

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

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16 **Abstract**

17 We describe two new sympatric species of the terrestrial-breeding genus *Phrynopus* 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 *Phrynopus*. 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 *Phrynopus* (*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*, *Phrynopus*, *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 “phrynopoid”
46 morphology) that led them to be historically considered part of a single natural group, the genus
47 *Phrynopus* (De la Riva 2020). *Phrynopus* 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 *Phrynopus* 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 *Phrynopus* was split into a number of genera, and the genus

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61 *Phrynopus* 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 *Phrynopus* 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. sancristobali*, 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; Díaz 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 *Phrynopus* 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 *Phrynopus* (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 *Phrynopus*. 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 adressed against Toe IV; lengths of Fingers I and II were estimated when adressed 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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gonads. All specimens were deposited in the herpetological collection of the Centro de Ornitología y Biodiversidad (CORBIDI), Lima, Peru. Institutional abbreviations follow Sabaj (2020). We obtained our research permit through the Dirección General Forestal y de Fauna Silvestre, Ministerio de Agricultura y Riego, Peru, which issued the Contrato de Acceso Marco a Recursos Genéticos, numbered 359-2013-MINAGRI-DGFFS-DGEFFS. Our research was approved by the Institutional Animal Care and Use Committee of University of Kansas (AUS 279-01). Specimens examined are listed in Appendix 1.

DNA extraction and sequencing

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

Phylogenetic analysis

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

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

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

Nomenclatural act

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

Results

Phylogenetic relationships

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

Phrynopus manuelriosi sp. nov.

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Figs. 2–4; Table 1

Holotype

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186 CORBIDI 7385 (Fig. 2), adult female, Santa Bárbara, Distrito de Huancabamba, Provincia de
 187 Oxapampa, Región de Pasco, Peru, (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. Landauro, and Lesly Lujan, on 25 August 2010.

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.

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.

213 **Comparisons**

214 Among the 37 described species of *Phrynopis*, 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). *Phrynopis 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.*

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236 *unchog*). Moreover, *P. manuelriosi* sp. nov. lacks tympanic membrane and annulus, which are
 237 present in *P. peruanus*.
 238
 239 The presence of tubercles on the heel and outer edge of tarsus is uncommon in the genus
 240 *Phrynopus*. Only 6 species (*P. bracki*, *P. dagmarae*, *P. kotosh*, *P. oblivious*, *P. tribulosus* and *P.*
 241 *vestigatus*) share the presence of tubercles on the heel and outer edge of tarsus. *Phrynopus*,
 242 *manuelriosi* sp. nov. differs from *P. bracki* and *P. dagmarae* by having tubercles on the upper
 243 eyelids (absent in *P. bracki* and *P. dagmarae*), and *P. dagmarae* has the Toe V shorter than Toe
 244 III, while the Toe V is larger than Toe III in the new species. *Phrynopus dagmarae*, *P. kotosh*
 245 and *P. oblivious* also differ by lacking \/-shaped fold on scapular region and a /\-shaped fold on
 246 the middle of the dorsum. Moreover, *P. manuelriosi* sp. nov. lacks dorsolateral folds, whereas
 247 dorsolateral folds are present in *P. dagmarae* (continuous), *P. kotosh* (discontinuous), and *P.*
 248 *vestigatus* (prominent and undulated). *Phrynopus tribulosus* has Toe V equal or slightly shorter
 249 than Toe III (von May et al. 2018), while in *P. manuelriosi* sp. nov. the Toe V is larger than Toe
 250 III. In addition, *P. barthlenae* and *P. mirosławae* have tubercles on the heel but lack tubercles on
 251 the outer edge of tarsus (present in *P. manuelriosi* sp. nov.) and Toe V is shorter than Toe III in
 252 *P. barthlenae*, and equal in length in *P. mirosławae*, while *P. manuelriosi* sp. nov. has Toe V
 253 longer than Toe III.
 254
 255 *Phrynopus 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. *Phrynopus 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 *Phrynopus 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; postrictal 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.
310 Skin on dorsum shagreen with scattered low tubercles posteriorly, bearing an interorbital fold, a
311 V-shaped fold on scapular region, and a / \-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.
328 **Coloration of holotype in preservative**

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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 *Phrynopus 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 Barbara 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 *Phrynopus* are known to occur in
363 the same area of Santa Bárbara. *P. mirosławae* and *P. tribulosus*, although we did not observe
364 them during our surveys. *Phrynopus mirosławae* 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 ***Phrynopus 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 *Phrynopus 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 *Phrynopus 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 *Phrynopus* 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. obliivius*, *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 *juninensis*, *P. kauneorum*, *P. lapidoides*, *P. lechriorhynchus*, *P. montium*, *P. obliivius*, *P.*
458 *peruanus*, *P. pesantesi*, *P. remotum*, *P. sancristobali*, *P. tautzorum*, *P. thompsoni*, *P. valquii*, and
459 *P. vestigiatus*. In the case of *P. bufoides* and *P. sancristobali*, 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. mariellaleo* (tympanic membrane and annulus present).

464

465 *Phrynopus 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). *Phrynopus pesantesi* differs from *P.*
468 *melanoinguinis* sp. nov. by having ulnar tubercles (absent) and the venter brown with gray
469 mottling (dull brown). *Phrynopus 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). *Phrynopus 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 *Phrynopus 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.).

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 postriotal 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.

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.

529 **Coloration of holotype in preservative**

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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 *Phrynopus 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. *Phrynopus melanoinguinis* sp. nov. might be syntopic
543 with *P. miroslawae* and sympatric with *P. tribulosus*.

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.

549 **Discussion**

550 Most species of *Phrynopus* 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.

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

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586 The occurrence of two or more species of *Phrynopos* in the same locality was considered rare,
 587 but recent surveys have reported many cases where species of *Phrynopos* 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 *Phrynopos manuelrosi* 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. manuelrosi* 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.
 596
 597 The description of two more endemic species of *Phrynopos* 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 *Phrynopos* known from Cordillera Yanachaga to seven (*Phrynopos auriculatus*, *P. badius*, *P.*
 602 *bracki*, *P. manuelrosi* sp. nov., *P. melanoinguinis* sp. nov., *P. mirosławae*, and *P. tribulosus*).
 603 This region shows the highest regional species diversity of *Phrynopos*, 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. valquii*). Moreover, Santa Bárbara is the only case of a type locality
 607 shared by three species of *Phrynopos* (*P. manuelrosi* sp. nov., *P. melanoinguinis* sp. nov., and *P.*
 608 *mirosławae*). 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*, *Qosqophryne*, *Psychrophrynella*, *Phrynopos*, and
 619 some *Pristimantis* have highly restricted distributions; thus, it is unlikely to find the new species
 620 species of *Phrynopos* 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
 625 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.

649 Conclusions

650 We describe two new species of the Andean genus *Phrynopus*, *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. mirosławae* and *P. tribulosus*, are also known to occur in the same locality. *Phrynopus*,
655 *melanoinguinis*, *P. mirosławae* 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 *Phrynopus* species. This highlights the
661 vulnerability of these frogs and the critical importance of conserving their fragile habitats.

663 Acknowledgements

664 For allowing access to herpetological collections, we are grateful to J. Córdova and C. Aguilar
665 (MUSM). We thank K. Siu-Ting and J.M. Padial for helpful comments on an earlier version of
666 the manuscript. We are grateful to the staff of the Servicio Nacional de Áreas Naturales
667 Protegidas por el Estado (SERNANP), especially the rangers and volunteers for their cooperation
668 and for the research permits. We also thank the staff of Consultores Asociados en Naturaleza y
669 Desarrollo (CANDES) and Willy Nañez of CORBIDI for the logistic support in the field, and
670 Daniel Matos and Abel Orihuela of Ministerio del Ambiente (MINAM) for the coordination with
671 the SERNANP.

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