

## To boldly go: on invasive goblin spiders in Brazil (Araneae, Oonopidae)

Antonio D. Brescovit<sup>1</sup> , Alexandre B. Bonaldo<sup>2</sup> , Ricardo Ott<sup>3</sup> & João L. Chavari<sup>1</sup>

1. Instituto Butantan, Laboratório Especial de Coleções Zoológicas. Av. Vital Brasil, 1500, 05503-900 São Paulo, SP, Brazil. (antonio.brescovit@butantan.gov.br)  
2. Museu Paraense Emílio Goeldi, Coordenação de Zoologia, Laboratório de Aracnologia, Campus de Pesquisa, Av. Perimetral, 1901, 66040-170 Belém, PA, Brazil.  
3. Museu de Ciências Naturais, Fundação Zoobotânica do Rio Grande do Sul, Rua Dr. Salvador França, 1427, 90690-000 Porto Alegre, RS, Brazil.

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**ABSTRACT.** Twelve non-native species of the spider family Oonopidae are recorded in Brazil: *Brignolia dasysterna* Platnick, Dupérré, Ott & Kranz-Baltensperger, 2011, *B. parumpunctata* (Simon, 1893), *Triaeris stenaspis* Simon, 1892, *Ischnothyreus peltifer* (Simon, 1892), *I. velox* Jackson, 1908, *Opopaea concolor* (Blackwall, 1859), *O. deserticola* Simon, 1892, *Pelcinus marmoratus* Simon, 1892, *Heteroonops spinimanus* (Simon, 1892), *Gamasomorpha parmata* (Thorell, 1890) (herein restored to its original combination in *Xestaspis*), *Orchestina pavesiiformis* Saaristo, 2007 and *O. dentifera* (Simon, 1893). Among these species, six new synonyms were detected: *Hytanis oblonga* Simon, 1893, *Xestaspis bipeltis* Thorell, 1895, *Gamasomorpha insularis* Simon, 1907 and *Opopaea lutzi* Petrunkevitch, 1929 with *Xestaspis parmata* Thorell, 1890; *Gamasomorpha humilis* Mello-Leitão, 1920 with *Opopaea concolor* (Blackwall, 1859) and *Gamasomorpha rufa* Banks, 1898 with *Opopaea deserticola* Simon, 1891. Since *Hytanis oblonga* is the type species of the genus, its synonymy with *Xestaspis parmata* imply in the synonymy of *Hytanis* Simon, 1893 with *Xestaspis* Simon, 1884. The partenogenetic species *T. stenaspis* is the most distributed synanthropic oonopid in Brazil, occurring all over the country, mainly in urban areas. Some species are reported here to occur in the interior of natural caves, a finding that should raise conservation concerns.

**KEYWORDS.** Spiders, synanthropy, distribution, conservation, goblin spiders.

**RESUMO.** Audaciosamente indo: aranhas gnomo invasoras no Brasil (Araneae, Oonopidae). Doze aranhas consideradas não-nativas da família Oonopidae são registradas em áreas urbanas no Brasil: *Brignolia dasysterna* Platnick, Dupérré, Ott & Kranz-Baltensperger, 2011, *B. parumpunctata* (Simon, 1893), *Triaeris stenaspis* Simon, 1892, *Ischnothyreus peltifer* (Simon, 1892), *I. velox* Jackson, 1908, *Opopaea concolor* (Blackwall, 1859), *O. deserticola* Simon, 1892, *Pelcinus marmoratus* Simon, 1892, *Heteroonops spinimanus* (Simon, 1892), *Gamasomorpha parmata* (Thorell, 1890) (aqui restaurada para sua combinação original em *Xestaspis*), *Orchestina pavesiiformis* Saaristo, 2007 and *O. dentifera* (Simon, 1893). Entre estas espécies, seis novos sinônimos foram detectados: *Hytanis oblonga* Simon, 1893, *Xestaspis bipeltis* Thorell, 1895, *Gamasomorpha insularis* Simon, 1907 e *Opopaea lutzi* Petrunkevitch, 1929 com *Xestaspis parmata* Thorell, 1890; *Gamasomorpha humilis* Mello-Leitão, 1920 com *Opopaea concolor* (Blackwall, 1859) e *Gamasomorpha rufa* Banks, 1898 with *Opopaea deserticola* Simon, 1891. Uma vez que *Hytanis oblonga* é a espécie-tipo do gênero, sua sinonímia com *Xestaspis parmata* implica na sinonímia de *Hytanis* Simon, 1893 com *Xestaspis* Simon, 1884. *Triaeris stenaspis*, uma espécie partenogenética, é o oonopídeo sinantrópico com distribuição mais ampla no Brasil, ocorrendo de norte a sul do país, principalmente em áreas urbanas. Algumas espécies relacionadas aqui ocorrem no interior de cavernas naturais, uma descoberta que deve levantar preocupações em relação à conservação das espécies nativas de cavernas.

**PALAVRAS-CHAVE.** Aranhas, animais sinantrópicos, distribuição, conservação, aranhas gnomo.

The theme of invasive species and the discussion on the artificial introduction of species from one continent to another have been amplified in recent years (THOMPSON, 2014; PEARCE, 2015), with most studies involving vertebrates (LONG, 1981, 2003; KRAUS, 2009) or invertebrates from terrestrial or aquatic urban ecosystems (DEE BOERSMA *et al.*, 2006).

Invasions of alien spiders on the Neotropical Region are yet poor studied. Most of the available papers are descriptive (CARVALHO *et al.*, 2007; ALMEIDA-SILVA & BRESCOVIT, 2008; OTT & BRESCOVIT, 2013; POMPOZZI *et al.*, 2013; RUIZ & BONALDO, 2013; TAUCARE-RÍOS & BUSTAMANTE, 2015) and just a few deals with the relations of the studied species with their environments (MALUMBRES-OLARTE, 2015; RODRÍGUEZ-RODRÍGUEZ *et al.*, 2015). Nevertheless, the handful studies

carried on in Brazil have been showing a large number of invasive or synanthropic spiders and the list of those spiders in Brazilian territory depicts today at least fifty species (BRESCOVIT, 2002; INDICATTI & BRESCOVIT, 2008; SIMÓ *et al.*, 2015).

The recognition of a given species as alien to a certain country or continent depends on the availability of reliable taxonomic data across large geographical regions. The taxonomic impediment to recognize patterns of biological invasions is particularly acute for megadiverse, poorly known arthropod groups. The spider family Oonopidae provides an excellent example in that context. This family has been subject of a worldwide coordinated effort to forward its taxonomic knowledge through the Goblin Spider Planetary Biodiversity Inventory project (PBI-Oonopidae), sponsored

by NSF/AMNH. Before this project, Oonopidae was one of the poorest-known spider families and faunistic papers invariably depicted extremely low systematic resolutions for this taxon (ABRAHIM *et al.*, 2012). During the activity peak of the PBI-Oonopidae between 2009 to 2014 the number of known species increased in nearly 300%. The efforts of the research group in that period raised the assigned species to the family from 472 to 1,600 (WORLD SPIDER CATALOG, 2019). Furthermore, due to the revisional approach adopted by the project, several previously unknown lineages were recognized and general patterns of geographic distribution of these lineages were brought to light (ÁLVAREZ-PADILLA *et al.*, 2012; THOMA *et al.*, 2014). While most Oonopidae species are microdistributed, presenting a great potential in the recognition of areas of endemism, a small set of species are seemingly synanthropic and have attained Pantropical or almost cosmopolitan distributions (PLATNICK & DUPÉRRÉ, 2009a; PLATNICK *et al.*, 2012a), boldly going where no other representative of their respective lineages has gone before.

Recently, studying substantial material of Brazilian oonopids, we detected twelve non-native species established today in Brazilian territory. Several of these species were apparently introduced relatively earlier (Tab. I) and are today established in large populations in urban areas, parks, houses and even in cave systems. The taxonomic status of the majority of these species was well defined in previous PBI-Oonopidae papers, but the examination of a large number of type specimens during the project allowed the discovery of six additional synonymies under the species here studied.

## MATERIAL AND METHODS

The material examined and types cited belongs to the following institutions: AMNH, American Museum of Natural History, New York (curator: L. Prendini); BMNH, Natural History Museum, London (J. Beccaloni); HUJ, Hebrew University of Jerusalem, Jerusalem (A. Chipman); IBSP, Instituto Butantan, São Paulo (A. D. Brescovit); INPA, Instituto Nacional de Pesquisas da Amazônia, Manaus (C. Magalhães); MCN, Museu de Ciências Naturais, Fundação Zoobotânica do Rio Grande do Sul, Porto Alegre (R. Ott); MCTP, Museu de Ciências e Tecnologia, Pontifícia Universidade Católica, Porto Alegre (R. A. Teixeira); MCZ, Museum of Comparative Zoology, Harvard University, Cambridge (G. Giribet); MNHN, Muséum National d'Histoire Naturelle, Paris (C. Rollard); MNRJ, Museu Nacional, Universidade Federal do Rio de Janeiro, Rio de Janeiro (A. B. Kury); MPEG, Museu Paraense Emílio Goeldi, Belém (A. B. Bonaldo); MRAC, Musée Royal de l'Afrique Centrale, Tervuren (R. Jocqué); MZSP, Museu de Zoologia, Universidade de São Paulo, São Paulo (R. Pinto da Rocha); MZUT, Zoological Museum, University of Turku, Turku (S. Koponen); SMNK, Staatliches Museum für Naturkunde Karlsruhe, Karlsruhe (H. Höfer).

Morphological observations were made using a Leica MZ12 stereomicroscope. Photographs were taken with a Leica DFC 500 digital camera mounted on a Leica MZ 16A stereomicroscope and extended focal range images were

composed with Leica Application Suite version 2.5.0, at IBSP. Other images were taken through a Zeiss Standard compound microscope with attached Canon A620 camera and processed with Helicon Focus 5.3 multi-range program (KOZUB *et al.*, 2012), at MCN. All figures were edited using Adobe Photoshop CS5 ver. 12.0. Maps were produced using QGIS 3.2.1 and edited in Adobe Photoshop CS6.

## TAXONOMY

### *Brignolia dasystema* Platnick, Dupérré, Ott & Kranz-Baltensperger, 2011 (Figs 1–6, 58)

*Brignolia dasystema* PLATNICK *et al.*, 2011:32, figs 95–141. Male holotype and female allotype taken in Berlese samples of young hammock forest litter at Deering Estate Park, SW 167<sup>th</sup> St. and SW 72<sup>nd</sup> Ave., South Miami, Dade Co., Florida, USA, 1.VI–25.VIII.1986, J. & S. Peck leg., in AMNH 37534, PBI\_OON 1270, not examined.

*Brignolia cubana*: FREITAS *et al.*, 2013:9–10, 15 (misidentification).

**Diagnosis.** *Brignolia dasystema* can be easily separated from autochthone Neotropical oonopids and from *B. parumpunctata* by the absence of ventral directed protrusion situated medially on the clypeus (see PLATNICK *et al.*, 2011, fig. 96), modified setae on the male sternum (Fig. 3), male palp with small protrusion on the palpal bulb (Figs 5, 6; PLATNICK *et al.*, 2011, figs 103–108) and females by the low protrusion situated between the epigastric furrow and the groove connecting the posterior spiracles (Fig. 4).

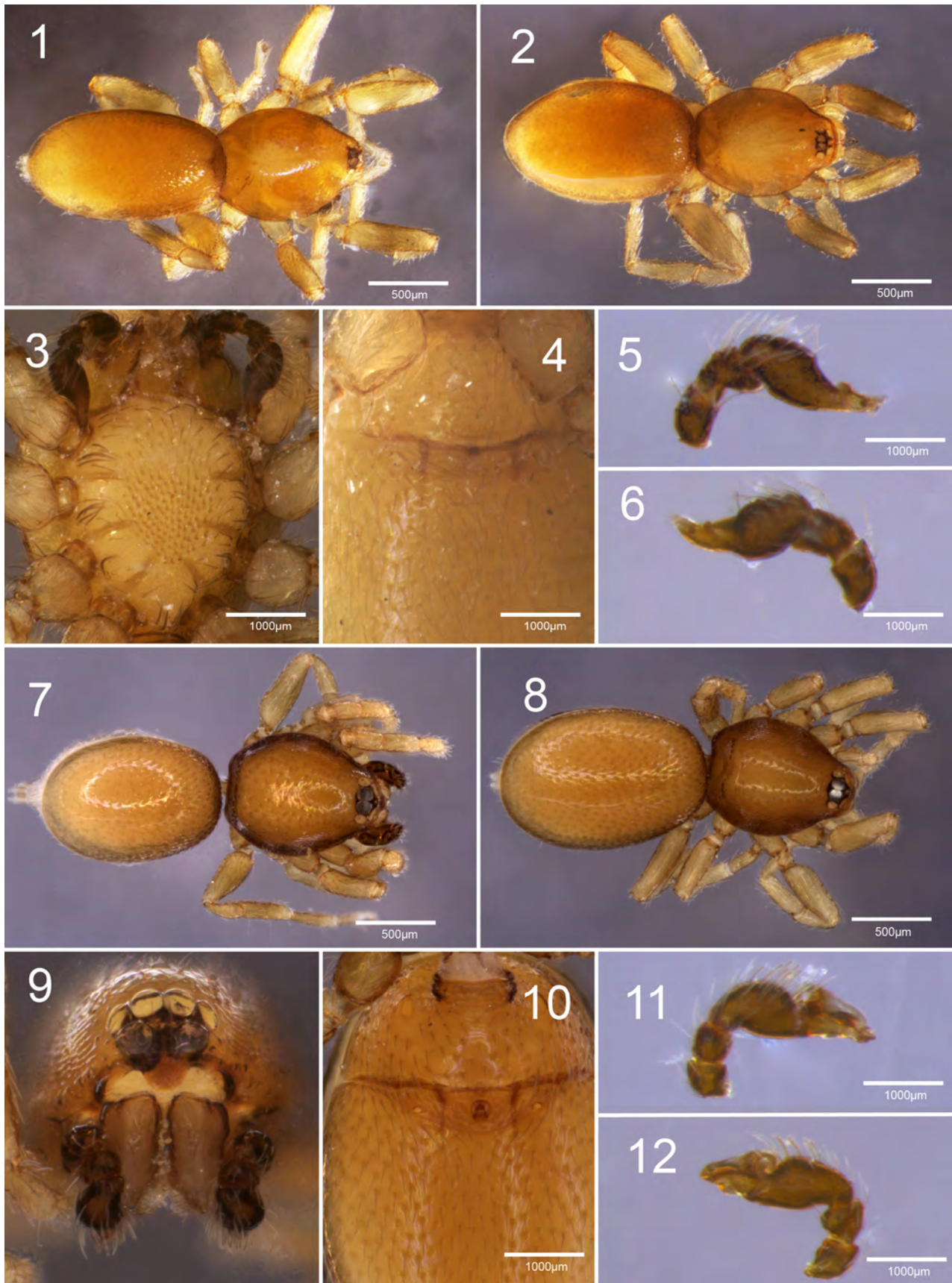
**Description.** Male and female, see PLATNICK *et al.*, 2011:32, figs 95–141.

**Distribution.** This species was recently described by PLATNICK *et al.* (2011) as being restrict to southern Florida, USA. It is recorded here to occur in at least four states of north and northeast Brazil (Fig. 58), where it may have been introduced earlier than in the USA. FREITAS *et al.* (2013) found a large population of *Brignolia dasystema* (sub *Brignolia cubana*) in Fernando de Noronha archipelago.

**Material examined.** BRAZIL. **Pará:** Uruará (03°43'03"S - 53°44'14"W), 3♀, 2014, R. L. Cajaiba leg. (IBSP 166582; IBSP 166583); Belém, Campus IPEAN (01°24'46.14" - 01°28'4.11"S - 48°20'4.6" - 48°20'31.84"W), 2♂ ♀, 2.V.1974, R. Schuh (AMNH PBI\_OON 485–487); **Pernambuco:** Fernando de Noronha, Arquipélago de Fernando de Noronha (03°50'57.2814"S - 32°24'10.7922"W), 51♂ 55♀, X.2005–IV.2006, G. Freitas leg. (IBSP 67112–67143; IBSP 67726–67736; IBSP 67852–67882); **Tocantins:** Porto Nacional, urban area (10°42'S - 48°25'W), ♀, 14–23. XI.2003, I. Knysak & R. Martins leg. (IBSP 92232); **Bahia:** Salvador, Baía de Aratu, Ponte do Fernandinho (12°47'32.8"S - 38°28'15.3"W); ♀, 13.IX.2004, K. Benati leg. (IBSP 58898); Ipiaú (14°8'18.8844"S - 39°45'47.6028"W), ♂, 2008, J. Cardoso leg. (IBSP 121852).

### *Brignolia parumpunctata* (Simon, 1893) (Figs 7–12, 58)

*Xestaspis parumpunctata* SIMON, 1893a:305. Female holotype from Sierra Leone, no specific locality, in MNHN.



Figs 1-12. Figs 1-6, *Brignolia dasystema* Platnick, Dupérré, Ott & Kranz-Baltensperger, 2011, male: 1, dorsal; 3, modified hairs in the sternum; 5, palp, retrolateral; 6, palp, prolateral. Female: 2, dorsal; 4, genitalia, ventral. Figs 7-12, *Brignolia parumpunctata* (Simon, 1893), male: 7, dorsal; 9, projection of clypeus; 11, palp prolateral; 12, palp retrolateral. Female: 8, dorsal; 10, genitalia, ventral.

*Gamasomorpha perplexa* BRYANT, 1942:325, figs 11-12, 18. Male holotype from Saint Croix, Virgin Islands, in MCZ; PLATNICK *et al.*, 2011:15 (Syn.).

*Opopaea recondita* CHICKERING, 1951:231, figs 18-19. Female holotype from Canal Zone Biological Area [= Barro Colorado Island], Panama, Panama, in MCZ; PLATNICK *et al.*, 2011:15 (Syn.).

*Brignolia cubana* DUMITRESCU & GEORGESCU, 1983:107, pl. 22, figs 1-8. Male holotype from Estación Ecológica Siboney, Santiago de Cuba, Cuba, in Institut de Spéologie, Bucharest; PLATNICK *et al.*, 2011:15 (Syn.).

*Brignolia recondita*: PLATNICK & DUPÉRRÉ, 2009a:4.

*Brignolia parumpunctata*: PLATNICK *et al.*, 2011:14.

**Diagnosis.** *Brignolia parumpunctata* can be easily separated from *B. dasystema* by the males having a ventrally directed protrusion situated medially on the clypeus (Fig. 9; PLATNICK *et al.*, 2011: fig. 21) and by the distinctive curled dorsal protrusion on the palpal bulb (Figs 11, 12; PLATNICK *et al.*, 2011: figs 31-32). Females can be easily recognized by the rounded, elevated protrusion situated between the epigastric furrow and by the groove connecting the posterior spiracles (Fig. 10).

**Description.** Male and female, see PLATNICK *et al.*, 2011:14, figs 1-94.

**Distribution.** This species has a Pantropical distribution, occurring in the Americas, Africa, Southern Asia, Pacific Islands and Australia. PLATNICK *et al.*, 2011:31, reported only one previously Brazilian record, in the state of Pará, norther Brazil. Here it is recorded in areas of north, northeast and southeast Brazil (Fig. 58). According to PLATNICK *et al.* (2011), this species may have an Asian origin, where the bulk of the genus diversity is from.

**Material examined.** BRAZIL. **Rondônia:** Porto Velho, Campo Experimental da EMBRAPA (08°48'9.7878"S - 63°49'48.1188"W), ♂, 2008, L. S. P. Trigueiro & C. A. D. Teixeira *et al.* leg. (IBSP 121507); **Pará:** Belém, *Campus IPEA*, 2♂, ♀, 02.V.1974, R. Schuh leg. (AMNH, PBI\_OON 485-487); **Pernambuco:** São Lourenço da Mata, Estação Ecológica de Tapacurá (08°0'39.8088"S - 35°5'2.8422"W), 2♂, 2008, C. Valença leg. (IBSP 97891, 97892); **Minas Gerais:** Belo Horizonte, *Campus UFMG, Estação Ecológica da UFMG* (19°52'S - 43°58'W), 3♂, 29.III.2000, É. S. S. Álvares & C. S. Azevedo leg. (IBSP 27202); 2♂, 10♀, IX.1999-II.2001, É. S. S. Álvares, E. O. Machado & C. S. Azevedo leg. (IBSP 32350); 4♂, XII.2000-I.2001, É. S. S. Álvares leg. (IBSP 66797); 5♂, I.2001, É. S. S. Álvares leg. (IBSP 66801); 2♂, II.2001 (IBSP 67303); Jaboticatubas, Serra do Cipó, Parque Nacional da Serra do Cipó (19°15'S - 43°31'W), 31♂, 21♀, 07-14.X.2002-26.VI.2003, Equipe Biota leg. (IBSP 69443, 69445-69476); Marliéria, Parque Estadual do Rio Doce (19°48'/19°29'S - 42°38'/42°28'W), 18♂, 20♀, 01-10.IX.2003, Equipe Biota leg. (IBSP 144583-144597); **Bahia:** Salvador, Baía de Aratu, Ponte do Fernandinho (12°47'32.8"S - 38°28'15.3"W), 18♂, ♀, 13.IX.2004, K. Benati leg. (IBSP 58842, 58843, 58852, 58863, 58869, 58874, 58875, 58877, 58883, 58884, 58901, 58906, 58907, 58915, 58917, 58922, 58938, 58970, 58976, 59013, 59019); Parque da Cidade Joventino Silva (12°59'57.6522"S - 38°28'11.1606"W), ♂, 22.VIII-03.

IX.2008, C. Queiroz & D. Mendes leg. (IBSP 125569); Baía de Aratu (12°47'32.8"S - 38°28'15.3"W), 42♂ 8♀, 2007-2008, K. Benati leg. (IBSP 125210, 125260, 125289, 126033, 126035-126039, 126044, 126047, 126051-126056, 126058, 126060, 126064-126065, 126067-126068, 126074, 126077-126078, 126080, 126083-126085, 126087, 126091-126092, 126094, 126097, 126100, 126104, 126106, 126111-126112, 126114, 126116, 126118-126125, 126128, 126130, 126134); Ilhéus, CEPLAC (14°47'45.5712"S - 39°11'25.2024"W, 5♂, 2♀, 11.IV.1998, A. D. Brescovit *et al.* leg. (IBSP 19206, 19211, 19220, 19290, 19302); 2♂, 14♀, 24.VII.1998 (IBSP 36122); 2♂, 18.IV.2007 - 06.IX.2007, P. P. Santos leg. (IBSP 98012, 98042); Una, Reserva Ecológica do Una (15°17'48"S - 39°04'28"W), ♂, X.1999 - IX.2000, M. F. Dias leg. (IBSP 64428); Itapetinga, Parque Municipal da Matinha (15°15'43.8618"S - 40°17'44.9376"W), ♂, III-IV.2003, J. P. S. Alves leg. (IBSP 66457); **Rio de Janeiro:** Volta Redonda, Mata da Cicuta (22°33'2.016"S - 44°5'31.7328"W), ♀, 11-18.VI.2003, F. S. Cunha leg. (IBSP 44865); 5♂, 2♀, 04-11.III.2002 - 11-18.VI.2001, Equipe Biota leg. (IBSP 136203-136208); ♂, 11.III.2002, E. F. Ramos leg. (IBSP 144105); Paraíba do Sul, Fazenda Maravilha, ♂, 17-21.VIII.2001, Equipe Biota leg. (IBSP 136639); Resende, Mata das Indústrias Nucleares (22°31'2.2326"S - 44°39'10.7748"W), ♂, VII-IX.2007, E. F. Ramos leg. (IBSP 97542); **São Paulo:** São Paulo, Campus USP, Mata do Cuaso (23°33'54.9432"S - 46°43'43.9608"W), ♂, 16-23.XI.1999, D. F. Candiani leg. (IBSP 42190); Campus Instituto Butantan, Mata do Butantan (23°33'59.493"S - 46°43'14.7612"W), ♂, 12-19.XII.1999, D. F. Candiani leg. (IBSP 68979).

### *Triaeris stenaspis* Simon, 1891

(Figs 13, 14, 60)

*Triaeris stenaspis* SIMON, 1891:561. Syntype female from Saint Vincent in MNHN and females syntypes from same locality in the BMNH; PLATNICK *et al.*, 2012a:22, figs 1-40.

*Triaeris patellaris* BRYANT, 1940:268, fig. 4. Female holotype from Soledad, Cienfuegos, Cuba, in MCZ; examined; CHICKERING, 1968b:358, figs 14-18 (Syn.).

*Triaeris berlandi* LAWRENCE, 1952:5, figs 3a-f. Female holotype from Thysville [= Mbanza-Ngungu], DR Congo, in Musée Royal de l'Afrique Centrale, Tervuren; PLATNICK *et al.*, 2012a:23 (Syn.).

*Triaeris lepus* SUMAN, 1965:235, figs 27-31. Female holotype from Puu Papaa Peak, Kaneohe, Oahu, Hawaii, in Bishop Museum; PLATNICK *et al.*, 2012a:23 (Syn.).

*Triaeris lacandona* BRIGNOLI, 1974a:208, figs 4A-F. Female holotype from Cueva de Yaxchilan, Peten, Guatemala, depository unknown; BRIGNOLI, 1975a:34, figs 1H-J; PLATNICK *et al.*, 2012a:23 (Syn.).

**Diagnosis.** *Triaeris stenaspis* is known only by females, easily recognized by their orange color, genitalia occupying the greater part of the postepigastric scutum, with a median longitudinal slit and internally with sinuous and distally squiggled duct, visible by transparence (Fig. 14).

**Description.** Female, see PLATNICK *et al.*, 2012a:22, figs 1-40

**Biology.** The taxonomic status of the pantropical goblin spider *Triaeris stenaspis* was defined by PLATNICK *et al.* (2012a). Presently, this species is well established in

many parts of the world (KOPONEN, 1997; KIELHORN, 2008; KORENKO *et al.*, 2009; PLATNICK *et al.*, 2012a) and apparently sustained populations are present in various continents. KORENKO *et al.* (2009) showed that at least one population from Czech Republic is parthenogenetic. Since no males were found with sampled females so far, including the 200 records here reported, parthenogenesis might actually be true for all known populations of this species. *Triaeris stenaspis* occur from north to south Brazil, being very common in the ground at urban parks. The invasion of this species in Brazilian territory might occurred long ago, giving its wide distribution in the country and the fact that it is not strictly peri-domiciliar, collected as well in forest litter and even in natural caves, but the first known record of this species in Brazil dates as recently as 1996. The large number of records reported here from natural caves in the states of Tocantins, Goiás and Minas Gerais might be cause for conservation concerns. Since this species appears to be a dominating faunistic component in those cavities, it may represent danger for autochthones species in these sensible environments.

**Distribution.** This Pantropical species have a wide distribution range in Brazil, occurring mainly in urban areas (Fig. 60). PLATNICK *et al.* (2012a) hypothesized that *Triaeris* belongs to a West African sub-group of the *Zyngoonops* group of genera, a large putatively monophyletic lineage with representatives in both the Neotropics and Africa. Under this assumption, *Triaeris stenaspis* would be an African species in origin, which attained a Pantropical distribution.

**Material examined.** BRAZIL, **Acre:** Senador Guiomard, Reserva Extrativista Catuaba (10°12'51.4584"S - 67°45'17.1216"W), 5♀, 2002, E. Morato *et al.* leg. (IBSP 86836, 86850, 86873, 86879, 86920); **Pará:** Uruará (03°43'4"S - 53°44'13"W), 2♀, 2014, R. L. Cajaiba leg. (IBSP 166581); **Tocantins:** Dianópolis, Gruta Vozinha (11°27'36"S - 46°51'00"W), 3♀, 04-09.XII.2007, R. Andrade *et al.* leg. (IBSP 97622); Gruta Areia (11°27'36"S - 46°51'00"W), 4♀, 04-09.XII.2007, R. Andrade *et al.* leg. (IBSP 97640); Gruta 3 Morros III (11°27'36"S 46°51'00"W), ♀, 21-29.V.2008, F. Pellegatti Franco *et al.* leg. (IBSP 134439); **Goiás:** São Domingos, Lapa de São Bernardo II (16°34'31.26"S - 49°10'32.307"W), 5♀, 18.IX.1997, P. Gnaspini *et al.* leg. (IBSP 23729, 23730); Lapa de Terra Ronca II (16°34'31.26"S - 49°10'32.307"W), 2♀, 20.IX.1997, P. Gnaspini *et al.* leg. (IBSP 23741, 23743); 3♀ 4 imm., 19.IX.1997, P. Gnaspini *et al.* leg. (IBSP 23746); Parque Estadual de Terra Ronca, Caverna Angélica (16°34'31.26"S - 49°10'32.307"W), 9♀, 6 imm., 06-09.IX.2000, F. P. Franco leg. (IBSP 26050, 26053, 26054, 26070, 26072); Parque Estadual de Terra Ronca, Caverna Passa Três (16°34'31.26"S - 49°10'32.307"W), 7♀, 5 imm., 03-05.IX.2000, C. A. Rheims & F. P. Franco leg. (IBSP 26064, 26077, 26079, 26080); Morrinhos, Parque Ecológico Jatobá Centenário (17°43'36"S - 49°07'55"W), 2♀, XII.2006 - VIII.2007, R. C. Santana leg. (IBSP 140921); **Mato Grosso,** Primavera do Leste, Gruta KNL-1-S1EMAL004 (181326E - 8383239mN), ♀, 12-20.VIII.2014, Equipe Spelayon leg. (IBSP 166573); **Mato Grosso do Sul:**

Campo Grande, Reserva UCDB (20°24'50.0256"S - 54°36'55.854"W); 2♀, 07.IV.1997, K. O. Vieira leg. (IBSP 38355); Inocência, Fazenda Lagoinha (19°17'03"S - 51°03'06"W), ♀, 18.XI.2004, J. Raizer leg. (IBSP 68030); Anastácio, Fazenda Jatiuca (20°31'54"S - 55°50'27"), ♀, 16-26.II.2008, R. Bessi leg. (IBSP 127638); Ivinhema (22°31'19.1856"S - 53°50'49.1676"W), 34♀, X.2004-XII.2006, K. Anjos leg. (IBSP 92829, 92833, 92839-92866); **Bahia:** Jaguaripe (13°10'01"S - 39°00'04"W), ♀, 02.III.2011, C. M. P. Leite leg. (IBSP 166609); Wenceslau Guimarães (13°34'32"S - 39°42'25"W), ♀, 21.I.2011, C. M. P. Leite leg. (IBSP 166610); Salvador, Baía do Aratu (12°47'32.8"S - 38°28'15.3"W), 5♀, 2008-2009, K. Benati (IBSP 126081, 126090, 126101, 126103, 126109); Ponte do Fernandinho (12°47'32.8"S - 38°28'15.3"W), 9♀, 13.IX.2004, K. Benati leg. (IBSP 58840-58841, 58860, 58893; 58897, 58902, 58918, 58939, 59023); Itapetinga, Parque Municipal da Matinha (15°15'43.8618"S - 40°17'44.9376"W), ♀, III-IV.2003, J. P. S. Alves leg. (IBSP 66524); Una, Reserva Ecológica de Una (15°17'48"S - 39°04'28"W), 4♀, X.1999-IX.2000, M. F. Dias leg. (IBSP 64221, 65268, 65350); **Minas Gerais:** Prudente de Morais, Fazenda Sapé (19°28'8.2344"S - 44°5'31.7328"W), 2♀, 13.X.2001, É. S. S. Álvares leg. (IBSP 44224, 44232); Uberlândia, Estação Ecológica do Panga (19°11'10"S - 49°23'30"W), 3♀, IV.2005-II.2006, M. F. Mineo leg. (IBSP 92663-92665); Belo Horizonte, Estação Ecológica da UFMG (19°52'S - 43°58'W), 8♀, I.2001, É. S. S. Álvares leg. (IBSP 66796, 66798, 66800); Estação Ecológica da UFMG, Mata do Sossego (19°52'S - 43°58'W), ♀, II.2001, É. S. S. Álvares, E. O. Machado & C. S. Azevedo leg. (IBSP 68573); Belo Horizonte, COPASA (20°10'S - 44°21'W), ♀, IV.2002, Equipe Biota (IBSP 136439); Jaboticatubas, Serra do Cipó, Parque Nacional da Serra do Cipó (19°15'S - 43°31'W), 40♀, 2002-2003, Equipe Biota leg. (IBSP 68258-68294); Mariana, Gruta FN-12 (20°10'18.3684"S - 43°25'13.029"W), 2♀, 16-21.I.2009, R. Andrade *et al.* leg. (IBSP 146210, 146216); Mina Alegria (20°10'18.3684"S - 43°25'13.029"W), 2♀, 16-17.IV.2009, R. Bessi *et al.* leg. (IBSP 149764); Prados, Gruta S3\_HOL\_010 (596862E - 7657188N), ♀, 29.VII.2014, L. T. Fonseca leg. (IBSP 166568); Córrego Fundo: Gruta MC\_SM\_015 (20°26'32"S; 45°35'57"W), ♀, 19-21.V.2014, Santos *et al.* leg. (IBSP 166569); Gruta MC\_SM\_016 (20°26'33"S; 45°35'56"W), ♀, 08-20.I.2014, Equipe Spelayon leg. (IBSP 166570); Pains (20°22'15"S; 45°39'39"W): ♀, 31.VII.2014, F. Bondezan leg. (IBSP 166571); Gruta Loca dos Negros (20°26'06"S; 45°39'33"W), ♀, 20.III.2009, R. Zampaulo leg. (IBSP 166584); Gruta Tamafi (20°23'39"S; 45°40'48"W), ♀, 09.III.2009, R. Zampaulo leg. (IBSP 166585); Gruta SPA\_015 (20°17'53"S; 45°25'38"W), ♀, 24.I.2014, M. Barcelos leg. (IBSP 166587); Cave SPA\_030 (20°18'24"S; 45°35'48"W), ♀, 05.II.2014, N. T. Pimentel & T. F. Ferreira leg. (IBSP 166588); Cave SPA\_037 (20°18'26"S; 45°35'45"W), ♀, 05.II.2014, F. O. Borges & M. Barcelos leg. (IBSP 166589); Cave SM-68 (20°20'47"S; 45°36'09"W), 2♀, 12.III.2012, Equipe Spelayon leg. (IBSP 166590); Cave SM-180 (20°22'21"S;

45°35'34"W), ♀, 13.II.2014, Equipe Spelayon leg. (IBSP 166591); Cave SM-287 (20°21'46"S; 45°35'51"W), ♀, 20.II.2014, Equipe Spelayon leg. (IBSP 166592), Cave SM-103 (20°21'40"S; 45°36'12"W), ♀, 18.VIII.2012, Equipe Spelayon leg. (IBSP 166596); Cave SM-246 (20°21'51"S; 45°35'46"W), ♀, 20.II.2014, Equipe Spelayon leg. (IBSP 166597); Cave SM-42 (20°21'50"S; 45°35'58"W), 2♀, 21.III.2012, Equipe Spelayon leg. (IBSP 166593); Cave SM-258 (20°21'52"S; 45°35'43"W), 2♀, 19.IX.2013, Equipe Spelayon leg. (IBSP 166602); Cave SM-153 (20°22'03"S; 45°35'38"W), ♀ (IBSP 166594); Cave SM-236 (20°22'11"S; 45°35'51"W), ♀ (IBSP 166595); Cave SM-178 (20°22'20"S; 45°35'23"W), ♀ (IBSP 166598); Cave SM-205 (20°22'23"S; 45°35'40"W), ♀ (IBSP 166599); Cave SM-157 (20°21'53"S; 45°35'39"W), ♀ (IBSP 166600); Cave SM-195 (20°22'23"S; 45°35'37"W), ♀ (IBSP 166601), all collected in 13.II.2014 by Equipe Spelayon; Matozinhos, Cave 100, 4♀, VI.2014, F. Bondezan leg. (IBSP 166575 - IBSP 166578); Cordisburgo, Gruta da Morena (19°8'25.584"S - 44°21'20.5272"W), ♀, 12.IX.2001, P. Gnaspini leg. (IBSP 71833); Lavras (21°13'43.9638"S - 44°58'1.6608"W), 17♀, 16-19.VI.2000, M. Andreizza leg. (IBSP 86329-86330, 86337-86339, 86342, 86346-86347, 86349-86350, 86352-86355, 86357, 86359, 86361); ♀, 2000, A. Almeida leg. (IBSP 121445); **Rio de Janeiro**: Rio de Janeiro, Bairro Laranjeiras, Parque Guinle (22°55'50.38"S - 43°11'1.12"W), 2♀, 20.X.2001, R. Baptista leg. (MNRJ 3759); Volta Redonda, Floresta da Cicuta (22°33'2.016"S - 44°5'31.7328"W), ♀, 11.III.2002, E. Folly leg. (IBSP 39605); 4♀, 04-11.III.2002-11-18.VI.2001, Equipe Biota (IBSP 136199-136202); Pinheiral, Fazenda Santa Helena, Fazenda Regional de Criação (Posto Zootécnico) (22°34'S - 44°21'W), ♀, 05.11.XI/1999, A. D. Brescovit leg. (IBSP 66897); **São Paulo**: Jacaref, Campus Vila Branca, Univap (23°12'28.7886"S - 45°56'52.746"W), ♀, 2007, N. M. C. Velho leg. (IBSP 143942); Cotia, Caucaia do Alto, Reserva do Morro Grande (23°40'52.4742"S - 46°59'25.4472"W), 2♀, III.2003, Equipe Biota leg. (IBSP 136732, 136733); São Paulo, Campus USP, Mata do Cuaso (23°33'54.9432"S - 46°43'43.9608"W), 16♀, 14-21.IX.1999, D. Candiani leg. (IBSP 69027-69032, 69034, 69036-69040); 10♀, 16-23.II.2000, D. F. Candiani leg. (IBSP 42154, 42156, 68768-68769); Parque da Previdência (23°34'47.9886"S - 46°43'38.8236"W), 19♀, 16-23.VIII.1999, D. F. Candiani leg. (IBSP 42159, 42162, 68785, 68787, 68797, 68806); 146♀, 14-21/IX/1999-19/XII/1999, D. Candiani leg. (IBSP 69041-69047, 69049-69082, 69084-69103); ♀, V.2000 - II.2001, D. F. Candiani (IBSP 76545); 7♀, 2000-2001, D. F. Candiani *et al.* leg. (IBSP 117046); 5♀, 1999, D. Candiani leg. (IBSP 69083, PBI\_OON 10912); Parque dos Príncipes (23°34'24.0162"S - 46°46'17.5944"W); ♀, 05-12.X.2004, P. A. M. Goldoni & S. Guizze leg. (IBSP 48525); Reservatório Guarapiranga, Jardim Ângela, Ilha dos Eucaliptos (23°45'9.5286"S - 46°44'23.3082"W), 17♀, 07-13.X.2003, I. Cizauskas & C. R. M. Garcia leg. (IBSP 61400-61413); Parque Ecológico Guarapiranga, 2♀, 2009, I. Cizauskas leg. (IBSP 155047-155048); Parque do Estado (23°39'8.625"S

- 46°36'57.9924"W), ♀, 21-28.I.2003, J. Valvassori leg. (IBSP 68057); Itapevi, Condomínio TransUrb (23°35'28.284"S - 46°58'29.121"W), ♀, 30.IV.1999, V. C. Onofrio leg. (IBSP 67591); Louveira (23°05'11"S - 46°57'02"W), 2♀, 16.I.1996, A. E. C. Farinha leg. (IBSP 12067); Pindamonhangaba (22°55'26"S - 45°27'42"W), ♀, 08-10.IV.1998, R. Martins & I. Knysak leg. (IBSP 20066); Peruíbe, Estação Ecológica Juréia-Itatins (24°33'015"S - 7°13'292"W), 2♀, 26.IX-03.V.1999, A. D. Brescovit *et al.* leg. (IBSP 25696, 25791); Iporanga, Parque Estadual Turístico Alto do Ribeira, (24°30'1.3968"S - 48°35'8.415"W), ♀, R. Andrade leg. (IBSP 29414); Dois Córregos, Sítio Guedes (22°19'5.3436"S - 48°21'45.6912"W), ♀, 01-06.IV.2002, G. Q. Romero leg. (IBSP 34974); Teodoro Sampaio, Parque Estadual do Morro do Diabo (22°31'S - 52°18'W), ♀, 24-31.III.2003, Equipe Biota leg. (IBSP 60189); Apiaí, Parque Estadual Turístico Alto do Ribeira, Núcleo Santana (24°27'36"S - 48°36'0"W), 4♀, 08-14.XII.2002, Equipe Biota leg. (IBSP 136440, 136515-136517); Jaboticabal (21°14'54.6288"S - 48°16'9.7278"W), 8♀, III-IV.2004 (IBSP 55862, 55909); Ubatuba, Parque Estadual da Ilha Anchieta (23°32'S - 45°03'W), 2♀, 23-30.VII.2001, Equipe Biota leg. (IBSP 56408, 56413); **Paraná**: Foz do Iguaçu, Parque Nacional de Foz do Iguaçu (25°36'S - 54°25'W), 8♀, 03-12.III.2002, Equipe Biota leg. (IBSP 60249, 60256, 136460-136465); Londrina, Parque Estadual Mata dos Godoy (23°27'1.0656"S - 51°14'11.0646"W), 6♀, 13.IV.1999-07.II.2000, J. Lopes leg. (IBSP 38208, 38240, 38258, 38266, 38269); Cornélio Procópio, Parque Estadual Mata São Francisco (23°9'30.2826"S - 50°33'52.416"W), 3♀, 08.V.2009, N. G. Cípola leg. (IBSP 150237, 150289, 150294); **Santa Catarina**: Brusque (27°05'53"S - 48°55'03"W), Bairro Bateias, campus Empresa Buettner, 10♀, 2014, D. L. Ronchi leg. (IBSP 166603-166608); Blumenau, Parque Natural Municipal Nascentes do Rio Garcia (26°55'10.7862"S - 49°05'36.5748"W), ♀, 23.I.2005, R. C. Francisco leg. (IBSP 68394); Palhoça (27°38'42"S - 48°40'04"W), Serra do Tabuleiro, ♀, I.2003, Equipe Biota leg. (IBSP 136435); Paulo Lopes, Parque Estadual da Serra do Tabuleiro (27°55'S - 48°42'W), ♀, 10-20.I.2003, Equipe Biota leg. (IBSP 60491); Santo Amaro da Imperatriz (27°41'0.9018"S - 48°43'50.9442"W), ♀, 2004, J. Steiner leg. (IBSP 91114).

### *Xestaspis parmata* Thorell, 1890, comb. rest.

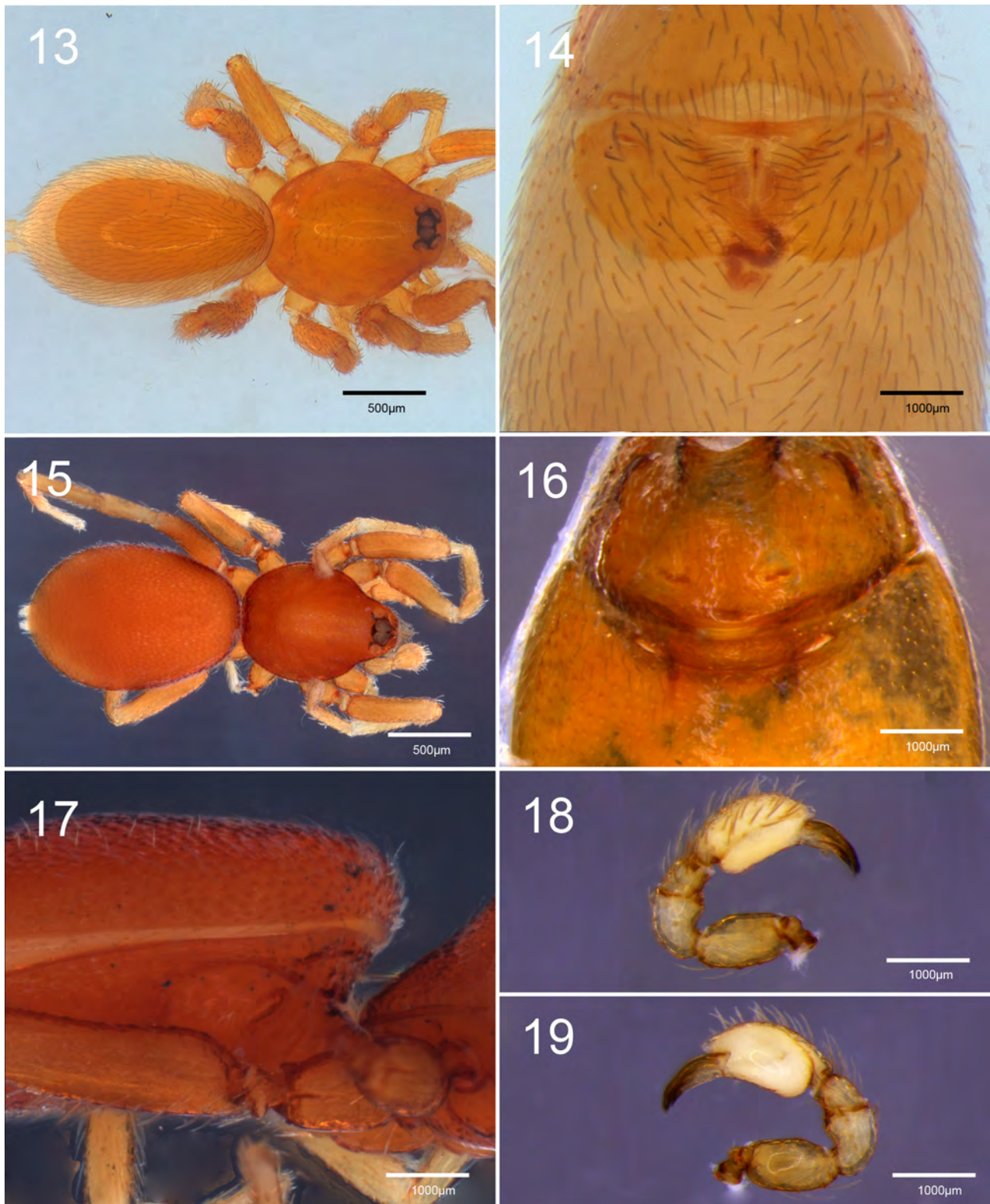
(Figs 15-19, 59)

*Xestaspis parmata* THORELL, 1890:391 (Holotype female from Ajer Mancior, Sumatra, Cel. Beccari leg., deposited in Museu Civico di Storia Naturela Giacomo Doria, examined).

*Hytanis oblonga* SIMON, 1893b:440 (Syntypes: one female, labeled "6537 *Hytanis oblonga* E. S. from Iquitos (de M.)", deposited in MNHN, AR 5692, PBI\_OON 4743, examined; one female deposited in MNHN, AR 5681, PBI\_OON 4742, labeled "14080 *Hytanis oblonga* E. S.; Free Town (Mq)", examined). **Syn. nov.**

*Gamasomorpha paramata*: SIMON, 1893a:301 (lapsus).

*Xestaspis bipeltis* THORELL, 1895:16 (Holotype male from Burma, Tharrawady & Oates col., in BMNH, 1895.9.21.43, PBI\_OON 25537, examined). **Syn. nov.**



Figs 13-19. Figs 13, 14, *Triarhis stenaspis* Simon, 1891, female: 13, dorsal; 14, genitalia, ventral. Figs 15-19, *Xestaspis parmata* Thorell, 1890, comb. rest., female: 15, dorsal; 16, genitalia, ventral; 17, abdomen, anterior, lateral. Male: 18, palp prolateral; 19, palp retrolateral.

*Gamasomorpha insularis* SIMON, 1907:231 (Syntypes: one female from Fernando Poo Island (presently Bioko), Gulf of Guinea, deposited in MNHN B524-23037 and one female from Mauritius Islands, Alluaud leg., deposited in MNHN B524-19782); BENOIT, 1977:38, figs 14a-e (Male "allotype", from Santa Helene, N. Rupert's Valley, 28/12/1965, P.

Basilaewsky, P. Benoit & N. Lelelup col., in MRAC 129.192; PBI\_OON 25595, examined); SAARISTO, 2001:311-314, figs 1A, 2A, 3A; 4-8); FREITAS *et al.*, 2013:15. **Syn. nov.**  
*Opopaea lutzi* PETRUNKEVITCH, 1929:61, figs 42-50 (Holotype male from Desecheo Island, Puerto Rico, West Indians, Feb. 18-20, 1914, in

AMNH 3507A; PBI\_OON 5512, examined); CHICKERING, 1969:153, figs 22-27. **Syn. nov.**

*Gamasomorpha bipeltis*: ROEWER, 1942:285.

*Gamasomorpha parmata*: ROEWER, 1954:1707.

*Gamasomorpha lutzii*: BRIGNOLI, 1975b:228; BURGER, 2011:149, figs 1C-E, 3F-H, 4A-B.

*Gamasomorpha madeirensis* WUNDERLICH, 1987:66, figs 40-43 (Holotype male and paratype female from Canico bei Funchal, Madeira Island, Dec. 1988, J. Wunderlich col., in SMF36925 and 36926, respectively); SAARISTO, 2001:311 (Syn. with *G. insularis*).

**Note.** All species here synonymized had their types examined and no morphological differences were found in relation to *Xestaspis parmata*. Simon's original label of *Hytanis oblonga* was removed from the vial with syntypes received from MNHN; the current labels were quite recently printed. SIMON (1893a) doesn't refer to specimens from Iquitos or Free Town (Africa?), however there is no other material of this species available at MNHN. Since *H. oblonga* is the type (and currently the sole) species of *Hytanis*, the genus is bound to be considered a junior synonym of *Xestaspis* (**Syn. nov.**).

**Diagnosis.** Specimens of *Xestaspis parmata* resembles those of *Brignolia* and *Opopaea* by the shape of the body, but can be distinguished from the species of these genera by the red coloration of the carapace and abdomen (Figs 15-17), larger size, less sclerotized palp and bulb, and, particularly from *Opopaea*, by the less enlarged patella.

**Description.** Male and female, see WUNDERLICH, 1987:66, figs 40-43 sub *Gamasomorpha madeirensis*.

**Distribution.** This species was probably introduced from south Asia, and is known to occur in just four areas in Brazil, in the north of the state of Piauí and in the Archipelago de Fernando de Noronha, which belongs to the state of Pernambuco, and from states of Minas Gerais and São Paulo (Fig. 59). It is an open question if this species has already managed to establish viable populations in the Brazilian territory.

**Material examined.** BRAZIL. **Piauí:** Parnaíba, Tabuleiros Litorâneos do Piauí (02°55'S - 41°50'W), Distrito de Irrigação, ♂, 2008, J. A. S. Santos col. (IBSP 144005); **Pernambuco:** Fernando de Noronha, Arquipélago de Fernando de Noronha (3°50'57.2814"S - 32°24'10.7922"W), ♂, ♀, IV/2006, G. Freitas col. (IBSP 67706-67707); **Minas Gerais:** Governador Valadares (18°51'03"S - 41°56'56"W), 15.X.1981, L. N. Sorkin, ♀ (AMNH, PBI\_OON 1167); **São Paulo:** São Paulo (23°33'S - 46°38'W), ♂, 2♀ (AMNH, PBI\_OON 37857).

### *Ischnothyreus peltifer* (Simon, 1891)

(Figs 20-25, 59)

*Ischnaspis peltifer* SIMON, 1891:562 (four female syntypes from Saint Vincent, Antilles, in BMNH, and one female syntype from St. Vincent, in MNHN).

*Ischnothyreus peltifer*: SIMON, 1893c:298 (not male, fig. 264, = *Camptoscapbiella simoni* Baehr); PLATNICK *et al.*, 2012b:7, figs 1-99.

*Dysderina antillana* BRYANT, 1942:324, figs 1, 7 (male holotype from Christiansted, Saint Croix, Virgin Islands, in MCZ; examined); CHICKERING, 1968a:80 (Syn.)

*Ischnothyreus omus* SUMAN, 1965:226, figs 1-8 (male holotype from Kailua, Oahu, Hawaii, in Bishop Museum, Honolulu). SAARISTO, 2001:345 (Syn.).

*Ischnothyreus formosus* BRIGNOLI, 1974b:80, figs 12-18 (male holotype from Akau, Taiwan, in Zoological Museum, Hamburg); SAARISTO, 2001:345 (Syn.).

*Ischnothyreus sechellorum* BENOIT, 1979:208, figs 7A-E (female holotype from Mahe, Seychelles Islands, in MRAC); SAARISTO, 1999:3 (Syn.).

**Diagnosis.** Males of *Ischnothyreus peltifer* can easily be separated from those of *I. velox* by the large protuberance on the base of the cheliceral fang in males (Fig. 23) and narrow, posteriorly situated, sinuous ridge on the postepigastric scutum (Fig. 22) in females.

**Description.** Male and female, see PLATNICK *et al.* 2012b:7, figs 1-99.

**Distribution.** According to PLATNICK *et al.* (2012b), this Pantropical species was probably introduced in Americas from Asia. Our records indicated that it was present only in three Brazilian states, the Northern State of Pará, the Northeastern State of Bahia and in the Southeastern State of São Paulo (Fig. 59). Judging by the records obtained, at the moment this species is not sympatric with *I. velox* in Brazil.

**Material examined.** BRAZIL. **Pará:** Belém, Ilha de Cotijuba (01°14'51.44"S - 48°32'47.14"W), ♀, XI.2004, L. Macambira leg. (MPEG 10271, PBI\_OON 40724); Bosque Rodrigues Alves (01°25'49"S - 48°27'23"W), 2♀, XI/2004, J. Barreiro leg. (MPEG 10478, 10479; PBI\_OON 40726, 40727); Belém, Mata de Várzea, ♂, 2♀, X.2004, L. Macambira leg. (MPEG 10709, PBI\_OON 40692); **Bahia:** Salvador, Baía do Aratu (12°47'32.8"S - 38°28'15.3"W), 3♂, ♀, 2007-2009, K. Benati leg. (IBSP 126042, 126086, 126131, 126135); Presidente Tancredo Neves (13°24'32"S - 39°20'05"W), 2♀, 07.II.2011, C. M. P. Leite (IBSP 166612, 166614); Itamaraju (16°57'58"S - 39°26'01"W), ♀, 14.IV.2011, C. M. P. Leite leg. (IBSP 166613); **São Paulo:** Ubatuba, Parque Estadual da Ilha Anchieta (23°32'S - 45°03'W), ♂, 23-30.VII.2001, Equipe Biota leg. (IBSP 69881).

### *Ischnothyreus velox* Jackson, 1908

(Figs 26-30, 59)

*Ischnothyreus velox* JACKSON, 1908:51, pl. 4, figs 9-13 (Lectotype male from greenhouse in Chester, England, in BMNH); SAARISTO, 2001:347 (desig. lectotype); PLATNICK *et al.*, 2012b:27, figs 100-132.

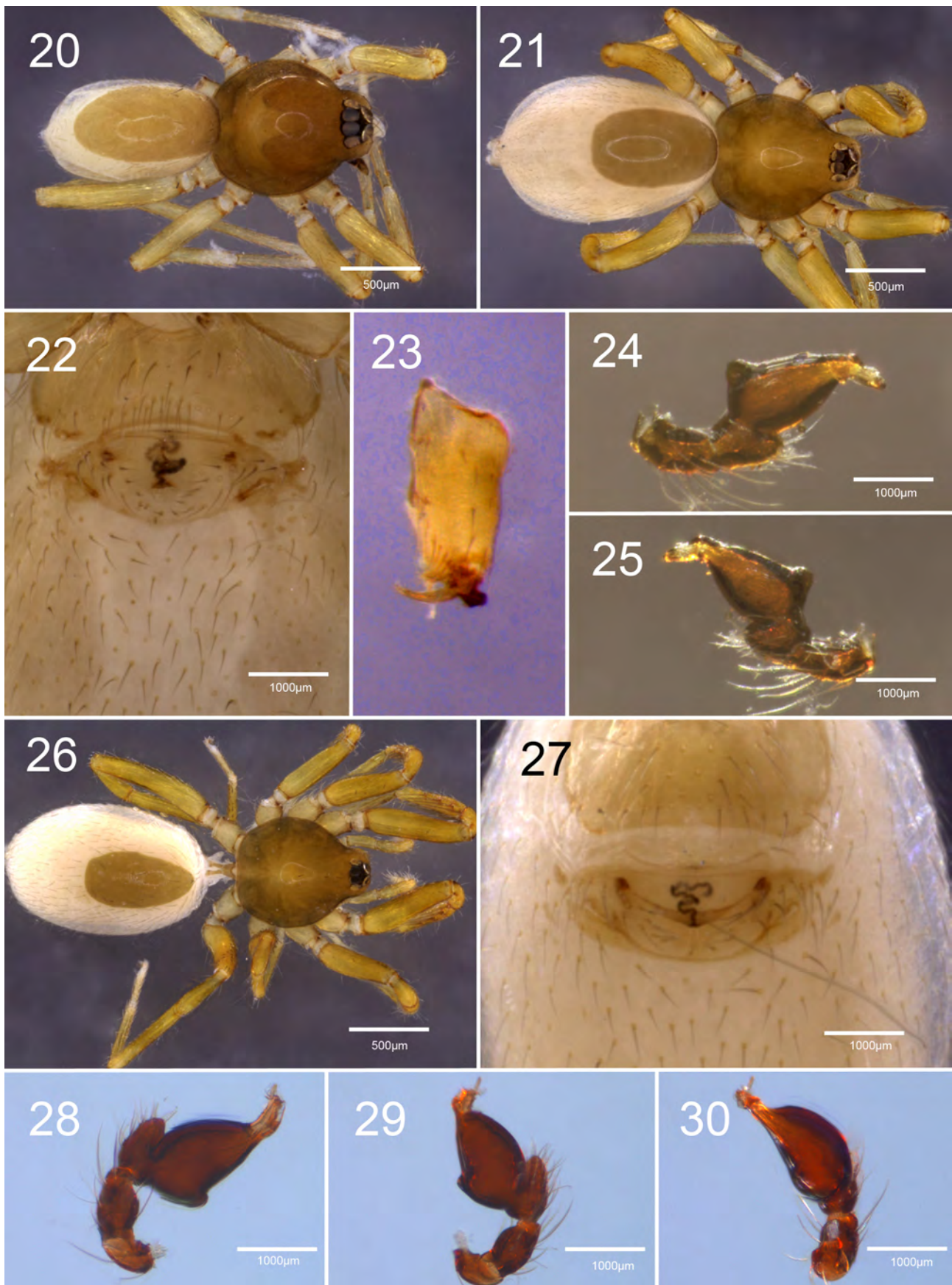
*Ischnothyreus barrowsi* CHAMBERLIN & IVIE, 1935:9, pl. II, fig. 7 (female holotype from Marco Island, Collier Co., Florida, USA, in AMNH). PLATNICK *et al.*, 2012b:27 (Syn.).

*Ischnothyreus indressus* CHICKERING, 1968a:84, figs 13-20 (male holotype from Nevis, Leeward Islands, in MCZ; examined); PLATNICK *et al.*, 2012b:27 (Syn.).

**Diagnosis.** *Ischnothyreus velox* differ from *I. peltifer* by the males lacking a protuberance on the base of the fang; palps with rounded bulb and sperm pore larger than that of *I. peltifer* (Figs 28-30). Females of *I. velox* have a distinctively procurved ridge occupying most of the width of the postepigastric scutum (Fig. 27).

**Description.** Male and female, see PLATNICK *et al.* 2012b:27, figs 100-132.





Figs 20-30. Figs 20-25, *Ischnothyreus peltifer* (Simon, 1891), male: 20, dorsal; 23, projection of chelicerae; 24, palp prolateral; 25, palp retrolateral. Female: 21, dorsal; 22, genitalia, ventral. Figs 26-30, *Ischnothyreus velox* Jackson, 1908, female: 26, dorsal; 27, genitalia, ventral. Male: 28, palp retrolateral; 29, palp prolateral; 30, palp ventral.

Distribution. According to PLATNICK *et al.* (2012b), who recorded this species in the state of Amazonas, this Pantropical species was probably introduced in Americas from Asia. The present records in Brazil are anecdotic but depict a wide distribution in the Western and Central Amazonia, as well as in the Northeastern state of Sergipe (Fig. 59).

Material examined. BRAZIL. **Acre:** Senador Guiomard, Reserva Extrativista Catuaba (10°12'51.4584"S - 67°45'17.1216"W), ♀, 2002, E. Morato *et al.* leg. (IBSP 86903); **Amazonas:** Benjamin Constant (04°22'58"S - 70°1'51"W), ♀, 2010, P. S. Pompeu *et al.* leg. (IBSP 166574); Coari, Rio Urucu, Base de Operações Geólogo Pedro de Moura (04°52'16"S - 65°20'04"W), ♂, ♀, 11-20.VII.2003, A. B. Bonaldo, J. Dias & D. Guimarães leg. (MPEG 10212, 10214; PBI\_OON 40693, 40725); **Sergipe:** Simão Dias, Toca da Raposa (10°43'46.992"S - 37°50'54.9702"W), 2♀, 20.VI.2014, no col. (IBSP 166579-166580).

### *Opopaea concolor* (Blackwall, 1859)

(Figs 31-35, 61)

*Oonops concolor* BLACKWALL, 1859:265 (Male and female syntypes from Madeira Island, Portugal, should be in Hope Department of Oxford University).

*Opopaea concolor*: KULCZYŃSKI, 1899:339, pl. 6, fig. 25; PLATNICK & DUPÉRRÉ, 2009a:22, figs 74-104.

*Myrmecoscapbiella borgmeyer* MELLO-LEITÃO, 1926:2 (Male holotype from Rodeio (26°55'22"S 49°21'59"W), Santa Catarina, Brazil, 1926, T. Borgmeyer col., in MNRJ 953); PLATNICK & DUPÉRRÉ, 2009a:22 (Syn.).

*Gamasomorpha humilis* MELLO-LEITÃO, 1920:172 (Female holotype from Pinheiro (actually Pinheiral), Barra do Pirai, Rio de Janeiro, Brazil, in MNRJ 955; PBI\_OON 10963, examined). **Syn. nov.**

*Opopaea devia* GERTSCH, 1936:5, fig. 13 (Female holotype from Edinburg, Hidalgo Co., Texas, USA, in AMNH); PLATNICK & DUPÉRRÉ, 2009a:22 (Syn.).

*Opopaea guaraniana* BIRABÉN, 1954:203, figs 30-36, 50 (Female holotype and male allotype from Manantiales, Corrientes, Argentina, lost); PLATNICK & DUPÉRRÉ, 2009a:22 (Syn.).

*Opopaea bandina* CHICKERING, 1969:147, figs 1-3 (Female holotype from Largo, Pinellas Co., Florida, USA, in MCZ); PLATNICK & DUPÉRRÉ, 2009a:22 (Syn.).

*Gamasomorpha atlantica* BENOIT, 1977:35, figs 13a-e (Male holotype from Prosperous Bay Plain, Saint Helena Island, in MRAC); SAARISTO & MARUSIK, 2008:20 (Syn.).

*Opopaea atlantica*: BRIGNOLI, 1983:188.

Diagnosis. *Opopaea concolor* can be distinguished from those of *O. deserticola* by the ventrally unexpanded palpal bulb in males (Figs 34, 35; PLATNICK & DUPÉRRÉ, 2009a, figs 99-104), and by having the small, dark knob marking the origin of the receptaculum situated quite far from the epigastric furrow in females (Fig. 33).

Description. Male and female, see PLATNICK & DUPÉRRÉ, 2009a:22, figs 73-104.

Distribution. According to PLATNICK & DUPÉRRÉ (2009a), this Pantropical species was probably introduced in Americas from the Old World. In the New World it is known to occur from the southern United States and Bahamas Islands south to Argentina. This species is synanthropic in Brazil and in many cases sympatric with *O. deserticola*, mainly in the southern portions of the country (Fig. 61). It is normally found inside houses and buildings, but can be also found in

caves, especially those most intensely disturbed by human activities.

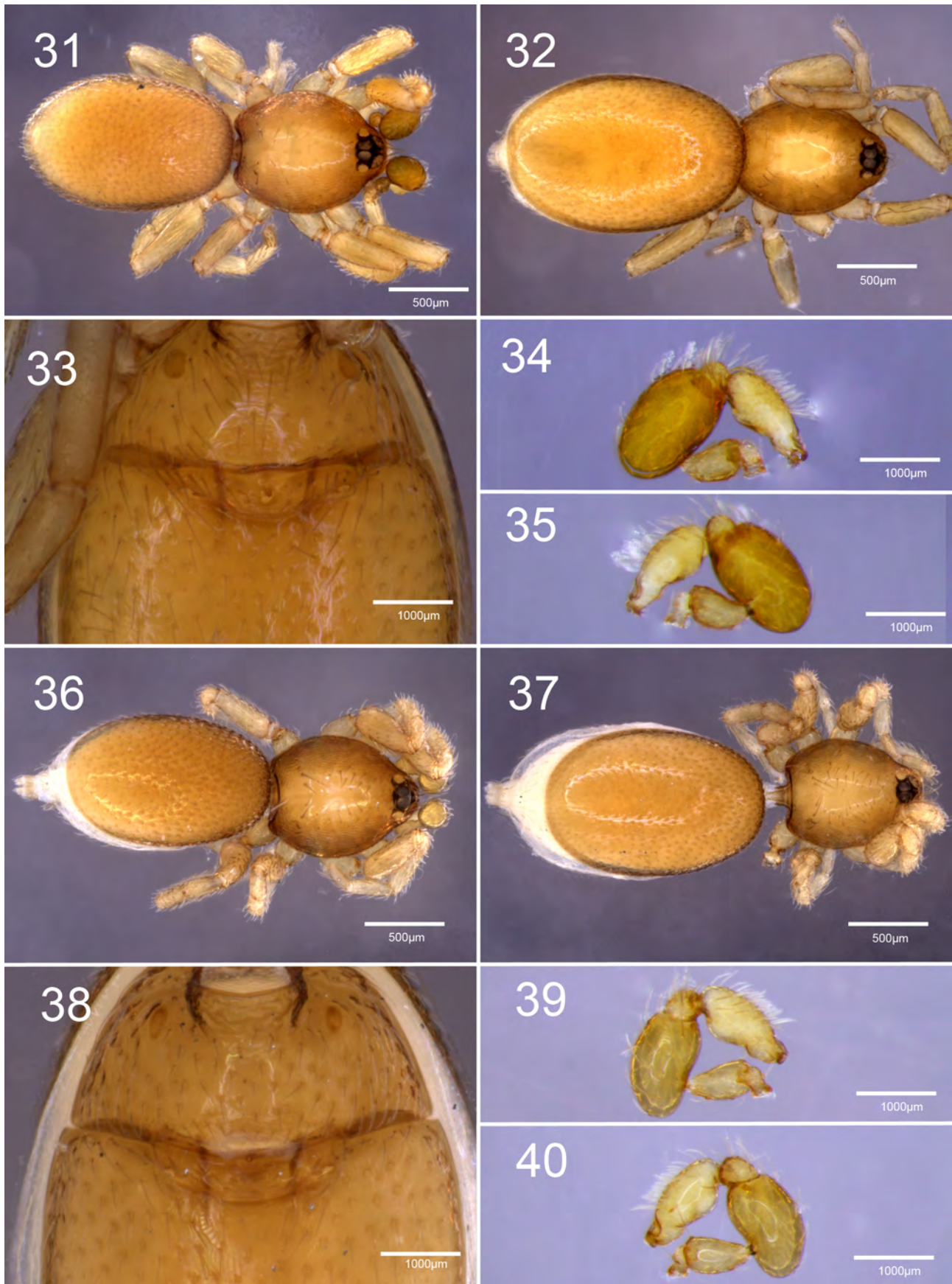
Material examined. BRAZIL, **Bahia:** Lajedinho (12°21'21"S - 40°54'21"W), Gruta da Mocozeira, ♀, 22.X.2014, F. Bondezan leg. (IBSP 166572); Presidente Tancredo Neves (13°22'44"S - 39°19'31"W), ♀, 07.II.2011, C. M. P. Leite leg. (IBSP 166611); **Mato Grosso do Sul:** Corumbá, Pantanal, Passo da Lontra (18°57'31.0674"S - 57°39'52.5918"W), ♀, I.1998-XI.1999, J. Raizer leg. (IBSP 68454); **Minas Gerais:** Jaboticatubas, Serra do Cipó, Parque Nacional da Serra do Cipó (19°15'S - 43°31'W), ♂, 03-09.I.2003, Equipe Biota leg. (IBSP 69444); Belo Horizonte, Campus da UFMG/ICB (19°52'S - 43°58'W), ♀, VII.2002, É. S. S. Álvares leg. (IBSP 73190); **Rio de Janeiro:** Teresópolis (22°26'26.2536"S - 43°5'13.9158"W), ♀, 28.XI.1976, Leny leg. (MNRJ 2094); Rio de Janeiro, Gávea (22°58'46.2324"S - 43°14'46.6224"W), 2♀, R. Arlé leg. (MNRJ 41760); **São Paulo:** São Paulo (23°32'51"S - 46°38'10"W), ♀, 13.II.2002, F. S. Cunha leg. (IBSP 33063); ♀, 20.IV.2007, G. Ruiz leg. (IBSP 122228); Parque da Previdência (23°34'49.7136"S - 46°43'38.4558"W), ♀, 17.XII.1999, D. F. Candiani leg. (IBSP 33007); Parque Alfredo Volpi (23°35'12.5766"S - 46°42'16.005"W), ♀, 19-23.III.2005, A. Bagio leg. (IBSP 59276); Campus da USP (23°33'44"S - 46°43'39"W), ♂, 13.V.2003, D. F. Candiani leg. (IBSP 131540); Mata do Cuaso (23°33'54.9432"S - 46°43'43.9608"W), ♀, 1999, D. Candiani leg. (IBSP 68981); campus Instituto Butantan, Mata do Butantan (23°33'59.493"S - 46°43'14.7612"W), ♂, 12-19.XII.1999, D. Candiani leg. (IBSP 68980); ♀, 18.VIII.2006, A. D. Brescovit leg. (IBSP 72788); Itapevi, Condomínio TransUrb (23°35'28.284"S - 46°58'29.121"W), ♀, 28.V.1999, V. C. Onofrio leg. (IBSP 67601); Campinas, Campus da Unicamp (22°48'45.9786"S - 47°4'12.198"W), ♂, 02.I.2006, A. J. Santos leg. (IBSP 72970); Guarujá, Ilha da Moela (24°02'57.5016"S - 46°15'48.0558"W), ♂, 17-19.VIII.2009, R. P. Indicatti & G. P. Perroni leg. (IBSP 144859); **Paraná:** Adrianópolis, Vale do Ribeira (24°47'0.726"S - 49°6'45.4962"W), ♀, 17.III.2005, D. M. B. Battesti leg. (IBSP 51581); Londrina, campus Universidade Estadual de Londrina, Fazenda Escala (23°20'23,45"5"S - 51°12'32,8"W), 2♂, VIII.2007, J. O. Fernando leg. (IBSP 118415, 118429); **Rio Grande do Sul:** Sapucaia do Sul (29°50'20"S 51°08'38"W), ♀, 20.VI.2013 (MCN 49333); Porto Alegre (30°01'58"S - 51°13'48"W), 5♀, 24.V.1981 - 21.II.1991, A. A. Lise, A. B. Bonaldo & L. Moura leg. (MCN 10350, 20746-20747, 28335); ♂, 6♀, 31.V.1935, P. Buck leg. (MNRJ 2090); Jardim Botânico (30°03'6.07"S - 51°10'37.95"W), ♂, 29.IV.1997, L. Moura leg. (MCN 28335); ♂, 25.III.2011, L. Moura leg. (MCN 49332).

### *Opopaea deserticola* Simon, 1891

(Figs 36-40, 61)

*Opopaea deserticola* SIMON, 1891:560, pl. 42, fig. 5 (female syntype from Saint Vincent, Antilles, in BMNH); PLATNICK & DUPÉRRÉ, 2009a:4, figs 1-72

*Gamasomorpha rufa* BANKS, 1898:211, pl. 13, fig. 12 (female holotype from Baja California, Mexico, G. Eisen & H. Vaslit, in the California Academy of Sciences, destroyed in the earthquake in 1900). **Syn. nov.**



Figs 31-40. 31-35. *Opopaea concolor* (Blackwall, 1859), male: 31, dorsal; 34, palp retrolateral; 35, palp prolateral. Female: 32, dorsal; 33, genitalia, ventral. Figs 36-40. *Opopaea deserticola* Simon, 1891, male: 36, dorsal; 39, palp retrolateral; 40, palp prolateral. Female: 37, dorsal; 38, genitalia, ventral.

*Opopaea darlingtoni* BRYANT, 1940:267, figs 5, 7 (male holotype from Maisí, Guantánamo, Cuba, in MCZ); DUMITRESCU & GEORGESCU, 1983:103 (Syn.).

*Gamasomorpha floridana*: BRYANT, 1945:199, figs 1, 2 (male, misidentified).

*Opopaea timida* CHICKERING, 1951:233, figs 20, 21 (male holotype believed to have been taken from a bat collected in the Chilibrillo Caves, Canal Zone, Panama, in MCZ); PLATNICK & DUPÉRRÉ, 2009a:4 (Syn.).

*Opopaea brasima* CHICKERING, 1969:148, figs 4-10 (male holotype from Kendall, Dade Co., Florida, USA, in AMNH, examined); DUMITRESCU & GEORGESCU, 1983:103 (Syn.).

*Gamasomorpha* BATTIROLA *et al.*, 2004:423, 426; MARQUES *et al.*, 2007:144.

Note. The figure of *Gamasomorpha rufa* presented by BANKS (1898) shows clearly a flattened oonopid, indicating in the description of the ventral face of abdomen “a dark stripe each side near furrow”, a characteristic structure found in the female genitalia of all *Opopaea*, but particularly distinctive in *O. deserticola*. In addition, this species is common in urban areas in Baja California (PLATNICK & DUPÉRRÉ, 2009a).

Diagnosis. *Opopaea deserticola* can be distinguished from *O. concolor* by the ventrally more expanded palpal bulb in males (Figs 39, 40; PLATNICK & DUPÉRRÉ, 2009a, figs 61-66) and females having small, dark knob marking the origin of the receptaculum situated very close to the epigastric furrow in genital area (Fig. 38).

Description. Male and female, see PLATNICK & DUPÉRRÉ (2009a:4, figs 1-72).

Distribution. According to PLATNICK & DUPÉRRÉ (2009a), this Pantropical species was probably introduced in Americas from the Old World. In the New World, it is known to occur from Florida, USA, to Southern Brazil. There were only two previous records from Brazil, one in the Northern State of Pará and other in the Southwestern State of Rio de Janeiro (PLATNICK & DUPÉRRÉ, 2009a), in a building and in a botanical garden, respectively (PLATNICK & DUPÉRRÉ (2009a:21). This species is synanthropic in Brazil, sympatric with *O. deserticola*, mainly in the states of Bahia and Mato Grosso (Fig. 61). Several specimens were collected in tree canopy at the district of Pirizal, Brazilian Pantanal, state of Mato Grosso. These specimens were collected mainly in *Vochysia divergens* (Vochysiaceae) and *Calophyllum brasiliense* Cambess. (Calophyllaceae) and determined erroneously as *Gamasomorpha* sp. at that time (BATTIROLA *et al.*, 2004; MARQUES *et al.*, 2007). However, this species is most commonly found inside houses, buildings and, exceptionally, in disturbed caves.

Material examined. BRAZIL. **Amapá**: Laranjal do Jari (0°50'31"S - 52°30'57"W), ♂, 14.XI.2003, J. A. P. Barreiros leg. (MPEG 10406); **Amazonas**: Manaus, Reserva Florestal Adolpho Ducke (02°57'42"S - 59°55'40"W), ♀, 15.IX.1995 (INPA); J. Adis leg. (INPA); ♀, 23.III.1996, J. Adis (INPA); **Pará**: Belém, campus Museu Paraense Emílio Goeldi, ♀, 10.II.1959, A. Nadler leg. (AMNH, PBI\_OON 10524); 5 Km East Belém, ♂, 02.V.1974, R. Schuh leg. (AMNH, PBI\_OON 10514); **Mato Grosso**: Nossa Senhora do Livramento, Pirizal, Pantanal de Poconé, Fazenda Retiro Novo (16°15'12"S - 56°22'12"W), 4♂, 10♀, XI-XII.1999, L. A. Battirola (IBSP 40724-40725, 40729, 40754, 40756, 40765-40767); Poconé (16°26'49.5924"S - 56°42'52.0518"W), 4♂,

7♀, 2003, I. Marques & Castilho leg. (IBSP 67014, 67025, 67027, 67029, 67032-67033, 67035, 67037); **Mato Grosso do Sul**: Dois Irmãos do Buriti, Piraputanga, Fazenda Correntes II (20°27'S - 55°30'W), ♀, 16-26.II.2008, R. Bessi leg. (IBSP 128724); **Bahia**: Paulo Afonso (9°24'28"S - 38°13'19'S), Ilha do Urubu, ♀, 2008, E. Daniele leg. (IBSP 125008); Salvador, Baía de Aratu (12°47'32,8"S - 38°28'15,3"W), 2♂, 73♀, 2007-2008, K. Benati leg. (IBSP 125202-125230, 125232-125236, 125238-125240, 125242-125243, 125245, 125248-125252, 125254-125259, 125261-125266, 125268-125270, 125272-125278, 125280, 125282-125291); Parque da Cidade Joventino Silva (12°59'57.6522"S - 38°28'11.1606"W), 2♀, 22.VIII-03.IX.2008, C. Queiroz & D. Mendes leg. (IBSP 125544-125545); **Minas Gerais**: Belo Horizonte (19°52'S - 44°06'W), ♀, 09.XII.2007, A. J. Santos leg. (IBSP 151663); Pains, Gruta Isaias (431402mE - 7747672mN), ♀, 19.VI.2009, R. Zampaulo leg. (IBSP 166586); **Rio de Janeiro**: Rio de Janeiro, Jardim Botânico (30°03'6.07"S - 51°10'37.95"W), ♀, 20.I.1959, A. Nadler leg. (AMNH, PBI\_OON 10526); Santa Teresa, ♀, 26.VI.1946, H. Sick leg. (AMNH, PBI\_OON 28177); Bairro Laranjeiras, Parque Guinle (22°55'50.38"S - 43°11'1.12"W), ♀, 20.X.2001, R. Baptista leg. (MNRJ 3760); **São Paulo**: São Paulo, Parque do Estado (23°39'8.625"S - 46°36'57.9924"W), ♂, 18-25.X.2003, J. Valvassori leg. (IBSP 68059); campus Instituto Butantan (23°33'59.493"S - 46°43'14.7612"W), ♀, 30.IX.2009, A. M. Giroti leg. (IBSP 151545); ♀, VII.2007, G. Perroni (IBSP 151566).

### *Pelicanus marmoratus* Simon, 1891

(Figs 41-45, 60)

*Pelicanus marmoratus* SIMON, 1891:559, fig. 4 (One male and two female syntypes from Saint Vincent Island, Antilles, in BMNH).

*Philesius marmoratus*: SIMON, 1893a:303.

*Myrmopopaea jacobsoni* REIMOSER, 1933:397, figs 1-3 (Male holotype and female paratype from Fort de Kock [= Bukittinggi], Sumatra, in Naturhistorischen Museum Wien); PLATNICK *et al.*, 2012c:18-19, figs 1-60, 121-144 (Syn).

*Gamasomorpha minima* BERLAND, 1942:5, fig. 1a (Male holotype from Canton Island, Phoenix Islands, in Bishop Museum, Honolulu); PLATNICK *et al.*, 2012c:18-19, figs 1-60, 121-144 (Syn).

*Hytanis pusilla* BRYANT, 1942:326, figs 13, 14 (Female holotype from Christiansted, Saint Croix, Virgin Islands, in MCZ); PLATNICK *et al.*, 2012c:18-19, figs 1-60, 121-144 (Syn).

*Scaphiella ula* SUMAN, 1965:230, figs 15-20 (Male holotype from Puu Papaa peak, Oahu, Hawaii, in Bishop Museum, Honolulu); PLATNICK *et al.*, 2012c:18-19, figs 1-60, 121-144 (Syn).

*Triaeris reticulatus* CHICKERING, 1968b:354, figs 6-13 (Male holotype from Saint Croix, Virgin Islands, in MCZ); CHICKERING, 1973:228 (Syn. with *Hytanis pusilla*).

*Triaeris pusillus*: CHICKERING, 1973:228.

*Silhouettella mahei* BENOIT, 1979:205, fig. 6A (male holotype from Morne Blanc, Mahe, Seychelles, in MRAC); PLATNICK *et al.*, 2012c:18-19, figs 1-60, 121-144 (Syn).

*Gamasomorpha gracilipes* WUNDERLICH, 1987:65, figs 37-39 (male holotype from Valle Gran Rey, La Gomera, Canary Islands, in Naturmuseum Senckenberg); SAARISTO, 2001:321 (Syn. with *Silhouettella mahei*).

*Pelicanus mahei*: SAARISTO, 2001:321, figs 40-46.

Diagnosis. Specimens of both sexes of *Pelicanus marmoratus* can be easily separated from all other Brazilian synanthropic species by the weakly sclerotized scuta on the abdomen (Fig. 42), light embolus with distal process in the

male palp (Figs 43, 44) and postepigastric scutum covering about 3/4 of abdominal length (Fig. 45).

Description. Male and female, see PLATNICK *et al.* (2012c:18, figs 1-60, 121-144).

Distribution. Considered Pantropical by PLATNICK *et al.* (2012c) who recorded this species in the African continent, in several Caribbean and Pacific islands and in the city of Ananindeua, state of Pará, Brazil (Fig. 60), where the specimens were collected manually in a house bathroom, along with synanthropic ants. This remains the single record of this species in Brazil.

Material examined. BRAZIL. **Pará:** Ananindeua (01°23'S - 48°24'W), ♀, 04.II.2009, B. Silva leg. (MPEG 18820, PBI\_OON 811); ♂, ♀, 08.II.2009, B. Silva leg. (MPEG 18821, PBI\_OON 810).

### *Heteroonops spinimanus* (Simon, 1891)

(Figs 46-48, 59)

*Oonops spinimanus* SIMON, 1891:563, pl. 42, fig. 6 (Female syntype from Saint Vincent Island, Antilles, in BMNH and five female syntypes from Caracas, Distrito Federal, Venezuela).

*Oonops bermudensis* BANKS, 1902:269, fig. 1a-c (Female holotype from the Bermuda Islands, no specific locality, in Peabody Museum, Yale); STERWALD, 1988:10 (Syn.).

*Heteroonops spinimanus:* DALMAS, 1916:203; PLATNICK & DUPÉRRE, 2009b:22, figs 79-139.

*Oonopinus humus* SUMAN, 1965:238, figs 35-37 (Female holotype from SE slope of Ulumawao Peak, Kailua, Oahu, Hawaii, in Bishop Museum, Honolulu); PLATNICK & DUPÉRRE, 2009b:22 (Syn.).

*Matyotia tetraspinosus* SAARISTO, 2001:351, figs 165-169 (Male holotype from Anse Cimitière, Silhouette, Seychelles Islands, in Zoological Museum at University of Turku, Turku); PLATNICK & DUPÉRRE, 2009b:22 (Syn.).

Diagnosis. Specimens of *Heteroonops spinimanus* can be easily separated from all other Brazilian synanthropic species by the enlarged posterior projection on the male endites (see PLATNICK & DUPÉRRE, 2009b:22, figs 84-86), angular embolus and conductor (see PLATNICK & DUPÉRRE, 2009b:22, figs 87, 88) and the umbrella-shaped anterior receptaculum in the female genitalia (Fig. 47).

Description. Male and female, see PLATNICK & DUPÉRRE (2009b:22, figs 79-139).

Biology. *Heteroonops spinimanus* has been considered to be parthenogenetic by some authors (SAARISTO, 2001), resembling, in that regard, *T. stenaspis*. Even though it was originally described from St. Vincent, it's currently known to have a Pantropical distribution. Although many populations of *H. spinimanus* may be parthenogenetic, apparently conspecific males have been collected together with females twice, in Seychelles Islands and in Florida (PLATNICK & DUPÉRRE, 2009b). All known specimens from Brazil are females, but the low number of samples prevents any claim on a parthenogenetic condition in Brazilian populations.

Distribution. According to PLATNICK & DUPÉRRE (2009b), this species has a Pantropical distribution, but only two records in the South American countries were provided so far, from Colombia and Venezuela. Here the first Brazilian records of *H. spinimanus* are presented, from states of Bahia, São Paulo and Rio Grande do Sul (Fig. 59).

Material examined. BRAZIL. **Bahia:** Itamaraju (16°59'26"S - 39°25'55"W, ♀, 14.IV.2011, C. M. P. Leite leg. (IBSP 166615); Jaguaripe (13°10'39"S - 38°59'51"W), 6♀, 02.III.2011, C. M. P. Leite leg. (IBSP 166616-166618); **São Paulo:** São Paulo, Vila Butantã, 2♀, 18.I.2000, F. S. Cunha leg. (IBSP 30397; IBSP 32967); Ubatuba, Parque Estadual da Ilha Anchieta (23°32'S - 45°03'W), ♀, 23-30.VII.2001, Equipe Biota (IBSP 69884); **Rio Grande do Sul:** Porto Alegre, Jardim Botânico, ♀, 02.I.2013 (MCN 49944); 2♀, 3.V.2013 (MCN 49945); 2♀, 5.VII.2013, all collected by G. O. Silva *et al.*

### *Orchestina pavesiiformis* Saaristo, 2007

(Figs 49-52, 61)

*Orchestina pavesiiformis* SAARISTO, 2007:124, figs 16, 18, 20, 21 (male holotype from Israel, Jerusalem, deposited in HUJ 14220; male paratype, same locality in MZUT 3703; female paratype, same locality, in MZUT 3704; female paratype, same locality, in HUJ 15326; female paratype, same locality, in HUJ 14184); IZQUIERDO & RAMÍREZ, 2017:24, figs 11, 17D-F, 19G, 20N, O, 22D, maps 1, 25.

Diagnosis. Both sexes have lateral setae on the carapace margins, a character exclusive in relation to other native American species. Males can be distinguished from those of the American species by the dorsal V-pattern on the abdomen (see IZQUIERDO & RAMÍREZ, 2017:24, fig. 11A-C) and by the shape of the embolus, sinuous and with distal extensions (Fig. 51; IZQUIERDO & RAMÍREZ, 2017, figs 17D-F, 20N, O). Females have genitalia with external pockets and an anterior receptaculum widened at its base with V-shaped anterior apodemes at the tip (Fig. 52; IZQUIERDO & RAMÍREZ, 2017:24, figs 19G, 22D).

Description. Male and female, see IZQUIERDO & RAMÍREZ (2017: 24).

Biology. According IZQUIERDO & RAMÍREZ (2017) this species has been introduced into many countries and its natural origin is still unknown. In Argentina, it has been collected in disturbed environments and in houses or other human buildings. Specimens were observed walking on walls, furniture and collected in leaf litter. In Brazil, it has been collected in the interior of houses as well as in the interior of disturbed caves.

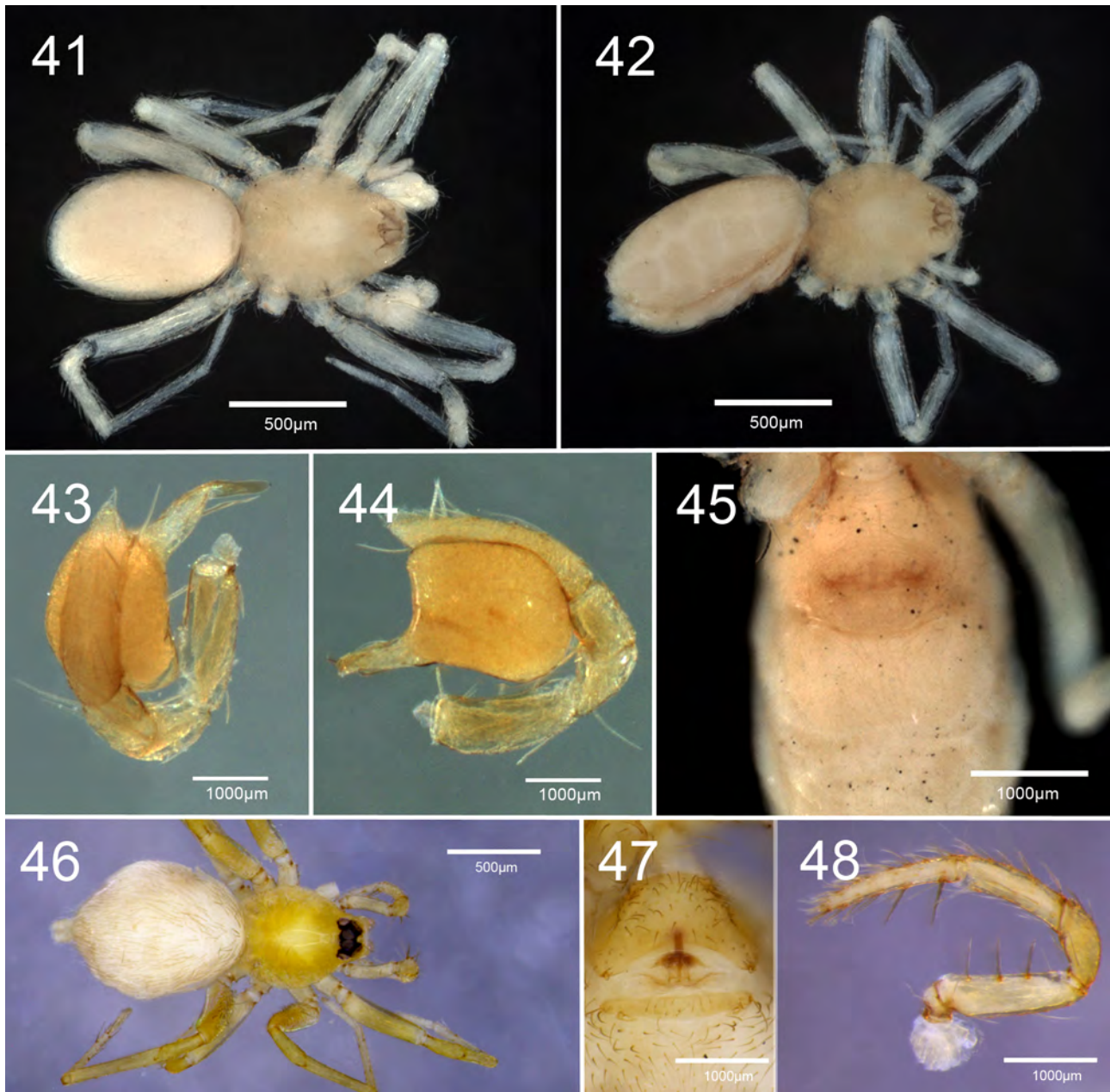
Distribution. According to IZQUIERDO & RAMÍREZ (2017), this species occurs in Israel, United States, Portugal, Brazil, and Argentina (Fig. 61).

Material examined. In addition to the Brazilian material recorded to the states of São Paulo and Minas Gerais by IZQUIERDO & RAMÍREZ (2017), only one specimen is reported here, from state of **São Paulo**, São Paulo, campus of Instituto Butantan, building of LECZ, collected manually on the table, ♀, 26.VII.2017, A. D. Brescovit leg. (IBSP 221179).

### *Orchestina dentifera* Simon, 1893

(Figs 53-57, 61)

*Orchestina dentifera* SIMON, 1893c:248 (one male, four females, and one juvenile syntypes from Sri Lanka (ex Ceylan), Kandy!, deposited in MNHN 6025 [additional number 15250], PBI\_OON 50024); SIMON, 1893d:294; DALMAS, 1916:232, figs 21, 27.



Figs 41-48. Figs 41-45, *Pelicinus marmoratus* Simon, 1891, male: 41, dorsal; 43, palp dorso-prolateral; 44, palp retrolateral. Female: 42, dorsal; 45, genitalia, ventral. Figs 46-48. *Heteroonops spinimanus* (Simon, 1891), female: 46, dorsal; 47, genitalia, ventral; 48, female palp, lateral.

*Orchestina justini* SAARISTO, 2001:356, figs 190-195 (male holotype from Seychelles Islands, Silhouette, Anse Cimitière, deposited in MZUT 1250, PBI\_OON 50030; 3 male and 3 female paratypes, same data, in MZUT 1251, PBI\_OON 50029). IZQUIERDO & RAMÍREZ, 2017:26 (Syn.).

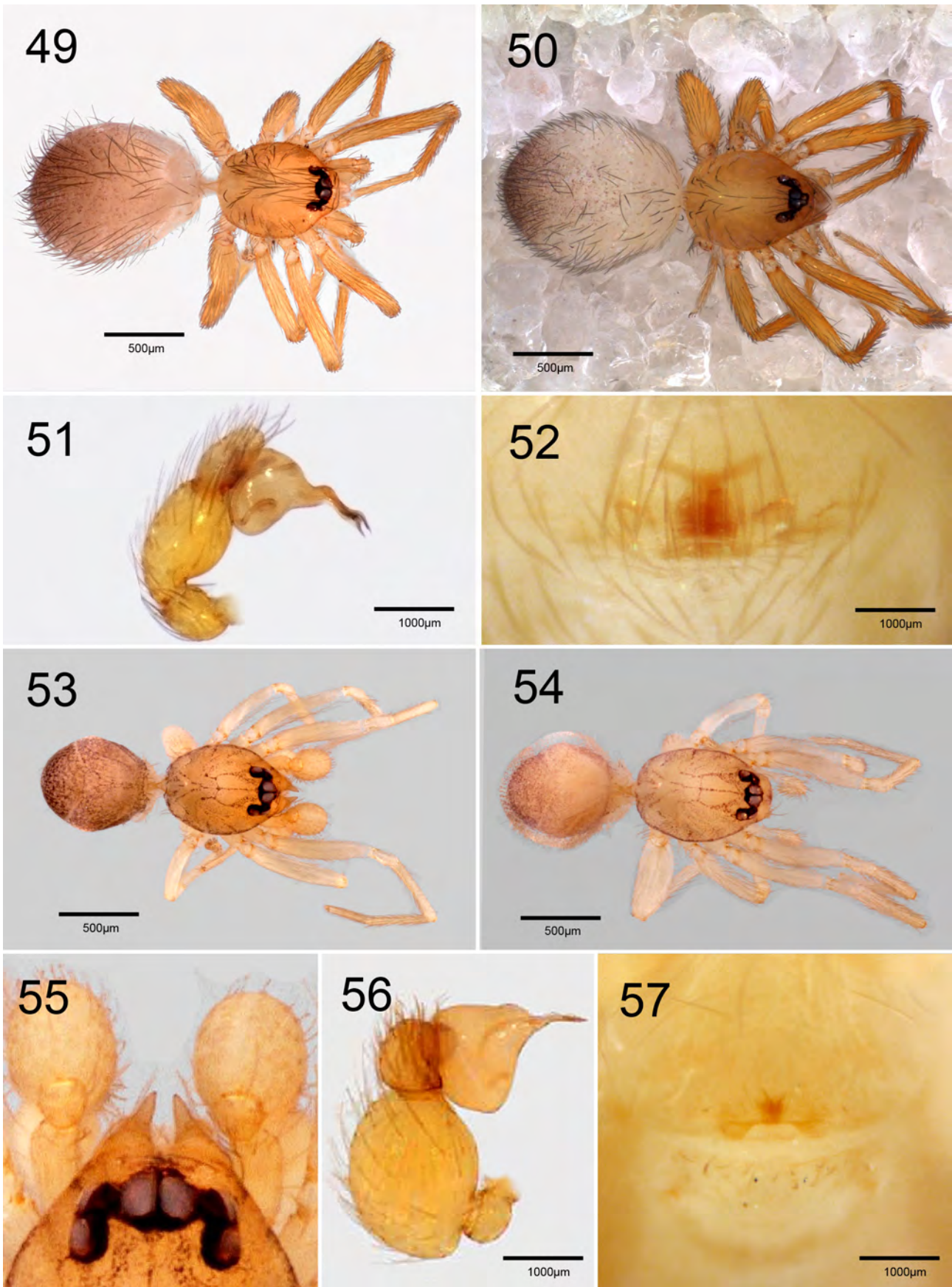
**Diagnosis.** *Orchestina dentifera* can be distinguished from *O. pavesiiformis* by the anterior face of male chelicerae with strong and conical projections on median line (Fig. 55; IZQUIERDO & RAMÍREZ, 2017, figs 25G, 26B, 27A, E) and labium with two pairs of large and flattened setae (see IZQUIERDO & RAMÍREZ, 2017:26, fig. 27B). Females are distinguished from *O. pavesiiformis* by the epigastric

region with highly convex ridge, anterior receptaculum with wide, rounded base, bifid tip; posterior receptaculum present, posterior apodeme bar-shaped (Fig. 57; IZQUIERDO & RAMÍREZ, 2017:26, figs 41A, 44C).

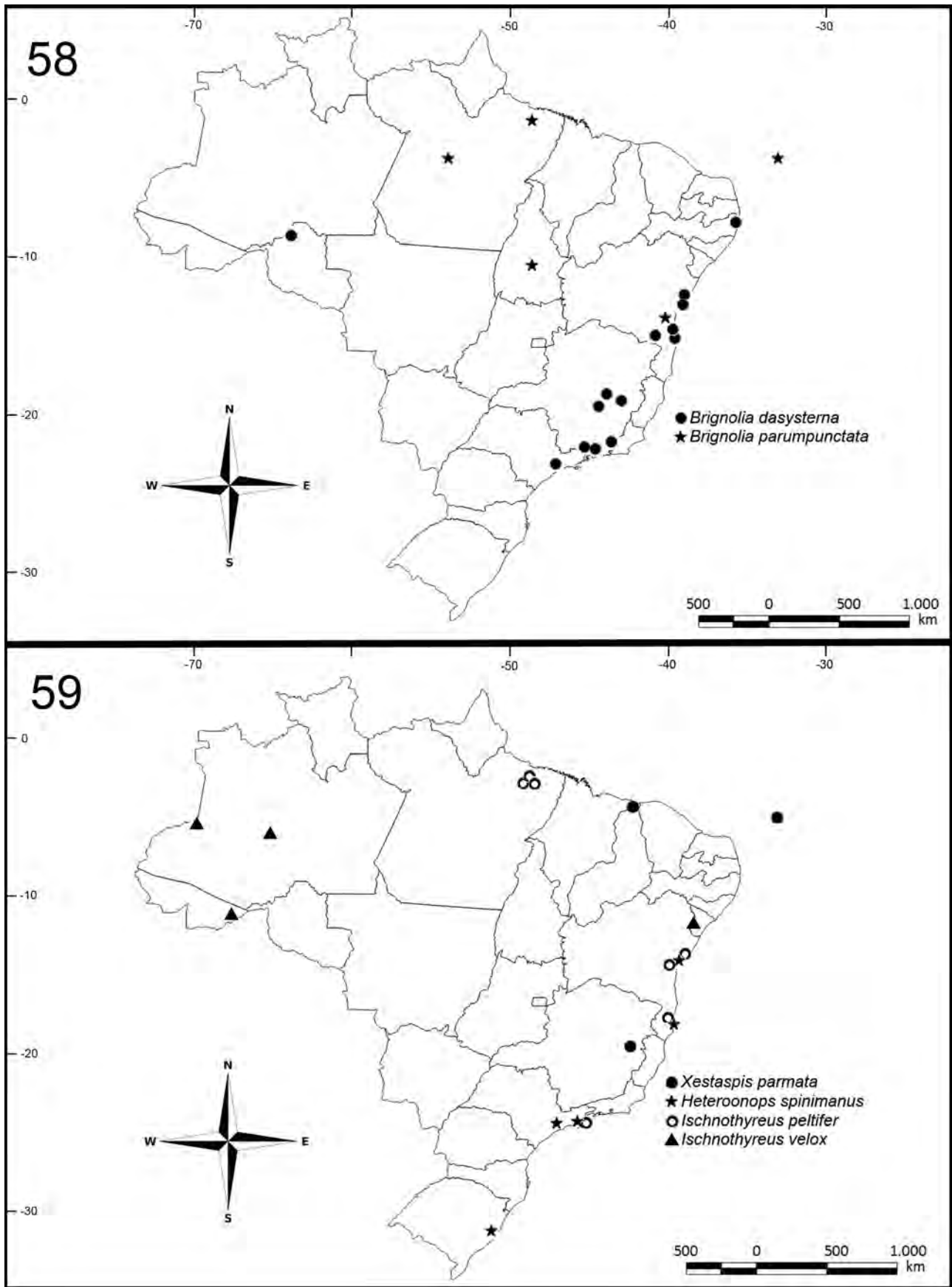
**Description.** Male and female, see IZQUIERDO & RAMÍREZ (2017:26, figs 25-27, 36D-F, 41A, 43, 44C).

**Biology.** In Brazil, specimens were found inside or around houses and in building constructions.

**Distribution.** According to IZQUIERDO & RAMÍREZ (2017), this species occurs in Sri Lanka, Seychelles, Isla Reunion (Indian Ocean), Tanzania, Haiti, Jamaica and Brazil (Fig. 61).

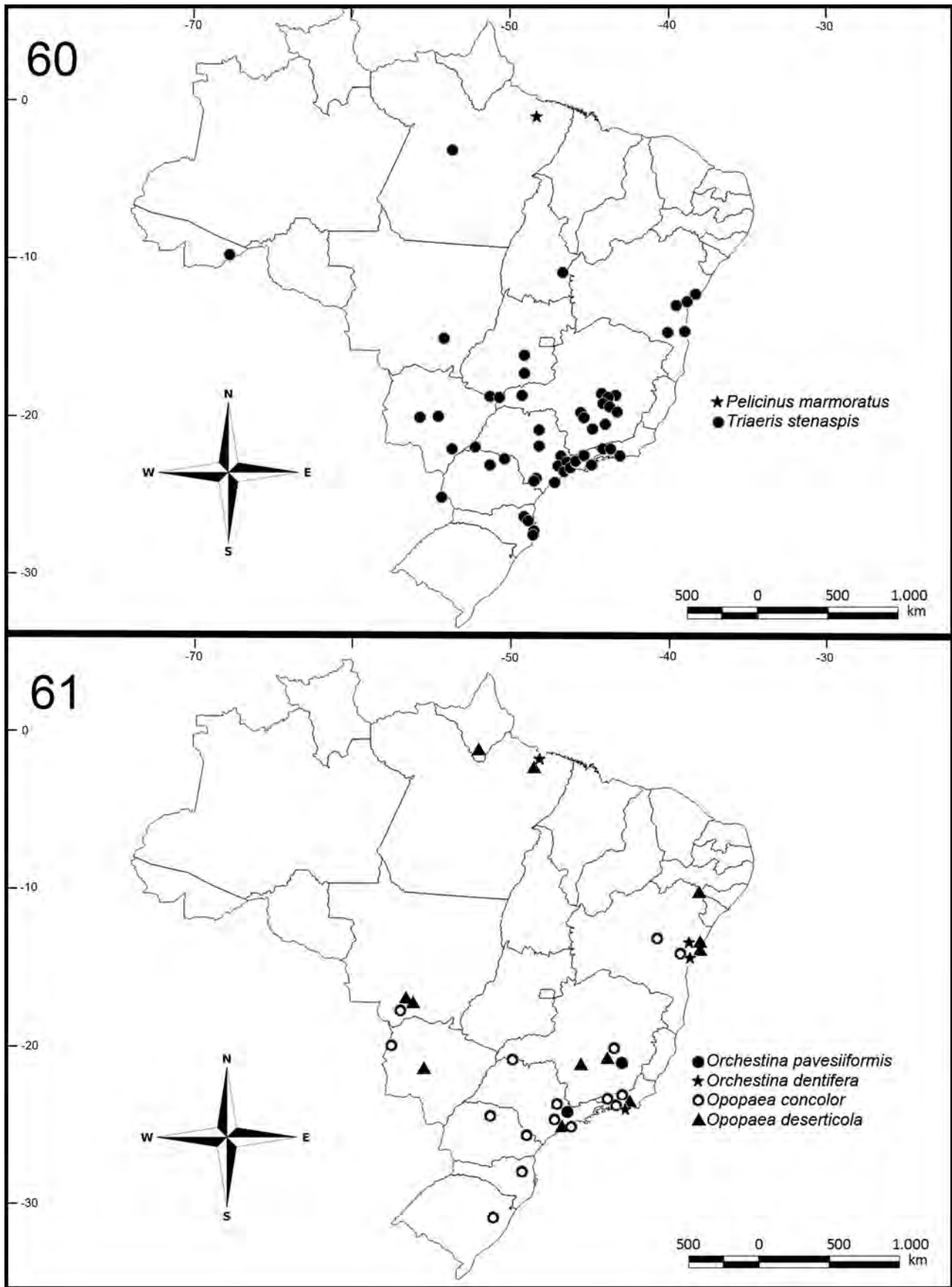


Figs 49-57. Figs 49-52, *Orchestina pavesiiformis* Saaristo, 2007, male: 49, dorsal; 51, palp retrolateral. Female: 50, dorsal; 52, genitalia, ventral; 53-57. *Orchestina dentifera* Simon, 1893, male: 53, dorsal; 55, projection of the chelicerae; 56, palp retrolateral. Female: 54, dorsal; 57, genitalia, ventral.



Figs 58, 59. Brazilian records of: 58, *Brignolia dasystema* and *B. parumpunctata*; 59, *Xestaspis parmata*, *Heteroonops spinimanus*, *Ischnothyreus peltifer* and *I. velox*.





Figs 60, 61. Brazilian records of: 60, *Pelicinus marmoratus* and *Triaeris stenaspis*; 61, *Orchestina pavesiiformis*, *O. dentifera*, *Opopaea concolor* and *O. deserticola*.

Material examined. In addition to the Brazilian material recorded to the states of Pará and Bahia by IZQUIERDO & RAMÍREZ (2017), only three specimens are recorded here: **Rio de Janeiro**, Rio de Janeiro, Campus UFRJ, 2♀, 23.IX.2010, Students from Universidade Federal do Rio de Janeiro leg. (IBSP 167792); **Bahia**, Salvador, Parque Metropolitano do Pituaçu, ♂, 19.I-26.II.2008, D. U. Sena leg. (IBSP 126003).

## DISCUSSION

Brazil has currently 149 known species of Oonopidae (WORLD SPIDER CATALOG, 2019), of which 12 are non-native, representing 8% of the described diversity of oonopids in the country. Some species appear to have been introduced in as early as the first half of the 20th century and have large distribution ranges, as shown in the data of the examined material and Table I. Noteworthy, among the oldest records are the ones of *Opopaea* species, *O. concolor* and *O. deserticola*, sampled as early as 1935 and 1946, respectively (Tab. I). These species are mainly peridomiciliary dwellers and records in native areas are very rare, despite being in the country by at least 70 years. The most intriguing case is perhaps that of *O. deserticola*, which is apparently well established in the Pirizal regions of the Pantanal, state of Mato Grosso, where they are commonly found in canopies (BATTIROLA *et al.*, 2004).

Judging only by the collecting dates here reported, the most recent introductions would be those of *Orchestina dentifera* and *Pelcinus marmoratus*, with records dated as recently as 2008 and 2009 respectively. The populations of these species are apparently small and restricted to a few scattered portions of the Brazilian territory: *P. marmoratus* was found only in the state of Pará, with a single record in an urban area, while *O. dentifera* was recorded along Atlantic coastal areas in three states (Pará, Bahia and Rio de Janeiro). It is not clear at this point if the Brazilian populations of these two species are not yet viable (being continually refilled with new specimens introduced anthropochorically) or represent already established populations that were undersampled in most environments.

The records of the remaining species are also recent, sampled in the last 10 years. The species *Triarhis stenaspis* appears to have the largest distribution range in the country, being the species that has invaded the largest number of natural areas. Assuming that the sampling dates are reliable indicators of a recent arrival, the success of this species in invading and colonizing both urban areas and natural habitats in all Brazilian biomes, including cave environments, could only be explained by the explosive reproduction rates allowed by parthenogenesis, a reproductive trait already proved by KORENKO *et al.* (2009) to occur in at least one population of this species. *Heteroonops spinimanus* also appears to have been introduced very recently and, to date, has been recorded in three Brazilian states, also along the Atlantic coast. Being considered another possible parthenogenetic species (see discussion in PLATNICK *et al.*, 2009b:3, 4), it may well reach quite quickly a wider distribution range in the country.

The richness and abundance of introduced Oonopidae species that eventually colonized urban areas in Brazil highlights the potential deleterious effects of these aliens on native spider populations and communities. Unfortunately, studies on the ecological effects of introduced spiders are sparse and there are a general lack of reliable data properly addressing critical questions for conservation, such as degrees of direct competition, rates of species substitution and the chain effects on the affected ecosystems, even the allegedly simplified ones, as is the case of caves. Cave environments could be particularly affected since they are prone to contain endemic faunistic components, maintaining species with varying degrees of troglomorphisms and population sizes. This issue deserves urgent attention regarding the implementation of studies to prevent, stop or at least reduce the expansion of these small invasive spiders in those environments.

Tab. I. Number of records and earliest recorded year of introduction of invasive oonopids in Brazil.

	Number of records	Earliest record
<i>Brignolia dasystema</i>	114	1974
<i>Brignolia parumpunctata</i>	220	1974
<i>Triarhis stenaspis</i>	515	1996
<i>Xestaspis parmata</i>	2	2006
<i>Ischnothyreus peltifer</i>	14	2001
<i>I. velox</i>	6	2002
<i>Opopaea concolor</i>	37	1935
<i>O. deserticola</i>	117	1946
<i>Pelcinus marmoratus</i>	3	2009
<i>Heteroonops spinimanus</i>	9	2000
<i>Orchestina pavesitiformis</i>	8	2002
<i>O. dentifera</i>	5	2008

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