

On the *Monacha* species of Lebanon (Gastropoda, Hygromiidae)

Eike Neubert¹, Michel Bariche²

¹ Naturhistorisches Museum der Burgergemeinde Bern, Bernastrasse 15, CH-3005 Bern, Switzerland

² Marine Biology & Ichthyology, Biology Dep., American University of Beirut

Corresponding author: Eike Neubert (eike.neubert@nmbe.ch)

Academic editor: Andrzej Wiktor | Received 25 April 2013 | Accepted 13 June 2013 | Published 20 June 2013

Citation: Neubert E, Michel Bariche M (2013) On the *Monacha* species of Lebanon (Gastropoda, Hygromiidae). ZooKeys 311: 1–18. doi: 10.3897/zookeys.311.5408

Abstract

In this paper, all seven hitherto known species of the hygromiid genus *Monacha* from Lebanon are briefly characterised and illustrated, and distribution maps are supplied (*Monacha (Monacha) syriaca* (Ehrenberg, 1831), *Monacha (Monacha) nummus* (Ehrenberg, 1831), *Monacha (Monacha) obstructa* (L. Pfeiffer, 1842), *Monacha (Monacha) crispulata* (Mousson, 1861), *Monacha (Monacha) solitudinis* (Bourguignat, 1852), *Monacha (Monacha) bari* Forcart 1981, and *Monacha (Monacha) cf. compingtae* (Pallary, 1929)). One species, *M. (M.) bari* Forcart, 1981, is recorded for the country for the first time, and its relationship to *M. (M.) compingtae* (Pallary, 1929) is discussed. Based on recently collected specimens, the genital organs of a long time ignored species, *Helix solitudinis* Bourguignat, 1852 could be investigated. It is here re-described as a *Monacha* species endemic for Lebanon.

Keywords

Hygromiidae, *Monacha*, Lebanon, re-description

Introduction

The malacofauna of Lebanon has been the focus of malacologists since quite early on. Among the earliest scientists collecting molluscs in this country, the famous German zoologists Friedrich Wilhelm Hemprich and Christian Gottfried Ehrenberg from Berlin, who visited the area during their second voyage from 1821–1825, have to be mentioned. Their results were subsequently published by Ehrenberg (1828–1833). Since

then, the country has been the target of many expeditions where zoological and/or botanical objects were collected. Already at that time, Beirut served as “headquarters” for collectors, and thus, nearby and easily accessible places like Wadi Beyrouth or Wadi Nahr el Kelb became type localities of dozens of species.

At the beginning of the 20th century, a French expedition lead by Henri Gadeau de KERVILLE collected an enormous amount of continental molluscs from Syria in the wider sense, which were treated in two large volumes by Germain (1921). These specimens are housed in the MNHN in Paris. Later, the French malacologist Pallary, a keen connoisseur of the malacofauna of Northern Africa and home-based in Oran, Algeria, published some papers on the wider Palestinian area (Pallary 1929, 1939). Finally, the famous Lebanese zoologists Henriette and George Thomé presented a small comprehensive volume covering the terrestrial malacofauna of Lebanon (Thomé and Thomé 1988). Since then, no papers dealing exclusively with Lebanon have been published, and occurrences of continental molluscs have only been mentioned as by-products in papers having another scope. Quite recently, Bössneck (2011) reported on the results of his field trip to the country, where he (among others) mentioned two interesting species of *Monacha* from the Lebanon Mountains. These findings stimulated the present authors to amend these records with the results of their own research.

In Turkey, there is a large variety of species of the genus *Monacha*, where currently 54 species are known, with still several species remaining undescribed because of the low number of specimens available, or of lack of anatomical data (Hausdorf 2000). Nonetheless, the mountains stretching towards the south house a few more species of the genus, some of them are endemic, not well studied as of today, and have been only inadequately illustrated thus far. It is also our aim to raise awareness of this more or less ignored group of invertebrate animals living in Lebanon to local people, laymen as well as students.

Material and methods

This paper summarises data obtained during two excursions by the authors in Lebanon in 2008 and 2011, and others by Bariche during the last years. Specimens were collected by hand, and stored in the respective institutional collections. The section “specimens examined” contains only information that has been personally confirmed by the authors, i.e. none of the records have been retrieved from the literature. The records of Boessneck have been verified by Hausdorf (and a few by Neubert), and have been added to the distribution data. Finally, it also includes all data related to the *Monacha* species from Lebanon, its vouchers can be found in the Pallary collection housed at the AUBM. All specimens were checked by Neubert during his visit at the AUBM in October 2011. Unfortunately, the whereabouts of the Thomé’s collection remains unclear. Inquiries at the MHNG, at the MNHN, and even personal contact of M. Bariche with the Thomés’ yielded no clarification. Thus, the voucher specimens listed in the book by Thomé and Thomé (1988) are not available at the moment.

All measurements in mm.

Abbreviations for institutions

AUBM	American University of Beirut, mollusc collection, Lebanon.
MHNB	Naturhistorisches Museum Basel, Switzerland.
MHNG	Muséum d'histoire naturelle de la ville de Genève, Switzerland.
MNHN	Museum national d'histoire naturelle Paris, France
MZL	Muséum d'Zoologie, Lausanne, Switzerland.
NMBE	Naturhistorisches Museum der Burgergemeinde Bern, Switzerland.
SMNS	Staatliches Museum für Naturkunde Stuttgart, Germany

Abbreviations for technical terms

H	shell height
D	shell diameter
PH	peristome height
PD	peristome diameter

Results

Familia Hygromiidae Tryon, 1866

Subfamilia Monachinae Wenz, 1930 (1904)

Tribus Monachaini Wenz, 1930 (1904)

Genus *Monacha* (*Monacha*) Fitzinger, 1833

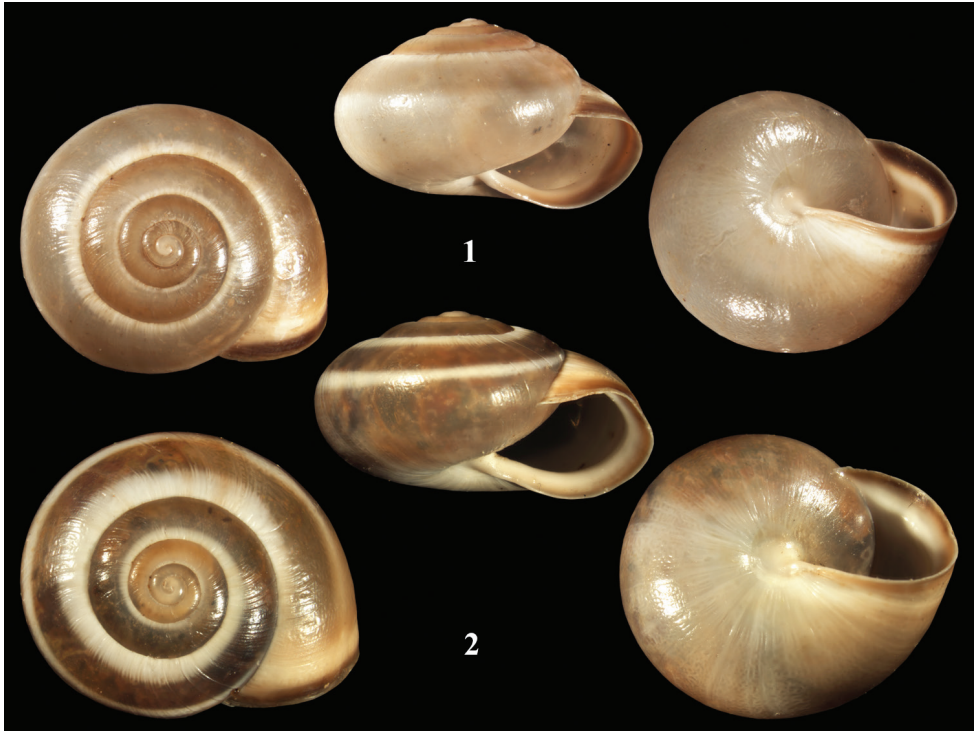
***Monacha* (*Monacha*) *syriaca* (Ehrenberg, 1831)**

http://species-id.net/wiki/Monacha_syriaca

Figs 1–3

1831 *Helix syriaca* Ehrenberg, Symbolae Physicae: 7 [Berytum].

Specimens examined: NMBE 506363/1, Hlaliye, 33.8393°N, 35.636°E, 475 m alt., 28.10.2011; NMBE 506362/2, Ajhbe, 34.0087°N, 35.7108°E, 1164 m alt., 27.10.2011; NMBE 506361/5, Aaramoun, 34.0153°N, 35.6973°E, 760 m alt., 27.10.2011; NMBE 506360/7, above Kfour, 34.0317°N, 35.6988°E, 895 m alt., 27.10.2011; NMBE 506359/3, Kfour, 34.0348°N, 35.6952°E, 817 m alt., 27.10.2011; NMBE 506358/2, Adma, 34.0166°N, 35.6489°E, 85 m alt., 27.10.2011; NMBE 505184/1, above Kfour, 34.0317°N, 35.6988°E, 895 m alt., 27.10.2011 (preserved); NMBE 25914/3, Trablous (= Tripoli), 34.43°N, 35.85°E, coll. Shuttleworth ex Verreaux; NMBE 508014/20, road between Ain Et Tine and Machgara, 33.516°N, 35.644°E, 1080 m alt., 21.08.2008; NMBE 508013/2, Jezzine, Mazrad el Mathané, 33.576°N, 35.519°E, 450 m alt., 21.08.2008; NMBE 508012/2, Nahr Bisri, 33.58°N, 35.535°E, 400 m alt., 21.08.2008;



Figures 1–2. *Monacha (Monacha) syriaca*. **1** NMBE 508006, Nahr Abu Ali close to Seraad, D = 10.12 mm. **2** NMBE 508001, Ain al Jdeidi, D = 11.12 mm. – $\times 4$, phot. E. Bochud.

NMBE 508011/64, Jezzine, W of village, steep slope, 33.546°N, 35.575°E, 940 m alt., 21.08.2008; NMBE 508010/15, Nahr Ibrahim, Chouene, trail Chouene–Chouwan Lake, 34.081°N, 35.785°E, 450 m alt., 20.08.2008; NMBE 508009/6, Nahr Ibrahim, close to the estuary, 34.065°N, 35.643°E, 20.08.2008; NMBE 508008/1, Qanater al Zbeideh, Nahr Beirut, 33.85°N, 35.556°E, 80 m alt., 19.08.2008; NMBE 508007/10, Byblos, 34.122°N, 35.652°E, 20 m alt., 19.08.2008; NMBE 508006/72, Nahr Abu Ali close to Seraad, 34.382°N, 35.93°E, 630 m alt., 19.08.2008; NMBE 508005/3, Tourza, 34.281°N, 35.89°E, 870 m alt., 19.08.2008; NMBE 508004/32, Jazire, Nahr al Qasimiyah (part of the Litani), 33.32°N, 35.288°E, 15 m alt., 18.08.2008; NMBE 508003/2, Nahr al Damour, 33.704°N, 35.452°E, 20 m alt., 18.08.2008; NMBE 508002/1, Nahr Kfar Matta at Jisr el Kadi, 33.721°N, 35.558°E, 260 m alt., 17.08.2008; NMBE 508001/75, Ain al Jdeidi, 33.809°N, 35.63°E, 918 m alt., 17.08.2008; NMBE 508000/1, Nahr el Kalb, upper reach of Nahr al-Kalb near Daraya, 33.955°N, 35.706°E, 700 m alt., 16.08.2008; NMBE 507999/7, Nahr el Kalb, Jeita Grotto, cave entrance, 33.944°N, 35.641°E, 90 m alt., 16.08.2008; NMBE 507998/2, Nahr el Kelb, road to Jeita Grotto, limestone slope, 33.947°N, 35.637°E, 115 m alt., 16.08.2008; AUBM-MOLL0470/5, Ghazir, 34.0302°N, 35.6701°E, 326 m alt., 25.10.2009; AUBM-MOLL0473/13, Habboub, 34.1284°N, 35.686°E,

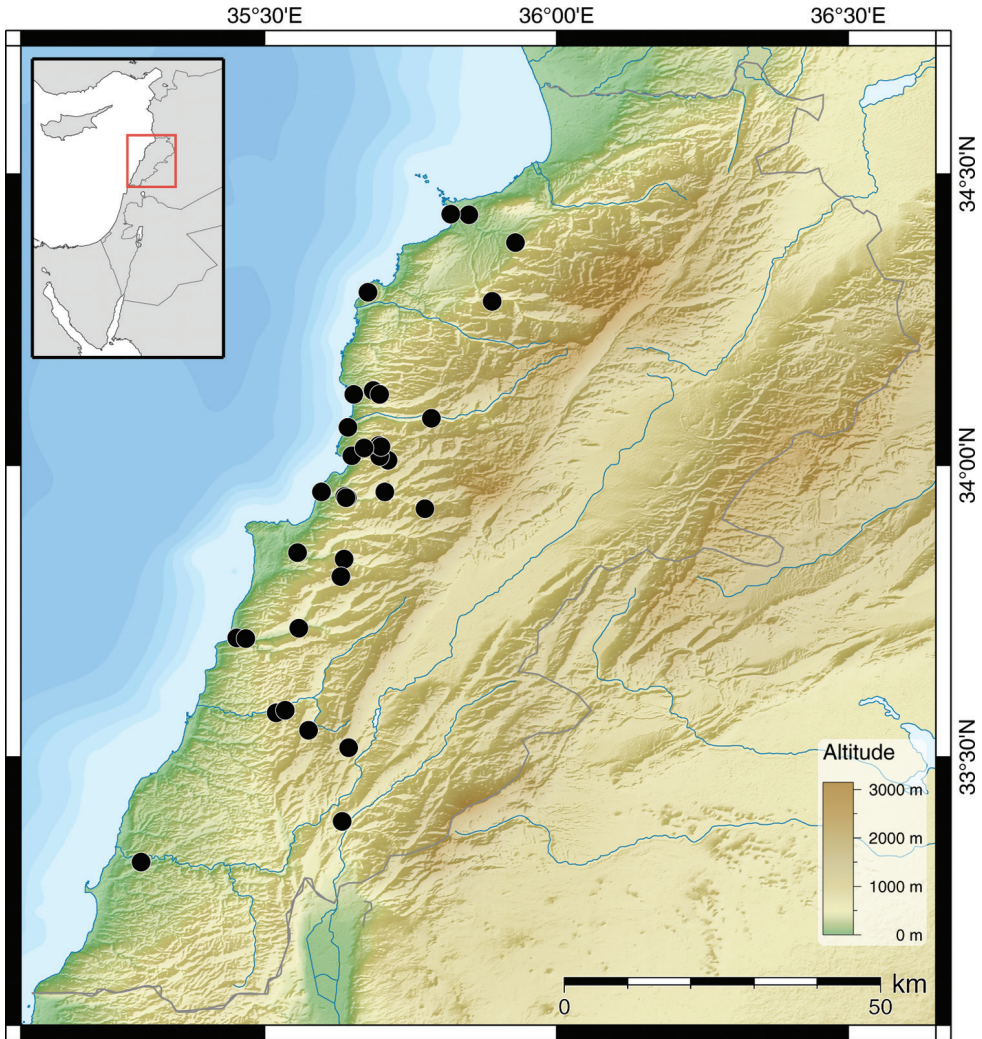


Figure 3. Distribution of *Monacha (Monacha) syriaca*.

420 m alt., 11.10.2009; AUBM-MOLL0476/7, Jeita, 33.9448°N, 35.6387°E, 88 m alt., 26.09.2009; AUBM-MOLL0478/1, Nahr el Kelb, 33.9547°N, 35.5975°E, 30 m alt., 26.09.2009; AUBM-MOLL0481/1, Hamat, 34.2967°N, 35.6768°E, 100 m alt., 16.01.2010; AUBM-MOLL0482/4, Kfar Kouass, 34.1222°N 35.6969°E, 630 m alt., 11.10.2009; AUBM-MOLL0485/3, Moultaa el Nahreyn, 33.7029°N, 35.4673°E, 110 m alt., 04.10.2009; AUBM-MOLL0487/5, Bteghrine, 33.9253°N, 35.7745°E, 1000 m alt., 07.10.2009; AUBM-MOLL0329/16, Jounieh, 33.3896°N, 35.6323°E, coll. Pallary; AUBM-MOLL0335/38, Tartous, 34.8905°N, 35.8707°E, coll. Pallary; AUBM-MOLL0337/40, Beirut, 33.8526°N, 35.4296°E, coll. Pallary; AUBM-MOLL0338/25, Tripoli, 34.4314°N, 35.8184°E, coll. Pallary.

Diagnosis. shell depressed, with two white spiral bands, malleate teleoconch sculpture, aperture reinforced by a strong white lip, apertural rim deep red, umbilicus closed by columellar callus.

Description. shell depressed, with an elevated broad conical spire, basic shell colour deep brown to yellowish brown, usually with a white subsutural band and a second white band at the shell's periphery; two protoconch whorls, smooth; teleoconch with a malleate sculpture (i.e. looks like markings of a small hammer) and very fine radial growth lines, surface with a glossy shine; last whorl rounded to slightly compressed forming a blunt shoulder; last whorl abruptly bent towards the aperture; aperture oval and depressed, reinforced by a strong white lip; apertural rim sharply bounded, usually deep red to brownish; umbilicus always closed by a small columellar callus.

Measurements (figured specimens). Fig. 1: H = 6.26; D = 10.12; PH = 3.17; PD = 5.38. Fig. 2: H = 6.18; D = 11.12; PH = 3.59; PD = 6.11.

Distribution. This species is found widespread along the coastal areas in Lebanon. It usually lives at lower altitudes, but can occasionally be found up to 1000 m in more sheltered areas such as in the bottom of deep valleys.

Remarks. This species cannot be mistaken for any other species in the area due to its characteristic colour pattern, teleoconch sculpture, and closed umbilicus.

Monacha (Monacha) nummus (Ehrenberg, 1831)

http://species-id.net/wiki/Monacha_nummus

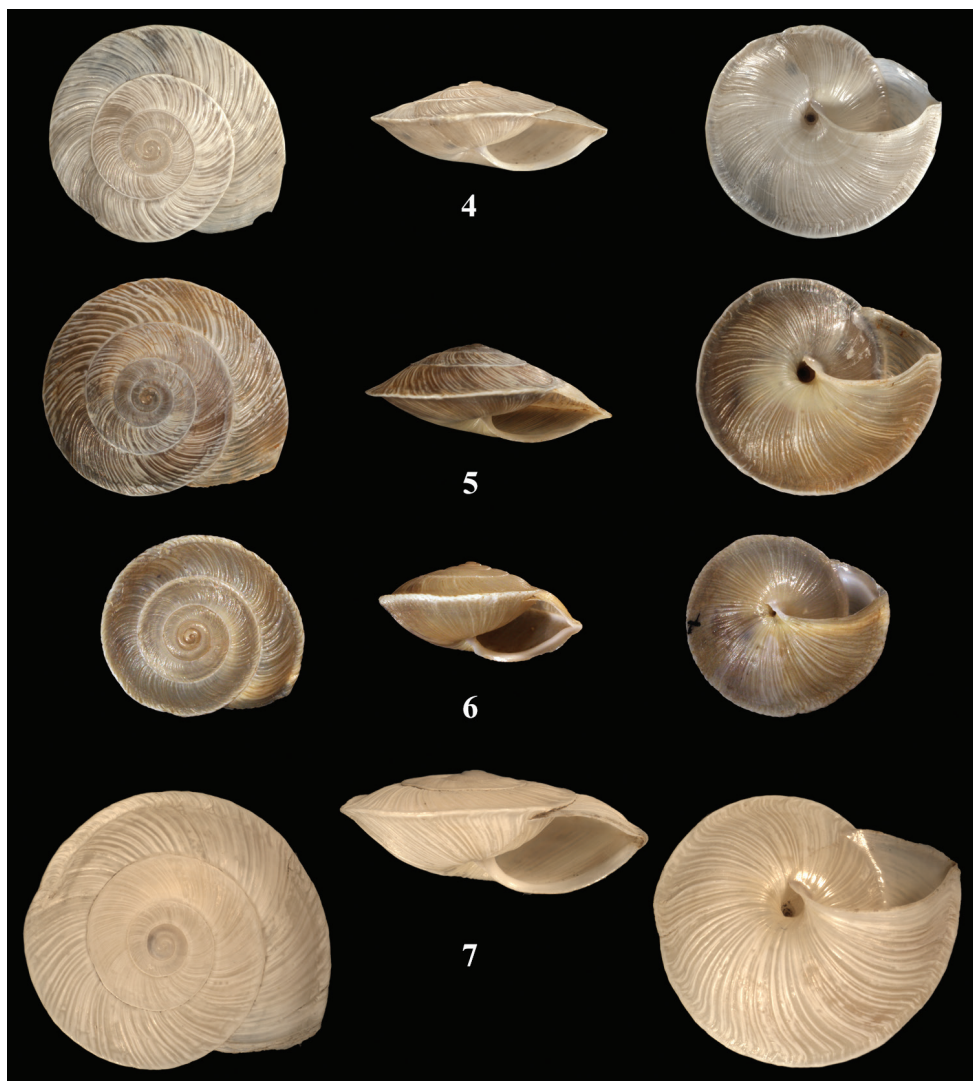
Figs 4, 5, 8

1831 *Caracolla nummus* Ehrenberg, Symbolae Physicae: 8 [Syrien].

1847 *Helix oxygyra* Charpentier, Zeitschrift für Malakozoologie, 4: 131 [Bei Nahr und Kelb [sic!], zwei Lieues nördlich von Beirut, an Felsen].

Type specimens. *oxygyra*: syntype in MZL.

Specimens examined. NMBE 508022/6, Nahr Ibrahim, Chouene, trail Chouene–Chouwan Lake, 34.081°N, 35.785°E, 450 m alt., 20.08.2008; NMBE 508021/2, Nahr Ibrahim, at the road between Chouaya and Yahchouch, 34.064°N, 35.728°E, 780 m alt., 20.08.2008; NMBE 508020/3, Nahr Ibrahim, 34.079°N, 35.679°E, 80 m alt., 20.08.2008; NMBE 508018/1, Qanater al Zbeideh, Nahr Beirut, 33.85°N, 35.556°E, 80 m alt., 19.08.2008; NMBE 508017/5, Nahr Abu Ali close to Seraad, 34.382°N, 35.93°E, 630 m alt., 19.08.2008; NMBE 508016/20, Nahr el Kelb, Jeita Grotto, cave entrance, 33.944°N, 35.641°E, 90 m alt., 16.08.2008; NMBE 508015/4, Nahr el Kelb, road to Jeita Grotto, limestone slope, 33.947°N, 35.637°E, 115 m alt., 16.08.2008; NMBE 506357/1, Wadi Abadie, 33.8434°N, 35.6344°E, 310 m alt., 28.10.2011; NMBE 506356/4, Kfar Hbab, 34.006°N, 35.6552°E, 113 m alt., 27.10.2011; NMBE 506355/3, Kfour, 34.0348°N, 35.6952°E, 817 m alt., 27.10.2011; NMBE 506354/12, Adma, 34.0166°N, 35.6489°E, 85 m alt., 27.10.2011; NMBE 25953/5, Beirut, 33.8526°N, 35.4296°E, coll. A. Vogt ex Bohny;



Figures 4–7. *Monacha* spp. **4** *Monacha (Monacha) nummus*, syntype *Helix oxygyra* Charpentier 1847, D = 15.55 mm **5** *Monacha (Monacha) nummus*, NMBE 508105, Beirut, Nahr el Kalb, Jeita Grotto, limestone rocks above entrance area to the cave, D = 16.15 mm **6** *Monacha (Monacha) spiroxia* (Bourguignat 1868), lectotype MHNG 15378, Turkey, Alexandrette (= Iskenderun), D = 13.4 mm **7** *Monacha (Monacha) carinata* Hausdorf, 2000, holotype SMNS 50426, Turkey, Iskenderun, canyon of Sariseki, Iskenderun, D = 20.2 mm. – $\times 2$, phot. Neubert, Richling.

AUBM-MOLL0466/9, AUBM-MOLL0478/1, AUBM-MOLL0480/13, Nahr el Kelb, 33.9547°N, 35.5975°E, 30 m alt., 26.09.2009; AUBM-MOLL0468/8, AUBM-MOLL0470/2, AUBM-MOLL0489/1, Ghazir, 34.0302°N, 35.6701°E, 326 m alt., 25.10.2009; AUBM-MOLL0476/5, Jeita, 33.9448°N, 35.6387°E, 88 m alt., 26.09.2009; AUBM-MOLL0552/2, Nahr Ibrahim, 34.0824°N, 35.802°E, 780 m alt.;

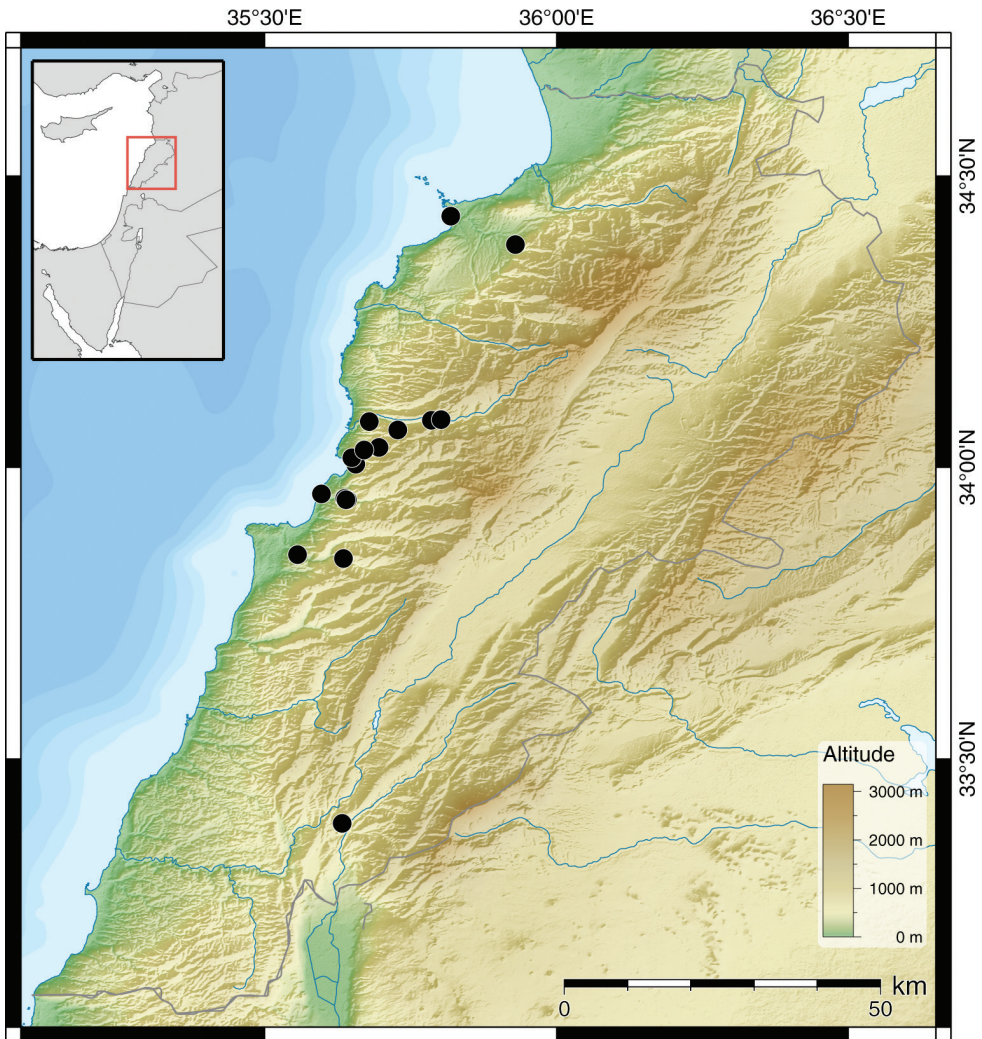


Figure 8. Distribution of *Monacha (Monacha) nummus*.

AUBM-MOLL0343/10, Nahr el Kelb, coll. Pallary; AUBM-MOLL0344/19, Tripoli, 34.4314°N, 35.8184°E, coll. Pallary; AUBM-MOLL0345/9, Jounieh, 33.3896°N, 35.6323°E, coll. Pallary.

Diagnosis. shell large, biconvex, periphery with a sharply bounded keel, umbilicus narrowly opened, broad periomphalum.

Description. shell large and flat, biconvex, shell colour grey to brown; protoconch consisting of two smooth whorls; teleoconch whorls with a dense pattern of white ribs, that extend to the glossy underside of the shell; suture marked by a white sutural thread; ca. six regularly increasing whorls, periphery with a sharply bounded crimped keel, sometimes spire stepped because of whorls attaching somewhat below the keel

during shell growth; last whorl not descending; aperture obliquely oval with an acute palatal edge, sometimes with a faint white lip; peristomial rim sharply bounded and simple; umbilicus always open, narrow, with a broad periomphalum.

Measurements. Syntype *oxygyra*: H = 5.91; D = 15.55; PH = 3.73; PD = 8.84. Fig. 5: H = 6.53; D = 16.15; PH = 3.39; PD = 8.67.

Distribution. This species seems to be restricted to coastal regions in Lebanon.

Remarks. This species can easily be confused with other keeled species known from the Levante area. To illustrate the differences, shells of *M. (M.) spiroxia* (Bourguignat, 1868) (Fig. 6) and *M. (M.) carinata* Hausdorf, 2000 (Fig. 7) are given. They differ in that the last has a larger shell with a more flat spire and a less pronounced keel. So far, these two species have only been recorded from the Hatay province of Turkey, while *M. (M.) nummus* seems to be restricted to Lebanon. *Monacha (M.) spiroxia* differs from *M. (M.) nummus* by being smaller, having a white lip in the aperture, a much narrower umbilicus and a teleoconch sculpture of small granules.

***Monacha (Monacha) obstructa* (L. Pfeiffer, 1842)**

http://species-id.net/wiki/Monacha_obstructa

Figs 9, 10

1842 *Helix obstructa* L. Pfeiffer, Symbolae ad Historiam Heliceorum II: 35 [Tripoli].

Specimens examined. NMBE 508032/12, Jazire, Nahr al Qasimiyah (part of the Litani), 33.32°N, 35.288°E, 15 m alt., 18.08.2008; NMBE 508031/32, Nahr al Damour, 33.704°N, 35.452°E, 20 m alt., 18.08.2008; NMBE 508030/7, Nahr el Kalb, Jeita Grotto, cave entrance, 33.944°N, 35.641°E, 90 m alt., 16.08.2008; AUBM-MOLL0492/1, Campus of the American University Beirut, 33.9016°N, 35.4781°E, 26.09.2009.

Diagnosis. shell small, cream white, aperture with thick white labial callus, umbilicus closed, forming a characteristic funnel-shaped “pseudo-umbilicus”.

Description. shell medium sized, spire broadly conical and somewhat elevated; protoconch consisting of 1.5 smooth whorls; teleoconch cream white, with faint riblets and a malleate sculpture (compare to *M. syriaca*), and evenly rounded whorls; last whorl slightly bending towards the aperture; aperture broadly oval, reinforced by a thick white labial callus, well discernible from the outside as a thick white callosity; peristomial rim sharply bounded, red; umbilicus closed, but last whorl forming a characteristic funnel-shaped “pseudo-umbilicus”.

Measurements (Fig. 9): H = 6.68; D = 11.83; PH = 4.2; PD = 5.85.

Distribution. In Lebanon this species seems to be quite restricted, probably because it is confined to more arid or steppe-like areas like central Syria, where it is one of the most abundant snail species. These environmental conditions are not present in the western part of Lebanon with its humid and steep mountain slopes. However, it is the most widespread species within *Monacha*. Its distribution ranges from Egypt



Figures 9, 10. *Monacha* spp. **9** *Monacha (Monacha) obstructa*. NMBE 508031, Nahr al Damour, D = 11.83 mm **10** *Monacha (Monacha) crispulata*, Jordan, Ajlune Castle, 25.06.2007, leg. Z. Amr, D = 8.3 mm. – $\times 4$, phot. E. Bochud.

to Pakistan, and Turkey to Saudi Arabia (Neubert 1998), probably because it can be easily distributed by human activities.

Remarks. This species is unmistakable for its funnel-shaped “pseudo-umbilicus”. Subadult shells can be differentiated from *M. syriaca* by the uniformity of its cream white shell.

***Monacha (Monacha) crispulata* (Mousson, 1861)**

http://species-id.net/wiki/Monacha_crispulata

Figs 10, 11

1861 *Helix crispulata* Mousson, Vierteljahresschrift der Naturforschenden Gesellschaft Zürich 6: 12 [Jerusalem ex Roth].

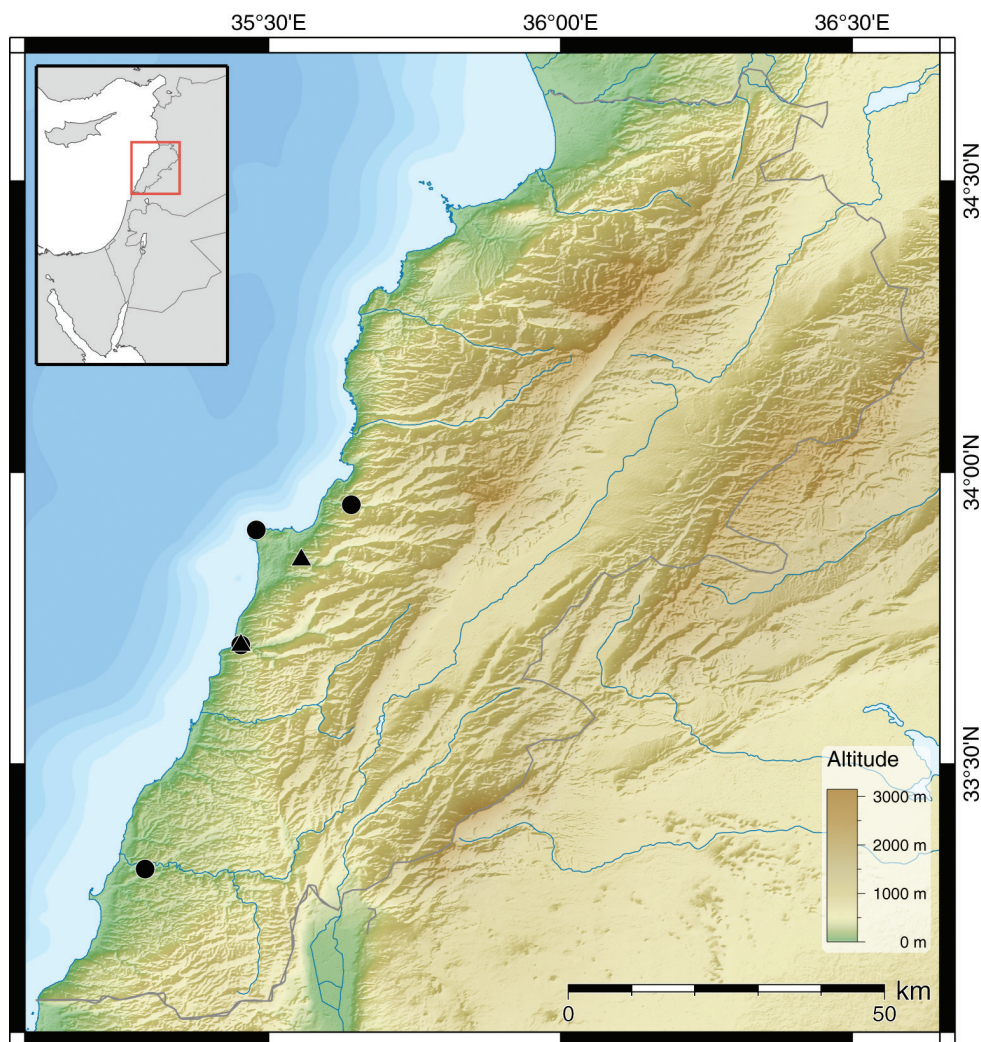


Figure 11. Distribution of *Monacha (Monacha) obstructa* (●) and *Monacha (Monacha) crispulata* (▲).

Specimens examined. NMBE 515473/1, Nahr al Damour, 33.704°N, 35.452°E, 20 m alt., 18.08.2008; NMBE 515474/1, Qanater al Zbeideh, Nahr Beirut, 33.85°N, 35.556°E, 80 m alt., 19.08.2008.

Diagnosis. shell small, teleoconch whorls with strong axial ribs, granules and long hairs.

Description. shell small, spire slightly elevated; protoconch consists of 1.5 smooth whorls; teleoconch greyish to brownish, with strong axial ribs, whorls covered by a dense sculpture of small granules and very long, soft hairs; hair scars visible even in eroded shells; suture moderately deep; last whorl slightly bending towards the aperture; aperture broadly oval, reinforced by a white lip (discernible from the outside);

peristomial rim sharply bounded; umbilicus open, partly covered by a triangular columellar callus.

Measurements (Fig. 11): H = 4.72; D = 8.3; PH = 2.08; PD = 4.76.

Distribution. This seems to be a rare species in Lebanon. It has probably been overlooked because of its small size, or even misidentified, because its hairs can easily fall off, and the remaining shell is less characteristic. It is possible that this species is more widespread.

Remarks. This species differs from all Lebanese *Monacha* species by its haired shell. Even eroded shells can be identified when using higher magnifications to see the axial ribs in combination with the granulated surface and remaining hair scars.

***Monacha (Monacha) solitudinis* (Bourguignat, 1852)**

http://species-id.net/wiki/Monacha_solitudinis

Figs 12–14, 17

1852 *Helix solitudinis* Bourguignat, Testacea novissimae...: 15 [Circa Heliopolim Syriae (Baalbek) habitat].

1859 *Helix solitudinis*, – L. Pfeiffer, Monographia heliceorum viventium IV: 280.

1868 *Helix solitudinis*, – L. Pfeiffer, Monographia heliceorum viventium V: 365.

1902 *Helicella (Monacha) solitudinis*, – Gude, Journal of Malacology, 9: 128.

1921 *Helix (Monacha) solitudinis* – Germain, Mollusques terrestres et fluviatiles de Syrie I: 156.

1939 *Metafruticicola solitudinis*, – Pallary, Mémoires présentés à l'Institut d'Égypte 39: 8.

2011 *Monacha* sp., – Boessneck, Zoology in the Middle East, 54: figs 5a, 42, 48.

Type specimens. Figured syntype NHMG 16032a; second syntype NHMG 16032b; Baalbek 34.0061°N, 36.2122°E.

Specimens examined. Bcharre, Jabal el Mekmel ca. 7 km SE of Bcharre, loose calcareous vegetation, 34.2138°N, 36.0683°E, 2600 m alt, leg. U. Boessneck et al.; NMBE 508041/1, NMBE 508044/12, AUBM-MOLL, Falougha above Soha water plant, 33.835°N, 35.756°E, 1610 m alt., 17.08.2008; NMBE 508043/3, Falougha next to Soha water plant, 33.835°N, 35.752°E, 1474 m alt., 17.08.2008; NMBE 508042/8, Falougha, 33.8352°N, 35.7561°E, 1591 m alt., 29.10.2011; NMBE 515472/1, ditto (preserved).

Diagnosis. shell medium sized, with a white spiral band at the periphery of the last whorl, aperture subrectangular, umbilicus open, cylindrical, penial papilla large and stout, flagellum longer than epiphallus.

Re-description. shell medium sized, spire depressed; protoconch consists of 2 smooth whorls; shell colour light brown to yellowish brown with a white spiral band at the periphery of the last whorl; teleoconch with a malleate sculpture and fine axial riblets; suture deep, simple; teleoconch with up to seven densely coiled whorls; last whorl slightly bending towards the aperture; aperture broadly subrectangular, reinforced by a

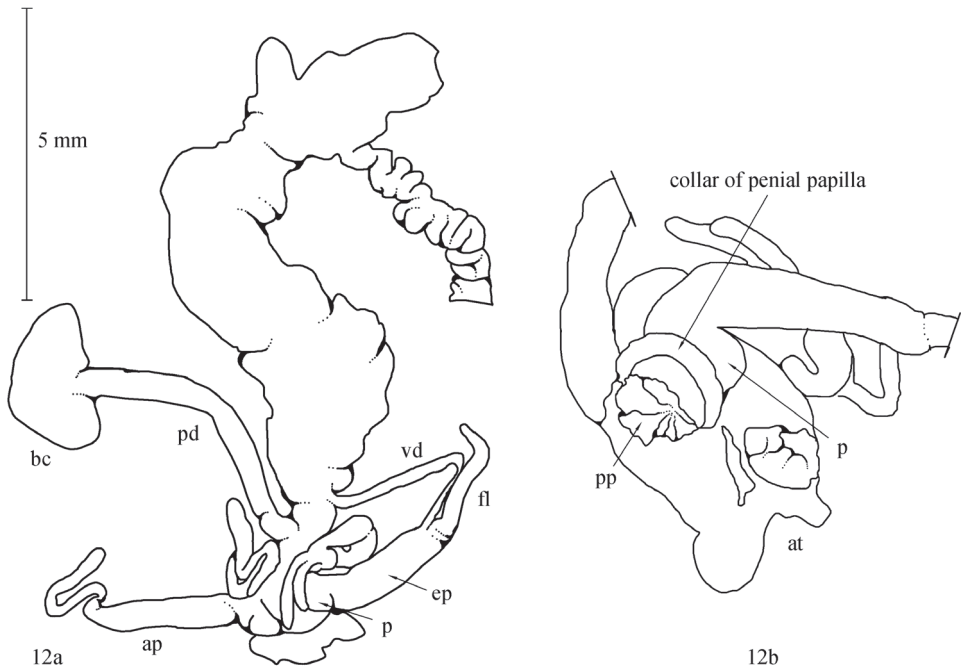


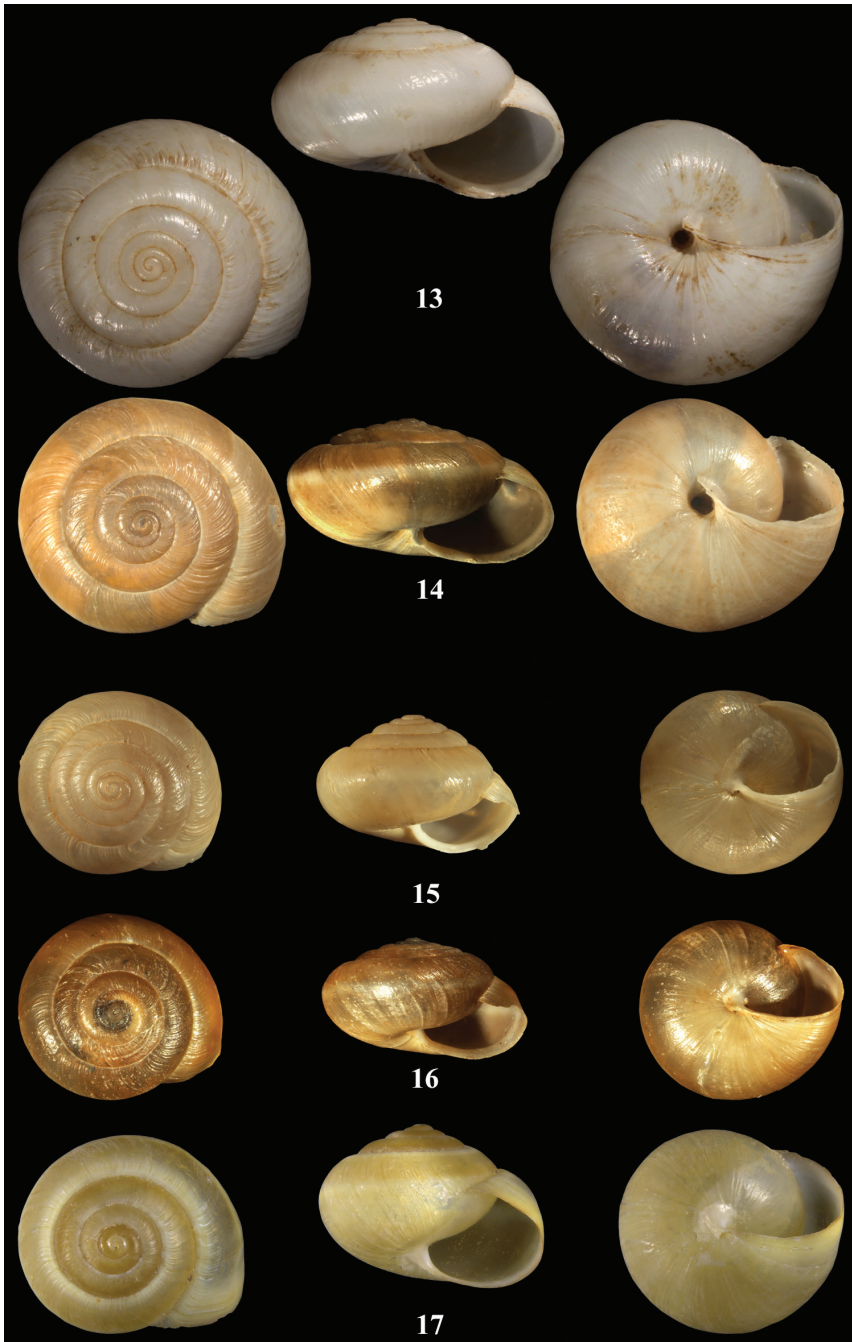
Figure 12. Anatomical drawing *Monacha (Monacha) solitudinis* **a** NMBE 515472, situs of genital organs, total length 10.9 mm **b** detail of atrium and penial papilla, not to scale. Abbreviations used: ap = appendicula; at = atrium; bc = bursa copulatrix; ep = epiphallus; fl = flagellum; ped = pedunculus; pp = penial papilla; vd = vas deferens.

small white lip; peristomial rim sharply bounded, simple; umbilicus open, cylindrical, a triangular columellar callus is indicated.

Genital organs (Figs 12a, b): The only specimen that could be investigated was quite strongly contracted; it is a species of *Monacha* sensu str., because the genitalia show an appendicula, but the penial retractor muscle is missing. Penis very short (0.8 mm), penial papilla large, stout with a central perforation completely filling the atrial lumen (preservation artefact?), basis of the papilla with a strong collar (Fig. 12b); epiphallus a thick-walled tube, its lumen filled with several finely crenulated pilasters (1.9 mm); flagellum short (2.1 mm), but surmounting the length of the epiphallus; appendicula (4.9 mm) branching off the atrium, subdivided in a thicker basal part, and a shorter part with a narrower lumen; two glandulae mucosae poorly ramified, inserting in the middle of the vagina; vesicle of bursa copulatrix large, hammer-like, pedunculus quite long (4.7 mm); atrial lumen with two pilasters, one of them large, knob-like, the other narrow and elongate; right ommatophoran retractor passes left to the genital organs.

Measurements. Syntype figured: H = 9.2; D = 14.6; PH = 4.81; PD = 7.92. Fig. 14: H = 7.28; D = 13.31; PH = 4.34; PD = 7.32.

Distribution. This species has only been recorded from two localities in Lebanon so far. The type locality Baalbek is not too far away and they were probably collected



Figures 13–17. *Monacha* spp. **13** *Monacha (Monacha) solitudinis*. 13 syntype NHMG 16032a, Baalbek, D = 14.6 mm **14** NMBE 508041, Falougha above Soha water plant, D = 13.31 mm **15** *Monacha (Monacha) bari*, paratype NHMB 11167a, Israel, Mt. Hermon, D = 10.13 mm **16** *Monacha (Monacha) bari*, NMBE 508040, below Barouk Cedar Forest, D = 10.19 mm **17** *Monacha (Monacha) cf. compingtae*, Aammig, D = XX mm. – $\times 4$, phot. E. Bochud.

on the way towards Baalbek. Both new localities are in the central chain of Mt. Lebanon at high altitudes. The habitats are characterised by coarse limestone boulders with interspersed subalpine grassland vegetation and is covered by snow during winter. In October 2011, two living animals were found actively crawling over the rocks (ca. 10°C and heavy rainfall), while in August 2008, only dead shells were encountered.

Differential diagnosis. This species differs from all other species in the Levante area by the shape of its umbilicus, which is cylindrical like a borehole. All other *Monacha* species with an open umbilicus deeply differ by either having a keeled (*M. nummus*) or a smaller and hairy shell (*M. crispulata*). *Monacha bari* and *M. compingtae* are conchologically similar to *M. solitudinis*, but these species have a narrow and almost closed umbilicus (refer to the discussion of *M. bari*). Information on the genital organs of the two latter species is scarce, only Hausdorf (2000: 84) reported on a subadult specimen of *M. compingtae* from southern Turkey. In this species, the penial papilla is short (large but stout in *M. solitudinis*), the epiphallus is longer than the flagellum (vice versa in *M. solitudinis*), and the appendicula is much shorter than in *M. solitudinis*. The specimen figured by Boessneck (2011) was re-investigated here; it matches the type specimen from Baalbek quite well, it is somewhat larger than the specimens from Falougha.

Remark. The correct generic affiliation of this species remained unclear until now. In his original description, Bourguignat compared it to his *Helix camelina*, which is a species of the Oxylidae; this text was uncritically copied by Pfeiffer (1859). Gude (1902) was the first to list it under the genus *Monacha*, but without any justification for doing so. Germain (1921), who remarked that this affiliation needs to be corroborated by an investigation of the genital organs, shared this point of view. Later, Pallary (1939) studied the type specimen and concluded that it should be classified within the genus *Metafruticiola*.

Monacha (Monacha) bari Forcart, 1981

http://species-id.net/wiki/Monacha_bari

Figs 15, 16, 18

1981 *Monacha (Monacha) bari* Forcart, Basteria, 45: 105, fig. 11a–d [Hermon, Abstieg von der unteren Skiliftstation zum Nahal Arar (= Wadi Assal) 1600–1400 m ü.M. (etwa 33°15'N, 35°15'E)].

Type specimens. *bari*: paratype MHN 11167a.

Specimens examined. NMBE 508040/11, below Barouk Cedar Forest, 33.7042°N, 35.6997°E, 1310 m alt., 29.10.2011; NMBE 508039/4, Qaroun Lake, 33.551°N, 35.682°E, 870 m alt., 21.08.2008.

Diagnosis. shell small, spire slightly elevated, teleoconch has a faint white spiral band and fine axial riblets, umbilicus narrow closed by columellar callus.

Description. shell small, spire conical, slightly elevated; protoconch consists of 1.5 smooth whorls; shell colour light brown to reddish brown, sometimes with a faint white spiral band at the periphery of the last whorl; teleoconch with fine axial riblets and a

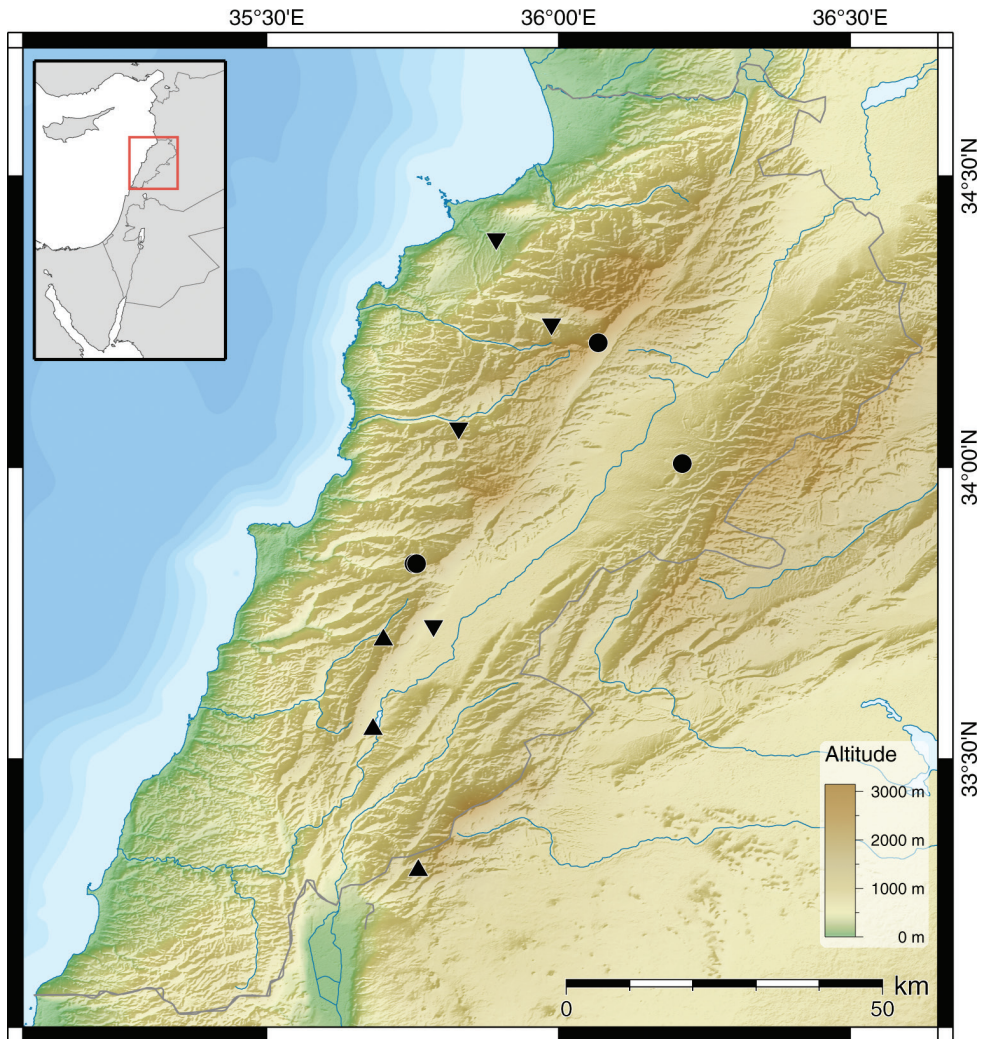


Figure 18. Distribution of *Monacha (Monacha) solitudinis* (●), *Monacha (Monacha) bari* (▲) and *Monacha (Monacha) cf. compingtae* (▼).

pattern of fine spiral threads (only visible under higher magnification); suture deep, simple; teleoconch with up to seven tightly coiled whorls; last whorl slightly bending towards the aperture; aperture broadly oval, reinforced by a white lip; peristomial rim sharply bounded, simple; umbilicus narrow, closed by a triangular columellar callus.

Measurements. Paratype *M. bari* figured: H = 6.9; D = 10.13; PH = 2.83; PD = 5.5. Fig. 16: H = 6.05; D = 10.19; PH = 2.94; PD = 6.05.

Distribution. This species is here recorded for the first time for Lebanon. It seems to be restricted to higher altitudes up to the alpine region of the highest mountains in the Eastmediterranean.

Remarks. The coordinates of the type locality of *M. bari* given by Forcart are erroneous, in fact they should be approximately at 33.3088°N, 35.7596°E. After careful examination of the paratypes of *M. bari* from the collection of Forcart (housed in the Natural History Museum in Basel, Switzerland), the recently collected specimens from Lebanon are here identified with this species. It is not possible to find any shell character that could justify a separation of the populations on specific level.

***Monacha* (*Monacha*) cf. *compingtae* (Pallary, 1929)**

Fig. 17

1929 *Theba compingtae* Pallary, Mémoires présentés à l'Institut d'Égypte 12: 7, pl. 1, figs 4–6.

2011 *Monacha* cf. *compingtae*, Boessneck, Zoology in the Middle East, 54: figs 5a, 42, 48.

Specimens examined. Zgharta (Prov. North Lebanon), dry ruderal vegetation with calcareous stones in side valley of Nahr Abou Ali above the village, 34.3922°N, 35.8929°E, 80 m alt., leg. U. Boessneck et al.; Hadchit (Prov. North Lebanon), slopes of the Nahr Quadicha, open calcareous fields, locally with loose vegetation and shrubs, partly ruderal impacted, 34.2459°N, 35.9881°E, 1000–1250 m alt., leg. U. Boessneck et al.; Aammiq (Prov. Bekaa), rocky open fields with loose vegetation at the edge of a wetland, ruderal impacted, 33.7284°N, 35.7858°E, 870 m alt., leg. U. Boessneck et al.; Afqa (Prov. Mount Lebanon), open calcareous fields with loose vegetation on single shrubs near the cave, locally wet, 34.0671°N, 35.8291°E, 1160 m alt., leg. U. Boessneck et al.

This *Monacha* species was identified by Hausdorf (in Boessneck 2011) as *M. (M.) cf. compingtae* (Pallary, 1929). We were able to check the specimen figured by Boessneck (2011, Fig. 5a) from Aammiq in order to compare it to the species we finally identified as *M. bari*. Both species differ by the possession of a white subsutural spiral band in *Monacha* cf. *compingtae* (not seen in any specimen of *M. bari*), and the telioconch sculpture, which is almost malleate in *Monacha* cf. *compingtae*, while *M. bari* shows axial riblets combined with spiral threads.

Monacha compingtae was described from Tartus and Safita, both localities from the coastal area in southern Syria. Later, Pallary (1939) extended the distribution area of *M. compingtae* with more records from Syria (“kilomètre 139 de la route de Lattaquié à Alep” = probably somewhere NE of Idlib), “Koubba Cheikh Mokbel”, “entre Lattaquié et Kerdaha” and “Slenfé (Nahib Younès” = Slinfah 35.583°N, 36.°E)). Hausdorf (2000: 84) reported this species from the Hatay area in southern Turkey, which constitutes the northern part of the eastmediterranean mountain ridge. In fact, this species is conchologically very similar to *M. compingtae*, the two white spiral bands are already mentioned by Pallary in the original description. However, he also mentions a transversal striation on the shell, which is not seen on the specimen from Aammiq. It might well be that this taxon from Lebanon represents another yet undescribed species.

Acknowledgements

For their support during the fieldwork, the authors are very grateful to Dr Nisreen Alwan and Nancy al-Sayar; to Dr Ulrich Boessneck (Efurt), who deliberately sent specimens collected during his visit to Lebanon in 2008; Estée Bochud (Bern) for taking pictures and shell measurements; to the curators of the institutions listed above who gave access to the type specimens under their responsibility.

References

- Böessneck U (2011) New records of freshwater and land molluscs from Lebanon (Mollusca: Gastropoda & Bivalvia). *Zoology in the Middle East* 54: 35–52. doi: 10.1080/09397140.2011.10648879
- Bourguignat J-R (1852) *Testacea novissima quae Cl. de Saulcy in itinere per Orientem annis 1850 et 1851 collegit*. Lutetiae, Paris (Baillière), 31 pp.
- Charpentier J de (1847) Uebersicht der durch Herrn Edm. Boissier von einer Reise nach Palästina mit zurückgebrachten Conchylien-Arten. *Zeitschrift für Malakozoologie* 4: 129–144.
- Ehrenberg CG (1831) *Animalia Mollusca*. In: Hemprich FG, Ehrenberg CG: *Symbolae Physicae, seu icones et descriptiones corporum naturalium novorum aut minus cognitorum, quae ex itineribus per Libyam ... I. Pars zoologica. Animalia Evertebrata*, Berlin, 46 pp., 2 pl.
- Forcart L (1981) Neubeschreibungen von Landschnecken aus Palästina. *Basteria* 45 (4/5): 97–108.
- Germain L (1921–1926) *Mollusques terrestres et fluviatiles de Syrie dans Voyage zoologique d'Henri Gadeau de Kerville en Syrie*. J.B. Baillière et fils (Paris).
- Gude GK (1902) A classified list of the Helicoid land shells of Asia, IV. *The Journal of Malacology*, 9: 112–129.
- Hausdorf B (2000) The genus *Monacha* in Turkey (Gastropoda: Pulmonata: Hygromiidae). *Archiv für Molluskenkunde* 128 (1/2): 61–151.
- Mousson A (1861) Coquilles terrestres et fluviatiles recueillies par M. le Prof. J. R. Roth dans son dernier voyage en Orient. *Vierteljahresschrift der Naturforschenden Gesellschaft Zürich* 6: 1–34.
- Neubert E (1998) Annotated checklist of the terrestrial and freshwater Molluscs of the Arabian Peninsula. *Fauna of Arabia* 17: 333–461.
- Pallary P (1929) Première addition à la faune malacologique de la Syrie. *Mémoires présentés à l'Institut d'Égypte* 12: 1–43, pl. 1–3.
- Pallary P (1939) Deuxième addition à la faune malacologique de la Syrie. *Mémoires présentés à l'Institut d'Égypte* 39: 1–141, pl. 1–7.
- Pfeiffer L (1859) *Monographia heliceorum viventium IV*. Brockhaus (Lipsiae). 920 pp.
- Pfeiffer L (1868) *Monographia heliceorum viventium V*. Brockhaus (Lipsiae). 565 pp.
- Thomé G, Thomé H (1988) *Les coquillages terrestres du Liban*. Beyrouth, 113 pp.

Three new species in the harvestmen genus *Acuclavella* (Opiliones, Dyspnoi, Ischyropsalidoidea), including description of male *Acuclavella quattuor* Shear, 1986

Casey H. Richart^{1,†}, Marshal Hedin^{1,‡}

¹ Department of Biology, San Diego State University, 5500 Campanile Drive, San Diego, California, 92182, USA

[†] [urn:lsid:zoobank.org:author:9110BBBC-A1E0-488E-83EF-87FFBFE27AA9](https://doi.org/urn:lsid:zoobank.org:author:9110BBBC-A1E0-488E-83EF-87FFBFE27AA9)

[‡] [urn:lsid:zoobank.org:author:A845B9F9-25BB-45BC-A8D0-DF1B1FDF5D80](https://doi.org/urn:lsid:zoobank.org:author:A845B9F9-25BB-45BC-A8D0-DF1B1FDF5D80)

Corresponding author: Casey H. Richart (pileated@gmail.com)

Academic editor: Adriano Kury | Received 13 February 2012 | Accepted 6 June 2013 | Published 20 June 2013

[urn:lsid:zoobank.org:pub:6814946F-05EF-4085-8BCE-1D65CE32C2B6](https://doi.org/urn:lsid:zoobank.org:pub:6814946F-05EF-4085-8BCE-1D65CE32C2B6)

Citation: Richart CH, Hedin M (2013) Three new species in the harvestmen genus *Acuclavella* (Opiliones, Dyspnoi, Ischyropsalidoidea), including description of male *Acuclavella quattuor* Shear, 1986. ZooKeys 311: 19–68. doi: 10.3897/zookeys.311.2920

Abstract

In Shear's (1986) cladistic analysis of the Ischyropsalidoidea, he described the new genus *Acuclavella* including four new species from the Pacific Northwest states of Washington and Idaho. Several of these species descriptions were based on very limited sample sizes. Our recent field work has increased by more than an order of magnitude both the number of specimens and known localities for *Acuclavella*. We use this new material to interpret species limits in *Acuclavella* using morphometric analyses and DNA sequence data from four gene regions. We sequence for the first time the protein-coding homolog of the Wnt2 gene for phylogenetic reconstruction in Opiliones. Our multi-locus phylogeny corroborates a sister relationship between *Acuclavella* and *Ceratolasma*, as hypothesized using morphology by Shear (1986). Within *Acuclavella*, morphometric clusters and reciprocal allelic monophyly allows recognition of three additional species: *Acuclavella leonardi* sp. n., *A. sheari* sp. n., and *A. makah* sp. n. This work also describes the previously unknown male of *Acuclavella quattuor*, from specimens collected at the type locality. Our research identifies a number of novel morphologies for *Acuclavella*, including females with four pairs of spines, individuals with three pairs of spines on scute areas I-III, and a population with two pairs of spines disjunct from *A. quattuor*, which was diagnosed with this spination character. We were unable to assign these populations to existing species, and conservatively do not yet recognize them as new. Intrageneric morphometrics and phylogenetic inference in *Acuclavella* were often concordant. However, we demonstrate that species delimitation signal would not be detected if only a single line of evidence were utilized.

Keywords

Molecular phylogenetics, morphometrics, Pacific Northwest, northern Idaho, Salmon River, Olympic Peninsula, species delimitation, cybertaxonomy

Introduction

The genus *Acuclavella* was described in 1986 by Shear in his revision of the superfamily Ischyropsalidoidea (Opiliones, Dyspnoi). In this work, four new species in the genus were described (Shear 1986). These descriptions were based in large part on very small sample sizes: *Acuclavella cosmetoides* Shear, 1986, the generic type species, was described from a single male and female; *A. shoshone* Shear, 1986 was described from two males and one female; and *A. quattuor* Shear, 1986 was described from a single female. Each of these three species was known only from their respective type localities in northern Idaho, north of the Salmon River. The fourth species, *A. merickeli* Shear, 1986, was represented by thirteen individuals. Twelve of these (four males and eight females) were from the type locality also in Idaho north of the Salmon River. The remaining individual was collected from the Cascade Mountains of western Washington, and due to morphological similarity, was considered by Shear to be conspecific with *A. merickeli* despite a geographic separation of over 500 kilometers.

The biogeographic situation in *Acuclavella*, with several short-range endemic species from a small geographic area, coupled with an apparently widespread (but disjunct) species, clearly invites further investigation. The biogeographic barriers separating the Cascade Mountains from the Rocky Mountains of northern Idaho have promoted speciation in a variety of taxa (e.g., amphibians: Nielson et al. 2001, Steele et al. 2005; arachnids: Derkarabetian et al. 2010; insects: Barr 2011). In addition, recent surveys have uncovered populations of *Acuclavella* from the Olympic Peninsula of northwestern Washington State, and from south of the Salmon River in Idaho. Cascade Mountain and Olympic Peninsula ecoregions of western Washington have been shown to house distinct species of non-vagile organisms (e.g., Good and Wake 1992). In Idaho, a habitat corridor of *Abies grandis*, north-facing and high above the south side of the Salmon River, may house unique populations of organisms that survived in an unrecognized compartment of a structured Pleistocene glacial refugium (Brunsfeld and Sullivan 2006).

The goal of this paper is to use morphometrics and molecular phylogenetics to investigate the validity of the four species hypothesized by Shear (1986), to delimit potentially new species within *Acuclavella*, and to revise the genus as necessary. We hypothesize that molecular phylogenies will reveal consistent reciprocal monophyly conforming to Idaho versus Washington taxa. Furthermore, we expect that *Acuclavella* populations from the two ecoregions in western Washington will show evidence of molecular divergence. In Idaho, expectations are that a morphologically distinct new population discovered south of the Salmon River will show molecular phylogenetic divergence from the four northern species described by Shear (1986).

Methods

Taxon sampling

Most fieldwork was conducted in the summer of 2008, with additional adult specimens collected from May to September in 2006, 2007, and 2009 (Appendix I - Collection Locality Information). *Acuclavella* are crenophilic denizens of small, perennial water features such as headwater streams and seeps in the *Tsuga heterophylla* Zone and the coastal *Picea sitchensis* Zone of the Pacific Northwest (Franklin and Dyrness 1988). Appropriate habitat and microhabitat was targeted for specimen collection throughout northern Idaho and western Washington, and collections include all type localities. When encountered, at least one specimen from each locality was preserved for molecular analyses by placing specimens into 100% EtOH, and subsequently transferring specimens to a -80° C freezer (following Vink et al. 2005).

Molecular data collection

Genomic DNA was extracted from two legs per specimen using the Qiagen DNeasy kit, per manufacturer's protocol. Currently, few genes are available for resolving shallow phylogenetic relationships in Opiliones (reviewed in Hedin et al. 2010). In this study, PCR amplification targeted four gene regions, including mitochondrial protein-coding cytochrome oxidase I (COI), the 28S large subunit ribosomal RNA (28S), nuclear protein-coding Elongation Factor-1 alpha (EF-1 α), and the nuclear protein-coding *wingless* (Wnt2). COI is a standard gene used for species-level delimitation in animals, and has been used extensively in opilionid phylogenetic studies (e.g. Boyer et al. 2005, 2007; Thomas and Hedin 2008; Hedin and Thomas 2010; Derkarabetian et al. 2010). 28S rRNA data has been used successfully to elucidate deeper relationships in opilionid phylogenies (e.g. Boyer et al. 2007, Derkarabetian et al. 2010, Giribet et al. 2010, Schönhöfer and Martens 2010, Sharma and Giribet 2011), but generally evolves too slowly to resolve shallow relationships (e.g. Hedin and Thomas 2010). EF-1 α has been shown to have phylogenetic utility at two levels (Hedin et al. 2010); in the region amplified for *Acuclavella*, more slowly evolving exons bracket a relatively rapidly evolving intron. The Wnt2 gene region has been used to reconstruct spider phylogenies (e.g., Blackledge et al. 2009, Satler et al. 2011), but is used here for the first time in Opiliones. PCR primer and amplification conditions are available as a supplementary file (Appendix II - PCR Primer Information). Amplicons were visualized on agarose gels, purified via polyethylene glycol (PEG) precipitation or on Millipore plates, and Sanger sequenced at the San Diego State University Microchemical Core Facility (http://www.sci.sdsu.edu/dna-core/sdsu_dnacore.html) or at MacroGen USA.

Phylogenetic analyses

Bi-directional Sanger reads were assembled into contiguous sequences using Sequencher v4.5 (Gene Codes Corporation, MI). EF-1 α haplotypes were reconstructed in PHASE (Stephens and Donnelly 2003) using an ingroup-only matrix with the following parameter settings: 100 iterations, thinning interval of 1, burn-in of 100, probability cut-off set to 0.70 (Harrigan et al. 2008). Haplotypes with nucleotides inferred with probability values < 0.70 were left ambiguous. The PHASE input file was converted from a FASTA alignment using SeqPHASE (Flot 2010). COI, EF-1 α exon, and Wnt2 gene regions were unambiguously aligned using amino acid translations in MacClade v4.06 (Maddison and Maddison 2003). EF-1 α intron and 28S rRNA data were aligned using MAFFT (Katoh et al. 2005), utilizing the Q-INS-i alignment algorithm strategy with parameters: BLOSUM62 alignment scoring matrix, 200PAM / K=2 scoring matrix value, an opening gap penalty of 1.53, and an offset value of 0.1. Regions of alignment uncertainty were removed from the 28S matrix with Gblocks (Castresana 2000) with the following parameter settings: minimum length of a block = 3, allowed gap position with half, minimum number of sequences used to identify a conserved position and a flanking position 11 and 17 respectively, and 8 as the maximum number of contiguous non-conserved positions.

Individual gene trees were reconstructed using maximum likelihood and Bayesian inference. Bayesian analyses were implemented using MrBayes v3.1.2 (Ronquist and Huelsenbeck 2003) run on an XSEDE utilizing the CIPRES portal (Miller et al. 2010). Models of DNA sequence evolution used in Bayesian analyses were determined using jModelTest v0.1.1 (Posada 2008, Guindon and Gascuel 2003). Likelihood scores were computed with three substitution schemes (24 models), unequal base frequencies, proportion of invariable sites, rate variation among sites, and used the BIONJ algorithm of Gascuel (1997). The goodness of fit of alternative models was determined using AIC (Akaike 1974) with CI set to 100%. Bayesian gene tree analyses were run for 5×10^6 generations sampling every 1000 trees, using best-fit models of molecular evolution. At this many generations, the average standard deviation of split frequencies between runs was < 0.01. The first 40% of the sampled trees were discarded as burn-in, and stationarity was confirmed visually using Tracer v2.4 (Rambaut and Drummond 2007). Maximum likelihood gene tree analyses were conducted using RAxML v7.2.8 (Stamatakis 2006), also on the CIPRES portal using the GTR + GAMMA model for tree inference and bootstrapping (Stamatakis et al. 2008). The EF-1 α data set was partitioned by intron and exon, and the COI data was partitioned by codon position. The Wnt2 and 28S matrices were not partitioned. Outgroup sequences were used to root all gene trees.

All available DNA sequences, with the exception of apparently nuclearized copies of COI (see Results), were concatenated for phylogeny reconstruction. The concatenated matrix was analyzed using both maximum likelihood (RAxML) and Bayesian approaches (MrBayes v3.1.2) using a seven-partition strategy (EF-1 α intron + exon, Wnt2, 28S, individual COI codon positions). The Bayesian analysis was run for 1 X

Table 1. Phylogenetic taxon sample and GenBank accession numbers.

Species / Voucher No.	COI	28S	EF-1 α	Wnt2
Acfquattuor_OP2230_DeVoto	KF181730	KF181744	KF181759	
Acfquattuor_OP2275_SplitCkTr	KF181731		KF181760	KF181779
Acfquattuor_OP2284_SelwayRvRd	KF181732		KF181761	KF181780
Acosmetoides_OP2281_2ShadowsCk		KF181745	KF181762	
Acosmetoides_OP2296_FS250		KF181746	KF181763	
Acosmetoides_OP2299_TribOrogrande				KF181781
Acosmetoides_OP2319_MeadowCk		KF181747		
Acosmetoides_OP2341_GooseCk		KF181748		
Aleonardi_OP2347_IronCk	GQ870648	KF181749	GQ872169	KF181782
Aleonardi_OP2349_NFGobleCk			KF181764	
Aleonardi_OP2712_KjesbuRd		KF181750	KF181765	
Aleonardi_OP2714_UpperIronCk	KF181728		KF181766	
Americkeli_OP2237_FS443	KF181733	KF181751	KF181767	
Americkeli_OP2245_FS443	KF181734			KF181783
Americkeli_OP2250_RedHorseCk	KF181735		KF181768	KF181784
Amakah_OP1699_RubyBeach	KF181736		KF181769	
Amakah_OP2345_CedarCk	GQ870647	KF181752	GQ872168	
Amakah_OP2715_HokoFalls	KF181737	KF181753	KF181770	KF181785
Amakah_OP2716_BrownesCk	KF181738		KF181771	
Amakah_OP2719_YahooLkRd			KF181772	
Aquattuor_OP2242_GrouseCk	KF181739		KF181773	KF181786
Aquattuor_OP2257_SlateCk	KF181740	KF181754	KF181774	
Aquattuor_OP2270_FS221	KF181729			KF181787
Asheari_OP2708_BurgdorfRd	KF181741	KF181755	KF181775	
Asheari_OP2709_BurgdorfRd	KF181742		KF181776	
Asheari_OP2720_FS592	KF181743	KF181756	KF181777	KF181788
Ashoshone_OP2316_EmeraldCkRd				KF181789
Ashoshone_OP2323_Hobo		KF181757	KF181778	
Ceratolasma	GQ912865	JX573543	AF240864	KF181790
Dendrolasma	KF181727	GQ912771	AF240865	
Hesperonemastoma	JX573642	JX573548	AF240869	KF181791
Ischyropsalis	JX573639	AF240870	JX573603	KF181792
Ortholasma	GQ912870	KF181758	GQ872161	
Sabacon	JX573670	JX573551	AF240877	KF181793
Taracus	JX573680	JX573592	AF240881	KF181794

10⁷ generations; tree sampling and burn-in were as above. Outgroup sequences for five of six ischyropsalidoid genera (*Ceratolasma*, *Taracus*, *Hesperonemastoma*, *Sabacon*, and *Ischyropsalis*) and two genera from Troguloidea (*Dendrolasma* and *Ortholasma*) were used in concatenated phylogenetic analyses. These sequences were generated in-house or downloaded from GenBank (see Table 1).

Morphometric analyses

Specimen measurements were taken using an Olympus SZX12 dissecting microscope with an ocular micrometer. Individuals with missing data (i.e., no Leg II) were excluded from analyses. Standard measurements (Acosta et al. 2007) of 16 characters were determined for 261 individuals (131 males, 133 females); the measurement data, including a figure showing our character measurements scheme, are available as a supplementary file (Appendix III). Total body length was not measured due to the variation in length allowed by free tergites and sternites in individuals that are gravid or have recently eaten; this measurement is known to vary with nutritional condition in Opiliones (Acosta and Machado 2007).

Principle components analyses (PCA) and discriminant function analyses (DFA) were carried out using SYSTAT 12 (Systat Software, Inc.). Though both PCA and DFA are multivariate analyses, DFA is a validation approach that deals specifically with the problem of separating predetermined groups. In taxonomy, DFAs are almost exclusively used to differentiate between morphologically similar species that are difficult to identify from single characteristics (see Klimov et al. 2004, Seifert 2003, Fisher 1936, Lubishew 1962). Multispecies delimitation via DFA, as done here, is becoming more common (see Schlick-Steiner et al. 2006, Schönhofner and Martens 2008, Gebiola et al. 2012).

In taxonomy, PCAs are regularly employed for species delimitation. This utilization has ranged from comparing the vector angles in a scatter plot of predefined species groups (Hamm 2010), to mathematical inference of groups without *a priori* assignment of samples to species (Ezard et al. 2010). Because a PCA has no intrinsic measure of group exclusivity, the parameters of group recovery in morphospace need to be defined. For purposes of this research we defined three PCA categories: a “recovered” group clustered together and did not overlap with samples from other groups; a “nearly recovered” group clustered together with minimal overlap with other groups (one or two individuals at the periphery of their respective group that are not enclosed within another); a “not recovered” group failed to cluster exclusively. For these analyses, three methods were employed to explore this morphospace: (1) hypothesized species were compared pairwise; (2) all combinations of principle components explaining a significant part of the variation in the data were explored; and (3) a nested approach where if clusters of hypothesized species were recovered by initial analyses, further analyses were conducted on those clusters. An in-depth discussion of morphometric methods is available as a supplementary file (Appendix IV).

Due to sexual dimorphism, male and female specimens were analyzed separately for all morphometric analyses. Since the data used to conduct PCAs were measured in the same units (mm), analyses were conducted on both correlation and covariance matrices. PCAs using covariance matrices tend to be dominated by characters showing the most variability (Jolliffe 2002). In order to reduce possible excessive influence on the principle components, variables were coded to have a mean of zero and variance of one; analyses on this data are based on a correlation matrix. Analyses run on covariance

matrices assume a multivariate normal distribution. This is often violated in taxonomic studies where data consists of multiple species of varying morphological distinctiveness; outliers could represent members of an under-sampled group. All unusual measurements were confirmed by repeated measurement, then included in analyses under the assumption they do not excessively affect interpretation of the data.

Species delimitation

We initiated analyses with seven *a priori* hypothesized species, with species-level distinctions based on geographic criteria or on preliminary specimen sorting (i.e., not morphometrics). Geographic criteria were based on the results of work on amphibians endemic to the disjunct mesic forests of the Pacific Northwest (Nielson et al. 2001), or the physiogeographic regions of western Washington (Good and Wake 1992). Areas of expected endemism included western Washington versus northern Idaho, as well as between Cascade Mountain and Olympic Peninsula regions in Washington, areas separated by the low valley of the Puget Trough. In Idaho, a morphologically diminutive population was discovered south of the Salmon River; this group was also tested as a species.

North of the Salmon River in Idaho, *A. merickeli* and *A. quattuor* were distinguished using the diagnostic features outlined by Shear (1986). Due to a gradation of intermediate morphologies spanning the diagnostic characters (Shear 1986) for *A. cosmetoides* and *A. shoshone* (Appendix V: males Figure 1, females Figure 2), coupled with the observation that the described morphologies of these species more or less occur only at the type localities (Appendix V: males, Figure 3; females, Figure 4), an initial attempt to distinguish these two species was abandoned. Since *A. cosmetoides* is the type species for the genus, all morphotypes north of the Middle Fork Clearwater were preliminarily identified as *A. cosmetoides*. A newly discovered population in northern Idaho matched the diagnostic morphology of *A. quattuor* of having pairs of spines on scute areas I and II. However, the new population is distributed between the Selway and Lochsa rivers, whereas *A. quattuor* occurs between the South Fork Clearwater and Salmon Rivers. This population was tested as a new species with the working name *Acuclavella* cf. *quattuor*.

Cybertaxonomy

A cybertaxonomic approach was undertaken for enhanced dissemination of this work (e.g., Miller et al. 2010). This work is available as an open access PDF (doi: 10.3897/zookeys.311.2920). Molecular sequence data has been uploaded to GenBank (<http://www.ncbi.nlm.nih.gov/Entrez>; see Table 1 for accession numbers). Some species of *Acuclavella* are difficult to diagnose using morphological characters alone. An effort has been made to facilitate DNA barcoding by uploading COI sequences to the Barcode of Life Data Systems (BOLD) v. 2.5 (<http://boldsystems.org>). Aligned

matrices and all phylogenetic trees have been deposited in the Dryad Digital Depository (doi: 10.5061/dryad.16737). Anatomical images included in this publication and many additional images have been deposited in MorphBank (<http://www.morphbank.net/?id=822371>). A Keyhole Markup Language file (KML) for interactive viewing of species distributions and collection information in Google Earth (<http://earth.google.com>) is available as a supplementary file (Appendix VI). All nomenclatural acts have been registered with ZooBank (<http://www.zoobank.org>). New species described in this study are provided to Encyclopedia of Life (<http://www.eol.org>) where interconnection of information is linked back to this open access, peer-reviewed work.

Data resources

The data underpinning the analyses reported in this paper are deposited in the Dryad Data Repository at doi: 10.5061/dryad.16737.

Results and discussion

Taxon sampling

Fieldwork resulted in the collection of 272 *Acuclavella* specimens from 61 localities. Populations were sampled from throughout northern Idaho, as well as in the Cascade Mountain and Olympic Peninsula physiogeographic regions of western Washington (Figure 1). The first state records for Montana were secured from the Bitterroot Mountains. Specimens were collected from the four type localities (Shear 1986), including male specimens from the type locality of *A. quattuor*, which Shear originally described from a single female. The vast majority of specimens were collected from beneath moist woody debris near small perennial water features.

Although the overall geographic distribution of *Acuclavella* is now reasonably well-known, additional collecting efforts in the following areas may prove fruitful: south of known localities in Idaho, east into more inland areas of Montana, the Coast Range of Oregon, and the Willapa Hills of Washington. For example, most animal taxa that include populations in the Olympic Peninsula and Cascade Mountains are also found in the Willapa Hills (e.g. *Rhyacotriton*, Good and Wake 1992; *Octoglena anura*, Shelley et al. 2010).

DNA sequence characteristics

Mitochondrial COI DNA sequences for a geographical subset of *Acuclavella* specimens were inconsistent with expectations of protein-coding gene evolution. These COI se-

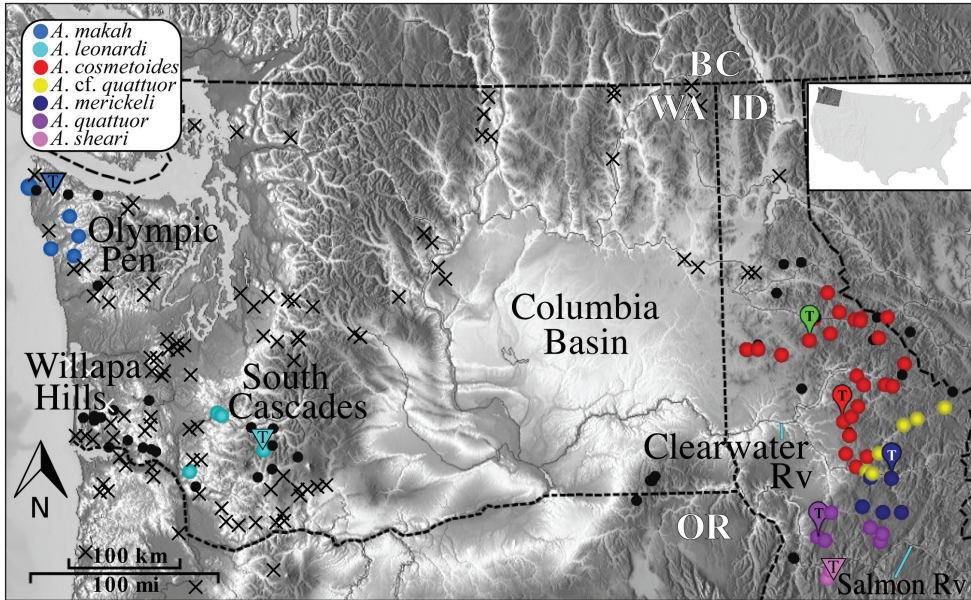


Figure 1. Distribution of *Acuclavella*. Black circles are localities where *Acuclavella* were specifically targeted, but not encountered. Black crosses indicate localities where general surveys of litter invertebrates (e.g., Opiliones, Diplopoda, terrestrial Gastropoda) were conducted and *Acuclavella* were not encountered. Blue lines point to labeled rivers. Type localities (T) from Shear (1986) are indicated with paddle icons; green paddle = *A. shoshone*. Type localities of species described in this work are indicated with inverted triangles. Image saved from Google Earth.

quence reads showed ambiguous nucleotides at individual sites, a large number of mutations resulting in replacement amino acid substitutions, as well as insertions and deletions. Some deletions resulted in frame-shift mutations resulting in stop codons. These patterns of variation suggest that nuclearized copies of the mitochondrial COI gene (NUMTs; White et al. 2008, Baldo et al. 2011) exist in *Acuclavella*.

Sequences showing evidence for nuclearization were discarded, resulting in the removal of all sequences for individuals collected north of the Middle Fork Clearwater River, including all *A. cosmetoides* and *A. shoshone* (*sensu stricto*). In addition to apparent authentic COI sequences, we have multiple nuclear markers providing a similar history for three new species, so our taxonomic conclusions should not be compromised. The final COI matrix included 18 ingroup sequences (L = 1224 base-pairs; Table 1). The first, second, and third codon positions included 34, 9, and 282 parsimony-informative characters in the ingroup. GenBank numbers for all mitochondrial gene sequences can be found in Table 1.

Gblocks removal of ambiguous sites in the 28S matrix resulted in a reduction of alignment length from 1211 to 1173 positions. The aligned matrix included 26 parsimony informative characters for 14 ingroup sequences. For the EF-1 α gene, eight of 22 *Acuclavella* sequences showed signs of heterozygosity. Three of these

individuals were ambiguous at a single base pair, two were ambiguous at two sites, and remaining individuals were ambiguous at 3, 4, or 5 sites respectively. All haplotypes were reconstructed using PHASE. The resulting aligned matrix consisted of 690 bp of exon data, and an 80 bp intron with gaps. The EF-1 α exon contained 30 parsimony informative characters in the ingroup. The intron sequences did not have outgroup representatives; there were 18 parsimony informative characters in the intron. The Wnt2 gene matrix (L = 370) included 7 ingroup sequences, with only 4 parsimony informative characters. GenBank numbers for all nuclear gene sequences can be found in Table 1.

Phylogenetic analyses

Models of evolution inferred from jModelTest are summarized in Table 2. These partition-specific models were used in a Bayesian analysis of the concatenated matrix. Figure 2 shows results from the Bayesian analysis and associated posterior probabilities; the phylogeny resulting from the RAxML analysis is available in a supplementary file (Appendix VII). The concatenated Bayesian and ML phylograms are congruent in their strong support for a monophyletic *Acuclavella*. *Ceratolasma* is recovered as sister to *Acuclavella* with strong support, confirming the hypothesis of Shear (1986). Deeper phylogenetic relationships within Ischyropsalidoidea are beyond the scope of this research, and no emendations to the current taxonomy are proposed, pending collection of additional data.

Within *Acuclavella*, samples from Washington and Idaho are reciprocally monophyletic in both ML and Bayesian analyses, with regional clades strongly-supported and subtended by relatively long branches (Figure 2, Appendix VIII). Within Washington, both analyses recover samples from the Olympic Peninsula and southern Cascade Mountains as monophyletic lineages separated by long branches with high support. Both ML and Bayesian analyses also strongly support *A. sheari* as monophyletic and sister to all other Idaho samples. *Acuclavella* cf. *quattuor* was recovered by both analy-

Table 2. Models of DNA sequence evolution as determined by jModelTest v0.1.1

Gene	Model
COI 1st partition	GTR+I+G
COI 2nd partition	GTR+I+G
COI 3rd partition	GTR+G
28S	GTR+I+G
EF-1 α intron	GTR+G
EF-1 α exon	SYM+G
Wnt2	SYM+I

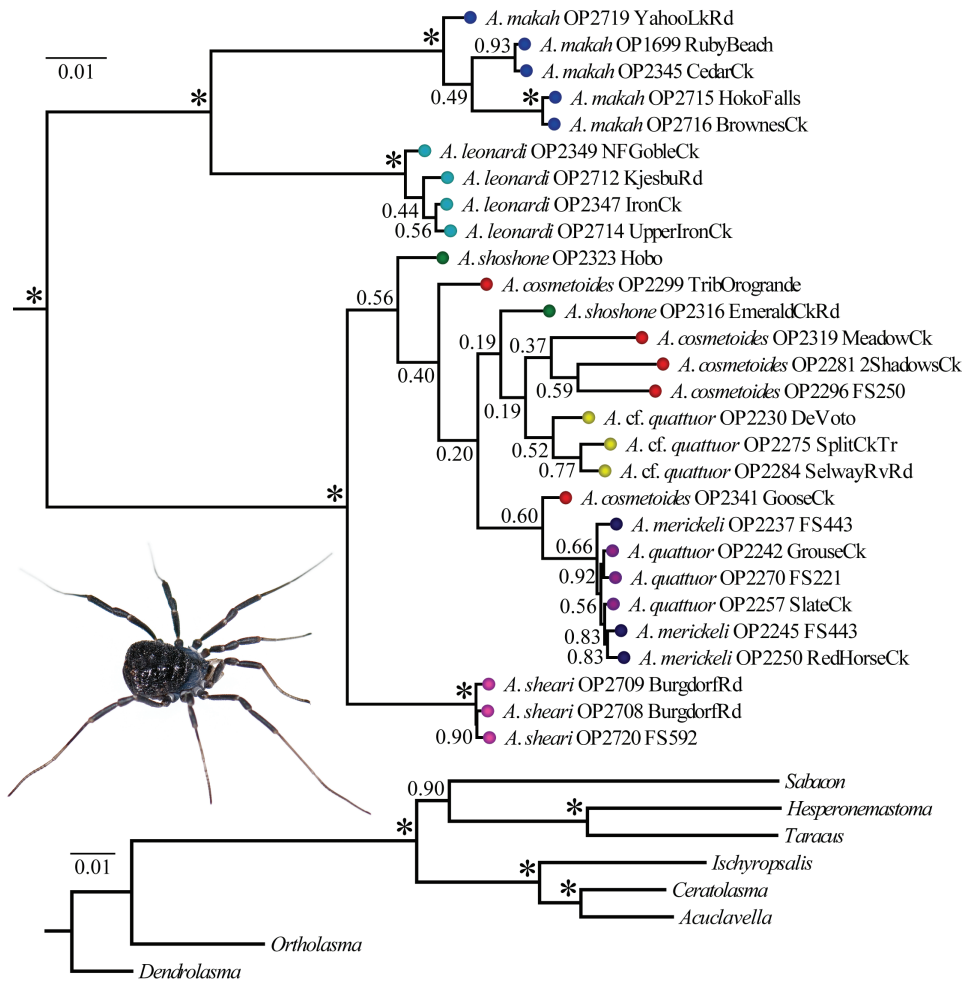


Figure 2. Bayesian phylogram resulting from analysis of concatenated dataset. Numbers at nodes correspond to Bayesian posterior probabilities. The outgroup topology is shown at bottom. The inset picture is a female *A. shoshone* (sensu stricto) collected from the type locality (Shear 1986).

ses, but is not supported by either (PP = 0.52, BS = 13). No other molecular clades are recovered conforming to other described species of northern Idaho *Acuclavella*. A KML file has been created to more easily visualize where sequences used to construct the Bayesian phylogeny occur in geographic space (Appendix VI).

All Bayesian individual gene trees had their longest branch separating Washington and Idaho genetic groups (Figure 3). The COI, EF-1 α , and 28S gene trees (Figure 3A, 3C, and 3D respectively) also show a deep split separating monophyletic *A. leonardi* and *A. makah* samples. *Acuclavella sheari* was also recovered in each of the individual trees, but with varied support and topological placement within a clade of

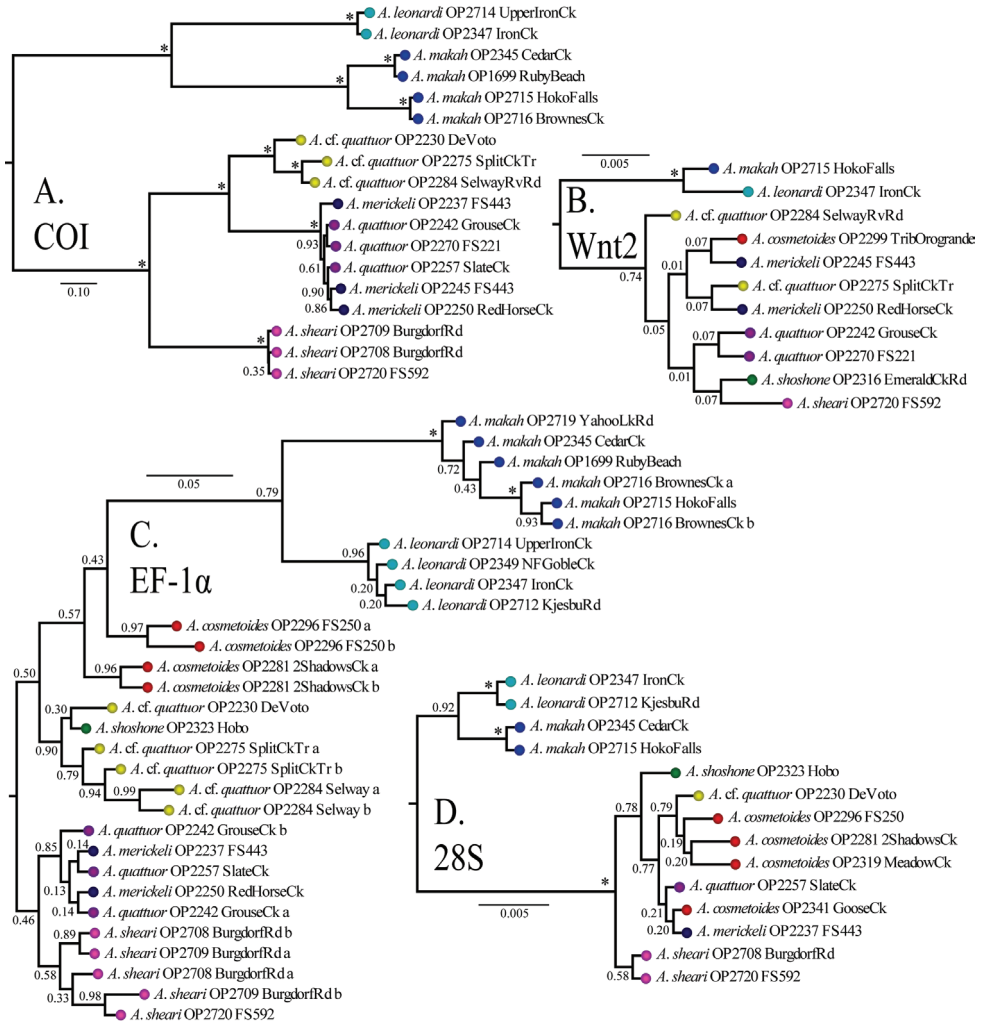


Figure 3. Bayesian gene trees with posterior probabilities: **A** COI **B** Wnt2 **C** EF-1α **D** 28S. All trees outgroup rooted (Appendix VII, Figure 1). RAxML concatenated phylogeny (Appendix VII, Figure 2) and gene trees (Appendix VII, Figures 3-5) are also available.

Idaho samples. The COI gene tree shows significant support for a monophyletic *A. sheari*, which is sister to all remaining Idaho sequences. This species is recovered as monophyletic but without support by EF-1α and 28S gene trees. The Wnt2 gene tree (Figure 3B) supports reciprocal monophyly of samples from Washington and Idaho, but lacks phylogenetic signal within Idaho. The lack of structure for shallow evolutionary events within *Aculeclavella*, coupled with high support for most nodes within Ischyropsalidoidea above the generic level, points to a deeper phylogenetic utility for Wnt2 in Opiliones.

Morphometric analyses

DFA and PCA morphometric analyses of male and female data sets most frequently recovered *Acuclavella sheari*, followed by *A. leonardi* and *A. makah*. Undescribed morphologies from northern Idaho clustered with samples of *A. cosmetoides* and *A. shoshone* (*sensu stricto*) of Shear, 1986. *Acuclavella merickeli* was also frequently recovered in these analyses. The two populations with two pairs of scutal spines on scute areas I and II (*A. quattuor* and *A. cf. quattuor*) were frequently recovered as distinct from other hypothesized species, but were not recovered as morphometrically distinct from each other.

Since the group frequencies of hypothesized species varied greatly (Table 3), jackknifed classification of some data sets reduced the number of individuals used to define a group to just two or three individuals. An assumption is made here that variation seen within hypothesized species is normal in terms of representing the cluster. Male *Acuclavella* are robustly discriminated by DFA analysis, with nearly all species correctly classified with 100% accuracy. An exception is *A. cf. quattuor*, with two specimens classified as an *A. quattuor* (classification error rate of 0.09). Results of DFAs are available. Results of DFAs are available in Appendix IV. The analysis was repeated with a jackknife classification resulting in successful discrimination of 94% of male individuals. Considering the high variation in group frequencies of hypothesized species (Table 3), analyses were rerun allowing prior probabilities of group membership. These analyses resulted in very similar groupings (not shown).

Groupings based on female specimens were not recovered as frequently as male-based groups. Females of *A. leonardi*, *A. merickeli*, and *A. sheari* were correctly discriminated in the classification matrix. There was support for female samples of *A. makah* and *A. cosmetoides*, with correct classification 93% and 97% of the time respectively. About 25% of *A. quattuor* and *A. cf. quattuor* females were misclassified as the other species. Results from the jackknifed classification matrix show similar, if slightly lower percentages of correctly classified individuals. In this matrix, one *A. sheari* is misclassified as *A. cosmetoides*, and three *A. cosmetoides* are misclassified as *A. sheari*. This is likely the result of convergent similarities – females of *A. sheari* and some *A. cosmetoides* (*A. shoshone sensu stricto*) lack scutal spines.

Principle components analyses regularly recovered all hypothesized species of *Acuclavella* with the exception of *A. cf. quattuor* from *A. quattuor*. Figure 4 shows a graph recovering or nearly recovering these species. *Acuclavella sheari* was most frequently recovered in morphospace, with strong evidence for *A. leonardi* and *A. makah* as morphometrically distinct. Also well-supported, *A. merickeli* and *A. cosmetoides* were recovered in PCA analyses. *Acuclavella cf. quattuor* and *A. quattuor* were regularly recovered as distinct from other hypothesized species, but were not distinguishable from each other. Generally, analyses run on covariance matrices had more success of clustering hypothesized species than analyses run on a correlation matrix. Similar to DFAs, males tended to be recovered more frequently than females. Graphs plotting principle components from all PCA analyses conducted (n=143), as well as a description of strategies employed to explore morphospace, are available as a supplementary file (Appendix IV).

Table 3. Sample sizes for PCA and DFA analyses.

Species	Males	Females	Total
<i>A. makah</i>	10	14	24
<i>A. leonardi</i>	6	4	10
<i>A. sheari</i>	4	3	7
<i>A. quattuor</i>	14	17	28
<i>A. merickeli</i>	19	19	38
<i>A. cf. quattuor</i>	22	18	40
<i>A. cosmetoides</i>	56	58	114
Total	131	133	261

The number of individuals from each hypothesized species used in morphometric analyses.

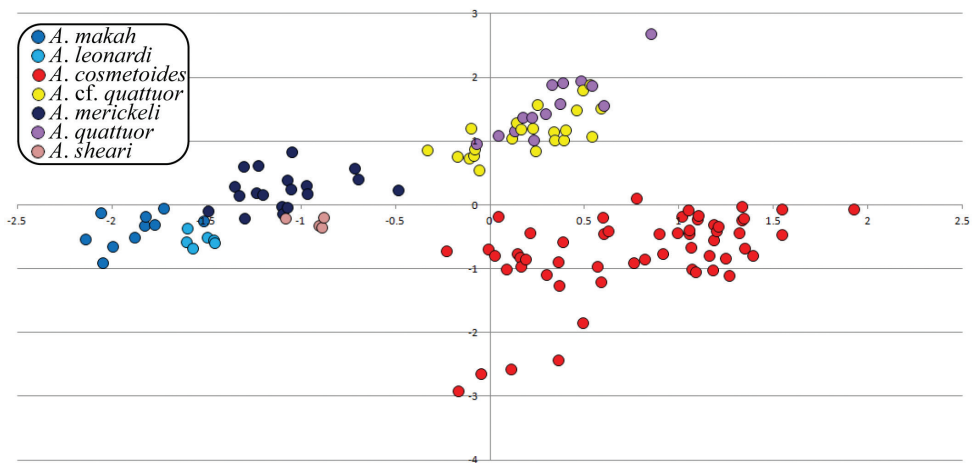


Figure 4. PCA bi-dimensional plot. PCA results of male data using a covariance matrix; plot of principle components 2 and 3.

Difficulties with previously described species

Both phylogenetic and morphometric analyses strongly support three new species: *Acuclavella leonardi*, *Acuclavella makah*, and *Acuclavella sheari*, but do not clearly support already recognized taxa within Idaho samples north of the Salmon River. Molecular and morphological data sets agree that phylogeographic structure associated with the Salmon, Selway, and Lochsa Rivers exists, and morphological data suggests differentiation across the South Fork Clearwater River. Although the number of individuals and known localities was greatly improved by this research, further specimen and genetic sampling is needed to adequately test species limits in northern Idaho. In particular, more rapidly-evolving genetic markers are needed for resolution of what is likely a relatively recent evolutionary radiation in this region. A lack of phylogenetic structure within most of northern Idaho, coupled with the finding that some of the

undescribed morphologies cluster together with *A. shoshone* and *A. cosmetoides*, make it possible that *A. cosmetoides* is a widespread and morphologically highly variable species that encompasses *A. shoshone*. The same hypothesis may apply to the *A. merickeli* plus *A. quattuor* lineage.

Though they lack molecular support, *A. merickeli* and *A. quattuor* are readily discriminated by morphometric analyses (Figure 4). Since male *A. quattuor* were collected from the type locality and agree with the diagnostics for females (Shear 1986), the male for this species is here described and figured for the first time.

Biogeography

The Western Hemlock Zone (WHZ) (Franklin and Dyrness 1988) of northern Idaho is disjunct from its coastal counterpart. This disjunction is explained by the orogeny of the Cascade Mountains 8–5 MYA (Mitchell and Montgomery 2006), and subsequent xerification of the Columbia Basin due to the rain shadow formed from this uplift (Leopold and Denton 1987). Since this isolation, the WHZ in Idaho has been subject to the numerous glacial oscillations including alpine glaciers and continental ice sheets (Gates 1983). During peak glaciation it is believed that organisms of the WHZ were forced into compartmented refugia, either in valleys or south of glaciated regions (Brunsfield and Sullivan 2006, Nielson et al. 2006).

Reconstructed molecular phylogenies show that *Acuclavella* biogeography is largely congruent with that seen in other dispersal-limited taxa of the Pacific Northwest. The deep phylogenetic split between Washington and Idaho (Figures 2, 3) conforms to the ancient vicariance hypothesis (Brunsfield et al. 2001) seen in amphibians including *Ascaphus* (Nielson et al. 2001), *Dicamptodon* (Steele et al. 2005), and *Plethodon* (Carstens et al. 2004). Relatively long branches separate *Acuclavella* species from the Cascade Mountain and Olympic Peninsula ecoregions of western Washington, which is congruent with patterns seen in *Rhyacotriton* salamanders (Good and Wake 1992).

Natural history

Acuclavella are riparian obligate forest-dwellers. The vast majority of specimens were collected within the channel wall of perennial headwater streams or in seep-like features. Most specimens were found underneath large woody debris, though excavation of woody sediment wedges (May and Gresswell 2003) or mats of moss adjacent to these streams also resulted in securing individuals. Coniferous canopy cover is typically consistent across these water features at northern Idaho localities. The dominant vegetation at Idaho sites included a canopy of *Tsuga heterophylla* and *Thuja plicata*, or *Abies grandis* in southern areas. Washington localities typically had a riparian vegetation corridor along small streams dominated by *Alnus rubra* and *Rubus spectabilis* (Appendix I).

The primary defense of *Acuclavella* appears to be one of crypsis. Their dark bodies are cryptic against moist woody debris, onto which specimens adhere with outstretched legs. When detected, specimens are easily collected by grabbing onto an outstretched leg; a hurried scramble is soon followed by limb rigidity. Autospasy was not encountered. A strategy to overcome the crypsis of *Acuclavella* is to expose appropriate areas and wait a few minutes for animals to come out of thanatosis to scurry for cover and appropriate microclimatic conditions. *Acuclavella* likely employ mechanical defense via spination. When turning a cover object the first author placed the pad of his finger directly on an individual; it felt like being pricked by a rose. It is likely that heavy sclerotization of the integument and the ubiquitous hemispherical warts add to the structural integrity of these spines. Though circumstantial, this proposal of heavy sclerotization for mechanical defense is consistent with recent findings in other harvestmen (Souza and Willemart 2011).

Acuclavella appear to have an annual life cycle, with seasonal adults. Penultimate stage subadult and adult animals have been collected in April and May; additionally, a young instar was collected in mid October (CHR3435; same locality as CHR 1409; Appendix I). Adults have been found from May to September. Mating was not encountered, nor was feeding. A single individual (CHR2403) (Appendix I) was found dead in the web of a *Pimoa* spider (CHR2404).

Poor-person's rainforest

Recent surveys of litter dwelling animals in mesic forests of the Pacific Northwest have led to the discovery of many new species. Taxa include terrestrial gastropods (Leonard et al. 2011, Leonard et al. 2003), millipeds (Shear and Leonard 2003, Shelley and Shear 2006, Shear and Leonard 2007, Shelley et al. 2010), pseudoscorpions (Mark Harvey personal communication), and Opiliones (this work, additional undescribed species). An appreciation of these creatures benefits society in numerous ways. Taxonomic knowledge provides a better understanding of forest ecosystems, by identifying potential nutrient recyclers and vertebrate prey items. Taxonomic knowledge also provides future researchers with a baseline from which to compare trends – in the words of Joseph Grinnell (1910) “this value will not, however, be realized until the lapse of many years, possibly a century...”. There is promise of novel chemical discovery in the repugnant secretions of harvestmen and millipeds. To some they embellish the aesthetic beauty of our forests. Though many of these newly described organisms are widespread, and without conservation concern, this is likely not true for all taxa (e.g. Shear and Leonard 2004). It is important that surveys continue in this geographic region and for litter faunas in particular. Litter dwellers are ideal organisms for establishing or adding to a metric of biodiversity, and it is important that true microendemics are identified; this information greatly informs conservation efforts (Harvey et al. 2011).

Systematics

The genus *Acuclavella*, and species *A. merickeli*, *A. cosmetoides*, *A. shoshone*, and female *A. quattuor* are described by Shear (1986); where descriptive information follows those entries, it should be considered an addendum to his descriptions. Depository abbreviations: American Museum of Natural History (AMNH), California Academy of Sciences (CAS), University of Washington Burke Museum (UWBM), personal collection of primary author (CHR).

Genus *Acuclavella* Shear, 1986

<http://species-id.net/wiki/Acuclavella>

Acuclavella Shear, 1986: 13. Type species *Acuclavella cosmetoides* Shear, 1986.

Description. The simple, distally tapering penis (Appendix VIII, Figure I) and short ovipositor (Appendix III, Figure 2) morphologies are conserved across species, with intraspecific variation seemingly as great as interspecific variation. Penis sheath with two sclerotized bands. Metatarsus of leg II with or without false leg articulations (Appendix IX). Distitarsi with three segments on legs I and II; distal end of legs III and IV with two constrictions, each comprised of two segments. All males with raised, glandular (Shear 1986), setose mound dorsally on basal article of chelicerae, setae often capped with secretions (MorphBank image 822802); mounds lacking in females. Chela teeth diaphanous, with two heavily sclerotized teeth distally on fingers (MorphBank image 822817). All leg coxae with prolateral and retrolateral tubercles in the form of distended clusters of warts (MorphBank image 828524). Epistome of stomotheca horn-like, projecting outward (MorphBank image 828520) to strongly decurved (MorphBank image 822815), without apparent interspecific trends. Pseudotrachea of pedipalpal coxaphysis sclerotized ctenoid (MorphBank image 822798).

Key to the species of *Acuclavella*. This dichotomous key should allow users to identify the new species described herein. However, discovery of new morphologies in northern Idaho not described by Shear (1986), and not ascribed to species herein, are currently of uncertain placement at the species level. These morphologies include females with pairs of spines on scute areas I-IV, individuals with three pairs of spines on scute areas I-III, and individuals with pairs of spines on scute areas I-II which are geographically and genetically distinct, but morphologically similar to *Acuclavella quattuor*. For the purposes of this work, the four species named by Shear (1986) are treated *sensu stricto*, and follow his diagnoses.

- | | | |
|----|--|---|
| 1a | Setose mounds dorsally on surface of cheliceral article I; males (Fig. 6) | 2 |
| 1b | Without raised mounds on chelicerae; females..... | 8 |

2a (1a)	Paramedian tubercles acute spines on scute area II only (Fig. 9).....	3
2b	Paramedian tubercles acute spines on multiple scute areas (Fig. 11)	6
3a (2a)	Ocularium height (ventral edge of eye to tip of ocularium) ≤ 0.60 mm; area II spine height ≤ 0.50 mm from surface of tergite.....	
 <i>Acuclavella sheari</i> sp. n.	
3b	Ocularium and area II spine heights ≥ 0.80 mm.....	4
4a (3b)	Distal ends of leg femora, patellae, and tibiae distinctly light, contrasting as light joints; palpi white or nearly so; with or without false leg articulations on leg II metatarsi (Appendix IX); with or without distal, dark, prolateral tubercle on palpal patellae; western Washington.....	5
4b	Distal ends of leg segments not light; palpi light brown or dark; without false leg articulations and palpal tubercles; Idaho	
 <i>A. merickeli</i> Shear, 1986	
5a (4a)	Leg II femur ≤ 3.76 mm; Cascade Mountains	<i>A. leonardi</i> sp. n.
5b	Leg II femur ≥ 3.76 mm; Olympic Peninsula	<i>A. makah</i> sp. n.
6a (2b)	Paramedian tubercles raised into acute spines on scute areas I and II only (Fig. 11); known distribution bracketed by the Salmon River and South Fork Clearwater River.....	<i>A. quattuor</i> Shear, 1986
6b	Paramedian tubercles with acute spines on three or more scute areas; scute area III always with such tubercles (Appendix V, Figure 1)	7
7a (6b)	Paramedian tubercles not raised into spines on scute area I, paired spines on areas II, III, and IV.....	<i>A. cosmetoides</i> Shear, 1986
7b	Paramedian tubercles raised into acute spines on scute areas I – IV (four pairs of abdominal spines.....	<i>A. shoshone</i> (Shear, 1986)
8a (1b)	Paramedian tubercles acute spines on scute area II only (Fig. 9).....	9
8b	Tubercles not as previous; if spines on area II only, greatly reduced.....	11
9a (8a)	As couplet 4a	10
9b	As couplet 4b.....	<i>A. merickeli</i>
10a (9a)	Known only from the Cascade Mountains of Washington State; no perceived discriminating features	<i>A. leonardi</i> sp. n.
10b	Known only from the Olympic Peninsula of Washington State; no perceived discriminating features	<i>A. makah</i> sp. n.
11a (8b)	As couplet 6a	<i>A. quattuor</i> Shear, 1986
11b	Paramedian tubercles not as previous	12
12a (11a)	As couplet 7a	<i>A. cosmetoides</i> Shear, 1986
12b	Paramedian tubercles not enlarged into spines, but paired, low, rounded tubercles on all scutal areas (Fig. 9)	13
13a (11b)	Palpal femur ≤ 0.88 mm; leg II tarsus ≤ 3.92 mm; leg II femur ≤ 2.75 mm; currently known south of Salmon River.....	<i>A. sheari</i> sp. n.
13b	Palpal femur ≥ 0.90 mm; leg II tarsus ≥ 3.95 mm; leg II femur ≥ 2.72 mm, north of Middle Fork Clearwater River.....	
 <i>A. shoshone</i> (Shear, 1986)	

***Acuclavella leonardi* sp. n.**

urn:lsid:zoobank.org:act:3F79C21E-E37A-4FC5-92E3-170CA8D9481C

http://species-id.net/wiki/Acuclavella_leonardi

MorphBank images of specimens considered this species include:

Paratype AMNH, MorphBank Specimen Id: 822643, 1 image

Paratype CASENT9039218, MorphBank Specimen Id: 822508, 4 images

Paratype CASENT9039224, MorphBank Specimen Id: 822516, 2 images

SDSU OP2347, MorphBank Specimen Id: 822509, 1 image

SDSU OP2349, MorphBank Specimen Id: 822511, 3 images

SDSU OP2712, MorphBank Specimen Id: 822515, 1 image

SDSU OP2714, MorphBank Specimen Id: 822517, 3 images

Figure 5 and 6, Appendix VIII: Figure 1, Figure 2

Acuclavella merickeli Shear, 1986 (in part: specimen from Pg 21-22)

Type material. Male **holotype** (AMNH), and male (CAS, CASENT9039218) and female (AMNH) **paratypes** from a tributary of Iron Creek, Forest Service Rd 25 4.6 miles south of FS Rd 300, Gifford Pinchot National Forest, Lewis County, Washington; male **paratype** (UWBM, WA2392/6319) and female **paratype** (CAS, CASENT9039224) from upper Iron Creek, FS Rd 28 0.1 miles E of FS 25, Gifford Pinchot National Forest, Skamania County, Washington; female **paratype** (UWBM, WA2391/6027) from a tributary of Goble Creek, Cowlitz County, Washington. Further information on type localities can be found in Appendix I with the exception of female paratype from Goble Creek. This specimen was collected by the first author 3 August 2005 and deposited in a research collection; this specimen was not used in morphometric analyses, but is used to characterize the species in the description below. This specimen was collected at a location accessed via S Goble Creek Rd; 2.0 miles east of Rose Valley Rd turn right, 1.6 miles turn right, 0.6 miles park; N46.0963°, W122.7607°, elevation 227 meters.

Etymology. The specific epithet is a patronym in honor of the naturalist and careful observer William P. Leonard for his work on litter-dwelling organisms in the poor-person's rainforest of the Pacific Northwest.

Diagnosis. Distinguished from all *Acuclavella* except *A. makah* by the combination of having paramedian tubercles as enlarged spines on area II only, and having light, strongly contrasting ends to sclerotized leg segments, giving the appearance of banding. Also distinguished from these taxa in that false leg articulations on the metatarsi of legs II are present, or single dark prolateral tubercles on the palpal patellae are present, but these features are not consistently found in *A. leonardi*. Scutes posterior to spines containing many raised mounds bearing warty tubercles, more distinct than in *A. makah*. Though the height of scutal spines is similar, the base of the spines in *A. leonardi* appears broader than in *A. makah*. Diagnostic COI sequences have been uploaded to the Barcode of Life Data Systems (BOLD: ACUOP005-13).



Figure 5. *Acuclavella leonardi*. Top: male, CHR3325; bottom: female, CHR 2492.

Description of male. Body arched and convex dorsally (Figure 5); sides parallel or nearly so when broader posteriorly. Nearly all of body heavily sclerotized black or brown, with densely scattered hemispherical warts which irregularly house short setae apically or posteriorly. Total length 4.18 mm ($n=3$, 3.88–4.35 mm), carapace length 1.22 mm ($n=7$, 1.10–1.35 mm), carapace width 2.53 mm ($n=3$, 2.44–2.65 mm), length of fused tergites I–V 2.24 mm ($n=7$, 2.10–2.41 mm), scutum length 2.73 mm ($n=3$, 2.65–2.85 mm), scutum width 2.63 mm ($n=3$, 2.55–2.75 mm).

Eye tubercle at anterior edge of carapace prolonged into an acute spine lacking hemispherical warts on distal half; standing 1.25 mm above the surface of the carapace ($n=3$, 1.13–1.35 mm), 1.05 mm ($n=7$, 0.88–1.19 mm) from the ventral edge of the eye to the tip of the tubercle. Eye color brown, brown-gray, or gray, located basally on tubercle. Metapeltidial paramedian sensory cones raised into a sharp, acute spine standing 0.26 mm ($n=7$, 0.23–0.28 mm) above surface of the carapace, curving slightly towards the midline; lateral to these spines clusters of warts form tubercles.

All scutal tergites with pairs of median tubercles, these prolonged into large spines on area II only; lateral tubercles distinct. Tergite I with paired median tubercles as raised mounds adorned with warts standing 0.09 mm ($n=7$, 0.08–0.13 mm) above the scutal surface; two additional pairs of warty mounds reduce in size laterally. Median tubercles on area II tergite greatly enlarged into erect spines standing 1.04 mm ($n=7$, 0.85–1.16 mm) above the surface of the scutum; lateral to these tubercles are raised mounds adorned with warts. Area III tergite with three pairs of relatively large tubercles in the form of raised mounds adorned with warts; these not decreasing in size laterally; apical setae on each mound; median pair 0.06 mm above scutum ($n=7$, 0.03–0.08 mm). Tergite areas IV and V with three or four (UWBM) pairs of tubercles in the form of raised mounds adorned with warts, these not decreasing in size laterally or posteriorly; setae as previous; area IV median tubercle height 0.06 mm ($n=7$, 0.05–0.08 mm). First free tergite (VI) with relatively large and numerous tubercles in the form of raised warty mounds. Second free tergite (VII) as previous in the holotype and paratype (CAS); paratype (UWBM) without tubercles. Free tergite VIII without distinguishable tubercles, or with a median pair only (CAS paratype).

Abdominal sternite warty sculpturing strongest laterally and on posterior margins; sternites brown. Sclerotized areas of genital operculum relatively setose. Prosomal sternum ($n=1$) length 0.19 mm, width 0.37 mm; brown; without setae. Labium weakly to moderately sclerotized, wider than long or longer than wide (CAS); lengths: 0.14, 0.20, 0.08 mm, widths: 0.28, 0.16, 0.18 mm; light brown to brown; without setae. Palpal endites setose, light brown or brown. Leg II endites bearing 1 or 2 (UWBM) setae, leg IV bearing 1 setae in paratype only, all other endites without setae. Horn-shaped process of epistome decurved, projecting 0.40 mm from sulcus ($n=1$).

Chelicerae brown or light brown (Figure 6); darker dorsally; article II with prolateral and retrolateral striations of darker, more sclerotized cuticle; weakly so in paratypes. Cheliceral measurements ($n=3$): article I length 1.14 mm (1.03–1.34 mm), width 0.41 mm (0.40–0.44 mm), article II length 1.32 mm (1.20–1.38 mm), width 0.38 mm (0.35–0.38 mm), article III length 0.56 mm (0.53–0.60 mm). Dorsal surface of article I with raised, glandular area dense with setae, these capped with a white secretion in the CAS paratype; proximal end of article I with boss-like tubercles on retrolateral and proventrolateral surfaces. Palpal coxae brown or light brown (UWBM paratype), with 2 seta-bearing tubercles. Palpal measurements given in Table 4. Trochanter (Figure 6) brown, light brown, or pale brown, with 3 or 4 (UWBM paratype) seta-bearing tubercles. Palpal femora brown or white; patella white without a dark diffuse band, with dark, prolateral tubercles distally, bearing small trichia and setae distally; tibia white,

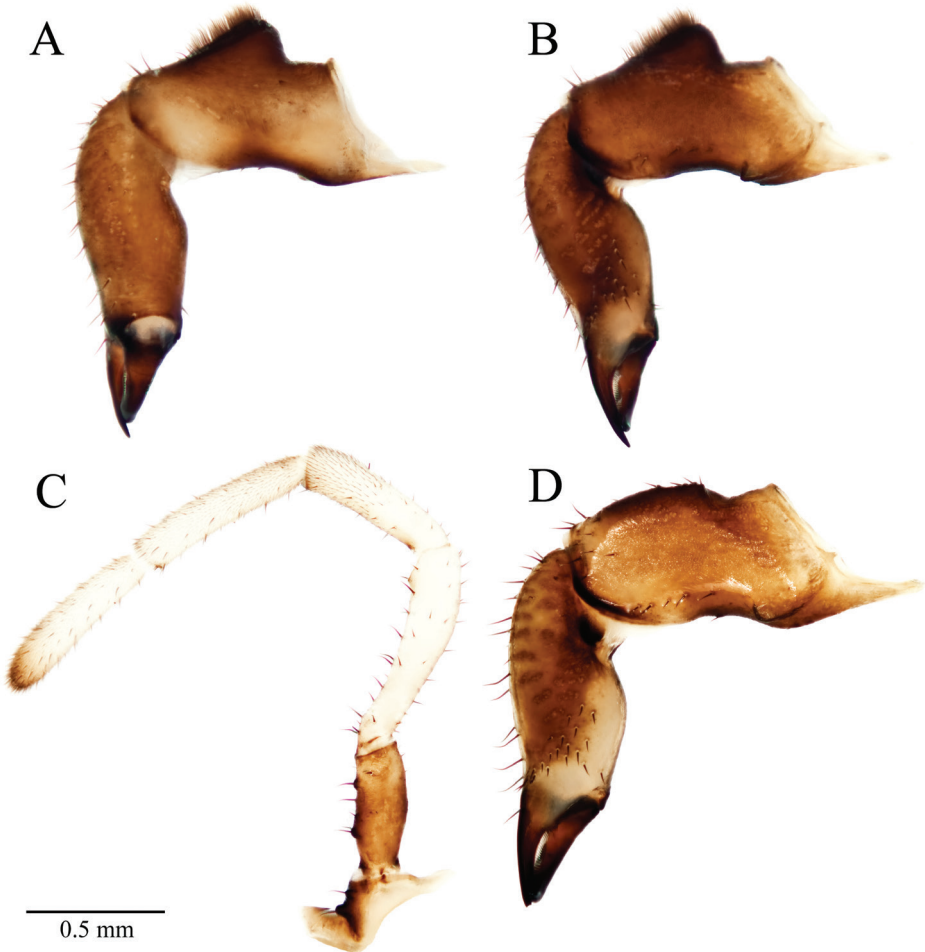


Figure 6. *Acuclavella leonardi* chelicerae and pedipalp. **A** right chelicera, retrolateral view **B** left chelicera, prolateral view **C** right pedipalp, retrolateral view (A–C, male CAsENT9039218) **D** left chelicera, retrolateral view (female CAsENT9039224).

with scattered setae and dense microtrichia; tarsus white, darkening distally, with vestiture of microtrichia and setae. Claw rudiment very small.

Leg measurements given in Table 4. Microsculpture of femora, patellae, and tibiae scattered, distally elevated scales, bilobed scales not observed; scales subtend setae, occasionally housing seta apically. Leg trochanters, femora, patellae, tibiae light brown, dark brown, or black, lighter at joints; metatarsi of leg III with proximal one-third to one-half light brown, brown, or black; leg IV with proximal three-quarters light brown, brown, or black; proximal end of metatarsi of legs I and II pale brown, brown, or black; remaining metatarsal areas pale brown; tarsi pale brown, darkening distally. Scaled microsculpture subequal to darkened areas, remainder with setae and microtrichia. Metatarsi of leg II with false leg articulations ($n=7$).

Table 4. *Acuclavella leonardi* palpus and leg measurements.

Appendage	Segment	Male			Female		
		mean	Range	<i>n</i>	Mean	Range	<i>n</i>
Palpus	trochanter	0.55	0.52–0.58	3	0.61	0.58–0.65	3
	femur	0.94	0.84–1.00	7	0.94	0.90–0.98	5
	patella	0.67	0.66–0.68	3	0.71	0.70–0.73	3
	tibia	0.74	0.73–0.74	3	0.73	0.67–0.80	3
	tarsus	0.69	0.67–0.72	3	0.66	0.64–0.68	3
Leg I	trochanter	0.49	0.48–0.52	3	0.50	0.45–0.56	3
	femur	2.37	2.28–2.44	3	2.25	1.96–2.60	3
	patella	0.88	0.88	3	0.87	0.84–0.92	3
	tibia	1.60	1.48–1.72	3	1.46	1.44–1.50	3
	metatarsus	2.17	1.84–2.36	3	2.19	2.04–2.28	3
	tarsus	2.89	2.72–3.08	3	2.80	2.68–2.88	3
Leg II	trochanter	0.57	0.52–0.60	7	0.55	0.52–0.56	5
	femur	3.53	3.32–3.76	7	3.46	2.88–3.92	5
	patella	1.02	0.96–1.08	7	1.00	0.92–1.08	5
	tibia	2.38	2.16–2.60	7	2.36	2.12–2.60	5
	metatarsus	3.70	3.44–4.05	7	3.69	3.28–3.96	5
	tarsus	4.93	4.74–5.19	7	4.58	4.40–4.70	5
Leg III	trochanter	0.48	0.48	3	0.54	0.52–0.60	3
	femur	2.23	2.12–2.28	3	2.10	1.88–2.25	3
	patella	0.85	0.80–0.88	3	0.84	0.80–0.88	3
	tibia	1.57	1.56–1.60	3	1.50	1.48–1.52	3
	metatarsus	2.49	2.40–2.56	3	2.38	2.24–2.45	3
	tarsus	3.13	3.08–3.20	3	2.95	2.80–3.05	3
Leg IV	trochanter	0.57	0.52–0.64	3	0.58	0.55–0.64	3
	femur	3.23	3.08–3.32	3	3.09	2.76–3.35	3
	patella	1.03	0.96–1.08	3	0.99	0.96–1.00	3
	tibia	2.01	1.96–2.08	3	1.99	1.96–2.05	3
	metatarsus	4.03	3.88–4.12	3	3.85	3.64–4.00	3
	tarsus	4.09	4.08–4.12	3	3.87	3.76–4.99	3

All measurements in millimeters; *n* = sample size.

Penis length 2.48 mm (*n*=1), glans plate 0.37 mm, stylus 0.09 mm, stylus slightly twisted, not decurved.

Description of female. Similar to male for nearly all characters. Total length 4.71 mm (*n*=3, 4.45–5.06 mm); carapace length 1.35 mm (*n*=5, 1.25–1.44 mm); carapace width 3.01 mm (*n*=3, 2.81–3.13 mm); scutum length 3.63 mm (*n*=3, 3.19–4.30 mm); scutum width 3.70 mm (*n*=3, 3.12–3.50 mm); length of fused sternites I–V 2.96 mm (*n*=5, 2.81–3.06 mm).

Eye tubercle height above surface of carapace 1.18 mm (*n*=3, 1.00–1.28 mm); distance from ventral edge of eye to tip of spine 1.06 mm (*n*=5, 0.90–1.13 mm).

Eye color dark brown, pink-gray, or light brown. Metapeltidial spine 0.22 mm ($n=5$, 0.18–0.25 mm).

Paramedian tubercles or tergite I height 0.08 mm ($n=5$, 0.05–0.13 mm) above surface of tergite; median tubercles raised mounds adorned with warts, two additional pairs of warty mounds reducing in size laterally. Tergite II paramedian tubercles greatly enlarged into erect spines standing 0.94 mm ($n=5$, 0.78–1.03 mm) above tergite surface; lateral to these are raised mounds adorned with warts. Paramedian tubercles of tergite III 0.07 mm ($n=5$, 0.03–0.10 mm); IV 0.06 mm ($n=5$, 0.03–0.10 mm). Tergite areas III, IV, and V with three pairs of relatively large tubercles in the form of raised mounds adorned with warts; these not decreasing in size laterally, decreasing slightly posteriorly across tergites. First free tergite (VI) adorned with raised warty mounds; tergites VII and VIII without discernable tubercles in paratypes (AMNH, CAS), tergite VII of paratype (UWBM) with tubercles as in VI.

Sternites brown. Transverse furrow and membranous lateral sutures of genital operculum less distinct than in other species. Prosomal sternum ($n=1$) length 0.18 mm, width 0.39 mm; pale-brown; without setae. Labium wider than long or subequal; lengths: 0.10, 0.19, 0.13 mm; widths: 0.16, 0.18, 0.19 mm; moderately or weakly sclerotized; brown; without setae. Palpal endites light brown. Leg II endites adorned with 2 setae.

Horn-shaped process of epistome decurved, projecting 0.42 mm from sulcus. Chelicerae brown or light brown; article I length 1.18 mm ($n=3$, 1.16–1.21 mm), width 0.43 mm (0.42–0.44 mm); article II length 1.39 mm (1.34–1.42 mm), width 0.41 mm (0.38–0.43 mm); article III length 0.59 mm (0.58–0.62 mm). Article I without raised glandular mound (Figure 6). Article II with 6 setae on prolateral dark area at cleavage of corpus and fixed finger of chela; 15 setae on ventral surface of article II; these patches discrete. Palpus dimensions in Table 4; coxae light brown or brown with two seta-bearing tubercles; trochanters brown or pale brown with 4 or 5 seta-bearing tubercles; only paratype (CAS) with tubercle on patella, only paratype (UWBM) with partial diffuse band on patella.

Leg measurements given in Table 4. Leg trochanters, femora, patellae, and tibiae brown to dark brown, lighter at joints; metatarsi of legs III with proximal one-half brown, of leg IV with proximal three-quarters light brown to brown, proximal ends of legs I and II light brown to brown, remaining metatarsal areas pale brown; tarsi pale brown, darkening distally.

Ovipositor length 0.80 mm, width 0.44 mm; corona of setae at furcal base surrounding lobes, apical setae on lobes; furca without dorsoventral differentiation.

Distribution and habitat. *Acuclavella leonardi* is found in the southern Cascade Mountains of Washington State in the Cowlitz River (includes Iron Creek) and Coweeman River (includes Goble Creek) watersheds in Lewis, Cowlitz, and Skamania Counties (Appendix I). Found in coniferous forests with small perennial water-features such as side-slope seeps, springs, and headwater streams; underneath stream-side woody debris.

***Acuclavella makah* sp. n.**

urn:lsid:zoobank.org:act:8C34D2F8-D211-42B7-B58A-A90285771589

http://species-id.net/wiki/Acuclavella_makah

MorphBank images of specimens considered this species include:

Holotype AMNH, MorphBank Specimen Id: 822644, 2 images

Paratype CASENT9039219, MorphBank Specimen Id: 822645, 3 images

CHR1536, MorphBank Specimen Id: 828519, 2 images

SDSU OP1699, MorphBank Specimen Id: 828516, 1 image

CHR2457.0, MorphBank Specimen Id: 822505, 4 images

CHR2457.1, MorphBank Specimen Id: 822506, 3 images

CHR2457.2, MorphBank Specimen Id: 822507, 3 images

CHR3387.0., MorphBank Specimen Id: 822647, 1 image

CHR3387.1, MorphBank Specimen Id: 822646, 1 image

Figure 7 and 8, Appendix VIII: Figure 1, Figure 2

Type material. Male **holotype** and female **paratype** (AMNH), and male and female **paratypes** (CAS, CASENT9039219) from Brownes Creek, Clallam County, Washington; male and female **paratypes** (UWBM, WA2393/7641) from Yahoo Lake Road 1.4 miles east of Hoh-Clearwater Road, Jefferson County, Washington (Appendix I). An additional specimen is housed at UWBM (WA0973/8147): Ahlstroms Prairie, Olympic National Park, Clallam County, Washington, (NAD 1927) N48.157°, W124.704°, elevation 40 meters; Rod Crawford (collected under permit), 16–17 July 1984.

Etymology. The specific epithet refers to the Makah Nation, which historically occupied much of the known distribution of the species. The name Makah was given to these people by their neighbors; it means “generous with food”. These people have shared with many people access to their beautiful land, next to the rocks and gulls. For more information on the Makah Nation see: <http://www.makah.com>.

Diagnosis. Distinguished from all *Acuclavella* except *A. leonardi* by the combination of having paramedian tubercles on area II only, having light, and strongly contrasting ends to sclerotized leg segments, giving the appearance of banding at joints. Though not always present, false leg articulations on the metatarsi of legs II, and single dark prolateral tubercles on the palpal patellae also diagnose it from these species. Scutal tubercles lateral to paramedian tubercles tend to be on more distinctive raised mounds in *A. leonardi*. Area II spines more narrow at base than in *A. leonardi*. Best diagnosed from *A. leonardi* using molecular data. Diagnostic COI sequences have been uploaded to the Barcode of Life Data Systems (BOLD: ACUOP007-13).

Description of male. Body arched and convex dorsally (Figure 7), sides parallel with equal scutum and carapace widths; nearly all of body heavily sclerotized, black, with densely scattered hemispherical warts which irregularly house short setae apically or posteriorly. Total length 4.11 mm ($n=3$; 4.05–4.20 mm), carapace length in midline 1.26 mm ($n=10$; 1.05–1.44 mm), greatest carapace width 2.57 mm ($n=3$; 2.50–2.72 mm); length of fused scutes I–V in midline 2.33 ($n=10$; 2.20–2.48 mm),



Figure 7. *Acuclavella makah*. Top: male, CHR2457.1; bottom: female, CASENT9039219.

scutum length in midline 2.74 ($n=3$; 2.60–2.90 mm) greatest scutum width 2.57 mm ($n=3$; 2.50–2.72 mm).

Eye tubercle at anterior edge of carapace, prolonged anteriorly into a sharp conical spine 1.27 mm ($n=10$; 1.05–1.45 mm) from ventral edge of eye to tip of spine; 1.58 mm above the carapace ($n=3$; 1.53–1.64 mm). Eyes light-brown to brown, lo-

cated basally on tubercle. Surface of carapace evenly curved, posterior margin arcuate. Metapeltidial paramedian sensory cones short, acute spines standing 0.19 mm ($n=10$; 0.13–0.25 mm) above the surface of the metapeltidium, shiny, lacking warts.

All scutal areas with pairs of paramedian tubercles. Area I paramedian tubercles cluster of cuticular warts standing 0.04 mm above the surrounding scute ($n=10$; 0.025–0.075 mm), two additional pairs of warty tubercles reduce in size laterally. Paramedian tubercles of area II rise to form large acute spines standing 1.37 mm above the scutum surface ($n=10$; 1.13–1.75 mm); lateral to spines a pair of tubercles typically small cluster of warts, though one individual (CHR1536) with lateral pair of short spines. Scute areas III, IV, and V with three pairs of wart-clustered tubercles each; paramedian pair largest, diminishing in size laterally along scute, and posteriorly across scute areas. Paramedian tubercles of area III stand 0.03 mm ($n=10$, 0.03–0.05 mm) above surface of the tergite; area IV tubercle height 0.04 mm ($n=10$, 0.03–0.08 mm). Holotype and paratype (CAS) with first free tergite (area VI) with barely distinguishable tubercle as median pair of enlarged warts, tergites VII and VIII without tubercles; paratype (UWBM) tergite VI with median and lateral pair of enlarged warty tubercles, tergite VII with barely distinguishable tubercles, these lacking on tergite VIII. Tergite IX divided, triangular, bracketing tergite X, which forms the anal operculum.

Abdominal sternites with infrequent setae; warty sculpturing strongest on posterior and lateral margins; brown to dark brown. Prosomal sternum length 0.14 mm ($n=2$, 0.13–0.16 mm), width 0.28 mm (0.26–0.30 mm); brown; without setae. Labium weakly sclerotized, wider than long, length 0.08 mm ($n=3$; 0.05–0.11 mm), width 0.15 mm (0.13–0.17 mm); light brown to yellow-brown, without setae. Palpal endites brown to yellow-brown, large, free, bearing many setae; leg II endite bearing 3 setae in holotype and paratype (CAS), 2 in paratype (UWBM); legs I, III, IV endites without setae. Epistome with horn-like anteriad projection, decurved or slightly decurved, projecting 0.34 mm ($n=2$, 0.33–0.36 mm) from sulcus. Chelicerae basal article (I) dark brown dorsally, middle article (II) with prolateral and retrolateral striations of darker, more sclerotized striations. Article I length 1.18 mm ($n=3$, 1.06–1.30 mm), width (I) at widest point 0.41 mm (0.38–0.43 mm), article II length 1.40 mm (1.34–1.46 mm), width (II) at widest point 0.40 mm (0.38–0.43 mm), article III length 0.53 mm (0.50–0.57 mm). Article I with raised, setose glandular area on dorsal surface (Figure 8). Article I with boss-like tubercles proximally on proventrolateral and retrolateral surfaces, one individual with proventrolateral tubercles bilobed; proventrolateral tubercle may function to macerate food or manipulate food items in conjunction with the epistome process. Cleavage of corpus and fixed finger housing 5 or 6 setae on prolateral side of article II; 12 or 15 setae on ventral surface of article II; these patches discrete. Palpal coxae yellow-brown to brown, with two seta-bearing tubercles ventrally (Figure 8). Palpal measurements given in Table 5. Trochanter brown to dark-brown with 3 (holotype) or 4 seta-bearing tubercles; femur white; patella white without dark band medially, with prolateral, distal darkened tubercles; bearing microtrichia and small setae distally; tibia white with scattered setae and dense microtrichia; tarsus white, usually darkening distally. Claw rudiment very small.

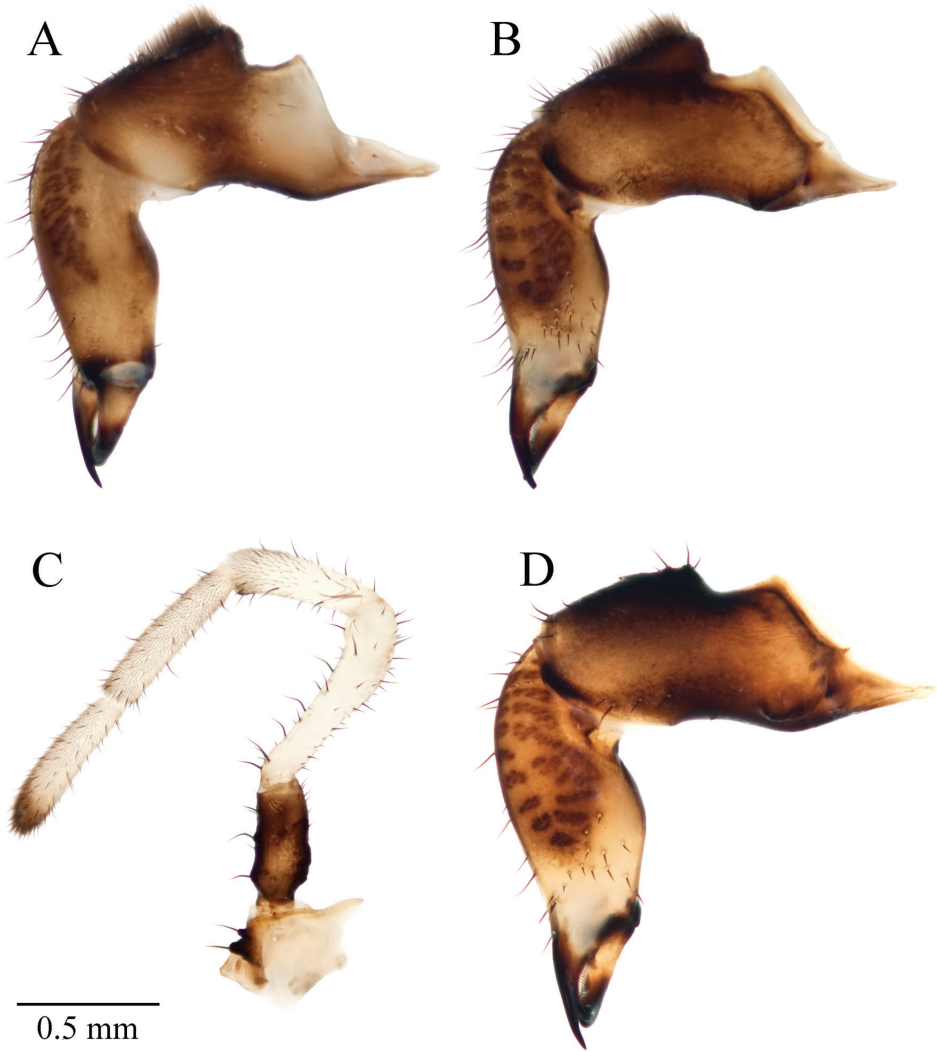


Figure 8. *Acuclavella makah* chelicerae and pedipalp. **A** right chelicera, retrolateral view **B** left chelicera, prolateral view **C** right pedipalp, retrolateral view (A–C, male CHR2457-0) **D** left chelicera, retrolateral view (female CHR2457-2).

Leg measurements given in Table 5. Trochanters, femora, patellae, tibiae black, lighter at joints, with scattered, distally elevated scales which subtend short setae, scales occasionally house setae apically or posteriorly. Metatarsi of leg III with proximal half black or darkened; leg IV proximal three-quarters black or darkened; legs I and II with proximal end darkened; remaining metatarsal areas pale-brown to yellow-brown. Scaled-microsculpture subequal to darkened areas. Metatarsi and tarsi without tubercles. Six of ten males with false leg articulations on metatarsi of leg II, including holotype. Leg claws single, black, not toothed, evenly curved.

Table 5. *Acuclavella* makah palpus and leg measurements.

Appendage	Segment	Male			Female		
		Mean	Range	<i>n</i>	Mean	Range	<i>n</i>
Palpus	trochanter	0.58	0.54-0.62	3	0.57	0.54-0.60	3
	femur	0.95	0.90-0.98	10	0.98	0.88-1.05	14
	patella	0.67	0.62-0.72	3	0.70	0.67-0.74	3
	tibia	0.79	0.73-0.84	3	0.72	0.68-0.76	3
	tarsus	0.73	0.71-0.76	3	0.68	0.67-0.68	3
Leg I	trochanter	0.57	0.56-0.60	3	0.53	0.50-0.56	3
	femur	2.67	2.60-2.72	3	2.46	2.36-2.56	3
	patella	0.97	0.92-1.00	3	0.90	0.88-0.92	3
	tibia	1.75	1.72-1.76	3	1.57	1.55-1.60	3
	metatarsus	2.36	2.24-2.44	3	2.13	2.08-2.20	3
	tarsus	2.93	2.76-3.04	3	2.72	2.56-2.80	3
Leg II	trochanter	0.60	0.56-0.64	10	0.61	0.56-0.64	14
	femur	4.01	3.76-4.35	10	3.85	3.60-4.20	14
	patella	1.12	1.00-1.20	10	1.14	1.04-1.28	14
	tibia	2.65	2.40-2.88	10	2.64	2.40-2.92	14
	metatarsus	3.91	3.60-4.25	10	3.80	3.52-4.75	14
	tarsus	5.11	4.55-5.50	10	4.62	4.25-4.94	14
Leg III	trochanter	0.55	0.52-0.56	3	0.57	0.55-0.60	3
	femur	2.47	2.32-2.56	3	2.21	2.12-2.32	3
	patella	0.93	0.92-0.96	3	0.96	0.88-1.05	3
	tibia	1.64	1.60-1.68	3	1.60	1.52-1.65	3
	metatarsus	2.64	2.56-2.76	3	2.48	2.40-2.64	3
	tarsus	3.16	3.00-3.32	3	2.95	2.85-3.12	3
Leg IV	trochanter	0.67	0.60-0.72	3	0.65	0.60-0.72	3
	femur	3.64	3.52-3.80	3	3.34	3.16-3.60	3
	patella	1.05	0.96-1.12	3	1.11	1.04-1.20	3
	tibia	2.32	2.32	3	2.30	2.16-2.44	3
	metatarsus	4.16	4.05-4.30	3	3.99	3.76-4.25	3
	tarsus	4.17	3.85-4.35	3	3.86	3.68-3.95	3

All measurements in millimeters; *n* = sample size

Penis 2.39 mm in length (*n*=2, 2.38–2.40 mm); glans plate length 0.34 mm (*n*=2, both 0.34 mm); stylus length 0.16 mm (0.14–0.17 mm). Shaft evenly tapered, broadening slightly at glans; glans with scattered small setae; stylus spirally twisted, very slightly decurved.

Description of female. Similar to male for nearly all characters. Total length 5.14 mm (*n*=3; 5.00–5.31 mm); carapace length 1.37 mm (*n*=14; 1.25–1.81 mm); carapace width 3.16 mm (*n*=3; 3.00–3.31 mm); scutum length 3.82 mm (*n*=3; 3.63–3.94

mm); scutum width 3.33 mm ($n=3$; 3.19–3.50 mm); length of fused sternites I–V 2.96 mm ($n=14$; 2.75–3.19 mm).

Eye tubercle height above surface of carapace 1.37 mm ($n=3$; 1.31–1.46 mm); distance from ventral surface of eye to tip of spine 1.21 mm ($n=14$; 1.08–1.33 mm). Eye color brown. Metapeltidial spine 0.16 mm ($n=14$; 0.10–0.23 mm).

Paramedian tubercles of tergite I height 0.06 mm ($n=14$; 0.03–0.10 mm) above surface of tergite. Tergite II paramedian tubercles greatly enlarged spines standing 1.10 mm ($n=14$; 0.93–1.28 mm) above tergite surface; tubercles lateral to spines warty mounds. Paramedian tubercle height of tergite III 0.07 mm ($n=14$; 0.05–0.10 mm); IV 0.06 mm ($n=14$; 0.03–0.08 mm). Tergites III, IV, V with three pair of tubercles with median pair largest, reducing in size laterally along tergite and posteriorly across tergites; the right median tubercle of tergite III of paratype (CAS) enlarged into mound with diminished frequency of warts standing 0.22 mm above carapace. First free tergite (IV) with median and lateral pair of enlarged warty tubercles, or barely distinguishable median pair of enlarged warts; tergite VII without tubercles, or tubercles barely distinguishable, or more noticeable; tergite VIII without distinguishable tubercles.

Sternites brown. Prosomal sternum length 0.17 mm ($n=2$, 0.17–0.18 mm), width 0.32 mm (0.30–0.33 mm); dark brown. Labium wider (0.17 mm, 0.16 mm, 0.14 mm) than long (0.09 mm, 0.09 mm, 0.14 mm), or nearly equal; yellow-brown to light brown. Palpal endites brown, yellow-brown, or light brown. Paratypes (AMNH, CAS) with 3 setae on leg II endite; paratype (UWBM) with 2 setae. Horn-shaped process of epistome decurved or projecting downward; length 0.48 mm ($n=2$, 0.47–0.50 mm).

Chelicerae article I length 1.35 mm ($n=3$; 1.20–1.47 mm), width 0.45 mm ($n=3$, 0.44–0.46 mm); article II length 1.49 mm ($n=3$; 1.48–1.50 mm), width 0.42 mm (0.42–0.43); article III length 0.59 mm ($n=3$; 0.56–0.60 mm). Article I without raised glandular mound (Figure 8); prolateral side of cheliceral article II with 2 or 4 setae at cleavage of corpus and fixed finger; 12 setae on ventral surface of article II; these patches discrete. Palpus dimensions in Table 5; coxae brown with two seta-bearing tubercles; trochanter brown to dark brown with 4 seta-bearing tubercles; only paratype with tubercle on patella.

Leg measurements given in Table 5. Leg trochanters, femora, patellae, and tibiae black, lighter at joints; metatarsi of leg III with proximal half black or darkened, leg IV with proximal three-quarters black or darkened, proximal end of legs I and II black or darkened, remaining metatarsal areas yellow-brown or pale brown; tarsi yellow-brown or pale brown, darkening distally.

Ovipositor length 0.74 mm ($n=2$, 0.72–0.77 mm), width 0.46 mm (0.44–0.47 mm); furca without dorsoventral differentiation; corona of setae at furcal base surround lobes; apical setae on lobes.

Distribution and habitat. Known from the northwest areas of the Olympic Peninsula in Clallam and Jefferson Counties, Washington State (Appendix I). Found in coniferous or riparian forests along small, perennial water-features such as headwater streams, springs, and seeps; underneath woody debris and moss.

***Acuclavella sheari* sp. n.**

urn:lsid:zoobank.org:act:FC134B12-DA1C-4184-B9AB-6AC36E990BB5

http://species-id.net/wiki/Acuclavella_sheari

MorphBank images of specimens considered this species include:

Holotype AMNH, MorphBank Specimen Id: 822513, 4 images

Paratype AMNH, MorphBank Specimen Id: 822512, 6 images

SDSU OP2708, MorphBank Specimen Id: 822514, 3 images

SDSU OP2720, MorphBank Specimen Id: 822518, 3 images

Figure 9 and 10, Appendix VIII: Figure 1, Figure 2

Type material. Male **holotype** (AMNH) and female **paratypes** (AMNH; CAS, CASENT9039225; UWBM) from Burgdorf Road 12.4 miles northwest of Warren Wagon Rd, tributary of Fall Creek, Payette National Forest, Idaho County, Idaho; two male **paratypes** (CAS, CASENT9039217; UWBM, ID0016/5360) from Burgdorf Road 16.6 miles northwest of Warren Wagon Road; otherwise as previous (Appendix I).

Etymology. The specific epithet honors Dr. William A. Shear, eminent millipede and opilionid taxonomist. His influence has been important to the authors' aspirations to be systematic biologists, and we thank him sincerely, and with pleasure.

Diagnosis. Generally reduced dimensions overall. Males with one pair of scutal spines on area II only; scutal spines ≤ 0.50 mm, distance from ventral edge of eye to tip of ocularium ≤ 0.60 mm distinguishes it from other males with single pair of spines. Females lack scutal spines. Diagnosed from spine-less *A. shoshone* (Shear, 1986 *sensu stricto*) females by having palpal femora ≤ 0.88 mm, leg II tarsi ≤ 3.92 mm, leg II femora ≤ 2.75 mm.

Description of male. Body arched and convex dorsally (Figure 9); sides broader posteriorly. Nearly all of body heavily sclerotized black or dark brown, with densely scattered hemispherical warts which irregularly house short setae apically or posteriorly; bilobed warts sporadic in holotype. Total length 3.90 mm ($n=3$, 3.68–4.06 mm), carapace length 1.24 mm ($n=4$, 1.15–1.32 mm), carapace width 2.34 mm ($n=3$, 2.31–2.36 mm); length of fused tergites I–V 2.26 mm ($n=4$, 2.16–2.31 mm), scutum length 2.61 mm ($n=3$, 2.32–2.81 mm), scutum width 2.60 mm ($n=3$, 2.55–2.63 mm).

Eye tubercle at anterior edge of carapace, hemispherical warts cover entirety of ocular spine, prolonged into an acute spine standing 0.67 mm above the surface of the carapace ($n=3$, 0.64–0.70 mm), 0.53 mm ($n=4$, 0.48–0.60 mm) from the ventral edge of the eye to the tip of the tubercle, eye spine less acute than in other species of *Acuclavella*. Eyes dark brown to brown, located basally on tubercle. Surface of carapace evenly curved, posterior margin arcuate. Metapeltidial paramedian sensory cones raised into sharp, acute spines standing 0.21 mm ($n=4$, 0.17–0.24 mm) above surface of the carapace, curving slightly towards the midline, lacking warty microsculpture, shiny; lateral to spines, clusters of warts form tubercles, missing in paratype (UWBM).

All scutal tergites with pairs of paramedian tubercles; relatively short, pointed spines on area II only. Fused tergite I with paramedian tubercles as raised mounds, relatively tall but not spike-like, standing 0.125 mm above the surrounding scute ($n=4$,



Figure 9. *Acuclavella sheari*. Top: male, CHR3254; bottom: female, CHR3404.

all 0.125), these adorned with warts; lateral to these, two additional raised mounds become smaller away from the midline, these lacking in holotype. Paramedian tubercles of area II form acute spine 0.39 mm above the scutum surface ($n=4$, 0.34–0.43 mm), curved slightly posteriad; lateral to these are raised tubercles adorned with warts. Fused tergites III, IV, and V with paramedian tubercles in the form of raised mound adorned with warts; lateral to these, area III with 2 or 3 additional pairs of tubercles, area IV with 1 or 2 additional tubercles, area V with 0 or 2 additional tubercles; these tend to diminish in size away from the midline and posteriorly across tergites. Area III tubercle height above tergite 0.081 mm ($n=4$, 0.075–0.10 mm); area IV tubercle height 0.075

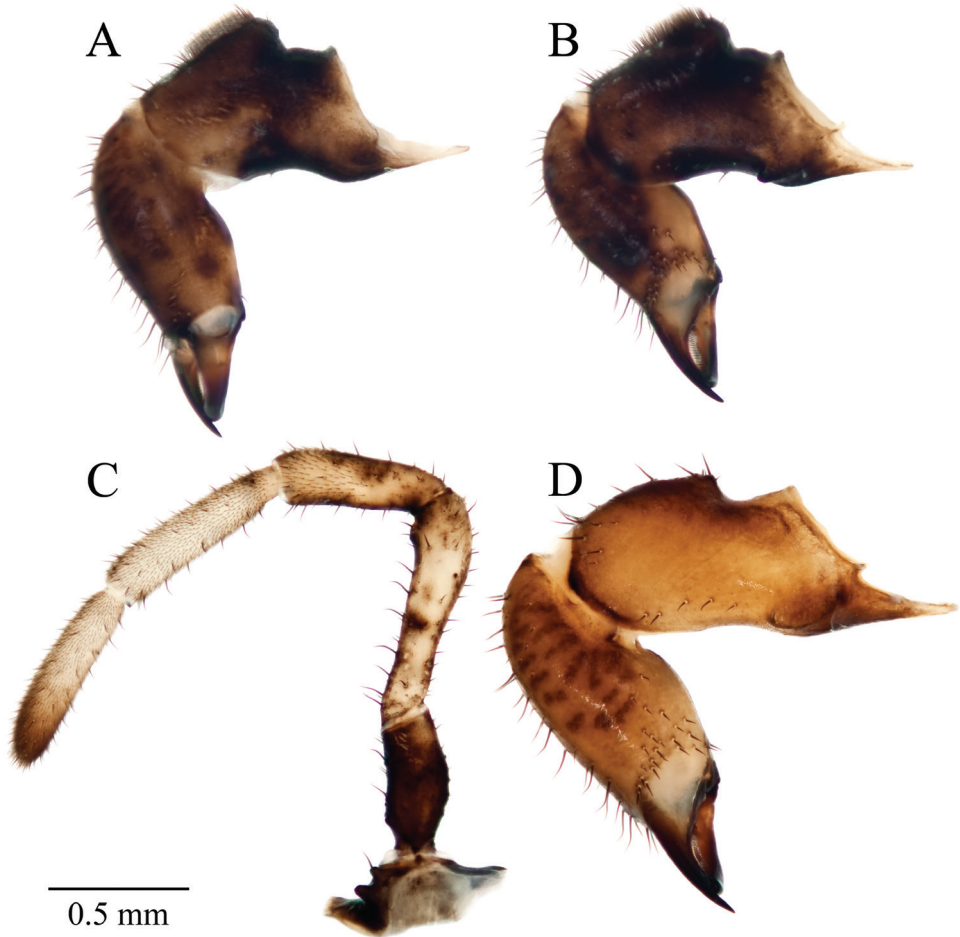


Figure 10. *Acuclavella sheari* chelicerae and pedipalp. **A** right chelicera, retrolateral view **B** left chelicera, prolateral view **C** right pedipalp, retrolateral view (A–C, male CASENT9039217) **D** left chelicera, retrolateral view (female AMNH).

mm (all 0.075 mm). Free tergites without discernable tubercles, or tubercles occur in single pairs as small warty mounds.

Abdominal sternites with infrequent setae; warty sculpturing strongest laterally and on posterior margin; sternites brown, dark brown, or black. Sclerotized areas of genital operculum relatively setose. Prosomal sternum ($n=1$) wider than long: length 0.19 mm, width 0.29 mm; dark brown; without setae. Labium weakly to well-sclerotized, wider than long ($n=3$); length 0.09 mm (0.08–0.10 mm), width 0.16 mm (0.13–0.19 mm), light brown to dark brown, darkness commensurate to sclerotization, without setae. All coxae with endites. Palpal endites setose, light brown, dark brown, or black. Leg II endite bearing 2 setae, other leg endites unadorned. Horn-shaped process of epistome decurved, projecting 0.34 mm from sulcus.

Chelicerae light brown, dark brown, or black; lighter individuals with article I darker dorsally, article II with darker, more sclerotized striations on prolateral and retrolateral surfaces (Figure 10). Article I length 1.11 mm ($n=3$, 1.00–1.18 mm), width 0.39 mm (0.36–0.42 mm), article II length 1.21 mm (1.12–1.27 mm), width 0.40 mm (all 0.40 mm), article III length 0.54 mm (0.52–0.56 mm). Dorsal surface of article I with raised, glandular area dense with setae. Palpal coxae pale-brown, light brown, or black, with 2 or 1 (UWBM) seta-bearing tubercles. Palpal measurements given in Table 6. Trochanter (Figure 10) pale-brown or dark brown, with 2 (CAS), 3 (holotype), or 4 (UWBM) seta-bearing tubercles; these reduced on later individual; femur pale-brown, brown, or dark brown; patella pale-brown or brown, with (CAS) or without a diffuse dark band on middle third, without darkened tubercle distally, bearing small setae and microtrichia distally; tibiae pale brown or white (holotype), with scattered setae and dense microtrichia; tarsus pale-brown or white (holotype), darkening to brown distally. Claw rudiment very small to absent.

Leg measurements given in Table 6. Microsculpture of femora, patellae, and tibiae scattered, distally elevated scales, bilobed scales not observed; scales subtend setae, occasionally housing seta apically. Trochanters, femora, patellae, tibiae black, or in holotype, trochanters light brown, femora, patellae, and tibiae brown; metatarsi of leg III proximal one-third to one-quarter brown or black; leg IV with proximal half black or brown; proximal end of metatarsi on legs I and II black or brown, remaining metatarsal areas pale brown or brown; all tarsi pale brown or brown, darkening distally. Scaled microsculpture subequal to darkened areas, remainder with setae and microtrichia. False leg articulations not observed. Leg claws single, black, not toothed, evenly curved.

Penis length 2.94 mm ($n=1$); glans plate 0.31 mm, scattered small setae; stylus 0.13 mm, slightly spirally twisted and decurved.

Description of female. Similar to male, differing in dimensions, secondary sexual characteristics, and dorsal armature. Total length 4.25 mm ($n=3$; 4.05–4.65 mm); carapace length 1.325 mm ($n=4$, 1.05–1.45 mm), width 2.62 mm ($n=3$, 2.35–3.05 mm); scutum length 3.03 ($n=3$, 2.90–3.25 mm), width 3.23 mm ($n=3$, 2.85–3.55 mm); fused tergites I–V length in midline 2.81 mm ($n=4$, 2.20–3.10 mm).

Eye tubercle height above surface of carapace 0.67 mm ($n=3$, 0.56–0.73 mm), distance from ventral surface of eye to tip of spine 0.54 mm ($n=4$, 0.46–0.60 mm). Eye color brown or gray. Metapeltidial spine 0.11 mm ($n=4$, 0.08–0.15 mm), curving slightly posteriad and towards the midline, lateral to these spines clusters of warts form tubercles.

Dorsal armature lacking. Dorsal adornment generally with median tubercles as raised mounds adorned with warts, decreasing in size away from midline and posteriorly across tergites. Two lateral pairs of warts on areas III and IV, area V with median pair of tubercles barely discernable, two pairs of lateral tubercles, or shallowly raised mounds adorned with warts. Free tergites with tubercles in the form of clusters of warts, or without discernable tubercles. Paramedian tubercles heights above the surface of the tergite ($n=4$): of area I 0.07 mm (0.05–0.08 mm); area II 0.10 mm (0.08–0.15 mm); area III 0.06 mm (0.05–0.08 mm); area IV 0.05 mm (0.03–0.08 mm).

Table 6. *Acuclavella sheari* palpus and leg measurements.

Appendage	Segment	Male			Female		
		Mean	Range	<i>n</i>	Mean	Range	<i>n</i>
Palpus	trochanter	0.51	0.49–0.54	3	0.50	0.43–0.54	3
	Femur	0.85	0.78–0.88	4	0.85	0.83–0.88	4
	Patella	0.63	0.62–0.65	3	0.66	0.60–0.70	3
	Tibia	0.74	0.72–0.75	3	0.73	0.65–0.79	3
	Tarsus	0.70	0.67–0.71	3	0.65	0.59–0.68	3
Leg I	trochanter	0.53	0.48–0.56	3	0.45	0.40–0.50	3
	Femur	1.80	1.76–1.84	3	1.73	1.48–1.94	3
	Patella	0.87	0.84–0.92	3	0.81	0.68–0.91	3
	Tibia	1.33	1.28–1.44	3	1.30	1.16–1.40	3
	metatarsus	1.97	1.96–2.00	3	1.88	1.68–2.09	3
	Tarsus	2.29	2.24–2.32	3	2.22	1.88–2.41	3
Leg II	trochanter	0.57	0.56–0.60	4	0.52	0.44–0.63	4
	Femur	2.53	2.48–2.64	4	2.52	2.16–2.75	3
	Patella	0.98	0.88–1.12	4	0.93	0.76–1.03	3
	Tibia	1.96	1.88–2.08	4	1.90	1.64–2.09	3
	metatarsus	2.99	2.88–3.12	4	3.00	2.56–3.24	3
	Tarsus	3.67	3.64–3.72	4	3.65	3.20–3.92	3
Leg III	trochanter	0.53	0.52–0.56	3	0.52	0.44–0.59	3
	Femur	1.72	1.64–1.80	3	1.71	1.56–1.84	3
	Patella	0.85	0.84–0.88	3	0.79	0.48–0.97	3
	Tibia	1.33	1.20–1.48	3	1.43	1.32–1.56	3
	metatarsus	2.21	2.16–2.28	3	2.18	1.84–2.38	3
	Tarsus	2.56	2.52–2.64	3	2.53	2.20–2.84	3
Leg IV	trochanter	0.59	0.52–0.68	3	0.59	0.52–0.69	3
	Femur	2.41	2.32–2.56	3	2.38	2.12–2.63	3
	Patella	0.96	0.88–1.04	3	0.89	0.80–0.94	3
	Tibia	1.84	1.76–1.88	3	1.77	1.56–2.03	3
	metatarsus	3.31	3.28–3.32	3	3.24	2.80–3.48	3
	Tarsus	3.23	3.08–3.32	3	3.29	2.80–3.68	3

All measurements in millimeters; *n*=sample size

Sternites brown, black, or light brown. Prosomal sternum (*n*=1) length: 0.26 mm, width 0.31 mm; light brown, without setae. Labium wider (0.13, 0.17, 0.16 mm) than long (0.08, 0.09, 0.08 mm); light brown or pale brown. Palpal endites brown or pale brown. Endites of leg II adorned with two or one (CAS) setae.

Chelicerae article I length 0.98 mm (*n*=3, 0.92–1.06 mm), width 0.41 mm (0.37–0.44 mm); article II length 1.31 mm (1.14–1.40 mm), width 0.43 mm (0.40–0.46 mm); article III length 0.57 mm (0.52–0.62 mm). Article I without raised glandular mound (Figure 10). Article I darker dorsally, chelicerae light brown or black with areas of dark brown. Palpus dimensions in Table 6; coxae light brown, dark brown, or

white, with two seta-bearing tubercles; trochanters light brown, dark brown, or white, with three (AMNH) or four (CAS, UWBM) seta-bearing tubercles; femora and patella pale brown, brown, or white, patella without both distal tubercles and diffuse bands; tibiae white or pale-brown; tarsus white or pale-brown, fading to brown distally. Claw rudiment very small (AMNH, UWBM), or absent.

Leg measurements given in Table 6. Leg trochanters brown, black, or pale-brown; femora, patellae, and tibiae dark brown, black, or pale brown; metatarsi of legs III with proximal third brown, black, or pale-brown; leg IV metatarsi with proximal half brown, black, or pale-brown; proximal end of legs I and II metatarsi brown, black, or pale-brown; remaining metatarsal areas pale-brown, brown, or white; tarsi pale brown or white, darkening distally.

Ovipositor length 0.82 mm, width 0.46 mm ($n=1$); otherwise as *A. makah*.

Distribution and habitat. Known from north-facing, horizontal band of *Abies grandis* forest above south shore of Salmon River. This habitat is mainly roadless, though intersected by FS 592 and Burgdorf Road in Payette National Forest, Idaho County, Idaho. Adults collected in June and September. *Abies grandis*, *Pseudotsuga menziesii*, and *Picea engelmannii* associated canopy cover; collected under woody debris adjacent to headwater streams.

Acuclavella quattuor Shear, 1986

http://species-id.net/wiki/Acuclavella_quattuor

MorphBank images of specimens considered this species include:

- AMNH, MorphBank Specimen Id: 822462, 3 images
- CHR2146.2, MorphBank Specimen Id: 822463, 3 images
- CHR2146.5, MorphBank Specimen Id: 822464, 3 images
- CHR2147.7, MorphBank Specimen Id: 822604, 1 image
- SDSU OP2256, MorphBank Specimen Id: 822496, 3 images
- SDSU OP2266, MorphBank Specimen Id: 822605, 1 image
- CHR2180.0, MorphBank Specimen Id: 822606, 2 images
- CHR2180.3, MorphBank Specimen Id: 822607, 1 image

Figure 11 and 12, Appendix VIII: Figure 1, Figure 2

Material examined. Males (AMNH; CAS, CASENT9039220; UWBM, ID0013/5661) from the type locality: Slate Creek Road 10.2 miles east of US 95, Nez Perce National Forest, Idaho County, Idaho.

Diagnosis. Diagnosed from all *Acuclavella* species except for *A. cf. quattuor* by having two pairs of erect spines on tergites I and II; similar morphologically to *A. cf. quattuor*, and is best diagnosed using molecular data: basepair 23 of the EF-1 α intron cytosine in *A. cf. quattuor*, adenine in *A. quattuor*; basepair 44 guanine in *A. quattuor*, thymine or adenine in *A. cf. quattuor* (thymine if sequences aligned due to a 3 basepair deletion in 40% of *A. cf. quattuor* at basepair 39–41).

Description of male. Body arched and convex dorsally (Figure 11); sides parallel or slightly broader posteriorly. Nearly all of body heavily sclerotized, black, with densely scattered hemispherical warts which irregularly house short setae apically or



Figure 11. *Acuclavella quattuor*. Male AMNH.

posteriorly. Total length 4.22 mm ($n=3$, 4.06–4.45 mm); carapace length 1.35 mm ($n=14$, 1.25–1.50 mm), width 2.60 mm ($n=3$, 2.56–2.64 mm); scutum length 2.84 mm ($n=3$, 2.56–3.25 mm), width 2.80 mm ($n=3$, 2.69–2.88 mm); length fused tergites I–V 2.69 mm ($n=14$, 2.50–3.10 mm).

Eye tubercle at anterior edge of carapace, erect, spine-like, standing 0.98 mm above the surface of the carapace ($n=3$, 0.94–1.05 mm), distance from ventral edge of eye to tip of spine 0.84 mm ($n=14$, 0.65–0.93 mm). Eye color brown, on basal part of ocularium. Surface of carapace evenly curved, posterior margin arcuate. Metapeltidial paramedian sensory cones acute or blunt spines standing 0.28 mm ($n=14$, 0.18–0.38 mm) above surface of carapace.

Scutum of opisthosoma rounded anteriorly, squared of posteriorly. All fused tergites with paramedian pair of tubercles, these in the form of erect spines on areas I and II. Spines of tergites I and II curve posteriad, stand 0.82 mm ($n=14$, 0.65–1.05 mm) and 0.91 mm ($n=14$, 0.70–1.12 mm) above the surface of the tergite respectively; tubercles lateral to spines as small raised wart mounds; one of 14 males with lateral spines on area II. Tergites III, IV, and V with four pairs of tubercles as raised areas adorned with warts; paramedian tubercles of area III height 0.06 mm ($n=14$, 0.03–0.20 mm); area IV paramedian tubercle height 0.05 mm ($n=14$, 0.03–0.08 mm); two individuals with one of paramedian tubercles of area III spine-like, left side on one, right side on other. First free tergite (VI) with small median tubercles in the form of enlarged shiny wart; remaining free tergites without discernable tubercles.

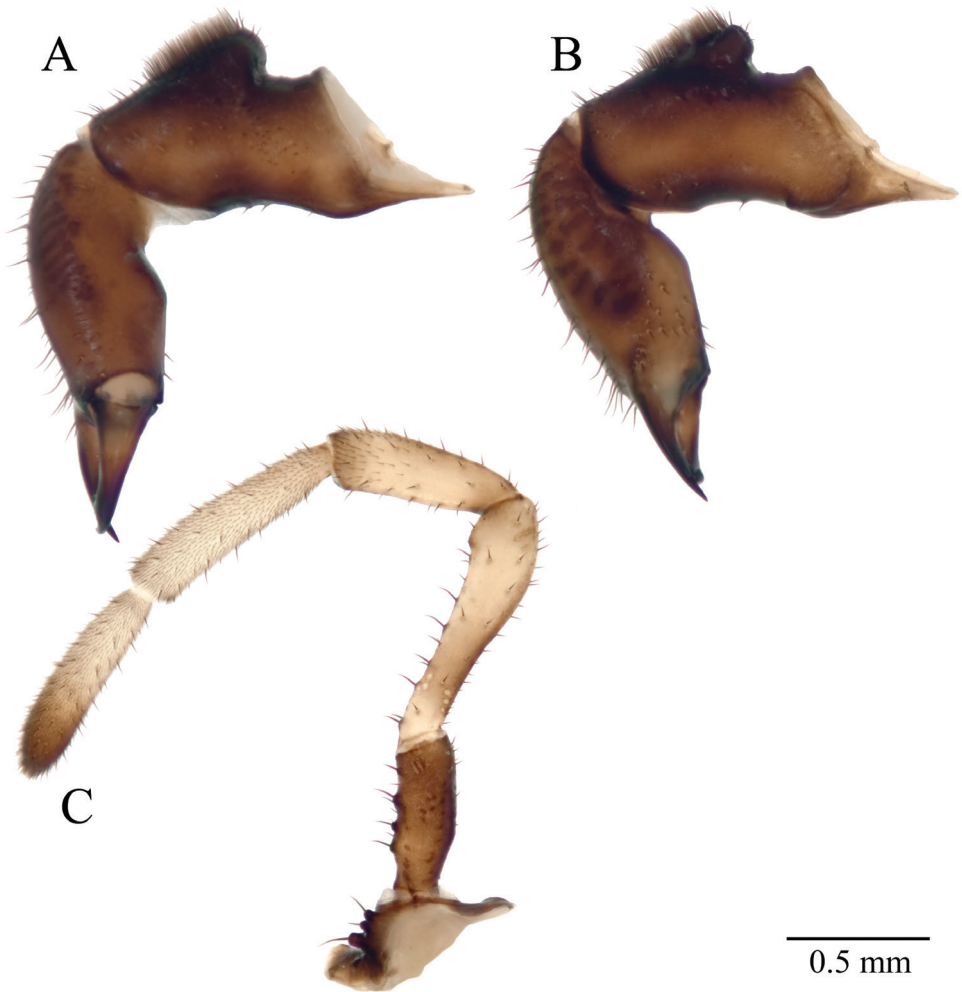


Figure 12. *Acuclavella quattuor* chelicerae and pedipalp. Male CHR2146.5, **A** right chelicera, retrolateral view **B** left chelicera, prolateral view **C** right pedipalp, retrolateral view.

Abdominal sternite warty sculpturing strongest laterally and on posterior margin; sternites brown or dark brown. Genital operculum tongue-shaped, clearly delineated by transverse furrow, lateral ends of furrow membranous suture, distal margin rebordered, glossy. Prosomal sternum wider than long; length 0.18 mm ($n=3$, 0.17–0.21 mm), width 0.29 mm ($n=3$, 0.26–0.31 mm); pale brown or brown; without setae. Labium moderately sclerotized, wider than long or dimensions subequal; length 0.11 mm ($n=3$, 0.09–0.13 mm), width 0.14 mm (0.13–0.16 mm); brown or dark brown; without setae. Palpal endites setose; brown. Leg II endites bearing 2 (AMNH, CAS) or 3 setae (UWBM), other leg endites glabrous. Horn-shaped process of epistome strongly decurved; projecting 0.43 mm ($n=4$, 0.41–0.44 mm) from sulcus.

Table 7. *Acuclavella quattuor* male palpus and leg measurements.

		Palpus	Leg I	Leg II	Leg III	Leg IV
Trochanter	Mean	0.53	0.61	0.61	0.51	0.64
	Range	0.52–0.54	0.60–0.62	0.56–0.68	0.48–0.56	0.56–0.75
	<i>n</i>	3	3	14	3	3
Femur	Mean	1.01	2.31	3.14	2.00	2.88
	Range	0.95–1.10	2.20–2.44	2.72–3.44	1.96–2.04	2.76–3.00
	<i>n</i>	14	3	14	3	3
Patella	Mean	0.67	0.97	1.08	0.95	1.10
	Range	0.64–0.73	0.92–1.03	1.00–1.20	0.92–1.00	0.96–1.20
	<i>n</i>	3	3	14	3	3
Tibia	Mean	0.83	1.54	2.21	1.54	2.11
	Range	0.80–0.88	1.48–1.58	2.00–2.41	1.48–1.64	2.00–2.20
	<i>n</i>	3	3	14	3	3
Metatarsus	Mean	-	2.36	3.55	2.61	3.98
	Range	-	2.16–2.63	2.96–4.08	2.48–2.80	3.68–4.25
	<i>n</i>	-	3	14	3	3
Tarsus	Mean	0.77	2.74	4.71	3.10	4.04
	Range	0.73–0.85	2.48–3.03	3.90–5.25	2.76–3.28	3.56–4.31
	<i>n</i>	3	3	14	3	3

All measurements in millimeters, *n*=sample size.

Chelicerae light brown or brown; darker dorsally; article II with darker, more sclerotized striations on prolateral and retrolateral surfaces (Figure 12). Article I length 1.21 mm (*n*=3, 1.18–1.25 mm), width 0.44 mm (*n*=3, 0.42–0.45 mm); article II length 1.38 mm (*n*=3, 1.35–1.40 mm), width 0.41 mm (*n*=3, 0.38–0.45 mm); article III length 0.59 mm (*n*=3, 0.58–0.60 mm). Article I dorsal surface with raised, glandular area densely setose (Figure 12). Prolateral side of article II with dark sclerotization at cleavage of corpus and fixed finger housing 6, 6, 5, 5 setae; ventral surface of article with 19, 20, 16, 23 setae; these patches discrete in three of four individuals examined. Palpal measurements given in Table 7. Palpal coxae light brown or brown (Figure 12), with 2 seta-bearing tubercles. Palpal trochanters light brown, brown, or dark brown; with 4 seta-bearing tubercles. Femora and patella white or yellow-brown; patella without distal prolateral tubercle, without diffuse dark band, bearing small setae and microtrichia distally. Tibia white or pale yellow, with scattered setae and dense microtrichia. Tarsi white or pale yellow, darkening distally; with setae and dense microtrichia. Claw rudiment very small.

Leg measurements given in Table 7. Microsculpture of femora, patellae, and tibiae scattered, distally elevated scales, bilobed scales not observed; scales infrequently house seta apically or distally. Trochanters, femora, patellae, and tibiae dark brown to black; metatarsi of leg III with proximal one-quarter brown or black, leg IV with proximal one-half brown or black, proximal end of metatarsi of I and II brown or black, remaining metatarsal areas brown; tarsi brown, darkening distally. Leg claws single, black, not toothed, evenly curved.

Penis length 2.49 mm ($n=4$, 2.35–2.55 mm); shaft evenly tapered, broadening slightly at glans. Glans bearing scattered small setae; glans length 0.32 mm ($n=3$, 0.29–0.36 mm). Stylus length 0.18 mm ($n=3$, 0.16–0.19 mm); stylus spirally twisted, decurved.

For female description see Shear (1986).

Distribution and habitat. *Acuclavella quattuor* populations are bracketed by the Salmon River to the south and the South Fork Clearwater River to the north, whereas *A. cf. quattuor* is found north of the Selway River and south of the Lolo Trail Ridge. *Acuclavella quattuor* habitats are dominated by *Abies grandis* and *Picea englemannii*; microhabitat in litter, moss, and moist woody debris adjacent to headwater streams.

Acuclavella merickeli Shear, 1986

http://species-id.net/wiki/Acuclavella_merickeli

MorphBank images of specimens considered this species include:

CHR2100.1, MorphBank Specimen Id: 822457, 2 images; type locality

CHR2100.2, MorphBank Specimen Id: 822597, 1 image

CHR2121.2, MorphBank Specimen Id: 822601, 3 images

CHR2121.4, MorphBank Specimen Id: 822602, 2 images

CHR2121.5, MorphBank Specimen Id: 822459, 3 images

CHR2140.0, MorphBank Specimen Id: 822460, 3 images

CHR2140.2, MorphBank Specimen Id: 822603, 1 image

Figures: Appendix IX

Diagnosis. For species descriptions see Shear 1986. Discriminated from other *Acuclavella* having a single pair of paramedian spines on tergite II by having relatively homogenously colored leg segments, not giving the appearance of banding at leg joints (from *A. makah* and *A. leonardi*) or by having scutal spines > 0.50 mm and ocularium height from ventral edge of eye to tip of spine > 0.60 mm (from *A. sheari*).

Distribution and habitat. *Acuclavella merickeli* populations are bracketed to the south by the South Fork Clearwater River and to the north by the Selway River; all localities in Nez Perce National Forest, Idaho County, Idaho. Coniferous habitat dominated by *Picea englemannii* or *Thuja plicata* and *Pseudotsuga menziesii*; microhabitats include moist woody debris and moss adjacent to small perennial water features such as side-slope seeps and headwater streams.

Acknowledgements

The American Arachnological Society (Vincent Roth Fund for Systematic Research) and the American Museum of Natural History (Theodore Roosevelt Memorial Grant) provided funding to CHR. Several institutional scholarships (Harry E. Hamber Memorial Scholarship, Frank Alverson Memorial Scholarship, and Jordan Dale Covin Memorial Scholarship) to CHR also helped contribute to this work. Specimens were

collected with help from Daniel Richart, William Leonard, Shahan Derkarabetian, Jeffery Underwood, Adrienne Richart, Paul Marek, and Dean Levitt. Hours of morphological measurements were expedited with help from Kristen Emata, David Carlson, Nicasia O'Neal, Adrienne Richart, and Kyle Wilson. Nicasia O'Neal, Roy Larimer, and Joseph Warfel helped with images. Maureen McCormack, Shahan Derkarabetian, and Robin Keith Hedin helped with laboratory training. Stas Vidyakin helped with accessioning MorphBank images. This research was enriched from discussions of concepts and methodologies with Ricardo Carretero, Rod Crawford, Shahan Derkarabetian, Andrew Gottscho, William Leonard, Dean Levitt, Maureen McCormack, Tod Reeder, Jordan Satler, Axel Schönhofer, Peter Scott, William Shear, Jack Sullivan, and others. The *Wnt2 Ischyropsalis* sequence came from the collection of Jochen Martens via Axel Schönhofer, which was collected by S. Huber. The manuscript was improved via suggestions and edits by Angela DiDomenico, Kristen Emata, Ryan Fitch, Erika Garcia, Marc Hayes, Jordan Satler, Axel Schönhofer, Prashant Sharma, William Shear, and an anonymous reviewer.

References

- Acosta LE, Machado G (2007) Diet and foraging. In: Pinto-da-Rocha R, Machado G, Giribet G (Eds) *Harvestmen: the Biology of Opiliones*. Harvard University Press, Cambridge, 309–338.
- Acosta LE, González AP, Tourinho AL (2007) Methods for taxonomic study. In: Pinto-da-Rocha R, Machado G, Giribet G (Eds) *Harvestmen: the Biology of Opiliones*. Harvard University Press, Cambridge, 494–505.
- Akaike H (1974) A new look at statistical model identification. *IEEE Transactions on Automatic Control* 19 (6): 716–723. doi: 10.1109/TAC.1974.1100705
- Baldo L, de Queiroz A, Hedin M, Hayashi C, Gatesy J (2011) Nuclear-mitochondrial sequences as witness of past interbreeding and population diversity in the jumping bristletail *Mesomachilis*. *Molecular Biology and Evolution* 28 (1): 195–210. doi: 10.1093/molbev/msq193
- Barr CE (2011) *Bryelmis* Barr (Coleoptera: Elmidae: Elminae), a new genus of riffle beetle with three new species from the Pacific Northwest, E.S.A. *The Coleopterists Bulletin* 65 (3): 197–212. doi: 10.1649/072.065.0301
- Blackledge TA, Scharff N, Coddington JA, Szüts T, Wenzel JW, Hayashi C, Agnarsson I (2009) Reconstructing web evolution and spider diversification in the molecular era. *Proceedings of the National Academy of Sciences* 106 (13): 5229–5234. doi: 10.1073/pnas.0901377106
- Boyer SL, Karaman I, Giribet G (2005) The genus *Cyphophthalmus* (Arachnida, Opiliones, Cyphophthalmi) in Europe: a phylogenetic approach to Balkan Peninsula biogeography. *Molecular Phylogenetics and Evolution* 36 (3): 554–567. doi: 10.1016/j.ympev.2005.04.004
- Boyer S, Baker JM, Gonzalo G (2007) Deep genetic divergences in *Aoraki denticulata* (Arachnida, Opiliones, Cyphophthalmi): a widespread 'mite harvestman' defies DNA taxonomy. *Molecular Ecology* 16 (23): 4999–5016. doi: 10.1111/j.1365-294X.2007.03555.x

- Brunsfeld SJ, Sullivan J, Soltis DE, Soltis PS (2001) Comparative phylogeography of north-western North America: a synthesis. In: Silvertown J, Antonovics J (Eds) *Intergrating Ecological and Evolutionary Processes in a Spatial Context*. Blackwell Science, Oxford, 319–339.
- Brunsfeld SJ, Sullivan J (2006) A multi-compartmented glacial refugium in the northern Rocky Mountains: evidence from the phylogeography of *Cardamine constancei* (Brassicaceae). *Conservation Genetics* 6: 895–904. doi: 10.1007/s10592-005-9076-7
- Carstens BC, Stevenson AL, Degenhardt JD, Sullivan J (2004) Testing nested phylogenetic and phylogeographic hypotheses in the *Plethodon vandykei* species group. *Systematic Biology* 53 (5): 781–792. doi: 10.1080/10635150490522296
- Castresana J (2000) Selection of conserved blocks from multiple alignments for their use in phylogenetic analysis. *Molecular Biology and Evolution* 17 (4): 540–552. doi: 10.1093/oxfordjournals.molbev.a026334
- Derkarabetian S, Steinmann DB, Hedin M (2010) Repeated and time-correlated morphological convergence in cave-dwelling harvestmen (Opiliones, Laniatores) from montane western North America. *PLoS ONE* 5 (5): e10388. doi: 10.1371/journal.pone.0010388
- Ezard T, Pearson P, Purvis A (2010) Algorithmic approach to aid species' delimitation in multidimensional morphospace. *BMC Evolutionary Biology* 10: 175. doi: 10.1186/1471-2148-10-175
- Fisher RA (1936) The use of multiple measurements in taxonomic problems. *Annals of Human Genetics* 7 (2): 179–188. doi: 10.1111/j.1469-1809.1936.tb02137.x
- Flot JF (2010) SEQPHASE: a web tool for interconverting PHASE input/output files and FASTA sequences alignments. *Molecular Ecology Resources* 10 (1): 162–166. doi: 10.1111/j.1755-0998.2009.02732.x
- Franklin JF, Dyrness CT (1988) *Natural Vegetation of Oregon and Washington*. Oregon University Press, Corvallis, 427 pp.
- Gascuel O (1997) BIONJ: an improved version of the NJ algorithm based on a simple model of sequence data. *Molecular Biology and Evolution* 14 (7): 685–695. doi: 10.1093/oxfordjournals.molbev.a025808
- Gates DM (1993) *Climate Change and its Biological Consequences*. Sinauer and Associates, Sunderland, Massachusetts, 280 pp.
- Gebiola M, Gómez-Zurita J, Monti MM, Navone P, Bernardo U (2012) Integration of molecular, ecological, morphological and endosymbiont data for species delimitation within the *Phygadeuon* complex (Hymenoptera: Eulophidae). *Molecular Ecology* 21 (5): 1190–1208. doi: 10.1111/j.1365-294X.2011.05428.x
- Giribet G, Vogt L, Pérez González A, Sharma P, Kury AB (2010) A multilocus approach to harvestman (Arachnida: Opiliones) phylogeny with emphasis on biogeography and the systematics of Laniatores. *Cladistics* 26 (4): 408–437. doi: 10.1111/j.1096-0031.2009.00296.x
- Good DA, Wake DB (1992) *Geographic Variation and Speciation in the Torrent Salamanders of the Genus Rhyacotriton* (Caudata: Rhyacotritonidae). University of California Press 126, Berkeley, 108 pp.
- Grinnell J (1910) The methods and uses of a research museum. *Popular Science Monthly* 77: 163–169.

- Guidon S, Gascuel O (2003) A simple, fast, and accurate algorithm to estimate large phylogenies by maximum likelihood. *Systematic Biology* 52 (5): 696–704. doi: 10.1080/10635150390235520
- Hamm CA (2010) Multivariate discrimination and description of a new species of *Tapinoma* from the western United States. *Annals of the Entomological Society of America* 103 (1): 20–29. doi: 10.1603/008.103.0104
- Harrigan RJ, Mazza ME, Sorenson MD (2008) Computation vs. cloning: Evaluation of two methods for haplotypes determination. *Molecular Ecology Resources* 8 (6): 1239–1248. doi: 10.1111/j.1755-0998.2008.02241.x
- Harvey MS, Rix MG, Framenau VW, Hamilton ZR, Johnson MS, Teale RJ, Humphreys G, Humphreys WF (2011) Protecting the innocent: studying short-range endemic taxa enhances conservation outcomes. *Invertebrate Systematics* 25 (1): 1–10. doi: 10.1071/IS11011
- Hedin M, Thomas S (2010) Molecular systematics of eastern North American Phalangodidae (Arachnida: Opiliones: Laniatores), demonstrating convergent morphological evolution in caves. *Molecular Phylogenetics and Evolution* 54 (1): 107–121. doi: 10.1016/j.ympev.2009.08.020
- Hedin M, Derkarabetian S, McCormack M, Richart C, Shultz JW (2010) The phylogenetic utility of the nuclear protein-coding gene EF-1 α for resolving recent divergences in Opiliones, emphasizing intron evolution. *Journal of Arachnology* 38: 9–20. doi: 10.1636/HA09-49.1
- Jolliffe IT (2002) *Principal Component Analysis – 2nd edition*. Springer, New York, 491 pp.
- Katoh K, Kuma K, Toh H, Miyata T (2005) MAFFT version 5: improvement in accuracy of multiple sequence alignment. *Nucleic Acids Research* 33 (2): 511–518. doi: 10.1093/nar/gki198
- Klimov PB, Lekveishvili M, Dowling APG, OConnor BM (2004) Multivariate analysis of morphological variation in two cryptic species of *Sancassania* (Acari: Acaridae) from Costa Rica. *Annals of the Entomological Society of America* 97: 322–345. doi: 10.1603/0013-8746(2004)097[0322:MAOMVI]2.0.CO;2
- Leonard W, Chichester L, Baugh J, Wilke T (2003) *Kootenaia burkei*, a new genus and species of slug from northern Idaho, United States (Gastropoda: Pulmonata: Arionidae). *Zootaxa* 355: 1–16.
- Leonard WP, Chichester L, Richart CH, Young TA (2011) *Securicauda hermani* and *Carinacauda stormi*, two new genera and species of slug from the Pacific Northwest of the United States (Gastropoda: Stylommatophora: Arionidae), with notes on *Gliabates oregonius* Webb 1959. *Zootaxa* 2746: 43–56.
- Leopold EB, Denton MF (1987) Comparative age of a grassland and steppe east and west of the northern Rocky Mountains. *Annals of the Missouri Botanical Garden* 74 (4): 841–867. doi: 10.2307/2399452
- Lubishew AA (1962) On the use of discriminant functions in taxonomy. *Biometrics* 18 (4): 455–477. doi: 10.2307/2527894
- Maddison DR, Maddison WP (2003) *MacClade version 4.06*. Sinauer, Sunderland, MA.
- May CL, Gresswell RE (2003) Large woody recruitment and redistribution in headwater streams in the southern Oregon Coast Range, U.S.A. *Canadian Journal of Forest Research* 33: 1352–1362. doi: 10.1139/x03-023

- Miller JA, Griswold CE, Haddad CR (2010) Taxonomic revision of the spider family Penestomidae (Araneae, Entelegynae). *Zootaxa* 2534: 1–36.
- Miller MA, Pfeiffer W, Schwartz T (2010) Creating the CIPRES Science Gateway for inference of large phylogenetic trees. *IEEE*: 1–8. doi: 10.1109/GCE.2010.5676129
- Mitchell SG, Montgomery DR (2006) Polygenetic topography of the Cascade Range, Washington State, USA. *American Journal of Science* 306 (9): 736–768. doi: 10.2475/09.2006.03
- Nielson M, Lohman K, Sullivan J (2001) Phylogeography of the tailed frog (*Ascaphus truei*): implications for the biogeography of the Pacific Northwest. *Evolution* 55 (1): 147–160.
- Nielson M, Lohman K, Daugherty CH, Allendorf FW, Knudsen KL, Sullivan J (2006) Allozyme and mitochondrial DNA variation in the tailed frog (Anura: *Ascaphus*): The influence of geography and gene flow. *Herpetologica* 62 (3): 235–258. doi: 10.1655/0018-0831(2006)62[235:AAMDVI]2.0.CO;2
- Posada D (2008) jModelTest: Phylogenetic model averaging. *Molecular Biology and Evolution* 25 (7): 1253–1256. doi: 10.1093/molbev/msn083
- Rambaut A, Drummond AJ (2007) Tracer v1.4, available from <http://beast.bio.ed.ac.uk/Tracer>
- Ronquist F, Huelsenbeck JP (2003) MrBayes 3: Bayesian phylogenetic inference under mixed models. *Bioinformatics* 19 (12): 1572–1574. doi: 10.1093/bioinformatics/btg180
- Satler J, Starrett J, Hayashi CY, Hedin M (2011) Inferring species trees from gene trees in a radiation of California trapdoor spiders (Araneae, Antrodiaetidae, Aliatypus). *PLoS ONE* 6 (9): e25355. doi: 10.1371/journal.pone.0025355
- Schlick-Steiner BC, Steiner FM, Moder K, Seifert B, Sanetra M, Dyreson E, Stauffer C, Christian E (2006) A multidisciplinary approach reveals cryptic diversity in western palearctic *Tetramorium* ants (Hymenoptera: Formicidae). *Molecular Phylogenetics and Evolution* 40 (1): 259–273. doi: 10.1016/j.ympev.2006.03.005
- Schönhofer AL, Martens J (2008) Revision of the genus *Trogulus* Latreille: the *Trogulus coriziformis* species-group of the western Mediterranean (Opiliones: Trogulidae). *Invertebrate Systematics* 22: 523–554. doi: 10.1071/IS08013
- Schönhofer AL, Martens J (2010) Hidden Mediterranean diversity: assessing species taxa by molecular phylogeny within the opilionid family Trogulidae (Arachnida, Opiliones). *Molecular Phylogenetics and Evolution* 54 (1): 59–75. doi: 10.1016/j.ympev.2009.10.013
- Seifert B (2003) *Hypoponera punctatissima* (ROGER) and *H. schauinslandi* (EMERY) – two morphologically and biologically distinct species (Hymenoptera: Formicidae). *Abhandlungen und Berichte des Naturkundemuseums Görlitz* 75: 61–81.
- Sharma PP, Giribet G (2011) The evolutionary and biogeographic history of the armoured harvestmen – Laniatores phylogeny based on ten molecular markers, with the description of two new families of Opiliones (Arachnida). *Invertebrate Systematics* 25 (2): 106–142. doi: 10.1071/IS11002
- Shear WA (1986) A cladistic analysis of the opilionid superfamily Ischyropsalidoidea, with description of the new family Ceratolasmatidae, the new genus *Acuclavella*, and four new species. *American Museum Novitates* 2844: 1–29.
- Shear WA, Leonard WP (2003) Microlympiidae, a new milliped family from North America, and *Microlympia echina*, new genus and species (Diplopoda: Chordeumatida: Brannerioidea). *Zootaxa* 243: 1–11.

- Shear WA, Leonard WP (2004) The milliped family Anthroleucosomatidae new to North America: *Leschius mcallisteri*, n. gen., n. sp. (Diplopoda: Chordeumatida: Anthroleucosomatoidae). *Zootaxa* 609: 1–7.
- Shear WA, Leonard WP (2007) Additions to the milliped family Caseyidae. I. *Caseya richarti*, n. sp., and new records of previously described species in the genus *Caseya* Cook and Collins 1895 (Diplopoda, Chordeumatida, Caseyidae). *Zootaxa* 1524: 23–34.
- Shelley RM, Shear WA (2006) *Leonardesmus injucundus*, n. gen., n. sp., an aromatic, small-bodied milliped from Washington State, USA, and a revised account of the family Nearctodesmidae (Polydesmida). *Zootaxa* 1176: 1–16.
- Shelley RM, Richart C, Bogan AE (2010) *Octoglena claraqua*, n. sp. (Polyzoniida: Hirudisomatidae), a new milliped from Idaho, USA; first record of the order from the western interior of North America. *Zootaxa* 2446: 55–64.
- Souza ES, Willemart RH (2011) Harvest-ironman: heavy armature, and not its defensive secretions, protects a harvestman against a spider. *Animal Behaviour* 81: 127–133. doi: 10.1016/j.anbehav.2010.09.023
- Stamatakis A (2006) RAxML-VI-HPC: maximum likelihood-based phylogenetic analyses with thousands of taxa and mixed models. *Bioinformatics* 22 (21): 2688–2690. doi: 10.1093/bioinformatics/btl446
- Stamatakis A, Hoover P, Rougemont J (2008) A rapid bootstrap algorithm for the RAxML web servers. *Systematics Biology* 57 (5): 758–771. doi: 10.1080/10635150802429642
- Steele CA, Carstens BC, Storfer A, Sullivan J (2005) Testing hypotheses of speciation timing in *Dicamptodon copei* and *Dicamptodon aterrimus* (Caudata: Dicamptodontidae). *Molecular Phylogenetics and Evolution* 36 (1): 90–100. doi: 10.1016/j.ympev.2004.12.001
- Stephens M, Donnelly P (2003) A comparison of Bayesian methods for haplotypes reconstruction from population genotype data. *American Journal of Human Genetics* 73 (5): 1162–1169. doi: 10.1086/379378
- Thomas SM, Hedin M (2008) Multigenic phylogeographic divergence in the paleoendemic southern Appalachian opilionid *Fumuntana deprehendor* Shear (Opiliones, Laniatores, Triaenonychidae). *Molecular Phylogenetics and Evolution* 46 (2): 645–658. doi: 10.1016/j.ympev.2007.10.013
- Vink CJ, Thomas SM, Paquin P, Hayashi CY, Hedin M (2005) The effects of preservatives and temperatures on arachnid DNA. *Invertebrate Systematics* 19 (2): 99–104. doi: 10.1071/IS04039
- White DJ, Wolff JN, Pierson M, Gemmell NJ (2008) Revealing the hidden complexities of mtDNA inheritance. *Molecular Ecology* 17 (23): 4925–4942. doi: 10.1111/j.1365-294X.2008.03982.x

Appendix I

Collection Locality Information (doi: 10.3897/zookeys.311.2920.app1) File format: Adobe PDF file (pdf).

Copyright notice: This dataset is made available under the Open Database License (<http://opendatacommons.org/licenses/odbl/1.0/>). The Open Database License (ODbL) is a license agreement intended to allow users to freely share, modify, and use this Dataset while maintaining this same freedom for others, provided that the original source and author(s) are credited.

Citation: Richart CH, Hedin M (2013) Three new species in the harvestmen genus *Acuclavella* (Opiliones, Dyspnoi, Ischyropsalidoidea), including description of male *Acuclavella quattuor* Shear, 1986. ZooKeys 311: 19–68. doi: 10.3897/zookeys.311.2920.app1

Appendix II

PCR Primer Information. (doi: 10.3897/zookeys.311.2920.app2) File format: Adobe PDF file (pdf).

Copyright notice: This dataset is made available under the Open Database License (<http://opendatacommons.org/licenses/odbl/1.0/>). The Open Database License (ODbL) is a license agreement intended to allow users to freely share, modify, and use this Dataset while maintaining this same freedom for others, provided that the original source and author(s) are credited.

Citation: Richart CH, Hedin M (2013) Three new species in the harvestmen genus *Acuclavella* (Opiliones, Dyspnoi, Ischyropsalidoidea), including description of male *Acuclavella quattuor* Shear, 1986. ZooKeys 311: 19–68. doi: 10.3897/zookeys.311.2920.app2

Appendix III

Male and Female Morphometric Data and Measurement Scheme. (doi: 10.3897/zookeys.311.2920.app3) File format: Microsoft Office Excel (xlsx).

Copyright notice: This dataset is made available under the Open Database License (<http://opendatacommons.org/licenses/odbl/1.0/>). The Open Database License (ODbL) is a license agreement intended to allow users to freely share, modify, and use this Dataset while maintaining this same freedom for others, provided that the original source and author(s) are credited.

Citation: Richart CH, Hedin M (2013) Three new species in the harvestmen genus *Acuclavella* (Opiliones, Dyspnoi, Ischyropsalidoidea), including description of male *Acuclavella quattuor* Shear, 1986. ZooKeys 311: 19–68. doi: 10.3897/zookeys.311.2920.app3

Appendix IV

PCA Methods and Results. (doi: 10.3897/zookeys.311.2920.app4) File format: Adobe PDF file (pdf).

Copyright notice: This dataset is made available under the Open Database License (<http://opendatacommons.org/licenses/odbl/1.0/>). The Open Database License (ODbL) is a license agreement intended to allow users to freely share, modify, and use this Dataset while maintaining this same freedom for others, provided that the original source and author(s) are credited.

Citation: Richart CH, Hedin M (2013) Three new species in the harvestmen genus *Acuclavella* (Opiliones, Dyspnoi, Ischyropsalidoidea), including description of male *Acuclavella quattuor* Shear, 1986. ZooKeys 311: 19–68. doi: 10.3897/zookeys.311.2920.app4

Appendix V

Morphologies of *Acuclavella* cf. *cosmetoides* and Geography of Morphotypes. Figure 1: male morphologies. Figure 2: female morphologies. Figure3: geographic distribution of male morphologies. Figure 4:geographic distribution of female morphologies. (doi: 10.3897/zookeys.311.2920.app5) File format: Adobe PDF file (pdf).

Copyright notice: This dataset is made available under the Open Database License (<http://opendatacommons.org/licenses/odbl/1.0/>). The Open Database License (ODbL) is a license agreement intended to allow users to freely share, modify, and use this Dataset while maintaining this same freedom for others, provided that the original source and author(s) are credited.

Citation: Richart CH, Hedin M (2013) Three new species in the harvestmen genus *Acuclavella* (Opiliones, Dyspnoi, Ischyropsalidoidea), including description of male *Acuclavella quattuor* Shear, 1986. ZooKeys 311: 19–68. doi: 10.3897/zookeys.311.2920.app5

Appendix VI

Acuclavella distribution. KMZ file of *Acuclavella* sampling effort, species distributions, and distribution of individuals used in concatenated phylogenetic reconstruction. (doi: 10.3897/zookeys.311.2920.app6) File format: Google Earth Keyhole Markup (kmz).

Copyright notice: This dataset is made available under the Open Database License (<http://opendatacommons.org/licenses/odbl/1.0/>). The Open Database License (ODbL) is a license agreement intended to allow users to freely share, modify, and use this Dataset while maintaining this same freedom for others, provided that the original source and author(s) are credited.

Citation: Richart CH, Hedin M (2013) Three new species in the harvestmen genus *Acuclavella* (Opiliones, Dyspnoi, Ischyropsalidoidea), including description of male *Acuclavella quattuor* Shear, 1986. ZooKeys 311: 19–68. doi: 10.3897/zookeys.311.2920.app6

Appendix VII

Phylogenetic Trees. Figure 1: Outgroup relationships from Bayesian gene trees. Figure 2: RAxML concatenated phylogeny. Figure 3: RAxML COI gene tree. Figure 4: RAxML EF-1 α gene tree. Figure 5: 28S and Wnt2 RAxML gene trees. (doi: 10.3897/zookeys.311.2920.app7) File format: Adobe PDF file (pdf).

Copyright notice: This dataset is made available under the Open Database License (<http://opendatacommons.org/licenses/odbl/1.0/>). The Open Database License (ODbL) is a license agreement intended to allow users to freely share, modify, and use this Dataset while maintaining this same freedom for others, provided that the original source and author(s) are credited.

Citation: Richart CH, Hedin M (2013) Three new species in the harvestmen genus *Acuclavella* (Opiliones, Dyspnoi, Ischyropsalidoidea), including description of male *Acuclavella quattuor* Shear, 1986. ZooKeys 311: 19–68. doi: 10.3897/zookeys.311.2920.app7

Appendix VIII

Acuclavella genitalia. Figure 1: penises. Figure 2: ovipositors. (doi: 10.3897/zookeys.311.2920.app8) File format: Adobe PDF file (pdf).

Copyright notice: This dataset is made available under the Open Database License (<http://opendatacommons.org/licenses/odbl/1.0/>). The Open Database License (ODbL) is a license agreement intended to allow users to freely share, modify, and use this Dataset while maintaining this same freedom for others, provided that the original source and author(s) are credited.

Citation: Richart CH, Hedin M (2013) Three new species in the harvestmen genus *Acuclavella* (Opiliones, Dyspnoi, Ischyropsalidoidea), including description of male *Acuclavella quattuor* Shear, 1986. ZooKeys 311: 19–68. doi: 10.3897/zookeys.311.2920.app8

Appendix IX

K. Leg II Morphologies. With and without femoral false leg articulations. (doi: 10.3897/zookeys.311.2920.app9) File format: Adobe PDF file (pdf).

Copyright notice: This dataset is made available under the Open Database License (<http://opendatacommons.org/licenses/odbl/1.0/>). The Open Database License (ODbL) is a license agreement intended to allow users to freely share, modify, and use this Dataset while maintaining this same freedom for others, provided that the original source and author(s) are credited.

Citation: Richart CH, Hedin M (2013) Three new species in the harvestmen genus *Acuclavella* (Opiliones, Dyspnoi, Ischyropsalidoidea), including description of male *Acuclavella quattuor* Shear, 1986. ZooKeys 311: 19–68. doi: 10.3897/zookeys.311.2920.app9

Three new species of the genus *Cymodusa* Holmgren (Hymenoptera, Ichneumonidae, Campopleginae) from Korea

Jin-Kyung Choi^{1,†}, Janko Kolarov^{2,‡}, Jong-Wook Lee^{1,§}

1 Department of Life Sciences, Yeungnam University, Gyeongsan, 712-749, Korea **2** Faculty of Pedagogy, University of Plovdiv, 24 Tsar Assen Str., 4000 Plovdiv, Bulgaria

† [urn:lsid:zoobank.org:author:23C34EE5-685C-4FF8-8E22-B560D94A44BE](https://doi.org/urn:lsid:zoobank.org:author:23C34EE5-685C-4FF8-8E22-B560D94A44BE)

‡ [urn:lsid:zoobank.org:author:EAA20BD6-53B6-4144-99B4-7C874B5E99D5](https://doi.org/urn:lsid:zoobank.org:author:EAA20BD6-53B6-4144-99B4-7C874B5E99D5)

§ [urn:lsid:zoobank.org:author:29B0EAD6-5F06-46DA-A384-69FDE8CBEF34](https://doi.org/urn:lsid:zoobank.org:author:29B0EAD6-5F06-46DA-A384-69FDE8CBEF34)

Corresponding author: Jong-Wook Lee (jwlee1@ynu.ac.kr)

Academic editor: Gavin Broad | Received 16 May 2013 | Accepted 18 June 2013 | Published 20 June 2013

[urn:lsid:zoobank.org:pub:A8BBF6FF-DCA7-4DD5-A26E-E19DC170DEAE](https://doi.org/urn:lsid:zoobank.org:pub:A8BBF6FF-DCA7-4DD5-A26E-E19DC170DEAE)

Citation: Choi, J-K Kolarov J, Lee J-W (2013) Three new species of the genus *Cymodusa* Holmgren (Hymenoptera, Ichneumonidae, Campopleginae) from Korea. ZooKeys 311: 69–82. doi: 10.3897/zookeys.311.15529

Abstract

Korean species of the genus *Cymodusa* Holmgren (Hymenoptera: Ichneumonidae: Campopleginae) are reviewed. Four species of *Cymodusa* (*Cymodusa*) are reported from Korea, including one newly recorded species, *Cymodusa aenigma* Dbar (1985), and three new species, *Cymodusa koreana* **sp. n.**, *Cymodusa yeungnamensis* **sp. n.** and *Cymodusa geolimi* **sp. n.** This genus is reported for the first time from Korea. Descriptions with photographs of new species, line drawings of propodeum and metasomal tergites of the Palearctic species of the “*australis*” group and a key to the Korean *Cymodusa* species are provided.

Keywords

Cymodusa aenigma Dbar, *Cymodusa koreana* sp. n., *Cymodusa yeungnamensis* sp. n., *Cymodusa geolimi* sp. n., taxonomy

Introduction

The genus *Cymodusa* Holmgren (1859) is a moderately large genus in the subfamily Campopleginae Förster with contains about 40 described species from the Eastern Palearctic, Nearctic, Neotropical, Oriental and Western Palearctic regions (Yu et al. 2012).

Since Holmgren (1859) described *Cymodusa leucocera*, 38 species have been recorded. Recently, Kolarov and Coruh (2008) and Kolarov and Yurtcan (2008) reported two new species from Turkey. In the Eastern Palaearctic region there are 15 species (Yu et al. 2012). This genus usually occurs among grasses and they are parasitoids of Lepidoptera (Kolarov and Yurtcan 2008). One of the collecting sites was clothed in trees, herbaceous plants and shrubs. The Palaearctic species have been reviewed and keyed by Dbar (1984, 1985).

Dbar (1985) recognizes four species groups, namely the “*jaceki*” group, “*leucocera*” group, “*australis*” group and “*convergator*” group. The “*australis*” group includes eight species, *C. australis* Smits van Burgst (1913), *C. orientalis* Uchida (1956), *C. longiterebra* Dbar (1985), *C. tibialis* Dbar (1985), *C. rufiventris* Dbar (1985), *C. aenigma* Dbar (1985), *C. parva* Dbar (1985) and *C. oculator* Dbar (1985), in the Palaearctic region as well as three species and one subspecies, *C. dravida* Gupta & Gupta (1974), *C. josephi* Gupta & Gupta (1974), *C. josephi malaise* Gupta & Gupta (1974) and *C. shiva* Gupta & Gupta (1974), in the Oriental region.

The “*australis*” group can easily be distinguished from the other species groups by the following characteristics: temple very narrowed; areolet with the 2nd recurrent vein before the middle; 6th and 7th tergites deeply emarginate in dorsal view.

In this study, we report the “*australis*” group of *Cymodusa* for the first time from Korea, including one newly recorded species and three new species.

Materials and methods

Materials used in this work were collected by sweeping and Malaise trapping, and were deposited in the animal systematic laboratory of the Yeungnam University (YNU, Gyeongsan, Korea). Some specimens examined in this study were loaned by the Naturhistoriska Riksmuseet, Sektionen for Entomologi (Swedish Museum of Natural History Department of Entomology) (NR, Stockholm, Sweden). Specimens were examined using a stereo microscope (Zeiss Stemi SV 11 Apo; Carl Zeiss, Göttingen, Germany) and key characters shown in the photographs were produced using a Delta imaging system (i-Delta 2.6; iMTechnology, Daejeon, Korea). The morphological terminology is mostly that of Gupta & Maheshwary (1977). Abbreviations are as follows. TD, type depository; TS, type species; ZI, Zoological Institute, Academy of Sciences, St. Petersburg 199034, Russia; GW, Gangwon-do; GB, Gyeongsangbuk-do; GN, Gyeongsangnam-do.

Results

Key to the Korean *Cymodusa*

- 1 Costula of propodeum complete (Fig. 5A); spiracle of propodeum very small circle, its diameter shorter than carina linking spiracle to pleural carina (Fig. 1I) *Cymodusa aenigma*

- Costula of propodeum incomplete or absent (Fig. 5I); spiracle of propodeum large, its diameter longer than carina linking spiracle to pleural carina (Fig. 2H)..... **2**
- 2 Mandible blackish brown (Figs. 4C and D); areolet of fore wing with stalk (Fig. 4M)..... ***Cymodusa geolimi* sp. n.**
- Mandible yellowish brown (Figs. 3C); areolet of fore wing without stalk (Fig. 3L)..... **3**
- 3 Spiracle of propodeum large (Fig. 3I); first to third flagellomeres to yellowish brown (Fig. 3M); notaulus absent (Fig. 3F) ... ***Cymodusa yeungnamensis* sp. n.**
- Spiracle of propodeum small (Fig. 2H); all antennal flagellomeres blackish brown; notaulus present as weak trace (Fig. 2F) ... ***Cymodusa koreana* sp. n.**

Family Ichneumonidae Latreille, 1802

Subfamily Campopleginae Förster, 1869

Genus *Cymodusa* (*Cymodusa*) Holmgren, 1859

<http://species-id.net/wiki/Cymodusa>

Cymodusa Holmgren, 1859: 327. TS: *Cymodusa leucocera* Holmgren

Thersilia Schmiedeknecht, 1907: 598. TS: (*Thersitia egregia* Schmiedeknecht) = *Cymodusa leucocera* Holmgren.

Diagnosis. Eyes densely setose. Inner margins of eyes strongly convergent ventrally. Clypeus small; apical margin rounded; weakly convex, not separated from face. Mandible short, upper tooth as long as or longer than lower tooth. Hind basitarsus without a median ventral row of close setae. Fore wing with areolet; nervulus opposite to basal vein or distad of it. Thyridium longitudinally elliptic. Ovipositor straight, with dorsal subapical notch.

Cymodusa (*Cymodusa*) *aenigma* Dbar, 1985

http://species-id.net/wiki/Cymodusa_aenigma

Figs 1, 5A, 6A

Cymodusa (*Cymodusa*) *aenigma* Dbar, 1985: 589. Holotype: female; TD: ZI.

Material examined. [Korea]: 2 females, Mureung valley, Samhwa-dong, Donghae-si GW, Korea, 15 July–1 August 2005, MT(Malaise trap), J.W. Lee; 1 female, ditto, 9–17 August 2005, MT, J.W. Lee; 1 female, ditto, 31 August–10 September 2005, MT, J.W. Lee.

Description. Female. Body length 6.1 mm.

Fore wing length 3.3–3.8 mm.

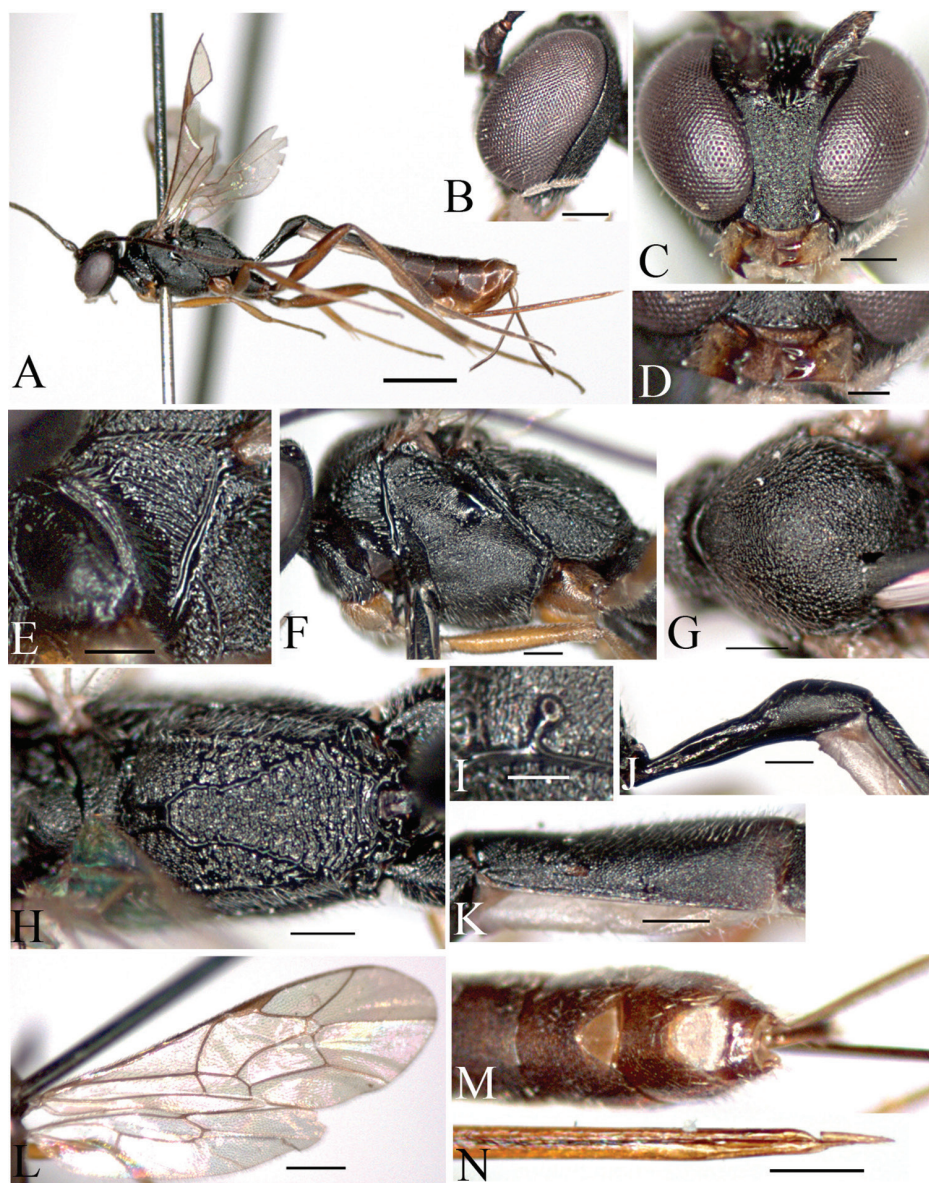


Figure 1. *Cymodusa (Cymodusa) aenigma* Dbar, 1985 (female). **A** habitus in lateral view **B** head in lateral view **C** head in frontal view **D** mandible **E** pronotum **F** mesopleuron in lateral view **G** mesoscutum in dorsal view **H** propodeum **I** spiracle of propodeum **J** petiole in lateral view **K** thyridium **L** wings **M** 6th and 7th tergites in dorsal view **N** ovipositor. (Scale bar 1 mm for A; 0.5 mm for L; 0.2 mm for B, C, E–H, J, K, M and N; 0.1 mm for D and I).

Antenna with 34–35 flagellomeres.

Color. Head black. Scape and pedicel blackish brown. Mandible yellow, apically brown; labial and maxillary palps pale yellow. Antenna blackish brown. Mesosoma

black. Tegula brown. Fore leg yellow; mid leg yellowish brown; hind coxa black, trochanter dark brown, femur and tibia brown, tarsus blackish brown. Metasoma black. Thyridium reddish brown. 5th to 7th metasomal segments reddish brown. Ovipositor yellowish brown.

Morphology. Head: Head finely and densely punctate. Occiput flat and polished. Temple broad and convex, finely punctate. Mandible very short, upper tooth longer than lower tooth (Fig. 1D). Ocelli slightly large; diameter of median ocellus 1.1 times as long as distance between ocellus and eye; lateral ocellus separated from eye by 0.9 times lateral ocellus diameter.

Mesosoma: Pronotum covered with transverse striae (Fig. 1E); epomia distinct. Mesoscutum finely and densely punctate; notaulus absent (Fig. 1G). Mesopleuron roughly punctate, reticulate; postpectal carina complete; epicnemial carina short; sternaulus absent; speculum convex, impunctate, polished (Fig. 1F); scutellum finely granulate. Propodeum with numerous transverse rugae and reticulate; basal area and areola separated by anterior transverse carina (Fig. 1H); areola and petiolar area not separated, impressed medially; costula complete (Fig. 5A); spiracle very small, round, connected to pleural carina; distance between spiracle and pleural carina 1.7 times as diameter of spiracle (Fig. 1I). Hind tibia with short spines; ratios between hind tarsal segments 4.5: 2.2: 1.6: 0.8: 0.9. Fore wing with areolet; basal vein opposite nervulus (Fig. 1L); nervellus not intercepted; discoidella absent.

Metasoma: Thyridium small, oval; separated from base of 2nd tergite by more than 3.0 times its diameter; distance between base of 2nd tergite and thyridium 0.6 times distance between base of 2nd tergite and spiracle (Fig. 1K). 6th and 7th tergites deeply emarginate apically (Fig. 1M). Ovipositor (Fig. 1N) shorter than metasoma, 1.4 times as long as hind tibia.

Male. Unknown.

Distribution. Korea (new record), Japan, Russia (Khabarovsk, Primor'ye Kray).

***Cymodusa (Cymodusa) koreana* Choi & Lee, sp. n.**

urn:lsid:zoobank.org:act:AEED4C3-D4B6-4E22-9E00-C20FB416D51E

http://species-id.net/wiki/Cymodusa_koreana

Figs 2, 5I, 6F

Material examined. Holotype: [Korea] (TD: YNU): 1 female, Unmunsa, Chungdo-gun, GB, Korea, 21.V.1989, I.S. Ye.

Paratypes: [Korea] (TD: YNU): 1 female, Gyeonbongsa, Ganseong, GW, Korea, 22 May 1992, J.W. Lee; 1 female, Mureung Valley, Samhwa-dong, Donghae-si, GW, Korea, 20 September–2 October 2006, MT, J.W. Lee; 1 female, ditto, 28 August–10 September 2006, MT, J.W. Lee; 1 female, Yeungnam Univ., Gyeongsan-si, GB, 13 May 1985, E.S. Kim; 1 female, Mirimsan, Bonghwa-gun, 4 May 1997, J.C. Jeong.

Description (female holotype). Body length 7.3 mm.

Fore wing length 4.1 mm.

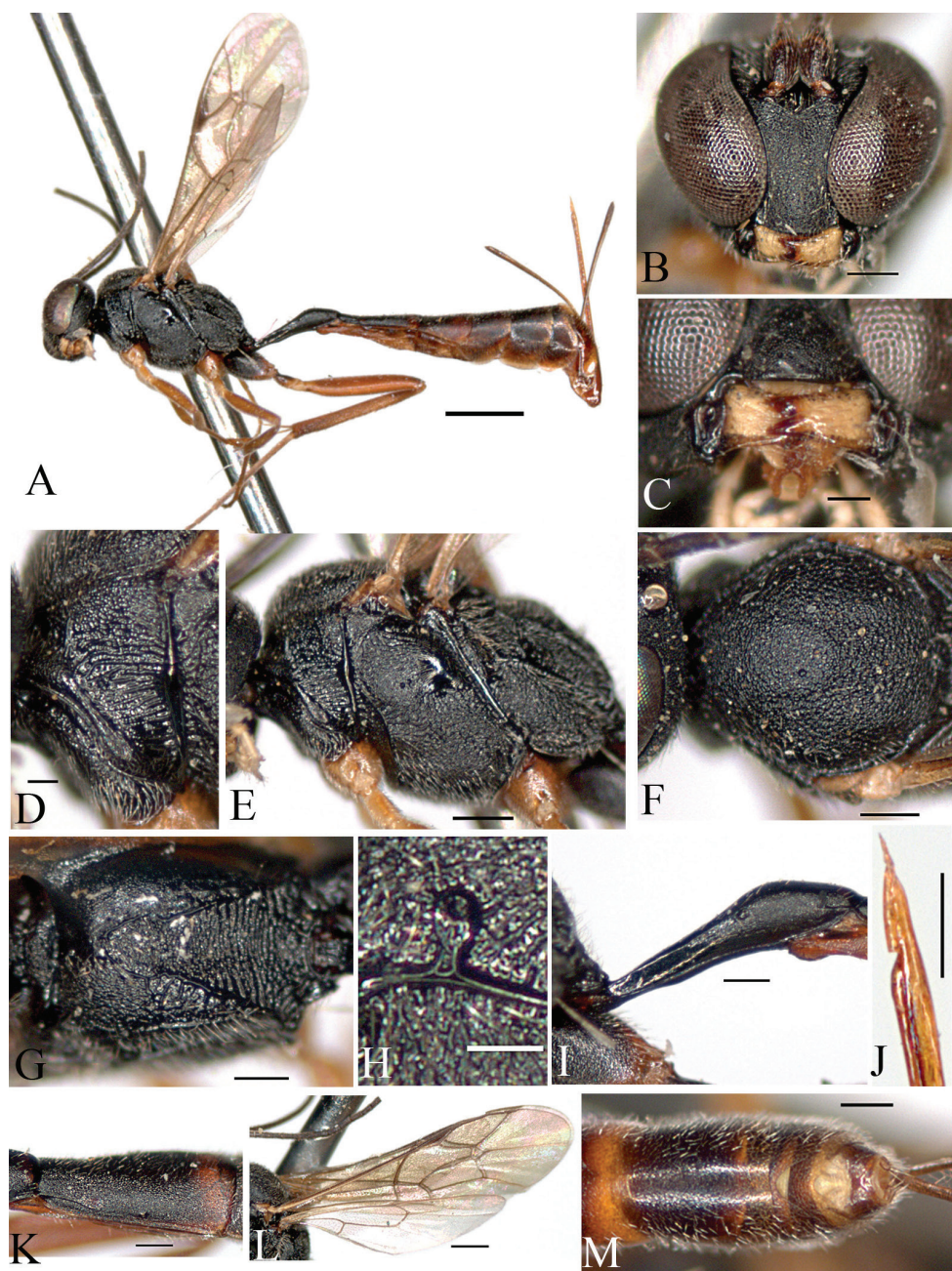


Figure 2. *Cymodusa (Cymodusa) koreana* Choi & Lee sp. n. (female). **A** habitus in lateral view **B** head in frontal view **C** mandible **D** pronotum **E** mesopleuron in lateral view **F** mesoscutum in dorsal view **G** propodeum **H** spiracle of propodeum **I** petiole in lateral view **J** ovipositor **K** thyridium **L** wings **M** 6th and 7th tergites in dorsal view. (Scale bar 1 mm for **A**; 0.5 mm for **L**; 0.2 mm for **B**, **E**–**G**, **I**–**K** and **M**; 0.1 mm for **C**, **D** and **H**).

Antenna with 31–32 flagellomeres.

Color. Head, scape and pedicel black. Mandible yellow, apically brown. Labial and maxillary palps pale yellow. Antenna blackish brown. Mesosoma black. Tegula yellow. Fore leg yellow; mid leg yellowish brown; hind coxa black, trochanter dark brown, trochantellus yellow, femur brown, tibia yellow medially, brown basally and apically, tarsus brown. Metasoma black, tergites apically narrowly reddish brown. Thyridium reddish brown.

Morphology. Head: Head finely and densely punctate. Occiput flat and polished. Temple broad and convex, finely punctate. Mandible very short, upper tooth as long as lower tooth (Fig. 2C). Minimum distance between eyes 0.6 times as long as maximum distance (Fig. 2B). Ocelli small; diameter of median ocellus 0.9 times as long as distance between ocellus and eye; lateral ocellus separated from eye by 1.3 times of lateral ocellus diameter. Antenna with 31–32 flagellomeres, 1st flagellomere 1.3 times as long as 2nd flagellomere.

Mesosoma: Pronotum with transverse striae ventrally (Fig. 2D); epomia weak. Mesoscutum finely and densely punctate; notaulus absent (Fig. 2F). Mesopleuron roughly punctate, reticulate; postpectal carinae complete, epicnemial carina short; sternaulus absent. Scutellum finely punctate; postscutellum broader than high. Propodeum with numerous transverse rugae and reticulate; basal area and areola separated by anterior transverse carina; costula absent (Fig. 5I); areola and petiolar area not separated, impressed medially (Fig. 2G); spiracle small, round, connected to pleural carina, distance between spiracle and pleural carina 0.9 times as long as diameter of spiracle (Fig. 2H). Hind tibia with short spines; ratio between hind tarsal segments 5.0: 2.2: 1.5: 0.9: 0.9. Fore wing with areolet; basal vein opposite nervulus; nervellus not intercepted; discoidella absent (Fig. 2L).

Metasoma: Thyridium small, oval; separated from base of 2nd tergite by more than 2.0 times its diameter; distance between base of 2nd tergite and thyridium 0.5 times as long as distance between base of 2nd tergite and spiracle (Fig. 2K). 6th and 7th tergites deeply emarginate apically (Fig. 2M). Ovipositor 1.5 times as long as hind tibia (Fig. 2J).

Male. Unknown.

Distribution. Korea.

Etymology. The specific name is derived from Korea, the locality of the type specimens.

Comments. This species is similar to *C. rufiventris* Dbar, 1985 in the structure of the propodeum and costula, but the basal area is different, areolet is sessile and pentagonal, and the metasoma is darker.

***Cymodusa (Cymodusa) yeungnamensis* Choi & Lee, sp. n.**

urn:lsid:zoobank.org:act:2BF6F784-E72A-411C-BB6D-44DDD147738B

http://species-id.net/wiki/Cymodusa_yeungnamensis

Figs 3, 5J, 6G

Material examined. Holotype: [Korea] (TD: YNU): 1 female, Yeungnam Univ., Gyeongsan-si, GB, Korea, 21 May 1990, M.J. Kim.



Figure 3. *Cymodusa (Cymodusa) yeungnamensis* Choi & Lee sp. n. (female). **A** habitus in lateral view **B** head in frontal view **C** mandible **D** pronotum **E** mesopleuron in lateral view **F** mesoscutum in dorsal view **G** ovipositor **H** propodeum **I** spiracle of propodeum **J** petiole in lateral view **K** thyridium **L** wings **M** antenna **N** 6th and 7th tergites in dorsal. (Scale bar 1 mm for **A**; 0.5 mm for **L**; 0.2 mm for **B–H**, **J**, **K**, **M** and **N**; 0.1 mm for **I**).

Paratype: [Korea] (TD: YNU): 1 female, Yeungnam Univ., Gyeongsan-si, GB, Korea, 19 June 1992, G.Y. Lee.

Description (female holotype). Body length 6.9 mm.

Fore wing length 4.0 mm.

Antenna with 18+ flagellomeres, apical flagellomeres missing. (antenna with 32 flagellomeres at paratype)

Color. Head black. Scape and pedicel blackish brown. Antenna black, except 3 antennal flagellomeres yellow. Mandible yellow, brown apically. Mesosoma black; tegula brown. Fore leg yellowish brown; mid coxa black, brown apically, trochanter and trochantellus yellow, femur to tarsus brown; hind coxa and trochanter black, trochantellus yellow, femur blackish brown, tibia reddish brown, blackish brown basally and apically, tarsus blackish brown. Metasoma blackish brown; petiole black. Thyridium reddish brown. Ovipositor brown.

Morphology. Head: Head densely finely punctate; Vertex slightly punctate. Occiput flat and polished. Temple finely punctate and flat. Mandible very short, upper tooth as long as lower one (Fig. 3C). Minimum distance between eyes 0.6 times as long as maximum distance (Fig. 3B). Ocelli small; diameter of median ocellus 0.8 times as long as distance between ocellus and eye; lateral ocellus separated from eye by 1.6 times of lateral ocellus diameter. Antenna with 32 flagellomeres, 1st flagellomere 1.5 times as long as 2nd flagellomere.

Mesosoma: Pronotum sparsely punctate; upper part reticulate; ventrally with transverse striae (Fig. 3D); epomia absent. Mesoscutum closely and finely punctate; notaulus absent (Fig. 3F). Mesopleuron reticulate; postpectal carinae complete; epicnemial carina reaching anterior margin of mesopleuron at its middle (Fig. 3E); epicnemial carina present; sternaulus absent. Scutellum closely punctate; postscutellum flat, broader than high. Propodeum reticulated; basal area and areola not separated by anterior transverse carina; costula present but incomplete (Fig. 5J); areola and petiolar area not separated, impressed (Fig. 3H); spiracle round; connected to pleural carina; distance between spiracle and pleural carina 0.7 times as long as diameter of spiracle (Fig. 3I). Hind tibia with short spines; ratio between hind tarsal segments 5.1: 2.4: 1.6: 0.9: 1.0. Fore wing with areolet (Fig. 3L); basal vein opposite nervellus; nervellus not intercepted; discoidella absent.

Metasoma: Thyridium separated from base of 2nd tergite by more than 3.0 times its diameter; distance between base of 2nd tergite and thyridium 0.5 times as long as distance between base of 2nd tergite and spiracle (Fig. 3K). 6th and 7th tergites deeply emarginate apically (Fig. 3N). Ovipositor (Fig. 3G) 1.6 times as long as hind tibia.

Male. Unknown.

Distribution. Korea.

Etymology. The specific name is derived from the locality of the type specimens.

Comments. This species is similar to *C. aenigma* Dbar, 1985, but the areolet is not sessile and the basal 1st to 3rd flagellomeres are yellow below (as in *C. leucocera* Holmgren and *C. distincta* Cresson).

***Cymodusa (Cymodusa) geolimi* Choi & Lee, sp. n.**

urn:lsid:zoobank.org:act:78152C58-FD90-4622-B380-53174A5C55BC

http://species-id.net/wiki/Cymodusa_geolimi

Figs 4, 5K, 6H

Material examined. Holotype: [Korea] (TD: YNU): 1 female, Mureung valley, Samhwa-dong, Donghae-si GW, Korea, 16-28 June 2005, MT, J.W. Lee.

Paratype: [Korea] (TD: YNU): 1 female, Gajoa-dong, Jinju-si, GN, Korea, 3-9 June 1989, J.W. Lee.

Description (female holotype). Body length 5.8 mm.

Fore wing length 3.4 mm.

Antenna with 31 flagellomeres.

Color. Head black. Scape and pedicel blackish brown. Antenna black. Mandible yellow, brown apically. Mesosoma black; tegula brown. Fore leg yellowish brown; mid coxa black, brown apically, trochanter and trochantellus yellow, femur to tarsus brown; hind coxa and trochanter black, trochantellus yellow, femur blackish brown, tibia reddish brown, blackish brown basally and apically, tarsus blackish brown. Metasoma blackish brown; petiole black. Thyridium reddish brown. Ovipositor brown.

Morphology. Head: Head closely and finely punctate. Vertex slightly punctate. Occiput flat and polished. Temple finely punctate and flat. Mandibles short, upper tooth as long as lower one. Minimum distance between eyes 0.6 times as long as maximum distance (Fig. 4C). Ocelli small; diameter of median ocellus 0.9 times as long as distance between ocellus and eye; lateral ocellus separated from eye by 1.0 times lateral ocellus diameter. Antenna with 31 flagellomeres, 1st flagellomere 1.3 times as long as 2nd flagellomere.

Mesosoma: Pronotum sparsely punctate; upper part reticulated; lower part with transverse striae; epomia absent. Mesoscutum closely and finely punctate; notaulus absent (Fig. 4G). Mesopleuron reticulate (Fig. 4F); postpectal carinae complete; epicnemial carina present; sternaulus absent. Scutellum closely punctate; postscutellum flat, broader than high. Propodeum reticulate; basal area and areola separated by anterior transverse carina (Fig. 5K); costula weak; areola and petiolar area not separated, impressed; spiracle small, round; distance between spiracle and pleural carina 1.00 times diameter of spiracle (Fig. 4I). Hind tibia with short spines; ratio between hind tarsal segments 4.4: 2.0: 1.3: 0.7: 0.9. Fore wing with areolet; basal vein opposite nervulus (Fig. 4M); nervellus not intercepted; discoidella absent.

Metasoma: Thyridium separated from base of 2nd tergite by more than 4.0 times its diameter; distance between base of 2nd tergite and thyridium 0.6 times as long as distance between base of 2nd tergite and spiracle (Fig. 4L). 6th and 7th terga deeply emarginate apically (Fig. 4N); 7th tergite longer than 6th tergite (Fig. 6H). Ovipositor (Fig. 4K) shorter than metasoma, 1.7 times as long as hind tibia.

Male. Unknown.

Distribution. Korea.

Etymology. The species is named after the nickname of Dr. Jong-Wook Lee, who collected the type specimens.

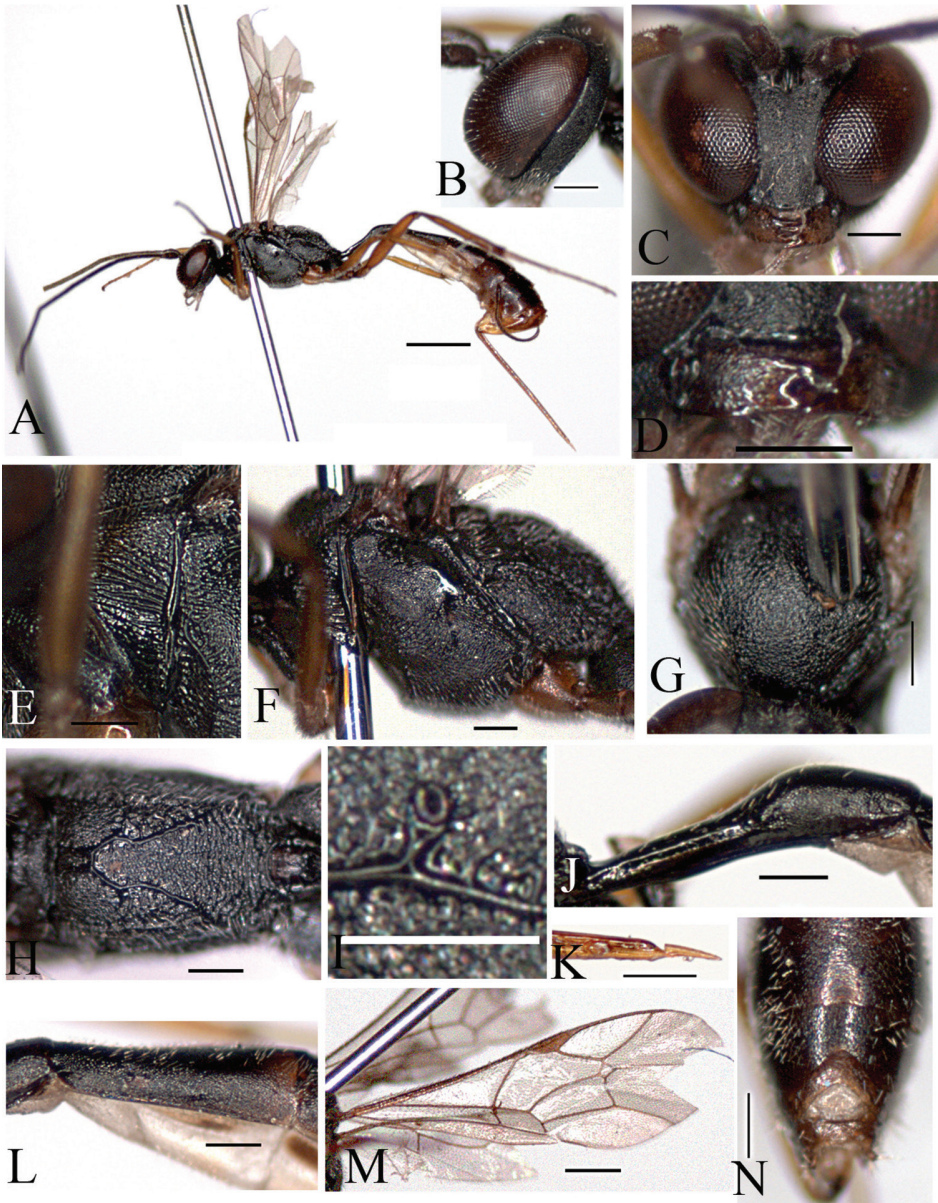


Figure 4. *Cymodusa (Cymodusa) geolimi* Choi & Lee sp. n. (female). **A** habitus in lateral view **B** head in lateral view **C** head in frontal view **D** mandible **E** pronotum **F** mesopleuron in lateral view **G** mesoscutum in dorsal view **H** propodeum **I** spiracle of propodeum **J** petiole in lateral view **K** ovipositor **L** thyridium **M** wings **N** 6th and 7th tergites in dorsal view. (Scale bar 1 mm for **A**; 0.5 mm for **M**; 0.2 mm for **B–L** and **N**).

Comments. This species is similar to *C. oculator* Dbar (1985) but differs by the number of flagellar segments, developed clypeal fovea, basal area and areola not separated, and a different color pattern.

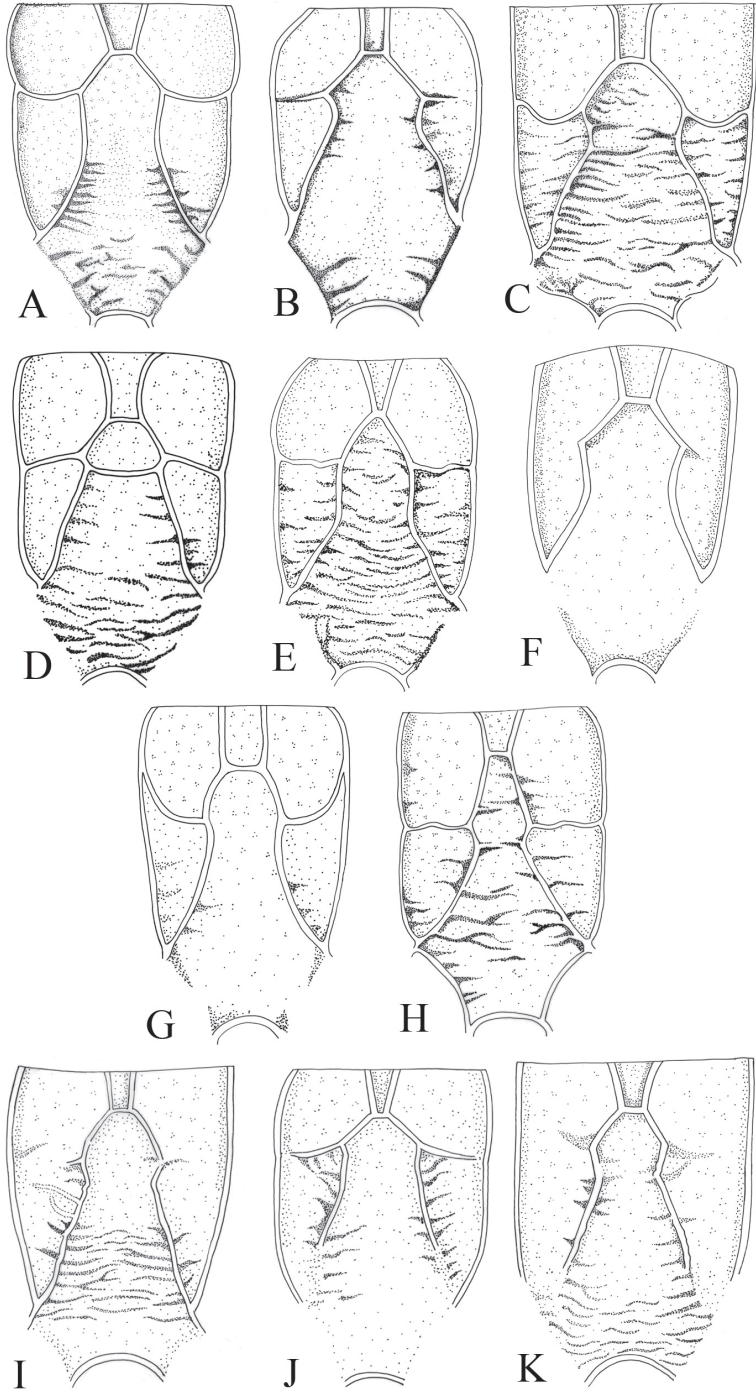


Figure 5. Propodeum, dorsal, of “australis” group. **A** *Cymodusa aenigma* **B** *C. australis* **C** *C. longiterebra* **D** *C. oculator* **E** *C. orientalis* **F** *C. parva* **G** *C. rufiventris* **H** *C. tibialis* **I** *C. koreana* sp. n. **J** *C. yeungnamensis* sp. n. **K** *C. geolimi* sp. n.

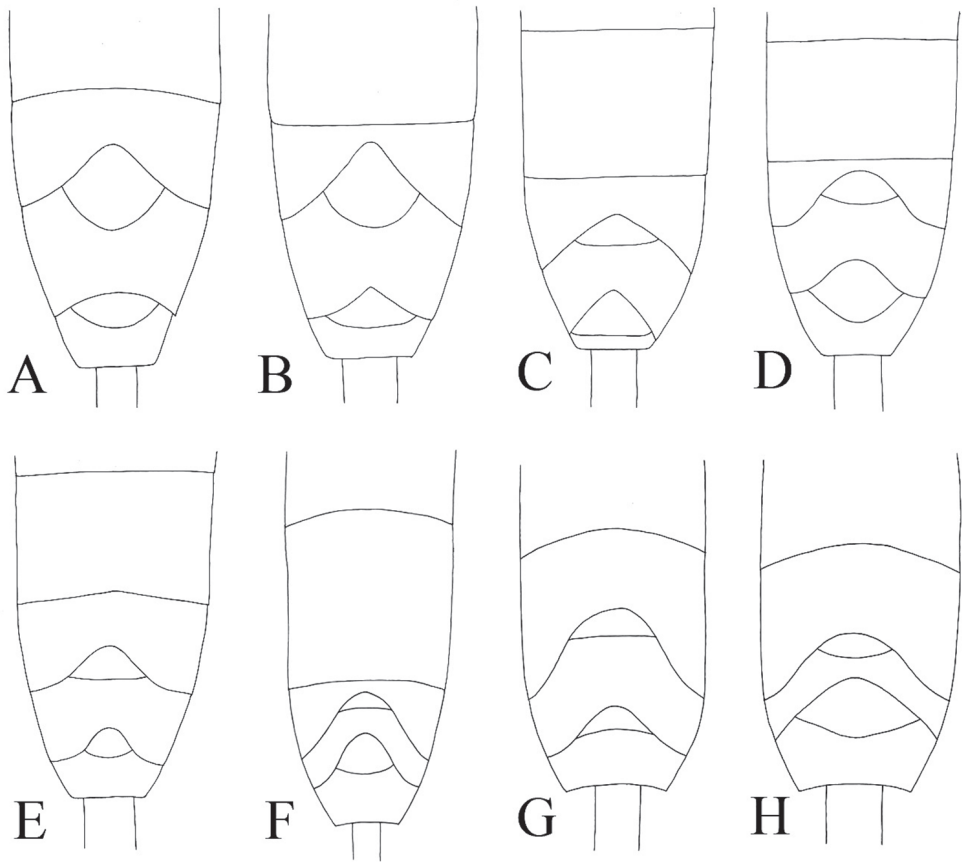


Figure 6. 6th and 7th metasomal terga, dorsal, of “*australis*” group. **A** *Cymodusa aenigma* **B** *C. longiterebra* **C** *C. orientalis* **D** *C. rufiventris* **E** *C. tibialis* **F** *C. koreana* sp. n. **G** *C. yeungnamensis* sp. n. **H** *C. geolimi* sp. n.

Acknowledgements

We are deeply grateful to Dr. Gavin Broad and anonymous reviewer for reviewing this manuscript, and thank Dr. Hege Vårdal of the Swedish Museum of Natural History Stockholm, Sweden, for providing type and voucher specimens used in this study. This work was supported by the 2013 Yeungnam University Research Grant and a grant from the National Institute of Biological Resources (NIBR), funded by the Ministry of Environment (MOE) of the Republic of Korea (NIBR No. 2013-02-001).

References

Dbar RS (1984) Revision of the Palearctic species of the genus *Cymodusa* Holmgren (Hymenoptera, Ichneumonidae). *Entomologicheskoye Obozreniye* 63(4): 802–812.

- Dbar RS (1985) Revision of the Palearctic species of *Cymodusa* Holmgren (Hymenoptera, Ichneumonidae). II. Entomologicheskoye Obozreniye 64(3): 588–600.
- Förster A (1869) Synopsis der Familien und Gattungen der Ichneumoniden. Verhandlungen des Naturhistorischen Vereins der Preussischen Rheinlande und Westfalens 25(1868): 135–221.
- Gupta VK, Gupta ML (1974) The Oriental species of *Cymodusa* Holmgren (Hymenoptera: Ichneumonidae). Oriental Insects 8: 1–14. doi: 10.1080/00305316.1974.10434434
- Gupta VK, Maheshwary S (1977) Ichneumonologia Orientalis, Part IV. The tribe Porozonitini (=Campoplegini) (Hymenoptera: Ichneumonidae). Oriental Insects Monograph. 5: 1–267.
- Holmgren AE (1859) Conspectus generum Ophionidum Sueciae. fversigt af Kongliga Vetenskaps-Akademiens Förhandlingar 15(1858): 321–330.
- Kolarov J, Coruh S (2008) A new species of *Cymodusa* (*Cymodusa*) Holmgren, [1859] (Hymenoptera: Ichneumonidae: Campopleginae) from Turkey. Entomological news 119(3): 291–294. doi: 10.3157/0013-872X(2008)119[291:ANSOCC]2.0.CO;2
- Kolarov J, Yurtcan M (2008) A new species of *Cymodusa* Holmgren, 1859 (Hymenoptera: Ichneumonidae: Campopleginae) from Turkey. Biologia 63(4): 548–549. doi: 10.2478/s11756-008-0088-x
- Latreille PA (1802) Histoire naturelle, générale et particuliere, des Crustacés et des Insectes. Tome troisième. Paris, 468 pp. [Ichneumonidae pp. 318–327]
- Momoi S (1970) Ichneumonidae (Hymenoptera) of the Ryukyu Archipelago. Pacific Insects 12(2): 327–399.
- Schmiedeknecht O (1907) Hymenopteren Mitteleuropas. Ichneumonidae. Gustav Fischer. Jena, 804 pp.
- Smits van Burgst CAL (1913) Tunesian Hymenoptera. Den Haag, 37 pp.
- Uchida T (1956) Die Ichneumoniden aus der Amami Inselgruppe. Insecta Matsumurana 19: 82–100.
- Viereck HL (1912) Descriptions of one new family, eight new genera, and thirty-three new species of Ichneumonidae. Proceedings of the United States National Museum 43: 575–593. doi: 10.5479/si.00963801.1942.575
- Yu DS, Van Achterberg, C, Horstmann K (2012) Taxapad 2012, Ichneumonoidea 2011. Database on flash-drive. www.taxapad.com, Ottawa, Ontario, Canada.

Description of *Nicrophorus efferens*, new species, from Bougainville Island (Coleoptera, Silphidae, Nicrophorinae)

Derek S. Sikes^{1,†}, Tonya Mousseau^{2,‡}

1 University of Alaska Museum, 907 Yukon Drive, Fairbanks, AK 99775-6960, USA **2** Mount Royal University, 4825 Mount Royal Gate Southwest, Calgary, AB T3E 6K6, CANADA

† [urn:lsid:zoobank.org:author:1448D556-54AB-4517-85C0-D76B7070027E](https://zoobank.org/urn:lsid:zoobank.org:author:1448D556-54AB-4517-85C0-D76B7070027E)

‡ [urn:lsid:zoobank.org:author:55DCF6A8-6E11-4046-85AB-45ACAF8DDA1B](https://zoobank.org/urn:lsid:zoobank.org:author:55DCF6A8-6E11-4046-85AB-45ACAF8DDA1B)

Corresponding author: Derek S. Sikes (dssikes@alaska.edu)

Academic editor: Michael Ivie | Received 1 March 2013 | Accepted 29 May 2013 | Published 20 June 2013

[urn:lsid:zoobank.org:pub:A6CA388D-1ECF-4D85-BFF6-F4EA3B7A7C6B](https://zoobank.org/urn:lsid:zoobank.org:pub:A6CA388D-1ECF-4D85-BFF6-F4EA3B7A7C6B)

Citation: Sikes DS, Mousseau T (2013) Description of *Nicrophorus efferens*, new species, from Bougainville Island (Coleoptera, Silphidae, Nicrophorinae). ZooKeys 311: 83–93. doi: 10.3897/zookeys.311.5141

Abstract

A new species of *Nicrophorus* in the *nepalensis* species-group, *Nicrophorus efferens* Sikes and Mousseau, is described from Bougainville Island in the Solomon Islands archipelago. It is distinguished from the known species of the genus *Nicrophorus* and its likely closest relative, *Nicrophorus reticulatus* Sikes and Madge, based on external morphology. A comparison among the four *Nicrophorus* species known from the Solomon Island archipelago and Papua New Guinea is presented.

Keywords

Nicrophorus, Silphidae, burying beetles, taxonomy, Solomon Islands, Papua New Guinea

Introduction

The *nepalensis* species group (Coleoptera, *Nicrophorus*) is the second largest species group within the genus *Nicrophorus*. Species of this group are primarily montane in regions of eastern Asia and the Malay Archipelago, ranging in longitude from 73°E

(Pakistan) to 160°E (Guadalcanal) and latitude from 51°N (Ussuri, Russia) to 9°48'S (Papua New Guinea) (Sikes et al. 2006). In the first descriptions of the species group, Portevin (1926), who originally aligned its members, and Hatch (1927), who formally recognized the species group, included five species: *Nicrophorus nepalensis* Hope, *Nicrophorus podagricus* Portevin, *Nicrophorus heurni* Portevin, *Nicrophorus quadripunctatus* Kraatz, and *Nicrophorus maculifrons* Kraatz. Růžička et al. (2000) added two more species to the group, *Nicrophorus montivagus* Lewis and *Nicrophorus sausai* Růžička, Háva, and Schneider (Růžička et al. 2000). Sikes (2003) removed *N. sausai*, but added *Nicrophorus apo* Arnett and *Nicrophorus insularis* Grouvelle. The most recent taxonomic and phylogenetic work on the *nepalensis* species group is that by Sikes et al. (2006) who included seven newly described species, bringing the total number of species in the group to 15.

Only two species of *Nicrophorus* were known from the Solomon Islands archipelago prior to this study: *N. reticulatus* Sikes and Madge and *N. kieticus* Mroczkowski (Sikes et al. 2006). In this work, a third *Nicrophorus* species for the region, from Bougainville Island, Papua New Guinea, is described. This new species is described within the context of an ongoing revision of the subfamily (see Sikes and Peck 2000, Sikes 2003, Sikes 2005, Sikes et al. 2002, 2006, 2008, Mousseau and Sikes 2011).

Methods

The 6 adult specimens comprising the type series were borrowed from the Bernice Pauahi Bishop Museum of Hawaii (BPBM). One paratype from this series will be deposited in the BMNH. These were compared with specimens from the following collections: FMNH – Field Museum of Natural History, Chicago, IL, USA; DSSC – Derek S. Sikes Collection, University of Alaska Fairbanks, Fairbanks, AK, USA; BMNH – Department of Entomology, The Natural History Museum, Cromwell Rd, London, UK. Four of the type specimens were broken with parts associated on separate pins. All *nepalensis*-group characters were coded independently by both authors and any differences of opinion were discussed to reach a consensus. Morphological data were managed using MacClade 4.04 OSX (Maddison and Maddison 2002) in one NEXUS file. The description was prepared by editing the output generated by use of the ‘export descriptions’ menu option of MacClade. Characters selected are relevant to the subfamily Nicrophorinae so include some characters invariant within the subfamily (synapomorphies of the subfamily) and invariant within the *nepalensis* species group. Also included are absence character states (for example “Frons black, without orange spot” – because an orange spot is common for species in this species group). Specimen data were managed in MANTIS (Naskrecki 2001). Google Earth build 6.1.0.5001 was used to georeference the type locality. The map was prepared using SimpleMappr (Shorthouse 2010). A red determination label bearing a unique alphanumeric code was placed on the pin of each specimen examined. These codes are listed in the material examined data to provide unambiguous association of speci-

mens with data. Habitus photos of *N. reticulatus* were captured using a Nikon D100 and a 60mm Nikkor lens at f22, using incandescent and fiber optic lights with a tripod. Beetles were positioned in front of an 18% grey photography card to simplify the exposure. Images of *N. efferens* were captured using a Leica DFC425 camera mounted on a Leica MZ16 stereomicroscope in combination with Leica Application Suite © software v.3.8.0; after the specimen had been washed in warm ddH₂O and Ammonium Hydroxide prior to a ~24h soak in acetone to remove grime and oils thereby brightening the elytral pattern. Images were edited using Adobe Photoshop v.7 to remove the background, stack images of multiple focal planes, and lighten the images. Observations were made with a Leica MZ16 stereomicroscope (7.1x-115× magnification, 1x planapochromatic objective/10x eyepieces, max resolution 420 Lp/mm, Leica Microsystems (Switzerland) Ltd.). Measurements were made using an ocular micrometer in the MZ16 scope. The scanning electron micrograph of the *N. reticulatus* elytron was taken with an Environmental SEM (system 2020, version 3.53, FEI company 1999), which, unlike standard SEM, does not require gold-coating of specimens and thus is ideal for imaging type specimens. The SEM image of *N. efferens* was taken with a Zeiss (LEO) EVO 60 using variable pressure as an alternative to high vacuum and gold coating.

Data resources

The data underpinning the analyses reported in this paper are deposited at GBIF, the Global Biodiversity Information Facility, http://ipt.pensoft.net/ipt/resource.do?r=type_specimen_data_for_new_species_nicrophorus_efferens.

Taxonomy

Nicrophorus efferens Sikes & Mousseau, sp. n.

urn:lsid:zoobank.org:act:7B4F8F2A-CC50-407A-BD15-CEC84EB706F6

http://species-id.net/wiki/Nicrophorus_efferens

Figs 1, 2A,B, 3, 4

Holotype. Male (in BPBM), here designated, labeled “Bougainville: NE Mutahi. 700m 18 km S. E. Tinputz” / “8–14. III. 1968” / “Tawi Collector BISHOP” / “*Nicrophorus kieticus* Mroczkowski [sic] det. S.B. Peck. 1993” / “HOLOTYPE *Nicrophorus efferens* Sikes & Mousseau 2013 BPBM124189Nic” [red paper].

Paratypes. 5 Specimens. **2 Females**, labeled “Bougainville: NE Mutahi. 700m 18 km S. E. Tinputz” / “15–21. III. 1968” / “Tawi Collector BISHOP” / “*Nicrophorus kieticus* Mroczkowski [sic] det. S.B. Peck. 1993” / “PARATYPE *Nicrophorus efferens* Sikes & Mousseau 2013 BPBM124190Nic” [red paper], this specimen’s head is mounted on a card below the body (deposited in London, BMNH), BPBM-

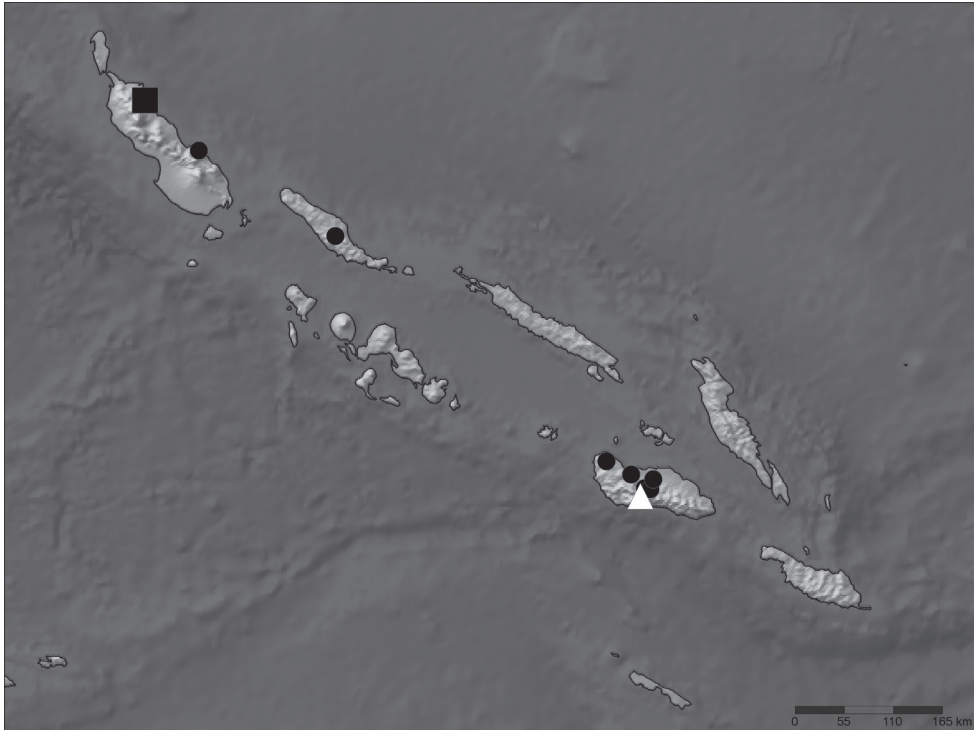


Figure 1. All georeferenced records for *N. effereus* (black square), *N. kieticus* (black circles) and *N. reticulatus* (white triangle) in the Solomon Islands archipelago.

124191Nic, this specimen's abdomen (genitalia missing) is mounted on a card on a separate pin and its left elytron is mounted below the body on a card. **1 Female**, labeled "Bougainville: NE Mutahi. 700m 18 km S. E. Tinputz" / "8–14. III. 1968" / "Tawi Collector BISHOP" / "*Nicrophorus kieticus* Mroczkowski [sic] det. S.B. Peck. 1993" / "PARATYPE *Nicrophorus effereus* Sikes & Mousseau 2013 BPBM124192Nic" [red paper], phoretic mites removed from body into genitalia vial on pin below body. **1 Female**, labeled "Bougainville: NE Mutahi. 700m 18 km S. E. Tinputz" / "1–7. III. 1968" / "& R. Straatman Collectors BISHOP MUSEUM" / "*Nicrophorus kieticus* Mroczkowski [sic] det. S.B. Peck. 1993" / "PARATYPE *Nicrophorus effereus* Sikes & Mousseau 2013 BPBM124193Nic" [red paper], this specimen's prothorax and head are mounted on a card below the body. **1 Male**, labeled "Bougainville: NE Mutahi. 700m 18 km S. E. Tinputz" / "1–7. III. 1968" / "Tawi Collector BISHOP" / "*Nicrophorus kieticus* Mroczkowski [sic] det. S.B. Peck. 1993" / "PARATYPE *Nicrophorus effereus* Sikes & Mousseau 2013 BPBM124194Nic" [red paper], specimen broken and mounted on two pins: hind legs and pygidium on card below body, on pin 2 genitalia in vial with glycerin, head and prothorax on card.

Type locality. [Papua New Guinea]: Solomon Islands [Archipelago]: Bougainville: NE Mutahi, SE Tinputz, 700m, [$\sim 5.716^{\circ}\text{S}$, 155.086°E +/- 2 km, WGS84] (Fig. 1).

Measurements. (2 males, 4 females), pronotal width: male 5.02 – 5.87, 5.45 ± 0.6 mm, female 5.34 – 6.42, 5.82 ± 0.45 mm.

Diagnosis. *Nicrophorus* with center of abdominal sternite 3 between metacoxae glabrous, with long erect brown setae lateral of center hidden above and exposed posterior of coxae (shared with only four *Nicrophorus* species: *N. distinctus* Grouvelle, *N. apo* Arnett, *N. heurni* Portevin, and *N. reticulatus* Sikes & Madge); epipleura black or black throughout with some dark orange in the posterior dorsal portion (of these four, shared with only *N. reticulatus*); elytral disc microsculpturing transverse, straight, narrow with breaks (Fig. 2A, B).

Description. Eight character states not shared with *N. reticulatus*, the likely closest relative of *N. efferens*, are indicated below by use of italics.

Head. Mandibles asymmetrical with dent receptive notch on right mandible. Mandibles preapically wide, short, stout; with dorsal, ridged groove. Maxillary palpus not elongate, concealed beneath mandible when mandibles open. Segment 1 of labial palpus as long as segment 2. Labrum with bilateral pair of elongate setae in clusters. Clypeal membrane orange, yellow or brown, not black. Female clypeal membrane fasciform, male clypeal membrane campanulate and produced posteriorly and laterally enclosed by clypeus. Gula bar of large males reduced to thin strip, replaced by membrane, not contiguous with submentum. Frontoclypeal (epistomal) suture present. Gular sutures confluent, reducing gula to small triangle posterior of gula bar. Antennomeres 2 and 3 fused making an 11 segmented antenna appear 10 segmented. Antennal scape with posterior face flattened (fitting against eyes). Ridge surrounding antennal socket forming distinctly ridged lateral edge of clypeus. Antennal club abrupt, large; basal segment black, apical 3 orange. Posterior of antennal club segments unlike anterior, with large raised ridge, with sharp lateral edges. Basal antennomere of club oval, transverse, not circular. Antennal club segment joints at edge of segments. Antennomeres of club weakly emarginate. Frons black, without orange spot. Epicranial sulci (grooves along inner margin of eyes) long, reaching posterior of eyes and usually joining. Postocular bulge present. Lateral margins of head of large males parallel, or subparallel, giving rear of head a square appearance. Width across postocular bulge of large males less than width across eyes. Post-ocular bulge of large males larger than females. Posterior margin of eye of large males in lateral view sinuate. Width of eyes of large males wide (width greater than or equal to half length). Dorsum of neck with two nonpunctate band(s). Microsculpture of frons absent (smooth, polished).

Thorax. Microsculpture of pronotum disc isodiametric. Pronotum of large males orbicular. Anterior margin of pronotum straight or weakly concave. Pronotal anterior impressions complete, with distinct inner arcs. *Setae on posteroventral portion of hypomeron long, erect, sparse; filling region extending a third or less length between trochantin and posterior margin, restricted to upper region of sclerite.* Anterior corner of hypomeron glabrous. Triangular depression at midpoint posterior margin of pronotum absent. Pronotum disk black. Posterior of pronotum with flat margin or border. *Setae present on pronotum, brown to light-brown, short (ca. < 0.1mm), sparse, and restricted to margins.* Pair of 'v'-shaped bumps present on posterior of pronotum. Medial groove

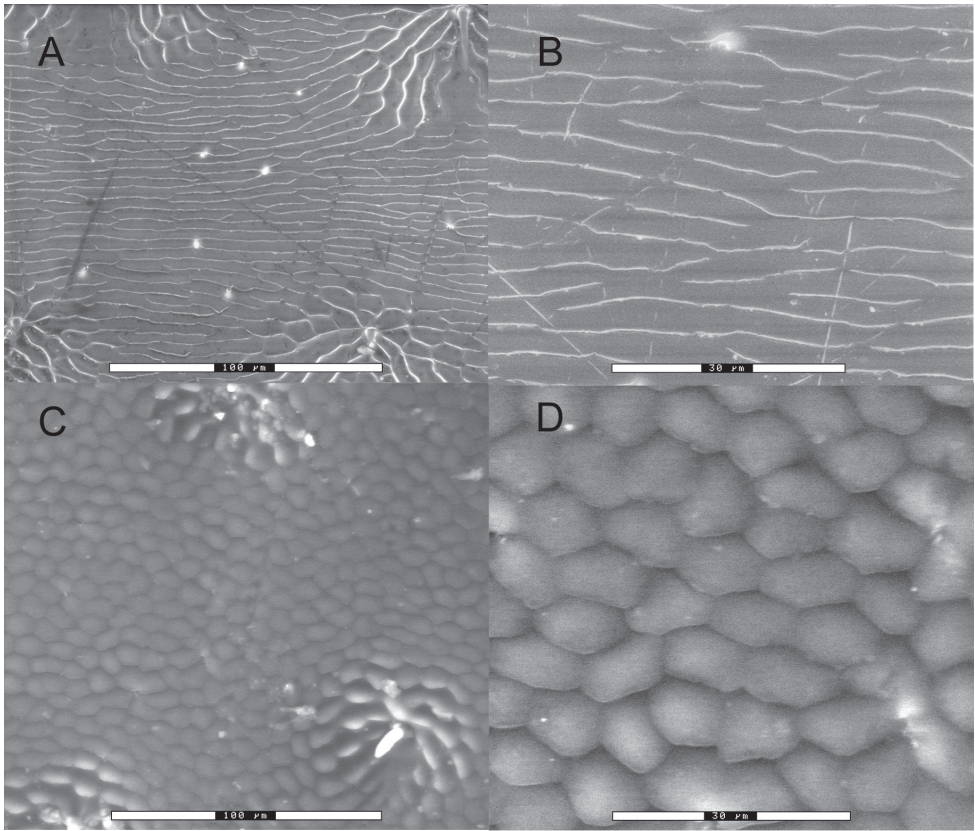


Figure 2. Scanning electron micrographs of elytral microsculpture. **A** *Nicrophorus efferens* paratype BPBM124191Nic, 500 \times , scale bar is 100 μ m **B** *N. efferens* 1500 \times , scale bar is 30 μ m **C** *Nicrophorus reticulatus* 500 \times scale bar is 100 μ m **D** *N. reticulatus* 1500 \times scale bar is 30 μ m.

of pronotum present. Antemesosternal sclerite cordiform and bilobate. Elytral locking device under apex of scutellum composed of single channel, without distinct, thin, raised median ridge. Humerus immediately dorsal of epipleural ridge black. Humeral setae present, short and not forming row. Epipleuron entirely black or black throughout with some dark orange in the posterior dorsal portion. Posterior epipleural ridge without isolated single-file row of contiguous preapical setae. Epipleural ridge distinct and short, to tip of scutellum. Epipleuron glabrous, or with very sparse, extremely small setae (ca. the size of a puncture). Lateral margin of elytron glabrous, or with very sparse, extremely small setae (ca. the size of a puncture). Elytral surface without long setae. Elytra bicolored and bifasciate. Elytron with costae not distinctly raised but visible to naked eye. Region adjacent to apex of elytral suture black. Profile of elytra in lateral view raised posteriorly. Anterior fascia of elytron without black spot, small, triangular-square, just reaching 2nd costa. Posterior fascia between costae, anterior margin u-shaped between costae 1 and 2 (with bottom of u towards posterior). Posterior fascia not touching lateral or posterior margins of elytron. Elytral posterior

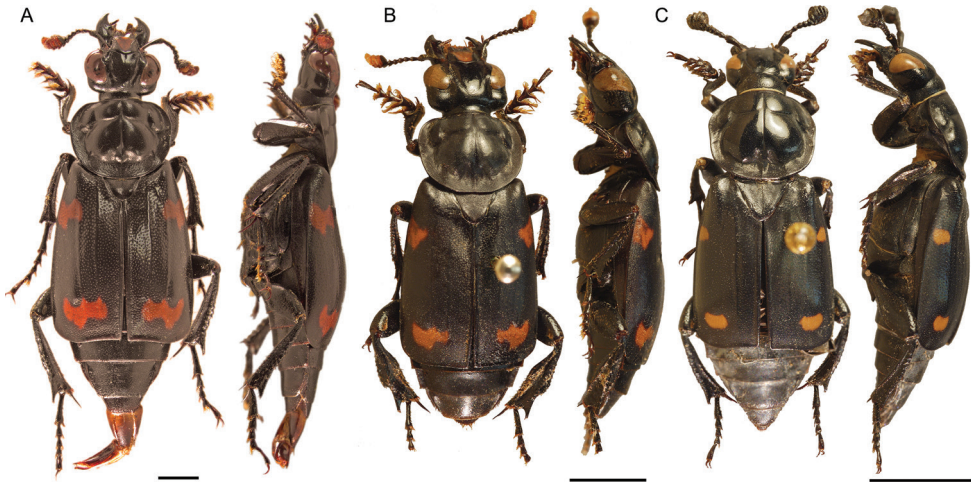


Figure 3. Dorsal and lateral habitus of adult males (at different scales). **A** small male (pronotal width 5.02 mm) with aedeagus everted, *N. efferens* (holotype BPBM124189Nic), scale bar is 2 mm, fifth protarsomeres are missing **B** large male (pronotal width 6.6 mm) *N. reticulatus* (paratype BMNH000826), scale bar is 5 mm **C** small male (pronotal width 5.19 mm) *N. kieticus* (BMNH000809Nic) scale bar is 5 mm.

fascia without black spot. Elytral posterior fascia greatly reduced, far from suture, jagged. *Elytral microsculpture* transverse straight, narrow with breaks. Elytral color variation – greatly increased orange and all black forms unknown. Ventral surface of elytron with golden sheen. Row of setae facing inward or downward on inner lateral margin of elytra in posterior quarter along lateral ridge of elytron in a distinct, single file row. *Posterior margin of elytron* with 5–7 clusters of very short, dark setae. Flange along mesepisternal anterior margin tapering gradually towards mesosternum. *Metanotal subalare* with sharply rising, internally margined ridge along anteriomedial edge, anteriomedial ridge narrow leaving a wide depression. Metepimeron constricted. Metepisternum with upper third impunctate and glabrous. Metasternal pubescence medially (between mesocoxae) and laterally, long, dark brown. Metasternum with long setae, bald patch posterior of mesocoxae absent. Metepimeral posterior lobe with sparse, short brown setae throughout lobe, or glabrous. Metasternum posterior margin edge glabrous. Furcal arms of metendosternal apophysis directed posteriorly (forming acute angle with main stalk). Laminae of metendosternal apophysis joining furcal arms to anterior projection of stem. Ventral laminae of metendosternal apophysis reaching from anterior projection to bend in main stalk (anterior 3 fourths of entire length). Radial hinge of wing notched. Apex of pterostigma of wing truncate-rectangular.

Abdomen. Pair of stridulatory files on tergite 5 present, files parallel, separated by 2 or more file widths, touching posterior margin of tergite. Stridulatory file scraper on venter of elytron near elytral apex (< 0.2mm). First abdominal spiracle slit short, less than one third length of spiracle, in straight line with center line of spiracle. First abdominal spiracle lobed, with small bulb projecting anteriorly. Posterior margins of abdominal segments 3–6 with short setae (2–3 times longer than the distance between

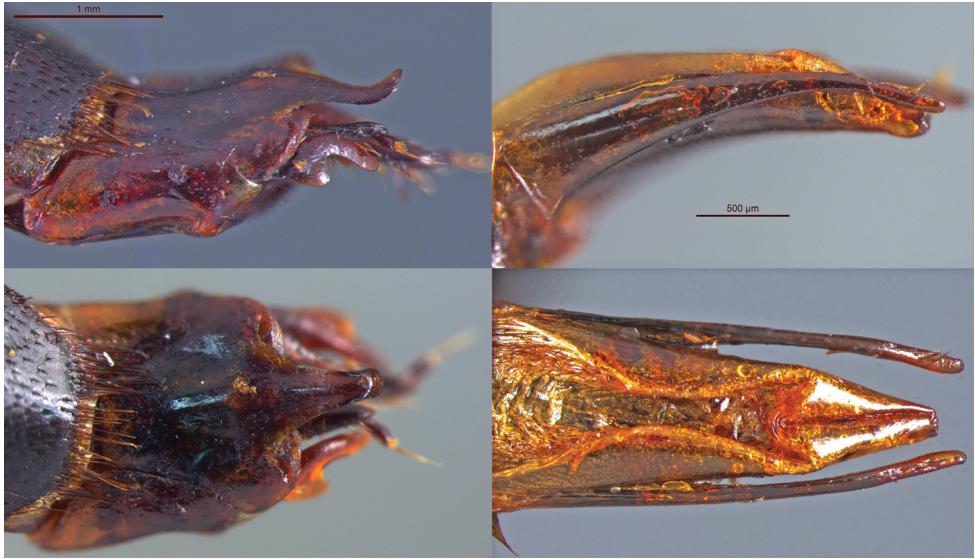


Figure 4. Genitalia. Female ovipositor (paratype BPBM124190Nic), **A** lateral **B** dorsal. Scale bar is 1 mm. Male aedeagus (holotype BPBM124189Nic) **C** lateral **D** dorsal. Scale bar is 500 µm.

adjacent setae). Center of sternite 3 between metacoxae glabrous, but with long erect setae lateral of center under coxae. Spiracle of tergite six elongate, slit-like, and parallel to lateral margin of tergite, or forming less than 30-degree angle with lateral margin. Tergite seven with short depressed setae. *Abdominal setae* dark brown. *Setae of tergite 9 of males (pygidium)* brown.

Legs. Venter of protibia apex with lateral process. Anterior face of protrochantin with regions of dense pubescence composed of short, recumbant golden setae. Anterior of procoxae with short setae on basal half. Lateral margin of anterior of procoxae smooth, without ridge or bump. Meso- and metatibia apical angle produced into lobe. Outer margin of mesotibia straight or curved outwards. Inner margin of metatibia of large males gradually curved outwards. Inner margin of metatibia with inner face not forming wide channel entirely filled with dense long setae. Middle of outer margin of metatibia slightly swollen in large males. Middle of inner face of metatibia of large males greatly widened (2.5 or greater than width at base). Metatibia straight. Inner face of apex of metafemora with circular or oval cluster of short setae occupying 1/4 to 1/2 of apical region. Inner face of base of metafemora with single ridge separating medial setose area (area not depressed) from lateral half. Female metafemora slender (length ≥ 2.5 times greatest width). Posterior/ventral face of metafemora flattened with ridged edges. Metatrochanter spine of males short and subapical, apex pointing parallel (or almost parallel) to leg, straight, not recurved dorsally. Metacoxae wider than long. Metacoxae with anterior line complete for half or more of metacoxa. Posterior margin of metacoxae without white microsetae. Tarsal empodium bisetose. Venter of metatarsomere 1 glabrous medially, especially distally, leaving narrow glabrous channel, with dense cluster of long setae at apex.

Ovipositor. (Fig. 4A, B) Valvifer with claw. Proctiger (T10) apex strongly lobed, spatulate, not bifurcate. Spatula on proctiger (T10) apex narrow (ca. equal to half the width across widest region of metatarsomere 5). Proctiger (T10) lobe with pubescent apex, ventral setae. Proctiger (T10) venter of spatula apex smooth, lacking medial keel-like ridge. Gonocoxite terminal claw absent. Dorsal ridge on proctiger (T10) absent. Paraproct (T9) apex glabrous, with well-defined, raised ridge. Paraproct without process. Valvifer claw glabrous, dentate; dentition composed of small rounded lobe situated in the middle to distal third of the valvifer.

Aedeagus. (Fig. 4C, D). Paramere apex rounded, with setae laterally not reaching the apical curve. Paramere ventral setal patch not composed of 5 evenly spaced setae of identical length and not overlapping with apicolateral patch. Without 3rd paramere setal patch or paramere flange. Parameres not constricted behind apex. Parameres evenly curved, tapering towards apex.

Variation. The holotype male and the largest female paratype has an epipleuron mostly black throughout, but with some dark orange faintly visible in the posterior dorsal portion. Two of the type specimens have no setae along the posterior margin of the elytra, presumably they have been lost to abrasion.

Geographic Distribution. This species is only known from the type locality (Fig. 1).

Etymology. From the latin verb '*effero*' to carry out for burial, bear to the grave, bury; present participle in the nominative singular.

Discussion

This new species is likely closely related to one if not both of the *Nicrophorus* previously known from the Solomon Islands archipelago (*N. reticulatus* [Figs 2, 3] and *N. kieticus*, [Figs 1,3]) and may be the key to untangling the mystery of how they are related. Two of its three diagnostic characters are likely synapomorphies between it and *N. reticulatus*. Although the holotype and one paratype of the four known specimens of *N. reticulatus* have entirely black epipleura, the two remaining paratypes have epipleura mostly black throughout, but with a small region of dark orange in the posterior dorsal portion. This weakly present orange region is present on two of the *N. efferens* type specimens. These two species, along with *N. kieticus*, also share greatly shortened setae along the posterior margins of abdominal sterna 3 to 6 and four highly reduced and similar looking fasciae on the elytra (Fig. 3). However, *N. efferens* differs strongly from *N. kieticus* in its possession of most synapomorphies of the *nepalensis* group (Sikes et al. 2006) including orange antennal clubs and an epipleural ridge no longer than the tip of the scutellum. *Nicrophorus kieticus* has black antennal clubs and a longer epipleural ridge, among a total of 12 character states we have found that *N. kieticus* does not share with members of the *nepalensis* group. Of 185 morphological characters coded for the subfamily, *N. efferens* shares 95.7% of its character states with *N. reticulatus*, 92.4% with *N. heurni*, a species restricted to the island of New Guinea, and 84.8% with *N. kieticus*. The primary character that differentiates *N. efferens* from *N. reticulatus* is that the

former has elytral microsculpturing typical of the *nepalensis* group (Fig. 2A, B) while the latter is the only species of the group with isodiametric microsculpturing (Fig. 2C, D). In the keys of Sikes et al. (2006) *N. efferens* would key out as *N. reticulatus* or *N. heurni* so an additional couplet is necessary to key out this new species. Given the rarity of these species in collections it would not be surprising if more new *Nicrophorus* species await discovery at higher elevations of the Melanesian and Malay Archipelago islands.

- 1 Elytra with epipleuron partially or entirely black (Fig. 3A, B) **2**
- Elytra with epipleuron entirely red–orange **4**
- 2 Elytral disc with strongly transverse microsculpturing (Fig. 2A, B) **2.5**
- Elytral disc with isodiametric microsculpturing (Fig. 2C, D) ... ***N. reticulatus***
- 2.5 Elytra with epipleuron black with entire medial (dorsal) half orange **3**
- Elytra with epipleuron entirely black (or black with some dark orange in the posterior dorsal portion) ***N. efferens***

Acknowledgments

Thank you to Shepherd Myers (BPBM) for loaning the *N. efferens* specimens studied and to Margaret Thayer (FMNH) and Max Barclay (BMNH) for making comparative material available. Thank you to Fred Falcao for the second author's visit to Hawaii. Thanks to lab manager and SEM technician Betty Strack at the FMNH for help with obtaining the *N. efferens* SEM images. We also thank our long time friend and collaborator, Ronald Madge, with whom we share a passion for silphid systematics, for his input on this new species.

References

- Hatch MH (1927) Studies on the Silphinae. Journal of the New York Entomological Society, 35: 331–370.
- Maddison DR, Maddison WP (2002) MacClade 4: Analysis of phylogeny and character evolution. Version 4.04. [<http://macclade.org>]
- Mousseau T, Sikes DS (2011). Almost but not quite a subspecies: A case of genetic but not morphological diagnosability in *Nicrophorus* (Coleoptera: Silphidae). The Biological Journal of the Linnean Society. 102: 311–333. doi: 10.1111/j.1095-8312.2010.01568.x
- Naskrecki P (2001) 'MANTIS: A Manager of Taxonomic Information and Specimens. File-MakerPro Database.' [<http://insects.oeb.harvard.edu/mantis/>]
- Portevin G (1926) *Les Grands Nécrophages du Globe. Silphini - Necrodini - Nicrophorini*. Encyclopédie Entomologique (A), Vol. 6, Lechevalier, Paris, 270 pp.
- Růžička J, Háva J, Schneider J (2000) Taxonomical and distributional notes on Oriental Silphidae, with description of *Nicrophorus sausai* sp. n. (Insecta: Coleoptera) Reichenbachia 33(2): 377–384.

- Shorthouse DP (2010) SimpleMapppr, an online tool to produce publication-quality point maps. [Retrieved from <http://www.simplemapppr.net>. Accessed 25 February, 13].
- Sikes DS (2003) Systematic Revision of the Subfamily Nicrophorinae (Coleoptera: Silphidae). Ph.D. Dissertation, University of Connecticut, xxiv + 938 pp.
- Sikes DS (2005) Silphidae. In: Beutel RG, Leschen RAB (Eds) Handbook of Zoology, Volume IV Arthropoda: Insecta Part 38 (Kristensen NP & Beutel RG Eds), Coleoptera, Beetles Volume I: Morphology and Systematics (Archostemmata, Adephaga, Myxophaga, Polyphaga partim). Waler de Gruyter, Berlin, New York, 288–296.
- Sikes DS, Peck SB (2000) Description of *Nicrophorus hispaniola*, new species, from Hispaniola (Coleoptera: Silphidae) and a key to the species of *Nicrophorus* of the New World. Annals of the Entomological Society of America. 93:391–397. doi: 10.1603/0013-8746(2000)093[0391:DONHNS]2.0.CO;2
- Sikes DS, Madge RB, Newton AF (2002) A catalog of the Nicrophorinae (Coleoptera: Silphidae) of the world. Zootaxa 65: 1–304.
- Sikes DS, Madge RB, Trumbo ST (2006) Revision of *Nicrophorus* in part: new species and inferred phylogeny of the *nepalensis*-group based on evidence from morphology and mitochondrial DNA (Coleoptera: Silphidae: Nicrophorinae). Invertebrate Systematics. 20: 305–365. doi: 10.1071/IS05020
- Sikes DS, Vamosi SM, Trumbo ST, Ricketts M, Venables C (2008) Molecular systematics and biogeography of *Nicrophorus* in part – The *investigator* species group (Coleoptera: Silphidae) using mixture model MCMC. Molecular Phylogenetics and Evolution 48: 646–666. doi: 10.1016/j.ympev.2008.04.034

