

1 *Psychrophrynella glauca* sp. n., a new species of terrestrial-breeding frogs (Amphibia,
2 Anura, Strabomantidae) from the montane forests of the Amazonian Andes of Puno, Peru

3

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12 **Abstract**

13 We describe a new species of small strabomantid frog (genus *Psychrophrynella*) from a humid
14 montane forest in the Peruvian Department Puno. Specimens were collected at 2225 m a.s.l. in
15 the leaf litter of primary montane forest near Thiuni, along the Macusani–San Gabán road, in the
16 province of Carabaya. The new species is assigned to *Psychrophrynella* on the basis of
17 morphological similarity, including presence of a tubercle on the inner edge of the tarsus, call
18 composed of multiple notes, and genetic data (16S rRNA partial sequences) showing similarity
19 with *P. bagrecito*, *P. chirihampatu* and *P. usurpator*. *Psychrophrynella glauca* sp. n. is readily
20 distinguished from the three other species of *Psychrophrynella* by its small size (snout–vent
21 length of 11.3 mm in one male, 18.2 and 19.8 mm in two females), and by having belly and legs
22 reddish-brown or red, and chest and throat brown to dark brown with a profusion of bluish-grey
23 flecks. The new species is only known from its type locality, and has not been collected in
24 surveyed montane forests at similar elevations in the middle Inambari Valley (Puno) or in the
25 Marcapata Valley (Cusco). With the discovery of *P. glauca*, the geographic distribution of
26 *Psychrophrynella* is again extended to Department Puno, where the genus had disappeared
27 following the description of the genus *Microkayla*. Furthermore, the Cordillera de Carabaya is
28 the only mountain range known to be home to four of the seven genera of Holoadeninae
29 (*Bryophryne*, *Microkayla*, *Noblella*, and *Psychrophrynella*), suggesting an intriguing
30 evolutionary history for this group in southern Peru.

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33 **Key words:** 16S rRNA, bioacoustics, Carabaya, cloud forest, frog, leaf litter amphibian,
34 Ollachea, taxonomy

Comentado [EA1]: I suggest remove the new name from the abstract and should be firstly named in Taxonomic account ("Results")

Comentado [EA2]: See above comment

35 Introduction

36 Frogs in the genus *Psychrophrynella* are small, terrestrial-breeding terraranas that had
37 originally been placed with the genus *Phrynopus* (Lynch 1986). These high-Andean terraranas
38 are very difficult to characterize morphologically (Hedges et al. 2008). Molecular analyses later
39 revealed that these species were closely related to *Barycholos*, *Bryophryne* and *Holoaden* of the
40 subfamily Holoadeninae, and not *Phrynopus* and related forms within Strabomantinae (Hedges et
41 al. 2008). Thus, these forms were assigned to the new genus *Psychrophrynella*, with *P. bagrecito*
42 as the type species (Hedges et al. 2008). Until recently, the genus contained 23 species, four
43 species from the Peruvian Departments Cusco and Puno (Catenazzi & Tito 2016), and 19 from
44 Bolivia. Following the description of the genus *Microkayla*, which contains all Bolivian species
45 (and the southern Peruvian species *M. boettgeri*) formerly assigned to *Psychrophrynella* (De La
46 Riva et al. 2017), the genus *Psychrophrynella* presently contains three species: *P. bagrecito*, *P.*
47 *chirihampatu* and *P. usurpator*.

48 These three species of *Psychrophrynella* are Peruvian endemics restricted to the
49 Amazonian slopes of the Andes in the upper Kosñipata, Quespillomayo, Japumayo and
50 Marcapata valleys in the Department Cusco, where they inhabit humid grasslands and montane
51 forests from 1770 to 3600 m.a.s.l. (Catenazzi & Tito 2016; Duellman & Lehr 2009; von May et
52 al. 2017). These small frogs inhabit the leaf litter and the layer of terrestrial mosses, and thus
53 require considerable effort to be detected, for example through intensive search within leaf litter
54 plots (Catenazzi et al. 2011). Most of the eastern valleys of the Andes in the southern Peruvian
55 Departments Cusco and Puno have been poorly explored, with few locations surveyed by using
56 leaf litter plots, and are likely to contain many undescribed species of *Psychrophrynella* and
57 other Holoadeninae (Catenazzi & von May 2014).

Comentado [EA3]: For three or fewer authors, list all author names

58 The taxonomy of Holadeninae has undergone frequent changes over the past decade
59 (reviewed in De La Riva et al. 2017), in part because of our limited understanding of
60 phylogenetic relationships among its members. As new species are discovered, and our
61 understanding is accrued, it is likely that phylogenetic relationships will be revised again.
62 Furthermore, the type species *P. bagrecito* shares several morphological traits with the type
63 species of *Noblella*, *N. peruviana* (De la Riva et al. 2008; Lehr 2006), suggesting that the two
64 species might be closely related. Future molecular analyses of these two species will help resolve
65 relationships among species of *Noblella* and *Psychrophrynella*. Until DNA sequences of these
66 two type species become available, new species can be assigned to either genus on the basis of
67 overall morphological similarity and genetic distance with species whose genes have been
68 sequenced.

69 During a rapid survey of the amphibian fauna of several tributaries of the Inambari River
70 in Department Puno, we surveyed a humid montane forest in the Ollachea Valley. As a result of
71 opportunistic, intensive search of the leaf litter, we found four specimens and recorded the call of
72 a new species of Holoadeninae. Because the advertisement call sounded similar to the calls of *P.*
73 *chirihampatu* and *P. usurpator* (Catenazzi & Tito 2016), we suspected that the new species was
74 a *Psychrophrynella*. Here we describe this new species, and provide morphological and
75 molecular evidence for its generic allocation.

77 Methods

78 We follow Duellman & Lehr (2009) and Lynch & Duellman (1997) for the format of the
79 diagnosis and description, except that the term dentigerous processes of vomers is used instead
80 of vomerine odontophores (Duellman et al. 2006). We follow Heinicke et al. (2017) for

Comentado [EA4]: The genetic distances are not suitable for the generic assignment, the genetic distances by its intrinsic feature could saturated due that only four states are possible. Is possible that two taxa will be sister in genetic distance but not sister taxa in phylogenetic relationships

81 taxonomic arrangement of genera within subfamilies. We derived meristic traits of similar
82 species from the specimens examined, from species descriptions, and from published
83 photographs of live or preserved specimens.

Comentado [EA5]: Due the discussion about the arrangement of Terrarana, I suggest that you state that "follows to Heinicke et al 2017 in the general taxonomic arrangement" because the Strabomantidae family is not broadly recognized (e.g. De la Riva et al. 2017 Zootaxa 4363:350-360)

Eliminado: we

84 We fixed and preserved specimens in 70% ethanol. We determined sex and maturity of
85 specimens by observing sexual characters and gonads through dissections. We measured the
86 following variables (Table 1) to the nearest 0.1 mm with digital calipers under a
87 stereomicroscope: snout–vent length (SVL), tibia length (TL), foot length (FL, distance from
88 proximal margin of inner metatarsal tubercle to tip of Toe IV), head length (HL, from angle of
89 jaw to tip of snout), head width (HW, at level of angle of jaw), eye diameter (ED), tympanum
90 diameter (TY), interorbital distance (IOD), upper eyelid width (EW), internarial distance (IND),
91 and eye–nostril distance (E–N, straight line distance between anterior corner of orbit and
92 posterior margin of external nares). Fingers and toes are numbered preaxially to postaxially from
93 I–IV and I–V respectively. We determined comparative lengths of toes III and V by adpressing
94 both toes against Toe IV; lengths of fingers I and II were determined by adpressing these fingers
95 against each other. We describe variation in coloration on the basis of field notes and
96 photographs of live frogs. We deposited photographs of live specimens (taken by A. Catenazzi)
97 at the Calphoto online database (<http://calphotos.berkeley.edu>).

98 We recorded advertisement calls of an unvouchered male of *Psychrophrynella glauca* sp.
99 n. at the type locality on 14 August 2017, and recorded air temperature with a quick reading
100 thermometer (recording ##### deposited at the Fonoteca Zoológica, Museo Nacional de Ciencias
101 Naturales, Madrid, www.fonozoo.com). We used a digital recorder (Zoom H2, recording at 48
102 kHz, 24-bit, WAV format) for field recording, and Raven Pro version 1.4 (Cornell Laboratory of
103 Ornithology, Ithaca, NY) to analyze call variables. We analyzed a single call. We measured the

Comentado [EA6]: See above comment

Comentado [EA7]: Change "type locality" by the name of the site

105 following variables from the oscillogram: note and duration and rate, interval between notes or
106 calls, number of pulses, and presence of amplitude modulation. We measured these variables
107 from the spectrogram: dominant frequency, and presence of frequency modulation or harmonics.
108 Spectral parameters were calculated through fast Fourier transform (FFT) set at a length of 512
109 points (Hann window, 50% overlap). Averages are reported $\pm$ SE.

110 We estimated genetic distances to confirm generic placement of the new species within
111 *Psychrophrynella* through analysis of the non-coding 16S rRNA mitochondrial fragment. We
112 used liver tissue from all type specimens (Table 1) to obtain DNA sequences for the new species
113 (Appendix 1 with Genbank accession codes; FASTA supplementary file). We compared our
114 sequences with those of other species of *Psychrophrynella*, and with those of Holoadeninae
115 species in related genera (*Barycholos*, *Bryophryne*, *Holoaden*, *Microkayla* and *Noblella*) from
116 GenBank (Appendix 1). We extracted DNA with a commercial extraction kit (IBI Scientific,
117 Peosta, USA). We followed standard protocols for DNA amplification and sequencing (Hedges
118 et al. 2008). We used the 16Sar (forward) primer (5'-3' sequence:
119 CGCCTGTTTATCAAAAACAT) and the 16Sbr (reverse) primer (5'-3' sequence:
120 CCGGTCTGAACTCAGATCACGT) (Palumbi et al. 2002). For the polymerase chain reaction
121 (PCR) we used these thermocycling conditions: 1 cycle of 96°C/3 min; 35 cycles of 95°C/30 s,
122 55°C/45 s, 72°C/1.5 min; 1 cycle 72°C/7 min. We used a Veriti thermal cycler (Applied
123 Biosystems). We purified PCR products with Exosap-IT (Affymetrix, Santa Clara, CA), and
124 shipped purified samples to MCLAB (San Francisco, CA) for sequencing. We aligned sequences
125 using Geneious R8, version 8.1.6 (Biomatters, <http://www.geneious.com/>) with the MAFFT
126 v7.017 alignment program (Katoh & Standley 2013). We estimated uncorrected p-distances (i.e.,
127 the proportion of nucleotide sites at which any two sequences are different) with the R package

128 “ape” (Paradis et al. 2004).

129 We list specimens examined in Appendix 2. For codes of collections we used the
130 following acronyms: CORBIDI = Herpetology Collection, Centro de Ornitología y
131 Biodiversidad, Lima, Peru; KU = Natural History Museum, University of Kansas, Lawrence,
132 Kansas, USA; MHNC = Museo de Historia Natural del Cusco, Peru; MUBI = Museo de
133 Biodiversidad del Perú, Cusco, Peru; MHNG = Muséum d’Histoire Naturelle, Genève,
134 Switzerland; and MUSM = Museo de Historia Natural Universidad Nacional Mayor de San
135 Marcos, Lima, Peru.

136 Our research was approved by the Institutional Animal Care and Use Committee of
137 Southern Illinois University Carbondale (protocol #13-027). The Peruvian Ministry of
138 Agriculture issued the permit authorizing this research (permit #292-2014-MINAGRI-DGFFS-
139 DGEFFS).

140 The electronic version of this article in Portable Document Format (PDF) will represent a
141 published work according to the International Commission on Zoological Nomenclature (ICZN),
142 and hence the new names contained in the electronic version are effectively published under that
143 Code from the electronic edition alone. This published work and the nomenclatural acts it
144 contains have been registered in ZooBank, the online registration system for the ICZN. The
145 ZooBank LSIDs (Life Science Identifiers) can be resolved and the associated information viewed
146 through any standard web browser by appending the LSID to the prefix <http://zoobank.org/>. The
147 LSID for this publication is: urn:lsid:zoobank.org:pub:B3E69C6D-2669-46A8-A4C5-
148 631C5F1160B2. The online version of this work is archived and available from the following
149 digital repositories: PeerJ, PubMed Central and CLOCKSS.

150

151 **Results**

152 *Psychrophrynella glauca* sp. n. lsid:zoobank.org:act:E815EC45-81B4-46BF-A9A7-

153 3E359DEBDB73

154 <http://zoobank.org/NomenclaturalActs/E815EC45-81B4-46BF-A9A7-3E359DEBDB73>

155 **Figure**

156 **Holotype** (Figs. 1–3, Table 1). CORBIDI 18729, an adult female (Figs. 2, 3) from 13.67603S;

157 70.46588W (WGS84), 2225 m a.s.l., near Thiuni, Distrito Ollachea, Provincia Carabaya,

158 Departamento Puno, Peru, collected by A. Catenazzi and A. Tito on 14 August 2017.

159 **Paratypes** (Fig. 4, Table 1). Three total: one adult male, CORBIDI 18730, one adult

160 female, MUBI 16322, and one juvenile, MUBI 16323 collected at the type locality by A.

161 Catenazzi and A. Tito on 14 August 2017.

162 **Generic placement.** A new species of *Psychrophrynella* as defined by De La Riva et al.

163 (2017). Frogs of the genus *Psychrophrynella* are morphologically similar and closely related to

164 *Barycholos*, *Bryophryne*, *Holoaden*, *Microkayla* and *Noblella* (De La Riva et al. 2017; Duellman

165 & Lehr 2009; Hedges et al. 2008; Heinicke et al. 2007; Padial et al. 2014). The new species is

166 assigned to *Psychrophrynella* rather than any of the other genera on the basis of molecular data

167 (Table 2), call composed of multiple notes, and overall morphological resemblance with the type

168 species *P. bagrecito*, including presence of a short fold-like tubercle on the inner edge of tarsus.

169 Our genetic distance data (Table 2) support the generic placement of the new species within

170 *Psychrophrynella*. The most closely related species is *P. usurpator* (16S rRNA uncorrected p-

171 distance: 12.3–12.5%), followed by *P. chirihampatu* (12.5–12.7%). Species from other genera

172 had genetic distances above 18%, with the exception of *N. heyeri* (16.7%) and *N. myrmecoides*

173 (16.8%).

Comentado [EA8]: Move the figures to above

Comentado [EA9]: Above state m.a.s.l., will be constant

Comentado [EA10]: Why not only “same data as the holotype”?

Comentado [EA11]: I strongly suggest that the generic placement should be based in phylogenetic position mainly due that the morphological evidence is not strong (you stated), is non-informative, biogeographically is complex, and due that is logical to think that is *Microkayla* due that all species from Cordillera de Carabaya are *Microkayla*, etc

I performed a preliminar phylogenetic analyses with your genetic data and the available in GenBank and I am convinced that this species is a *Psychrophrynella* but you need convinced to the reader, please consider make a phylogenetic analysis, I understand that there are few sequences for *Psychrophrynella* but for me the most important is demonstrate that not is *Microkayla*

Con formato: Resaltar

Con formato: Resaltar

Con formato: Resaltar

174 **Diagnosis.** A species of *Psychrophrynella* characterized by (1) skin on dorsum smooth to
175 finely shagreen; skin on venter smooth, discoidal fold present; (2) tympanic membrane not
176 differentiated, anteroventral part of tympanic annulus visible below skin; (3) snout very short,
177 bluntly rounded in dorsal view and in profile; (4) upper eyelid lacking tubercles, narrower than
178 IOD; cranial crests absent; (5) dentigerous process of vomers absent; (6) vocal slits present;
179 nuptial pads absent; (7) Finger I slightly shorter than Finger II; tips of digits bulbous, not
180 expanded laterally; (8) fingers lacking lateral fringes; (9) ulnar tubercles absent; (10) heel
181 lacking tubercles; inner edge of tarsus bearing a short, oblique fold-like tubercle; (11) inner
182 metatarsal tubercle elliptical, of similar relief and length of prominent, ovoid, outer metatarsal
183 tubercle; supernumerary plantar tubercles absent; (12) toes lacking lateral fringes; webbing
184 absent; Toe V slightly shorter, or about the same length as Toe III; tips of digits not expanded,
185 weakly pointed; (13) dorsum reddish-brown to tan, with dark brown markings; one juvenile with
186 an orange middorsal line extending from tip of snout to cloaca and to posterior surface of thighs;
187 interorbital bar present; flanks brown with dark markings or entirely dark; chest dark brown with
188 bluish-grey flecks; throat and palmar and plantar surfaces grayish-brown with small, bluish-grey
189 flecks; belly and legs red or reddish-brown with bluish-grey flecks; (14) SVL 11.3 mm in one
190 male, 18.2 and 19.8 mm in two females.

191 **Comparisons.** The new species differs from the three known species of
192 *Psychrophrynella* by its unique combination of red coloration on legs and belly, and profusion of
193 bluish-grey flecks on ventral surfaces of head, body and legs. Morphologically, it is most similar
194 to *P. bagrecito* in having a short fold-like tubercle on the inner edge of tarsus, a prominent ovoid
195 outer metatarsal tubercle, discoidal fold present, an elliptical pupil, small size reaching ~19 mm,
196 and dark brown flanks in at least some specimens. It can be distinguished from *P. bagrecito*

Comentado [EA12]: Maybe you like re-wrote this

197 (characters in parenthesis) by having smooth skin on venter (areolate), dorsal coloration with
 198 broad markings (longitudinal stripes), snout short and bluntly rounded (snout moderately long,
 199 rounded in dorsal view and in profile), and ventral coloration in preservative brown with light
 200 grey flecks (white to cream with brown mottling). *Psychrophrynella glauca* sp. n. can be
 201 distinguished from *P. chirihampatu* by having reddish-brown to dark brown coloration and
 202 bluish-grey flecks on ventral parts (yellow coloration with reddish-brown or grey flecks). Finger
 203 I slightly shorter or same length as Finger II (Finger I shorter than Finger II), inner metatarsal
 204 tubercle the same length of outer metatarsal tubercle (inner metatarsal tubercle at least three
 205 times the size of outer metatarsal tubercle), more bluntly rounded head (slender and longer head),
 206 smaller size reaching 19.8 mm in females (27.7 mm in females?), and by the advertisement call
 207 having 26 notes and a fundamental frequency of 3027 Hz (up to 68 notes, 2712 Hz). The new
 208 species differs from *P. usurpator* by its reddish-brown ventral coloration (dull brown, gray or
 209 black with cream flecks), smaller SVL reaching 19.8 mm in females (SVL up to 30.5 mm in
 210 females?), and by the fold-like tubercle on the inner edge of tarsus being short (long and
 211 prominent tubercle).

212

213 **Description of holotype.** Adult female (18.2 mm SVL); head narrower than body, its
 214 length 34.6% of SVL; head slightly longer than wide, head length 108.6% of head width; head
 215 width 31.9% of SVL; snout very short, bluntly rounded in dorsal and lateral views (Fig. 2), eye
 216 diameter 31.7% of head length, its diameter 1.3 times as large as its distance from the nostril;
 217 nostrils not protuberant, close to snout, directed laterally; canthus rostralis slightly concave in
 218 dorsal view, convex in profile; loreal region flat; lips rounded; upper eyelids lacking tubercles;
 219 upper eyelid width 65.0% of interorbital distance; interorbital region flat, lacking cranial crests;

Comentado [EA13]: Who in parenthesis? Maybe you like think this and state the characteristics of *P. glauca* in parenthesis

Comentado [EA14]: Only the first time that you state the new name use sp. n.

Comentado [EA15]: There are not differences

Comentado [EA16]: Change by profile

Comentado [EA17]: Might want to think about how to reword this to improve flow.

220 eye-nostril distance 75.0% of eye diameter; supratympanic fold weak; tympanic membrane not
221 differentiated, anteroventral part of tympanic annulus visible below skin; postrictal tubercles
222 absent. Choanae round, very small, positioned far anterior and laterally, widely separated from
223 each other; dentigerous processes of vomers and vomerine teeth absent; tongue large, ovoid, not
224 notched.

225 Skin on dorsum smooth to finely shagreen; barely visible dorsolateral folds anteriorly;
226 skin on flanks and venter smooth; no pectoral fold, no discoidal fold; cloaca not protuberant,
227 cloacal region without tubercles. Ulnar tubercles and folds absent; palmar tubercle flat and oval,
228 approximately same length but twice the width of elongate, thenar tubercle; supernumerary
229 palmar tubercles absent; subarticular tubercles prominent, ovoid in ventral view, rounded in
230 lateral view, largest at base of fingers; fingers lacking lateral fringes; relative lengths of fingers 3
231 > 4 > 2 ≥ 1 (Fig. 3); tips of digits bulbous, not expanded laterally (Fig. 3); forearm lacking
232 tubercles.

233 Hindlimbs moderately long, tibia length 46.2% of SVL; foot length 45.1% of SVL; upper
234 and posterior surfaces of hindlimbs smooth; heel without tubercles; inner edge of tarsus bearing a
235 short, oblique fold-like tubercle, outer edge of tarsus lacking tubercles; inner metatarsal tubercle
236 elliptical, of similar relief and length of prominent, ovoid, outer metatarsal tubercle; minute
237 plantar supernumerary tubercles weakly defined; subarticular tubercles rounded, ovoid in ventral
238 view; toes lacking lateral fringes, not webbed; toe tips weakly pointed, not expanded laterally;
239 relative lengths of toes 4 > 3 > 5 > 2 > 1 (Fig. 3); foot length 45.1% of SVL.

240
241 **Measurements of holotype (in mm).** SVL 18.2, TL 8.4, FL 8.2, HL 6.3, HW 5.8, ED
242 2.0, IOD 2.0, EW 1.3, IND 1.8, E-N 1.5.

Comentado [EA18]: Consider move this phrase to above paragraph

Comentado [EA19]: Consider an initial phrase about the general form of the forelimbs

Comentado [EA20]: Change by profile

Comentado [EA21]: Why not use the Roman numbers?

Comentado [EA22]: What about supernumerary tubercles? I see one supernumerary tubercle in Finger III. (For me supernumerary tubercle are those between subarticular tubercles)

What about webbing in hands?

Pads? Grooves in tip Fingers?

Comentado [EA23]: Same as above

Comentado [EA24]: What about TY?

Con formato: Sangría: Primera línea: 1.27 cm

Eliminado: :

Con formato: Fuente:Negrita

Con formato: Fuente:Negrita

244

245 **Coloration of holotype in alcohol.** Dorsal surfaces of head, body, and limbs grayish tan,
246 with a dark brown X-shaped middorsal mark bordered laterally by a cream line. The interorbital
247 bar is a narrow dark stripe and is bordered anteriorly by a poorly defined cream stripe. There is a
248 dark brown stripe extending from the tip of the snout to above the tympanum and the insertion of
249 forelimb; furthermore, there are two longitudinal dark markings along the line separating the
250 dorsum from the flanks, and dark markings on each side of dorsum near the point of hind limb
251 insertion. The iris is dark gray. The throat is brown coloration anteriorly, fading into pale brown
252 with light grey flecks posteriorly. Chest and belly brown with light grey mottling and large
253 flecks. Ventral parts of limbs reddish-brown with cream mottling and flecks on brachium and
254 thighs, and tan with light grey flecks on antebrachium, crus and pes. The dorsal surfaces of hind
255 limbs have transverse dark bars. The posterior surfaces of thighs are reddish-brown with a large,
256 dark tan marking surrounding the cloaca and reaching one third the length of thigh, bordered
257 anteriorly by a narrow, cream stripe; the plantar and palmar surfaces are tan, fading into light
258 grey along fingers and toes.

259

260 **Coloration of holotype in life.** Similar to coloration in alcohol, but the dorsal coloration
261 varies from beige to brown, and the thighs are reddish-brown with brown mottling. Ventrally,
262 flecks are bluish-grey, largest and most noticeable on chest, and the belly and ventral surfaces of
263 limbs are red or reddish-brown. The iris is dark tan with golden flecks, forming a ring around the
264 pupil.

265

266 **Variation.** Coloration in life is based on field notes and photographs taken by A.
267 Catenazzi of the three **paratypes** (Fig. 4). These three **paratypes** have two subocular dark brown
268 spots, which are not visible in the holotype. Furthermore, all three have more extensive dark
269 coloration on flanks, either forming a nearly continuous dorsolateral line connected to the
270 supratympanic marking as in MUBI 16322, or having several dark markings as in CORBIDI
271 18730, or the entire flank dark as in MUBI 16323. The latter specimen, juvenile MUBI 16323
272 also has much darker dorsum than the other type specimens, as well as an orange middorsal line
273 extending from the tip of snout to the cloaca and to the posterior surface of thighs. The ventral
274 coloration of thighs varies from reddish-brown with brown mottling in the two females (the
275 holotype and MUBI 16322), to bright red with little mottling in the male, and orange with little
276 brown mottling in the juvenile.

277 **Advertisement call.** A single call of an unvouchered specimen was recorded at 19h45 on
278 14 August 2017 (Fig. 5). At a $T_{\text{air}} = 13.7^{\circ}\text{C}$, the advertisement calls lasted 2188 ms, and
279 consisted of 26 single-pulsed notes, produced at a rate of 11.88 notes/s. Low amplitude and poor
280 recording quality prevented analysis of the first five notes. In the remaining 21 notes, peak
281 frequency averaged 3027 ± 22 Hz (range 2756–3100 Hz) and increased during calls ($F_{1,19} = 21.3$,
282 $p < 0.001$); peak frequency averaged 2900 ± 29 Hz for the sixth to 11th note, and 3078 ± 16 Hz
283 for the last three notes of each call. Amplitude also increased during each call ($F_{1,19} = 6.7$, $p <$
284 0.017), reaching peak amplitude for notes that had highest frequency and longest duration (notes
285 19 through 21). Average note duration was 15.6 ± 2.4 ms (range 9–53 ms), and the 19th, 20th and
286 21st notes had longer duration (38.3 ± 7.9 ms) than all other notes (11.9 ± 1.0 ms).

287 **Etymology.** The specific name *glauca* is the feminine form of the Latin adjective
288 *glaucus*, from the ancient Greek noun *glauikos*, meaning “bluish-grey”, in reference to the bluish-
289 grey flecks on the ventral parts of body and limbs.

290

291 **Distribution, natural history and threats.** The four specimens were found in the leaf
292 litter along a descending ridge separating two creeks in the humid montane forest along the road
293 from Thiuni to Ollachea. Sympatric species detected during our quick survey included
294 *Gastrotheca testudinea*, *Pristimantis platydactylus*, and an undescribed *Pristimantis* sp. Much of
295 the original forest vegetation has been replaced by cultivated fields and pasture along the road,
296 but this remnant forest extended from nearly the side of road to the upper ridge of the mountain.
297 Further advance of agriculture, or clearing of forest might threaten this species if its distribution
298 is restricted to the Ollachea Valley. In absence of more detailed data regarding its extent of
299 occurrence, and according to the IUCN Red List criteria and categories (IUCN 2013), we suggest
300 this species to be in the “Data Deficient” category of the Red List.

301

302 **Discussion**

303 The diversity of small, terrestrial-breeding frogs in the humid grasslands and montane
304 forests of the Tropical Andes has until recently been grossly underestimated (De La Riva et al.
305 2017). A similar pattern has occurred in the Atlantic forest of Brazil, where the diversity and
306 micro-endemism of the minute terrestrial-breeding *Brachycephalus* was long unappreciated (Pie
307 et al. 2017; Pie & Ribeiro 2015; Ribeiro et al. 2017). In Peru, most species of Holoadeninae have
308 been described since 2008 (Catenazzi & Ttito 2016; [Catenazzi et al. 2015](#); Catenazzi et al.
309 2017b; [De la Riva et al. 2008](#); De La Riva et al. 2017; Lehr & Catenazzi 2008; Lehr & Catenazzi

Con formato: Sangría: Primera línea: 1.27 cm

Eliminado: Catenazzi et al. 2015;

Eliminado: De la Riva et al. 2008;

2009a; Lehr & Catenazzi 2009b; Lehr & Catenazzi 2010). Additional, undescribed species of *Psychrophrynella* have already been identified (Catenazzi et al. 2013; von May et al. 2017), and museum material indicates several more species might exist among misidentified specimens, such as in the type series of *P. bagrecito* (De la Riva et al. 2008; Duellman & Lehr 2009). Therefore, we can expect that additional field work, specimen comparisons, bioacoustics and genetic or genomic analyses will reveal many more species of *Psychrophrynella* and related Holoadeninae genera from the Tropical Andes.

De La Riva et al. (2017) recently allocated all Bolivian species previously assigned to *Psychrophrynella*, and the Peruvian species *boettgeri* from Department Puno, to the new genus *Microkayla*. Accordingly, the genus *Psychrophrynella* was left with only three species, *P. bagrecito*, *P. chirihampatu* and *P. usurpator* all distributed around the Vilcanota massif and its associated cordilleras in the Peruvian Department Cusco. Using phylogenomic approaches, Heinicke et al. (2017) validated the allocation of *Barycholos*, *Bryophryne*, *Euparkerella*, *Holoaden*, *Microkayla*, *Noblella* and *Psychrophrynella* (but not of *Niceforonia*, *Lynchi*, *Oreobates* and *Phrynopus*) within Holoadeninae; we follow their proposed taxonomic arrangement here. With the description of *P. glauca*, the geographic distribution of *Psychrophrynella* is again extended to Department Puno, and to a tributary of the Inambari River in the Cordillera de Carabaya. The Cordillera de Carabaya also contains the type localities of *Noblella peruviana*, the type species of *Noblella*, at Santo Domingo in the upper reaches of a small tributary of the Inambari River, of *Microkayla boettgeri* at Phara in Province Sandia, and of *Bryophryne togra* and *B. wilakunka* in Province Carabaya. The Cordillera de Carabaya is thus unique in being home to four of the seven genera of Holoadeninae: *Bryophryne*, *Microkayla*, *Noblella*, and *Psychrophrynella*. Only three genera are known to occur in the northern Cordillera

335 de Vilcanota and associated cordilleras (*Bryophryne*, *Noblella* and *Psychrophrynella*), and only
336 two genera in the northern Cordillera de Urubamba (*Bryophryne* and *Noblella*), and in the
337 southern Cordillera de Apolobamba (*Microkayla* and *Noblella*). Therefore, the Cordillera de
338 Carabaya appears to host substantial beta diversity of Holadeninae, suggesting an intriguing
339 evolutionary history for this group in southern Peru.

340 Our generic allocation remains tentative in light of an unresolved taxonomic situation
341 regarding *Noblella* and *Psychrophrynella*, as previously reviewed (Catenazzi & Tito 2016; De la
342 Riva et al. 2008; De La Riva et al. 2017). In short, the type species of both genera, which have
343 not been included in phylogenetic analyses for lack of DNA sequences, share several
344 morphological traits such that the hypothetical scenario of them forming part of the same clade is
345 a plausible hypothesis. Here we have assigned the new species to *Psychrophrynella* on the basis
346 of general body shape and appearance, overall similarity with the type species *P. bagrecito*, and
347 relatively small uncorrected p-distances of 16S rRNA and similarities in advertisement call with
348 *P. chirihampatu* and *P. usurpator*. As a priority, future work should sample tissues and record
349 advertisement calls of *P. bagrecito* and *N. peruviana*, so that multiples approaches can be
350 pursued to determine the phylogenetic relationships of species of *Noblella* and *Psychrophrynella*.

351 Anuran communities in the humid montane forests of southern Peru have undergone
352 sharp reductions in species richness and abundance following epizootics of chytridiomycosis
353 (Catenazzi et al. 2011; Catenazzi et al. 2014). The disease/host dynamics now seem to be
354 enzootic (Catenazzi et al. 2017a), and although experimental infection trials have shown that
355 terrestrial-breeding frogs can be highly susceptible to chytridiomycosis (Catenazzi et al. 2017a),
356 populations of Strabomantidae generally have not declined as sharply as those of aquatic-
357 breeding sympatric frogs. Thus, chytridiomycosis might not directly threaten *P. glauca*. A more

Eliminado: ; De la Riva et al. 2008

Comentado [EA25]: I understand your argument and agree that is necessary more evidence including other species and genera but I think that the generic placement should be the stronger possible with the available evidence

Con formato: Resaltar

359 immediate threat to *P. glauca* is embodied by hydroelectric projects that are planned or under
360 construction along the San Gabán River. The new dams might directly flood montane forest, or
361 intercept water from streams and rivers that drain the forest, thus reducing habitat quality. These
362 projects are part of many planned dams in the Inambari watershed that threaten to alter fish
363 migrations, biodiversity and geochemical cycles locally and downstream throughout the Amazon
364 basin (Forsberg et al. 2017; Latrubesse et al. 2017). These consequences that might not have
365 been properly taken into consideration during the decision making progress evaluating financial
366 interests and the findings of the Environmental Impact Assessment (Rode et al. 2015). We hope
367 that the timely description of new species such as *P. glauca* will contribute to the conservation of
368 the humid montane forest, and promote mitigating solutions including restoration of degraded
369 forest habitat.

370

371 **Conclusions**

372 We describe a new species of terrestrial-breeding frog of the family Strabomantidae, and provide
373 evidence for its allocation within the genus *Psychrophrynella*. The new species *P. glauca* is only
374 known from its type locality, similarly to most other small Holoadeninae known to occur at high
375 elevations in the Andes of southern Peru and Bolivia. With our description we contribute to a
376 better knowledge of the diversity of this group, and reveal the presence of four genera of
377 Holoadeninae in the Cordillera de Carabaya of southern Peru, suggesting that phylogeographic
378 studies of the Holoadeninae species of this mountain range may shed insights into radiation in
379 this group.

380

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385

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Comentado [EA26]: Please review the guide for Reference

Remove the connector "and"
Include the DOI for the references cited

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