

1 **Two new sympatric species of *Phrynobius* (Anura:  
2 **Strabomantidae) from the Elfin Forests of Cordillera  
3 de Yanachaga in central Peru****

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6 Pablo J. Venegas<sup>1,2</sup>, Luis A. García-Ayachi<sup>1,2</sup>, Lesly Lujan<sup>2</sup>, Vilma Duran<sup>2</sup>, Ana Motta<sup>3</sup>

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8 <sup>1</sup> Rainforest Partnership, Austin, Texas, USA.

9 <sup>2</sup> Instituto Peruano de Herpetología, Lima, Peru.

10 <sup>3</sup> Biodiversity Institute and Natural History Museum, University of Kansas, Lawrence, USA.

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12 Corresponding Author:

13 Ana Motta

14 1345 Jayhawk Boulevard, Lawrence, Kansas, 66045-7561, USA

15 Email address: motta@ku.edu

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16 **Abstract**

17 We describe two new sympatric species of the terrestrial-breeding genus *Phrynobius* from elfin  
18 forests (3280 m a.s.l.) of the Cordillera Yanachaga, Yanachaga-Chemillén National Park, in  
19 central Peru. We used Maximum Likelihood approach to infer a molecular phylogeny on a  
20 dataset composed of 97 terminals, including three terminals representing the new species, and  
21 4271 bp of concatenated mtDNA and nuDNA fragments. Our phylogenetic analyses support the  
22 placement of the two new species in the genus *Phrynobius*. The first new species is a medium-  
23 sized species characterized by having upper eyelids bearing small tubercles, heel bearing one or  
24 two subconical tubercles, outer edge of tarsus bearing a row of broad conical tubercles, and red  
25 coloration in the groin, anterior and posterior surface of thighs, and concealed surface of shanks.  
26 The second new species is characterized by lacking tubercles on the heel and tarsus and by black  
27 coloration in the groin and the hidden surfaces of the hind limbs. Both new species are only  
28 known from the type locality and occur sympatrically with two other species of *Phrynobius* (*P.*  
29 *miroslawae* and *P. tribulosus*).

30

31 **Introduction**

32 Terrestrial breeding frogs of the family Strabomantidae are highly diverse, with more than 800  
33 species distributed in tropical and subtropical South America and lower Central America (Frost  
34 2024). Species of this group deposit their eggs in terrestrial sites *where they* undergo direct  
35 development, lacking the aquatic tadpole stage. This mode of life history, not associated with  
36 aquatic environments, *is* responsible for their success in inhabiting a variety of environments,  
37 from cloud forest to *humid* grasslands (Duellman & Lehr 2009; Frost 2024; Hedges et al. 2008).  
38 Strabomantid species comprise about half of all species of frogs known to inhabit Peru, *where*  
39 *they* are distributed in 12 genera (*Bryophryne*, *Lynchius*, *Microkayla*, *Niceforonia*, *Noblella*,  
40 *Oreobates*, *Phrynobius*, *Phyllonastes*, *Pristimantis*, *Qosqophryne*, *Strabomantis* and  
41 *Yunganastes*), occupying a variety of habitats such as Pacific dry forest, humid lowland  
42 tropical forests, montane forests, puna and paramo (Duellman & Lehr 2009; Frost 2024; von  
43 May et al. 2024).

44

45 The high Andean strabomantid frogs share a similar external morphology (the “phrynoboid”  
46 morphology) that led them to be historically considered part of a single natural group, the genus  
47 *Phrynobius* (De la Riva 2020). *Phrynobius* was re-erected *by Lynch (1975)* to include 14 species  
48 of small frogs, with short limbs and simple digits, broadly distributed in the high Andes from  
49 Colombia to Bolivia (Lynch 1975). Subsequently, the genus experienced a rapid increase in  
50 species *described* and drastic changes in its composition (De la Riva 2007; De la Riva et al.  
51 2008; Lehr 2006).

52

53 The monophyly of *Phrynobius* sensu Lynch (1975) was rejected by molecular phylogenetic  
54 analyses, revealing a scenario where high-elevation lineages have independent origins (Hedges et  
55 al. 2008). The non-monophyletic *Phrynobius* was split into a number of genera, and the genus

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61 *Phrynobius* was restricted to a clade of 21 species that occur in upper humid forests and  
62 grasslands of the Cordillera Oriental in Peru (Hedges et al. 2008). In the following decade, 16  
63 new species of *Phrynobius* were described and the redefined genus faced another increase in its  
64 number of species (Chaparro et al. 2008; Chávez et al. 2015; Lehr et al. 2012; Lehr & Rodríguez  
65 2017; Lehr et al. 2017; Mamani & Malqui 2014; Rodríguez & Catenazzi 2017; Trueb & Lehr  
66 2008; Venegas et al. 2018; von May et al. 2018). With *P. curator* and *P. nicoleae* being  
67 considered synonyms of *P. tribulosus* (von May et al. 2018), *P. ayacucho* (Lehr, 2007) being  
68 transferred to the genus *Oreobates* (Padial et al. 2012), and the recent description of *P.*  
69 *apumantarum*, *P. remotum*, and *P. sanctistobali*, the genus now comprises 37 species,  
70 distributed in the Cordillera Oriental and Central of Peru, restricted to a region between 6° and  
71 13° of latitude (Chávez et al. 2020; 2023; Diaz et al. 2023; Venegas et al. 2018). Species have  
72 been recorded at elevations between 2,600–4,490 m, and most of them show very restricted  
73 distributions, both horizontally and vertically (Chávez et al. 2023; Rodríguez & Catenazzi 2017).

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75 Even with the discovery of numerous new species resulting from recent fieldwork, the diversity  
76 of species of *Phrynobius* is still considered underestimated, as many remote regions of the slopes  
77 of the Andes in Peru have not yet been explored (De la Riva et al. 2018). The Yanachaga-  
78 Chemillén National Park in the Pasco Department of Peru has a remarkable diversity of  
79 amphibians (41 species have been recorded for the area, including undescribed species; Angulo  
80 et al. 2016), and fieldwork in the area has led to the discovery of new species of strabomantid  
81 frogs, including new species of *Phrynobius* (Duellman & Hedges 2005; Chaparro et al. 2008;  
82 Duellman & Hedges 2008; Hedges 1990; Lehr et al. 2012; 2017). Fieldwork in this region  
83 conducted by some of the authors of this study revealed the existence of two unnamed species of  
84 *Phrynobius*. We used phylogenetic analyses of nuclear and mitochondrial genes to assess their  
85 phylogenetic relationships and combine morphological and molecular data to support the  
86 recognition of the two species that we name and describe herein.

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## 88 Materials & Methods

### 89 Morphology

90 Character definition and terminology follow that of Duellman & Lehr (2009). Measurements,  
91 taken with digital calipers (to nearest 0.1 mm), are as follow: **SVL** (snout–vent length), **TL** (tibia  
92 length), **FL** (foot length, distance from proximal margin of inner metatarsal tubercle to tip of Toe  
93 IV), **HL** (head length, obliquely from angle of jaw to tip of snout), **HW** (head width, at level of  
94 angle of jaw), **ED** (eye diameter), **IOD** (interorbital distance), **EW** (upper eyelid width), **IND**  
95 (internarial distance), and **E-N** (eye–nostril distance, straight line distance between anterior  
96 corner of orbit and posterior margin of external nares). Fingers are numbered preaxially to  
97 postaxially from I–IV. Comparative lengths of Toes III and V were determined when both were  
98 adpressed against Toe IV; lengths of Fingers I and II were estimated when adpressed against  
99 each other. Specimens were preserved in 10% formalin and stored in 70% ethanol and were  
100 sexed externally by the presence or absence of vocal slits and internally by the condition of the

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104 gonads. All specimens were deposited in the herpetological collection of the Centro de  
105 Ornitología y Biodiversidad (CORBIDI), Lima, Peru. Institutional abbreviations follow Sabaj  
106 (2020). We obtained our research permit through the Dirección General Forestal y de Fauna  
107 Silvestre, Ministerio de Agricultura y Riego, Peru, which issued the Contrato de Acceso Marco a  
108 Recursos Genéticos, numbered 359-2013-MINAGRI-DGFFS-DGEFFS. Our research was  
109 approved by the Institutional Animal Care and Use Committee of University of Kansas (AUS  
110 279-01). Specimens examined are listed in Appendix 1.

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#### 111 DNA extraction and sequencing

112 We extracted total DNA from ethanol preserved tissues, following standard high-salt protocol  
113 adapted for microcentrifuge tubes (Lyra et al. 2017; Maniatis et al. 1982). We amplified two  
114 mitochondrial encoded gene fragments: one including the partial sequences of 12S rRNA,  
115 tRNA-val and 16S rRNA genes (H1 fragment) and a fragment of the cytochrome c oxidase I  
116 (COI); and two nuclear genes: partial sequences of tyrosinase (TYR), and partial sequences of  
117 recombination activating 1 (RAG-1). Amplifications were carried out in a 22  $\mu$ l reaction using  
118 Ampliqon Taq DNA Polymerase Master Mix (Amplicon A/S, Odense M, Denmark), with  
119 primers listed in Table S1. For the mitochondrial genes we followed polymerase chain reactions  
120 (PCR) conditions described in Lyra et al. (2017) (UP reaction protocol). For nuclear genes we  
121 used the following PCR cycling protocol: 3min of denaturation at 95°C, followed by 45 cycles of  
122 20s of denaturation at 95°C plus 20s of annealing at 56°C plus 1 min of extension at 68°C,  
123 followed by 3 min of final extension at 68°C, and stored at 12°C. PCR products were purified  
124 using an enzymatic reaction containing 1 unit of Exonuclease I and 0.5 unit of Alcaline  
125 phosphatase (Thermo Fisher Scientific Inc.) and were sent to Macrogen, Inc., Seoul, Republic of  
126 Korea, for sequencing. Sequence files were checked for quality and contigs were assembled  
127 using Geneious R11 (Biomatters).

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#### 128 Phylogenetic analysis

129 We used phylogenetic trees to assess generic assignment and investigate the relationship of the  
130 two new species. We chose the mitochondrial 12S rRNA and partial sequence of 16S rRNA  
131 genes, and the protein-coding gene cytochrome c oxidase subunit I (COI) as well as nuclear  
132 genes recombination-activating gene 1 (RAG1) and tyrosinase precursor (tyr) to perform our  
133 analyses. We supplemented our sequences with all sequences available on GenBank belonging to  
134 species of *Phrynoporus* and for which the chosen gene fragments were available. One new species  
135 is represented by the terminal CORBIDI 7379, and the other one by the terminals CORBIDI  
136 7382 and CORBIDI 7385. Our ingroup sample includes 63 terminals of *Phrynoporus* representing  
137 24 nominal species, two unnamed species (*Phrynoporus* spI and *Phrynoporus* sp. of von May et al.  
138 2018), and the two species we name herein. As outgroups we included one terminal per species  
139 of the related genera *Lynchiush* (n = 8) and *Oreobates* (n = 25), and rooted all our analyses with  
140 the distant species *Haddadus binotatus* (Padial et al. 2014). Specimen voucher numbers for  
141 newly produced sequences and accession numbers for all sequences used in this study are listed  
142 in Appendix 2.

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146 We performed multiple sequence alignments in MAFFT online v7 using the G-INS-i strategy,  
147 which is considered appropriate for alignments that consist of large numbers of sequences  
148 (Katoh & Standley 2013).

149  
150 We generated Maximum Likelihood (ML) phylogenograms using IQ-TREE v1.6 (Nguyen et al.  
151 2015) from the concatenated sequence of the five gene fragments included in our dataset. We  
152 determined the best-fit substitution model for each gene via ModelFinder, implemented within  
153 IQ-TREE (Kalyaanamoorthy et al. 2017) and performed a partitioned analysis according to  
154 codon position within the protein-coding genes (COI, RAG, and tyr). We calculated branch  
155 support with 10,000 bootstrap replicates using the Ultrafast Bootstrapping algorithm (Hoang et  
156 al. 2018). Alignments, script, and output files, including partitions and trees, are available as  
157 supplementary files.

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158 **Nomenclatural act**

159 The electronic version of this article in Portable Document Format (PDF) will represent a  
160 published work according to the International Commission on Zoological Nomenclature (ICZN),  
161 and hence the new names contained in the electronic version are effectively published under that  
162 Code from the electronic edition alone. This published work and the nomenclatural acts it  
163 contains have been registered in ZooBank, the online registration system for the ICZN. The  
164 ZooBank LSIDs (Life Science Identifiers) can be resolved and the associated information viewed  
165 through any standard web browser by appending the LSID to the prefix <http://zoobank.org/>. The  
166 LSID for this publication is: urn:lsid:zoobank.org:pub:513EDEF4-3F07-4097-8902-  
167 8DFED468E4C3. The online version of this work is archived and available from the following  
168 digital repositories: PeerJ, PubMed Central SCIE and CLOCKSS.

169

170 **Results**

171 **Phylogenetic relationships**

172 The optimal similarity-alignment of our concatenated dataset comprises 4271 character columns  
173 for 97 terminals. Our analysis recovered the genera *Phrynoporus* as monophyletic, with 100%  
174 bootstrap support and supported the placement of the two new species in the genus *Phrynoporus*  
175 (Fig. 1). The two new species are recovered as part of a highly supported clade (0.90) that  
176 includes *P. apumantarum*, *P. badius*, *P. barthlenae*, *P. bracki*, *P. bufooides*, *P. horstpauli*, *P. inti*,  
177 *P. kauneorum*, *P. miroslawae*, *P. pesantesi*, *P. sanctistobali*, *P. tautzorum* and *Phrynoporus* sp.  
178 The phylogenetic position and morphological distinctiveness of the newly collected specimens  
179 support the description of the two new species, which we name and diagnose below.

180

181 ***Phrynoporus manuelriosi* sp. nov.**

182 urn:lsid:zoobank.org:act:86453AE1-9A6A-4ABD-9681-CF9E372B877D

183 Figs. 2–4; Table 1

184 **Holotype**

186 CORBIDI 7385 (Fig. 2), adult female, Santa Bárbara, Distrito de Huancabamba, Provincia de  
187 Oxapampa, Región de Pasco, Perú, (10°20'29.1"S, 75°38'27.1"W, 3280 m a.s.l.), collected by  
188 Pablo J. Venegas, Vilma Duran, Caroll Z. Landrauro, and Lesly Lujan, on 25 August 2010.

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#### 189 **Paratypes (9)**

190 Nine specimens in total, four adult females (CORBIDI 7380, 7382-83, 7387) and five adult  
191 males (CORBIDI 7381, 7386, 7388-90) ~~same data as~~ the holotype.

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#### 192 **Diagnosis**

193 (1) ~~skin~~ on dorsum shagreen with scattered low tubercles, more abundant and prominent on  
194 flanks and hind limbs; ~~usually~~ bearing interorbital fold, \ /-shaped fold on scapular region, and /  
195 \-shaped fold on the middle of dorsum; skin on venter areolate; groins smooth or weakly  
196 areolate; dorsolateral folds absent; supratympanic fold conspicuous and long, slightly curved  
197 above the tympanic region; discoidal fold present only as thoracic fold or completely absent; (2)  
198 tympanic membrane and annulus absent; (3) snout moderately short, bluntly rounded in dorsal  
199 view and in profile; (4) upper eyelid bearing small tubercles, narrower than IOD; cranial crests  
200 absent; (5) vomerine teeth absent; (6) vocal slits present and nuptial pads absent; (7) Finger I  
201 shorter than Finger II; tips of fingers rounded and narrow; (8) fingers lacking lateral fringes;  
202 subarticular tubercles small and rounded in dorsal view, and flat on lateral view; (9) ulnar  
203 tubercle present, more evident in males; (10) heel bearing one or two subconical tubercles and  
204 outer edge of tarsus bearing a row of broad conical tubercles; inner tarsal fold absent; (11) inner  
205 metatarsal tubercle ovoid, about equal in size to rounded outer metatarsal tubercle; subarticular  
206 tubercles small and rounded in dorsal view, flat on lateral view; supernumerary plantar tubercles  
207 present; (12) toes lacking lateral fringes; webbing absent; Toe V longer than Toe III; tips of toes  
208 rounded; (13) in life, dorsum of head, body, and limbs pale brown, yellowish-brown or grayish-  
209 brown with dark brown markings; groin, anterior and posterior surface of thighs, and concealed  
210 surface of shanks red; ventral surface yellowish-brown, grayish-brown or yellow with or without  
211 dark brown flecks ~~on~~ the throat and belly; (14) SVL in five males 11.3-17.4 mm, in five females  
212 19.4-27.1 mm.

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#### 213 **Comparisons**

214 Among the 37 described species of *Phrynobius*, only *P. badius*, *P. bracki*, *P. daemon*, *P.*  
215 *heimorum*, *P. inti*, *P. paucari*, *P. peruanus*, *P. unchog* and *P. vestigiatus* show reddish coloration  
216 in the groins (Duellman & Lehr 2009; Lehr 2001; Lehr et al. 2012; Lehr & Oroz 2012; Lehr &  
217 Rodríguez 2017). *Phrynobius manuelriosi* sp. nov. differs from *P. badius*, *P. bracki*, *P. inti*, *P.*  
218 *paucari* and *P. vestigiatus* by having uniformly red groin (groin dark brown with bright orange  
219 flecks in *P. badius*; brown with red spots in *P. bracki*; pale grayish with salmon-colored flecks in  
220 *P. inti*; greenish yellow with diffuse salmon blotches in *P. paucari*; and dark brown with red  
221 well-defined blotches in *P. vestigiatus*). Of the species sharing a uniformly reddish coloration in  
222 the groin, *P. manuelriosi* sp. nov. can be distinguished by the presence of heel tubercle (absent in  
223 *P. daemon*, *P. heimorum*, and *P. peruanus*), tarsal tubercles (absent in *P. heimorum* and *P.*  
224 *peruanus*), and eyelid tubercles (absent in *P. daemon*, *P. heimorum*, *P. peruanus*, and *P.*

236 *unchog*). Moreover, *P. manuelriosi* sp. nov. lacks tympanic membrane and annulus, which are  
237 present in *P. peruanus*.

238 The presence of tubercles on the heel and outer edge of tarsus is uncommon in the genus  
239 *Phrynobius*. Only 6 species (*P. bracki*, *P. dagmarae*, *P. kotosh*, *P. oblivious*, *P. tribulosus* and *P.*  
240 *vestigiatu*s) share the presence of tubercles on the heel and outer edge of tarsus. *Phrynobius*  
241 *manuelriosi* sp. nov. differs from *P. bracki* and *P. dagmarae* by having tubercles on the upper  
242 eyelids (absent in *P. bracki* and *P. dagmarae*), and *P. dagmarae* has the Toe V shorter than Toe  
243 III, while the Toe V is larger than Toe III in the new species. *Phrynobius* *dagmarae*, *P. kotosh*  
244 and *P. oblivious* also differ by lacking \ /-shaped fold on scapular region and a / \ -shaped fold on  
245 the middle of the dorsum. Moreover, *P. manuelriosi* sp. nov. lacks dorsolateral folds, whereas  
246 dorsolateral folds are present in *P. dagmarae* (continuous), *P. kotosh* (discontinuous), and *P.*  
247 *vestigiatu*s (prominent and undulated). *Phrynobius* *tribulosus* has Toe V equal or slightly shorter  
248 than Toe III (von May et al. 2018), while in *P. manuelriosi* sp. nov. the Toe V is larger than Toe  
249 III. In addition, *P. barthlenae* and *P. miroslawae* have tubercles on the heel but lack tubercles on  
250 the outer edge of tarsus (present in *P. manuelriosi* sp. nov.) and Toe V is shorter than Toe III in  
251 *P. barthlenae*, and equal in length in *P. miroslawae*, while *P. manuelriosi* sp. nov. has Toe V  
252 longer than Toe III.

253  
254  
255 *Phrynobius horstpauli* is structurally similar to *P. manuelriosi* sp. nov., also being found in leaf  
256 and branches of the understory (Duellman & Lehr 2009; Lehr et al. 2000) and sharing the slender  
257 limbs and relative long narrow fingers and toes. *Phrynobius* *manuelriosi* sp. nov. can be easily  
258 distinguished from *P. horstpauli* by having one or two subconical tubercles on the heel and a row  
259 of conical tubercles on the outer edge of the tarsus (absent in *P. horstpauli*), and a smaller size  
260 with a SVL of 11.3 to 17.4 mm in males and 19.4 to 27.1 mm in females (17.7 to 25.6 mm in  
261 males and 30.8 to 39.7 mm in females of *P. horstpauli*).  
262

263 *Phrynobius melanoinguinis* sp. nov., described below, occurs in sympatry with *P. manuelriosi* sp.  
264 nov. and it can be easily distinguished by lacking heel and tarsal tubercles (present in *P.*  
265 *manuelriosi* sp. nov.), and by having dorsolateral and supratympanic folds (absent in *P.*  
266 *manuelriosi* sp. nov.).  
267

268 Due to the variable coloration of *P. manuelriosi* sp. nov., we consider possible to confuse it with  
269 *Noblella duellmani*, a geographically close species. *Noblella* *duellmani* occurs in Pasco, at  
270 elevations between 2900 and 3500 m, the same elevational range of *P. manuelriosi* sp. nov.  
271 *Noblella* *duellmani* can be easily distinguished from *P. manuelriosi* sp. nov. by having Toe V  
272 shorter than Toe III (Toe V longer than Toe III in *P. manuelriosi* sp. nov.), tips of digit slightly  
273 expanded and those of Toes III-V slightly acuminate (tips of toes narrow and rounded in *P.*  
274 *manuelriosi* sp. nov.), and skin of belly smooth (areolate in *P. manuelriosi* sp. nov.).  
275

### Description of holotype

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289 Adult female (Fig. 2, 3); body moderately robust; head about as wide as body, nearly as long as  
290 wide; snout bluntly rounded in dorsal view and in profile; canthus rostralis slightly curved in  
291 dorsal view, rounded in profile; loreal region nearly flat; lips rounded; nostrils barely  
292 protuberant, directed laterally; internarial region barely depressed; top of head flat; width of  
293 upper eyelid narrower than interorbital distance (EW/IOD 0.84); eye large, its diameter much  
294 greater than its distance from nostril (E-N/ED 0.70); tympanic membrane and annulus absent;  
295 supratympanic fold distinct, angling posteroventrally from ~~a~~ point behind the tympanic region  
296 close to the arm insertion; postorbital tubercles present, rounded. Tongue longer than broad,  
297 notched posteriorly, posterior half free; choanae small, round, not concealed by palatal shelf of  
298 maxillary; dentigerous processes of vomers absent.

299 Forelimb slender; ulnar tubercles low, diffuse; palmar tubercle low, round, about same size as  
300 thenar tubercle; subarticular tubercles distinct, small and rounded in dorsal view, and flat in  
301 lateral view; supernumerary tubercles present, weakly defined; fingers slender and long, lacking  
302 lateral fringes; relative lengths of fingers I<II<IV<III; tips of fingers narrow, rounded, lacking  
303 circumferential grooves. Hind limb long and slender; heel bearing two small subconical tubercles  
304 and tarsus bearing three low ~~conical tubercles~~, broad at ~~their~~ base; inner tarsal fold present on  
305 distal half of tarsus; inner metatarsal tubercle elevated, round, about twice the size of subconical  
306 outer metatarsal tubercle; subarticular tubercles distinct only at the base of toes, small and  
307 rounded in dorsal view, ~~flat~~ in lateral view; supernumerary tubercles present, distinct; toes  
308 slender, lacking lateral fringes; relative lengths of toes I<II<III<V<IV; tips of toes narrow,  
309 rounded, lacking circumferential grooves.

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310 Skin on dorsum shagreen with scattered low tubercles posteriorly, bearing an interorbital fold, a  
311 V-shaped fold on scapular region, and a  $\backslash$ -shaped fold on the sacrum; flanks and hind limbs  
312 tuberculate; upper eyelids bearing low rounded tubercles; skin on venter areolate; thoracic fold  
313 present; skin ventral and ventrolateral to cloaca granular.

314 Measurements (in mm) and proportions of holotype: SVL 26.1; TL 11.5; HW 9.3; HL 8.9; IOD  
315 3.1; IND 1.8; EW 2.6; FL 14.1; ED 2.7; E-N 1.9; TL/SVL 0.44; FL/SVL 0.54; HL/SVL 0.34;  
316 HW/SVL 0.36; HW/HL 1.04; E-N/ED 0.70; EW/IOD 0.84.

### 317 **Coloration of holotype in life**

318 Dorsal coloration of head, body, and limbs pale brown with pale and dark markings that include  
319 a dark brown interorbital bar with a cream border in the anterior margin, a brownish-cream V-  
320 shaped fold with a dark brown border in the posterior margin, one dark brown chevron with  
321 brownish-cream borders in the middle of the dorsum, one diagonal stripe on the flanks with a  
322 pale cream border, and two transverse bars on the hind limbs; dark brown head markings include  
323 a bold canthal and supratympanic stripes with cream borders, and a bold labial bar below the  
324 eyes with a cream border in the posterior margin (Fig. 2A); groin, anterior and posterior surface  
325 of thighs, and concealed surface of shanks red; ventral surface yellowish-brown with palms,  
326 soles and ventral surface of thighs brown (Fig. 2B); iris dark or light bronze with fine black  
327 reticulations and a faint reddish stripe across the middle.

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### 328 **Coloration of holotype in preservative**

335 Dorsum of head, body, limbs, and sides of head grayish-brown with the same dark brown  
336 markings and the same pale borders (now grayish-cream) described above; groin, anterior and  
337 posterior surface of thighs, and concealed surface of shanks grayish-brown; ventral surface on  
338 throat and chest pale tan, belly grayish-cream, with palm, soles, and limbs brown (Fig. 3).

#### 339 Variation

340 Sexual dimorphism is evident in respect to snout-vent length, with males smaller than females:  
341 SVL 11.3–17.4 mm in males and 19.4–27.1 mm in females (Table 1). In life, the ventral  
342 coloration of males and females can be variable from brownish-cream to yellowish-cream (Figs.  
343 4C, 4F, and 4J); only one male (CORBIDI 7381) has a grayish venter with dark brown flecks in  
344 the throat and belly (Fig. 4H). The dorsal coloration is variable: one female (CORBIDI 7380) has  
345 a dull brown dorsum without distinct markings except for a dark brown interorbital bar (Fig.  
346 4A); one female (CORBIDI 7383) has a fine pale middorsal stripe (Fig. 4E); one male  
347 (CORBIDI 7381) has a grayish-brown dorsum with dark brown blotches, a black labial bar  
348 below the eyes and lack canthal stripe (Fig. 4G); one male (CORBIDI 7381) has dark yellow  
349 flanks (Fig. 4L); one female (CORBIDI 7387) has an orange hue on dorsum with the markings  
350 similar to those of the holotype except for the canthal stripe (Fig. 4K); one male (CORBIDI  
351 7386) has a distinct black canthal and supratympanic stripe (Fig. 4I).

#### 352 Distribution and natural history

353 *Phrynobius manuelriosi* sp. nov. is only known from the type locality in the west margin of Río  
354 Huancabamba at an elevation of 3280 m a.s.l., Provincia de Oxapampa, Departamento de Pasco,  
355 on the eastern slope of Cordillera Oriental in central Peru (Fig. 5). Eighteen individuals, of which  
356 ten were collected, were found in 5 hours surveying amphibians at night by four collectors. All  
357 individuals were found on the ground in the elfin forest, and perched on leaves and branches  
358 about 20–100 cm above the ground in the forest and in the riparian vegetation. The  
359 herpetological survey in Santa Bárbara occurred in the dry season and no rainfall was recorded  
360 during the four days of survey in this locality. The two new species described herein were found  
361 in sympatry in the elfin forest of Santa Bárbara. *Gastrotheca griswoldi* also occurs in the area,  
362 but above tree line in the Puna grasslands. Two other species of *Phrynobius* are known to occur in  
363 the same area of Santa Bárbara, *P. miroslawae* and *P. tribulosus*, although we did not observe  
364 them during our surveys. *Phrynobius miroslawae* is found in the elfin forest and might be  
365 syntopic with *P. manuelriosi* sp. nov., whereas *P. tribulosus* inhabits the Puna grasslands  
366 (Chaparro et al. 2008). The sympatric *P. auriculatus* and *P. bracki* occur 38 km airline SE of  
367 Santa Bárbara, 5.5 km E Oxapampa 2600 m, on mountains at the opposite side of the Oxapampa  
368 valley (Chaparro et al. 2008), while while *P. badius* is closer at the eastern margin of Río  
369 Huancabamba (20 km airline SE of Santa Bárbara).

#### 370 Etymology

371 The name is a patronym for Manuel Ríos, a Peruvian forest engineer and professor at the Faculty  
372 of Forestry at Universidad Nacional Agraria La Molina (UNALM), Lima, Peru, from 1970 to  
373 2017, who has dedicated his life to preserving the natural heritage of his country. As professor,  
374 Manuel trained hundreds of students, inspiring them to become committed advocates for

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393 resource conservation, wildlife management and the protection of natural areas. He is also a  
394 founder and life member of the Board of Directors of the Peruvian Foundation for the  
395 Conservation of Nature (Pro Naturaleza), an organization that has played a key role in the  
396 preservation and protection of the environment in Peru. Likewise, he was Director of the  
397 Conservation Data Center (CDC-UNALM) between 1983 and 1998, and his legacy is present in  
398 the creation and planning of some of the most emblematic protected areas in Peru: the Paracas  
399 National Reserve, the Titicaca National Reserve, the Lachay National Reserve, the Abiseo  
400 National Park, and the Tabaconas Namballe National Sanctuary, among many others.

401

402 ***Phrynobius melanoinguinis* sp. nov.**

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404 Figs, 6–7

405 **Holotype**

406 CORBIDI 7379 (Fig. 6), adult female, Santa Bárbara, Distrito de Huancabamba, Provincia de  
407 Oxapampa, Región de Pasco, Peru (10°20'29.1''S, 75°38'27.1''W, 3280 m a.s.l.), collected by  
408 Pablo J. Venegas on 25 August 2010.

409 **Diagnosis**

410 (1) Skin on dorsum smooth with scattered granules; flanks and venter areolate; dorsolateral folds  
411 present, short; discoidal fold present only as thoracic fold; supratympanic fold conspicuous and  
412 long, slightly curved above the tympanic region; (2) tympanic membrane and annulus absent; (3)  
413 snout moderately short, bluntly rounded in dorsal view and in profile; (4) upper eyelids narrower  
414 than IOD, bearing low small tubercles; cranial crests absent; (5) vomerine teeth absent; (6) males  
415 unknown; (7) Finger I shorter than Finger II; tips of fingers rounded and narrow; (8) fingers  
416 lacking lateral fringes; subarticular tubercles small, rounded, weakly defined in dorsal view and  
417 flat in lateral view; supernumerary tubercles present, weakly defined; (9) ulnar tubercles absent;  
418 (10) heel and outer edge of tarsus lacking tubercles; inner tarsal fold absent; (11) inner metatarsal  
419 tubercle ovoid, prominent, about equal in size to lower, rounded, outer metatarsal tubercle;  
420 subarticular tubercles small, round, distinct only at the base of toes in dorsal view and flat on  
421 lateral view; supernumerary plantar tubercles absent; (12) toes lacking lateral fringes; webbing  
422 absent; Toe V slightly longer than Toe III; tips of toes rounded; (13) in life, dorsum dark brown  
423 without marks; groin, anterior and posterior surface of thighs, and concealed surface of shanks  
424 black; ventral surface brown with a black blotch on the throat; iris bluish-gray with fine black  
425 reticulation; (14) SVL of single female 23.6 mm.

426 **Comparisons**

427 *Phrynobius melanoinguinis* sp. nov. is strikingly different from any other known species in the  
428 genus by having the groin and the hidden surfaces of the hind limbs black, and bluish-gray iris.  
429 *Phrynobius melanoinguinis* sp. nov. occurs in sympatry with *P. manuelriosi* sp. nov., *P.*  
430 *miroslawae*, and *P. tribulosus*, and it differs from those by lacking heel and tarsal tubercles and  
431 by having dorsolateral and supratympanic folds (absent in *P. manuelriosi* sp. nov. and *P.*  
432 *tribulosus*). Three other species of *Phrynobius* occur within the Yanachaga-Chemillén National

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445 Park: *P. auriculatus*, *P. badius*, and *P. bracki* (Chaparro et al. 2008; Duellman & Lehr 2009;  
446 Lehr & Oroz 2012). *P. melanoinguinis* sp. nov. differs from those species by lacking heel and  
447 tarsal tubercles (heel tubercle present in *P. auriculatus* and *P. bracki*; tarsal tubercle present in *P.*  
448 *bracki*), by having eyelid tubercles (absent in *P. auriculatus*, *P. badius*, and *P. bracki*), and by  
449 having dorsolateral (absent in *P. bracki*) and supratympanic folds (absent in *P. badius*, and *P.*  
450 *bracki*).

451

452 The absence of heel and tarsal tubercles also distinguishes *P. melanoinguinis* sp. nov. from *P.*  
453 *daemon*, *P. dagmarae*, *P. interstinctus*, *P. vestigiatus*, *P. kotosh*, *P. oblivious*, *P. remotum*, and *P.*  
454 *unchog*. The combination of dorsolateral and long supratympanic folds also differentiates *P.*  
455 *melanoinguinis* sp. nov. from many other species in the genus: *P. barthlenae*, *P. bufoides*, *P.*  
456 *capitalis*, *P. chaparroi*, *P. daemon*, *P. dagmarae*, *P. heimorum*, *P. interstinctus*, *P. inti*, *P.*  
457 *julinensis*, *P. kauneorum*, *P. lapidooides*, *P. lechriorhyncus*, *P. montium*, *P. oblivious*, *P.*  
458 *peruanus*, *P. pesantesi*, *P. remotum*, *P. sanctistobali*, *P. tautzorum*, *P. thompsoni*, *P. valquii*, and  
459 *P. vestigiatus*. In the case of *P. bufoides* and *P. sanctistobali*, both species are also easily  
460 distinguished from *P. melanoinguinis* sp. nov. by the presence of conspicuous large round or  
461 elongate pustules on dorsum and flanks (dorsum smooth with areolate flanks in the new species).  
462 Moreover, the absence of tympanic membrane and annulus distinguishes the new species from *P.*  
463 *auriculatus*, *P. peruanus* and *P. mariellaeo* (tympanic membrane and annulus present).

464

465 *Phrynoporus paucari* differs from *P. melanoinguinis* sp. nov. by having larger subconical tubercles  
466 forming discontinuous longitudinal ridges dorsolaterally (dorsal skin smooth) and venter  
467 greenish-yellow with brown reticulation (dull brown). *Phrynoporus pesantesi* differs from *P.*  
468 *melanoinguinis* sp. nov. by having ulnar tubercles (absent) and the venter brown with gray  
469 mottling (dull brown). *Phrynoporus horstpauli* differs from *P. melanoinguinis* sp. nov. by having  
470 the skin of dorsum slightly tuberculate (smooth), Toe V much longer than Toe III (Toe V slightly  
471 longer than Toe III), and venter cream with brown blotches (dull brown). *Phrynoporus barthlenae*,  
472 *P. heimorum* and *P. tautzorum* differ from *P. melanoinguinis* sp. nov. by having Toe III larger  
473 than Toe V, while Toe III is shorter than Toe V in *P. melanoinguinis* sp. nov. Furthermore, the  
474 dorsum is coarsely tuberculate in *P. barthlenae* and *P. apumantarum*, and smooth in *P.*  
475 *melanoinguinis* sp. nov. In the case of *P. apumantarum*, this species also has the venter coarsely  
476 areolate, while in *P. melanoinguinis* sp. nov. is areolate peripherally and smooth in the center.  
477 *Phrynoporus kauneorum* and *P. lechriorhynchus* differ from *P. melanoinguinis* sp. nov. by the  
478 presence of dentigerous processes of vomers, whereas these are absent in *P. melanoinguinis* sp.  
479 nov.; in addition, *P. kauneorum* lacks dorsolateral fold (dorsolateral fold present in the new  
480 species), while *P. lechriorhynchus* has the snout spatulate, long and depressed, broadly rounded  
481 in dorsal view and sloping anteroventrally in profile (snout short and bluntly rounded in dorsal  
482 view and in profile in *P. melanoinguinis* sp. nov.).

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491 In addition, *P. personatus* from the Río Abiseo National Park (Departamento San Martín) in  
492 northern Peru is similar to *P. melanoinguinis* sp. nov. in that both species have the groins and  
493 hidden surfaces of hind limbs black (Rodríguez & Catenazzi 2017). However, *P. personatus* has  
494 the black surfaces of groins and hind limbs adorned by conspicuous white blotches and the skin  
495 of dorsum shagreen with scattered tubercles (dorsum smooth in *P. melanoinguinis* sp. nov.).

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#### 496 **Description of the holotype**

497 Adult female (Fig. 6, 7); body moderately robust; head narrower than body, nearly as long as  
498 wide; snout bluntly rounded in dorsal view and in profile; canthus rostralis slightly curved in  
499 dorsal view, rounded in profile; loreal region nearly flat; lips rounded; nostrils barely  
500 protuberant, directed laterally; internarial region flat; top of head flat; width of upper eyelid  
501 narrower than IOD (EW/IOD 0.74); eye large, its diameter greater than its distance from nostril  
502 (E-N/ED 0.63); tympanic membrane and annulus, absent; supratympanic fold conspicuous and  
503 long, slightly curved above the tympanic region; ovoid postocular tubercles present, minute.  
504 Tongue slightly longer than broad, not notched posteriorly, posterior half free; choanae small,  
505 rounded, not concealed by palatal shelf of maxillary; dentigerous processes of vomers absent.  
506

507 Forelimb slender; ulnar tubercles absent; palmar tubercle low, round, slightly longer than thenar  
508 tubercle; subarticular tubercles small and rounded in dorsal view, and flat on lateral view; two  
509 supernumerary tubercles present, weakly defined; fingers short and slender, lacking lateral  
510 fringes; relative lengths of fingers I<II<IV<III; tips of fingers narrow, rounded, lacking  
511 circumferential grooves. Hind limb slender; heel and tarsus lacking tubercles; inner tarsal fold  
512 absent; inner metatarsal tubercle prominent, round, about twice as much of round outer  
513 metatarsal tubercle; subarticular tubercles small, rounded, weakly defined in dorsal view and flat  
514 in lateral view; supernumerary tubercles absent; toes slender, lacking lateral fringes; relative  
515 lengths of toes I<II<III<V<IV; tips of toes narrow, rounded, lacking circumferential grooves.  
516 Skin on dorsum smooth with scattered low and round tubercles, flanks areolate, and hind limbs  
517 tuberculate; dorsolateral fold present and short; upper eyelids bearing low small tubercles; skin  
518 of throat and chest areolate, belly weakly areolate in the center and coarsely areolate  
519 peripherally; discoidal fold absent, thoracic fold present; skin ventral and ventrolateral to cloaca  
520 granular.

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521 Measurements (in mm) and proportions of holotype: SVL 23.6; TL 8.9; HW 8.7; HL 9.1; IOD  
522 2.7; IND 1.8; EW 2; FL 9.3; ED 2.7; E-N 1.7; TL/SVL 0.38; FL/SVL 0.39; HL/SVL 0.39;  
523 HW/SVL 0.37; HW/HL 0.96; E-N/ED 0.63; EW/IOD 0.75.

#### 524 **Coloration of holotype in life**

525 Dorsal surface of head, body and limbs, and flanks reddish-brown (Fig. 6A); anterior and  
526 posterior surface of thighs, groin, and concealed surface of tibia black (Fig. 6B); ventral surface  
527 pale reddish-brown, with a dark brown blotch on the throat (Fig. 6C); iris bluish-gray with fine  
528 black reticulation.

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#### 529 **Coloration of holotype in preservative**

536 Dorsum dark brown without marks; groin, anterior and posterior surface of thighs and concealed  
537 surface of shanks black; ventral surface brown with a black blotch on the throat (Fig. 7).

### 538 Distribution and natural history

539 *Phrynobius melanoinguinis* sp. nov. is known only from the type locality and is syntopic with *P.*  
540 *manuelriosi* sp. nov.. The type locality for both new species described here is within the Peruvian  
541 Yanachaga-Chemillén National Park (Fig. 5). The single specimen collected was found on mossy  
542 ground at night in elfin forest habitat. *Phrynobius melanoinguinis* sp. nov. might be syntopic  
543 with *P. miroslawae* and sympatric with *P. tribulosus*.

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### 544 Etymology

545 The specific name is an adjective derived from the Greek *melano* (meaning black) and the Latin  
546 *inguinis* (meaning groin) and is used as a noun in apposition. The name refers to the species'  
547 distinctive black groin.

548

### 549 Discussion

550 Most species of *Phrynobius* have a cryptic mode of life restricted to leaf litter and moss layers,  
551 making it difficult to find and observe individuals in the field (Lehr & Oroz 2012).  
552 Consequently, many species are known from a limited number of specimens, reflecting the rarity  
553 of some species in this genus (Lehr & Oroz 2012; Rodríguez & Catenazzi 2017), which leads to  
554 descriptions based on a few or even a single specimen, including our description of *P.*  
555 *melanoinguinis* sp. nov. The challenge of describing a species based on few specimens is related  
556 to the lack of understanding of the intraspecific variation in diagnostic characters (Köhler &  
557 Padial 2016). However, that can be mitigated by using multiple lines of evidence (e.g.  
558 morphological and molecular in the case of *P. melanoinguinis* sp. nov.) to name well-supported  
559 singleton species.

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560 While *Phrynobius melanoinguinis* sp. nov. is known only from the female holotype, the type  
561 series of *P. manuelriosi* sp. nov. is comparatively large (five females and five males). Limited  
562 number of male specimens seems to be recurrent in *Phrynobius*, as for three other species only  
563 females are known (*P. miroslawae*, *P. thompsoni* and *P. vestigiatus*), while in eight species only  
564 one male has been collected (*P. lapidoidea*, *P. unchog*, *P. anancites*, *P. capitalis*, *P. personatus*,  
565 *P. daemon* and *P. chaparroi*). This could represent actual female and male ratio in populations  
566 but could also be related to natural history aspects of the species. Even though little is known  
567 about the reproductive behavior of most species of *Phrynobius*, the “secretiveness of males” can  
568 explain why males are harder to find (Lehr 2001). For example, males of *P. bracki* have been  
569 reported to call from hidden places in leaf litter and moss vegetation (Hedges 1990) and males of  
570 *P. peruanus* were found calling from inside grass tussocks (Lehr 2007). Moreover, males of *P.*  
571 *badius* and *P. tribulosus* were heard calling, but not located in the dense vegetation, despite the  
572 collectors’ efforts (Lehr et al. 2012).

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586 The occurrence of two or more species of *Phrynobius* in the same locality was considered rare,  
587 but recent surveys have reported many cases where species of *Phrynobius* co-occur in the same  
588 region (see Chaparro et al. 2008; Chávez et al. 2015; Duellman & Hedges 2008; Lehr 2001; Lehr  
589 & Aguilar 2002; Lehr et al. 2002; Lehr et al. 2005; Lehr et al. 2012; Lehr & Oroz 2012; Lehr &  
590 Rodríguez 2017; Rodríguez & Catenazzi 2017; Von May 2017). In many cases, sympatric  
591 species segregate by elevation or by microhabitat type (Rodríguez & Catenazzi 2017).  
592 *Phrynobius manuelriosi* sp. nov. and *P. melanoinguinis* sp. nov. were found in sympatry on the  
593 ground in the elfin forest at 3280 m elevation, but show some fine habitat segregation, since only  
594 *P. manuelriosi* sp. nov. is known to perch on leaves and branches about 20–100 cm above the  
595 ground in the forest and in the riparian vegetation.

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596  
597 The description of two more endemic species of *Phrynobius* supports the idea that the genus  
598 shows a high regional diversity and endemism in the eastern slopes of the Andes in Peru (Fig. 8),  
599 and the region likely has many species still to be discovered (Rodríguez & Catenazzi 2017). The  
600 addition of the two new species described herein also increases the number of species of  
601 *Phrynobius* known from Cordillera Yanachaga to seven (*Phrynobius auriculatus*, *P. badius*, *P.*  
602 *bracki*, *P. manuelriosi* sp. nov., *P. melanoinguinis* sp. nov., *P. miroslawae*, and *P. tribulosus*).  
603 This region shows the highest regional species diversity of *Phrynobius*, along with Cordillera de  
604 Carpish (*P. daemon*, *P. dagmarae*, *P. interstinctus*, *P. kauneorum*, *P. lapidoides*, *P. unchog*, and  
605 *P. vestigiatus*), and followed by Río Abiseo National Park (*P. anancites*, *P. capitalis*, *P.*  
606 *dumicola*, *P. personatus*, *P. valqui*). Moreover, Santa Bárbara is the only case of a type locality  
607 shared by three species of *Phrynobius* (*P. manuelriosi* sp. nov., *P. melanoinguinis* sp. nov., and *P.*  
608 *miroslawae*). Santa Bárbara is located in the northwestern extreme of the Peruvian Yanachaga-  
609 Chemillén National Park (see Fig. 5) and, like in other regions throughout the Peruvian Andes,  
610 montane habitats are continuously destructed due to the increase of agricultural land and cattle  
611 ranching (Dillon et al. 1995; Venegas 2007; Weigend et al. 2005), which constitutes a serious  
612 threat to the species occurring therein. During our survey of amphibians and reptiles for four  
613 days in Santa Bárbara, we recorded several forest fires in the buffer zone of Yanachaga-  
614 Chemillén National Park, and we also found remains of forest fire in the grasslands and the tree  
615 line within the national park limits (an area deemed banned for resource use). Although both  
616 species are present within the park, evidence of habitat destruction in the area raises concerns  
617 about whether these species are truly protected. Many Peruvian species of high Andean  
618 strabomantid frogs in the genera *Bryophryne*, *Qosgophryne*, *Psychrophrynella*, *Phrynobius*, and  
619 some *Pristimantis*, have highly restricted distributions; thus, it is unlikely to find the new species  
620 species of *Phrynobius* described herein in other parts of the national park or its surroundings.  
621 Although we agree that it is important to keep records of species that are included in threatened  
622 categories lists (e.g., IUCN Red List, Peruvian government threatened species list) or in natural  
623 protected areas (Aguilar et al. 2010; von May et al. 2008), we believe that the fact that these  
624 species are legally protected will not grant their survival, especially if there are no means to  
secure the park boundaries from intrusions. This needs to be considered when classifying these

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646 frogs in species categories or analyzing if they will become a priority for protection in  
647 governmental conservation plans.

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## 649 **Conclusions**

650 We describe two new species of the Andean genus *Phrynobius*, *P. manuelriosi* sp. nov. and *P.*  
651 *melanoinguinis* sp. nov., based on robust morphological and molecular evidence. The new  
652 species occur sympatrically ~~in~~ the Andean elfin forest at an elevation of 3280 m, in the  
653 Yanachaga-Chemillén National Park in Pasco Department, Peru. Two other species of the genus,  
654 *P. miroslawae* and *P. tribulosus*, are also known to occur in the same locality. *Phrynobius*  
655 *melanoinguinis*, *P. miroslawae* and *P. tribulosus* are terrestrial, as is typical for most members of  
656 the genus, whereas *P. manuelriosi* possesses arboreal habitus. Despite their discovery within the  
657 boundaries of a national park, the long-term survival of these new species is not guaranteed if the  
658 borders of the protected area are not well protected. Fires observed both within the park and in its  
659 buffer zone ~~underscore~~ the urgent need for effective protection measures, particularly given the  
660 restricted distribution ranges characteristic of *Phrynobius* species. This highlights the  
661 vulnerability of these frogs and the critical importance of conserving their fragile habitats.

662

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672

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