

***Susanlimae ianwhittingtoni* gen. nov., sp. nov. (Monogenoidea: Dactylogyridae), a dweller of the gill rakers of *Pseudeutropius moolenburghae* (Siluriformes: Schilbeidae) from Sumatra**

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ABSTRACT. A new genus and species of Monogenoidea, *Susanlimae ianwhittingtoni* gen. nov., sp. nov., are proposed for dactylogyrids collected from the “Nuayang tipis”, *Pseudeutropius moolenburghae* Weber & de Beaufort, 1913, which inhabits freshwater in Sumatra. While clearly a member of a putative clade that includes Asian and African catfish parasites, *S. ianwhittingtoni* sp. nov. differs from most members of this clade by having a bifurcated haptor that embraces the gill rakers of its host. This haptoral morphology and mode of attachment also occur in species of *Bifurcohaptor* Jain, 1958. However, species of *Susanlimae* gen. nov. are easily distinguished by the comparative morphology of their haptoral armature. In *Susanlimae* gen. nov., the ventral and dorsal bars are single, elongated, and inverted u-shaped (ventral bar short and two short dorsal bars in *Bifurcohaptor* spp.); the dorsal anchor is robust with well-defined roots (reduced roots and elongate shaft in *Bifurcohaptor* spp.); and the ventral anchor has an elongated, deep root (inconspicuous in *Bifurcohaptor* spp.).

KEY WORDS. Catfish; Dactylogyridea; ectoparasite; Platyhelminthes; Polyonchoinea.

During a survey of monogenoidean parasites from the freshwaters of Sumatra (Indonesia), specimens of a new genus and species were collected from “Nuayang tipis”, *Pseudeutropius moolenburghae* Weber & de Beaufort, 1913. This fish is a demersal species inhabiting rivers and lakes in Sumatra and Borneo (FROESE & PAULY 2015). The new species is unique among the Dactylogyridae in having haptoral structures adapted to embrace the gill rakers of the host. The organization and morphology of the hard structures within the haptor warrant the proposal of a new genus to accommodate the new species. A similar attachment style, however, occurs in species assigned to other monogenoid genera and families.

MATERIAL AND METHODS

Host fish used in this research were captured by local fishermen in May 2005 from the Batang Hari River near the village of Kubu Kandang, Sous-district Pemayung, Province of Jambi, Sumatra, Indonesia. They were acquired from the local fish market and were immediately preserved in ethanol (70–80%) for subsequent processing.

In the laboratory, the gills of each fish were removed and their parasites were collected from the sediment with the aid of probes and forceps under a dissecting scope. Some specimens were stained with Gomori's trichrome and mounted in Damar's gum for study of their soft anatomy; other specimens were cleared and mounted in Hoyer's mounting medium (prepared as in HUMASON 1979) for study of their hard structures. Illustrations were prepared using a camera lucida on an Olympus BX51 microscope equipped with phase contrast. Measurements, all in micrometers, were taken following the procedures of MIZELLE & KLUCKA (1953). In the descriptions, the mean is followed by the range and the number of structures measured (n), in parentheses; the length of the body includes the haptor (longitudinal axis of haptor). The length of the male copulatory organ (MCO) and bars represent their actual length measured with the software ImageJ (SCHNEIDER et al. 2012). The hooks are numbered according to MIZELLE (1936). The type specimens are deposited in the parasitological collections of the Instituto Oswaldo Cruz (Rio de Janeiro, Brazil) (CHIOC) and the Muséum national d'Histoire naturelle (Paris, France) (MNHN).

TAXONOMY

Class Monogenoidea Bychowsky, 1937
 Subclass Polyonchoinea Bychowsky, 1937
 Dactylogyroidea Bychowsky, 1937
 Dactylogyridae Bychowsky, 1933
Susanlimae gen. nov.

Diagnosis. Body fusiform, strongly flattened dorsoventrally, comprising body proper (cephalic region, trunk) and haptor; peduncle inconspicuous. Tegument smooth. Cephalic lobes inconspicuous; cephalic glands unicellular, lateral or posterolateral to pharynx. Eyespots 4; granules small, ovate. Mouth subterminal, midventral, prepharyngeal; pharynx muscular, glandular; oesophagus short or inconspicuous; intestinal caeca two, non-confluent, lacking diverticula. Common genital pore midventral at level of intestinal bifurcation. Gonads intercaecal, tandem, germanium posterior to testis. Vas deferens looping left intestinal caecum; seminal vesicle a simple dilation of vas deferens. Copulatory complex comprising MCO, lacking accessory piece. MCO sclerotized, tubular. Seminal receptacle pregerminal; vaginal pore single, ventral, intercaecal, non-sclerotized; vaginal duct intercaecal, non-sclerotized. Vitellaria in trunk, absent from regions of other reproductive organs. Haptor bifurcated, composed by bilateral "arms"; with dorsal, ventral anchor/bar complexes, seven pairs of similar hooks (5 pairs ventral, 2 pairs dorsal). Hook with shank comprising single subunit. Bars elongate, inverted U-shaped. Ventral anchor with elongate deep root. Superficial root of dorsal anchor provided with accessory sclerite (= *cuneus*). Parasites of the gill rakers of species of Siluriformes (Actinopterygii).

Type species. *Susanlimae ianwhittingtoni* sp. nov.

Etymology. The generic epithet is in honor of Dr. Lee Hong Susan Lim (1952-2014) (University of Malaya in Kuala Lumpur), a Malaysian parasitologist and a good friend of the first two authors. Dr. Lim is greatly responsible for most of our knowledge of the diversity of Monogenoidea from Asian Siluriformes.

Susanlimae ianwhittingtoni gen. nov., sp. nov.

Figs. 1-9

Description (based on 6 specimens – 3 stained, 3 cleared). Body 330 (227-394, n = 3) long; greatest width 92 (82-99, n = 3) at level of gonads. Eyespots 4, accessory granules scarce in cephalic region. Pharynx ovate, 38 (35-44, n = 3) long, 28 (26-29, n = 3) wide. Testis ovate, often transverse, 25 (13-39, n = 3) long, 49 (39-57, n = 3) wide; vas deferens looping left intestinal caecum; seminal vesicle elongate; prostatic reservoir not observed. MCO 61 (53-77, n = 3) long, wide, curved on itself; base of MCO wide. Germarium ovate, 42 (23-57, n = 3) long, 32 (27-39, n = 3) wide. Uterus not observed. Vaginal pore single, ventral, at midlength of the body. Vitellaria dense, overlapping caeca. Hooks similar in shape, with slightly erected thumb,

straight shaft, short point; hooks 16 (15-19, n = 6) long; shank proximally bulbous; 5 ventral pairs, one dorsal pair (hook pair 6) located at the distal extremity of the haptoral arms; one dorsal hook (hook pair 7) located at midlength of each bilateral arm. Ventral anchor 86 (73-94, n = 3) long, base 10 (8-11, n = 3) wide, delicate, with well differentiated roots, deep root about 10 times longer than superficial root, straight shaft, point recurved, about ½ shaft length. Dorsal anchor 73 (60-82, n = 3) long, base 51 (42-61, n = 3) wide, robust base, with short deep root, robust superficial root with small cuneus, shaft and point evenly curved, shaft with small external indentation near midlength. Ventral bar 389 (279-492, n = 3), delicate, tapering towards extremities, medially hinged. Dorsal bar 250 (192-303, n = 3) long, robust, with slightly tapering ends. Eggs not observed.

Type host. *Pseudeutropius moolenburghae* Weber & de Beaufort, 1913 (Siluriformes: Schilbeidae)

Site of infection. Gill rakers.

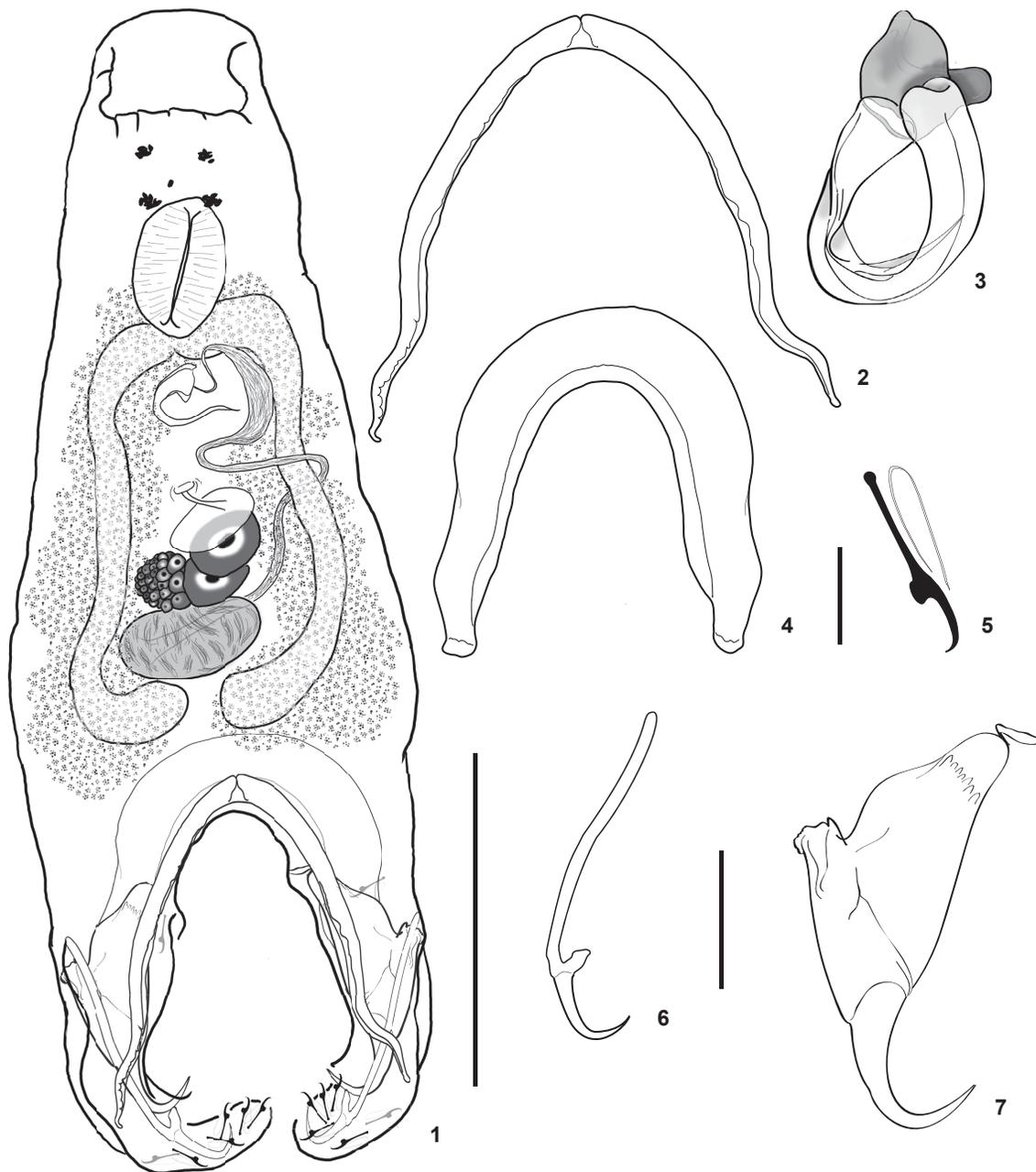
Type locality. Batang Hari river, near the Village de Kubu Kandang, Sous-district Pemayang, Province Jambi, Indonesia. Latitude: 1°36'17.51"S, Longitude: 103°19'16.65"E. Date: May 2005.

Specimens deposited. CHIOC 38202 a (holotype); 5 paratypes CHIOC 38202 b,c, MNHN HEL545, HEL546. One paratype kept in the private collection of AP.

Etymology. The specific epithet is proposed in honor of the late Dr. Ian Whittington (1960-2014), South Australian Museum and University of Adelaide, Australia. Dr. Whittington passed away too soon, but not before making an impressive contribution to the biology, taxonomy, and phylogeny of Monogenoidea, especially those of Capsalidae.

DISCUSSION

Despite its apparently unique morphology, *S. ianwhittingtoni* shares many features with other dactylogyrid parasites of siluriform fishes from Asia and Africa. For instance, the ventral anchors are significantly smaller than the dorsal anchors, each dorsal anchor has an accessory sclerite (= *cuneus*) associated with its superficial root, and the ventral bar tends to be V-shaped or composed of two smaller bars, articulated or not. These features are common in species of several genera, such as *Thaparocleidus* Jain, 1952, *Cornudisoides* Kulkarni, 1969 (Fig. 10), *Bifurcohaptor* Jain, 1958 (Fig. 11), *Mizelleus* Jain, 1957, *Bychowskyella* Achmerow, 1952, *Quadriacanthus* Paperna, 1961, *Malayanodisoides* Lim & Furtado, 1986, *Notopterodisoides* Lim & Furtado, 1986, *Pseudancylodisoides* Yamaguti, 1963, and *Paraquadriacanthus* Ergens, 1988. Together with other characteristics, they may represent synapomorphies providing support for the shared ancestry of these genera. LIM et al. (2001) for example, regarded the dactylogyrids mentioned above, together with other species lacking the aforementioned combination of characters, as members of Ancylo-discoidinae Gussev, 1961. In this study, however, we accept the hypotheses of KRITSKY

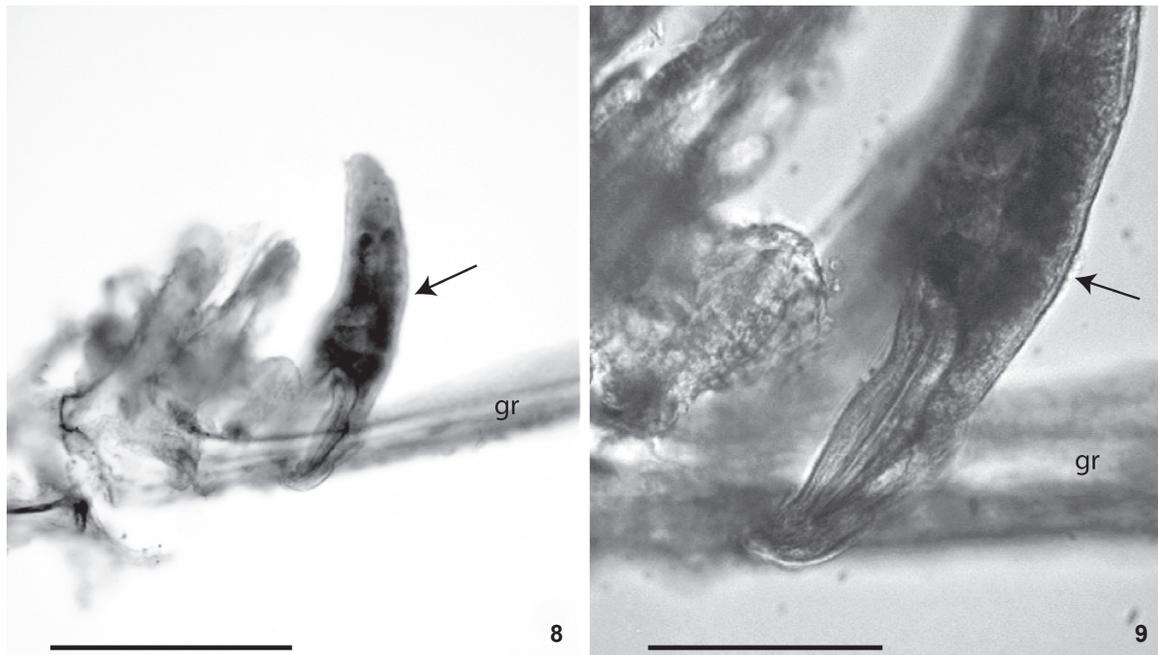


Figures 1-7. *Susanlimae ianwhittingtoni* sp. nov.: (1) holotype, ventral view; (2) male copulatory complex; (3) ventral bar; (4) dorsal bar; (5) hook; (6) ventral anchor; (7) dorsal anchor. Scale bars: 1 = 100 μ m, 2 = 25 μ m, 3, 4, 6, 7 = 50 μ m, 5 = 5 μ m.

& BOEGER (1989) and ŠIMKOVÁ et al. (2003) and refrain from recognizing Ancylo-discoidinae (as well as Ancyrocephalinae) as valid, since this group is not monophyletic according to published phylogenies.

Unlike most species of Monogenoidea of catfishes, *S. ianwhittingtoni* attaches to its host by embracing the gill rakers

and penetrating the epithelium with anchors and hooks located at the distal extremities of the bifurcated haptor (haptor arms) (Figs. 8, 9). A similar kind of attachment is found in species of *Bifurcohaptor*, but in the latter the attachment is to the gill filament rather than the gill rakers (KEARN & BIJUKUMAR 1997). *Bifurcohaptor indicus* Jain, 1958 (and likely all other spe-



Figures 8-9. Photomicrographs of specimens of *Susanlimae ianwhittingtoni* sp. nov. (arrows) attached to the on the gill rakers (gr) of *Pseudeutropius moolenburghae*. Scale bars: 8 = 250 μ m, 9 = 100 μ m.

cies of this genus) embraces only part of the gill filament, the afferent border (see KEARN & BIJUKUMAR (1997). However, comparative analysis of the hard parts involved in the attachment in *S. ianwhittingtoni* and *B. indicus* indicates that this mode of attachment likely originated from independent evolutionary events (see Figs. 11, 12). In the new species, both ventral and dorsal bars are inverted V or U-shaped, the ventral anchor depicts an unusually long deep root, while the dorsal anchor is robust with conspicuous roots. In species of *Bifurcohaptor*, the dorsal bar is small and robust; the ventral bar is split into two separate parts, each positioned by each ventral anchor and located distally in the haptor arms; and the dorsal anchor is greatly elongated, almost as long as each respective haptor arm, lacking conspicuous roots.

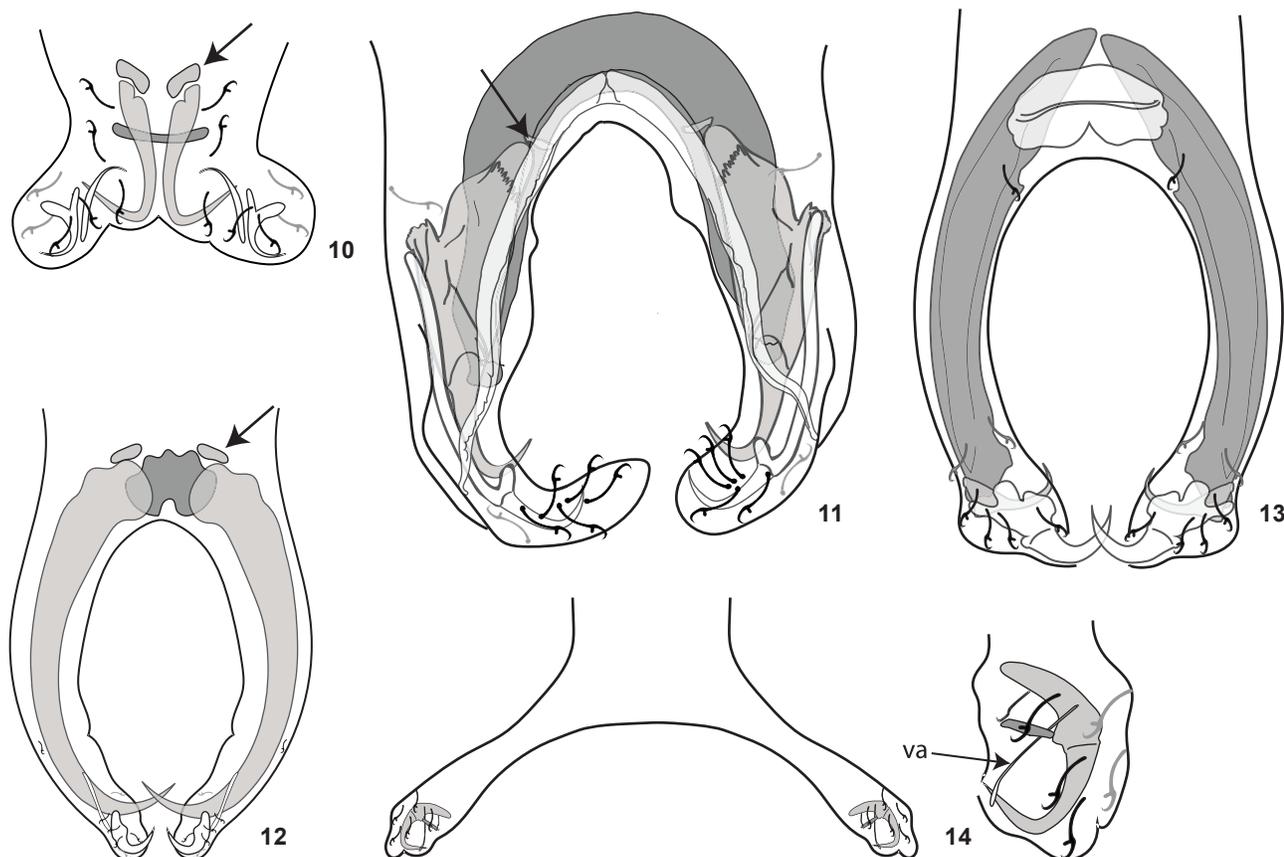
Even though haptors adapted to embrace the gill filaments or gill rakers are unusual among the Monogenoidea, species of other genera of Polyonchoinea have similar functional morphology, including those of Dactylogyridae and Diplectanidae. The single species of *Furcohaptor*, *F. cynoglossi* Bijukumar & Kearn, 1996 (Fig. 13), parasite of the gill filaments of the flatfishes *Cynoglossus macrostomus* Norman, 1928 and *Cynoglossus puncticeps* (Richardson, 1846), has haptor arms but the haptor armature is reduced. The genus was originally assigned to the Ancyrocephalinae, but is most likely a member of Diplectanidae. This hypothesis is supported by the morphology of the anchor and divided bar, greatly resembling those of Diplectanidae; by the presence of a male copulatory organ directed posteriorly; and

the morphology of the head area and head organs. A definitive transfer of *Furcohaptor* to Diplectanidae may not happen until analysis of molecular data and description of the internal organs (not provided in the original description of the species) (BIJUKUMAR & KEARN 1996). In *F. cynoglossi*, the haptor arms are elongate and the hard structures (hooks, bars, and anchors) are located solely at the distal portion of these arms.

Another diplectanid, *Aetheolabes goldiensis* Boeger & Kritsky, 2009 (Fig. 14), a parasite of the freshwater Sciaenidae *Plagioscion* sp. (and its likely congeneric *Diplectanum umbrinum* Tripathi, 1959 – see BOEGER & KRITSKY 2009) also presents an embracing haptor. In species of *Aetholebes* Boeger & Kritsky, 2009 the ventral bar is short, robust, and located at the base of the haptor arms; the dorsal bars are long, almost as long as the haptor arms, and articulate proximally; and the anchors are located distally within the arms. As *Susanlimae* gen. nov. and *Aetholebes* are members of distinct families within the Polyonchoinea, the general similarities in haptor morphology are simply the result of convergent evolution.

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Figures 10-14. Morphology and organization of the haptor armature of different groups of Dactylogyrirea: (10) haptor armature of *Cornudiscoides bagri* Lim, 1987 modified from LIM (1987); (11) haptor armature of *Susanlimae ianwhittingtoni* sp. nov.; (12) haptor armature of *Bifurcohaptor* modified from KEARN & BIJUKUMAR (1997); (13) haptor armature of *Furcohaptor cynoglossi* modified from BIJUKUMAR & KEARN (1996); (14) haptor armature of *Aetheolabes goldiensis* modified from BOEGER & KRITSKY (2009). Ventral sclerites are clear; light gray indicates the dorsal anchor; dark gray indicates dorsal bar; black hooks are ventral; gray hooks are dorsal. Arrows indicate the accessory sclerite; va = reduced ventral anchor. Figures are not to scale. Number and/or distribution of hooks in figures 10, 12, 13 are merely illustrative, as they were not confirmed in type or voucher specimen.

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ERRATA

Page 532:

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