

1 **On the Andean genus *Leschenius* (Coleoptera:
2 Curculionidae: Entiminae): Updated phylogeny, with
3 a new species from Ecuador, discovery of males, and
4 larval description of the potato weevil *Leschenius
5 vulcanorum***

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7 María Guadalupe del Río, Adriana Elena Marvaldi
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9 División Entomología, Museo de La Plata, FCNyM, UNLP, CONICET, La Plata, Buenos Aires,
10 Argentina.

11
12 Corresponding Authors:
13 María G. del Río
14 Paseo del Bosque s/n, La Plata, 1900, Buenos Aires, Argentina.
15 Email address: gdelrio@fcnym.unlp.edu.ar
16
17 Adriana Marvaldi
18 Paseo del Bosque s/n, La Plata, 1900, Buenos Aires, Argentina.
19 Email address: marvaldi@fcnym.unlp.edu.ar

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35 **Abstract**

36 The weevil genus *Leschenius* del Río (Curculionidae: Entiminae: Naupactini) is distributed in the
37 northern Andes, in Colombia and Ecuador. Among its species, *Leschenius* L.
38 *vulcanorum* stands out as an important pest of potatoes in its parthenogenetic form, known as
39 “tiroteador de la papa”. In this study, the adult male and the larval stage (first and mature larvae)
40 of *L. vulcanorum* are described and illustrated for the first time. A description of the male
41 of *Leschenius bifurcatus* is also provided. A new bisexual species was discovered, *Leschenius*
42 *ventrilingulatus* sp. nov., and is described from Ecuador. An updated phylogenetic analysis was
43 performed, including the new species, with results indicating a sister group relationship
44 between *Leschenius* L. *ventrilingulatus* sp. nov. and *L. vulcanorum*. They can be distinguished
45 because the former is usually of smaller size and is covered by denser and thicker setae, it has
46 shorter antennae, a subcylindrical shape of the pronotum, shorter elytra (about 1.5x longer than
47 wide at base), the female has ventrite 4 with a posterior rounded projection, and posterior margin
48 of ventrite 5 subacute, not excavated. This paper also includes lectotype designations, a revised
49 key to all known species of *Leschenius*, habitus photos of males and females, illustrations of
50 genitalia, and a distribution map.

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52

53 **Introduction**

54 The weevils of the South American genus *Leschenius* del Río (Curculionidae, Entiminae,
55 Naupactini) range in the highlands of Colombia and Ecuador, at approximately 1800 to 5000
56 m.a.s.l. (del Río, Marvaldi & Lanteri, 2012). According to the phylogeny of the Naupactini tribe
57 by Lanteri & del Río (2017), the genus *Leschenius* belongs to the so-called “Andean group”, a
58 clade of genera distributed in the Andes: (*Asymmathetes* Wibmer & O’Brien (*Amphideritus*
59 *Schoenherr* (*Leschenius* (*Amitrus* Schoenherr (*Obrieniolus* del Río (*Melanocyphus* Jekel,
60 (*Trichocyphus* Heller))))))), supported by the pro-femora about as wide as meta-femora and by
61 some features of the vestiture like the scarcity of scales and the elytral setae being either erect
62 and long or absent. The “Andean group” belongs to a larger clade, defined by the reduction of
63 the elytral humeri and metathoracic wings, that includes most genera often related to *Pantomorus*
64 *Schoenherr* *sensu lato*, like *Atrichonotus* Buchanan, *Aramigus* Horn, *Phacepholis* Horn,
65 *Parapantomorus* Emden (Lanteri & del Río 2017).

66 **The genus** *Leschenius* is recognized by the black, denuded, and shiny integument, the well-
67 developed denuded ridge at the apex of the rostrum, bordering the mandibles (pre-epistome), the
68 pronotal base "V" shaped, the elytral base curved backward, the reduction of the metathoracic
69 wings, and by the procoxae separated and situated much closer to the anterior than to the
70 posterior margin of the prosternum.

71 some specimens which differ in some diagnostic characters such as the length of the elytra, the
72 density of the vestiture, and the shape of the female ventrite 4. After close examination, we
73 concluded that these specimens correspond to a new bisexual species, *Leschenius L. ventrilingulatus* del Río
74 & Marvaldi sp. nov., close to *Leschenius L. vulcanorum*. We also found male specimens, previously
75 unnoticed, as belonging to *L. vulcanorum*. Finally, and despite its great economic importance as
76 a potato pest, we realized there was not a detailed larval description for this species, or any
77 representative of *Leschenius*.

78 Mixed in the series of *Leschenius-L. vulcanorum* (Kirsch), we found some specimens which
79 differ in some diagnostic characters such as the length of the elytra, the density of the vestiture,
80 and the shape of the female ventrite 4. After close examination, we concluded that these
81 specimens correspond to a new bisexual species, *Leschenius L. ventrilingulatus* del Río &
82 Marvaldi sp. nov., close to *Leschenius L. vulcanorum*. We also found male specimens,
83 previously unnoticed, as belonging to *L. vulcanorum*. Finally, and despite its great economic
84 importance as a potato pest, we realized there was not a detailed larval description for this
85 species, or any representative of *Leschenius*.

86 The purpose of this contribution was to provide a systematic update of the
87 genus *Leschenius*, including descriptions of a new species, the larva of *L. vulcanorum*, the males
88 of *L. vulcanorum* and *L. bifurcatus* del Río, Marvaldi & Lanteri, along with lectotype
89 designations, updated phylogenetic analysis and a revised key to all known species of the genus.

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93 **Materials & Methods**

94 The study was based upon the examination of adult specimens borrowed from the following
95 institutions: Charles O'Brien collection, now housed at Arizona State University (ASUCOB,
96 Tempe, USA), The Natural History Museum (BMNH, London, UK), Museo de La Plata (MLPC,

97 La Plata, Argentina), Muséum National d'Histoire Naturelle, (MNHN, Paris, France), Museum
98 für Tierkunde, (MTD, Dresden, Germany), Museum für Naturkunde (ZMB, Berlin, Germany).

99 Immature stages. The slide-mounted larval specimens are deposited at the MLPC,
100 labelled with the data of this article. Techniques for dissection of larvae, terminology and
101 abbreviations herein applied corresponds to *Marvaldi* (1998).

102 Dissections of genitalia were made according to standard entomological techniques, and
103 characters were drawn using a camera lucida adapted to a stereoscopic microscope (Nikon
104 SMZ800). All measurements were taken with an ocular micrometer attached to the same
105 microscope, and their abbreviations used in the description are as follows: L, maximum length;
106 LA, length of antennae; LB, length of body; LE, length of elytra; LP, length of pronotum; W,
107 maximum width; WRa, width of rostrum measured across apex (excluding scrobes); WRb, width
108 of rostrum at anterior margins of eyes. The terminology used for the morphological structures
109 follows *Marvaldi et al.* (2014), *Lanteri & del Río* (2017) and the glossary of weevil characters by
110 *Lyal* (2021). The terminology used for the sculpture follows that of *Harris* (1979).

111 The electronic version of this article in Portable Document Format (PDF) will represent a
112 published work according to the International Commission on Zoological Nomenclature (ICZN),
113 and hence the new names contained in the electronic version are effectively published under that
114 Code from the electronic edition alone. This published work and the nomenclatural acts it
115 contains have been registered in ZooBank, the online registration system for the ICZN. The
116 ZooBank LSIDs (Life Science Identifiers) can be resolved and the associated information viewed
117 through any standard web browser by appending the LSID to the prefix <http://zoobank.org/>. The
118 LSID for this publication is: urn:lsid:zoobank.org:pub:348600A7-0721-4BC9-A3FD-
119 CB5CBDC55954. The online version of this work is archived and available from the following
120 digital repositories: PeerJ, PubMed Central SCIE and CLOCKSS.

121

122 **Phylogenetic analysis**

123 For the phylogenetic analysis of *Leschenius*, the list of morphological characters and the data
124 matrix by *del Río, Marvaldi & Lanteri* (2012) were modified to include the new species as a
125 terminal taxon as well as new information about the males of *Leschenius vulcanorum* and *L.*
126 *bifurcatus*.

127 For the inclusion of the new species in the present analysis, four characters from the list by del
128 *del Río, Marvaldi & Lanteri (2012)* were redefined (chars. 5, 41, 47, and 48), and a new one for
129 the male genitalia was added (Table 1). The new list consisted of 50 morphological characters of
130 the adults, including 36 from the external morphology and 14 from the female (9) and male
131 terminalia (5). The data matrix herein analyzed includes 12 terminal taxa, corresponding to seven
132 species of *Leschenius* plus five outgroup taxa (Table 2) closely related to *Leschenius* according
133 to *Lanteri & del Río (2017)*. All characters were treated as non-additive and analyzed under
134 equal weights.

135 A cladistic analysis was conducted with TNT v1.5 (*Goloboff & Catalano, 2016*), using
136 the “traditional search” algorithm, with 100 random addition sequences, Tree Bisection and
137 Reconnection (TBR) branch swapping, holding 10 trees during each replication. The most
138 parsimonious tree was rooted with *Melanocyphus lugubris*. Clade stability was evaluated with
139 1000 replication Bootstrap (BT) (*Felsenstein, 1985*), support values over 50 % were indicated
140 below branches. The total length (L), the consistency index (CI) (*Kluge & Farris, 1969*), and the
141 retention index (RI) (*Farris, 1989*) of the most parsimonious trees (MP tree) were calculated
142 excluding the uninformative characters. The character changes were mapped on the tree using
143 fast (ACCTRAN) optimization with WINCLADA1.00.08 (*Nixon, 2002*).

144
145

146 **Results**

147 **Cladistics.** The analysis yielded one most parsimonious tree (L=155 steps, CI= 0.56, RI= 0.53) (Fig. 1). In the cladogram *Asymmathetes pascoei* is the sister group of *the genus Leschenius*, relationship that is supported by several synapomorphies (at least 10 exclusive and one homoplastic character changes, shown in Fig. 1). *The genus Leschenius* is monophyletic and includes the new species, *Leschenius L. ventrilingulatus*, sharing the synapomorphies of the genus: the well-developed pre-epistome (character 10.1); the ‘V’-shaped pronotal base (character 17.2); the slightly prominent to absent elytral humeri (character 23.1); and the procoxae almost contiguous with anterior margin of prosternum (character 29.0). *Leschenius* is also supported by five homoplastic characters states: antennal scape slightly exceeding posterior margin of eyes (character 12.2, with reversal to 12.1 in *L. nigrans*–*L. manueli* clade); funicle segments 2 and 1 subequal (character 13.2, with reversal to 13.1 in *L. vulcanorum* and *L. rugicollis*); elytral base straight to slightly curved backwards (character 21.1) with evolutionary

159 transition to 21.2 in *L. nigrans*–*L. manueli* clade (apomorphic with a reversal to 21.1 in *L.*
160 *nigrans*). The latter is an important character for *Leschenius* because all the other genera of the
161 “Andean group” have the elytral base bisinuate (21.0).

162 The genus *Leschenius* has two main clades, named A and B in Fig. 1. Clade A is well
163 supported and includes the new species, *L. ventrilingulatus* as sister of *L. vulcanorum*, a
164 relationship supported by the very short rostrum (character 5.2), the relatively wide pronotum
165 (character 14.1), the short cornu of spermatheca (character 44.0); the penis with its apex tapering
166 into a long acute projection (character 47.0), dorsally slightly recurved (character 49.1), with a
167 long flagellum like sclerotization in the endophallus (character 48.2). Clade B includes the
168 remaining five species of the genus, supported by the rostral sulcus exceeding posterior margin
169 of eyes (character 7.1), corbel of metatibial apex narrow, setose or denuded (character 31.2);
170 penis apodemes half as long as penis body (character 45.1), and by some homoplastic characters
171 such as medium sized body (character 0.1), sides of rostrum slightly convergent towards apex
172 (character 6.0), and presence of apical projection of elytra (character 24.1). In clade
173 B, *Leschenius* *L. bifurcatus* is the sister of the remaining species, which form a clade defined by
174 the elytral base strongly curved backwards (character 21.2) and by the homoplastic characters:
175 antennal scape reaching posterior margin of eye (character 12.1), pronotum slightly conical
176 (character 15.1) with lateral angles projected in males (character 18.1), and humeral angle of
177 males anteriorly projected (character 22.1). They are grouped in two sister subclades, one
178 including *L. nigrans* and *L. silviae*, defined by one synapomorphy, the obtuse angle between the
179 longitudinal axis of penis and its apodemes (character 46.1), and four homoplastic character
180 states: funicle segment 2 more than 1.5 times longer than segment 1 (character 13.0), maximum
181 width of elytra at anterior third (character 20.2), apical comb of metatibiae longer than dorsal
182 comb (character 32.0), and blunt posterior margin of ventrite 5 in males (character 35.3). The
183 other subclade includes *L. rugicollis* and *L. manueli* and is supported by two synapomorphies,
184 strongly convex eyes (character 11.0), and apex of median lobe evenly rounded (character 47.3),
185 plus three homoplastic character states, the rostrum and forehead coarsely strigose (character
186 3.3), pronotum foveolate-granulose (character 4.2), and penis apodemes slightly shorter than
187 penis body (character 45.0).

188

189 **Taxonomy**

190 *Leschenius* del Río 2012

191 *Leschenius* del Río in del Río, Marvaldi & Lanteri 2012: 55.

192

193 Most characters of the following larval description, based on specimens of *Leschenius* *L.*

194 *vulcanorum*, may apply to the genus *Leschenius*. Terminology as in Marvaldi (1998).

195

196 **Description of larvae. Mature larva.** Body robust, widest at thorax and first abdominal
197 segments. Cuticle asperities present on ventral areas; and absent on lateral and dorsal areas.
198 **Head** (Fig. 2A). Deeply retracted into thorax, longer than wide; posterior margin ogival;
199 posterior half unpigmented, with softer integument and without setae; all setae shifted anteriorly,
200 placed on anterior third. Epicranial line more than 0.5 x the length of head capsule. Frontal lines
201 and endocarina absent. Hypopharyngeal bracon with paramedian maculae. Postoccipital
202 condyles obtuse, hyaline. Setae (Fig. 2A): *fs*4,5, *des*5, and *les*2 long, subequal situated on
203 anterior cephalic fifth; *des*1, *des*3 shorter but well developed; *fs*1,2,3, *des*4, *pes*1-4 minute; *les*1
204 reduced; *vcs*1 longer than *vcs*2, both short. Stemmata absent. Antenna (Fig. 2B) with sensorium
205 about 2.5 x wider than long, truncate at apex. Labrum (Fig. 2C) with *lms*1,2,3, subequal, *lms*1
206 slightly less widely separated than *lms*2. Epipharynx (Fig. 2D) with *mes*1 less separated than
207 *mes*2; epipharyngeal sensilla as single units (not as sensillum clusters), one pair between *mes*1
208 and *mes*2 but closer to *mes*2, and another pair close to bases of labral rods; labral rods (Fig. 2D)
209 ax shaped, bifurcate, with one arm reaching *mes*1 and the other *mes*2. Mandibles (Fig. 2E) with
210 *mds*1 slightly longer than *mds*2, both transversely placed within the scrobe; *mds*2 exterior and
211 slightly basal to *mds*1. Maxillae (Fig. 2F-G) with spinules on dorsal surface of external margin of
212 stipes and below mala and palpus; maxillary mala with a row of eight *dms* and with four *vms*.
213 Labium (Fig. 2F) with premental sclerite well sclerotized, with posterior extension truncate and
214 expanded at apex, anterior extension slender. **Thorax** (Fig. 3A). Spiracle (Fig. 3E) ellipsoidal,
215 without airtubes. Pronotum (Fig. 3A) with nine setae. Meso- and metathorax with *pds*3 distinctly
216 longer than others; alar area with two *as*. Pedal areas of thoracic segments (Figs. 3A-B) each
217 with nine setae: seta *z* conspicuous; setae *x* and *y* subequal; *u* smaller than *v*; *v* smaller than *w*;
218 seta *v'* present and rather conspicuous; small *x'* distinct; a pair of additional anterosternal
219 microsetae occasionally present in front of each pedal area. **Abdomen** (Figs. 3C-D, F): Spiracles
220 (Fig. 3E) elliptical, 2x smaller than thoracic one, without airtubes. Segments AI-VII (Figs. 3A,

221 C) with five *pds*, *pds3* and *pds5* the longest; spiracular area with only *ss2* distinct and
222 progressively placed closer to postdorsum, *ss1* vestigial or absent. AVIII with four *pds*, lacking
223 the homologous *pds2* of preceding segments; *ss* indistinct. Abdominal apex (Fig. 3D) modified,
224 with transverse posterior sclerotized ridges on dorsum, pleura and sternum of AIX; AIX with
225 four *ds* including a seta *ds'*, placed lateral to *ds1*; laterally with two *ls* strongly unequal, the
226 longest on sclerotized ridge; AX terminal, four-lobed, each lateral anal lobe with three minute
227 setae, the outermost larger.

228 **First instar larva** (Figs. 2H–J, 3E). **Head** (Figs. 2H–I) only slightly retracted into thorax,
229 slightly longer than wide; major cephalic setae placed on anterior half, *des2* and *les1* less reduced
230 than in older larvae; *des1* minute (smaller than in mature larvae). Frontal lines weakly distinct.
231 Anterior and posterior stemmata distinct, as dark pigmented spots. Antennal sensorium
232 prominent and projected outwards. Mandibles (Fig. 2J) with *mds1,2* strongly unequal, *mds1*
233 about 5x longer than *mds2*. **Thorax**. Spiracle (Fig. 3F) bicameral with annulated airtubes; pedal
234 area with setae *z*, and *v'* relatively small, seta *w* relatively very long and spatulate or blunt at
235 apex. **Abdomen**. Spiracles (Fig. 3F) bicameral, with airtubes having a smaller number of rings
236 than thoracic one; abdominal apex not distinctly sclerotized.

237
238 **Remarks.** The characters mentioned above for the first larva, newly hatched, are exclusive of the
239 first instar (see also Marvaldi & Loiácono, 1994). Additional differences between early and older
240 instar larvae involve relative dimension of structures, like the antennal sensorium, which are
241 relatively much larger in the first instar; the pigmentation and level of sclerotization of body
242 areas tend to increase in successive instars; there are larger differences in length between setae of
243 different body areas in early instars.

244
245 **Comparative notes.** The larval characters given herein for the genus *Leschenius* are in full
246 agreement with those defining the tribe Naupactini (Marvaldi & Loiácono, 1994) or *Naupactus*
247 and allied genera (Marvaldi, 1998). Two apparently distinct features of the mature larva studied
248 of *L. vulcanorum* are the head capsule with *des1* well developed (in known mature larvae of
249 other Naupactini *des1* is minute or very small); also, setae *x* and *y* of pedal areas of thoracic
250 segments are subequal (in other Naupactini as *Naupactus*, seta *x* is distinctly smaller than *y*).
251 Larval characters deemed as diagnostic for the species are given below.

252

253 **Notes on type material.** Four paratypes of the species of *Leschenius* described in *del Río,*
254 *Marvaldi & Lanteri* (2012) were finally deposited in the MLPC instead of the collection
255 mentioned in the original publication: one paratype of *Leschenius-L. bifurcatus* del Río, Marvaldi
256 & Lanteri, with labels 'ECUADOR, PICHINCHA, POMASQUI, RUNICUCHO, 2400 m, 6–
257 XII–1993, K. Volbracht'; two paratypes of *L. manueli* del Río, Marvaldi & Lanteri, with labels
258 'ECUADOR, AZUAY, VIA CUENCA LOJA, 5 km DE ONA, 13–I–1997, A. Paucar'; and one
259 paratype of *L. silviae* del Río, Marvaldi & Lanteri, with label 'Cuenca Jesta'.
260

261 ***Leschenius vulcanorum* (Kirsch, 1889)**

262 (Figs. 2, 3, 4, 5A, 6G, K, 7)
263 *Canephorus vulcanorum* Kirsch 1889: 17; Strand 1943: 96 (*Canephorulana*);
264 Kuschel 1955: 277 (*Amitrus*); Kuschel in Wibmer & O'Brien 1986: 53 (*Asymmathetes*) (Fig.
265 4A).

266 *Amphideritus brevis* Oliff 1891: 68; DallaTorre, Emden & Emden 1936: 14 (*Macrostylus*);
267 Kuschel 1955: 277 (*Amitrus*) (syn. of *A. vulcanorum*); Kuschel in Wibmer & O'Brien 1986: 53
268 (*Asymmathetes*); *del Río, Marvaldi & Lanteri* 2012: 60 (*Leschenius*) (Fig. 4B).
269 *Amphideritus pigmaeus* Oliff 1891: 68; Dalla Torre, Emden & Emden 1936: 14 (*Macrostylus*);
270 Kuschel 1955: 277 (*Amitrus*) (syn. of *A. vulcanorum*); Kuschel in Wibmer & O'Brien 1986: 53
271 (*Asymmathetes*) (Fig. 4C).

272 *Caulostrophus aequatorialis* Kirsch 1889: 13; Dalla Torre, Emden & Emden 1939: 319
273 (*Macrostylus* ~~(*Amphideritus*)~~); Kuschel in Wibmer & O'Brien 1986: 53 (*Asymmathetes*) (Fig.
274 4D).
275

276 Diagnosis and description of female in *del Río, Marvaldi & Lanteri* (2012).

277

278 **Description of male** (Fig. 5A). Smaller than female (4.0–6.3 mm; females 5.3–8.7 mm); rostrum
279 shorter (L/Wa: 0.76–0.84); less convergent towards apex (Wb/Wa, 1.15–1.30); antennal club
280 more elongate (L/W, 2.5–2.8); pronotum (W/L: 1.25–1.35), wider than the elytra and longer than
281 in females, with sides more arcuate; elytra slightly shorter (L/W, 1.32–1.40) with apex not
282 divided, more rounded; metatibiae with larger mucro than in females; posterior margin of

Con formato: Fuente: Cursiva

283 ventrite 5 blunt. Genitalia (Figs 6G, K). Median lobe slightly curved in lateral view, tapering
284 towards apex, with subacute, dorsad slightly recurved hook-like apex; penis as long as abdomen;
285 apodemes slightly shorter than median lobe (0.7x); endophallus armed with minute spicules and
286 with a slightly sclerotized flagellum.

287

288 **Larval stage.** One mature larva and four submature larvae, as well as associated adults of *L.*
289 *vulcanorum*, were collected from the following locality: Colombia, Municipio
290 de Sibaté, vereda el Romeral, 4°26'3" N, 74°14'8" O (3100 masl), J.E.C. Gomez leg., 2009.
291 Additionally, 10 first instar larvae were obtained from eggs deposited by some of the collected
292 adults kept in captivity. After comparison with larvae known for other species in tribe Naupactini
293 (Marvaldi & Loaiácono, 1994; Marvaldi, 1998) the following combination of characters can be
294 suggested as diagnostic for the species *L. vulcanorum*.

295 **Mature larva** (Figs. 2–G, 3A–E). Maximum head width 2.2 mm. Setae fine, brown. Head
296 yellowish, intense yellow on anterior margin of frons, about 1.3 x longer than wide. Cephalic
297 setae (Fig. 2A): *des1* well developed (although shorter than *des3* and those placed on anterior
298 third, *fs4*, *fs5*, *des5*, and *les2*). Epipharynx with spinules anteriad and posteriad to the labral
299 rods; epipharyngeal sensilla not in clusters but apparently fused into single units. Pronotum
300 pigmented with pattern of brownish maculae (Fig. 3A). Abdominal apex (AIX) with transverse
301 sclerotized ridges in dorsum, pleura and sternum (Fig. 3D).

302 **Larva 1** (Figs. 2H–J, 3F). Maximum head width 0.2 mm. Head (Fig. 2A) with *des1* minute (like
303 *des 4*), cephalic setae well developed are *des2*, *des3*, *des5*, *fs4*, *fs5*, *les1*, and *les2*; *des1* slightly
304 more widely separated than *fs4*. Clypeus with setae subequal. Labrum with *lms2* somewhat more
305 widely separated than *lms1*.

306

307 **Type material examined.** Lectotype of *Canephorus vulcanorum* Kirsch, female, Ecuador
308 Tunguragua, 3800, *Canephorus vulcanorum* typus Kirsch, MTD, (Fig. 4A), here designated.
309 Paralectotype of *Canephorus vulcanorum*, Ecuador, Sangay, 3500m, cotypus, MTD. Lectotype
310 of *Amphideritus brevis* Oliff, female, Ecuador feet, Ed. Whymper, MNHN, here designated (Fig.
311 4B). Lectotype of *Amphideritus pigmaeus* Oliff, female, Chimborazo, Ecuador, 12-13000 feet,
312 Ed Whymper, MNHN, here designated (Fig. 4C). Lectotype of *Caulostrophus aequatorialis*
313 Kirsch, female, Cotopaxi, 5688, Typus, MTD, here designated (Fig. 4D).

314

315 **Other material examined.** **COLOMBIA.** No loc., int. Miami, 5-2-85, with cut flowers of
316 *Dianthus* sp. (1f USNM). **ECUADOR.** No loc., 11-11-93 (2f USNM). **Chimborazo:**
317 Chimborazo, S side of Mt, elev 11600 nr Snowline, 19 june 1975 (1f USNM); Colta, 3-VI-05,
318 Ohaus, 9-VII-05 (27f ZMB); Interandin-Hochland, Colta 3500-4000m, 8-10-VII-1905 Ohaus (1f
319 ZMB), 2,8-VII-1905 Ohaus (2f ZMB); Faldas del Chimborazo, jan-1983, in pine leaves *Pinus*
320 *radiata*, adults feed, Lopez col (1f USNM); Guamote, 3-7-1969, en cocoons of alfalfa plants (1f
321 USNM); Guaslam prov, 1-22-60-on bucts of young corn, Merino (2f USNM); Quimiag on
322 maize, Jan 1979 Kirckhy (2f BMNH); Riobamba, 3-VII-1905, unter steinen, Ohaus (17f ZMB),
323 20-XI-05 (1f ZMB), 27-XI-05 (2f ZMB); Riobamba, Ause de Cubillin, 3500, 5-Vii-05 Ohaus
324 (42f ZMB). **Cotopaxi:** 71 km W Latacunga under stones May 1, 1978, O'Brien & Marshall (1f
325 1m MLP; 45 km W Latacunga, under stones May1 1978, O'Brien & Marshall (8f MLP); 21
326 km S Latacunga, April 25 1978, CW&L O'Brien & Marshall (1f MLP); 6 km W Latacunga,
327 under stones, May1 1978, O'Brien & Marshall (3f MLP); Latacunga, XI 1981 Onore Brit Mus
328 1990-214 (1f BMNH); 15 km W entrance PN Cotopaxi, April 30 1978, O'Brien & Marshall (4f
329 MLP); 14 km W entrance PN Cotopaxi, April 30 1978, O'Brien & Marshall (1f MLP);
330 Cotopaxi, P. 13 km S Latacunga along PanAma, XI-3-77, G Noonan, M. Moffett, under clumps
331 soil and grass, rocks, debris-in green grassy field w green short grass ca 2600m. (13f 1m MLP);
332 Tilipulo, V-III-1981, G. Onore Brit Mus 1985-254 (4f BMNH). **Bolívar:** Guaranda, X-I-1955,
333 on new corn (7f USNM). **Loja.** Loja, Ohaus (2f ZMB). **Pichincha:** 38.8km NE Quito on PanAm
334 XI-8-77, G Noonan, M. Moffett, under rocks on dirt clumps, in areas with sparse to very sparce
335 short grass ca 2200m. (1f MLP). **Tungurahua:** SE end Ambato, XI-1-77, G Noonan, M.
336 Moffett, under rock in fields w short sparce grass, soil dry under stones, ca 2500m. (2f MLP);
337 13km NE Baños, April 26 1978, O'Brien & Marshall (1f MLP); Baños, 1800m, 9-V-37
338 Brundage (2f USNM); Baños, X-4-44, EJ Hambleton (1f USNM); Pomasqui, X-7-54, Merino,
339 orange trees (4f USNM); Totoras, 7 km SE Ambato, April 26 1978, O'Brien & Marshall (3f
340 MLP). Plus, the material listed in *del Río, Marvaldi & Lanteri (2012)*.
341
342 **Remarks.** In the revision of the genus *Leschenius* (*del Río, Marvaldi & Lanteri, 2012*), the type
343 material of the species *Asymmathetes aequatorialis* (Kirsch) was not seen and we mentioned
344 that this species may also belong to the genus *Leschenius*. Herein, based on the observation of

345 the type material of all the names related to *Leschenius vulcanorum* (Fig. 4A–D), including
346 *Caulostrophus aequatorialis* Kirsch (Fig. 4D), we establish the synonymy of this name with
347 *Leschenius L. vulcanorum* (Fig. 4A). This species is only known from the type material and
348 corresponds to a phenotype within the great variation observed in *L. vulcanorum* (see Fig. 4A–
349 D).

350 Bisexual populations of *Leschenius L. vulcanorum* ~~were have been~~ so far only seen in
351 Ecuador, near Latacunga locality (Cotopaxi province), and in Ambato locality (Tungurahua
352 province) (Fig. 7). Differences noted between the bisexual and the parthenogenetic populations
353 are related with the body size and morphometrics of the elytra. The bisexual form is usually
354 smaller with slightly shorter elytra.

355

356 **Host plants.** *Leschenius vulcanorum* was found in association with cabbage *Brassica oleracea*
357 L. (Brassicaceae), alfalfa *Medicago sativa* L. (Fabaceae), young corn *Zea mays* L. (Poaceae),
358 pine *Pinus radiata* D. Don, orange trees, and with cut flowers of carnation *Dianthus* sp. It is
359 considered an important pest of potato *Solanum tuberosum* L. (Solanaceae) in Colombia, and is
360 known as ‘~~p~~Potato shooter’ (del Rio, Marvaldi & Lanteri 2012).

361

362 *Leschenius ventrilingulatus* del Río & Marvaldi, sp. n.

363 urn:lsid:zoobank.org:act:9A70B8AE-74BF-4631-98A7-A1C814113833

364 (Figs 5B–F; 6A–F, H, L, 7)

365

366 **Diagnosis.** *Leschenius ventrilingulatus* is easily distinguished from the remaining species of
367 *Leschenius* (except *L. vulcanorum*) by possessing a shorter and less conical rostrum with sides
368 not thickened and elevated, and apex not projected. It is very similar to *L. vulcanorum*, but
369 distinguished by its size usually small, vestiture of denser and thicker setae (mainly on head, legs
370 and elytra), shorter antennae (with funicular segments 1 and 2 subequal); shape of the pronotum,
371 subcylindrical with anterior margin as wide as posterior margin; elytra shorter (about 1.5x longer
372 than wide at base; 2x in *L. vulcanorum*); the metatibial apex with narrow corbel; female with
373 ventrite 4 with a posterior rounded projection; and posterior margin of ventrite 5 subacute, not
374 excavated. Female genitalia with plate of sternite VIII sub-rhomboidal not elongate with longer
375 apodeme.

377 **Description. Female** (Figs. 5B, D–F). Species medium-sized (LB, 5.0–6.0 mm). Tegument
378 visible, dark brown to reddish brown, shiny. Vestiture composed of disperse, pale ochre to cream
379 colored setae, moderately dense, cream-colored decumbent setae-like scales, absent on middle of
380 pronotum (forming wide lateral stripes) on pronotum, grouped on patches on elytra, in some
381 cases devoid of scales along middle line; also present on venter and legs (more abundant on
382 distal third of femora). Rostrum very short (Fig. 5F) (L/Wa, 0.73–0.83), sides moderately
383 convergent towards apex (Wb/Wa, 1.31–1.43), dorsum moderately convex. Forehead foveolate–
384 strigose, with longitudinal striae. Vertex sparsely punctate. Antennae (Fig. 6A) of medium length
385 (LB/LA, 2.50–2.85); scape reaching to slightly exceeding posterior margin of eyes. Funicle with
386 segment 2 about as long as segment 1, both elongated; funicle segments 3–6 slightly longer than
387 wide, and funicle segment 7 as long as wide; club slightly fusiform (L/W, 2.3–2.4). Pronotum
388 (Fig. 5B) slightly subcylindrical, moderately wider than long (W/L, 1.30–1.35), with anterior
389 margin as wide as posterior margin; median groove absent. Scutellar shield subtriangular, large
390 and wide (surrounded by elevated edges), denuded. Elytra (Fig. 5B, E) short (L/W, 1.23–1.33),
391 with maximum width on anterior third, slightly convex; base slightly curved backwards on
392 middle; intervals about twice as long as striae, flat; striae with medium-sized punctures, 9–10
393 slightly closer along posterior two-thirds; elytral apex acute not projected or bifurcate, entire.
394 Legs. Procoxae much closer to anterior than posterior margin of prosternum; protibiae with
395 medium-sized mucro, and row of acute small denticles (six or seven, on distal two-thirds of
396 tibiae); meso- and metatibiae with small mucro and without denticles; metatibial apex with
397 narrow corbel covered with disperse small elongate cream scales; apical and dorsal combs
398 subequal. Abdomen (Fig. 5D). Intercoxal portion of ventrite 1 slightly broader than metacoxal
399 cavities (1.10–1.15x); ventrite 2 longer than ventrites 3 and 4 combined (1.60x without
400 projection; 1.10 along midline); ventrite 4 with a posterior rounded projection; posterior margin
401 of ventrite 5 subacute, not excavated; tergites I–VII membranose. Terminalia. Sternite VIII (Figs.
402 6B–C) with plate sub-rhomboidal, not elongate, with tuft of medium-sized and coarse setae, and
403 with shorter setae on apical third; ‘V’-shaped sclerotization with lateral arms reaching two-thirds
404 of plate, and lateral margins sclerotized; apodeme 2.8–3.0x longer than plate. Ovipositor (Figs.
405 6D–E) as long as ventrites 1–5; with scattered fine short setae on sides of baculi on anterior third;
406 ventral baculi subparallel; styli wide. Spermathecal body (Fig. 6F) sub-cylindrical; collum (duct-

407 lobe) conical, short; ramus (gland-lobe) indistinct; cornu long; spermathecal duct (Fig. 6D) short,
408 half as long as ovipositor, or 3x the maximum width of spermatheca, membranous, moderately
409 wide.
410 Male (Fig. 5C). Same size as female (4.8–5.5 mm); rostrum less conical (Wb/Wa, 1.32–1.37);
411 antennal club more elongate (L/W, 2.6–2.7); elytra slightly more elongate (L/W, 1.30–1.35);
412 metatibia with larger mucro than in female; ventrite 4 without posterior projection; posterior
413 margin of ventrite 5 blunt. Genitalia (Figs. 6H, L). Penis (median lobe) slightly curved in lateral
414 view, tapering towards apex, with dorsad recurved hook-like apex; penis as long as abdomen;
415 penis apodemes slightly shorter than penis body (0.8x); endophallus armed with minute spicules
416 and with a sclerotized flagellum, with a denticulated blade (Fig. 6H).
417 Morphometrics. Holotype, female: rostrum L/Wa: 0.77, Wb/Wa: 1.31; antenna LB/LA: 2.85,
418 antennal club L/W: 2.31; pronotum W/L: 1.34; elytra L/W: 1.33; LE/LP: 2.6.

419

420 **Etymology.** The specific name refers to the tongue-like projection of the female venter.

421

422 **Material examined. Holotype.** Female, 5 mm long, with labels as follows “ECUADOR,
423 Totoras,/ 7km SE. Ambato/Apr. 26, 1978 C&L/ O’Brien & Marshall” (MLPC). **Paratypes.**
424 Same data as holotype (1m ASUCOB); ECUADOR, 6 km W/ Latacunga under/ stones May 1,
425 1978/ O’Brien & Marshall (1f, 1m MLP, dissected with genitalia in vial with glicerine);
426 ECUADOR/ Latacunga/ IX- 1981 Onore/ Brit. Mus./ 1990-214 (3f 2m BMNH); ECUADOR,
427 5km SE./ Pelileo, April/ 26, 1978 CW&LB/ O’Brien & Marshall (1m ASUCOB); ECUADOR/
428 Catamayo B./ Loja 5.9.05/ F. Ohaus (1f ZMB).

429

430 **Geographic distribution (Fig. 7).** *Leschenius ventrilingulatus* is endemic of Ecuador, known for
431 Cotopaxi, Loja, and Tungurahua provinces. It is distributed on the interandean region on river
432 basins, between 1250 and 2750 meters above sea level (unlike *L. vulcanorum* which is
433 distributed in higher altitudes 2600-5000 m. a. s. l.). It is sympatric with *L. vulcanorum* (Fig. 7)
434 in Latacunga (Cotopaxi province) and Totoras (Tungurahua province) at 2600-2750 m. a. s. l.

435

436 **Remarks.** *Leschenius ventrilingulatus* is the sister species of *Leschenius-L. vulcanorum*,
437 according to results of the cladistics analysis herein undertaken (Fig. 1).

438

439 *Leschenius bifurcatus* del Río, Marvaldi & Lanteri 2012

440 (Figs 5G-H, 6 I, J, M, 7)

441

442 Diagnosis and description of female in *del Río, Marvaldi & Lanteri (2012)*.

443

444 **Description of male** (Fig. 5H). Similar size as female, slightly smaller (Fig. 5G) (10–13 mm);
445 rostrum (L/Wa, 1.0–1.1; Wb/Wa, 1.30–1.45); antennal club slightly more elongate (L/W, 3.1–
446 3.4); pronotum (W/L, 1.17–1.19); elytra (L/W, 1.60–1.65) with projected apex but not
447 bifurcated, only slightly divided; ventrites 3–5 not bulged as female, posterior margin of ventrite
448 5 blunt. Genitalia (Figs 6I, J, M). Penis body slightly curved in lateral view, tapering towards
449 apex (Fig. 6M), with dorsal strongly recurved hook-like apex (Fig. 6J); penis longer than
450 abdomen (1.25–1.30); penis apodemes much shorter than median lobe (0.4x); endophallus armed
451 with minute spicules and with two long wing-shaped sclerites (Fig. 6I).

452

453 **Material examined. ECUADOR.** No loc, intercept. Port Miami, 16-VI-2004 (1f USNM).

454 Imbabura: 3.9 km N. Ibarra on Pan Am, XI-II-77, Moffet collr, under rocks by rd. in dry area,
455 with sparse grass ca. 2300 (1f, 3m ASUCOB; 1f, 2m MLPC); Urcuqui, 14-III-62, Merino &
456 Vasquez, in soil nr cotton plants (2m USNM); El Chotar, Mr. Juncal, 1 june-1961, Merino &
457 Vasquez, reared from larvae doing damage to roots of bean plants (1m USNM). Pichincha:
458 Conocoto, 31-Jan-1992, Alvaro Barragan (1m MLP); Pomasqui, Runicucho, 2400 m., 6 Dec-
459 1993, E. Volbracht (1m PUCE). Imbabura: 3.9 km N. Ibarra on Pan Am, XI II-77, Moffet collr,
460 under rocks by rd. in dry area, with sparse grass ca. 2300 (1f, 3m ASUCOB, 1f, 2m MLPC);
461 Urcuqui, 14 III-62, Merino & Vasquez, in soil nr cotton plants (2m USNM); El Chotar, Mr.
462 Juncal, 1 june 1961, Merino & Vasquez, reared from larvae doing damage to roots of bean plants
463 (1m USNM). Plus, the material listed in *del Río, Marvaldi & Lanteri (2012)*.

464

465 **Remarks.** The specimens of the population from Ibarra (Imbabura) are slightly different from
466 the type, the female (Fig. 5G) has a wider and shorter pronotum and the elytral apex only slightly
467 bifurcated but strongly projected posteriad; the males have also a wider and shorter pronotum
468 and the penis with a shorter ostium area and a more recurved apex (Fig. 6J).

Con formato: Español (Méjico)

469

470 **Host plants.** The larvae of *Leschenius L. bifurcatus* was found in association with bean plants,
471 *Phaseolus vulgaris* L. (Fabaceae).

472

473 **Key to species of *Leschenius***

474 Modified from *del Río, Marvaldi & Lanteri (2012)*.

475

476 1. Size 4–8 mm (usually 5–7 mm, exceptionally more than 8 mm but never more than 9 mm); *r-*
477 *Rostrum* very short (L/W less than 0.95, usually near 0.8); pronotum with setae forming two
478 feeble lateral stripes; elytral apex not
479 projected.....

480 2

481 1'. Size 8–12.5 mm; *r-* *Rostrum* moderately short (L/W more than 0.96, usually 1); pronotum
482 without or with scattered setae, never forming lateral stripes; elytral apex projected
483 backwards.....

484 2. Elytra with cream-colored decumbent setae like scales grouped on patches on entire elytra; *p-*
485 *Pronotum* subcylindrical, with anterior margin as wide as posterior margin; *m-* *Metatibial*
486 apex with narrow corbel; *f-* *Female* with ventrite 4 with a posterior rounded projection (Fig.
487 5D)..... *L. ventrilingulatus, n. sp.*

488 (Figs. 5B–F)

489 2'. Elytra devoid of cream-colored decumbent setae like scales, or if present, limited to margins
490 (intervals 9–10); *p-* *Pronotum* slightly subconical, with posterior margin wider than anterior
491 margin; *m-* *Metatibial* apex with moderately broad corbel (width: 1/3 of the length of the tibial
492 apex); *f-* *Female* with ventrite 4 without posterior projection.....

493 *L. vulcanorum* (Figs. 4A–D, 5A, see fig. 1 in *del Río, Marvaldi &*
494 *Lanteri, 2012*)

495 3. Pronotum with posterior margin as wide as anterior margin; *e-* *Elongate* elytra (3x or more the
496 length of pronotum); *e-* *Elytral* apex, in females strongly projected backwards and distinctly
497 bifid or divided; in males, moderately projected and rounded to slightly
498 divided..... *L. bifurcatus* (see figs. 2–3 of *del Río,*
499 *Marvaldi & Lanteri, 2012*)

Con formato: Fuente: Negrita, Cursiva

500 3'. Pronotum with posterior margin distinctly wider than anterior margin; o-oval elytra (less
 501 than 2.8x the length of pronotum, usually between 2.2–2.7x); e-Elytral apex of both males
 502 and females moderately projected, entire to slightly
 503 divided.....4

504 4. Corbel plate of metatibia broad (width: almost half the diameter of apex of tibiae).....
 505L. nigrans (see figs. 9–10 of *del Río, Marvaldi & Lanteri, 2012*)

506 4'. Corbel plate of metatibia narrow (less than quarter apex of tibiae).....5

507 5. Body length 8.4–9.3 mm, moderately sized; p-Pronotum as wide as elytra in males; e-Elytra
 508 2.2–2.45x the length of pronotum; s-Spermatheca with collum not
 509 constricted.....6

510 5'. Body length 7.8–8.4 mm, small; p-Pronotum distinctly narrower than elytra in both sexes; e-
 511 Elytra 2.5–2.75x the length of pronotum; s-Spermatheca with tubular collum, constricted near
 512 spermathecal body;L. silviae (see figs. 11–12 and 42 of *del Río, Marvaldi & Lanteri, 2012*)

513 6. Pronotum slightly wider than long (W/L: 1.10–1.25); o-Ovipositor without setae on sides of
 514 baculi; a-Apex of penis slightly
 515 pointed.....
 516L. rugicollis (see figs. 4–5 of *del Río, Marvaldi & Lanteri, 2012*)

517 6'. Pronotum of males strongly wider than long (W/L: 1.30–1.35); o-Ovipositor with
 518 conspicuous setae on sides of baculi; a-Apex of penis
 519 rounded.....L.
 520 manueli (see figs. 6–8 of *del Río, Marvaldi & Lanteri, 2012*)

521

522

Con formato: Fuente: Sin Negrita, Sin Cursiva

523 Discussion

524

525 The cladistics analysis performed led us to uncover evidence for the monophyly of the genus
 526 *Leschenius* and for the relationships among its species. It is worth to note that *Asymmathetes*
 527 *pascoei* is supported as the sister group of *Leschenius*, in agreement with *del Río, Marvaldi &*
 528 *Lanteri (2012)*, but differing from results of the analysis by *Lanteri & del Río (2017)*:

Con formato: Fuente: Cursiva

531 (*Asymmathetes* (*Amphideritus* (*Leschenius* (*Amitrus* (*Obrieniolus* (*Melanocyphus*,
532 *Trichocyphus*)))))). However, this is not unexpected as these works had different objectives and
533 thus differ in the taxon and character sampling (e.g. in the analysis of *Lanteri & del Río* there is
534 only one species representing the genus *Leschenius* and, on the other hand, the present study was
535 not designed to test intergeneric relationships). Further research on the systematics and
536 phylogeny of the “Andean Group” will require expanding the outgroup and ingroup taxon
537 sampling (e.g., including *Obrienolus* and more species representative of each genera) and using
538 additional sources of evidence like molecular data.

539 The description of a new species and more complete information about male characters,
540 allowed us to propose an updated phylogenetic hypothesis of *Leschenius*, which differs from the
541 previous study by *del Río, Marvaldi & Lanteri (2012)* regarding the position of *L. bifurcatus* and
542 relationships within clade B (Fig. 1). In the new phylogeny *L. bifurcatus* is the sister species of a
543 subclade that contains *L. nigrans* sister of *L. silviae* and *L. rugicollis* sister of *L. manueli*. Also,
544 the present study led to the discovery of the sister species of *L. vulcanorum* and the pair *L.*
545 *ventrilingulatus* – *L. vulcanorum* is proposed as the sister group of all remaining species of
546 *Leschenius*.

547 The genus *Leschenius* is distributed in the *n*Northern Andes of Ecuador and northern
548 Colombia, approximately 1800–5000 m.a.s.l (Fig. 7), corresponding to the *North Andean*
549 Páramo province of the South American *T*ransition Zone (*Morrone, 201406*). All species
550 included have a narrow distribution range in Ecuador, except *L. vulcanorum* that is widely
551 distributed from central to northern Ecuador and extend to southern Colombia. The latter species,
552 *L. vulcanorum*, along with *L. ventrilingulatus*, *L. bifurcatus*, and *L. nigrans* inhabit in northern
553 Ecuador, and they have been found in sympatry (Fig. 7). The other three species, *L. silviae*, *L.*
554 *manueli*, and *L. rugicollis* are distributed in the southern provinces of Ecuador.

555 The new species *Leschenius-L. ventrilingulatus* is distributed on the inter-Andean region on
556 river basins (Fig. 7) and is sympatric with *L. vulcanorum* at 2500–2750 m.a.s.l., that corresponds
557 to the lowest altitude for *L. vulcanorum*.

558 Of particular interest is the occurrence of parthenogenesis within the genus *Leschenius*,
559 and this mode of reproduction was suggested by *del Río, Marvaldi & Lanteri (2012)* for the
560 species *L. vulcanorum* and *L. bifurcatus*, based on the apparent absence of males in their
561 populations. The examination of additional material in the present study provided new evidence

562 that suggest *L. bifurcatus* is not parthenogenetic, leaving *Leschenius vulcanorum* as the only
563 parthenogenetic species. The parthenogenesis of *L. vulcanorum* was confirmed by laboratory
564 rearing of unmated females that produced viable offspring (del Río, Marvaldi & Lanteri, 2012).
565 Nonetheless, in the present study, we discovered males of *L. vulcanorum*, indicating that this
566 species also has bisexual populations, so far only seen in three localities in Cotopaxi and
567 Tungurahua provinces of Ecuador. The existence of both sexual and parthenogenetic populations
568 within the species supports the idea of a special kind of parthenogenesis, called “geographical
569 parthenogenesis” (Lanteri & Normark, 1995). In *L. vulcanorum*, as in other species having this
570 kind of reproduction, the sexual and parthenogenetic forms have different distribution ranges,
571 being the parthenogenetic one more widespread than the respective sexual forms (Vandel, 1931).
572 So far, exemplars of both sexes of this species were collected in just three localities.
573 Morphological differences between the bisexual and the parthenogenetic form are noticed in
574 body size and morphometrics of the elytra, being the bisexual form usually smaller and with
575 slightly shorter elytra. It is important to remark that within the parthenogenetic form the variation
576 in body size and morphometrics is extremely wide (Figs. 4A–D). Same as other parthenogenetic
577 weevils, including the around 30 parthenogenetic species of the tribe Naupactini (Lanteri &
578 Normark 1995), in *L. vulcanorum* the parthenogenesis is also associated with the wingless
579 condition and xeric habitats.

580 An interesting feature observed in females of the new species described, *L.*
581 *ventrilingulatus* (sexually dimorphic) is the ventrite 4 with its posterior margin produced
582 medially into a tapered lamina or tongue-like projection. This characteristic seems to be unique
583 within the tribe Naupactini, and so far, also unknown among Neotropical Entiminae. Although
584 unusual, a modified ventrite 4 is also present in genera of Entiminae inhabiting montane areas of
585 other regions of the world (see Brown, 2017 and reference therein): New Zealand
586 [*Austromonticola* Brown 2017, *Chalepistes* Brown 2017, and *Nicaeana* Pascoe 1877], Solomon
587 Islands [*Platyacus* Faust, 1897 (Celeuthetini)], Mauritian Islands [*Syzygops* Schönherr, 1826
588 (Ottistirini)]; Chinese Himalayas [*Trichalophus caudiculatus* (Fairmaire, 1886) (Tropiphorini)],
589 Kashmir and Himalayas [*Leptomias* Faust, 1886 (Tanytropini)], and Central
590 America [*Sciomias* Sharp, 1911 (Sciaphilini)]. These structures are hypothesized to have evolved
591 in response to oviposition needs in and beside cushion plants or selected to mitigate the female
592 costs of prolonged mating (Brown, 2017). Although the function has not been studied yet, the

593 first suggestion that these ventral structures may assist in the preparation of oviposition sites in
594 close-packed vegetational structures seems highly plausible: the cushion growth form is a
595 common feature of the alpine vegetation worldwide where the weevil species with this trait are
596 found.

597 Concerning the immature stages, the study of the larvae of *Leschenius* lead us to confirm
598 that the tribe Naupactini is very homogeneous in larval morphology, as suggested by *Marvaldi*
599 (1998). Also, there seem to be no particular features that could be suggestive as adaptive traits to
600 arid environments. This is not unexpected, since *Leschenius* larvae, like those of most entimines,
601 are subterranean and then “preadapted” to live in harsh environments.

602 Finally, additional research with molecular tools will be very important to find out the
603 genetic divergence and evolution of the group, and the role of the parthenogenesis. Moreover,
604 the study of the biology and behavior of *L. ventrilingulatus* will offer insights into the function of
605 the abdominal structures of the female.

606
607

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613

614 **References**

615 Brown SD. 2017. *Austromonticola*, a new genus of broad-nosed weevil (Coleoptera,
616 Curculionidae, Entiminae) from montane areas of New Zealand. *ZooKeys* (707): 73.
617 del Río MG, Marvaldi AE, Lanteri, A. 2012. Systematics and cladistics of a new Naupactini
618 genus (Coleoptera: Curculionidae: Entiminae) from the Andes of Colombia and
619 Ecuador. *Zoological Journal of the Linnean Society* 166(1): 54–71.
620 Farris JS. 1989. The retention index and the rescaled consistency index. *Cladistics* 5: 417–419.
621 Felsenstein J. 1985. Confidence limits on phylogenies: An approach using the bBootstrap.
622 *Evolution* 39: 783–791.
623 Goloboff PA, Catalano S. 2016. TNT version 1.5, including a full implementation of phylogenetic
624 morphometrics. *Cladistics* 32(3): 221–238. DOI 10.1111/cla.12160.

625 Harris RA. 1979. A glossary of surface sculpturing. *Occasional papers in Entomology* 28: 1–31.

626 Kluge AG, Farris JS. 1969. Quantitative phyletics and the evolution of anurans. *Systematic*
627 *Biology* 18(4): 1–32.

628 Lanteri AA, del Río MG. 2017. Phylogeny of the tribe Naupactini (Coleoptera: Curculionidae)
629 based on morphological characters. *Systematic Entomology* 42(2): 429–447.

630 Lanteri AA, Normark BB. 1995. Parthenogenesis in the tribe Naupactini (Coleoptera:
631 Curculionidae). *Annals of the Entomological Society of America* 88(6): 722–731.

632 Lyal CHC. (Ed.) Glossary of Weevil Characters. International Weevil Community
633 Website. Available at <http://weevil.info/glossary-weevil-characters> (accessed 10 November 10,
634 2021). Con formato: Sin subrayado
Con formato: Sin subrayado

635 Marvaldi AE. 1998. Larvae of South American Entimini (Coleoptera: Curculionidae), and
636 phylogenetic implications of certain characters. *Revista Chilena de Entomología* 25: 21–44.

637 Marvaldi AE, Loiácono MS. 1994. First instar larvae in the tribe Naupactini (Coleoptera:
638 Curculionidae). *Revista Brasileira de Entomologia* 38(2): 453–466.

639 Marvaldi AE, Lanteri AA, del Río MG, Oberprieler RG. 2014. Chapter 3.7.5 Entiminae
640 Schoenherr, 1823. In: Leschen RAB, Beutel RG, eds. *Handbook of Zoology. Arthropoda:*
641 *Insecta: Coleoptera, Beetles. Volume 3: Morphology and Systematics (Phytophaga)*.
642 Berlin: de Gruyter, 503–523. Con formato: Fuente: (Predeterminada) Times New Roman

643 Con formato: Fuente: (Predeterminada) Times New Roman

644 Morrone JJ. 2014. *Biogeographical regionalisation of the Neotropical region. Zootaxa* 3782:
645 1–110 *Biogeographic areas and transition zones of Latin America and the Caribbean islands*
646 *based on panbiogeographic and cladistic analyses of the entomofauna. Annual Review of*
647 *Entomology* 51: 467–494. Con formato: Fuente: (Predeterminada) Times New Roman, 12 pto
Con formato: Fuente: (Predeterminada) Times New Roman, 12 pto

648 Nixon KC. 2002. *WinClada, version 1.00. 08*. Ithaca, NY, Published by the author 734, 745.

649 Vandel A. 1931. *La parthenogénèse*. Paris: G. Doin.