

1 **Feeding ecology of Omo River guerezas (*Colobus guereza guereza*) in natural versus**
2 **plantation forests in the central highlands of Ethiopia**

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INCLUSION AND DIVERSITY STATEMENT

The author list includes contributors from the location where the research was conducted, who participated in study conception, study design, data collection, analysis, and/or interpretation of the findings.

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AUTHORS CONTRIBUTIONS

DY, AB, PJF, NN and AMe originally formulated the idea and designed this study, DY carried out the fieldwork, organized and analyzed the data, and DY, AB, PJF, NN, HI, AMo, TME, and AMe wrote the manuscript.

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51 **Abstract:** Understanding the impacts of habitat modification on primate feeding ecology is
52 essential for designing effective conservation management strategies. The dietary guild (e.g.,
53 frugivore, folivore, insectivore, and omnivore) of primates and their degree of ecological
54 flexibility impacts their ability to cope with human-modified habitats. The Omo River guereza
55 (*Colobus guereza guereza*) is a subspecies of eastern black-and-white colobus monkey endemic
56 to the western Rift Valley forests of Ethiopia, where it faces increasing anthropogenic change.
57 While there is some understanding of how this subspecies copes with anthropogenic pressures,
58 we aimed to compare the feeding ecology of Omo River guerezas in natural and human-modified
59 habitats. Specifically, we collected data on two neighbouring guereza groups inhabiting adjacent
60 plantation and natural forest habitats over 12 months in Wof-Washa Natural State Forest in the
61 central highlands of Ethiopia. Furthermore, we conducted vegetation surveys on the botanical
62 composition and vertical structure of both habitat types. The monthly food availability index of
63 young leaves was higher in the natural forest than in plantation forest habitat. We observed
64 guerezas feeding on 30 plant species in the natural forest but only 18 species in the plantation
65 forest. Guerezas in both forest types consumed mostly young leaves, but the natural forest group
66 relied more on mature leaves and shoots, and less on fruits and stems, than the plantation forest
67 group. *Maesa lanceolata* leaves contributed a greater proportion of the overall diet for the
68 plantation forest group, while *Vernonia leopoldi* accounted for the largest proportion of the
69 guereza diet for the natural forest group. The top five species consumed comprised 83% of the
70 diet in the plantation forest group and 70% in the natural forest group, indicating that relatively
71 few plant species dominate guereza diets in these habitats. Conservation of both natural and
72 plantation forests, especially the plant species most intensively exploited by guerezas, should be
73 prioritized to assist in Omo River guereza conservation efforts.

74 **Keywords:** Colobus monkeys, Ethiopia, guereza, human-modified habitat, Wof-Washa Natural
75 State Forest.

76

77 **INTRODUCTION**

78 The exponential growth of human populations and the consequent impact on natural
79 environments have led to degraded and fragmented habitats across landscapes, driving native
80 fauna to either adapt or become locally extirpated (Estrada et al. 2017; Estrada et al. 2019;
81 Mekonnen et al. 2017; Mekonnen et al. 2018a). Among the diverse fauna faced with
82 anthropogenic habitat loss and fragmentation are nonhuman primates (hereafter primates), a large
83 taxonomic order whose populations are declining globally, with many taxa listed within the
84 threatened categories (i.e., Vulnerable, Endangered and Critically Endangered) of the IUCN Red
85 List (Estrada et al. 2017; Fernández et al. 2022; Torres-Romero et al. 2023)

86 Understanding how animals cope with habitat degradation and fragmentation is urgently
87 needed given the alarming rate at which natural habitats are being altered, ultimately limiting the
88 ability of some species to either persist within a habitat fragment or move between fragments
89 (Marsh and Chapman 2013; Galan-Acedo et al. 2019). The type and intensity of land-use changes
90 (both historical and present day) are major determinants of biodiversity in many landscapes
91 (Galan-Acedo et al. 2021; Redei et al. 2020; Torres-Romero et al. 2023). Extensive
92 transformation of natural habitats to agroecosystems (e.g., cultivation, plantations, etc.) and
93 urbanization invariably erodes wild food resources (McLennan and Hockings 2014). In turn, this
94 may pose a threat to dietary specialists putting them at higher risk of local extirpation when
95 compared to dietary generalists (Boyle and Smith 2010; Eppley et al. 2020; Machado et al. 2022;
96 Mekonnen et al. 2018b). Despite these challenges, some species can persist in human-modified

97 habitats by incorporating agricultural crops and exotic (non-native) flora into their diet (Eppley et
98 al. 2017; Eppley and Goodman 2022; Estrada et al. 2012; McLennan and Hockings 2014). Still,
99 natural forests harbour higher wildlife biodiversity than plantations. For example, many
100 plantation forests have only one or a few tree species per hectare, while natural forests may have
101 ≥ 300 species, with the latter supporting greater faunal diversity (Brockerhoff et al. 2008; Kessler
102 et al. 2005; Onyekwelu et al. 2008). While plantation forests can provide timber and other
103 utilitarian materials, natural forests are often considered critical for ecosystem services which are
104 not effectively met by plantations (Sobuj and Rahman 2011).

105 Whether a species is able to cope with various anthropogenic and natural pressures can
106 potentially be predicted by their dietary guild (Boyle and Smith 2010; Eppley et al. 2020, 2022;
107 Machado et al. 2022). While primates as a whole consume a diverse array of resources, including
108 leaves, fruits/seeds, flowers, gum/sap, bark, and insects (Ibrahim et al. 2023; Lim et al. 2021;
109 Mekonnen et al. 2010; Tesfaye et al. 2021), many species can be narrowly classified as belonging
110 to a specialized dietary guild (Eppley et al. 2020; Hawes and Peres 2014; Kappeler and Heymann
111 1996; Mekonnen et al. 2018b). Dietary specialists typically have anatomical and gastrointestinal
112 specializations that allow them to consume and digest food items that may not be as easily
113 digested by other species (Lambert 1998, 2011). For example, folivorous monkeys often consume
114 young leaves and leaf buds as they have gastrointestinal specializations allowing them to more
115 easily digest leaves compared to frugivorous taxa which often possess shortened, simple digestive
116 tracts (Chapman and Chapman 2002; Cristóbal-Azkarate and Arroyo-Rodríguez 2007; Hanya
117 and Chapman 2013). Beyond a species' dietary guild, however, preferred food resources vary in
118 their nutritional composition and geographic distribution, with habitat quality potentially playing
119 an influential role in a species' feeding ecology (Lambert 2011; Rothman and Bryer 2019).

120 Among the most well-known primate folivores are the colobines, large-bodied monkeys
121 from the subfamily Colobinae that are geographically distributed across sub-Saharan Africa and
122 southern Asia. For instance, in Africa, this taxonomic group is represented by three genera:
123 *Colobus*, *Procolobus*, and *Piliocolobus* (Fashing 2022). Among the most easily recognizable is
124 the eastern black-and-white colobus (*Colobus guereza*), also referred to as the guereza; it is a
125 large-bodied, forest-dependent monkey with a wide, yet patchy, distribution throughout
126 equatorial Africa with, according to most experts, eight subspecies (Fashing and Oates 2013;
127 Zinner et al. 2019). It lives in groups of variable size of up to 23 individuals, and has specialized
128 digestive adaptations to exploit foliage, like other colobines (Chivers 1994; Fashing 2022).
129 Guerezas appear to cope relatively well with low to moderate habitat disturbance, surviving in
130 forest fragments, selectively logged forests, and plantation forests, in addition to natural forest
131 habitats (Fashing et al. 2012; Fashing and Oates 2013; Oates 1977a,b; Onderdonk and Chapman
132 2000). However, the feeding strategies they follow to cope with some of these disturbed and non-
133 native habitats, particularly plantation forests, are not well known (Fashing et al. 2012). Two of
134 the eight subspecies of guerezas are endemic to Ethiopia: the Omo River guereza (*C. g. guereza*)
135 and the Djaffa Mountains guereza (*C. g. gallarum*). Compared to other guereza subspecies, the
136 ecology and behaviour of these Ethiopian taxa are relatively little known, with previous research
137 having been carried out in only a few localities (Dunbar and Dunbar 1974; Dunbar 1987; Tesfaye
138 et al. 2021). The most intensive study of Omo River guerezas to date found that they relied on
139 more whole fruits and flowers, and devoted more feeding time to exotic species, when inhabiting
140 anthropogenically-disturbed forest habitats (forest fragments and disturbed continuous forest)
141 than in large, undisturbed continuous forest habitat (Tefaye et al. 2021).

142 Over the past century, Ethiopia has experienced rapid deforestation, which has resulted in
143 habitat fragmentation, landscape/soil degradation, and biodiversity loss (Fashing et al. 2022;

144 Nyssen et al. 2014). The country's forest cover shrunk from 40% at the beginning of 20th century
145 to 2.4% in 2000 and 1.1% in 2010 (Gebru 2016). Furthermore, from 2010-2020, Ethiopia's
146 annual percentage of forest wood removal (3.0%) was the highest in Africa (FAO 2020). With
147 anthropogenic pressures increasing, and relatively little known about the feeding ecology of the
148 endemic Omo River guereza (though see Tesfaye et al. 2021), it is imperative to obtain broad
149 ecological data for this subspecies, and to evaluate how it is coping in this changing landscape.
150 Accordingly, we aimed to 1) determine dietary preferences of Omo River guerezas in two
151 habitats, i.e., a plantation forest and a natural forest, and 2) determine food availability in these
152 two habitats. We hypothesized that the expected lower plant species diversity in the disturbed
153 plantation forest would force guerezas to feed on a more limited array of plant species than in the
154 more intact natural forest. Based on this, we predicted that plantation forest would have lower
155 food availability than natural forest, and 2) guerezas in plantation forest would have lower dietary
156 diversity than conspecifics in natural forest.

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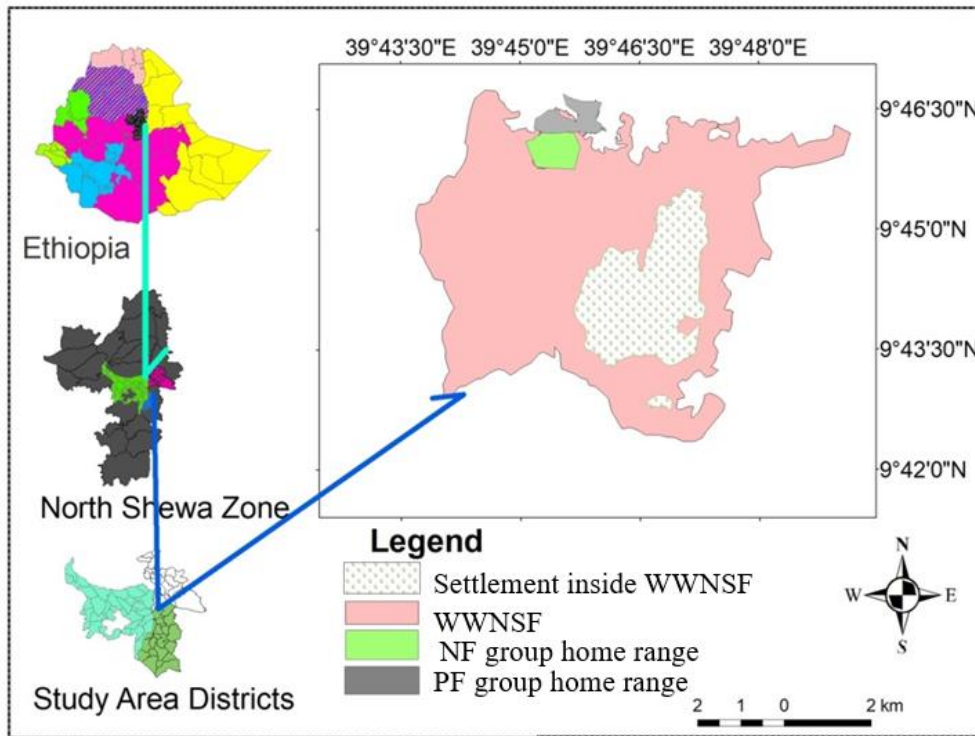
158 **METHODS**

159 **Study area**

160 We conducted this study from May 2015-April 2016 in Wof-Washa Natural State Forest
161 (WWNSF), located in the central highlands of Ethiopia. This protected area is located on a
162 forested escarpment that forms part of the Awash River catchment, which drains into the Danakil
163 plains in the northern section of the Rift Valley (Bekele 1993; Yazezew et al. 2022).
164 Geographically, it extends between 9°42' and 9°47' N latitude and between 39°43' and 39°49' E
165 longitude, situated at elevations between 1650–3700 m asl (Fig. 1). The study area experiences

166 mean annual low and high temperatures of 6.3°C and 22.0° C, respectively, and a typically
167 seasonal rainfall pattern, with mean annual precipitation of 1840 mm.

168



169

170 **Fig. 1** Map of the study area and the home ranges of two study groups (NF=Natural Forest and
171 PF=Plantation Forest) of Omo River guerezas (*Colobus guereza guereza*) at Wof-Washa Natural
172 State Forest, Ethiopia, in 2019.

173

174 WWNSF is home to one of the few remaining dry evergreen Afromontane forests and the
175 oldest natural state forests in the Ethiopian central highlands. It was set aside by the Shewan King
176 Zera Yaqob, one of the country's most important early rulers, as the King's forest (or Crown
177 forest) in the 15th century (ca. 1434-1468). The steep terrain along with the history of royal
178 protection have limited forest access throughout modern times and contributed to its preservation
179 (Veronika 2008). Furthermore, Emperor Menelik II established the first forest policy in Ethiopia
180 and declared Wof-Washa Forest as a State Reserve Forest in the 1880s. Though agricultural

181 encroachment and tree felling for fuel and construction have been problems for centuries, the
182 forest has never been commercially exploited. Unable to exploit the forest for fuelwood, local
183 farmers supplement their private fuelwood with cow dung, which unfortunately decreases its
184 availability for use as fertilizer. Accordingly, local agricultural production is low due to poor soil
185 fertility and unable to meet the population's subsistence needs, a situation which has been
186 exacerbated by recurrent drought (Ayalew 2018; Veronika 2008).

187 There is more natural forest (3,197 ha) than plantation forest (61 ha) at WWNSF (Ayalew
188 2018), with the plantation forest serving as a buffer between the remaining natural forest and
189 local settlements. The plantation forest at WWNSF was established between 1985-2000 by
190 planting exotic tree species, including *Cupressus lucitanica* (Cupressaceae), *Eucalyptus globulus*
191 (Myrtaceae), and *Pinus patula* (Pinaceae), as part of a strategy to rehabilitate degraded areas of
192 natural forest (Ayalew 2018). It thus represents an anthropogenically altered area of forest
193 consisting of a mix of naturally growing indigenous and planted exotic species.

194 **Study groups**

195 We selected two groups of Omo River guerezas, one from a plantation forest and the
196 other from the relatively intact natural forest, for behavioural ecology data collection. The
197 potential caveat of only observing two groups is the direct comparison limits our ability to
198 determine whether any dietary variation is caused by habitat type, other variables, or simply
199 reflects variation between groups. Researchers and trained local field assistants habituated
200 guerezas over three months (February-April 2015) via daily follows. We initially identified the
201 study groups by individual members that had unique pelage markings or other identifiable
202 features. The two groups were of similar size. At the start of the study, the plantation forest group
203 consisted of seven individuals (two adult males, two adult females, one sub-adult male and two
204 juveniles), and this increased to nine individuals after two infants were born in September and

205 October 2015. The natural forest group consisted of six individuals (three adult males, two adult
206 females and one sub-adult male) at the beginning of the study, then increased to eight individuals
207 after two infants were born in October and November 2015. The home ranges of the guereza
208 groups slightly overlapped, with the home range of the plantation forest group being smaller
209 (2.98 ha) than that of the natural forest group (5.40 ha) (Yazew et al. in prep.).

210

211 **Vegetation composition**

212 To characterize the botanical composition and diversity of the home ranges for each
213 group, we conducted vegetation surveys along randomly created transects (Yazew et al. in
214 prep.). Specifically, we created two transects totalling 400-500 m in length and then generated six
215 50 m x 10 m (0.3 ha) vegetation quadrats to systematically sample all plant species present
216 (Teelen 2007). Within each quadrat, we recorded all trees with DBH ≥ 10 cm and all climbers
217 with DBH ≥ 5 cm, and (when possible) identified them to species level. Furthermore, we
218 measured tree height (m), canopy size/diameter (m) (i.e., mean of canopy diameters
219 measurements along two perpendicular axes from 180 and 234 tree measurements in the natural
220 and plantation forests, respectively), and canopy/crown cover (%) (i.e., visual estimation of the
221 level of canopy coverage that obstructs sunlight from reaching the forest floor) (Buchi et al.
222 2018; Gallegos and Glimskär 2009). We identified plant species *in situ*, and collected and
223 preserved (i.e., pressed) specimens of unidentified or questionable taxa. For the latter, we
224 recorded local names and transported specimens to the Addis Ababa University National
225 Herbarium for further taxonomic identification.

226 We used quadrats within the home range of each study group to quantify and characterize
227 the vegetation in each habitat type. We calculated plant species density by dividing the total

228 number of stems recorded per hectare. We calculated plant species diversity of trees ≥ 10 cm
229 DBH using the Shannon-Wiener diversity index (H'), Simpson's Dominance index (D), and the
230 evenness index (J) (Krebs 1999). We used Sørensen indices (Ss coefficient) to assess the
231 similarity in plant species richness between the two home ranges. Ss coefficients range from 0 to
232 1, with 0 representing zero species shared and 1 representing all species shared (Krebs 1999). We
233 calculated the basal area (BA) of each tree species to estimate the biomass of each species in each
234 home range (Fashing 2001; Felton et al. 2008) and determine the dominant tree species in each
235 home range (Kool 1989). We used the Importance Value Index (IVI) to quantify the dominance,
236 occurrence, and abundance of a given plant species in relation to other species in each home
237 range (Kent and Coker 1992). $IVI = RD + RF + RDO$, where RD is relative density, RF is
238 relative frequency, and RDO is relative dominance of the corresponding species i in the group's
239 home range.

240

241 **Phenology**

242 We assessed phenology for selected food plant species in the home ranges of both
243 plantation and natural forest groups monthly during the 12-month study. Based on preliminary
244 observations during the habituation period, we selected and marked the 10 most frequently
245 consumed plant species (trees/shrubs ≥ 10 cm DBH and climbers ≥ 5 cm DBH) abundant (>10
246 individuals/species) in each group's home range. We recorded phenological data from these
247 marked species 1-2 days per month, after we had collected monthly dietary data for each group.
248 We monitored each marked tree for the relative abundance of young leaves, mature leaves,
249 flowers, and whole fruits (Fashing 2001; Ganzhorn et al. 2011; Tesfaye et al. 2021). We assigned
250 each plant food item a relative abundance value (score) that ranged from 0 to 8, in intervals of 1
251 (Mekonnen et al. 2017). An abundance score of zero indicates that a tree showed 0% of its

252 potential abundance (i.e., the item was absent from the plant) during the assessment, while 8
253 indicates an abundance of 87.5-100% (where 100% indicates the plant was fully laden with the
254 item).

255 Based on tree species diversity in both habitats and on our behavioural observations during
256 the habituation phase, we analysed phenological data from 13 botanical species. Specifically, we
257 monitored 10 trees (*Olinia rochetiana*, *Cupressus lusitanica*, *Podocarpus falcatus*, *Allophylus*
258 *abyssinicus*, *Galiniera saxifraga*, *Ilex mitis*, *Juniperus procera*, *Maesa lanceolata*, *Pittosporum*
259 *viridiflorum*, and *Bersama abyssinica*), two shrubs (*Vernonia leopoldi* and *Discopodium*
260 *penninervium*), and one climber (*Embelia schimperi*). We calculated food availability from the
261 mean availability scores of the different food item categories (i.e., young leaves, mature leaves,
262 flowers, and fruits) for each of the 13 marked tree species. Specifically, we calculated the
263 monthly food availability index (FAI) for each food item by multiplying the mean phenology
264 scores of species *i* with the mean basal area of species *i* and density of species *i* per ha (Fashing
265 2001; Mekonnen et al. 2018b; Tesfaye et al. 2021).

266

267 **Feeding ecology**

268 We collected feeding ecology data on each guereza group for five consecutive days per
269 month from 06:00 to 18:00 h. We commenced daily observations at the sleeping site where we
270 left the group on the previous evening. We recorded the activities of individuals using
271 instantaneous scan sampling every 15-minutes (Altmann 1974) with sampling periods of up to 5
272 minutes (Eustace et al. 2015; Fashing 2001; Mekonnen et al. 2018b; Pinheiro and Mendes 2015).
273 We recorded the first behaviour engaged in for ≥ 5 seconds during each scan. We recorded
274 feeding when an individual manipulated food items, including when they obtained the item,
275 moved the item(s) towards their mouth, or masticated it (Eustace et al. 2015; Pinheiro and

276 Mendes 2015). For each feeding scan, we collected data on food species consumed, plant part,
277 and maturity of the item. We recorded plant parts as young leaves, mature leaves, stems, flowers,
278 fruits, shoots, and bark, as well as insects as animal prey. We identified and recorded plant
279 species consumed *in situ* if known, and collected unknown species so that botanists at the Addis
280 Ababa University National Herbarium could identify them later.

281 We collected 22,618 individual behavioural records during 1,268 observation hours
282 (plantation forest group = 650 h; natural forest group= 618 h). We evaluated dietary composition
283 by calculating the daily and monthly proportions of different dietary items and plant species in
284 the feeding scans (Felton et al. 2008; Mekonnen et al. 2010). To determine the dietary preference
285 or selection ratio for specific plant species, we divided the percentage of food items from each
286 species by its percentage density in the transect sample. Ratios above 1 indicate positive selection
287 (Fashing et al. 2014; Dunham 2017; Mekonnen et al. 2018b; Tesfaye et al. 2021). We calculated
288 dietary diversity using the Shannon-Wiener index, H' , and evenness via the evenness index, J
289 (Krebs 1999). We also calculated the percentage overlap in the consumption of each dietary item
290 for each plant species between the two groups (Fashing 2001; Dunham 2017).

291

292 **Statistical Analysis**

293 We compared the diversity indices of food plant species between the two habitats using
294 the Diversity t test. We tested the relationship between the availability indices of plant food items
295 and the percentage of feeding time on the same items using Pearson correlations. We also tested
296 for differences in FAI between habitats using Mann–Whitney U tests. We set statistical
297 significance level at $P \leq 0.05$. We conducted all statistical tests using PAST software version 3.26
298 (Hammer et al. 2001) or SPSS software version 26 (IBM SPSS Inc., Chicago, IL, USA).

299

300 **Ethical note**

301 The Ethiopian Wildlife Conservation Authority and Amhara Region Forest and Wildlife
302 Enterprise granted permission to conduct this research. This project also adhered to the legal
303 requirements of Ethiopia and complied with the American Society of Primatologists' Principles
304 for the Ethical Treatment of Nonhuman Primates.

305

306 **Data availability:** The data sets summarized and analysed for this study are available from the
307 corresponding author on reasonable request.

308

309 **RESULTS**

310 **Habitat description and resource availability**

311 The home range of the plantation forest group contained 12 species from 10 families (8
312 trees, 1 liana/climber, 1 tree/shrub and 2 shrubs), while the home range of the adjacent natural
313 forest group had 21 species from 19 families (12 trees, 6 shrubs and 3 tree/shrubs). Six plant
314 species were common in the home range of the natural forest group but did not occur in the
315 plantation forest group's home range (Table S1). The home range of the natural forest group had
316 a higher stem density than the plantation forest group's home range (Table 1).

317 The plant species similarity index in the two home ranges was moderate (9 of 24 species;
318 Sørensen S_s coefficient 0.55). The vegetation in the home range of the natural forest group was
319 more diverse than in that of the plantation forest group (Table 1). The Shannon-Wiener diversity
320 index (PAST diversity t test: $t = -7.1$, $df = 23$, $p = 0.019$) and Simpson's diversity index ($t = 6.3$,

321 df = 23, p = 0.013) were significantly higher in the home range of the natural forest group than in
 322 that of the plantation forest group (Table 1 and Table S1). Species evenness was higher and
 323 dominance was lower within the home range of the natural forest group than in the home range of
 324 the plantation forest group (Table 1 and Table S1).

325

326 **Table 1.** Overview of vegetation characteristics in the home ranges of Omo River guereza groups
 327 inhabiting plantation and natural forests in Wof-Washa Natural State Forest, Ethiopia, from May
 328 2015-April 2016.

Habitat variables	Plantation forest	Natural forest
Taxa recorded	12	21
Large tree (≥ 10 cm DBH) stem density per ha	600.0	780.0
Large tree (≥ 10 cm DBH) species richness	40.0	70.0
Shannon-Wiener	1.60	2.30
Simpson's species diversity index	0.70	0.90
Large tree (≥ 10 cm DBH) species evenness	0.64	0.77
Large tree (≥ 10 cm DBH) species dominance index	0.29	0.12

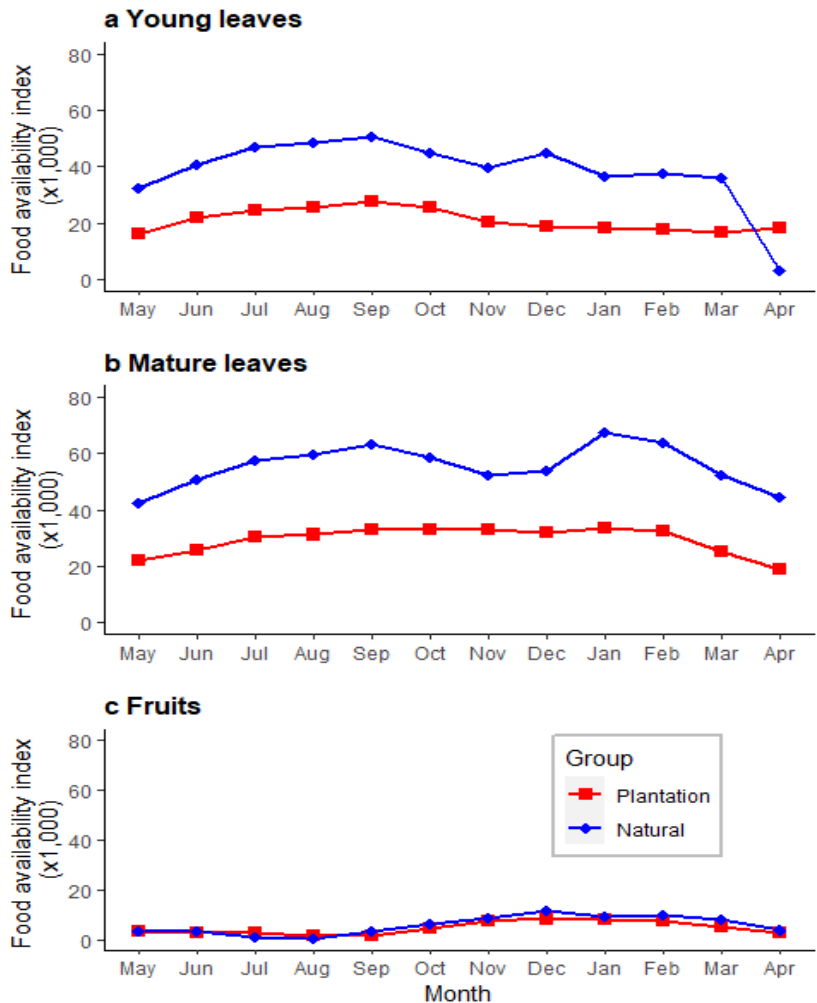
329

330 *Maesa lanceolata*, *Juniperus procera*, *Podocarpus falcatus*, and *Cupressus lusitanica*
 331 were the most dominant species in the home range of the plantation forest group, while *Olinia*
 332 *rochetiana*, *Erica arborea*, *Allophylus abyssinicus*, *J. procera*, and *M. lanceolata* were the most
 333 dominant species in the home range of the natural forest group (Table S2).

334

335 **Phenology**

336 Plant food item availability varied over time (Fig. 2). Young, and to a lesser extent mature
 337 leaves, were the most abundant items throughout the year, while fruit was more seasonal. The
 338 monthly food availability indices for young leaves



339

340 **Fig. 2** Monthly food availability indices (young leaves, mature leaves and fruits) for plantation
 341 and natural forest groups of Omo River guerezas in Wof-Washa Natural State Forest, Ethiopia,
 342 from May 2015-April 2016.

343
 344 (Mann Whitney U; $Z = -4.16$, $P < 0.001$) and mature leaves ($Z = -4.15$, $P < 0.001$) were

345 statistically significantly higher in the home range of the natural forest group than in that of the
 346 plantation forest group. However, there was no statistically significant difference in fruit FAI

347 between the home ranges of the two groups (Mann Whitney U; $Z = 0.98$, $P = 0.350$). Fruit was at

348 peak availability during December in both groups' home ranges. The overall annual young leaf

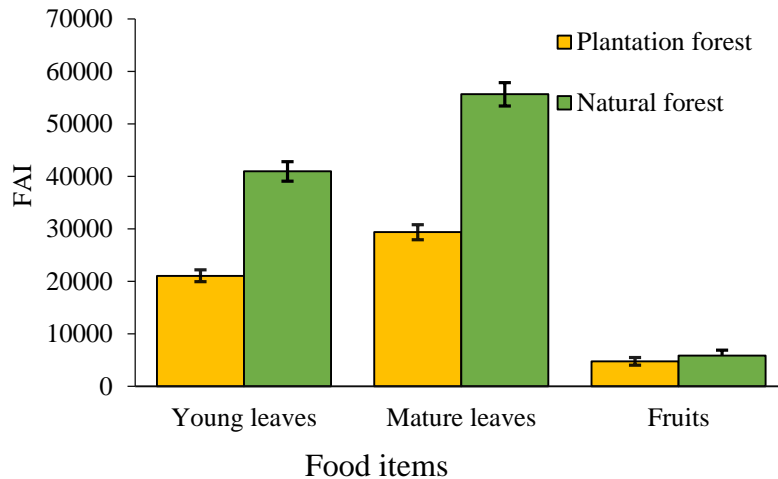
349 availability per hectare was 32% higher in the home range of the natural forest group than in that

350 of the plantation forest group, and mature leaf availability per hectare was 30% higher. Moreover,

351 the overall annual availability of fruit was 10% higher in the home range of the natural forest
352 group than in that of the plantation forest group (Fig. 3).

353

354



355

356 **Fig. 3** Comparison of total food availability index (FAI) values of food items in the home ranges
357 of Omo River guereza groups in plantation and natural forest (mean \pm SE) in Wof-Washa Natural
358 State Forest, Ethiopia, from May 2015-April 2016.

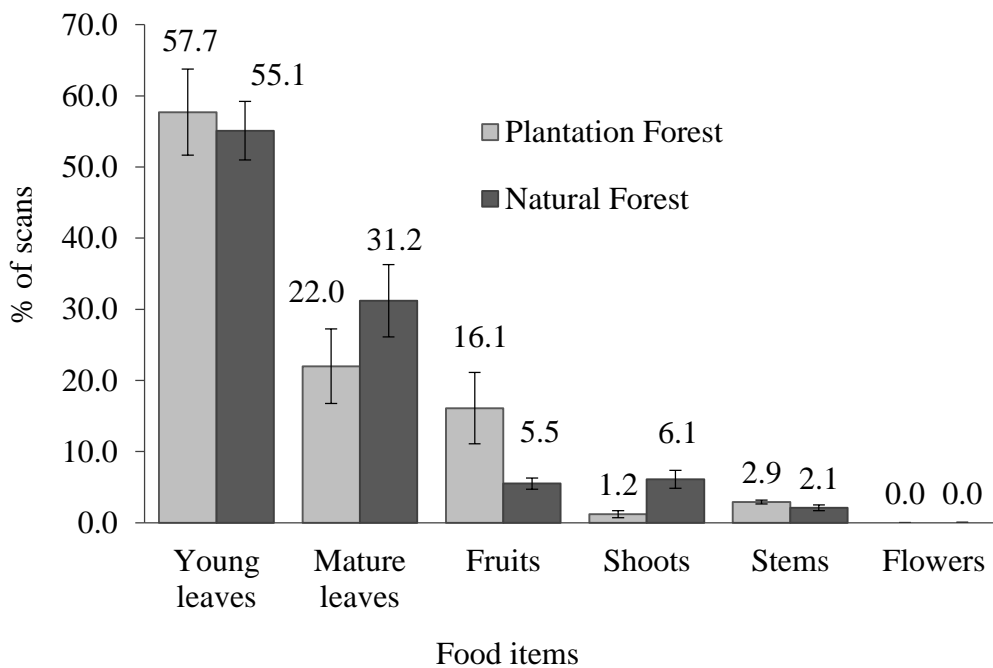
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360 *Juniperus procera* contributed the highest abundance of both young and mature leaves in
361 the home ranges of both groups, although its consumption rank was only 6th in plantation forest
362 and 9th in natural forest (Table 2). Young leaves and mature leaves of *Maesa lanceolata* were the
363 second most abundant foods in the home ranges of both groups and the species ranked first in
364 plantation forest and second in natural forest in consumption (Table 2). *Vernonia leopoldi* was
365 not an abundant item in either group's home range even though it accounted for the highest
366 percentage of feeding time scans in the natural forest group and ranked fourth in the diet of the
367 plantation forest group.

368

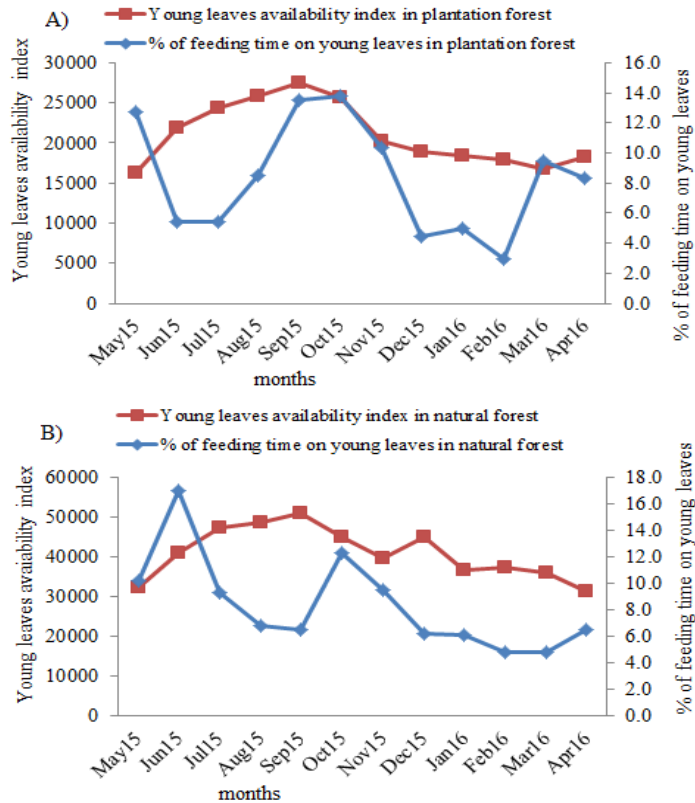
369 **Food item consumption**

370 Overall, leaves accounted for most of the guereza diet in both the plantation forest (80%)
 371 and natural forest (86%) groups. There were no significant differences between groups in the
 372 monthly consumption of young leaves ($t = 0.5$, $df = 11$, $p = 0.644$), mature leaves ($t = 1.5$, $df =$
 373 11 , $p = 0.151$), and stems ($t = 2$, $df = 11$, $p = 0.056$) (Fig. 4). However, the plantation forest group
 374 spent significantly more time monthly feeding on fruits (16.1% vs. 5.5%; $t = 2.2$, $df = 11$, $p =$
 375 0.037) and less time on shoots (1.2% vs. 6.1%; $t = 3.6$, $df = 11$, $p = 0.001$) than the natural forest
 376 group.



377
 378 **Fig. 4** Annual percentage of plant part contribution to the diets of two groups of Omo River
 379 guerezas (mean \pm SE) inhabiting different forest types in Wof-Washa Natural State Forest,
 380 Ethiopia, from May 2015-April 2016.

381
 382 Food availability indices of young leaves in phenology tree species and total feeding time
 383 scans on young leaves (for all phenology plant species in each group's range) were significantly
 384 correlated for the natural forest group ($r=0.63$, $P = 0.004$; Fig. 5B) but not for the plantation
 385 forest group ($r=0.33$, $P = 0.06$; Fig. 5A).



386

387 **Fig. 5** Comparison of the availability of young leaves and their consumption by Omo River
 388 guerezas in (A) plantation forest and (B) natural forest in Wof-Washa Natural State Forest,
 389 Ethiopia, from May 2015-April 2016.

390 *Ilex mitis* was the most selected for plant species by the plantation forest group, followed by
 391 *Vernonia leopoldi*, and *Ficus sur* (Table 2). In contrast, *V. leopoldi*, *Ilex mitis* and *Pittosporum*
 392 *viridiflorum* were the top three most selected for plant food species by the natural forest group.
 393 *Maesa lanceolata* had a low selection ratio for both groups despite having the highest percentage
 394 contribution to the overall diet of the plantation forest group and the second highest contribution
 395 to the diet of the natural forest group.

396 **Table 2.** Dietary selection ratios based on stem density (individuals/ ha) and percentage of time spent feeding by groups of Omo River guerezas
 397 in plantation and natural forest habitats, in Wof-Washa Natural State Forest, Ethiopia, from May 2015-April 2016.

Plantation Forest					Natural Forest				
Rank	Species consumed	Annual diet %	% of stem density	Selection Ratio (rank)	Rank	Species consumed	Annual diet %	% of stem density	Selection Ratio (Rank)
1	<i>Maesa lanceolata</i>	30.6	44.4	0.7 (8)	1	<i>Vernonia leopoldi</i>	29.8	1.7	17.4 (1)
2	<i>Podocarpus falcatus</i>	16.1	8.3	1.9 (7)	2	<i>Maesa lanceolata</i>	15.6	12.8	1.2 (10)
3	<i>Ilex mitis</i>	14.1	1.1	12.8 (1)	3	<i>Pittosporum viridiflorum</i>	12.0	2.1	5.6 (4)
4	<i>Vernonia leopoldi</i>	13.5	1.7	7.9 (2)	4	<i>Olinia rochetiana</i>	7.3	19.2	0.4 (13)
5	<i>Discopodium penninervium</i>	8.6	2.2	3.9 (4)	5	<i>Ilex mitis</i>	5.7	0.4	13.3 (2)
6	<i>Juniperus procera</i>	3.9	27.8	0.1 (10)	6	<i>Galiniara saxifraga</i>	4.9	2.6	1.9 (6)
7	<i>Olinia rochetiana</i>	3.7	1.1	3.4 (5)	7	<i>Discopodium penninervium</i>	4.5	3.4	1.3 (9)
8	<i>Ficus Sur</i>	3.5	0.6	5.8 (3)	8	<i>Allophylus abyssinicus</i>	3.8	14.5	0.3 (14)
9	<i>Bersama abyssinica</i>	2.9	1.1	2.6 (6)	9	<i>Juniperus procera</i>	3.4	14.5	0.2 (15)
10	<i>Cupressus lusitanica</i>	1.5	8.3	0.2 (9)	10	<i>Cupressus lusitanica</i>	2.5	0.4	6.3(3)
					11	<i>Ficus sur</i>	1.9	0.6	3.2 (5)
					12	<i>Olea europaea</i>	1.7	2.6	0.7 (11)
					13	<i>Bersama abyssinica</i>	1.5	0.9	1.8 (7)
					14	<i>Halleria lucida</i>	1.4	0.9	1.6 (8)
					15	<i>Podocarpus falcatus</i>	1.0	1.7	0.6 (12)

398 *Note:* SR=Selection Ratio is equals the percentage of a specific species in the diet divided by the availability percentage of that species (% of
 399 stem density) along sampled transects enumerated.

400

401 **Dietary species richness and diversity**

402 The two groups of guerezas consumed a total of 31 food plant species. The natural
 403 forest group obtained food from 30 species, while the plantation forest group obtained food
 404 from 18 species (Tables S3 and S4). The five most consumed species accounted for 83% of
 405 the overall diet of the plantation forest group, and 70% of the overall diet of the natural forest
 406 group. Ten plant species each accounted for $\geq 1.0\%$ of the overall annual diet for the
 407 plantation forest group (Table 3) whereas 15 species each accounted $\geq 1.0\%$ of the overall
 408 annual diet for the natural forest group (Table 4). *Maesa lanceolata* was the top food species
 409 for the plantation forest group (30.65% of the diet) and the second most consumed species for
 410 the natural forest group (15.64%). *Vernonia leopoldi* was the most consumed species by the
 411 natural forest group (29.77%) and the fourth most consumed species by the plantation forest
 412 group (13.55%).

413

414 **Table 3.** Percentage contribution of food items from the top 10 plant species in the diet of
 415 Omo River guerezas in plantation forest in Wof-Washa Natural State Forest, Ethiopia, from
 416 May 2015-April 2016.

Family	Species	*Growth form	Young leaves	Mature leaves	Fruit	Shoot	Stem	Total contribution
Myrsinaceae	<i>Maesa lanceolata</i>	T	20.51	8.48	0.05	0.55	1.06	30.65
Podocarpaceae	<i>Podocarpus falcatus</i>	T	3.61	0.38	12.06	0.08	-	16.12
Aquifoliaceae	<i>Ilex mitis</i>	T	10.62	2.77	-	0.28	0.45	14.13
Asteraceae	<i>Vernonia leopoldi</i>	S	8.98	3.96	-	0.13	0.48	13.55
Solanaceae	<i>Discopodium penninervium</i>	S	5.17	2.83	-	-	0.63	8.63
Cupressaceae	<i>Juniperus procera</i>	T	2.17	1.14	0.48	0.10	0.03	3.91
Oliniaceae	<i>Olinia rochetiana</i>	T	2.55	0.98	-	0.05	0.13	3.71
Moraceae	<i>Ficus sur</i>	T	0.03	-	3.51	-	-	3.53
Melanthaceae	<i>Bersama abyssinica</i>	T/S	1.79	0.96	-	-	0.10	2.85
Cupressaceae	<i>Cupressus lusitanica</i>	T	1.11	0.40	-	0.03	-	1.54

417 Note: * T= Tree, T/S= Tree /Shrub, S= Shrub,

418

419

420

421 **Table 4.** Percentage contribution of food items from the top 15 plant species consumed by
 422 Omo River guerezas in natural forest in Wof-Washa Natural State Forest, Ethiopia, from May
 423 2015-April 2016.

Family	Species	*Growth form	Young leaves	Mature leaves	Fruit	Shoot	Stem	Total contribution
Asteraceae	<i>Vernonia leopoldi</i>	S	16.05	11.83	0.19	0.90	0.81	29.77
Myrsinaceae	<i>Maesa lanceolata</i>	T	8.40	4.71	0.99	1.15	0.40	15.64
Pittosporaceae	<i>Pittosporum viridiflorum</i>	T	7.13	2.94	1.18	0.59	0.15	11.99
Oliniaceae	<i>Olinia rochetiana</i>	T	4.46	2.11	0.12	0.46	0.12	7.28
Aquifoliaceae	<i>Ilex mitis</i>	T	3.00	2.29	-	0.37	-	5.67
Rubiaceae	<i>Galiniera saxifraga</i>	T	2.79	1.24	0.50	0.31	0.09	4.93
Solanaceae	<i>Discopodium penninervium</i>	S	2.04	1.49	0.19	0.46	0.28	4.46
Sapindaceae	<i>Allophylus abyssinicus</i>	T	2.51	0.90	0.09	0.22	0.03	3.75
Cupressaceae	<i>Juniperus procera</i>	T	2.29	0.65	0.25	0.15	0.03	3.38
Cupressaceae	<i>Cupressus lusitanica</i>	T	1.08	0.87	-	0.59	-	2.54
Moraceae	<i>Ficus sur</i>	T	-	-	1.89	-	-	1.89
Oleaceae	<i>Olea europaea</i>	T	1.02	0.53	0.03	0.12	-	1.70
Melanthaceae	<i>Bersama abyssinica</i>	T/S	0.81	0.37	0.06	0.22	0.06	1.52
Scrophulariaceae	<i>Halleria lucida</i>	T/S	0.59	0.34	-	0.34	0.09	1.36
Podocarpaceae	<i>Podocarpus falcatus</i>	T	0.84	0.09	-	-	0.03	0.96

424 Note: * T= Tree, T/S= Tree /Shrub, S= Shrub.

425

426

427 Overall, about half of the food items from the top nine plant species were consumed

428 by both guereza groups. The overlap was highest for young leaves, followed by mature leaves

429 (Table 5).

430

431 **Table 5.** Percentage overlap in species-specific food item (n=9 species) between two Omo
 432 River guereza groups in Wof-Washa Natural State Forest, Ethiopia, from May 2015-April
 433 2016.

Species	Young leaves	Mature leaves	Fruit	Stems	Shoots	Total
<i>Maesa lanceolata</i>	8.4	4.7	0.1	0.8	0.6	14.6
<i>Vernonia leopoldi</i>	9.0	4.0	0.0	0.5	0.1	13.6

<i>Ilex mitis</i>	3.0	2.3	0.0	0.0	0.3	5.6
<i>Discopodium penninervium</i>	2.0	1.5	0.0	0.3	0.0	3.8
<i>Olinia rochetiana</i>	2.6	1.0	0.0	0.1	0.1	3.8
<i>Juniperus procera</i>	2.2	0.7	0.3	0.0	0.1	3.3
<i>Ficus sur</i>	0.0	0.0	1.9	0.0	0.0	1.9
<i>Cupressus lusitanica</i>	1.1	0.4	0.0	0.0	0.0	1.5
<i>Bersama abyssinica</i>	0.8	0.4	0.0	0.1	0.0	1.3
Total	29.1	15.0	2.3	1.8	1.2	49.4

434

435 The mean monthly diversity of food species was significantly lower in the plantation
436 forest group than the natural forest group (Mann Whitney U; Z = -2.3, P = 0.019; Table 6).
437 Dietary diversity was highest in September and October and lowest in February for the
438 plantation forest group, while it was highest in June and lowest in December for the natural
439 forest group (Table 6). The mean monthly dietary evenness was also significantly lower for
440 guereza groups inhabiting the plantation forest than for those in the natural forest (Mann
441 Whitney U; Z = -4.2, P < 0.001; Table 6).

442

443 **Table 6.** Food species diversity and evenness indices of guerezas over 12 months study period
444 in plantation and natural forest in Wof-Washa Natural State Forest, Ethiopia, from May 2015-
445 April 2016.

Month	Shannon-Wiener diversity index, H'			Evenness index, J		
	plantation forest	natural forest	mean	plantation forest	natural forest	Mean
May15	1.77	2.44	2.11	0.36	0.44	0.40
Jun15	1.60	2.54	2.07	0.37	0.42	0.39
Jul15	1.66	2.31	1.99	0.40	0.48	0.44
Aug15	1.66	1.84	1.75	0.40	0.48	0.44
Sep15	2.19	2.40	2.30	0.36	0.49	0.43
Oct15	2.19	2.45	2.32	0.36	0.43	0.40
Nov15	2.00	2.00	2.00	0.37	0.44	0.41
Dec15	1.76	1.46	1.61	0.39	0.47	0.43
Jan16	1.85	1.98	1.92	0.36	0.46	0.41
Feb16	1.18	1.95	1.57	0.41	0.48	0.44
Mar16	1.50	1.68	1.59	0.38	0.46	0.42
Apr16	1.58	2.05	1.82	0.40	0.45	0.42
Mean	1.75	2.09	1.92	0.38	0.46	0.42

446 **DISCUSSION**

447 In our comparative study of Omo River guereza groups in natural forest versus
448 plantation forest at WWNSF, we found that plant species richness and overall food
449 availability were lower in the home range of the group inhabiting plantation forest than in the
450 group inhabiting natural forest. The natural forest group often consumed food items from
451 large tree species like *Pittosporium viridiflorum*, *Galiniera saxifraga*, and *Allophylus*
452 *abyssinicus*, which did not occur in the plantation forest and were thus unavailable to the
453 guerezas living there. Although plantations are generally less suitable habitats for many
454 primates and other animals than natural forest habitats (Brockerhoff et al. 2008; Fashing et al.
455 2012; Merker and Yustian 2008), guereza groups nevertheless survived in both environments
456 at WWNSF, consistent with previous studies showing that this species can persist in a variety
457 of degraded and human-modified habitats (Fashing 2012; Chapman et al. 2000; Tesfaye et al
458 2021; Oates 1977b; Wasserman and Chapman 2003). In the plantation forest, tree species
459 such as *Juniperus procera*, *Podocarpus falcatus*, *Olinia rochetiana*, *Maesa lanceolata*, *Ilex*
460 *mitis*, and the exotic *Cupressus lusitanica* provide benefits to local people as timber and to
461 guerezas as habitat and food sources (Gerard et al. 2015; Grimes and Paterson 2000;
462 Konersmann et al. 2021). Like guereza, several other forest primate species, including slow
463 lorises (*Nycticebus bengalensis*), southern bamboo lemurs (*Hapalemur meridionalis*), tarsiers
464 (*Tarsius diana*), howler monkeys (*Alouatta* spp.), and siamangs (*Hylobates syndactylus*), are
465 also capable of inhabiting plantations, in some instances containing high densities of non-
466 native trees (Eppley et al. 2015; Merker and Yustian 2008; Nowak and Lee 2013;
467 Pliosungnoen et al. 2010).

468 Large tree stem density, species diversity, and species richness were all higher in the
469 home range of the natural forest group than in that of the plantation forest group where a
470 relatively small number of species accounted for most of the trees. These differences likely

471 stem from the plantation forest's history as a heavily degraded area of natural forest to which
472 several exotic species were added several decades ago (Ayalew 2018). In western Kenya,
473 even forests established by planting a variety indigenous species over a half century earlier
474 are known to not entirely mirror the tree composition of older natural forest nearby and
475 contain lower densities of several monkey species, including guereza, than the natural forest
476 (Fashing et al. 2012).

477 Seasonal changes in resource abundance and availability have fundamental effects on
478 the behaviour and ecology of primates (Dunbar 1988). Guerezas in both forest types at Wof-
479 Washa proved to be highly folivorous and their diet closely followed local resource
480 phenology patterns. Guereza feeding time on their preferred resource (i.e., young leaves) was
481 strongly influenced by temporal variation in their abundance and availability. Food abundance
482 was significantly higher in the natural forest habitat than in the plantation forest, which may
483 largely be due to the legacy of intensive human encroachment and habitat degradation in the
484 plantation forest habitat (Ayalew 2018). Such factors reduce plant species richness, diversity,
485 and structure, and are known to negatively affect many primate species (Boyle et al. 2012;
486 Eppley et al. 2020). Although the guerezas spent most of their time feeding on *Maesa*
487 *lanceolata*, *Vernonia leopoldi*, *Ilex mitis*, *Podocarpus falcatus*, *Discopodium penninervium*
488 and/or *Pittosporum viridiflorum*, when the abundance of favoured food items on these plant
489 species was reduced, guerezas fed more on fallback species (Marshall et al. 2009) such as
490 *Juniperus procera* and *Allophylus abyssinica*. Some of these desirable plant species for
491 guerezas (specifically, *M. lanceolata*, *J. procera*, *C. lusitanica*, and *P. falcatus*) had high
492 importance value indices within plantation forest habitat, revealing that these tree species
493 have been relatively resistant to the various anthropogenic pressures imposed by local
494 communities (Ayalew et al. 2015).

495 Overall, we identified a total of 31 plant species as guereza foods in WWNSF,
496 comprising 31 genera and 26 families. Comparable values have been reported in several
497 studies of other *Colobus guereza* subspecies (Bocian 1997; Fashing 2001; Oates 1977a), as
498 well as in other *C. g. guereza* populations in southern Ethiopia (Tefaye et al. 2021). These
499 low dietary species richness values seem to be characteristic of guerezas, including for
500 populations inhabiting species-rich rain forest habitats (Table 7). While they typically exhibit
501 a preference for young leaves (Oates 1977a; Tefaye et al. 2021), guerezas feed on other food
502 items when preferred resources are scarce, and there are sites where they seem to prefer fruits
503 when available (Fashing 2001; Fashing et al. 2007; Oates 1977a; Plumptre 2006). This
504 ecological flexibility to expand their diet when under pressure can be considered an asset for
505 African colobines coping with the resource scarcity resulting from anthropogenic disturbance,
506 including populations in forest fragments (Chapman and Chapman 1999; Eppley et al. 2017;
507 Tefaye et al. 2021) and plantation forests (this study).

508 .

509 **Table 7I.** The diet of black-and-white colobus monkeys, *Colobus* spp., across their range in Africa.

Study species	Study site	Food items									Sampling type	References
		YL	ML	UL	TotL	FR	FL	SH	ST	#sp p		
<i>C. guereza guereza</i> (plantation forest)	WWNSF, Ethiopia	57. 7	22	-	79.7	16.1		1.2	2. 9	18	Scan	This study
<i>C. g. guereza</i> (natural forest)	WWNSF, Ethiopia	55. 1	31. 2	-	86.3	5.5		6.1	2. 1	30	Scan	This study
<i>C. g. guereza</i>	Aregash Forest, Ethiopia	51	11	0	62	20	5	-	-	37	Scan	Tesfaye et al. (2021)
<i>C. g. guereza</i>	Munessa, Ethiopia	57	26	0	83	3	1	-	-	27	Scan	Tesfaye et al. (2021)
<i>C. g. guereza</i>	Wondo Genet, Ethiopia	50	15	0	65	6	13	-	-	32	Scan	Tesfaye et al. (2021)
<i>C. g. occidentalis</i>	Kalinzu, Uganda	87	0	0	87	5	1	-	-	39	Scan	Matsuda et al. (2020)
<i>C. g. occidentalis</i>	Kibale, Uganda	65	14	1	80	12	6	-	-	-	Scan	Wasserman and Chapman (2003)
<i>C. g. occidentalis</i>	Kibale Forest, Uganda	57. 7	12. 4	2.5	72.6	13.6	2.1	-	-	43	Scan	Oates (1977a)
<i>C. g. occidentalis</i>	Ituri Forest, DRC	26. 2	3.8	24.2	54.2	24.6	2.9	-	-	31	Scan	Bocian (1997)
<i>C. g. matschiei</i>	Kakamega Forest, Kenya	20. 4	6.6	22.5	49.5	38.6	0.5	-	-	28+	Scan	Fashing (2001)
<i>C. angolensis cottoni</i>	Ituri Forest, DRC	23. 5	2.4	22	47.9	5.4	7.2	-	-	37	Scan	Bocian (1997)
<i>C. a. palliatus</i>	Diani Forest, Kenya	58	13	0	71	5	14	-	-	110	Scan	Dunham (2017)

<i>C. a. palliatus</i>	Diani Forest, Kenya	21. 1	35. 4	4.1	60.6	9.5	27. 6	-	-	42	Scan	Jansson (2011)
<i>C. polykomos</i>	Tiwai, Sierra Leone	29. 7	26. 4	-	56.1	34.6	1.7	-	-	56	Scan	Dasilva (1994)

510 YL = young leaves; ML = mature leaves; UL = unclassified leaves; TotL= total leaves; FR = fruit; FL = flowers; SH = shoots; ST = stems; # spp.

511 = number of species consumed.

512

513 Primates often select foods based on their accessibility and availability throughout the
514 year, as well as their nutritional content (Clink et al. 2017; Eppley et al. 2017; Fashing et al.
515 2007; Lambert and Rothman 2015). It is likely that guerezas in WWNSF selected food species
516 based on a combination of these factors. Young leaves comprised a significant portion of the
517 diet in both groups, similar to what has been reported for other *Colobus guereza* populations,
518 including *C. g. guereza* inhabiting continuous and fragmented forests in southern Ethiopia
519 (Bocian 1997; Oates 1977a; Tesfaye et al. 2021), a strategy thought to meet their nutritional
520 requirements by ensuring high protein intake (Dasilva 1994; Ganzhorn et al. 2017). However,
521 while young leaves tend to contain higher concentrations of protein, their overall nutritional
522 quality can be highly variable (Ganzhorn et al. 2017; Ryan et al. 2013).

523 The mean monthly H' values of the two groups were similar to those in previous studies
524 of other black-and-white colobus monkey species, such as *Colobus guereza occidentalis* ($H' =$
525 1.9) and *Colobus angolensis cottoni* ($H' = 1.8$) in Ituri, D.R. Congo (Bocian 1997) and *C. g.*
526 *matschiei* (T-group, $H' = 1.6$; and O group, $H' = 1.7$) in Kakamega, Kenya (Fashing 2001).
527 However, the mean food species evenness indices for both groups (plantation forest group, $J =$
528 0.38; natural forest group, $J = 0.46$) were considerably lower than those reported for *C. g.*
529 *matschiei* in Kakamega (T-group, $J = 0.71$; O-group, $J = 0.72$; Fashing 2001) and *C. g.*
530 *guereza* in southern Ethiopia ($J = 0.85$; Tesfaye et al. 2021). This difference might be due to
531 the impact of deforestation in WWNSF, leading to the decline of some food tree species and
532 dominance by others. If this is the case, reducing anthropogenic pressures on the forest is a
533 critical part of an effective conservation management strategy at WWNSF. Indeed, previous
534 research has shown that the basal area of big trees in WWNSF declined from 100.3 m²/ha in
535 1993 to 64.32 m²/ha in 2013 (Fisaha et al. 2013). This decline was likely due to selective
536 logging for timber and other construction purposes resulting in the dominance of secondary
537 vegetation, such as small-sized trees and shrubs, especially in the plantation forest and close

538 to villages. These large trees are often food resources for guerezas, thus the continuation of
539 this selective logging and progressive changes in resource abundance and availability will
540 likely adversely affect guerezas, as well as other wildlife dependent on this habitat (Dunbar,
541 1988; Kamilar and Paciulli, 2008; Konersmann et al. 2021). For this reason, it will be
542 important to create awareness campaigns on sustainable forest use and management targeting
543 resource-dependent community members. There is a growing awareness that in many cases,
544 including in Ethiopia, community participation in natural resource management can be more
545 effective than traditionally strict protected areas (Ashenafi and Leader-Williams 2005; Estrada
546 et al. 2022; Fashing et al. 2022). Facilitating critical reflection about livelihood priorities
547 through participatory conservation approaches can effectively align community actions with
548 natural resource management strategies that may improve conservation outcomes (Eppley et
549 al. 2023; Wali et al. 2017).

550

551 **Conclusion**

552 While our results on feeding ecology show that Omo River guerezas at Wof-Washa
553 were able to survive on the resources available largely in either natural or plantation forest,
554 they also highlight the critical need for further studies of the dietary and habitat preferences of
555 this subspecies so we can understand how it may respond to future climatic and anthropogenic
556 pressures. Considering the increasing anthropogenic pressures and habitat degradation
557 throughout the tropics, the ability of Omo River guerezas to persist in the less botanically
558 diverse plantation forest habitat must be viewed as a positive. In terms of conservation
559 strategies, plantation forests can provide effective (and critical) habitat buffer zones and
560 corridors, allowing for genetic connectedness across the landscape. We strongly encourage
561 community-led efforts aimed at reducing deforestation while increasing landscape-level

562 reforestation, integrating both primate food and utilitarian tree species (Konersmann et al.
563 2021; Mekonnen et al. 2022).

564

565 REFERENCES

- 566 Altmann, J. (1974). Observational study of behavior: sampling methods. *Behaviour*, 49, 227–
567 267.
- 568 Ashenafi, Z., & Leader-Williams, N. (2005) Indigenous common property resource
569 management in the central highlands of Ethiopia. *Human Ecology*, 33, 539–563.
- 570 Ayalew, A.T. (2018). Vegetation ecology and carbon stock of Wof-Washa Forest, North
571 Shewa Zone, Amhara Region, Ethiopia. PhD thesis, Addis Ababa University, Addis
572 Ababa, Ethiopia.
- 573 Ayalew, A. T., Soromessa, T., & Kelbessa, E. (2015). Structure and regeneration status of
574 Menagesha Amba Mariam Forest in central highlands of Shewa, Ethiopia. *Agriculture,
575 Forestry and Fisheries*, 4(4), 184–194.
- 576 Bekele, T. (1993). *Vegetation Ecology of Renunant Afromontane Forests on the Central
577 Plateau of Shewa, Ethiopia*. Uppsala University, Uppsala, Sweden.
- 578 Bocian, C. (1997). *Niche separation of black-and-white colobus monkeys (Colobus angolensis
579 and C. guereza) in the Ituri Forest*. Ph.D. dissertation, City University of New York,
580 New York.
- 581 Boyle, S., & Smith, A. (2010). Can landscape and species characteristics predict primate
582 presence in forest fragments in the Brazilian Amazon? *Biological Conservation*, 143,
583 1134–1143.
- 584 Boyle, S., Zartman, C., Spironello, W., & Smith, A. (2012). Implications of habitat
585 fragmentation on the diet of bearded saki monkeys in central Amazonian forest.
586 *Journal of Mammalogy*, 93, 959–976.
- 587 Brockerhoff , E. G, Jactel, H., Parrotta, J. A., Quine, C. P., & Sayer, J. (2008). Plantation
588 forests and biodiversity: oxymoron or opportunity? *Biodiversity and Conservation*, 17,
589 925–951.
- 590 Buchi, L., Wendling, M., Mouly, P., & Charles, R. (2018). Comparison of Visual Assessment
591 and Digital Image Analysis for Canopy Cover Estimation. *Agronomy Journal*, 110,
592 1289–1295. doi:10.2134/agronj2017.11.0679.
- 593 Chapman, C.A ., & Chapman, L. J. (1999). Implications of small scale variation in ecological
594 conditions for the diet and density of red colobus monkeys. *Primates*, 40, 215–231.
- 595 Chapman, C., & Chapman, L. J. (2002). Foraging challenges of red colobus monkeys: inatural
596 forest luence of nutrients and secondary compounds. *Comparative Biochemistry and
597 Physiology Part A: Molecular & Integrative Physiology*, 133(3), 861–875.
- 598 Chapman, C.A., Balcomb, S.R., Gillespie, T.R., Skorupa, J.P., & Struhsaker, T.T.
599 (2000). Long-term effects of logging on African primate communities: a 28-
600 year comparison from Kibale National Park, Uganda. *Conservation Biology*, 14,
601 207–217.
- 602 Chivers, D. J. (1994). Functional anatomy of the gastrointestinal tract. In A. G. Davies & J. F.
603 Oates (Eds.), *Colobine monkeys: Their ecology, behavior and evolution* (pp. 205–227).
604 Cambridge: Cambridge University Press.

605 Clink, D. J., Dillis, C., Feilen, K. L., Beaudrot, L., & Marshall, A. J. (2017). Dietary diversity,
606 feeding selectivity, and responses to fruit scarcity of two sympatric Bornean primates
607 *Hylobates albibarbis* and *Presbytis rubicunda rubida*). *PLoS ONE*, 12(3), e0173369.

608 Cristóbal-Azkarate, J., and Arroyo-Rodríguez, V. (2007). Diet and activity pattern of howler
609 monkeys (*Alouatta palliata*) in Los Tuxtlas, Mexico: Effects of habitat fragmentation
610 and implications for conservation. *American Journal of Primatology*, 69, 1013–1029.

611 Dasilva, G. L. (1994). Diet of *Colobus polykomos* on Tiwai Island: Selection of food in
612 relation to its seasonal abundance and nutritional quality. *International Journal of*
613 *Primatology*, 15(5), 655–680.

614 Dunbar, R. (1987). Habitat quality, population dynamics, and group composition in
615 colobus monkeys (*Colobus guereza*). *International Journal of Primatology*, 8,
616 299–329.

617 Dunbar, R. (1988). *Primate Social Systems: Studies in Behavioural Adaptation*. Springer
618 Science & Business Media.

619 Dunbar, R., & Dunbar, E. (1974). Ecology and population dynamics of *Colobus*
620 *guereza* in Ethiopia. *Folia Primatologica*, 21, 188–208.

621 Dunham, N.T. (2017). Feeding ecology and dietary flexibility of *Colobus angolensis palliatus*
622 in relation to habitat disturbance. *International Journal of Primatology*, 38, 553–571.

623 Eppley, T. M., Donati, G., Ramanamanjato, J-B., Randriatafika, F., Andriamandimbiarisoa,
624 L.N., Rabehevitra, D., et al. (2015). The use of an invasive species habitat by a small
625 folivorous primate: implications for lemur conservation in Madagascar. *PLoS ONE*
626 10(11): e0140981. doi:10.1371/journal.pone.0140981.

627 Eppley, T. M., Balestri, M., Campera, M., Rabenantoandro, J., Ramanamanjato, J. B.,
628 Randriatafika, F., et al. (2017). Ecological flexibility as measured by the use of
629 pioneer and exotic plants by two lemurids: *Eulemur collaris* and *Hapalemur*
630 *meridionalis*. *International Journal of Primatology*, 38, 338–357.

631 Eppley, T. M., Borgerson, C., Patel, E. R., Herrera, J. P., Kirkby, A. E., Golden, C. D., et al.
632 (2023). A habitat stronghold on the precipice: A call-to-action for supporting lemur
633 conservation in northeast Madagascar. *American Journal of Primatology*, e23483.

634 Eppley, T. M., & Goodman, S. M. (2022). Non-native habitat use by endemic terrestrial
635 Malagasy mammals. In S. M. Goodman (Ed.), *The New Natural History of*
636 *Madagascar* (pp. 1818–1821). Princeton, USA: Princeton University Press.

637 Eppley, T. M., Santini, L., Tinsman, J. C., & Donati, G. (2020). Do functional traits offset the
638 effects of fragmentation? The case of large-bodied diurnal lemur species. *American*
639 *Journal of Primatology*, 82(4), e23104.

640 Eppley, T. M., Hoeks, S., Chapman, C. A., Ganzhorn, J. U., Hall, K., et al. (2022).
641 Descending from the trees: Factors favoring transitions to terrestriality in primates of
642 Madagascar and the Americas. *Proceedings of the National Academy of Sciences USA*
643 119(42), e2121105119. <https://doi.org/10.1073/pnas.2121105119>.

644 Estrada, A., Raboy, B. E., & Oliveira, L. C. (2012). Agroecosystems and primate conservation
645 in the tropics: a review. *American Journal of Primatology*, 74, 696–711.

646 Estrada, A., Garber, P. A., & Chaudhary, A. (2019). Expanding global commodities trade and
647 consumption place the world’s primates at risk of extinction. *PeerJ*, 7, e7068.

648 Estrada, A., Garber, P. A., Rylands, A. B., Roos, C., Fernandez-Duque, E., Di Fiore, A., et al.
649 (2017). Impending extinction crisis of the world’s primates: Why primates matter.
650 *Science Advances*, 3(1), e1600946.

651 Estrada, A., Garber, P. A., Gouveia, S., Fernández-Llamazares, Á., Ascensão, F., Fuentes, A.,
652 et al. (2022). Global importance of Indigenous Peoples, their lands, and knowledge
653 systems for saving the world’s primates from extinction. *Science Advances*, 8(31),
654 eabn2927.

- 655 Eustace, A., Kisingo, A., Kahana, L., & Lyimo, E. (2015). Activity patterns of black-and-
656 white Colobus monkey (*Colobus guereza caudatus*) in Rau Forest Reserve, Tanzania.
657 *Research and Reviews: Journal of Ecology and Environmental Sciences*, 3(4), 17–24.
- 658 FAO, Global Forest Resources Assessment (2020). Main report (Food and Agriculture
659 Organization of the United Nations, Rome, 2020).
- 660 Fashing, P. (2001). Feeding ecology of guerezas in the Kakamega Forest, Kenya: the
661 importance of Moraceae fruit in their diet. *International Journal of Primatology*,
662 22(4), 579–609.
- 663 Fashing, P. J. (2022). Natural history of black-and-white colobus monkeys. In I. Matsuda, C.
664 C. Grueter, & J. A. Teichroeb (Eds.), *The Colobines: Natural History, Behaviour, and*
665 *Ecology Diversity* (pp. 128–145). Cambridge: Cambridge University Press.
- 666 Fashing, P. J., & Oates, J. F. (2013). *Colobus guereza*. In J. Kingdon, D. Happold, & T.
667 Butynski (Eds.), *Mammals of Africa* (pp. 111–119). London: Bloomsbury Press.
- 668 Fashing, P. J., Dierenatural forest eld, E. S., & Mowry, C. B. (2007). Inatural forest luence of
669 plant and soil chemistry on food selection, ranging patterns, and biomass of *Colobus*
670 *guereza* in Kakamega Forest, Kenya. *International Journal of Primatology*, 28(3),
671 673–703.
- 672 Fashing, P.J., Nguyen, N., Luteshi, P., Opondo, W., Cash, J.F., and Cords, M. (2012).
673 Evaluating the suitability of planted forests for African forest monkeys: A case study
674 from Kakamega Forest, Kenya. *American Journal of Primatology*, 74, 77–90.
- 675 Fashing, P. J., Nguyen, N., Demissew, S., Gizaw, A., Atickem, A., Mekonnen, A., et al.
676 (2022). Ecology, evolution, and conservation of Ethiopia’s biodiversity. *Proceedings*
677 *of the National Academy of Sciences*, 119(50), e2206635119.
- 678 Fashing, P., Nguyen, N., Venkataraman, V., & Kerby, J. (2014). Gelada feeding ecology in an
679 intact ecosystem at Guassa, Ethiopia: variability over time and implications for
680 theropith and hominin dietary evolution. *American Journal of Physical Anthropology*,
681 155(1), 1–16.
- 682 Felton, A. M., Felton, A., Wood, J. T., & Lindenmayer, D. B. (2008). Diet and feeding
683 ecology of *Ateles chamek* in a Bolivian semihumid forest: the importance of *Ficus* as a
684 staple food resource. *International Journal of Primatology*, 29(2), 379–403.
- 685 Fernández, D., Kerhoas, D., Dempsey, A., Billany, J., McCabe, G., & Argirova, E. (2022).
686 The current status of the world’s primates: Mapping threats to understand priorities for
687 primate conservation. *International Journal of Primatology*, 43(1), 15–39.
- 688 Fisaha, G., Hundera, K., & Dalle, G. (2013). Woody plants' diversity, structural analysis and
689 regeneration status of Wof-Washa natural forest, North-east Ethiopia. *African Journal*
690 *of Ecology*, 51(4), 599–608.
- 691 Galan-Acedo, C., Arroyo-Rodríguez, V., Cudney-Valenzuela, S. J., & Fahrig, L. (2019). A
692 global assessment of primate responses to landscape structure. *Biological Reviews*, 94,
693 1605–1618.
- 694 Galan-Acedo, C., Spaan, D., Bicca-Marques, J. C., de Azevedo, R. B., Villalobos, F., &
695 Rosete-Vergés, F. (2021). Regional deforestation drives the impact of forest cover and
696 matrix quality on primate species richness. *Biological Conservation*, 263, 109338.
- 697 Gallegos, T. A., & Glimskar, A. (2009). Computer-aided calibration for visual estimation of
698 vegetation cover. *Journal of Vegetation Science*, 20, 973–983. doi:10.1111/j.1654-
699 1103.2009.01111.x.
- 700 Ganzhorn, J. U., Arrigo-Nelson, S. J., Carrai, V., Chalise, M. K., Donati, G., Droescher, I., et
701 al. (2017). The importance of protein in leaf selection of folivorous primates.
702 *American Journal of Primatology*, 79(4), e22550.

- 703 Ganzhorn, J. U., Rakotondranary, S. J., & Ratovonamana, Y. R. (2011). Habitat description
704 and phenology. In J. M. Setchell & D. J. Curtis (Eds.), *Field and laboratory methods*
705 *in primatology* (pp. 40-56). Cambridge: Cambridge University Press.
- 706 Gebru, T. D. (2016). Deforestation in Ethiopia: Causes, Impacts and Remedy. *International*
707 *Journal of Engineering Development and Research*, 4, (2), 204–209.
- 708 Gerard, A., Ganzhorn, J. U., Kull, C. A., & Carrière, S. M. (2015). Possible roles of alien
709 plants in native fauna conservation: Madagascar as a case study. *Restoration Ecology*,
710 doi:[10.1111/rec.12246](https://doi.org/10.1111/rec.12246).
- 711 Grimes, K., & Paterson, J. D. (2000). Colobus guereza and exotic plant species in the Entebbe
712 Botanical Gardens. *American Journal of Primatology*, 51, 59–60.
- 713 Hammer, Ø., Harper, D. A., & Ryan, P. D. (2001). PAST: Paleontological statistics software
714 package for education and data analysis. *Palaeontologia Electronica*, 4(1), 9.
- 715 Hanya, G., & Chapman, C. A. (2013). Linking feeding ecology and population abundance: a
716 review of food resource limitation on primates. *Ecological research*, 28(2), 183–190.
- 717 Harris, T., Chapman, C. A., & Monatural forest ort, S. L. (2010). Small folivorous primate
718 groups exhibit behavioural and physiological effects of food scarcity *Behavioral*
719 *Ecology*, 21, 46–56.
- 720 Hawes, J. E., & Peres, C. A. (2014). Ecological correlates of trophic status and frugivory in
721 neotropical primates. *Oikos*, 123(3), 365–377.
- 722 Ibrahim, H., Bekele, A., Fashing, P.J., Nguyen, N., Yazezew, D., Moges, A., et al. (2023).
723 Feeding ecology of a highland population of hamadryas baboons (*Papio hamadryas*)
724 at Borena-Sayint National Park, northern Ethiopia. *Primates*,
725 <https://doi.org/10.1007/s10329-023-01077-6>.
- 726 Jansson, C. (2011). *The major food trees of the Angola black-andwhite colobus (Colobus*
727 *angolensis palliatus) in Diani Forest, Kenya* (No. 1652-280X). Sweden: Swedish
728 University of Agricultural Sciences.
- 729 Kamilar, J. M., & Paciulli, L. M. (2008). Examining the extinction risk of specialized
730 folivores: a comparative study of colobine monkeys. *American Journal of*
731 *Primatology*, 70, 816–827. doi: [10.1002/ajp.20553](https://doi.org/10.1002/ajp.20553) PMID: [18521872](https://pubmed.ncbi.nlm.nih.gov/18521872/).
- 732 Kappeler, P. M., & Heymann, E. W. (1996). Nonconvergence in the evolution of primate life
733 history and socio-ecology. *Biological Journal of the Linnean Society*, 59, 297–232.
- 734 Kent, M., & Coker, P. (1992). *Vegetation Description and Analysis. A practical approach*
735 London. New York: Bolhaven Printing Press, John Wiley and Sons. Inc.
- 736 Kessler, M., Kebler, P. J. A., Gradstein, S. R., Bach, K., Schnull, M., & Pitopang, R. (2005).
737 Tree diversity in primary forest and different land use systems in Central Sulawesi,
738 Indonesia. *Biodiversity Conservation*, 14, 547–560.
- 739 Konersmann, C., Noromiarilanto, F., Ratovonamana, Y. R., Brinkmann, K., Jensen, K.,
740 Kobbe, S., et al. (2021). Using utilitarian plants for lemur conservation. *International*
741 *Journal of Primatology*, 43, 1026–1045.
- 742 Kool, K. M. (1989). Behavioural ecology of the silver leaf monkey, *Trachypithecus auratus*
743 *sondaicus*, in the Pangandaran Nature Reserve, West Java, Indonesia. *PhD thesis, the*
744 *University of New South Wales*.
- 745 Krebs, C. J. (1999). *Ecological Methodology*. England: Longman Inc., Harlow.
- 746 Lambert, J. E. (1998). Primate digestion: interactions among anatomy, physiology, and
747 feeding ecology. *Evolutionary Anthropology*, 7(1), 8–20.
- 748 Lambert, J. E. (2011). Primate nutritional ecology: feeding biology and diet at ecological and
749 evolutionary scales. In C. Campbell, A. Fuentes, K. C. MacKinnon, M. Panger, S.
750 Bearder, & R. Stumplantation forest (Eds.), *Primates in perspective* (pp. 512–522).
751 2nd edn. Oxford: Oxford University Press.

- 752 Lambert, J. E., & Rothman, J. M. (2015). Fallback foods, optimal diets, and nutritional
753 targets: primate responses to varying food availability and quality. *Annual Review of*
754 *Anthropology*, 44, 493–512.
- 755 Lim, J. Y., Wasserman, M. D., Veen, J., Despres-Einspenner, M. L., & Kissling, W. D.
756 (2021). Ecological and evolutionary significance of primates' most consumed plant
757 families. *Proceedings of the Royal Society B*, 288(1953), 20210737.
- 758 Machado, F. F., Jardim, L., Dinnage, R., Brito, D., & Cardillo, M. (2023). Diet disparity and
759 diversity predict extinction risk in primates. *Animal Conservation*, 26, 331–339.
- 760 Marsh, L. K., & Chapman, C. A. (2013) *Primates in Fragments: Ecology and Conservation*.
761 New York: Springer.
- 762 Marshall, A., Boyko, C., Feilen, K., Boyko, R., & Leighton, M. (2009). Defining fallback
763 foods and assessing their importance in primate ecology and evolution *American*
764 *Journal of Physical Anthropology* 140, 603–614.
- 765 Matsuda, I., Ihobe, H., Tashiro, Y., Yumoto, T., Baranga, D., & Hashimoto, C. (2020). The
766 diet and feeding behavior of the black-and-white colobus (*Colobus guereza*) in the
767 Kalinzu Forest, Uganda. *Primates*, 61, 473–484.
- 768 McLennan, M., & Hockings, K. (2014). Wild chimpanzees show group differences in
769 selection of agricultural crops. *Scientific Reports*, 4(1), 5956.
770 <https://doi.org/10.1038/srep05956>.
- 771 Mekonnen, A., Bekele, A., Fashing, P. J., Hemson, G., & Atickem, A. (2010). Diet, activity
772 patterns, and ranging ecology of the Bale monkey (*Chlorocebus djamdjamensis*) in
773 Odobullu Forest, Ethiopia. *International Journal of Primatology*, 31(3), 339–362.
- 774 Mekonnen, A., Fashing, P. J., Sargis, E. J., Venkataraman, V. V., Bekele, A., Hernandez-
775 Aguilar, R. A., et al. (2018a). Flexibility in positional behavior, strata use, and
776 substrate utilization among Bale monkeys (*Chlorocebus djamdjamensis*) in response to
777 habitat fragmentation and degradation. *American Journal of Primatology*, 80, e22760.
- 778 Mekonnen, A., Fashing, P. J., Bekele, A., Hernandez-Aguilar, R. A., Rueness, E. K., &
779 Stenseth, N. C. (2018b). Dietary flexibility of Bale monkeys (*Chlorocebus*
780 *djamdjamensis*) in southern Ethiopia: effects of habitat degradation and life in
781 fragments. *BMC Ecology*, 18(1), 4.
- 782 Mekonnen, A., Fashing, P. J., Bekele, A., Hernandez-Aguilar, R. A., Rueness, E. K., Nga, N.,
783 et al. (2017). Impacts of habitat loss and fragmentation on the activity budget, ranging
784 ecology and habitat use of Bale monkeys (*Chlorocebus djamdjamensis*) in the
785 southern Ethiopian Highlands. *American Journal of Primatology*, 79(7), e22644.
- 786 Mekonnen, A., Fashing, P. J., Chapman, C. A., Venkataraman, V. V., & Stenseth, N. C.
787 (2022). The value of flagship and umbrella species for restoration and sustainable
788 development: Bale monkeys and bamboo forest in Ethiopia. *Journal for Nature*
789 *Conservation*, 65, 126117.
- 790 Merker, S., & Yustian, I. (2008). Habitat use analysis of Dian's tarsier (*Tarsius diana*) in a
791 mixed-species plantation in Sulawesi, Indonesia. *Primates*, 49, 161–164.
- 792 Nowak, K., & Lee, P. C. (2013). "Specialist" primates can be flexible in response to habitat
793 alteration. In L. K., Marsh & Chapman, C. A. (Eds.), *Primates in fragments:*
794 *complexity and resilience* (pp. 199–211). New York: Springer Science & Business
795 Media.
- 796 Nyssen, J., Frankl, A., Haile, M., Hurni, H., Descheemaeker, K., Crummey, D., et al. (2014).
797 Environmental conditions and human drivers for changes to north Ethiopian mountain
798 landscapes over 145 years. *Science of the Total Environment*, 485, 164–179.
- 799 Oates, J. (1977a). The guereza and its food. In T. H. Clutton-Brock (Ed.), *Primate Ecology:*
800 *Studies of Feeding and Ranging Behavior in Lemurs, Monkeys and Apes* (pp. 275–
801 321). New York: Academic Press.

- 802 Oates, J.F. (1977b). The guereza and man. In H.R.H. Prince Ranier III and G.H. Bourne
803 (eds.), *Primate Conservation*. New York: Academic Press.
- 804 Onderdonk, D. A., & Chapman, C. A. (2000). Coping with forest fragmentation: the primates
805 of Kibale National Park, Uganda. *International Journal of Primatology*, 21, 587–611.
- 806 Onyekwelu, J. C., Mosandl, R., & Stimm, B. (2008). Tree species diversity and soil status of
807 primary and degraded tropical rainatural forest orest ecosystems in South-Western
808 Nigeria. *Journal of Tropical Forest Science*, 20, 193–204.
- 809 Pinheiro, H. L. N., & Pontes, A. R. (2015). Home range, diet, and activity patterns of common
810 marmosets (*Callithrix jacchus*) in very small and isolated fragments of the Atlantic
811 forest of northeastern Brazil. *International Journal of Ecology*, 2015, 685816.
- 812 Pliosungnoen, M., Gale, G., & Savini, T. (2010). Density and microhabitat use of Bengal slow
813 loris in primary forest and non-native plantation forest. *American Journal of*
814 *Primatology*, 72(12), 1108–1117.
- 815 Plumptre, A. J. (2006). The diets, preferences, and overlap of the primate community in the
816 Budongo Forest Reserve, Uganda: effects of logging on primate diets. In *Primates of*
817 *western Uganda* (pp. 345–371). New York, NY: Springer New York.
- 818 Redei, T., Csecserits, A., Lhotsky, B., Barabás, S., Kröel-Dulay, G., Ónodi, G., & Botta-
819 Dukát, Z. (2020). Plantation forests cannot support the richness of forest specialist
820 plants in the forest-steppe zone. *Forest Ecology and Management*, 461, 117964.
- 821 Rothman, J. M., & Bryer, M. A. (2019). The effects of humans on the primate nutritional
822 landscape. In: A. Behie, J. A. Teichroeb, & N. Malone (Eds.), *Primate Research and*
823 *Conservation in the Anthropocene* (pp. 199–215). Cambridge: Cambridge University
824 Press.
- 825 Ryan, A. M., Chapman, C. A., & Rothman, J. M. (2013). How do differences in species and
826 part consumption affect diet nutrient concentrations? A test with red colobus monkeys
827 in Kibale National Park, Uganda. *African Journal of Ecology*, 51(1), 1–10.
- 828 Sobuj, N. A., & Rahman, M. (2011). Comparison of plant diversity of natural forest and
829 plantations of Rema-Kalenga Wildlife Sanctuary of Bangladesh. *Journal of Forest*
830 *Science*, 27 (3), 127–134.
- 831 Teelen, S. (2007). Primate abundance along five transect lines at Ngogo, Kibale National
832 Park, Uganda. *American Journal of Primatology*, 69(9), 1030–1044.
- 833 Tesfaye, D., Fashing, P. J., Meshesha, A. A., Bekele, A., & Stenseth, N. C. (2021). Feeding
834 ecology of the Omo River guereza (*Colobus guereza guereza*) in habitats with varying
835 levels of fragmentation and disturbance in the southern Ethiopian Highlands.
836 *International Journal of Primatology*, 42(1), 64–88.
- 837 Torres-Romero, E. J., Nijman, V., Fernández, D., & Eppley, T. M. (2023). Human-modified
838 landscapes driving the global primate extinction crisis. *Global Change Biology*,
839 <https://doi.org/10.1111/gcb.16902>.
- 840 Veronika, S. (2008). *Wof Washa – cave of bird Dynamics of a Forest at the Eastern*
841 *Escarpment of the Ethiopian Highlands*. University of Zurich, Zurich.
- 842 Wali, A., Alvira, D., Tallman, P. S., Ravikumar, A., & Macedo, M. O. (2017). A new
843 approach to conservation: using community empowerment for sustainable well-being.
844 *Ecology and Society* 22(4):6. <https://doi.org/10.5751/ES-09598-220406>.
- 845 Wasserman, M. D., & Chapman, C. A. (2003). Determinants of colobine monkey abundance:
846 The importance of food energy, protein and fiber content. *Journal of Animal Ecology*,
847 72, 650–659.
- 848 Yazezew, D., Bekele, A., Fashing, P. J., Nguyen, N., Ibrahim, H., Amara Moges, et al. (In
849 Prep.). *Activity budget and ranging ecology of Omo River Guereza (Colobus guereza*
850 *guereza) in natural and plantation forest of Wof-Washa, central highlands of Ethiopia.*

- 851 Yazezew, D., Bekele, A., Fashing, P. J., Nguyen, N., Moges, A., Ibrahim, H., et al. (2022).
852 Population size and habitat preference of the Omo River guereza (*Colobus guereza*
853 *guereza*) in a multi-habitat matrix in the central highlands of Ethiopia. *Primates*, 63,
854 151–160.
- 855 Zinner, D., Tesfaye, D., Stenseth, N. C., Bekele, A., Mekonnen, A., et al. (2019). Is *Colobus*
856 *guereza gallarum* a valid endemic Ethiopian taxon? *Primate Biology*, 6, 7–16.
857

858 **Table S1** Results of enumeration of plants ≥ 10 cm DBH in the quadrats of transects within the home ranges of the natural and plantation forest
 859 study groups of Omo River guerezas at Wof-Washa Natural State Forest, Ethiopia (rank is based on abundance).

Rank	Family	Species	Stems/ ha	% total trees	Basal area/ ha(cm ² /ha)	Origin	Utilitarian
Natural Forest (n=234 trees)							
1	Oliniaceae	<i>Olinia rochetiana</i>	150.0	19.2	801.4	Ind	Fn, FT, Fw, Tim
2	Ericaceae	<i>Erica arborea</i>	130.0	16.7	565.4	Ind	Ch, Fd, Fn, Fw
3	Sapindaceae	<i>Allophylus abyssinicus</i>	113.3	14.5	995.0	Ind	FT, Fw, Tim
3	Cupressaceae	<i>Juniperus procera</i>	113.3	14.5	4228.6	Ind	Fur, Fw, Tim
5	Myrsinaceae	<i>Maesa lanceolata</i>	100.0	12.8	1728.3	Ind	Fn, Fw
6	Solanaceae	<i>Discopodium penninervium</i>	26.7	3.4	402.3	Ind	FT, Fw,
7	Rubiaceae	<i>Galiniera saxifraga</i>	20.0	2.6	305.2	Ind	Fw, Tim
7	Loganiaceae	<i>Nuxia congesta</i>	20.0	2.6	1015.5	Ind	Ch, Fn, Fw
7	Oleaceae	<i>Olea europaea</i>	20.0	2.6	738.5	Ind	Fur, Fw, Tim
10	Pittosporaceae	<i>Pittosporum viridiflorum</i>	16.7	2.1	1557.9	Ind	FT, Fw, Tim
11	Podocarpaceae	<i>Podocarpus falcatus</i>	13.3	1.7	3024.9	Ind	Fur, Fw, Tim
11	Asteraceae	<i>Vernonia leopoldi</i>	13.3	1.7	847.8	Ind	Fw, MU
13	Santalaceae	<i>Osyris quadripartita</i>	10.0	1.3	277.6	Ind	Fw, Tim
14	Meliantaceae	<i>Bersama abyssinica</i>	6.7	0.9	376.8	Ind	Fn, Fw
14	Scrophulariaceae	<i>Halleria lucida</i>	6.7	0.9	364.3	Ind	Fw, MU
16	Cupressaceae	<i>Cupressus lusitanica</i>	3.3	0.4	588.8	Ex	Fur, Fw, Tim
16	Flacourtiaceae	<i>Dovyalis abyssinica</i>	3.3	0.4	261.7	Ind	Fn, Food, MU
16	Aquifoliaceae	<i>Ilex mitis</i>	3.3	0.4	669.9	Ind	Ch, FT, Fw, MU, Tim
16	Celastraceae	<i>Maytenus arbutifolia</i>	3.3	0.4	376.8	Ind	Fd, Fn, FT, Fw

16	Myricaceae	<i>Myrica salicifolia</i>	3.3	0.4	2936.6	Ind	Fw, Tim
16	Myrsinaceae	<i>Myrsine africana</i>	3.3	0.4	550.2	Ind	Fn, Fw, MU
Total			780.0	100.0	22613.3		
Plantation Forest (n = 180 trees)							
1	Myrsinaceae	<i>Maesa lanceolata</i>	266.7	44.4	1099.7	Ind	Fn, Fw, MU
2	Cupressaceae	<i>Juniperus procera</i>	166.7	27.8	1755.3	Ind	Fur, Fw, MU, Tim
3	Cupressaceae	<i>Cupressus lusitanica</i>	50.0	8.3	1519.8	Ex	Fur, Fw, Tim
3	Podocarpaceae	<i>Podocarpus falcatus</i>	50.0	8.3	462.9	Ind	Fur, Fw, MU, Tim
5	Solanaceae	<i>Discopodium penninervium</i>	13.3	2.2	277.6	Ind	FT, Fw
5	Fabaceae	<i>Erythrina brucei</i>	13.3	2.2	4906.0	Ind	Fd, Fn, Fur, Fw, MU
7	Asteraceae	<i>Vernonia leopoldi</i>	10.0	1.7	462.9	Ind	Fn, Fw, MU
8	Melanthaceae	<i>Bersama abyssinica</i>	6.7	1.1	261.7	Ind	Fn, Fw
8	Myrsinaceae	<i>Embelia schimperi</i>	6.7	1.1	167.5	Ind	Fw, MU
8	Aquifoliaceae	<i>Ilex mitis</i>	6.7	1.1	18463.2	Ind	Ch, FT, Fw, MU, Tim
8	<i>Oliniaceae</i>	<i>Olinia Rochetiana</i>	6.7	1.1	3168.9	Ind	Fn, FT, Fw, Tim
9	Moraceae	<i>Ficus Sur</i>	3.3	0.6	49112.2	Ind	Food, Fur, Tim
Total			600.0	100.0	81657.5		

860 Key: Ex=Exotic; Ind = Indigenous; Fw = Firewood; Tim = Timber; Fur = Furniture; FT = Farm Tools; Ch = Charcoal; Fd = Fodder; Fn = Fence,

861 MU=Medicinal Use

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866 Table S2. Important Value Index (IVI) of plant species occurring in the home ranges of Omo
 867 River guereza study groups in plantation forest and natural Forest at Wof-Washa Natural State
 868 Forest, Ethiopia.

Plantation Forest				Natural Forest			
Species	Relative Dominance	IVI	%IVI	Species	Relative Dominance	IVI	% IVI
<i>Maesa lanceolata</i>	35.1	80.0	54.5	<i>Olinia rochetiana</i>	8.5	27.9	22.8
<i>Juniperus procera</i>	8.8	36.7	25.0	<i>Erica arborea</i>	3.5	20.3	16.6
<i>Cupressus lusitanica</i>	0.9	9.3	6.3	<i>Allophylus abyssinicus</i>	2.9	17.5	14.3
<i>Podocarpus falcatus</i>	1.0	9.4	6.4	<i>Juniperus procera</i>	3.2	17.8	14.6
<i>Discopodium penninervium</i>	0.1	2.3	1.6	<i>Maesa lanceolata</i>	2.8	15.7	12.8
<i>Erythrina brucei</i>	0.1	2.3	1.6	<i>Discopodium penninervium</i>	0.2	3.6	3.0
<i>Vernonia leopoldi</i>	0.0	1.7	1.2	<i>Galiniera saxifraga</i>	0.1	2.7	2.2
<i>Bersama abyssinica</i>	0.0	1.1	0.8	<i>Nuxia congesta</i>	0.1	2.7	2.2
<i>Embelia schimperi</i>	0.0	1.1	0.8	<i>Olea europaea</i>	0.1	2.7	2.2
<i>Ilex mitis</i>	0.0	1.1	0.8	<i>Pittosporum viridiflorum</i>	0.1	2.2	1.8
<i>Olinia rochetiana</i>	0.0	1.1	0.8	<i>Podocarpus falcatus</i>	0.0	1.8	1.4
<i>Ficus Sur</i>	0.0	0.6	0.4	<i>Vernonia leopoldi</i>	0.0	1.8	1.4
		146.8		<i>Osyris quadripartita</i>	0.0	1.3	1.1
				<i>Bersama abyssinica</i>	0.0	0.9	0.7
				<i>Halleria lucida</i>	0.0	0.9	0.7
				<i>Cupressus lusitanica</i>	0.0	0.4	0.4
				<i>Dovyalis abyssinica</i>	0.0	0.4	0.4
				<i>Ilex mitis</i>	0.0	0.4	0.4
				<i>Maytenus arbutifolia</i>	0.0	0.4	0.4
				<i>Myrica salicifolia</i>	0.0	0.4	0.4
				<i>Myrsine africana</i>	0.0	0.4	0.4

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878 Table S3. Contribution of food items from each plant species consumed by Omo River guerezas in plantation forest at Wof-Washa Natural State
 879 Forest, Ethiopia (n=3972).

Family	Species	*Growth form	Young leaves	Mature leaves	Fruit	Shoot	Stem	Total spp. contribution
Myrsinaceae	<i>Maesa lanceolata</i>	T	20.51	8.48	0.05	0.55	1.06	30.65
Podocarpaceae	<i>Podocarpus falcatus</i>	T	3.61	0.38	12.06	0.08	-	16.12
Aquifoliaceae	<i>Ilex mitis</i>	T	10.62	2.77	-	0.28	0.45	14.13
Asteraceae	<i>Vernonia leopoldi</i>	S	8.98	3.96	-	0.13	0.48	13.55
Solanaceae	<i>Discopodium penninervium</i>	S	5.17	2.83	-	-	0.63	8.63
Cupressaceae	<i>Juniperus procera</i>	T	2.17	1.14	0.48	0.1	0.03	3.91
Oliniaceae	<i>Olinia rochetiana</i>	T	2.55	0.98	-	0.05	0.13	3.71
Moraceae	<i>Ficus sur</i>	T	0.03	-	3.51	-	-	3.53
Melianthaceae	<i>Bersama abyssinica</i>	T/S	1.79	0.96	-	-	0.1	2.85
Cupressaceae	<i>Cupressus lusitanica</i>	T	1.11	0.4	-	0.03	-	1.54
Oleaceae	<i>Jasminum abyssinicum</i>	L/C	0.55	-	-	-	-	0.55
Ranunculaceae	<i>Clematis hirsuta</i>	L/C	0.3	0.03	-	0.03	-	0.35
	Unidentified mosses	E	0.18	-	-	-	-	0.18
Myrsinaceae	<i>Embelia schimperi</i>	L/C	0.08	-	-	-	-	0.08
Rosaceae	<i>Rubus steudneri</i>	S	0.05	0.03	-	-	-	0.08
Scrophulariaceae	<i>Halleria lucida</i>	T/S	0.05	-	-	-	-	0.05
Myrsinaceae	<i>Myrsine africana</i>	S	0.03	-	-	-	0.03	0.05
Oleaceae	<i>Olea europaea</i>	T	0.03	-	-	-	-	0.03
Total			57.8	21.94	16.12	1.24	2.9	100

880 Note: *Growth form T= Tree, T/S= Tree /Shrub, S= Shrub, H= Herb, L/C= Liana/Climber, E= Epiphyte.

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882 Table S4. Contribution of food items from each plant species consumed by Omo River guerezas in natural forest at Wof-Washa Natural State
 883 Forest, Ethiopia (n=3228).

Family	Species	*Growth form	Young leaves	Mature leaves	Fruit	Shoot	Stem	Total spp. contribution
Asteraceae	<i>Vernonia leopoldi</i>	S	16.05	11.83	0.19	0.9	0.81	29.77
Myrsinaceae	<i>Maesa lanceolata</i>	T	8.4	4.71	0.99	1.15	0.40	15.64
Pittosporaceae	<i>Pittosporum viridiflorum</i>	T	7.13	2.94	1.18	0.59	0.15	11.99
Oliniaceae	<i>Olinia rochetiana</i>	T	4.46	2.11	0.12	0.46	0.12	7.28
Aquifoliaceae	<i>Ilex mitis</i>	T	3.00	2.29	-	0.37	-	5.67
Rubiaceae	<i>Galiniera saxifraga</i>	T	2.79	1.24	0.50	0.31	0.09	4.93
Solanaceae	<i>Discopodium penninervium</i>	S	2.04	1.49	0.19	0.46	0.28	4.46
Sapindaceae	<i>Allophylus abyssinicus</i>	T	2.51	0.90	0.09	0.22	0.03	3.75
Cupressaceae	<i>Juniperus procera</i>	T	2.29	0.65	0.25	0.15	0.03	3.38
Cupressaceae	<i>Cupressus lusitanica</i>	T	1.08	0.87	-	0.59	-	2.54
Moraceae	<i>Ficus sur</i>	T	-	-	1.89	-	-	1.89
Oleaceae	<i>Olea europaea</i>	T	1.02	0.53	0.03	0.12	-	1.70
Melanthaceae	<i>Bersama abyssinica</i>	T/S	0.81	0.37	0.06	0.22	0.06	1.52
Scrophulariaceae	<i>Halleria lucida</i>	T/S	0.59	0.34	-	0.34	0.09	1.36
Podocarpaceae	<i>Podocarpus falcatus</i>	T	0.84	0.09	-	-	0.03	0.96
Loganiaceae	<i>Nuxia congesta</i>	T	0.28	0.50	-	0.09	-	0.87
Myrsinaceae	<i>Myrsine africana</i>	S	0.43	0.12	-	0.03	-	0.59
Flacourtiaceae	<i>Dovyalis abyssinica</i>	S	0.34	0.03	0.03	-	-	0.40
Ranunculaceae	<i>Clematis hirsuta</i>	L/C	0.28	0.06	-	-	-	0.34
Meliaceae	<i>Turraea holstii</i>	S	0.19	0.03	-	-	-	0.22
	Unidentified mosses	E	0.22	-	-	-	-	0.22
Myrsinaceae	<i>Embelia schimperi</i>	S	0.09	0.03	-	-	-	0.12
Ericaceae	<i>Erica arborea</i>	T/S	0.06	0.03	-	0.03	-	0.12
Celastraceae	<i>Maytenus arbutifolia</i>	T/S	-	0.06	-	-	-	0.06
Anacardiaceae	<i>Rhus natalensis</i>	T	0.06	-	-	-	-	0.06
Balsaminaceae	<i>Impatiens tinctoria</i>	H	0.03	-	-	-	-	0.03
Oleaceae	<i>Jasminum abyssinicum</i>	L/C	-	-	-	0.03	-	0.03

Lamiaceae	<i>Plectranthus lactiflorus</i>	H	0.03	-	-	-	-	0.03
Urticaceae	<i>Urera hypselodendron</i>	L/C	0.03	-	-	-	-	0.03
Cucurbitaceae	<i>Zehneria scabra</i>	L/C	0.03	-	-	-	-	0.03
Total			55.08	31.23	5.51	6.07	2.11	100

884 Note: * Growth form T= Tree, T/S= Tree /Shrub, S= Shrub, H= Herb, L/C= Liana/Climber, E= Epiphyte.