A second species of Cheleion from Johor, Malaysia (Coleoptera, Scarabaeidae, Aphodiinae, Stereomerini)

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Academic editor: F. Krell | Received 10 July 2015 | Accepted 6 October 2015 | Published 5 November 2015

Citation: Král D, Hájek J (2015) A second species of Cheleion from Johor, Malaysia (Coleoptera, Scarabaeidae, Aphodiinae, Stereomerini). ZooKeys 532: 87–97. doi: 10.3897/zookeys.532.6116

Abstract

A new species of the genus Cheleion Vårdal & Forshage, 2010, Cheleion jendeki sp. n., from Johor, Malaysia is described, illustrated and compared with the type species of the genus, C. malayanum Vårdal & Forshage, 2010. Photographs of the two species are presented. The adaptation to inquilinous lifestyle of Cheleion is compared with those in other beetle groups and briefly discussed.

Keywords

Cheleion, new species Coleoptera, Scarabaeidae, Aphodiinae, Stereomerini, Malaysia, Oriental Region

Introduction

Scarabaeoidea (Lamellicornia) represent a distinct, cosmopolitan group of beetles, comprising approximately 2,500 genera and 35,000 species worldwide. They occupy a vast range of various niches, including inquilinous, either myrmecophilous or termitophilous, lifestyles (for a review, see Scholtz and Grebennikov 2005). Among the most peculiar presumably inquilinous scarabs are members of the small, rarely collected tribe Stereomerini of the subfamily Aphodiinae. The 21 presently known species of Stereomerini are currently assigned to nine genera. All representatives of the tribe are allegedly associated with termite nests. Seven of them, Adebrattia Bordat & Howden,
1995, Australoxenella Howden & Storey, 1992, Bruneixenus Howden & Storey, 1992, Daintreeola Howden & Storey, 2000, Danielssonia Bordat & Howden, 1995, Pseudostereomera Bordat & Howden, 1995, and Stereomera Arrow, 1905 are restricted to insular southeastern Asia and Australia (Howden and Storey 1992, 2000; Bordat and Howden 1995; Storey and Howden 1996; Maruyama and Nomura 2011). At present, only the genera Cheleion Vårdal & Forshage, 2010 and Rhinocerotopsis Maruyama, 2009 are known from the Peninsular Malaysia (Maruyama 2009, Vårdal and Forshage 2010). The genus Cheleion so far has contains only one species, C. malayanum Vårdal & Forshage, 2010, described from Pahang. A second species, C. jendeki sp. n., discovered from primary tropical forest of Endau Rompin NP in another Malaysian state, Johor, is described in the present paper.

**Material and methods**

The specimens were examined with an Olympus SZ61 stereomicroscope. Measurements were taken with an ocular graticule. The habitus photographs were taken using a Canon MP-E 65mm f/2.8 macro lens with 5:1 optical magnification on bellows attached to a Canon EOS 550D camera. Partially focused images of specimen were combined using Helicon Focus 3.20.2Pro software. External morphology of both species was also examined with a Hitachi S-3700N environmental electron microscope in the Department of Paleontology, National Museum in Prague (in both cases using uncoated specimens). Exact label data are cited for the type material. Our remarks and addenda are found in brackets, separate label lines are indicated by a slash (/), separate labels by a double slash (//). The holotype of the newly described species is deposited in the collection of National Museum, Prague, Czech Republic (NMPC). For comparison, the holotype of Cheleion malayanum (deposited in Swedish Museum of Natural History, Stockholm, Sweden) was studied. For morphological terms used in the description we largely follow Howden and Storey (1992) and Vårdal and Forshage (2010).

**Taxonomy**

*Cheleion jendeki* sp. n.

http://zoobank.org/4DF33FD9-FF5D-47A1-8967-06EAE42349D

Figures 1, 3, 5, 7, 9, 11, 13–15, 17

**Type locality.** Malaysia, Johor, Endau Rompin National Park, 02°37’12”N 103°21’00”E, 120–300 m a. s. l.

**Type material.** Holotype: ♀, “Malaysia, Johor / Endau Rompin NP / N2.62, E103.35 / 28-31.v.2013, 120-300 m / E. Jendek & O. Šauša leg. [printed] // Cheleion jendeki sp. nov. / HOLOTPYUS ♀ / David Král & Jiří Hájek det. 2015 [red, printed]”. 
Description of female holotype. Slightly convex, integument chestnut brown; head appendages and tarsi amber coloured; whole dorsal surface more or less covered with appressed lanceolate scales (Fig. 1).

Head (Figs 1, 3, 7) remarkably transverse, subrectangular in dorsal view, clypeus shiny, impunctate, apically pointed and reflexed under head, frons slightly convex with five straight, anteriorly divergent furrows; posterior transverse furrow across head between posterolateral corners of eyes; occiput with numerous small, longitudinal pits. Surface covered with dense appressed, lanceolate, approximately regularly spaced scales, individual scales separated from each other by less than their diameter (Fig. 7). Antennae long, length equal to width of head, with long macrosetae. Maxillary palpi length equal to length of head, with securiform ultimate palpomere. Labial palpi with long macrosetae apically. Eyes small but visible in dorsal view (Fig. 7).

Pronotum (Figs 3, 9, 13) large and transverse, anterior edge shallowly bisinuate, sides regularly, broadly rounded, posterior edge with broad medial protrusion. Pronotal disc with seven furrows medially, converging towards middle in hourglass...
Figures 3–6. Habitus of *Cheleion*. 3, 5 *C. jendeki* sp. n. (♀ holotype) 4, 6 *C. malayanum* (♀ holotype; Malaysia, Pahang, Bukit Fraser) 3, 4 dorsal view 5, 6 ventral view.

pattern, mid furrow shallower than lateral furrows (Figs 3, 9). Anteromedial disc with distinctly raised knob, posteromedial disc and posterolateral sides with slightly lower, bulbous areas; anterolaterally of the furrows with large, flat elliptical depressions, de-
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Figures 7–12. Details of *Cheleion*. 7, 9, 11 *C. jendeki* sp. n. (♀ holotype) 8, 10, 12 *C. malayanum* (♀ holotype; Malaysia, Pahang, Bukit Fraser) 7–8 head 9–10 pronotum 11–12 base of right elytron.

lineated by furrows. Knob posteriorly and bulbous areas anteriorly with tufts of long dense microtrichiae (= trichomes) (Fig. 13); surface covered with dense apressed, lanceolate, approximately regularly spaced scales, individual scales separated by less their diameter from each other anterolaterally and laterally; scales on knob and bulbous areas smaller and sparser; flat lateral areas with several sparse rather irregularly spaced scales only (Fig. 13).

Scutellar shield triangular, notably small (Fig. 11).
Elytra approximately as broad as pronotum and only slightly longer than pronotum and head combined; tapering posteriad, rounded apically. Each elytron with five longitudinal ridges before the lateral edge (Figs 1, 3, 11); ridges of approximately same height, elevated and almost continuous, consisting of longitudinal rows of almost confluent tubercles (Fig. 11); intervals (between ridges) flat, rugose, with irregularly circular pads, each pad bearing lanceolate scale on posterior edge, individual pads sepa-
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rated by less their diameter from each other discally, becoming confluent into small rows or groups laterally, especially in humeral area (Figs 11, 14). Epipleura broadly inflexed; posterior two thirds of lateral edge slightly recurved (to allow free movement of metathoracic legs).

Legs short with broad femora and tibiae; tarsi short, tetramerous; claws weak, short, almost straight (Figs 5, 15, 17). Femora shiny, covered with coarse, dense, almost regularly spaced macrosetigerous punctures (Figs 15, 17). Protibiae moderately wide, with finely serrated outer edge and one strong apical lateral tooth, tarsus inserted well before apex (Fig. 15). Meso- and metatibiae broad with finely serrated outer edge and concave apex; each with two inconspicuous terminal spurs and two rows of thick short macrosetae on outer edge (Fig. 17).

Macropterous.

Pygidium exposed, strongly punctate proximally, less strongly apically, with small emargination on proximal pygidial border.

Venter. Prosternal process remarkably elevated, strongly expanded anteriad and posteriad (Figs 5, 15); anterior part grooved longitudinally and sinuate apically, posterior part hastate, surface rugose with marginal bead (Fig. 15). Metaventrite narrow with alutaceous surface (Fig. 17). Mesoventral plate flat, triangular, tapering, widest anteriorly, grooved along midline, surface alutaceous, covered with coarse, dense, almost regularly spaced macrosetigerous punctures (Fig. 17).

Five visible abdominal ventrites apparently fused, covered with coarse, dense, almost regularly spaced macrosetigerous punctures.

**Measurements.** Total body length: 1.9 mm, width at broadest point 0.9 mm.

**Differential diagnosis.** The new species is classified in the genus *Cheleion* mainly by the combination of the following characters: strongly tuberculate and rugose body surface, noticeably long antennae, pronotum with distinct anteromedial knob and bulbous areas medially and laterally and hastate posterior prosternal process. *Cheleion jendeki* sp. n. is similar and probably closely related to *C. malayanum*, the only other known species of the genus, but clearly differs mainly as follows:

- lateral longitudinal grooves on head straight (Figs 1, 3, 7) (weakly s-shaped in *C. malayanum* (Figs 2, 4, 8));
- sides of pronotum regularly rounded, maximum width of pronotum in midlength; posterior angles subrectangular (Figs 3, 9, 15) (sides of pronotum more attenuated in basal third, maximum width of pronotum in anterior third; posterior angles obtuse with apparent spiniform scales in *C. malayanum* (Figs 4, 10, 16);
- elytral surface at first sight moderately rugose (Figs 1, 3, 11) (more strongly rugose in *C. malayanum* (Figs 2, 4, 12);
- elytral ridges distinctly elevated (Figs 3, 11) (almost flat in *C. malayanum* (Figs 4, 12);
- elytral ridges continuous, consisting of longitudinal rows of almost confluent tubercles; lateral ridges indistinct (Fig. 3) (rather discontinuous, consisting of tubercles with scanty longitudinal groups of tubercles with scales; all elytral ridges distinct in *C. malayanum* (Fig. 4));
– pads on elytral intervals separated by less than their diameter and mostly arranged as triseriate discally (Fig. 11) (confluent to subconfluent and mostly arranged as biseriate in *C. malayanum* (Fig. 12));

– marginal bead of posterior part of prosternal process rounded apically (Fig. 15) (angulate apically in *C. malayanum* (Fig. 16)).

In spite of clear differences mentioned above, we are aware that only single specimens are known for each *Cheleion* species. In addition, both type localities are placed only about 200 km apart, without any distinct barrier between them. Thus, we cannot exclude the possibility that morphological differences of *C. jendeki* sp. n. represent only an intraspecific variability of *C. malayanum*, but we consider it quite improbable.

**Etymology.** Patronymic; named in honour of our colleague and friend Eduard Jendek (Ottawa, Canada), excellent student in Buprestidae and collector of the holotype.

**Distribution.** So far known only from the type locality in the Johor Province of continental Malaysia.

**Collecting circumstances.** Flight intercept trap exposed inside lowland primary tropical forest (Fig. 19; E. Jendek, pers. comm. 2015).

**Discussion**

Virtually nothing is known about the biology of Stereomerini. Beetles were repeatedly supposed to be termitophilous, based on single finding of *Termitaxis holmgreni*
Krikken, 1970 with termites in Peru (Krikken 1970). However this genus no longer belongs to the tribe Stereomerini as it was excluded by Bordat and Howden (1995). All other members of the Stereomerini were usually collected by flight intercept traps (FIT) in primary forests, more rarely they were also sifted or attracted at UV light (Storey and Howden 1996), collected with window trunk traps, or with yellow pan traps (Howden and Storey 2000).

We have not been able to trace any “typical characters” distinguishing myrmecophilous and termitophilous beetles. For example, Crowson (1981) noted that “termitophilous beetles tend to show rather less extreme structural modifications than comparable myrmecophilous ones”, and that “termitophilous beetles do not as a rule develop the elaborate trichomes seen in some of the more specialized myrmecophiles”.

It is far beyond the scope of this paper to solve this problem, but we would like to point out several facts that may suggest myrmecophilous association of *Cheleion* and other Stereomerini.

1) There exist numerous well known myrmecophilous aphodiines, especially of the tribe Eupariini (see, e.g., Stebnicka 2009; Maruyama 2010). Those beetles usually live in debris in ant nests, fly well and are frequently collected with FIT or attracted at light.

2) In rather rare cases of presence of trichomes in termitophilous scarabaeids, those structures are not recorded from the pronotum and have a quite different appearance from *Cheleion* (see, e.g., Maruyama 2012a,b).

3) The peculiar structure on the pronotum in *Cheleion*, consisting of a central pit surrounded by numerous long microtrichia (= trichomes), is surprisingly similar to the pronotal structure of myrmecophilous ptinids (see, e.g., Bell and Philips 2008a,b), paussine carabids (Geiselhardt et al. 2007), elytral structures of myrmecophilous chlamidopsine histerids (e.g., Caterino and Degallier 2007), or elytral and abdominal structures of pselaphine (clavigerine) staphylinids (e.g., Nomura 1997, Baňař and Hlaváč 2014).

**Acknowledgements**

We thank Johannes Bergsten (NHRS) who allowed us to examine the holotype of *Cheleion malayanum* under his care and Eduard Jendek (Canadian Food Inspection Agency, Ottawa, Canada) for the donating of the type specimen of the new species to NMPC, his valuable information about the habitat where the specimen has been collected and for kindly providing the type locality photograph. Special thanks to Michal Tkoč (NMPC) for his help with taking the SEM micrographs. We are obliged to Peter Hlaváč (Prague, Czech Republic) for valuable comments to discussion, and to Robert B. Angus (School of Biological Sciences, Royal Holloway University of London, United Kingdom) for correcting the English of an earlier version of the manuscript. We also thank Mattias Forshage (Uppsala University, Sweden) and an anonymous reviewer for their helpful comments on the manuscript. David Král would like to acknowledge the
institutional support from resources of the Ministry of Education, Youth and Sports of the Czech Republic. The work of J. Hájek was partly supported by the Ministry of Culture of the Czech Republic (DKRVO 2015/14, National Museum, 0002327201).

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