

# The montane trees of the Cameroon Highlands, West-Central Africa, with the Endangered, *Deinbollia onanae* sp. nov. (Sapindaceae), a new primate-dispersed, Endangered species (#54299)

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# The montane trees of the Cameroon Highlands, West-Central Africa, with the Endangered, *Deinbollia onanae* sp. nov. (Sapindaceae), a new primate-dispersed, Endangered species

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We test the hypothesis that the tree species previously known as *Deinbollia* sp. 2. is a new species for science. We formally characterise and name this species as *Deinbollia onanae* (Sapindaceae-Litchi clade) and we discuss it in the context of the assemblage of montane tree species in the Cameroon Highlands of West-Central Africa. The new species is a shade-bearing, non-pioneer understory forest tree species reaching 15 m high and a trunk diameter that can attain over 40 cm at 1.3 m above the ground. Seed dispersal has been recorded by chimpanzees (*Pan troglodytes ellioti*) and by putty-nose monkeys (*Cercopithecus nictitans*) and the species is used by chimpanzees for nesting. Cameroon has the highest species-diversity and species endemism known in this African-Western Indian Ocean genus of 42, mainly lowland species. *Deinbollia onanae* is an infrequent tree species known from six locations in surviving islands of montane (sometimes also upper submontane) forest along the line of the Cameroon Highlands, including one at Ngel Nyaki in Mambilla, Nigeria. *Deinbollia onanae* is here assessed as Endangered according to the IUCN 2012 standard, threatened by severe fragmentation of its mountain forest habitat due to extensive and ongoing clearance for agriculture. The majority of the 28 tree species of montane forest (above 2000 m alt.) in the Cameroon Highlands are also widespread in East African mountains (i.e. are Afromontane wide). *Deinbollia onanae* is one of only seven species known to be endemic (globally restricted to) these highlands. It is postulated that this new species is in a sister relationship with *Deinbollia oreophila*, a frequent species at a lower (submontane) altitudinal band of the same range. Detailed ecological data on *Deinbollia onanae* from the Nigerian location, Ngel Nyaki where it has been known under the name *Deinbollia "pinnata"* is reviewed.

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2 Endangered, *Deinbollia onanae* sp. nov. (Sapindaceae), a new primate-dispersed,  
3 Endangered species

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## ABSTRACT

We test the hypothesis that the tree species previously known as *Deinbollia sp. 2.* is a new species for science. We formally characterise and name this species as *Deinbollia onanae* (Sapindaceae-Litchi clade) and we discuss it in the context of the assemblage of montane tree species in the Cameroon Highlands of West-Central Africa. The new species is a shade-bearing, non-pioneer understorey forest tree species reaching 15 m high and a trunk diameter that can attain over 40 cm at 1.3 m above the ground. Seed dispersal has been recorded by chimpanzees (*Pan troglodytes ellioti*) and by putty-nose monkeys (*Cercopithecus nictitans*) and the species is used by chimpanzees for nesting. Cameroon has the highest species-diversity and species endemism known in this African-Western Indian Ocean genus of 42, mainly lowland species. *Deinbollia onanae* is an infrequent tree species known from six locations in surviving islands of montane (sometimes also upper submontane) forest along the line of the Cameroon Highlands, including one at Ngel Nyaki in Mambilla, Nigeria. *Deinbollia onanae* is here assessed as Endangered according to the IUCN 2012 standard, threatened by severe fragmentation of its mountain forest habitat due to extensive and ongoing clearance for agriculture. The majority of the 28 tree species of montane forest (above 2000 m alt.) in the Cameroon Highlands are also widespread in East African mountains (i.e. are Afromontane wide). *Deinbollia onanae* is one of only seven species known to be endemic (globally restricted to) these highlands. It is postulated that this new species is in a sister relationship with *Deinbollia oreophila*, a frequent species at a lower (submontane) altitudinal band of the same range. Detailed ecological data on *Deinbollia onanae* from the Nigerian location, Ngel Nyaki where it has been known under the name *Deinbollia "pinnata"* is reviewed.

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## 105 INTRODUCTION.

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As part of the project to designate Important Plant Areas (IPAs) in Cameroon (also known as Tropical Important Plant Areas or TIPAs), we are striving to name, assess the conservation status and include in IPAs (Darbyshire et al., 2017) rare and threatened plant species in the surviving, threatened natural habitat of the Cross-Sanaga interval (Cheek et al., 2001).

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Several of these species were previously designated as new to science but not formally published in a series of checklists (see below) ranging over much of the Cross-Sanaga interval. The Cross-Sanaga has the highest vascular plant species diversity per degree square in tropical Africa (Barthlott et al., 1996) but natural habitat is being steadily being cleared, predominantly for agriculture.

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In this paper we test the hypothesis that the high-altitude tree species formerly designated as “*Deinbollia* sp. 2” (Harvey et al., 2004, Cheek et al., 2004, Cheek et al., 2010), “*Deinbollia* sp.” (Chapman & Chapman, 2001) or “*Deinbollia pinnata*” (Abiem et al., 2020), is a new species to science, and we describe, characterise and name it as *Deinbollia onanae* Cheek. The species is discussed in the context of the assemblage of the other montane forest tree species (occurring above 2000 m alt.) in the Cameroon Highlands (see Discussion below).

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The genus *Deinbollia* Schum. & Thonn. is traditionally place in the tribe Sapindeae DC. and is characterised by its 1-pinnate, imparipinnate leaves, flowers with petals well developed and about the same in number as the imbricate sepals, the petals with a well-developed ligule (or appendage) on the adaxial surface and with stamens 9–30 in number, the intrastaminal disc central, the edge with more than 5 shallow ridges. The fruits develop 1–3 indehiscent, apocarpous fleshy mericarps (Fouilloy & Hallé, 1973).

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Molecular Phylogenetic sampling of Sapindaceae is incomplete with many African genera not represented, as can be seen in Buerki et al., (2009). In that study *Deinbollia* is represented by six samples of four species, all from Madagascar (on which limited basis it appears monophyletic) and is resolved in the informally named ‘Litchi Group’ of genera, where it is in a sister relationship to a subclade comprising the genera *Lepisanthes* Blume (Africa to Asia) *Atalaya* Blume (American) and *Pseudima* Radlk. (American) (Buerki et al., 2009). The delimitation of

138 Sapindaceae in this paper follows the evidence of *Buerki et al.*, (2010), that is, excluding  
139 Aceraceae, Hippocastanaceae and Xanthoceraceae which have sometimes been included within  
140 it.

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142 *Deinbollia* has 42 accepted species, one shared between Africa, Reunion and Madagascar, 5  
143 endemic to Madagascar, and 35 species restricted to subsaharan continental Africa. The species  
144 predominantly occur in lowland evergreen forest and are absent from countries that lack this  
145 habitat such as Rwanda, Burundi, Swaziland and Lesotho (high altitude) and Namibia, Botswana,  
146 Eritrea, Mali and Burkina Faso (low rainfall and lacking significant evergreen forest). The  
147 highest species diversity is found in Cameroon, with 16 species (Plants of the World Online  
148 accessed May 2020). Cameroon has the highest levels of country-level endemism in the genus.  
149 Ten of the Cameroon species are globally threatened with extinction (Cheek in *Onana & Cheek*  
150 *2011: 314–316; Cheek, 2004a; Cheek, 2017a; Cheek, 2017b*). In contrast only 10 species are  
151 recorded for the whole of West Tropical Africa (*Keay, 1958*). Since the Flore Du Cameroun  
152 account was published (*Fouilloy & Hallé, 1973*), several further species apart from those listed  
153 below, were published for Cameroon by *Thomas (1986)*. The genus was last revised by  
154 *Radlkofer (1932)*.

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156 In the 21<sup>st</sup> century only two new species to science have been published in the genus, *Deinbollia*  
157 *mezilii* D.W.Thomas & D.J.Harris (*Thomas & Harris, 2000*) and *D. oreophila* Cheek (*Cheek &*  
158 *Etuge 2009*), both from Cameroon. But specimens often remain unidentified in herbaria. For  
159 example, 16 specimens unidentified to species are listed in the Gabon Checklist (*Sosef et*  
160 *al., 2005*). The genus has no major uses but the fruits of several species are reported as being  
161 edible by humans, and the seeds are probably primate-dispersed or dispersed by large  
162 frugivorous birds, and the flowers probably bee-pollinated (*Cheek & Etuge, 2009*). Several  
163 species are recorded to be useful locally in West Africa especially medicinally, e.g. the bark of *D.*  
164 *grandifolia* Hook.f. is used for treating jaundice and the wood for planks (*Burkill, 2000:17–19*).

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## 167 METHODS & MATERIALS

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169 *The electronic version of this article in Portable Document Format (PDF) will represent a*  
170 *published work according to the International Code of Nomenclature for algae, fungi,*  
171 *and plants (ICN), and hence the new names contained in the electronic version are*  
172 *effectively published under that Code from the electronic edition alone. In addition, new*  
173 *names contained in this work which have been issued with identifiers by IPNI will*  
174 *eventually be made available to the Global Names Index. The IPNI LSIDs can be*  
175 *resolved and the associated information viewed through any standard web browser by*  
176 *appending the LSID contained in this publication to the prefix "<http://ipni.org/>". The*  
177 *online version of this work is archived and available from the following digital*  
178 *repositories: PeerJ, PubMed Central, and CLOCKSS.*

179 Fieldwork in Cameroon resulting in the specimens cited in this paper was conducted under the  
180 terms of the series of Memoranda of Collaboration between IRAD (Institute for Agronomic  
181 Research and Development)-National Herbarium of Cameroon and Royal Botanic Gardens,  
182 Kew beginning in 1992, the most recent of which is valid until 5<sup>th</sup> Sept. 2021. The most recent




183 research permit issued for fieldwork under these agreements was  
184 000146/MINRESI/B00/C00/C10/C12 (issued 28 Nov 2019), and the export permit number was  
185 098/IRAD/DG/CRRA-NK/SSRB/12/2019 (issued 19 Dec 2019). At the Royal Botanic Gardens,  
186 Kew, fieldwork was approved by the Institutional Review Board of Kew entitled the Overseas  
187 Fieldwork Committee (OFC) for which the most recent registration number was OFC 807-3  
188 (2019). The most complete set of duplicates for all specimens made was deposited at YA, the  
189 remainder exported to K for identification and distribution following standard practice. Field  
190 work methodology followed was *Cheek & Cable (1997)*.

191 Herbarium citations follow Index Herbariorum (*Thiers et al., 2020*). Specimens indicated “!”  
192 were seen by one or more authors, they were studied at K, P, WAG, and YA. The National  
193 Herbarium of Cameroon, YA, was also searched for additional material of the new taxon as was  
194 Tropicos (<http://legacy.tropicos.org/SpecimenSearch.aspx>). During the time that this paper was  
195 researched in 2019–2020, it was not possible to obtain physical access to material at WAG (due  
196 to the transfer of WAG to Naturalis, Leiden, subsequent construction work, and covid-19 travel  
197 and access restrictions). However images for WAG specimens were studied at  
198 <https://bioportal.naturalis.nl/?language=en> and those from P at  
199 [https://science.mnhn.fr/institution/mnhn/collection/p/item/search/form?lang=en\\_US](https://science.mnhn.fr/institution/mnhn/collection/p/item/search/form?lang=en_US). Specimens  
200 of *Deinbollia* at FHO could not be accessed due to covid-19 and are not available digitally.  
201 Specimens at FHI are also not available digitally. We also searched *JStor Global Plants (2020)*  
202 for additional type material of the genus not already represented at K.

203  
204 Binomial authorities follow the International Plant Names Index (*IPNI, 2020*). The conservation  
205 assessment was made using the categories and criteria of *IUCN (2012)*. GeoCAT was used to  
206 calculate red list metrics (*Bachman et al., 2011*). Herbarium material was examined with a Leica  
207 Wild M8 dissecting binocular microscope fitted with an eyepiece graticule measuring in units of  
208 0.025 mm at maximum magnification. The drawing was made with the same equipment using  
209 Leica 308700 camera lucida attachment. Flowers from herbarium specimens of the new species  
210 described below were soaked in warm water to rehydrate the flowers, allowing dissection,  
211 characterisation and measurement. The terms and format of the description follow the  
212 conventions of (*Cheek & Etuge, 2009*). Georeferences for specimens lacking latitude and  
213 longitude were obtained using Google Earth ([https://www.google.com/intl/en\\_uk/earth/versions/](https://www.google.com/intl/en_uk/earth/versions/)  
214 ). The map was made using SimpleMappr (<https://www.simplemappr.net>).

## 215 RESULTS

### 216 TAXONOMIC TREATMENT

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219 *Deinbollia* sp. 2 (Fig. 1), because it has leaves of flowering branches less than 1 m long, only  
220 sparsely hairy on the lower surface, leaflets more than 15 cm long and sepals adaxially glabrous,  
221 flower buds very sparsely hairy and less than 5 mm diam., borne on a branched inflorescence  
222 10–30 cm long, keys out in the Flore Du Cameroun treatment of *Deinbollia* (*Fouilloy & Hallé,*  
223 *1973*) to a couplet leading to *D. grandifolia* Hook.f. and *D. maxima* Gilg. However, it differs  
224 from these two species in having (2–)8–11-jugate (not 4–7-jugate), and in other characters shown  
225 in table 1. In its Nigerian location  its species has been referred to as *D. pinnata* (*Abiem et al.,*  
226 *2020*). *Deinbollia pinnata* Schum. & Thonn. is a common lowland West African species that

227 occurs from Guinea to Nigeria, it differs in being densely hairy, so that the lower surface of the  
228 leaflets are softly hairy to the touch due to dense, patent, translucent hairs, and it is usually a  
229 small shrub of disturbed habitats, with an unbranched, raceme-like inflorescence that is  
230 pendulous in fruit, with hairy fruits 12-13 mm wide  
231 (see [https://commons.wikimedia.org/wiki/File:Deinbollia\\_pinnata\\_MS6765.jpg](https://commons.wikimedia.org/wiki/File:Deinbollia_pinnata_MS6765.jpg); Keay 1958:  
232 714–715). In contrast, *Deinbollia sp. 2* has only a very few, sparse, red, subappressed hairs along  
233 the midrib and secondary nerves, is a tree of intact high elevation forest, the inflorescence is  
234 erect, with numerous long branches bearing glabrous fruits 20 mm or more wide (see description  
235 below). Additional characters separating *Deinbollia sp. 2* from *Deinbollia pinnata* are included  
236 in table 1.

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239 The affinities of *Deinbollia sp. 2* may be with the recently described *D. oreophila* since this  
240 species also occurs at altitude in the Cameroon Highlands and both species share numerous  
241 raised lenticels and also leaflets with high length: breadth ratios and with high numbers of  
242 secondary nerves. Both species share an unusual structure which is also seen in *Deinbollia*  
243 *pinnata*: the adaxial surface of the leaf rhachis is not rounded as in the other West African  
244 species, but flattened, the margins slightly raised forming acute angles with the sides, with a  
245 distinct, raised midline (Cheek & Onana, 2009: Fig. 1C). In fact, at two locations, Mt Kupe and  
246 Bali Ngemba, the two species *D. oreophila* and *Deinbollia sp. 2* are sympatric and their  
247 altitudinal ranges can overlap (Cheek et al., 2004; Harvey et al., 2004). As the only two species  
248 of the genus to grow at altitude in the Cameroon Highlands, there is a possibility that they might  
249 be confused with each other. The two species can be separated using the morphological  
250 characters presented in table 2.

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253 ***Deinbollia onanae* Cheek sp. nov.** – Fig. 1–3

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Similar to but differing from *Deinbollia oreophila* Cheek in the length of leaves of flowering  
stems (14–)60–70 cm, number of leaflets per leaves (4–)16–23, width of leaflets (2.1–)2.5–4 cm,  
number of secondary nerves on each side of midrib (15–)17–18, versus respectively, 25–63 cm,  
(4–)6–8(–10), (2.1–)2.5–4 cm and (15–)17–18; versus stems with lenticels brown, concolorous  
and inconspicuous, not discolorous, bright white and conspicuous; ovary bilocular not trilocular.

256 Typus: Cameroon, *Mt Oku and the Ijim Ridge*, Aboh to Tum, 2400 m alt., fl. 22 Nov. 1996,  
257 *Etuge 3600* (holotype K000337729! Fig. 2, isotypes MO!, WAG0336084!, WAG0336083!,  
258 YA0057050!);

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261 *Deinbollia cf. pinnata* Schum. & Thonn., sensu Cheek, in Cheek et al., (2000:162).

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263 *Deinbollia sp. 2* sensu Cheek in Harvey et al., (2004: 125); Cheek & Etuge in Cheek et al., (2004:  
264 399); Cheek in Cheek et al., (2010: 143, fig 23).

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266 *Deinbollia sp.* Chapman & Chapman 2001: c41

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270 Monoecious tree or treelet (4–)5–15 m tall, trunk 14.5–40 cm diameter at 1.3 m from the ground,  
271 lacking exudate or scent when wounded, sparingly branched, nearly glabrous, apart from the  
272 inflorescence. Stems of flowering branches terete 1–1.5 cm diameter, solid (not hollow), second  
273 internode below apical inflorescence 2–2.5 cm long, outer epidermis pale grey-brown,  
274 contrasting with the darker brown bases of the adjoining petiolar pulvini, lenticels dense, raised,  
275 elliptic, 0.6–1.1 mm long, concolorous, inconspicuous, glabrescent, hairs sparse to dense, dark  
276 brown, cylindrical 0.1–0.5 mm long.

277

278 Leaves alternate, pinnately compound, (14–)60–70 cm long; leaflets (4–)16–23 per leaf on  
279 flowering stems, leaflets 10–14 per leaf on leaves of juvenile trees. Petiole (4–)9.5–20.8 cm long,  
280 terete, c. 4 mm diameter at midpoint, drying pale yellow; basal pulvini dark brown; rhachis (4.5–  
281 )32–44 cm long, (2–)8–11-jugate on flowering stems, 5–7-jugate on non-flowering stems of  
282 juvenile trees, the upper surface of the distal half flattened with two thin lateral wings and with a  
283 central dark hairy rounded central ridge, the rest of the rhachis glabrescent with sparse  
284 inconspicuous hairs (*de Wilde* 4555), or with sparse dark brown appressed hairs (*Cable* 3386).  
285 Leaflets mostly oblong (6.6–)14–19.5 x (2.1–)2.5–4 cm, (but leaflets of sterile branches to 6.5  
286 cm wide), acumen c. 1 cm long, base broadly acute, slightly asymmetric, (basalmost leaflets  
287 lanceolate and about half the length of the other leaflets) lateral nerves and midrib yellow, raised  
288 above and below, convex, (15–)17–18 on each side of the midrib, nearly brochidodromous, the  
289 lateral nerve apices forming a weak irregular submarginal nerve, stronger branches uniting with  
290 the secondary nerve above, intersecondary nerves strong, parallel to the secondaries, tertiary and  
291 quaternary nerves reticulate raised yellow and conspicuous, on both surfaces, contrasting with  
292 the pale grey-green areolae (except in *Cable* 3386(K) where they are concolorous and so  
293 inconspicuous above, possibly an artefact of poor drying); upper surface glabrous, lower surface  
294 with inconspicuous, minute, cylindrical, subappressed glossy dark-brown hairs c. 0.25 mm long,  
295 distributed very sparsely along the midrib and secondary nerves, absent from mature leaves of  
296 non-flowering specimens (e.g. *Cheek* 8709) but then the same hair type present on axillary buds  
297 and young leaves; petiolules yellow, 2–5 mm long, glabrous.

298

299 Inflorescence a 80–120-flowered, loose, terminal panicle 25 x 10 cm; auxiliary inflorescences  
300 sometimes present in the axils of the distal 1–4 leaves (*Cheek* 13625); peduncle of terminal  
301 inflorescences 0–2 cm long; rhachis internodes (1–)2–3 cm long, shortest in the distal portion;  
302 first order bracts caducous; indumentum brown hairy; primary branches 10–20 per inflorescence,  
303 2–8 cm long, each bearing (1–)2–5 partial-inflorescences; partial-peduncles 0–5 mm long, apex  
304 with a cluster of 3–5 bracteoles; bracteoles subulate to narrowly lanceolate, 2–3 mm long, apex  
305 narrowly acute, partial-inflorescences (1–)3-flowered in glomerules, pedicels erect, terete, 3–4 x  
306 1.5 mm (female), 4–5 x 1 mm (male), sparsely puberulent, hairs 0.1–0.5 mm long.

307

308 Flowers white, scent not recorded, flower buds c. 4 mm diam., open flowers c. 6 x 7 mm. Calyx  
309 with sepals 5(–6), orbicular to broadly ovate, concave, green colour, 4–5 x 3.5–4.5 mm apex  
310 obtuse. Corolla apex slightly exerted from calyx, petals rhombic or spatulate. Male flowers (Fig.  
311 1C). Petals 5(–6), white, rhombic c. 5 x 3 mm, apex obtuse-acute, base cuneate, margins  
312 densely ciliate, hairs 0.3 mm long, outer surface glabrous, inner surface glabrous in distal half,  
313 proximal half compressed funneliform with ventral appendage adnate at margins, retuse (notched)

314 for 0.5 mm at midline, adaxial surface moderately densely hairy, hairs c. 0.3 mm long. Extra-  
315 staminal disc torus-like, glabrous, irregular, outer wall convex, lacking constrictions or teeth with  
316 c. 15 poorly defined lobes, 2.5–3 mm wide, c. 0.8 mm high. Stamens c. 15, erect, slightly  
317 exerted by 1–2 mm at anthesis, c. 5–6.5 mm long; filament 4–5 mm long, straight, densely  
318 puberulent the entire length (Fig. 1D); anthers yellow, ovate-ellipsoid, 1–1.3 mm long. Ovary  
319 (vestigial, Fig. 1E) bilobed, c. 1 x 1.5 mm densely appressed hairy, hairs c. 0.5 mm; style 0.7 mm  
320 long, glabrous.

321

322 Female flowers (Fig. 1G), with sepals and petals as the male flowers, but petals c. 6 x 2.6–2.9  
323 mm, usually detaching with a stamen attached, probably due to interlocking hairs (see Fig. 1 J),  
324 proximal two-thirds claw-like, c. 0.7 mm wide, margin sparsely and irregularly ciliate; ventral  
325 appendage with apex deeply bilobed, lobes c. 1 mm x 1 mm; disc as in male flower. Stamens c.  
326 10 (see Fig. 1I), included at anthesis, filament c. 2.5 mm long, proximal half to quarter glabrous,  
327 distal part densely hairy; anther as male flowers but indehiscent; ovary bilobed (see Fig. 1 H),  
328 3.2 x 5 mm, indumentum as male flower, style c. 5 mm long, apical 1 mm, curved, surface  
329 papillate-minutely puberulent, apex subcapitate. Infructescence, of same dimensions as  
330 infructescence, erect. Fruit colour not recorded, probably yellow when ripe (as in other species of  
331 the genus), mericarps 1 or 2, transversely ellipsoid, c. 1.8 x 2.1 x 1.2 cm (hydrated), the surface  
332 leathery, shallowly and finely muricated, glabrous, mesocarp spongy and juicy, 1-seeded. Seed  
333 ellipsoid, c. 1.8 x 1.1 x 0.8 cm, testa thin, parchment like, endosperm absent, cotyledons fleshy.

334

335 **Phenology:** flowering in November-December; fruiting in February and April, immature fruit  
336 recorded in December and June.

337

338 **Local name and uses:** none are known. “Onana’s *Deinbollia* is suggested as a common name.

339

340 **Etymology:** The specific epithet of *Deinbollia onanae* means ‘of Onana’ commemorating Dr  
341 Jean-Michel Onana, currently Lecturer in Botany at University of Yaoundé I, Cameroon,  
342 champion of plant conservation in Cameroon, specialist in Sapindales (Burseraceae, author of  
343 Flore Du Cameroun Burseraceae (*Onana, 2017*), co-chair of the IUCN Central African Red List  
344 Authority for Plants, former Head of the National Herbarium of Cameroon (2005–2016), co-  
345 author of the Red Data Book of the Plants of Cameroon (*Onana & Cheek, 2011*) and the  
346 Taxonomic Checklist of the Vascular Plants of Cameroon Cameroon (*Onana, 2011*). He led field  
347 teams of YA staff working with those of K that resulted in the collection of several of the  
348 specimens of this species and personally collected this species in the field (*Onana 1600, K, YA*).

349

350 **Distribution & ecology:** known only from the Cameroon Highlands of Cameroon (one location  
351 in the adjoining Mambilla Plateau, Nigeria) Fig. 3. Upper submontane & montane evergreen  
352 forest, sometimes in gallery forest; (1200–)2050–2200 m alt.

353

354 Detailed information on the ecology of *Deinbollia onanae* (under the name *D. pinnata*) is  
355 available from several studies led by Hazel Chapman at Ngel Nyaki, the largest surviving forest  
356 in the Mambilla Plateau, a branch of the Cameroon Highlands that extends into Nigeria (see map,  
357 Fig. 3). At this submontane forest patch, area c.5.7 km<sup>2</sup>, 1588–1690 m altitude, *Deinbollia* is  
358 recorded as one of the 20 most abundant woody plant species, with 158.68 stems above 1 cm  
359 diam. per ha (*Abiem et al., 2020*). We consider that many and probably most of the smaller of

360 these numerous stems may not be the usually infrequent *D. onanae*, but the much smaller (0.8–  
361 3(–5) m tall) *D. oreophila* which at this altitude, over the border in Cameroon, is vastly more  
362 frequent in submontane forest (Cheek & Etuge 2009). In contrast, the 1970s the 1 ha  
363 enumeration plot at Ngel Nyaki (Chapman & Chapman 2001: 25–26) yielded five stems of  
364 “*Deinbollia* sp.” in the C strata (understorey trees 7–13 m high) with diameter at 1.3 m above the  
365 ground exceeding 14.5 cm, of which two exceeded 28 cm and one 40 cm. This is more consistent  
366 with frequencies observed in Cameroon for *Deinbollia onanae*. Moreover, many of the  
367 observations of animals feeding on *Deinbollia* at Ngel Nyaki have been made using binoculars of  
368 primates trained on the crowns of trees so are, indeed likely to be of the larger, less frequent *D.*  
369 *onanae* which is evidenced at this location by two herbarium specimens (see “additional  
370 specimens” below) while *D. oreophila* has not yet been so recorded. Studies on the dietary  
371 preferences of the rare Nigerian-Cameroon chimpanzee (*Pan troglodytes ellioti*) by Dutton &  
372 Chapman (2014) at Ngel Nyaki found that among the 52 plant species consumed mainly as fruit,  
373 *Deinbollia* was the 4<sup>th</sup> (wet season) or 5<sup>th</sup> (dry season) species preferred of the 17 tree species  
374 over 10 cm diameter at breast height that were identified as seeds from 495 fecal samples. This  
375 record is certainly *D. onanae* since *D. oreophila* does not form trunks of such large diameters  
376 (Cheek & Etuge, 2009). However, *Deinbollia* was found in only one of these fecal samples, in  
377 which 47 of its seeds were recorded, collected in February 2011 (Dutton & Chapman, 2014).  
378 Only 16 weaned individuals of chimpanzee are known at Ngel Nyaki. More numerous and so  
379 probably more effective at seed dispersal are putty-nosed monkeys (*Cercopithecus nictitans*).  
380 Studies by Chapman *et al.*, (2010) found that *Deinbollia* seeds are both swallowed, passing  
381 through the gut (average 2 per fecal sample) and are sucked and spat by the putty-nosed monkeys  
382 (averaging 5 seeds per spitting event). Germination takes place about 30 days later, starting  
383 earliest and with highest success (60–70%) in defecated seeds. Artificially cleaned seeds showed  
384 much lower success (c. 40%) and germinated slightly later, and the lowest success (c.35 %) and  
385 slowest germination of the three treatments were of those seeds that had been spat (Chapman *et*  
386 *al.*, 2010). In addition, leaves but not fruit of *Deinbollia* have been recorded as being consumed  
387 by tanzanian monkeys (*Chlorocebus tantalus tantalus*), but only in very low quantities (Agmen *et*  
388 *al.*, 2010). Studies of dispersal of seeds of about 40 Ngel Nyaki forest species up to 30 m into  
389 grassland from the forest edge using seed traps showed that *Deinbollia* was one of the small  
390 number of forest species that do not disperse seeds out of the forest, but that within forest,  
391 natural regeneration from seed does occur. The species has been classified as a shade-bearer and  
392 is not a pioneer (Barnes & Chapman, 2014). *Deinbollia* “*pinnata*” was one of three species of  
393 tree used to test the Janzen-Connell hypothesis at this site. Five hundred and seventy seedlings  
394 were raised and planted at distances of up to 25 m from 19 mature conspecific “mother” trees  
395 and monitored over three months. Predation was significantly higher closer to the mother trees  
396 than distant from them (c. 30% vs. 20%), but there was no support for Janzen-Connell effects in  
397 seedling height growth. About 80% of the seedlings survived, and they grew 4.5–5.5 cm over the  
398 3 months (Matthesius *et al.*, 2011). *Deinbollia* “*pinnata*” is one of 28 identified tree species used  
399 by chimpanzees as nesting trees at Ngel Nyaki, but is not among the preferred top five (Dutton *et*  
400 *al.*, 2016).

401

402 **Additional specimens: CAMEROON. South West Region, Mt Kupe, near main summit,**  
403 **immature fr., 26 June 1996, Cable 3386 (K000197863!, YA!);\_North West Region.**

404 *Bali Ngemba Forest Reserve, fr. April 2002, Onana 1600 (YA!); Mt Oku and the Ijim Ridge:*

405 *above Laikom, st. 21 Nov..1996, Cheek 8709 (K000337728! YA!); Dom, Kinjinjang Rock, st. 25*

406 Sept. 2006, *Cheek* 13436 (K000580433!; YA!); *ibid.* Forest Patch 1, fl. buds, 27 Sept. 2006,  
407 *Cheek* 13625 (K000580434!, MO!,US!, YA!); *ibid.*, Javelong Forest, st. 29 April 2005, *Pollard*  
408 1400 (K000580432!; YA!); **Adamaoua Region**, c. 120 km E of Ngaoundéré, 15 km NE of Belel,  
409 falls in Koudini River, alt.  $\pm$  1200 m, fl. 4 Dec. 1964, *W.J.J.O. & J.J.F.E. de Wilde, B.E.E. de*  
410 *Wilde-Duyffes* 4555 (K000593309!; K000593310!, WAG1269760! , YA). **NIGERIA. Taraba**  
411 **State, Mambilla Plateau**, Ngel Nyaki Forest Reserve, near camp, fr. 2 Dec. 2003, *H.M.*  
412 *Chapman* 481 (FHI, K!); *ibid.* female fl. 4 Dec. 2002, *H.M. Chapman* 484 (FHI, K!).

413

414 **Notes:** *Deinbollia onanae* first came to our attention in 2000 when completing the “Plants of  
415 Kilum-Ijim” (*Cheek et al., 2000*). Two specimens of *Deinbollia* matched no other and were  
416 named *Deinbollia cf. pinnata* (*Cheek et al., 2000*). In subsequent surveys this taxon was more  
417 explicitly referred to as a new species: *Deinbollia* sp. 2 (*Harvey et al., 2004, Cheek et al., 2004,*  
418 *Cheek et al., 2009*). However, the earliest known collection was made in 1964 (*W.J.J.O. &*  
419 *J.J.F.E. de Wilde, de Wilde-Duyffes* 4555(K)).

420 This species is remarkable for the very large number of pairs of unusually long and slender  
421 leaflets, and for the comparatively large size of the individuals which often 10-15 m in height,  
422 among the largest trees known in the genus.

423

424

425 **Conservation:** *Deinbollia onanae* is rare at each of its six known locations so far as is known,  
426 although at Ngel Nyaki this is difficult to establish due to potential confusion with *Deinbollia*  
427 *oreophila*. Despite many thousands of herbarium specimens being collected at Kilum-Ijim, at  
428 Mt Kupe and the Bakossi Mts, at Ngel Nyaki and at Bali Ngemba (*Cheek et al., 2000; Cheek et*  
429 *al., 2004; Harvey et al., 2006*) only two specimens of this species at two sites, were made at each  
430 of the first three locations and only one at the third location. Surveys at other sites in the  
431 Cameroon Highlands and elsewhere, e.g at Mt Cameroon and at the Lebialem Highlands, failed  
432 to find this species (*Cheek et al., 1996; Cable & Cheek 1998; Harvey et al., 2010; Cheek et al.,*  
433 *2011*). However, at Dom, where a targetted search for this species was made by the first author,  
434 three specimens were made, each representing single, isolated trees (*Cheek et al., 2010*). No  
435 more individuals than these were found. At Adamaoua it has only been collected once, and only  
436 a single tree was then noted (*W.J.J.O. & J.J.F.E. de Wilde, B.E.E. de Wilde-Duyffes* 4555(K)).  
437 None of these locations is formally protected for nature conservation. Tree cutting for timber  
438 and habitat clearance for agriculture has long been known to be a threat at all but the last of these  
439 locations (references cited above). Severe habitat fragmentation has resulted over many hundreds  
440 of years, forest patches being now distant from each other by tens of kilometres, isolated in  
441 oceans of cultivation and secondary fire-maintained grassland. Ecological evidence from Ngel  
442 Nyaki is that while *Deinbollia* regenerates in that forest patch, its primate dispersers do not, or  
443 seldom cross to other forest patches (Dutton & Chapman, 2014, see ecology notes above). We  
444 assess the area of occupancy of *Deinbollia onanae* as 34 km<sup>2</sup> using the IUCN preferred 4 km<sup>2</sup>  
445 cell size. Therefore, we assess this species as Endangered, EN B2ab(iii) using the *IUCN (2012)*  
446 standard. We suggest that this species be included in forest restoration plantings within its  
447 natural range to partly reverse its move to extinction. However, the likely large (c. 1 cm diam.),  
448 thin-walled seeds are probably recalcitrant, so not suitable for conventional seed-banking, and  
449 should not be allowed to be dried before sowing. Experience at Ngel Nyaki (Matthesius et al.,  
450 2011) shows that it is possible to raise hundreds of seedlings in nurseries and to establish them in  
451 natural forest.

452

453 **DISCUSSION**

454

455 The discovery of a threatened, new species to science from surviving natural habitat in the  
456 Cameroon Highlands is not unusual. At most of the six locations from which we here describe  
457 *Deinbollia onanae*, additional new or resurrected species to science, all highly localised, range-  
458 restricted and threatened with extinction, have been documented in recent years. At Mt Kupe for  
459 example, *Coffea montekupensis* Stoffelen (Stoffelen *et al.*, 1997) and more recently the new  
460 species and genus to science *Kupeantha kupensis* Cheek & Sonké (Cheek *et al.*, 2018a). At Bali  
461 Ngemba, *Leptonychia kamerunensis* Engler & K. Krause (Cheek *et al.*, 2013), *Psychotria*  
462 *babatwoensis* Cheek (Cheek *et al.*, 2009) and *Allophylus ujori* Cheek (Cheek & Etuge, 2009b), at  
463 Mt Oku and the Ijim Ridge *Kniphofia reflexa* Marais (Maisels *et al.*, 2000), *Scleria cheekii*  
464 Bauters (Bauters *et al.*, 2018), while at Dom, the endemic epiphytic sedge *Coleochloa domensis*  
465 Musaya & D.A Simpson (Musaya *et al.*, 2010). No additional such new species are known from  
466 the Adamaoua location, probably because it is less completely sampled than the preceding four.

467

468 However, *Deinbollia onanae* is exceptional among these aforementioned species in that it is a  
469 new species of tree predominantly of montane forest. The many other newly discovered for  
470 science, resurrected or rediscovered plant species of the Cameroon Highlands have been  
471 overwhelmingly either been herbs or shrubs or are derived from submontane habitats (800–2000  
472 m altitude).

473

474 **Montane Forest Trees of the Cameroon Highlands**

475

476 The Cameroon Highlands extend through four tropical African countries. Beginning in the south  
477 on the volcano island of Bioko (Equatorial Guinea) they continue on the mainland with the  
478 Mount Cameroon active volcano, heading NNE along a major fault, forming the ridges, plateaux  
479 and isolated peaks of the Bakossi Mts and Mt Kupe, Muanenguba, Bamboutos Mts, the Lebialem  
480 and Bamenda Highlands, Mt Oku, Tchabal Mbabo, then heading eastwards and forming the  
481 lower and drier Adamaoua Highlands which extend into the Central African Republic. Two  
482 westward extending arms from the central section in Cameroon extend into Nigeria, forming the  
483 Obudu and Mambilla Plateaux. The altitudinal division between montane and submontane forest  
484 is well-marked in the Cameroon Highlands. Most species of montane tree only occur above the  
485 2000 m contour and not below it (Table 3), while tree species from the submontane forest belt  
486 rarely exceed the 2000 m contour (Cheek *et al.*, 1996; Cheek *et al.*, 2000; Cheek *et al.*, 2004),  
487 although some species of tree, like *Deinbollia onanae* can occur on either side of the 2000 m  
488 contour. Since most of the Cameroon Highlands do not ascend above 2000 m alt., montane forest  
489 is not ubiquitous along their length. Moreover, even where altitude is sufficient to support it and  
490 where it formerly occurred, montane forest has seen massive clearance for agriculture, and has  
491 been totally lost at the Bamboutos Mountains of West Region Cameroon (Ngoufou, 1992). Indeed,  
492 the Bamenda Highlands of Cameroon, long since cleared of their montane forest, are now known  
493 in Cameroon as “The grasslands” because they are blanketed in secondary grassland, perpetuated  
494 by fire. It has been estimated that as much as 96.5% of the original montane forest of the  
495 Bamenda Highlands has been lost (Cheek *et al.*, 2000: 49-50). The tallest mountain in the range,  
496 Mt Cameroon (4040 m), despite its height and lack of human activity above 2000 m alt., has  
497 surprisingly little forest above this contour due to the free-draining nature of its predominantly

498 volcanic cinder substrate (*Thomas & Cheek, 1992; Cheek et al., 1996; Cable & Cheek, 1998*).  
499 The single largest block of montane forest that survives by far in the Cameroon Highlands is  
500 that at Mt Oku and the Ijim Ridge (Kilum-Ijim) where about 70 km<sup>2</sup> has been estimated to  
501 survive and to have a measure of protection. Here it extends from the 2000 m contour to the  
502 summit at 3011 m alt. (*Cheek et al., 2000: 20*). Elsewhere in the Cameroon Highlands, such as at  
503 Mt Kupe, Muanenguba, Bali-Ngamba, Ngel Nyaki and Dom, surviving patches of montane  
504 forest consists of only a few hectares.  
505

506 The tree species diversity of the montane forest of the Cameroon Highlands is low (28 species,  
507 see table 3) compared with submontane forest which has hundreds of species, and also in great  
508 contrast, montane forest contains few Cameroon Highland endemic tree species (only seven: 25%  
509 of the total, see table 3). The majority of the canopy contains even fewer species. It was  
510 estimated that just ten species made up 90% of the montane forest canopy at Mt Oku and the Ijim  
511 Ridge, three of which are endemics (*Cheek et al., 2000:20*). The majority of montane tree  
512 species of the Cameroon Highlands are widespread in montane forest in Africa (Afromontane)  
513 occurring also east of the Congo Basin in the rift mountains of East Africa and several, such as  
514 *Ilex mitis*, extend north to Ethiopia and south to South Africa. The East African montane forest is  
515 more species-diverse, and only a subset of its species extend west to the Cameroon Highlands,  
516 and an even smaller subset, just seven species, extend even further west from the Cameroon  
517 Highlands, to the Guinea Highlands (*Couch et al., 2019:54*).  
518

519  
520 The high altitudinal range of *Deinbollia onanae* is unrivalled west of the Congo basin by any  
521 other species of the genus. Elsewhere in Africa it is matched only by *Deinbollia*  
522 *kilimandscharica* Taub., of mountains from Ethiopia to Malawi, reported to achieve 2250 m  
523 elevation in Tanzania (*Davies & Verdcourt, 1998*). Most species of the genus in tropical Africa  
524 are lowland forest shrubs, in the Cameroon Highlands only *Deinbollia oreophila* also occurs  
525 regularly at altitude over 800 m, and is largely confined to the submontane forest band being  
526 recorded from (880–)1000–1900(–2050) m altitude where it is often relatively frequent (*Cheek &*  
527 *Etuge, 2009*). We postulate based on their shared morphological characters that these two may be  
528 sister species (see results above) that have segregated between two adjacent altitudinally based  
529 vegetation types in a similar way to certain clades of bird species in the Cameroon Highlands  
530 such as the Turaco (*Njabo & Sorensen, 2009*). This hypothesis needs testing. It would most  
531 readily done by a comprehensive species-level molecular phylogenomic study of *Deinbollia* as  
532 has been achieved in several other genera, such as *Nepenthes* L.f. (*Murphy et al., 2020*).  
533

534 The fruits of *Deinbollia onanae* are similar to those of other species of the genus, i.e., fleshy,  
535 indehiscent and large-seeded, suggesting that the now intermittent distribution of this species,  
536 along a line c. 570 km along peaks of the Cameroon Highland line, was likely due to dispersal in  
537 the gut by animals. Chimpanzees (*Pan troglodytes ellioti*) and putty-nose monkeys  
538 (*Cercopithecus nictitans*) are known to disperse the species at one location however these  
539 species do not cross from one forest patch to another, especially when as now these patches can  
540 be separated by tens of kilometres of secondary grassland. Formerly the range of *Deinbollia*  
541 *onanae* may have once been more continuous along the mountain range than today, but it was  
542 likely greatly reduced when forest was cleared for agriculture as reported above.  
543



544  
545  
546

## 547 CONCLUSIONS

548

549 Such cases as *Deinbollia onanae* underline the urgency for publishing further discoveries while  
550 species still survive, since threats to such rare species are often clear and current, putting these  
551 species at high risk of extinction. About 2000 new species of vascular plant have been  
552 discovered by science each year for the last decade or more (Cheek *et al.*, 2020). Until species  
553 are delimited and known to science, it is more difficult to assess them for their conservation  
554 status and so the possibility of protecting them is reduced (Cheek *et al.*, 2020). Documented  
555 extinctions of plant species are increasing, e.g. *Oxygyne triandra* Schltr. of Cameroon is now  
556 known to be globally extinct (Cheek *et al.*, 2018b) as is *Afrothismia pachyantha* Schltr. (Cheek *et*  
557 *al.*, 2019). In some cases species appear to be extinct even before they are known to science,  
558 such as *Vepris bali* Cheek, once sympatric with *Deinbollia onanae* at Bali Ngemba (Cheek *et al.*,  
559 2018c), and elsewhere, *Nepenthes maximoides* Cheek (King & Cheek, 2020). Most of the >800  
560 Cameroonian species in the Red Data Book for the plants of Cameroon are threatened with  
561 extinction due to habitat clearance or degradation, especially of forest for small-holder and  
562 plantation agriculture e.g. oil palm, following logging (Onana & Cheek, 2011). Efforts are now  
563 being made to delimit the highest priority areas in Cameroon for plant conservation as Tropical  
564 Important Plant Areas (TIPAs) using the revised IPA criteria set out in Darbyshire *et al.*, (2017).  
565 This is intended to help avoid the global extinction of additional endemic species such as the  
566 Endangered *Deinbollia onanae* which will be included in the proposed IPA s of Mt Kupe, Bali  
567 Ngemba, Kilum-Ijim and Dom.

568

569

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584

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**Table 1** (on next page)

Characters separating *Deinbollia onanae* from *D. grandifolia*, *D. maxima* and *D. pinnata*.

Characters taken from *Fouilloy & Hallé (1973)* and *Keay (1958)*.



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	<i>Deinbollia grandifolia</i>	<i>Deinbollia maxima</i>	<i>Deinbollia onanae</i>	<i>Deinbollia pinnata</i>
Leaves	(5-)7-10-jugate	4-6-jugate	(2-)8-11-jugate	(2-)5-9-(-12)-jugate
Leaf rachis adaxial surface	Rounded	Rounded	Flattened, with margins angled-winged, midline with raised ridge	Flattened, with margins angled-winged, midline with raised ridge
Indumentum of abaxial surface of leaflet	Glabrous, or with a few scattered inconspicuous hairs	Glabrous, or with a few scattered inconspicuous hairs	Softly hairy with numerous translucent, patent hairs	Glabrous, or with a few scattered inconspicuous hairs
Leaflet width	5-8 cm	6-8(-10) cm	(2.1-)2.5-4 cm	2.3-7.5(-10) cm
N°s pair of secondary nerves (distal leaflets)	12-14(-16)	8 - 10	(12-)17-18	6-12
Fruit breadth, indumentum	1.5 cm, glabrescent	Dimensions unknown, glabrescent	1.3-1.5 cm, tomentose	2 cm, glabrous

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**Table 2** (on next page)

The more significant differences between *Deinbollia onanae* and *Deinbollia oreophila*.

Data on *Deinbollia oreophila* from (Cheek & Etuge, 2009)

1 **Table 2:**  
 2 **Characters separating *Deinbollia onanae* and *Deinbollia oreophila*.**  
 3  
 4 Characters for *Deinbollia oreophila* taken from *Cheek & Etuge (2009)*.  
 5

	<i>Deinbollia oreophila</i>	<i>Deinbollia onanae</i>
Height at maturing	0.8–3(–5) m	(4–)5–10(–15) m
Stem indumentum	Glabrous	Simple hairy, sparse to dense, glabrescent.
Lenticels	Highly conspicuous, bright white, contrasting with epidermis	Inconspicuous, grey-brown, concolorous with epidermis
Length of leaves (flowering stems)	25–63 cm	(14–)60–70 cm
Number of leaflets per leaf (flowering stems)	(4–)6–8(–10)	(4–)16–23
Width of leaflets (flowering stems).	(3–)5.5–9(–10.2) cm	(2.1–)2.5–4 cm
N° secondary nerves each side of midrib	(7–)9–14(–17)	(15–)17–18
Indumentum of lower surface of leaf blade	Glabrous	Inconspicuously sparsely simple hairy on secondary nerves and midrib
Sepals	Orbicular, margins glabrous	Ovate, margins hairy
Petals	Oblong or obovate, base cuneate; adaxial appendage surface glabrous	Rhombic or spatulate, basal claw (stalk); adaxial appendage surface hairy
Staminal filaments of male flowers	Proximal half glabrous.	Entire length densely hairy.
Ovary of female flowers	3-lobed, surface with very sparse, stout hairs	2-lobed, densely hair with fine hairs
Altitudinal range	(880–)1000–2050 m	(1400–)2050–2200 m

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**Table 3**(on next page)

The 28 montane forest trees of the Cameroon Highlands.

Data mainly from *Cheek et al., (2000)*, updated with subsequent literature e.g. *Kenfack (2011)*, *Cheek & Ngolan (2007)*, *Cheek et al., (2017)* and *POWO (2019)*

1

Currently accepted species name	Former name used in Cameroon Highlands, if any (e.g. <i>Cheek et al., 2000</i> )	Endemic to Cameroon Highlands (Y/N)	Occurring also below 2000 m alt. (Y/N)	Species forming 90% of the canopy (Y/N)	Forest edge species = E Infrequent species = R
<i>Schefflera abyssinica</i> (A.Rich.)		N	N	Y	
<i>Schefflera mannii</i> (Hook.f.)		Y	N	Y	
<i>Prunus africana</i> (L.) Kalkman	<i>Pygeum africanum</i> Hook.f.	N	N	Y	
<i>Syzygium staudtii</i> (Engl.) Mildbr.	<i>Syzygium guineense</i> subsp. <i>bamendae</i> F.White	N	N	Y	
<i>Myrsine melanophloeos</i> (L.) Sweet	<i>Rapanea melanophloeos</i> (L.) Mez	N	N	Y	
<i>Diclis bambuseti</i> R.E.Fr.	<i>Arundinaria alpina</i> K.Schum.	N	N	Y	
<i>Carapa oreophila</i> Kenfack	<i>Carapa grandiflora</i> Sprague	Y	N	Y	
<i>Bersama abyssinica</i> Fresen.		N	N	Y	
<i>Ixora foliosa</i> Hiern		Y	N	Y	
<i>Clausena anisata</i> (Willd.) Benth,		N	Y	Y	
<i>Nuxia congesta</i> Fresen.		N	N	N	E
<i>Lasiosiphon glaucus</i> Fresen.	<i>Gnidia glauca</i> (Fresen.) Gilg	N	N	N	E
<i>Hypericum revolutum</i> Vahl subsp. <i>revolutum</i>		N	N	N	E
<i>Maesa lanceolata</i> G.Don		N	N	N	E
<i>Alsophila dregei</i> (Kunze) R.M. Tryon	<i>Cyathea dregei</i> Kunze	N	N	N	E
<i>Podocarpus latifolius</i> (Thunb.) Mirb.		N	Y	N	R
<i>Croton macrostachyus</i> Delile		N	Y	N	R

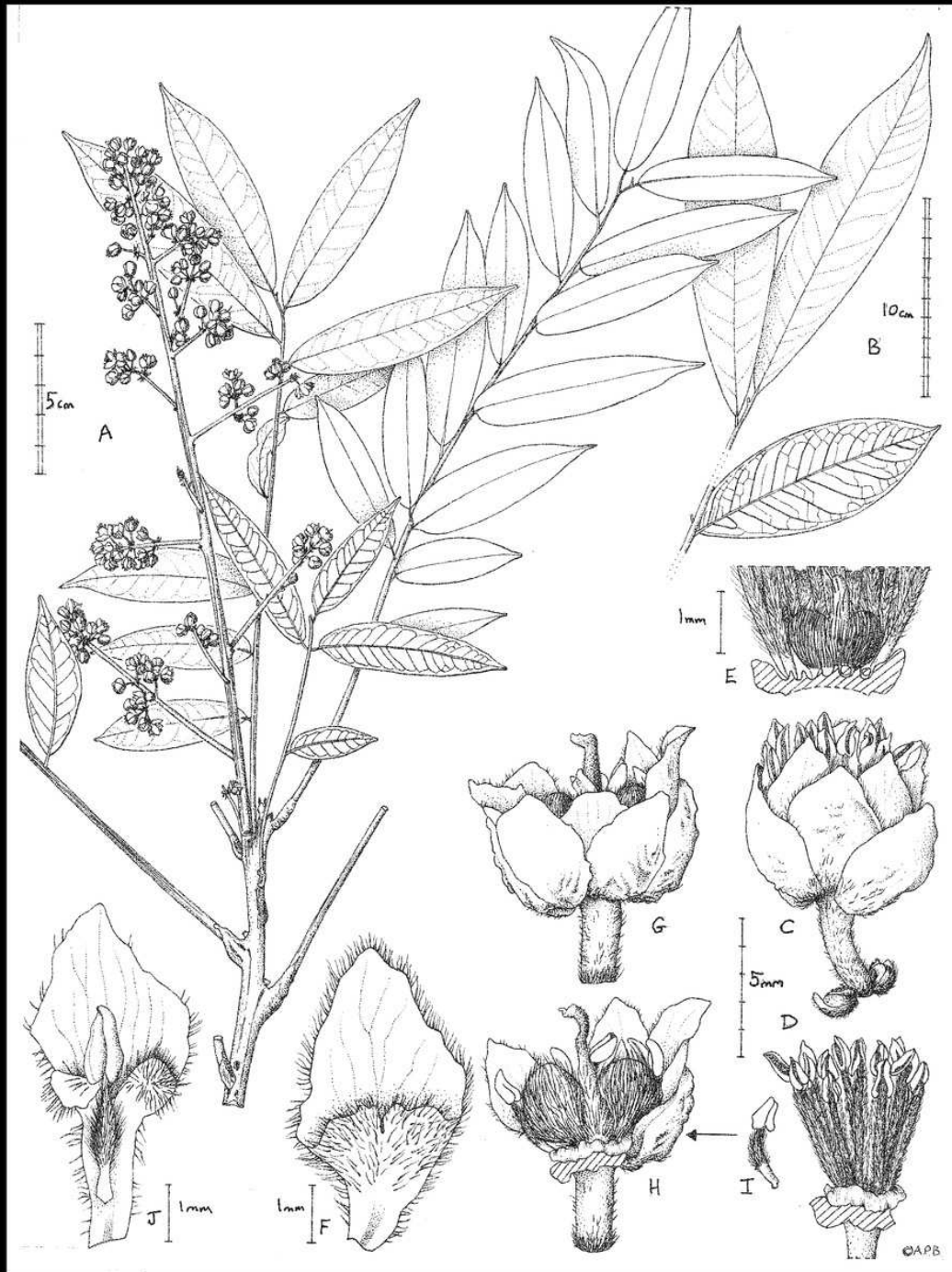
<i>Albizia gummifera</i> (J.F.Gmel)C.A.Smith		N	Y	N	R
<i>Cassipourea malosana</i> (Baker)Alston		N	N	N	R
<i>Brucea antidysenterica</i> J.F.Mill.		N	N	N	R
<i>Ilex mitis</i> (L.)Radlk.		N	N	N	R
<i>Neoboutonia mannii</i> Benth. & Hook.f.	<i>Neoboutonia glabrescens</i> Prain	N	Y	N	R
<i>Olea capensis</i> subsp. <i>macrocarpa</i> (C.H.Wright)I.Verd.	<i>Olea capensis</i>	N	N	N	R
<i>Eugenia gilgii</i> Engl. & Brehmer		Y	N	N	R
<i>Agarista salicifolia</i> (Lam.)G.Don	<i>Agaurea salicifolia</i> (Lam.)Oliv.	N	N	N	R
<i>Dovyalis cameroonensis</i> Cheek & Ngolan	<i>Dovyalis sp.nov.</i>	Y	N	N	R
<i>Ternstroemia cameroonensis</i> Cheek	<i>Ternstroemia polypetala</i> Melch.	Y	Y	N	R
<i>Deinbollia onanae</i> Cheek	<i>Deinbollia sp. 2</i>	Y	Y	N	R

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# Figure 1

*Deinbollia onanae*.

(A) habit, flowering branch; (B) detail from a large leaf showing apex and distal leaves (adaxial surfaces) and second leaf from the base (abaxial surface); (C) male flower lateral view; (D) male flower, petals and sepals removed to show the extra staminal disc and androecium; (E) base of D (male flower) showing the vestigial gynoecium and disc cut to show notches holding filament bases; (F) petal, adaxial surface, male flower; (G) female flower, lateral view; (H) female flower (with 3 sepals, 2 petals and anterior stamens removed) to show gynoecium and disc; (I) stamen from female flower; (J) petal, adaxial surface, of female flower with stamen. A, C-J from *de Wilde et al.* 4553 (K); B from *Cheek* 13436 (K). Drawn by Andrew Brown.





## Figure 2

*Deinbollia onanae*.Photo of the holotype: *Etuge* 3600 (holotypus K000593309). Photo by X van der Burgt.

## Figure 3

*Deinbollia onanae*. Global distribution map. By Xander van der Burgt.

By Xander van der Burgt.

