



Rowlandius dumitrescoae species group: new diagnosis, key and description of new cave-dwelling species from Brazil (Schizomida, Hubbardiidae)

Alessandro Ponce de Leão Giupponi¹, Gustavo Silva de Miranda², Osvaldo M. Villarreal³

l Laboratório de Referência Nacional em Vetores das Riquetsioses, LIRN-IOC-FIOCRUZ, Manguinhos, 21040-360, Rio de Janeiro, RJ, Brazil 2 Center for Macroecology, Evolution and Climate, Natural History Museum of Denmark (Zoological Museum), University of Copenhagen, Universitetsparken 15, Copenhagen, Denmark, 2100 3 Departamento de Invertebrados, Pós-Graduação em Zoologia, Museu Nacional, Universidade Federal do Rio de Janeiro, Quinta da Boa Vista, São Cristóvão, 20.940-040, Rio de Janeiro, RJ, Brazil

Corresponding author: Alessandro Ponce de Leão Giupponi (agiupponi@gmail.com)

Academic editor: P. Stoev | Received 25 May 2016 | Accepted 20 October 2016 | Published 16 November 2016

http://zoobank.org/C384D6F4-13F3-4AD8-BFC9-845957671FED

Citation: Giupponi APL, Miranda GS, Villarreal OM (2016) *Rowlandius dumitrescoae* species group: new diagnosis, key and description of new cave-dwelling species from Brazil (Schizomida, Hubbardiidae). ZooKeys 632: 13–34. doi: 10.3897/zookeys.632.9337

Abstract

The *Rowlandius dumitrescoae* species group is reviewed and rediagnosed, and its composition is revised. The group now includes *R. cousinensis*, *R. decui*, *R. dumitrescoae*, *R. insignis*, *R. linsduarte*, *R. monensis*, *R. peckorum*, *R. potiguar*, *R. sul*, *R. ubajara*, and *R. pedrosoi* **sp. n.** A new species is described from a cave in northeast Brazil (Santa Quitéria, Ceará). Identification keys and distributional maps are provided for the species of the group. Sixteen species of Schizomida, including five of *Rowlandius*, are currently recognized from Brazil.

Keywords

Diversity, Hubbardiinae, Neotropics, Schizomids, Short-tailed whipscorpion, taxonomy

Introduction

Rowlandius Reddell & Cokendolpher, 1995, is the most diverse Neotropical genus of Schizomida with 63 described species (Reddell and Cokendolpher 1995; Teruel 2012; Teruel et al. 2012; Delgado-Santa and Armas 2013; Santos et al. 2013). Reddell and Cokendolpher (1995) proposed the genus with a broad concept, using characters that could also fit other genera, and was redefined by Teruel (2004). Reddell and Cokendolpher (1995) assumed Rowlandius as monophyletic, but this has never been tested in a phylogenetic analysis; on the contrary, the presence of several variable characters within the genus (e.g., the number of setae on the propeltidium and the shape of the spermathecae) indicates the opposite (Teruel et al. 2012). Within Rowlandius, the biconourus species group was also proposed as monophyletic, but this hypothesis has not been tested either (Teruel et al. 2012).

The species that compose *Rowlandius* were recognized as a species group long before the genus was erected, when almost all species of Schizomida were placed in *Schizomus* Cook, 1899 (the historical "trash can" of the order). The first attempt to subdivide *Schizomus* into species groups was made by Rowland and Reddell (1979a) who proposed seven; one of them, the *dumitrescoae* group, was divided in three complexes: *dumitrescoae*, *primibiconourus* and *viridis* complex. All *Schizomus* species of these complexes were transferred to *Rowlandius* by Reddell and Cokendolpher (1995). Later, new endeavors to detect and define groups within *Rowlandius* were made by Armas (2002), Teruel (2012) and Teruel et al. (2012), but these included only Cuban species and did not cover all morphological variation within the genus.

Almost 80% of *Rowlandius* species with a known male have striking secondary sexual dimorphism, i.e., the male pedipalp segments are much longer than that of the conspecific females. An interesting case of dimorphism is present in *R. gracilis* Teruel, 2004 and *R. potiguar* Santos, Ferreira & Buzzato, 2013, where the same population has both heteromorphic males with long pedipalp articles and homeomorphic males with shorter, female-sized pedipalp articles (Teruel 2004; Teruel et al. 2012; Santos et al. 2013; Oliveira and Ferreira 2014).

Rowlandius has an extensive geographic distribution, occurring from Cuba to Brazil. A major radiation of the genus seems to have occurred in the Greater Antilles, where the vast majority of the known species are found (Harvey 2003). In contrast, only five species have been described so far from continental South America (R. arduus Armas, Villarreal & Colmenares-García, 2009, R. linsduarte Santos, Dias, Brescovit & Santos, 2008, R. potiguar Santos, Ferreira & Buzzato, 2013, R. sul Cokendolpher & Reddell, 2000 and R. ubajara Santos, Ferreira & Buzzato, 2013). The genus has been recorded from different biomes, including the Brazilian Amazonia, the Brazilian Atlantic forest, and the Venezuelan cloud forest (Santos et al. 2008; Armas et al. 2009). Recently, some species were discovered inhabiting caves or patches of forest inserted in dry areas of Brazil, the Caatinga (Santos et al. 2008; Santos et al. 2013).

In the present article, a new species of *Rowlandius* is described and illustrated from the state of Ceará, northeast Brazil. Additionally, the *Rowlandius dumitrescoae* group is rediagnosed, an identification key to its species is provided, and the relationships of the new species are discussed.

Material and methods

The material studied is deposited in Museu Nacional, Universidade Federal de Rio de Janeiro (MNRJ) and FIOCRUZ, Instituto Oswaldo Cruz (CAVAISC). Terminology of pedipalps, legs and spermathecae follows Reddell and Cokendolpher (1995) and Moreno-González et al. (2014); flagellum setation follows terminology of Harvey (1992), modified by Cokendolpher and Reddell (1992), Villarreal et al. (2014), Moreno-González et al. (2014) and Monjaraz-Ruedas et al. (2016); cheliceral setation terminology is based on Lawrence (1969) modified by Villarreal et al. (2016). Description format follows Villarreal et al. (2016). The terms α - and β -males are used here for the two different sizes of heteromorphs. Those with extremely long palp segments are α -heteromorphic males, and those with palp lengths intermediate between those of females and those of α -males are called β -heteromorphic males.

The keys were built based on the material analyzed and the original descriptions (in the case of species with no specimens available for examination). Males are unknown for *Rowlandius sul* and this species was not included in the male key. The preparation and illustrations of the spermathecae follow Villarreal et al. (2016). Dorsal, ventral, and lateral photos were made with a Leica MZ16 microscope attached to a FujiFilm X10 camera. Pictures of live specimens (courtesy of Denis Rafael Pedroso; Fig. 8) were taken with a Canon Power-Shot SX130 IS. To generate the SEM images, the specimens were critical point dried and mounted on stubs using an adhesive copper aluminum tape. The mounted stubs were then coated with platinum-palladium and scanned with a JEOL JSM-6390 LV.

Acronyms used:

AMN anterior median notch of the chitinized arch;

Dm dorso-median setae of abdomen and flagellum;

Dl dorso-lateral setae of the abdomen and flagellum;

LL lateral lobe of spermathecae; ML median lobe of spermathecae;

Msp microsetae patch of the male flagellum;

VI ventro-lateral setae of the abdomen and flagellum.

Additional material examined

Rowlandius ubajara Santos, Ferreira & Buzzato, 2013: Brazil, Ceará, Ubajara, Ubajara National Park, 11–14.i.2013, 3°50'24.42"S 40°54'3.96"W, 869m a.s.l., Carlos

Frankl Sperber, Thiago Gechel Kloss, Fabiene Maria de Jesus and Gabriel de Oliveira Lobregart *leg.* (1 male, MNRJ 4270).

Rowlandius potiguar Santos, Ferreira & Buzzato, 2013: Brazil, Rio Grande do Norte, Martins, 6°5'7.87"S 37°55'6.62"W, 319m a.s.l., C. Fukushima and A. Giupponi leg. (8 females, MNRJ 4269).

Taxonomy

Hubbardiidae Cook, 1899 Hubbardiinae Cook, 1899 *Rowlandius* Reddell & Cokendolpher, 1995

Rowlandius dumistrocae species group

Diagnosis. Male pedipalps of some species sexually dimorphic, with femur and patella extremely elongated, and femur strongly bent proximally (Figs 3D-E, 4A, B). Male flagellum lanceolate (as in R. cousinensis (Rowland & Reddell, 1979), R. dumitrescoae (Rowland & Reddell, 1979), R. insignis (Hansen in Hansen & Sorensen, 1905), R. monensis (Rowland & Reddell, 1979) and R. pedrosoi sp. n.), subquadrate (as in R. linsduarte and R. potiguar) or ovoid (as in R. peckorum (Rowland & Reddell, 1979) and R. ubajara); male flagellum with rounded dorsal projections (with exception of R. dumitrescoae), never surpassing the lateral borders; male flagellum with posterior border surface (between setae Dl3) elevated or flat (more rare). Spermathecae with four lobes, lateral pair long with a curved stalk and a terminal enlarged bulb; median lobes short and digitiform or subconical (Figs 7A, B). Chitinized arch very short (relation width/length = 3.7) with acute lateral tips (R. cousinensis, R. linsduarteae, R. monensis, R. pedrosoi sp. n., R. potiguar and R. ubajara) or rounded lateral tip (R. dumitrescoae, R. insignis, R. peckorum and R. sul); anteromedian notch contacting the posterior branch in some species. Gonopod absent. The species included in this group can be checked in Table 4.

Distribution. Brazil, Costa Rica, Cuba, Jamaica, Martinique (Windward Islands) and Puerto Rico (Fig. 9).

Rowlandius pedrosoi sp. n.

http://zoobank.org/D6088B71-0770-44CD-8283-2CE412AE608C Figures 1–8, Tables 3 and 4

Diagnosis. Large specimens, male body total length 4.01mm, females 3.85mm (chelicerae and flagellum not included). Spermathecae similar to *R. potiguar*, but stalk of LL thicker and curved in the apical third; *R. pedrosoi* sp. n. with stalk of LL and ML with several glandular pores. Lateral tip of chitinized arch "V-shaped", with obtuse angle,

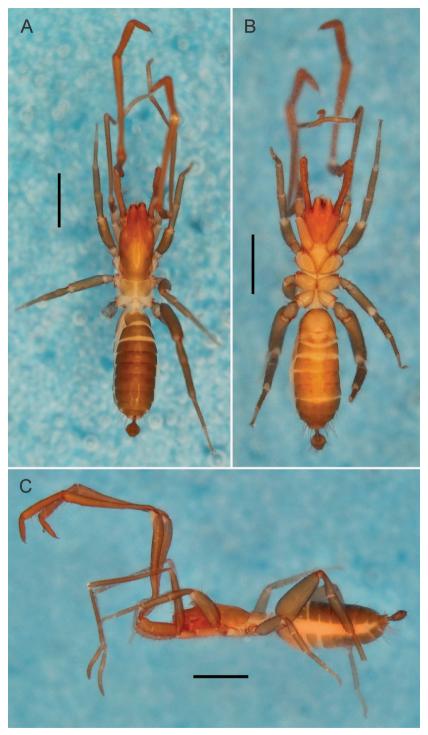


Figure 1. Habitus of an α -heteromorphic male of *Rowlandius pedrosoi* sp. n. (MNRJ 04266). **A** Dorsal view **B** Ventral view **C** Lateral view. Scale bars 1 mm.

greater than 150°, which distinguishes R. pedrosoi sp. n. from R. potiguar and R. linsduarte. Heteromorphic males present, with α (long pedipalps) and β (shorter pedipalps, but longer than those of females) heteromorphics, similar to R. potiguar. Male flagellum with setae Dm1 exactly between the main globose area of the flagellum and the stalk, such as in R. linsduarte and differently from R. potiguar and R. ubajara.

Type material. Holotype: Brazil, *Ceará*, Santa Quitéria, Gruta P-08, 41529 mE / 9495881 mN SAD'69S, 15–21.vii.2014, Pellegatti and Pedroso *leg*. (1 male, MNRJ 04266). **Paratypes:** same data as holotype (1 male, 7 females and 10 juveniles, MNRJ 04267); same data as holotype (1 female and 1 juvenile, CAVAISC-ARAC 0008); same data as holotype, 03–13.ii.2014 (4 females and 8 juveniles, MNRJ 04268).

Etymology. The species name is in honor of arachnologist Denis Rafael Pedroso, friend and collector of the type series (of this and many other new species of arachnids).

Description. *Male holotype.* Color (Fig. 8E–F): live animals with abdominal tergites and sternites olive-brown; pleura white. Pedipalps reddish-brown; legs light brown with the extremities dark-brown. Prosoma light brown; ventral region lighter than the dorsal. Alcohol preserved specimens (Fig. 1) with propeltidium and chelicerae reddish-brown, meso and metapeltidium yellowish-brown (lighter than the chelicerae and propeltidium), legs light brown, abdominal tergites brown and sternites yellowish-brown, flagellum medium-brown. Ventrally coxae I-IV and sternal region yellowish. All body setation light reddish-brown.

Prosoma (Fig. 1). Anterior process of propeltidium with two setae (one behind the other) followed by two pairs of dorsosubmedian transversally oriented setae; eyespot suboval; metapeltidium divided. Anterior sternum with 11+2 setae and posterior sternum with 5 setae. Anterior process as wide as long, with a wide base, narrowing abruptly, forming an almost right triangle; the tip of the process is curved downwards.

Opisthosoma (Fig. 1). Setae: Tergite I with two pairs of anterior microsetae and one pair of large *Dm* setae. Tergite II with three pairs of anterior microsetae parallel to each other, and one large pair of *Dm* setae. Tergites III–IX and XII each with one pair of large *Dm* setae; VIII with small *Dl2*; IX without *Dm*, but pairs *Dl1* and *Dl2* present; X without dorsal setae; XI with *Dl1* and without *Dl2*; XII with short rounded posterodorsal process and with setae *Dl1* and *Dl2*. Abdominal apodemes with coloration identical to the rest of the sternites. Sternites I–II with many scattered microsetae. Sternite III with 22 microsetae. Sternite IV with *Vl2*, *Vl1* and *Vm2* plus four *AS* microsetae. Sternite V with *Vl2*, *Vl1A*, *Vl1B* and *Vm2*, plus six *AS*. Sternite VI with *Vm1*, *Vm2*, *Vl1A*, *Vl1B*, *Vl2*, plus six *AS*. Sternite VII with *Vm2*, *Vl1* (A and B), *Vl2*, six *AS* and without *Vm1*. Sternite VIII with *Vm2*, *Vl1*, *Vl2*, plus six *AS*. Sternite IX with *Vm1*, *Vm2*, *Vl1* and *Vl2* plus one pair of supranumeric setae between *Vl1* and *Vm2*. Sternite XII with six setae plus four microsetae.

Flagellum (Fig. 2). In dorsal view flagellum diamond shaped, as wide as long, with rounded lateral and apical tips; with three bulges: a pair positioned dorsosubmedian (each bulge seated on opposite sides), without setae, separated by a depression, and one bulge in the central distal region (posteromedian), with the setae *Dm4* on its apex;

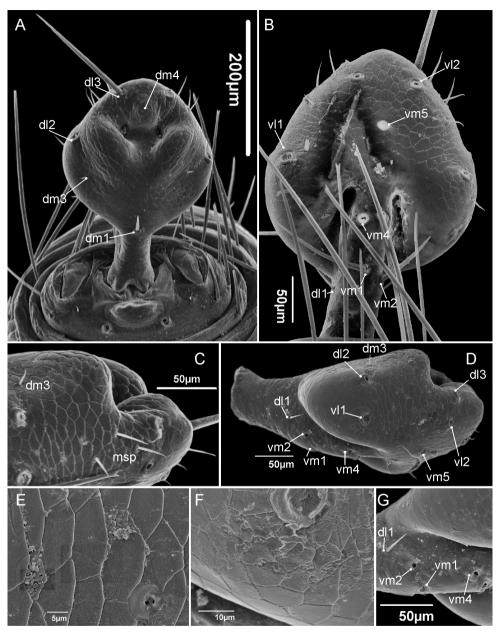


Figure 2. Male flagellum of *Rowlandius pedrosoi* sp. n. (MNRJ 04267). **A** Dorsal view **B** Ventral view **C** Detail in distolateral view **D** Lateral view **E** Uropygi gland opening **F** A set of glands below *VL1* **G** Detail of the position of the proximal ventral and lateral setae.

the central distal bulge is not connected to the lateral ones, with a depression between them. Dm1 is exactly on the edge between the diamond-shaped part and the stalk. Dl3 is positioned distally in relation to Dm4. Ventrally, Vm5 is closer to Vl2 than to Vl1

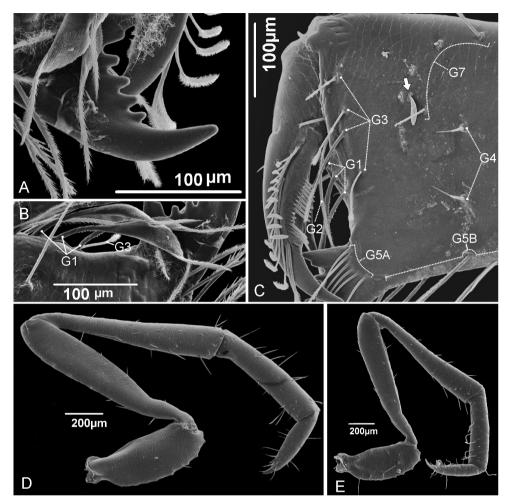


Figure 3. Details of the chelicera and pedipalps of *Rowlandius pedrosoi* sp. n., male (MNRJ 04267). **A** Fixed finger of chelicera **B** Cheliceral setae G1 **C** Mesal view of right chelicera showing setal groups; the arrow indicates the *Basidiobolus* fungus **D** Right pedipalp of a β-heteromorphic, ectal view **E** Left pedipalp of a β-heteromorphic, mesal view.

and *Vm4*. *Vm1* is closer to *Vm4* than to *Vm2*. Three microsetae on the lateral of the flagellum (msp), between the pairs *Dl2/Vl1* and *Dl3/Vl2*, closer to the latter. *Dl1*, *Vl1* and *Vl2* forming a straight line in the frontal axis. *Female flagellum* (Fig. 6A–C) with four flagellomeres (I=II=III>IV), wider between the second and third flagellomeres. Dorsally with a small *Dm1* close to the distal margin of the first flagellomere, placed in the middle line; a pair of larger *Dl1* on the wider portion of flagellum, in the point between the second and third flagellomere; one large *Dm4* in the apical portion of the third flagellomere; a pair of small *Dl4* on the fourth flagellomere in mediolateral position; a pair of large *Dl3* apically on the terminal position of the flagellum. Ventrally with a small basal *Vm1* on the first flagellomere, positioned near the distal border; a pair of median *Vm4* in the second flagellomere; one large medial placed *Vm5* on the

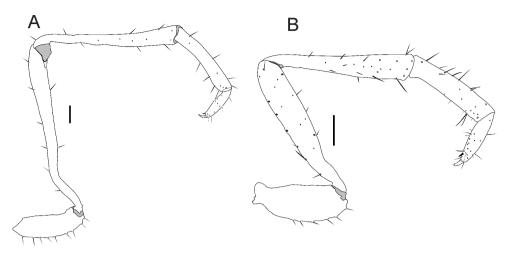


Figure 4. Right pedipalps of heteromorphic males of *Rowlandius pedrosoi* sp. n., ectal view (MNRJ 04267). **A** α -heteromorphic **B** β -heteromorphic. Scale bars 0.2 mm.

third flagellomere; a pair of a large *Vl1* on wider portion of the flagellum, between the second and third flagellomeres; a pair of large *Vl2* on the fourth flagellomeres, apically.

Chelicerae (Fig. 3A–C). Movable finger sharp and curved; serrula with 16 hyaline teeth increasing in size towards distal region; guard tooth rounded. Lamella smooth. Fixed finger with bifid basal tooth, followed by four small subequal teeth; last tooth is the biggest, recurved, with an acute apex, subequal to the basal cusp of bifid tooth. Setation: G1 (setae group 1) with 3 spatulate setae; G2 with 4 feathered setae; G3 with 4 setae, all feathered dorsally and with serrated ventral surfaces; G4 with 2 setae, smooth, short and thick with thin apex; G5A with 6 similar sized feathered setae; G5B with 9 setae larger than G5A; G6 with 1 smooth setae longer than half of movable finger length; G7 with 6 setae decreasing in size from proximal to distal, feathered from the middle to its end. Setal group formula: 3–4–4–2–6–9–1–6.

Pedipalp (Figs 3D–E, 4). All segments without spinose setae. **Trochanter:** subcylindrical in α -heteromorphic males (in lateral view), longer than wide, with apical portion curved upward; short trapezoid in β -heteromorphic males and even shorter in females (Fig. 5); without apical spur (frontal projection); one ventral row of eight large setae with an intermediate row of three small setae. **Femur:** subcylindrical, club-shaped, with distal portion two times wider than the basal part; in α -heteromorphic males the femur is longer than the total length of the prosoma (pro-, meso- and metapeltidium together); in α -heteromorphic males the femur is longer than the patella (in β -heteromorphic males the femur and patella are subequal and in females the patella is longer); with few setae, only one ventral and one dorsal row of setae; on the ectal surface only one apical setae; on the mesal surface, one row of three setae. **Patella:** subcylindrical, club-shaped, with distal portion two times wider than the basal part; more setae than the femur, with two dorsal and two ventral rows, and four setae on the ectal surface. **Tibia:** cylindrical, α -heteromorphic males with distal portion slightly wider; shorter than half the length of



Figure 5. Habitus of a female of *Rowlandius pedrosoi* sp. n. (MNRJ 04267). **A** Dorsal view **B** Ventral view **C** Lateral view. Scale bars 1 mm.

the femur and patella; in β -heteromorphic males and females, the tibia, femur and the patella have similar length. The tibia has the largest number of setae on the pedipalps, with some feather-like setae on the ventral region. **Tarsus:** conical, shorter than the tibia, with lots of setae in the distal third, with two dorsolateral and two ventrolateral

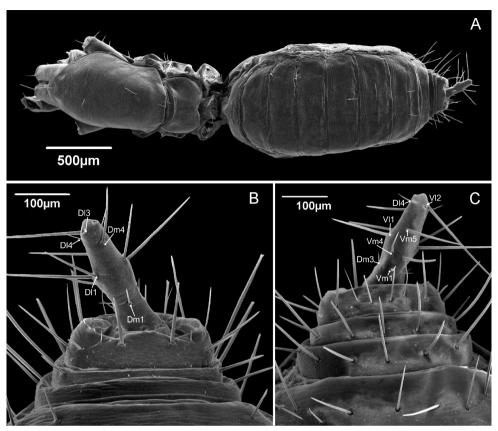


Figure 6. Details of prosoma, opisthosoma and abdomen of a female of *Rowlandius pedrosoi* sp. n. (MNRJ 04267). **A** Dorsal view of prosoma and opisthosoma **B** Dorsal view of female flagellum **C** Ventral view of female flagellum.

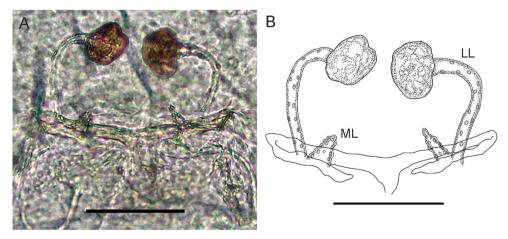


Figure 7. Spermathecae of *Rowlandius pedrosoi* sp. n. (MNRJ 04267). **A** Dorsal view picture **B** Schematic drawing. Scale bars 100 μ m.

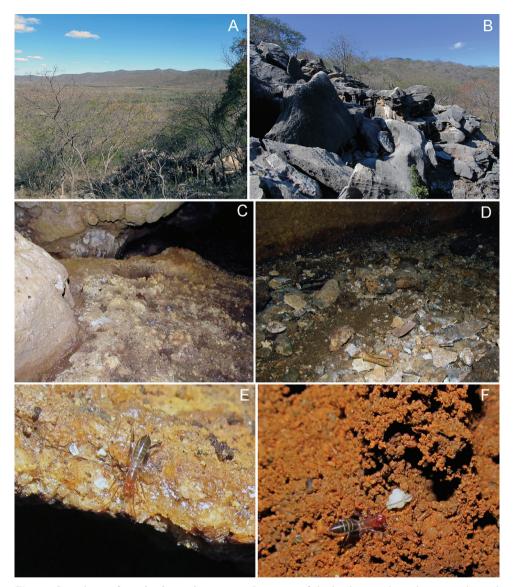


Figure 8. Habitat of *Rowlandius pedrosoi* sp. n. **A** A view of the landscape where the cave is located **B** Entrance of the cave **C–D** Microhabitat inside the cave where the specimens were collected **E** Female wandering on the cave floor **F** Female walking over some eggs.

rows of setae; two ventrodistal spines pointing forward; tarsal claw sharp and curved, slightly larger than half the tibia length; tarsal spur present.

Spermathecae of paratype (Fig. 7). Two pairs of lobes; stalk of the lateral lobe (LL) long, curved (the tips close to each other) and very light colored (almost transparent); with few granules along the structure. Tip of the LL with a wrinkled, rounded structure (resembling a walnut), brown colored (which means it is sclerotized), of about half width of the

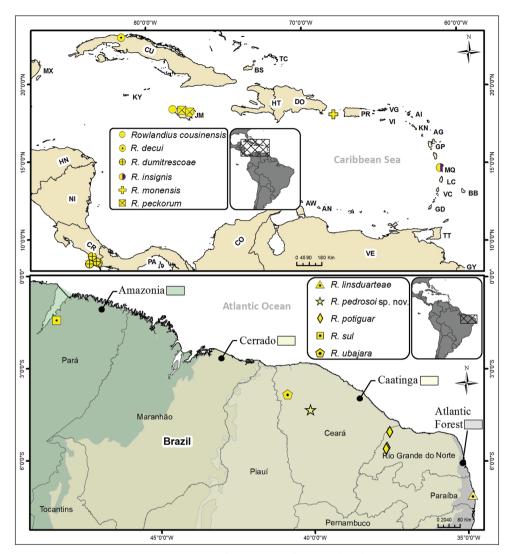


Figure 9. Map showing the distribution of the species of *Rowlandius dumitrescoae* group. The background colors in the Brazilian map represent the biomes.

stalk. The bases of LLs are separated by a distance similar to their lengths. The median lobes (ML) are short, cone-shaped, with a wide base and thin apex; its length is less than a third the size of the LL stalk; the integument is wrinkled with folds on its surface. Bases of the two lobes in contact. The chitinized arch is wider than long, cordiform (or as a "V", as described by Santos et al. 2013), similar to *R. potiguar*, however, in *R. pedrosoi* sp. n. the arch is strongly flattened. In *R. potiguar*, the vertex of the "V" has about 90–100° (a right angle tending towards the obtuse); in *R. pedrosoi* sp. n. the same vertex is clearly more obtuse than 150°.

Distribution (Fig. 9). Only known from the type locality: Brazil, Ceará, Santa Quitéria.

Natural history. The type locality is the largest cave in the state of Ceará, formed as a sloping crack (Fig. 8A–B) and with no more than seven square meters of floor space. The specimens were found in one of the few spots with some moisture in the ground. The soil was composed of damp earth of fine sediment agglomerated with gravel, small stones, shells of gastropods and bones from small mammals (Fig. 8C–F). When captured, the schizomids were walking on stones, gravel and debris, where the light barely reached (twilight zone).

Noteworthy of mention is a rare find of a secondary capilliconidium of a (probable) *Basidiobolus* sp. fungus among the cheliceral G7 setae (Fig. 3C, arrow; cf Blackwell and Malloch (1989)). The capilliconidium produces an apical droplet of extracellular material that helps the fungus to attach to and disperse with the host (Dykstra and Bradley-Kerr 1994).

Identification keys to the species of the dumitrescoae group

Key to the males (R. sul male unknown)

1	Occurs in Brazil
_	Occurs in the Caribbean or Central America5
2	Male pedipalp trochanter trapezoid in mesal view, with biggest edge facing downwards; apical region of trochanter with a small protrusion that does not touch the articulation of the trochanter-femur; pedipalps showing sexual dimorphism, i.e. larger than those of females; males with heteromorphs; posterodorsal process-XII long
_	Male pedipalp trochanter cylindrical in mesal view; apical region of trochanter without a small protrusion (all apical region is the articulation trochanter-femur); males without heteromorphs; pedipalps without sexual dimorphism; posterodorsal process-XII short
3	Posterodorsal process on abdominal segment XII with wide base (exceeding the width of the flagellum pedicel), with rhombus apex, almost as wide as the base of the process; flagellum wider in the basal third; in dorsal view, the dorsal projections of the flagellum reach the lateral border of the flagellum (see Santos et al. 2013, fig. 3A)
_	Posterodorsal process on abdominal segment XII with narrow base (not exceeding the width of the flagellum pedicel), with thin apex (much narrower than the base); flagellum wider in the median region; in dorsal view the dorsal projections do not reach or surpass the lateral borders of the flagellum (see Santos et al. 2008, fig. 1)
4	Posterodorsal process on abdominal segment XII wider than long; base of the male flagellum dorsal projections not connected, i.e. with a median projection between them (see Santos et al. 2013, fig. 4A, 5A)

5 - 6 - 7	Posterodorsal process on abdominal segment XII longer than wide (Fig. 6); base of the flagellum dorsal projections connected, i.e. without the median projection between them (Fig. 2)
_	Dorsum of flagellum, in lateral view, flat-shaped between lateral and poste-
0	rior bulge
8	Pedipalp dimorphic (elongated segments); flagellum in lateral view with median region and stalk at the same level
_	Pedipalp not dimorphic; flagellum in lateral view with median region higher than the level of the stalk
9	Flagellum lanceolate; flagellum in lateral view with flat posterior region
-	Flagellum nearly globose; flagellum in lateral view with elevated posterior region
Key to t	the females
Key to t	the females Occurs in Brazil
1 –	Occurs in Brazil
	Occurs in Brazil
1 –	Occurs in Brazil
1 –	Occurs in Brazil
1 - 2	Occurs in Brazil
1 –	Occurs in Brazil
1 - 2	Occurs in Brazil
1 - 2	Occurs in Brazil
1 - 2	Occurs in Brazil
1 - 2 - 3	Occurs in Brazil
1 - 2 - 3	Occurs in Brazil
1 - 2 - 3	Occurs in Brazil
1	Occurs in Brazil

6	Median lobes of spermathecae close to the base of the chitinized arch and
	distant to the base of the lateral lobes; lateral lobes long, stalk curved, apex
	discoid
_	Median lobes of spermathecae distant to the base of the chitinized arch and
	close to the base of the lateral lobes; lateral lobes long or short, stalk curved or
	not, and apex rounded or discoid7
7	Lateral lobes of spermathecae short; median and lateral lobes with their bases
	in the same line (one is not anterior or posterior to the other)
_	Lateral lobes of spermathecae long; base of the lateral and median lobes not
	in the same line8
8	Posterior region of the chitinized arch of spermathecae straight
_	Posterior region of the chitinized arch of spermathecae curved9
9	Lateral and median lobes of spermathecae close to the anterior region of the
	chitinized arch; median and lateral lobes with their bases in the same line
_	Lateral and median lobes of spermathecae in the center of the chitinized arch;
	median lobes positioned anteriorly to lateral lobes <i>R. insignis</i> (Martinique)

Discussion

In general, species groups facilitate comparisons and identifications in speciose genera as they comprise a subset of a genus, and make the process of understanding relationships

Table 1. Species groups and complexes proposed by Rowland and Reddell (1979a) and Reddell and Cokendolpher (1995) to the *dumitrescoae* group (when the species were still in *Schizomus* (R&R79) and after being transferred to *Rowlandius* (R&C95)).

Group	Complex	Species
	dumitrescoae complex	R. dumitrescoae
		R. decui
	primibiconourus complex	R. cousinensis
		R. primibiconourus
dumitrescoae group		R. longipalpus
		R. brevipatellatus
	viridis complex	R. gladiger
		R. monensis
		R. desecho
		R. biconourus
		R. insignis
		R. peckorum
		R. viridis

more comprehensible (Passos et al. 2015). Initially, only few genera were recognized in Schizomida and some of these (e.g., *Schizomus* Cook, 1899; *Trithyreus* Kraepelin, 1899) accumulated a number of species, but eventually they were subdivided, first into species groups, some of which were later recognized as new genera (Rowland and Reddell 1979a, b, 1980, 1981). The *dumitrescoae* group is an example of species group that was raised to genus. The group was defined by Rowland and Reddell (1979a) and was later transferred to *Rowlandius* by Reddell and Cokendolpher (1995). At that time, all species were from Central America (see Table 1 and Fig. 9) and were defined by the large body size, carapace with two to four pairs of dorsal and one pair of apical setae, female flagellum with four flagellomeres, spermathecae elongated laterally and reduced in the middle, and a few other characters (Rowland and Reddell 1979a). Afterwards, Armas (2002) proposed other species groups based on Cuban species and defined them using mainly characters of the pedipalp and the spermathecae (Table 2).

Studies on South American Schizomida revealed *Rowlandius* species inhabiting Brazil (Cokendolpher and Reddell 2000; Santos et al. 2008; Santos et al. 2013) and those species have a set of characters shared with some Caribbean (*R. cousinensis*, *R. decui*, *R. insignis*, *R. monensis* and *R. peckorum*) and Central American species (*R. dumitrescoae*), suggesting that the Brazilian *Rowlandius* fauna also belong to the *dumitrescoae* group. The characters present in all these species are: 1) female spermathecae with long lateral lobes and with a broad distal expansion, 2) median lobes short, digitiform without distal expansion, 3) gonopod absent, 4) chitinized arch with opened anterior branch (without AMN) and posterior branch rounded (*R. cousinensis*, *R. dumitrescoae*, *R. insignis*, *R. monensis* and *R. peckorum*), or anterior branch closed and posterior branch retrocurved (*R. pedrosoi* sp. n. and *R. potiguar*)

Table 2. Rowlandius species groups and complexes proposed by Armas (2002).

Groups	Subgroups	Species	Diagnostic character	
		R. biconourus	"D. C. I.	
I		R. ramosi	"Presence of a dorsal spur on the heteromorphic pedipalp trochanter of the male."	
		R. recuerdo	of the mate.	
II		R. abeli	"Spermathecae differs significantly from the general pattern present in congeners."	
Ш		R. decui	"Spermathecae with the terminal bulb underdeveloped and short	
111		R. digitiger	middle lobe."	
IV		R. cubanacan	"Long and subequal spermathecae with the terminal bulb	
1 V		R. labarcae	underdeveloped."	
	V-1	R. negreai		
		R. monticola		
	V-2	R. baracoae		
V	V-3	R. toldo	"Spermathecae with terminal bulbs well developed, with lateral lobes	
		R. gladiger	clearly longer and with a larger bulb."	
		R. alayoni		
		R. siboney		
		R. terueli		

basitarsus

telotarsus

Body	Male holotype MNRJ 4266	Female paratype MNRJ 4267
Total body: L	4.01	3.85
Propeltidium: L	1.25	0.98
Propeltidium: W	0.67	0.61
Metapeltidium: L	0.62	0.24
Metapeltidium: W	0.25	0.29
Abdomen: L	2.3	2.00
Abdomen: W	0.9	0.92
Flagellum: L	0.37	0.25
Flagellum: W	0.23	0.07
Pedipalp: L		
trochanter	0.82	0.58
femur	2.06	0.56
patella	1.84	0.62
tibia	0.91	0.53
tarsus + claw	0.54	0.41
Leg: I L		
coxa	0.42	0.61
trochanter	0.33	0.33
femur	1.18	1.23
patella	1.55	1.53
tibia	1.07	0.99
basitarsus	0.33	0.21
telotarsus	0.55	0.3
Leg: IV L		
femur	1.06	1.24
patella	0.51	0.55
tibia	0.82	0.89
1 :	0.67	0.67

Table 3. Measurements of *Rowlandius pedrosoi* sp. n. specimens.

or rounded (*R. linsduarte*, *R. ubajara* and *R. sul*), **5)** males with pedipalp elongated (such as *R. decui*, *R. dumitrescoae*, *R. insignis*, *R. potiguar* and *R. pedrosoi* sp. n.), and **6)** male flagellum never trilobate in dorsal view, but diamond-shaped and with dorsal projection (absent in *R. dumitrescoae* and reduced in *R. decui*). Santos et al. (2008) already noted that *R. linsduarte* and *R. sul* are more closely related to each other than to any other species based on the female genitalia, but did not include them in any group. Here a new composition of the *dumitrescoae* group is proposed based on the above-mentioned characters (see also Table 4).

0.67

0.46

0.67

0.47

Some *Rowlandius* illustrated in the literature are potentially part of the *dumitres-coae* group, but are not presently included, once no material was accessible during the preparation of the work. One of them is an undescribed species from Tortuguero,

Species maintained	Species removed	Species added
R. cousinensis	R. primibiconourus*	R. linsduarte
R. decui	R. longipalpus	R. potiguar
R. dumitrescoae	R. gladiger	R. sul
R. monensis	R. desecho	R. ubajara
R. peckorum	R. biconourus	R. pedrosoi sp. n.
R. insignis	R. viridis	

Table 4. List of species maintained, removed, and added to the *dumitrescoae* group.

Costa Rica, illustrated by Armas (2009) (see fig. 3D); the spermathecae of the specimen fits the present definition of the *dumitrescoae* group, but as the species was not formally described and the male is not known, the correct relationship of the morphospecies cannot be assured by now. Another species that can potentially be part of the group is *R. viridis*; Rowland and Reddell (1979a) illustrated this species from four localities, and one of them (from Pedro Great Cave, Clarindon Parish) is similar to the standard shape of the *dumitrescoae* group, but since there is a huge variation in the size and shape of the lobes in this species, further studies are needed before reaching a conclusion on those populations.

An interesting character observed in some species of *Rowlandius* (e.g. *R. dumitres-coae*, *R. insignis*, *R. potiguar* and *R. pedrosoi* sp. n.) is the strong sexual dimorphism of the palps. The femur and patella of the pedipalps are extremely long in α -heteromorphic males compared to females and homeomorphic males, as reported by Santos et al. (2013). Other cases of elongated male-dimorphic appendages in arachnids are found in harvestmen (Orrico and Kury 2009; Buzatto et al. 2011; Zatz et al. 2011) and whip spiders (Vasconcelos et al. 2014). It is possible that the elongate pedipalps of *R. pedrosoi* sp. n. evolved due to sexual selection pressures, similarly to that found in *R. potiguar* (Santos et al. 2013).

Rowlandius is the only short-tailed whip scorpion genus found in the dry biome of Caatinga (Santos et al. 2008; Santos et al. 2013). The four schizomid species found in that harsh environment (R. linsduarte, R. pedrosoi sp. n., R. potiguar and R. ubajara) are restricted to protected places, such as forests or caves, where the temperature is mild, the humidity is high and the variation these environmental conditions is lower. These species appear to be limited to these hypogean habitats, but they do not have apparent troglomorphisms and their presence in caves may be a recent invasion after climate change in Northeastern Brazil and retraction of the forest (Santos et al. 2007). The small size and the relatively thin cuticle of schizomids makes them sensitive to dehydration and caves serve as a suitable habitat for these animals (Oliveira and Ferreira 2014). The exotic species Stenochrus portoricensis Chamberlim, 1922, for example, has already been found in caves in Central Brazil (Gallão et al. 2015).

^{*}This species was removed because its documentation in the literature is insufficient and we had no access to specimen; see discussion for details.

Acknowledgments

We thank Denis Rafael Pedroso (MNRJ) and Flávia Pellegatti Franco for collecting the material and providing specimens for description; Henrik Enghoff for helping with the identification of the Basidiobolus fungus; Lars Vilhelmsen for checking the English; Plataforma de Microscopia Eletrônica Rudolf Barth (FIOCRUZ - IOC) and its technologists and technicians Roger, Wendell and Rômulo for helping with the SEM. We are also grateful to Mario Gatti and Rodolfo Armando da Cunha from *Coleção* Micológica Trichocomaceae (FIOCRUZ - IOC) for making available the microscope for image capture, to José Augusto Martins Roxinol (UFV; SISBIOTA 563360/2010-0) for donating a specimen of R. ubajara and Caroline Fukushima (IBSP) and Wilmar Silva (IBSP) for donating specimens of R. potiguar (this material was collected in the scope of the project "Biodiversidade de Aranhas Migalomorfas (Araneae, Mygalomorphae) de diferentes formações vegetais do Rio Grande do Norte, Brasil" (CAPES #23038.00814/2011-83)). We are also grateful to the reviewers James Cokendolpher, Mark Harvey and Jairo A. Moreno-González for the important corrections and comments that helped improve the quality of the article. This study was partly supported by grants from Coordenação de Aperfeicoamento de Pessoal de Nível Superior (CAPES, www.capes.gov.br/) to OVM (process number 5900115 CAPES/PEC-PG) and to GSM (process number 8922-13-6 CAPES/Science Without Borders). GSM is also grateful for the Danish National Research Foundation for support to the Center for Macroecology, Evolution and Climate (grant number DNRF96).

References

- Armas LFd (2002) Nuevas especies de *Rowlandius* Reddell & Cokendolpher, 1995 (Schizomida: Hubbardiidae) de Cuba. Revista Ibérica de Aracnología 6: 149–167. http://www.sea-entomologia.org/PDF/RIA_6/R06-016-149.pdf
- Armas LFd, Villarreal OM, Colmenares-García PA (2009) Nuevo *Rowlandius* Reddell & Cokendolpher, 1995 (Schizomida: Hubbardiidae) de la Sierra San Luis, Venezuela noroccidental. Papeis Avulsos de Zoologia 49: 361–368. doi: 10.1590/S0031-10492009002800001
- Blackwell M, Malloch D (1989) Similarity of *Amphoromorpha* and secondary capilliconidia of *Basidiobolus*. Mycologia 81: 735–741. doi: 10.2307/3759878
- Buzatto BA, Requena GS, Lourenço RS, Munguía-Steyer R, Machado G (2011) Conditional male dimorphism and alternative reproductive tactics in a Neotropical arachnid (Opiliones). Evolutionary Ecology 25: 331–349. doi: 10.1007/s10682-010-9431-0
- Cokendolpher JC, Reddell JR (1992) Revision of Protoschizomidae (Arachnida: Schizomidae) with notes on the phylogeny of the order. Texas Memorial Museum Speleological Monographs, Austin 3: 31–74. http://www.nsrl.ttu.edu/personnel/JCCokendolpher/Cokendolpherpubs/publications/1992Cokendolpher_Reddell_Protoschizomidae.pdf
- Cokendolpher JC, Reddell JR (2000) New and rare Schizomida (Arachnida: Hubbardiidae) from South America. Amazoniana 16: 187–212. http://www.nsrl.ttu.edu/personnel/

- $\label{local-control} JCC oken dolpher/Coken dolpher pubs/publications/2000 Coken dolpher_Reddell_Schiz-South Amer.pdf$
- Delgado-Santa L, Armas LF (2013) Tres nuevos Hubbardiinae (Schizomida: Hubbardiidae) de Colombia. Revista Ibérica de Aracnología 22: 37–45. http://www.sea-entomologia.org/PDF/RIA22/037045RIA22HubbardinaeColombiaST.pdf
- Dykstra MJ, Bradley-Kerr B (1994) The Adhesive Droplet of Capilliconidia of *Basidiobolus ranarum* Exhibits Unique Ultrastructural Features. Mycologia 86: 336–342. doi: 10.2307/3760563
- Gallão JE, Bichuette ME, Giupponi APL (2015) First record of *Stenochrus portoricensis* Chamberlin, 1922 (Arachnida: Schizomida: Hubbardiidae) for caves in Brazil: evidence for a troglophile status of an exotic species. Check List 11: 1546. doi: 10.15560/11.1.1546
- Harvey MS (1992) The Schizomida (Chelicerata) of Australia. Invertebrate Taxonomy 6: 77–129. doi: 10.1071/IT9920077
- Harvey MS (2003) Catalogue of the smaller arachnid orders of the world. Amblypygi, Uropygi, Schizomida, Palpigradi, Ricinulei and Solifugae. CSIRO Publishing, Collingwood, 398 pp.
- Lawrence RF (1969) The trichoid structures on the chelicarae of the short-tailed whip-scorpions (Schizomida; Arachnida). Transactions of the Royal Society of Africa 38: 123–132. doi: 10.1080/00359196909519080
- Monjaraz-Ruedas R, Francke OF, Cruz-López JA, Santibánez-López CE (2016) Annuli and setal patterns in the flagellum of female micro-whipscorpions (Arachnida: Schizomida): Hypotheses of homology across an order. Zoologischer Anzeiger 263: 118–134. doi: 10.1016/j.jcz.2016.05.003
- Moreno-González JA, Delgado-Santa L, Armas LF (2014) Two new species of *Piaroa* (Arachnida: Schizomida, Hubbardiidae) from Colombia, with comments on the genus taxonomy and the flagellar setae pattern of Hubbardiinae. Zootaxa 3852: 227–251. doi: 10.11646/zootaxa.3852.2.4
- Oliveira MPAd, Ferreira RL (2014) Aspects of the Behavior and Activity Rhythms of *Rowlan-dius potiguar* (Schizomida: Hubbardiidae). PLoS ONE 9: e91913. doi: 10.1371/journal. pone.0091913
- Orrico VGD, Kury AB (2009) A cladistic analysis of the Stygnicranainae Roewer, 1913 (Arachnida, Opiliones, Cranaidae) where do longipalp cranaids belong? Zoological Journal of the Linnean Society 157: 470–494. doi: 10.1111/j.1096-3642.2009.00543.x
- Passos MI, Miranda GS, Nessimian JL (2015) Three new species of *Macrelmis* Motschulsky (Coleoptera: Elmidae: Elminae) from Southeastern Brazil with new definition of species groups to the genus. Zootaxa 4058: 195–210. doi: 10.11646/zootaxa.4058.2.3
- Reddell JR, Cokendolpher JC (1995) Catalogue, bibliography, and generic revision of the order Schizomida (Arachnida). Texas Memorial Museum Speleological Monographs, Austin 4: 1–170. http://www.nsrl.ttu.edu/personnel/JCCokendolpher/Cokendolpherpubs/publications/1995Reddell_Cokendolpher_catalogue.pdf
- Rowland JM, Reddell JR (1979a) The order Schizomida (Arachnida) in the New World. I. Protoschizomidae and *dumitrescoae* group (Schizomidae: *Schizomus*). Journal of Arachnology 6: 161–196. http://www.jstor.org/stable/3705088
- Rowland JM, Reddell JR (1979b) The order Schizomida (Arachnida) in the New World. II. *simonis* and *brasiliensis* groups (Schizomidae: *Schizomus*). The Journal of Arachnology 7: 89–119. http://www.americanarachnology.org/JoA_free/JoA_v7_n2/JoA_v7_p89.pdf

- Rowland JM, Reddell JR (1980) The order Schizomida (Arachnida) in the New World. III. *mexicanus* and *pecki* groups (Schizomidae: *Schizomus*). Journal of Arachnology 8: 1–34. http://www.jstor.org/stable/3705202
- Rowland JM, Reddell JR (1981) The order Schizomida (Arachnida) in the New World . IV. goodnightorum and briggsi groups and unplaced species (Schizomidae, *Schizomus*). Journal of Arachnology 9: 19–46. http://www.americanarachnology.org/JoA_free/JoA_v9_n1/JoA_v9_p19.pdf
- Santos AJ, Dias SC, Brescovit AD, Santos PP (2008) The arachnid order Schizomida in the Brazilian Atlantic Forest: a new species of *Rowlandius* and new records of *Stenochrus portoricensis* (Schizomida: Hubbardiidae). Zootaxa 1850: 53–60. doi: 10.11646/%25x
- Santos AJ, Ferreira RL, Buzzato BA (2013) Two new cave-dwelling species of the short-tailed whipscorpion genus *Rowlandius* (Arachnida: Schizomida: Hubbardiidae) from Northeastern Brazil, with comments on male dimorphism. PLoS ONE 8: e63616. doi: 10.1371/journal. pone.0063616
- Santos AMM, Cavalcanti DR, Silva JMC, Tabarelli M (2007) Biogeographical relationships among tropical forests in north-eastern Brazil. Journal of Biogeography 34: 437–446. doi: 10.1111/j.1365-2699.2006.01604.x
- Teruel R (2004) Nuevas adiciones a la fauna de esquizómidos de Cuba Oriental, con la descripción de cuatro nuevas especies (Schizomida: Hubbardiidae). Revista Ibérica de Aracnología 9: 31–42. http://www.sea-entomologia.org/Publicaciones/RevistaIbericaAracnologia/RIA09/R09-006-031.pdf
- Teruel R (2012) Un nuevo Rowlandius Reddell & Cokendolpher 1995 del Macizo de Guamuhaya, Cuba central (Schizomida: Hubbardiidae). Revista Ibérica de Aracnología 21: 61–64. http://www.sea-entomologia.org/PDF/RIA21/061064RIA21Rowlandiusmantine zinspcolor.pdf
- Teruel R, Armas LFd, Rodríguez T (2012) Adiciones a los esquizómidos de Cuba central, con la descripción de cuatro nuevos *Rowlandius* Reddell & Cokendolpher, 1995 (Schizomida: Hubbardiidae). Revista Ibérica de Aracnología 21: 97–112. http://www.sea-entomologia.org/PDF/RIA21/097112RIA21SchizomidanuevosCubacolorbr.pdf
- Vasconcelos ACO, Giupponi APL, Ferreira RL (2014) A new species of *Charinus* from Minas Gerais State, Brazil, with comments on its sexual dimorphism (Arachnida: Amblypygi: Charinidae). The Journal of Arachnology 42: 155–162. doi: 10.1636/H14-01.1
- Villarreal OM, Armas LF, García LF (2014) A new species of *Piaroa* (Schizomida: Hubbardiidae) from Venezuela, with taxonomic notes on the genus. Zootaxa 3765: 371–381. doi: 10.11646/zootaxa.3765.4.5
- Villarreal OM, Miranda GS, Giupponi AP (2016) New Proposal of Setal Homology in Schizomida and Revision of *Surazomus* (Hubbardiidae) from Ecuador. PLoS ONE 11: e0147012. doi: 10.1371/journal.pone.0147012
- Zatz C, Werneck RM, Macías-Ordóñez R, Machado G (2011) Alternative mating tactics in dimorphic males of the harvestman *Longiperna concolor* (Arachnida: Opiliones). Behavioral Ecology and Sociobiology 65: 995–1005. doi: 10.1007/s00265-010-1103-0