Prospective in (Primate) Dental Analysis through Tooth 3D Topographical Quantification. *PLoS ONE* 8:e66142.
- Guy, F., V. Lazzari, E. Gilissen, and G. Thiery. 2015. To What Extent is Primate Second Molar Enamel Occlusal Morphology Shaped by the Enamel-Dentine Junction? *PLOS ONE* 10:e0138802.
- Guy, F., G. Thiery, and V. Lazzari. 2017. 3D quantification of the occlusal enamel curvature: a decisive feature in dental function analysis and diet in Primates. 17th International Symposium on Dental Morphology (ISDM) & 2nd congress of International Association for Paleodontology (IAPO).

- Happel, R. 1988. Seed-eating by West African cercopithecines, with reference to the possible evolution of bilophodont molars. *Am. J. Phys. Anthropol.* 75:303–327.
- Harrison, T. 2005. The zoogeographic and phylogenetic relationships of early catarrhine primates in Asia. *Anthropol. Sci.* 113:43–51.
- Hladik, C. M. 1977. A comparative study of two sympatric species of leaf monkeys: *Presbytis entellus* and *Presbytis senex*. Pp. 333–353 in T. H. Clutton-Brock, ed. *Primate Ecology*. Academic Press, London.
- Isbell, L. A. 1998. Diet for a small primate: Insectivory and gummivory in the (large) patas monkey (*Erythrocebus patas pyrrhonotus*). *Am. J. Primatol.* 45:381–398.
- Isbell, L. A., J. D. Pruettz, and T. P. Young. 1998. Movements of vervets (*Cercopithecus aethiops*) and patas monkeys (*Erythrocebus patas*) as estimators of food resource size, density, and distribution. *Behav. Ecol. Sociobiol.* 42:123–133.
- Jablonski, N., D. Su, J. Kelley, L. Flynn, and X. Ji. 2011. The Mio-Pliocene colobine monkey, *Mesopithecus*, in China. *Am J Phys Anthr.* 144.
- Janis, C. M., J. Damuth, and J. M. Theodor. 2000. Miocene ungulates and terrestrial primary productivity: Where have all the browsers gone? *Proc. Natl. Acad. Sci.* 97:7899–7904. National Academy of Sciences.
- Kay, R. F. 1977. The evolution of molar occlusion in the Cercopithecidae and early catarrhines. *Am. J. Phys. Anthropol.* 46:327–352.
- Kay, R. F. 1975. The functional adaptations of primate molar teeth. *Am. J. Phys. Anthropol.* 43:195–215.
- Kay, R. F., and K. M. Hiiemae. 1974. Jaw movement and tooth use in recent and fossil primates. *Am. J. Phys. Anthropol.* 40:227–256.

- Kaya, F., F. Bibi, I. Žliobaitė, J. T. Eronen, T. Hui, and M. Fortelius. 2018. The rise and fall of the Old World savannah fauna and the origins of the African savannah biome. *Nat. Ecol. Evol.* 2:241–246.
- King, T., L. C. Aiello, and P. Andrews. 1999. Dental microwear of *Griphopithecus alpani*. *J. Hum. Evol.* 36:3–31.
- Kool, K. M. 1993. The diet and feeding behavior of the silver leaf monkey (*Trachypithecus auratus sondaicus*) in Indonesia. *Int. J. Primatol.* 14:667–700.
- Kostopoulos, D. S. 2009. The Pikermian Event: Temporal and spatial resolution of the Turolian large mammal fauna in SE Europe. *Palaeogeogr. Palaeoclimatol. Palaeoecol.* 274:82–95.
- Koufos, G., L. Bonis, D. Kostopoulos, L. Viriot, and T. Vlachou. 2004. New material of *Mesopithecus* (Primates, Cercopithecidae) from the Turolian locality of Vathylakkos 2, Macedonia, Greece. *Paläontol. Z.* 78:213–228.
- Koufos, G. D. 2009. The Neogene cercopithecids (Mammalia, Primates) of Greece. *Geodiversitas* 31:817–850.
- Lambert, J. E., C. A. Chapman, R. W. Wrangham, and N. L. Conklin-Brittain. 2004. Hardness of cercopithecine foods: Implications for the critical function of enamel thickness in exploiting fallback foods. *Am. J. Phys. Anthropol.* 125:363–368.
- Li, P., P. E. Morse, and R. F. Kay. 2020. Dental topographic change with macrowear and dietary inference in *Homunculus patagonicus*. *J. Hum. Evol.* 144:102786.
- Li, Y., and N. Spassov. 2017. A new species of *Paramachaerodus* (Mammalia, Carnivora, Felidae) from the late Miocene of China and Bulgaria, and revision of *Promegantereon Kretzoi*, 1938 and *Paramachaerodus* Pilgrim, 1913. *PalZ* 91:409–426.
- Lucas, P. 2000. Mechanical Defences to Herbivory. *Ann. Bot.* 86:913–920.

- Lucas, P., and M. F. Teaford. 1994. Functional morphology of colobine teeth. Pp. 173–204 in A. G. Davies and J. F. Oates, eds. *Colobine Monkeys: Their Ecology, Behaviour and Evolution*. Cambridge University Press.
- Matsuda, I., C. A. Chapman, and M. Clauss. 2019. Colobine forestomach anatomy and diet. *J. Morphol.* 280:1608–1616.
- Matsuda, I., S. Higashi, Y. Otani, A. Tuuga, H. Bernard, and R. T. Corlett. 2013. A short note on seed dispersal by colobines: the case of the proboscis monkey. *Integr. Zool.* 8:395–399.
- McGraw, W. S., A. van Casteren, E. Kane, E. Geissler, B. Burrows, and D. J. Daegling. 2016. Feeding and oral processing behaviors of two colobine monkeys in Tai Forest, Ivory Coast. *J. Hum. Evol.* 98:90–102.
- McGraw, W. S., A. E. Vick, and D. J. Daegling. 2014. Dietary variation and food hardness in sooty mangabeys (*Cercocebus atys*): Implications for fallback foods and dental adaptation: Seasonality Of Feeding In Sooty Mangabeys. *Am. J. Phys. Anthropol.* 154:413–423.
- Merceron, G., E. Berlioz, H. Vonhof, D. Green, M. Garel, and T. Tütken. 2021. Tooth tales told by dental diet proxies: An alpine community of sympatric ruminants as a model to decipher the ecology of fossil fauna. *Palaeogeogr. Palaeoclimatol. Palaeoecol.* 562:110077.
- Merceron, G., C. Blondel, L. D. Bonis, G. D. Koufos, and L. Viriot. 2005. A New Method of Dental Microwear Analysis: Application to Extant Primates and *Ouranopithecus macedoniensis* (Late Miocene of Greece). *PALAIOS* 20:551–561.
- Merceron, G., M. Colyn, and D. Geraads. 2018. Browsing and non-browsing extant and extinct giraffids: Evidence from dental microwear textural analysis. *Palaeogeogr. Palaeoclimatol. Palaeoecol.* 505:128–139.
- Merceron, G., L. Costeur, O. Maridet, A. Ramdarshan, and U. B. Göhlich. 2012. Multi-proxy approach detects heterogeneous habitats for primates during the Miocene climatic optimum in Central Europe. *J. Hum. Evol.* 63:150–161.

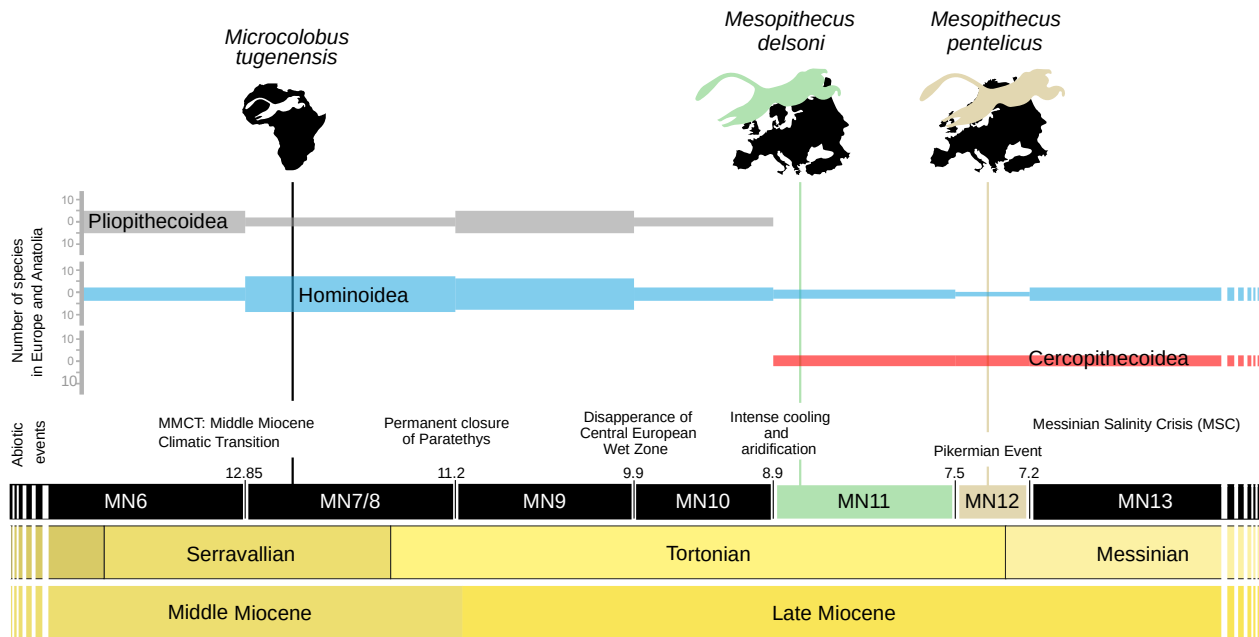
- Merceron, G., T. M. Kaiser, D. S. Kostopoulos, and E. Schulz. 2010. Ruminant diets and the Miocene extinction of European great apes. *Proc. R. Soc. B Biol. Sci.* 277:3105–3112.
- Merceron, G., D. S. Kostopoulos, L. de Bonis, F. Fourel, G. D. Koufos, C. Lécuyer, and F. Martineau. 2013. Stable isotope ecology of Miocene bovids from northern Greece and the ape/monkey turnover in the Balkans. *J. Hum. Evol.* 65:185–198.
- Merceron, G., G. D. Koufos, and X. Valentin. 2009a. Feeding habits of the first European colobine, *Mesopithecus* (Mammalia, Primates): evidence from a comparative dental microwear analysis with modern cercopithecids. *Geodiversitas* 31:865–878.
- Merceron, G., A. Ramdarshan, C. Blondel, J.-R. Boisserie, N. Brunetiere, A. Francisco, D. Gautier, X. Milhet, A. Novello, and D. Pret. 2016. Untangling the environmental from the dietary: dust does not matter. *Proc. R. Soc. B Biol. Sci.* 283:20161032.
- Merceron, G., E. Schulz, L. Kordos, and T. M. Kaiser. 2007. Paleoenvironment of *Dryopithecus brancoi* at Rudabánya, Hungary: evidence from dental meso- and micro-wear analyses of large vegetarian mammals. *J. Hum. Evol.* 53:331–349.
- Merceron, G., J. Scott, R. S. Scott, D. Geraads, N. Spassov, and P. S. Ungar. 2009b. Folivory or fruit/seed predation for *Mesopithecus*, an earliest colobine from the late Miocene of Eurasia? *J. Hum. Evol.* 57:732–738.
- Montañez, I. P., N. J. Tabor, D. Niemeier, W. A. DiMichele, T. D. Frank, C. R. Fielding, J. L. Isbell, L. P. Birgenheier, and M. C. Rygel. 2007. CO₂-Forced Climate and Vegetation Instability During Late Paleozoic Deglaciation. *Science* 315:87–91.
- Moyà-Solà, S., D. M. Alba, S. Almécija, I. Casanovas-Vilar, M. Köhler, S. D. Esteban-Trivigno, J. M. Robles, J. Galindo, and J. Fortuny. 2009. A unique Middle Miocene European hominoid and the origins of the great ape and human clade. *Proc. Natl. Acad. Sci.* 106:9601–9606.

- Nüchel, J., P. K. Bøcher, W. Xiao, A.-X. Zhu, and J.-C. Svenning. 2018. Snub-nosed monkeys (*Rhinopithecus*): potential distribution and its implication for conservation. *Biodivers. Conserv.* 27:1517–1538.
- Oates, J. F. 1994. The natural history of African colobines. Pp. 75–128 in A. G. Davies and J. F. Oates, eds. *Colobine monkeys: Their ecology, behaviour and evolution*. New York, NY.
- Osborne, C. P. 2008. Atmosphere, ecology and evolution: what drove the Miocene expansion of C4 grasslands? *J. Ecol.* 96:35–45.
- Osborne, C. P., and D. J. Beerling. 2006. Nature's green revolution: the remarkable evolutionary rise of C4 plants. *Philos. Trans. R. Soc. Lond. B. Biol. Sci.* 361:173–194.
- Pampush, J. D., J. P. Spradley, P. E. Morse, D. Griffith, J. T. Gladman, L. A. Gonzales, and R. F. Kay. 2018. Adaptive wear-based changes in dental topography associated with atelid (Mammalia: Primates) diets. *Biol. J. Linn. Soc.* 124:584–606.
- Pound, M. J., A. M. Haywood, U. Salzmann, and J. B. Riding. 2012. Global vegetation dynamics and latitudinal temperature gradients during the Mid to Late Miocene (15.97–5.33Ma). *Earth-Sci. Rev.* 112:1–22.
- Pradella, C., and L. Rook. 2007. *Mesopithecus* (Primates: Cercopithecoidea) from Villafranca d'Asti (Early Villafranchian; NW Italy) and palaeoecological context of its extinction. *Swiss J. Geosci.* 100:145–152.
- R Core Team. 2021. *R: A language and environment for statistical computing*. R Foundation for Statistical Computing, Vienna, Austria.
- Ramdarshan, A., C. Blondel, N. Brunetière, A. Francisco, D. Gautier, J. Surault, and G. Merceron. 2016. Seeds, browse, and tooth wear: a sheep perspective. *Ecol. Evol.* 6:5559–5569.
- Rossie, J. B., C. C. Gilbert, and A. Hill. 2013. Early cercopithecoid monkeys from the Tugen Hills, Kenya. *Proc. Natl. Acad. Sci.* 110:5818–5822.

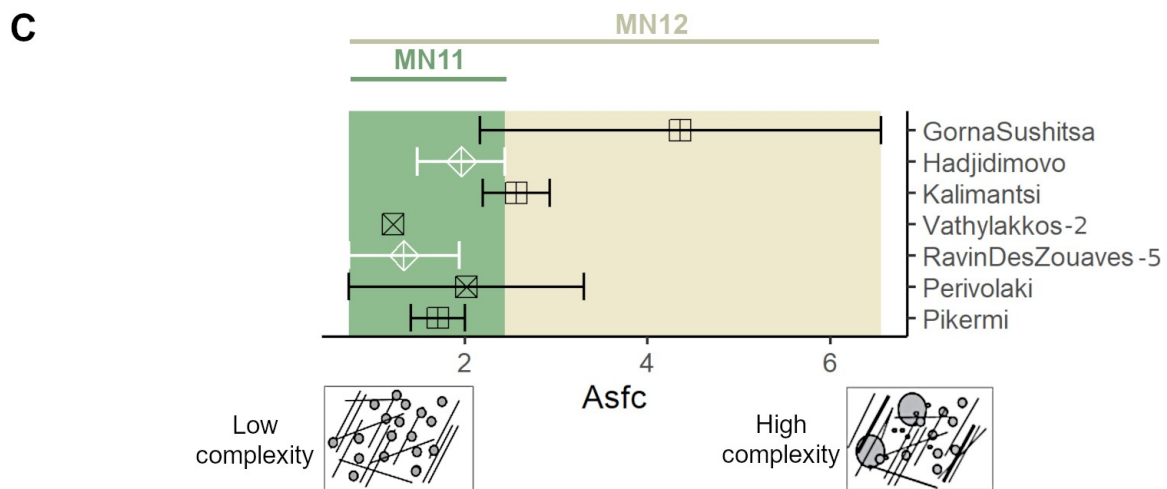
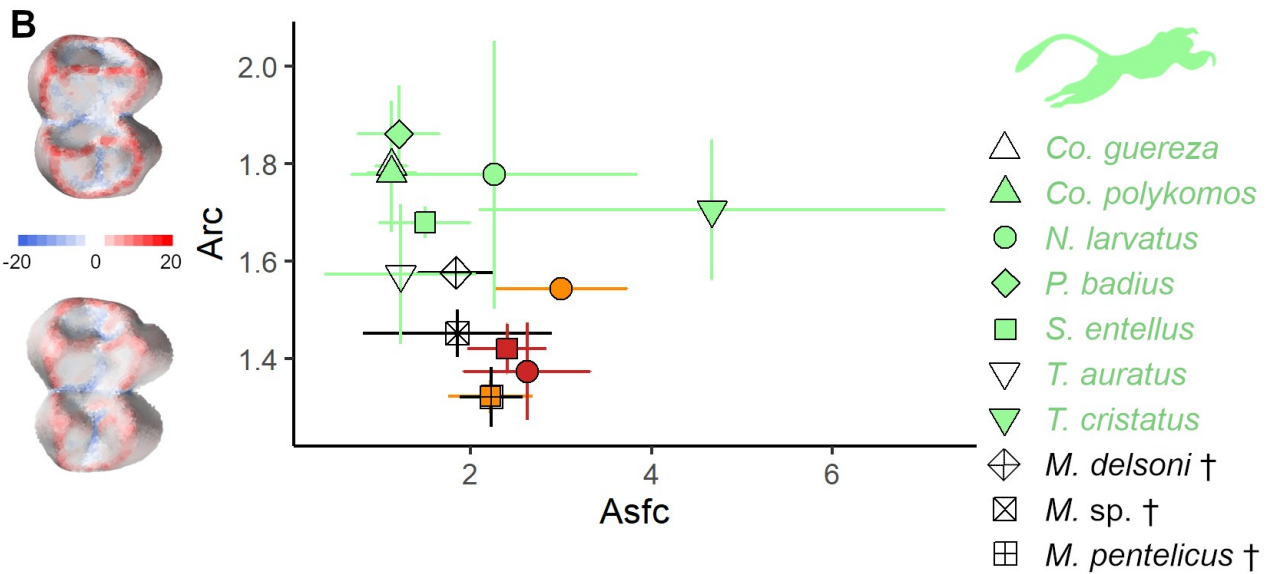
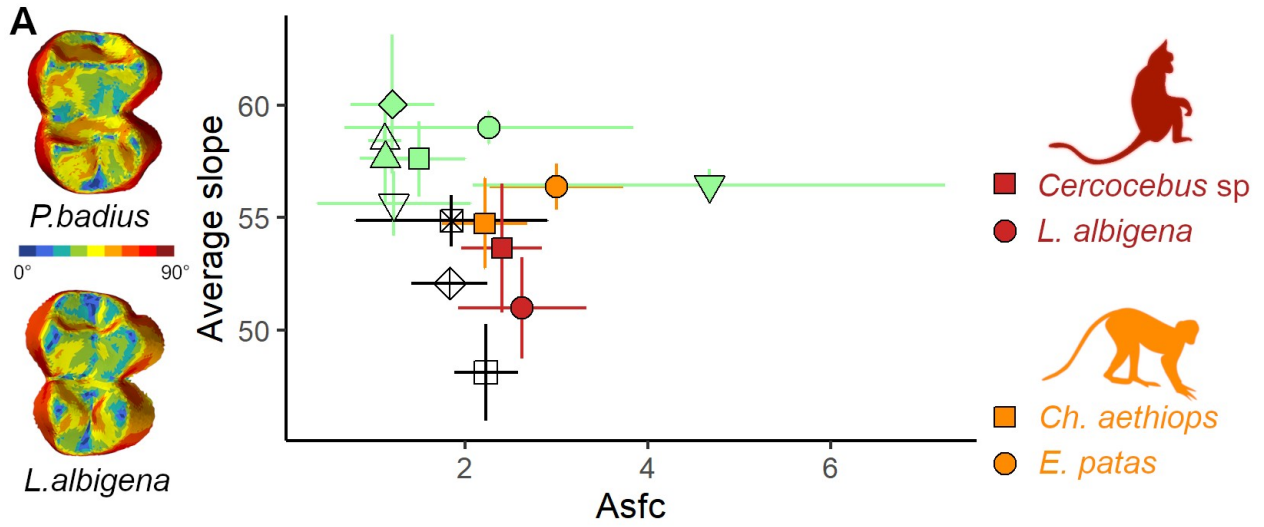
- Sahney, S., M. J. Benton, and H. J. Falcon-Lang. 2010. Rainforest collapse triggered Carboniferous tetrapod diversification in Euramerica. *Geology* 38:1079–1082.
- Sayers, K., and M. A. Norconk. 2008. Himalayan *Semnopithecus entellus* at Langtang National Park, Nepal: Diet, Activity Patterns, and Resources. *Int. J. Primatol.* 29:509–530.
- Schulz-Kornas, E., J. Stuhlträger, M. Clauss, R. M. Wittig, and K. Kupczik. 2019. Dust affects chewing efficiency and tooth wear in forest dwelling Western chimpanzees (*Pan troglodytes verus*). *Am. J. Phys. Anthropol.* 169:66–77.
- Scott, R. S., M. F. Teaford, and P. S. Ungar. 2012. Dental microwear texture and anthropoid diets. *Am. J. Phys. Anthropol.* 147:551–579.
- Scott, R. S., P. S. Ungar, T. S. Bergstrom, C. A. Brown, B. E. Childs, M. F. Teaford, and A. Walker. 2006. Dental microwear texture analysis: technical considerations. *J. Hum. Evol.* 51:339–349.
- Spassov, N., D. Geraads, L. Hristova, and G. N. Markov. 2019. The late Miocene mammal fauna from Gorna Sushitsa, southwestern Bulgaria, and the early/middle Turolian transition. *Neues Jahrb. Fr Geol. Palontologie - Abh.* 291:317–350.
- Spassov, N., D. Geraads, L. Hristova, G. N. Markov, B. Garevska, and R. † Garevski. 2018. The late Miocene mammal faunas of the Republic of Macedonia (FYROM). *Palaeontogr. Abt. A* 1–85. Schweizerbart'sche Verlagsbuchhandlung.
- Spassov, N., D. Geraads, L. Hristova, G. N. Markov, G. Merceron, T. Tzankov, K. Stoyanov, M. Böhme, and A. Dimitrova. 2012. A hominid tooth from Bulgaria: The last pre-human hominid of continental Europe. *J. Hum. Evol.* 62:138–145.
- Sukselainen, L., M. Fortelius, and T. Harrison. 2015. Co-occurrence of pliopithecoid and hominoid primates in the fossil record: An ecometric analysis. *J. Hum. Evol.* 84:25–41.
- Teaford, M. F., and O. J. Oyen. 1989. In vivo and in vitro turnover in dental microwear. *Am. J. Phys. Anthropol.* 80:447–460.

- Teaford, M., P. Ungar, A. Taylor, C. Ross, and C. Vinyard. 2017. In vivo rates of dental microwear formation in laboratory primates fed different food items. *Biosurface Biotribology* 3.
- Thiery, G., G. Gillet, V. Lazzari, G. Merceron, and F. Guy. 2017. Was *Mesopithecus* a seed eating colobine? Assessment of cracking, grinding and shearing ability using dental topography. *J. Hum. Evol.* 112:79–92.
- Thiery, G., F. Guy, and V. Lazzari. 2019. A comparison of relief estimates used in three-dimensional dental topography. *Am. J. Phys. Anthropol.* 170:260–274.
- Ungar, P. S., and R. F. Kay. 1995. The dietary adaptations of European Miocene catarrhines. *Proc. Natl. Acad. Sci.* 92:5479–5481.
- Ungar, P., and M. Williamson. 2000. Exploring the effects of tooth wear on functional morphology: a preliminary study using dental topography. *Palaeontol. Electron.* 3:1–18.
- Utescher, T., A. A. Bruch, A. Micheels, V. Mosbrugger, and S. Popova. 2011. Cenozoic climate gradients in Eurasia — a palaeo-perspective on future climate change? *Palaeogeogr. Palaeoclimatol. Palaeoecol.* 304:351–358.
- van Dam, J. A. 2006. Geographic and temporal patterns in the late Neogene (12–3 Ma) aridification of Europe: The use of small mammals as paleoprecipitation proxies. *Palaeogeogr. Palaeoclimatol. Palaeoecol.* 238:190–218.
- Van Dam, J. A., L. Alcalá, A. A. Zarza, J. P. Calvo, M. Garcés, and W. Krijgsman. 2001. The Upper Miocene Mammal Record from the Teruel-Alfambra Region (Spain). The MN System and Continental Stage/Age Concepts Discussed. *J. Vertebr. Paleontol.* 21:367–385. [Society of Vertebrate Paleontology, Taylor & Francis, Ltd.].
- Whitten, P. L. 1983. Diet and dominance among female vervet monkeys (*Cercopithecus aethiops*). *Am. J. Primatol.* 5:139–159.

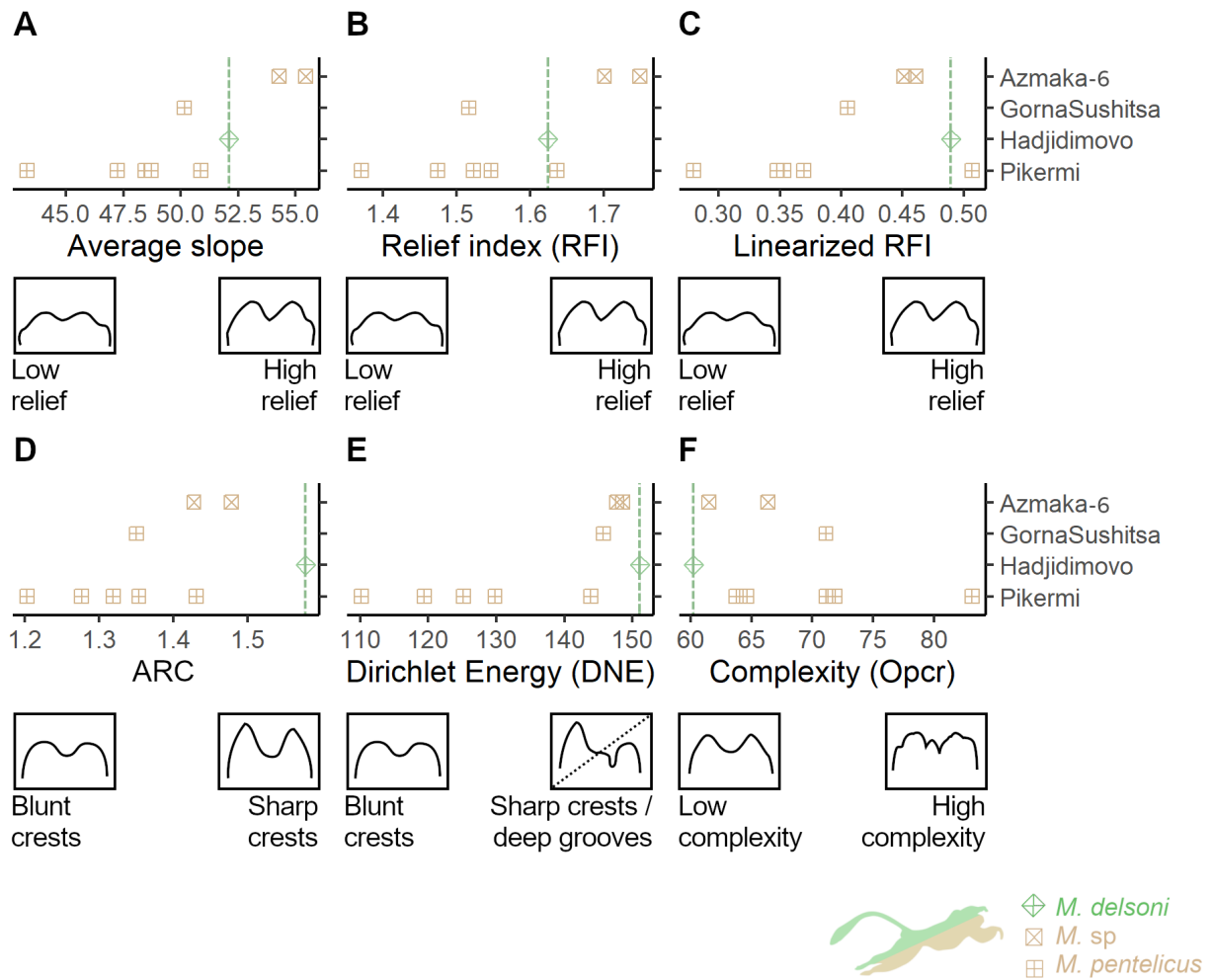
- Wilson, G. P., A. R. Evans, I. J. Corfe, P. D. Smits, M. Fortelius, and J. Jernvall. 2012. Adaptive radiation of multituberculate mammals before the extinction of dinosaurs. *Nature* 483:457–460.
- Winchester, J. M., D. M. Boyer, E. M. St. Clair, A. D. Gosselin-Ildari, S. B. Cooke, and J. A. Ledogar. 2014. Dental topography of platyrrhines and prosimians: Convergence and contrasts: Dental Topography of Noncatarrhines. *Am. J. Phys. Anthropol.* 153:29–44.
- Wrangham, R. W., and P. G. Waterman. 1981. Feeding Behaviour of Vervet Monkeys on *Acacia tortilis* and *Acacia xanthophloea*: With Special Reference to Reproductive Strategies and Tannin Production. *J. Anim. Ecol.* 50:715.
- Yamashita, N., F. P. Cuzzo, and M. L. Sauter. 2012. Interpreting food processing through dietary mechanical properties: A *Lemur catta* case study. *Am. J. Phys. Anthropol.* 148:205–214.
- Yeager, C. P. 1989. Feeding ecology of the proboscis monkey (*Nasalis larvatus*). *Int. J. Primatol.* 10:497–530.
- Zhou, X., B. Wang, Q. Pan, J. Zhang, S. Kumar, X. Sun, Z. Liu, H. Pan, Y. Lin, G. Liu, W. Zhan, M. Li, B. Ren, X. Ma, H. Ruan, C. Cheng, D. Wang, F. Shi, Y. Hui, Y. Tao, C. Zhang, P. Zhu, Z. Xiang, W. Jiang, J. Chang, H. Wang, Z. Cao, Z. Jiang, B. Li, G. Yang, C. Roos, P. A. Garber, M. W. Bruford, R. Li, and M. Li. 2014. Whole-genome sequencing of the snub-nosed monkey provides insights into folivory and evolutionary history. *Nat. Genet.* 46:1303–1310. Nature Publishing Group.



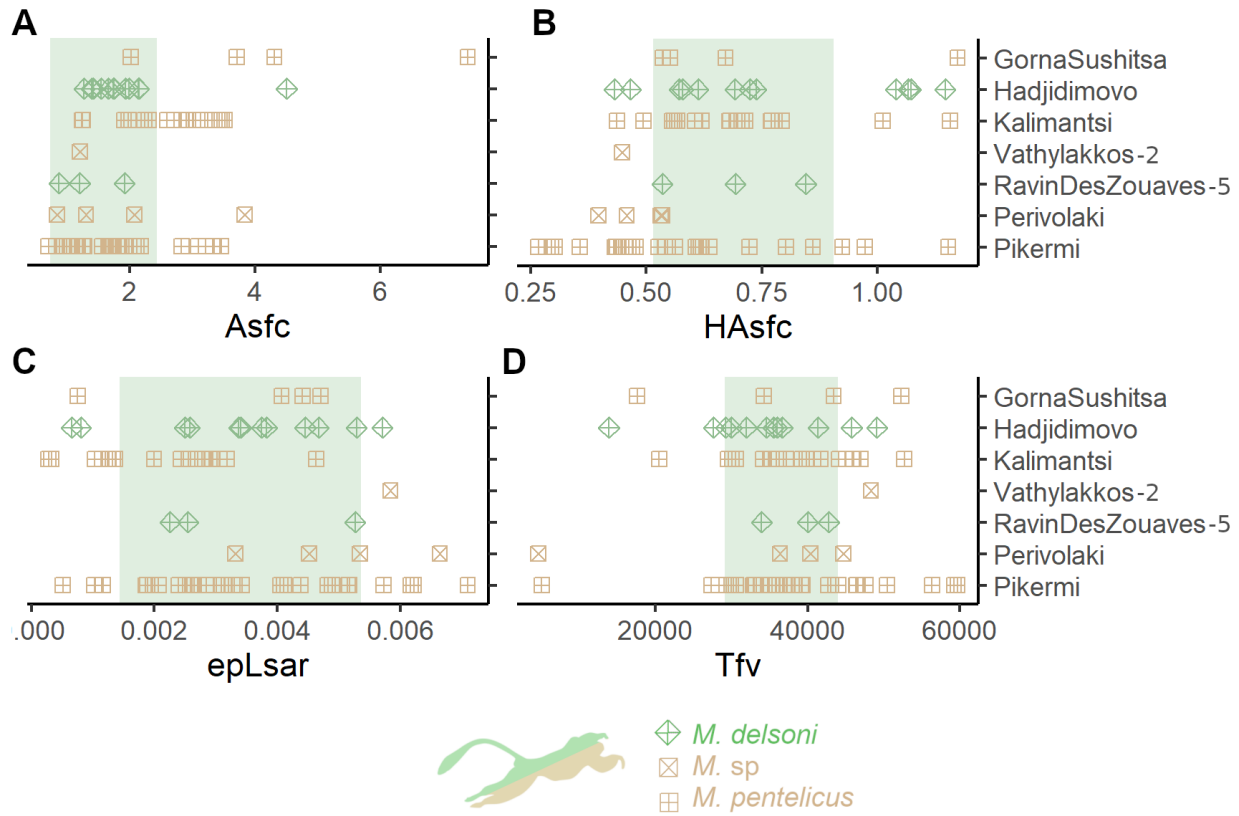
503 Figure 1. Chronology of the successive primate radiations that took place in western Eurasia
 504 during the Miocene (see Table S1). The progressive disappearance of the Central European Wet
 505 Zone (Utescher et al. 2011) coincides with a decrease in biodiversity of European pliopithecoids
 506 (gray) and hominoids (blue). Although the first part of the evolutionary history of cercopithecids
 507 took place in Africa, they appear in the fossil record of Europe during the late Miocene with the
 508 genus *Mesopithecus* (red). This study compares *M. delsoni* and *M. pentelicus* specimens found in
 509 the Balkans within MN11 and MN12 Mammal Neogene zones (Guérin, 1980, 1982), with
 510 populations from the MN12 sharing features with both *M. delsoni* and *M. pentelicus*.
 511



513 Figure 2. The dietary niches of *M. delsoni* and *M. pentelicus* compared to extant catarrhines.
514 *Mesopithecus* sp. groups specimens that are similar to *M. pentelicus* but whose taxonomic
515 affinities are not clear (*M. cf. pentelicus* from Azmaka and *M. delsoni/pentelicus* from Perivolaki
516 and Vathylakkos-2). Data points correspond to specific means with error bars marking confidence
517 intervals (standard error of the mean with $\alpha = 0.05$). (A) Bivariate plot of the complexity of
518 enamel wear facets (Asfc) versus molar average slope. (B) Bivariate plot of Asfc versus average
519 molar area-relative curvature (ARC). (C) Asfc of extinct specimens ordered by site, from the
520 northernmost (Gorna Sushitsa) to the southernmost location (Pikermi).



521 Figure 3. Dental topographic analysis of *Mesopithecus* from the Balkans. Variables are ordered by
 522 site, from the northernmost (Gorna Sushitsa) to the southernmost location (Pikermi). Dental relief
 523 estimated from (A) average slope; (B) relief index sensu Ungar and Williamson (2000); and (C)
 524 relief index sensu Boyer (2008); dental sharpness estimated from (D) average area-relative
 525 curvature (ARC) and (E) Dirichlet normal energy (DNE). (F) Complexity estimated from
 526 Orientation Patch Count Rotated (OPCR).



527 Figure 4. Dental microwear texture analysis of *Mesopithecus* from the Balkans. Variables are
 528 ordered by site, from the northernmost (Gorna Sushitsa) to the southernmost location (Pikermi).
 529 (A) Textural complexity (Asfc). (B) Heterogeneity of Asfc (HASfc). (C) Anisotropy (epLsar). (D)
 530 Textural fill volume (Tfv). The green area shows the confidence interval for *M. delsoni* values
 531 (standard error of the mean with $\alpha = 0.05$).

532 Table 1. Results of post hoc LSD tests on texture parameters obtained from Phase II facets of
 533 extant cercopithecids.

	<i>L.a.</i>	<i>C.a.</i>	<i>Ch.a.</i>	<i>E.p.</i>	<i>Co.g.</i>	<i>Co.p.</i>	<i>P.b.</i>	<i>N.l.</i>	<i>S.e.</i>	<i>T.a.</i>	<i>T.c.</i>
<i>L. albigena</i>	■										
<i>C. atys</i>	Hasfc	■									
<i>Ch. aethiops</i>	Hasfc		■								
<i>E. patas</i>			Hasfc	■							
<i>Co. guereza</i>	Asfc epLsar Tfv	Asfc epLsar Tfv	Asfc epLsar Tfv	Asfc epLsar Tfv	■						
<i>Co. polykomos</i>	Asfc epLsar Tfv	Asfc epLsar Tfv	Asfc, epLsar Tfv	Asfc epLsar Tfv		■					
<i>P. badius</i>	Asfc epLsar	Asfc epLsar	Asfc epLsar	Asfc epLsar		Tfv	■				
<i>N. larvatus</i>		Hasfc	Hasfc	Asfc	epLsar	EpLsar Tfv		■			
<i>S. entellus</i>	Asfc	Asfc		Asfc					■		
<i>T. auratus</i>	Asfc epLsar	Asfc epLsar	Asfc epLsar	Asfc epLsar		Tfv		Asfc epLsar		■	
<i>T. cristatus</i>			Hasfc		Asfc epLsar Tfv	Asfc epLsar Tfv	Asfc epLsar	Asfc	Asfc	Asfc epLsar	■

534 Only variables with a significant P-value for the LSD test are shown ($\alpha = 0.05$). Comparison
 535 between extant colobines and cercopithecines is highlighted in grey. See supplementary materials
 536 (Table S8) for more details.

537 Table 2. Ability of topographic variables to separate colobines from other cercopithecids,
538 evaluated using Wilcoxon's rank-sum test. Variables classified from the most discriminant (top) to
539 the least discriminant (bottom).

Wilcoxon		
Variable	U	P-value
Average Arc	2	0.000
Average slope	15	0.000
Rfi	21	0.000
Dne	41	0.014
Lrfi	49	0.043
Area	50	0.048
Opcr	81	0.645

540