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Corentin Jouault, André Nel. A new genus and species of Aulacidae (Hymenoptera: Evanioidea) from mid-Cretaceous Burmese amber. Cretaceous Research, 2021, 124, pp.104799. 10.1016/j.cretres.2021.104799. insu-03161875

HAL Id: insu-03161875 https://insu.hal.science/insu-03161875

Submitted on 8 Mar 2021

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PII: S0195-6671(21)00046-X

DOI: https://doi.org/10.1016/j.cretres.2021.104799

Reference: YCRES 104799

To appear in: Cretaceous Research

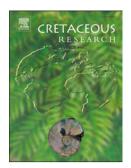
Received Date: 10 July 2020

Revised Date: 8 December 2020 Accepted Date: 23 February 2021

Please cite this article as: Jouault, C., Nel, A., A new genus and species of Aulacidae (Hymenoptera: Evanioidea) from mid-Cretaceous Burmese amber, *Cretaceous Research*, https://doi.org/10.1016/j.cretres.2021.104799.

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- A new genus and species of Aulacidae (Hymenoptera: Evanioidea) from
- 2 mid-Cretaceous Burmese amber

3

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11

12 ABSTRACT

- 13 The mid-Cretaceous amber from Kachin State of Myanmar provides a well-diversified fauna of au-
- 14 lacid wasps. Here we describe a new aulacid genus and species Paleoaulacus minutus. This discov-
- 15 ery highlights the underestimated diversity of aulacid wasps in Burmese Ablian–Cenomanian amber.
- 16 Interestingly, *Paleoaulacus minutus* gen. et sp. nov. shares wing venation and body characters with
- 17 the tribes Electrofoenini and Aulacini, challenging a clear placement. Future discoveries of addi-
- 18 tional specimens are necessary to precise its position or erect a new tribe to accommodate this pecu-
- 19 liar genus. This fossil shows a developed wing venation, similar to that of the extant genus Aulacus
- 20 Jurine, 1807.

21

22 **Keywords.** Hymenoptera, Evanioidea, Aulacidae, Cenomanian, fossil record

23

24 1. Introduction

The superfamily Evanioidea comprises the families Aulacidae, Evaniidae, Gasteruptiidae, †Praeaulacidae, †Anomopterellidae, †Andreneliidae, †Baissidae, and †Othniodellithidae. These wasps were clearly more diverse during the Mesozoic than in the extant fauna, with records of fos-sils of all these families. Notice that Engel et al. (2016), Engel (2017), and Turrisi & Ellenberger (2019) considered the praeaulacid subfamily †Nevaniinae as a different family but did not provide a clear phylogenetic argument or synapomorphy justifying such a placement. Thus we prefer to fol-low Zhang & Rasnitsyn (2007) and Li et al. (2018), keeping the †Nevaniinae as a subfamily of †Praeaulacidae.

Among the Evanioidea, the aulacid wasps are easily recognizable by an anterior margin of metasomal segment 1 distinctly separated from metanotum, and metatibia at most slightly widened toward apex (vs. metasomal segment 1 with anterior margin almost touching metanotum, and metatibia gradually enlarged or markedly swollen toward apex, appearing club-shaped) according to Goulet & Huber (1993). The Aulacidae comprise extant and fossil species, distributed in two subfamilies †Hyptiogastritinae (Engel, 2006, 2017) and Aulacinae (Shuckard, 1841). Furthermore, the monophyly of the family is well-supported (Jennings & Austin, 2000; Smith, 2001; Turrisi, 2006, 2007, 2014; Turrisi et al., 2009; Turrisi & Madl, 2013). Mesozoic Aulacidae are not uncommon but only known from Burmese amber (burmite, Kachin, northern Myanmar) inclusions (Turrisi & Ellenberger, 2019) with 11 described species (Cockerell, 1917a, 1917b; Engel, 2017; Li et al., 2018; Turrisi & Ellenberger, 2019; Jouault et al., 2020). Cockerell (1916a, 1916b, 1917a, 1917b, 1920a, 1920b) provided the first scientific studies on this amber. This resin is now renowned for its plethoric insect inclusions (Ross, 2019, 2020). One more time, it has delivered a new genus and species which is described an illustrated herein.

2. Material and methods

- 50 The amber piece containing the specimen studied herein derives from the deposits of Noije Bum in
- 51 the Hukawng Valley (26° 29′ N, 96° 35′ E), Kachin State, northern Myanmar (Fig. 1). Radiometric

02	data established an early Cenomanian age (98.79 \pm 0.62 Ma) for Kachin amber, based on zircons
53	from volcanic clasts found within the amber-bearing sediments (Shi et al., 2012). Some ammonites
54	found in the amber-bearing bed and within amber corroborate a late Albian-early Cenomanian age
55	(Cruickshank and Ko, 2003; Yu et al., 2019).
56	The new aulacid specimen is embedded in a piece of clear yellow amber. It is a nearly com-
57	plete individual but with damaged antennae, with the apical flagellomeres missing on right antenna
58	and left antenna broken. The type specimen is housed in the amber collection of the Geological De-
59	partment and Museum (IGR) of the University of Rennes, France. The specimen was examined and
50	photographed with a Nikon SMZ25 with an attached Nikon D800 camera. All images were digitally
51	stacked photomicrographic composites of several individual focal planes, which were obtained us-
52	ing HeliconFocus 6.7. The figures were composed with Adobe Illustrator CC2019 and Photoshop
53	CC2019 softwares. The wing venation terminology follows Jouault et al. (2020).
54	Published work and nomenclatural acts are registered in ZooBank (http://www.zoobank.org/, last
55	access: 07/12/2020), with the following LSID (reference): urn:lsid:zoobank.org:pub:478D05C9-
56	6F3D-4395-A7B0-45030873215B
57	
58	3. Systematic paleontology
59	Order Hymenoptera Linnaeus, 1758
70	Suborder Apocrita Gerstaecker, 1867
71	Superfamily Evanioidea Latreille, 1802
72	Family Aulacidae Shuckard, 1841
73	Subfamily Aulacinae Shuckard, 1841
74	See summary of fossil species in Table 1
75	
76	Genus <i>Paleoaulacus</i> gen. nov.
77	Type species: <i>Paleoaulacus minutus</i> sp. nov.

- 78 urn:lsid:zoobank.org:act:5FD961F6-7D8C-4D30-A80E-86F56A95A9B9
- 79 Etymology. Named after 'paleo' in reference to its antiquity, and the extant genus Aulacus Jurine,
- 80 1807. Gender masculine.
- 81 Diagnosis. Head hypognathous; compound eye very large, prominent, ovoid; forewing long and
- 82 narrow; venation well developed, with veins 1rs-m, 2rs-m present but spectral and 2m-cu tubular;
- 83 cell 1M narrow with parallel vein 1Rs+M and 1cu; vein 1M short (more than six times shorter than
- 1Rs); vein 2Rs+M short (twice as short as 2Rs); vein 1cu and M+Cu practically aligned.

- 86 Paleoaulacus minutus sp. nov.
- 87 urn:lsid:zoobank.org:act:7745BB59-C62E-40A1-BF7F-BB738EF1A7A4
- 88 Figs 1–2.
- 89 Material. Holotype IGR.BU-014 (male; preserved in a flat triangular piece of amber measuring
- 90 $8\times5\times3$ mm with a Psychodidae).
- 91 Locality and horizon. Noije Bum Hill, Hukawng Valley, Kachin State, Myanmar; upper Albian to
- 92 lower Cenomanian, mid-Cretaceous.
- 93 Etymology. From the Latin word *minutus* meaning 'small', referring to the small size of the species.
- 94 The specific epithet is to be treat as an adjective in a genitive case.
- 95 *Diagnosis*. As for the genus.
- 96 Description. Head globulous, practically hypognathous, higher than wide (in full face view); occipi-
- 97 tal carina absent; malar space narrow, almost inexistant; gena angled and slightly protruding; com-
- 98 pound eye very large, prominent, ovoid, without circum-ocular carina, inner margins slightly angled
- 99 (maximal distance between eyes near upper margin); ocelli small, arranged in a small triangle on
- top of vertex above compound eyes and separated by at least their diameter. Antennal toruli broadly
- separated, situated above head mid-length, distinctly separated from clypeus, near eyes apex; scape
- robust, 0.13 mm long, longer than wide, more than twice as long as pedicel, enlarged medially,
- slightly longer than or sub-equal to first flagellomere; pedicel almost as long as wide, 0.06 mm long;

104 flagellum filiform, with nine preserved flagellomeres, all longer than wide and cylindrical. Mandi-105 bles present but closed and not projected forward; palpal formula not clearly visible, at least com-106 posed by maxillary palps six-segmented and labial palps bi-segmented. 107 Mesosoma longer than high; pronotum short, smooth, without defined, raised posterior dorsal 108 surface bordering mesoscutum, posterior border concave; propleuron projected into a conspicuous 109 neck; mesoscutum flat in lateral view, with conspicuous edge along anterior border, with notauli not 110 connected; mesosctellum with slightly developed axilla; tegula visible; mesopleuron short. 111 Legs long compared to mesosoma, with slender podites; protibial spur clearly curved; tibial spur 112 formula 1-2-2; meso- and metacoxae closer to each other than to procoxa; trochantellus distinct but 113 fused to associated femur; profemur shortest; basitarsus elongate but sub-equal to combined lengths 114 of remaining tarsomeres; pretarsal claws small, simple; arolium minute. 115 Forewing with moderately wide costal space of comparatively consistent width along its length, 116 slightly narrower than pterostigma; pterostigma longer than wide, expanded in width medially, tapering gradually in width to acute apex; r-rs vein originating at widest part of pterostigma (slightly 117 118 distad pterostigmal midlength); cell 1Cu long, separated from cell 2Cu by angled and curved vein 119 1cu-a; cell R slightly wider than 1Cu; cell 1R1 large compared to other cells, vein 1Rs slightly 120 longer than 2Rs, both slightly curved, vein 2Rs+M short (more than twice as short as 2Rs), vein 121 1Rs+M straight and parallel to 1Cu, more than twice as long as 2Rs+M and enclosing cell 1M rec-122 tangular shaped, conspicuously longer than wide; cell 1M separated from 2M by a small vein 1m-cu sub-parallel to 2Rs and as long as 1M; cell 2Cu wider than long, slightly elongated basally with 123

Metasoma elongated in profile, cylindrical, longer than mesosoma; inserted well after metanotum; first metasomal segment long with anterior part modified into a petiole, remaining part cone

2Rs. Hind wing hidden under forewing, without visible venation except C+R.

presence of vein 2cu-a and separated from 2M by vein 2Cu; cell 2M elongated, enclosed by vein

2m-cu and 3Cu; cell 2R1 broad and elongated (twice as long as wide), with greatest width near vein

2rs-m; vein 1rs-m and 2rs-m spectral and subequal, slightly curved; cell 1Rs shorter but as wide as

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shaped; first tergite and sternite longer than remaining ones; tergites and sternites equal to sub-equal in lengths and separated by conspicuous sutures; last segment shorter than preceding ones. Hypopygium divided ventrally to form an elongate, apically rounded lobe on each metasomal side. Apex of gaster projected apically into rounded parameres.

Measurements (in mm). Total length as preserved (excluding antennae) 3.1; head length (excl. mandibles) 0.37; compound eye 0.31 long, 0.24 wide; mesosomal length 1.0, maximum height 0.47; dorsal length of propodeum (between metanotum and petiolar insertion) 0.19; metasomal length 1.35; profemur length 0.46, protibial length 0.43, probasitari length 0.31, combined lengths of remaining tarsomeres 0.23; mesofemur length 0.53, mesotibial length at least 0.5, mesobasitarsus length 0.27, combined lengths of remaining tarsomeres not measurable; metafemur length 0.53, metatibial length 0.62; metabasitarsus length 0.27, combined lengths of remaining tarsomeres 0.35; forewing length 2.1; hind wing length 1.2.

4. Discussion.

Paleoaulacus gen. nov. is assigned to the Aulacidae based on the geniculate antenna with a distinct elbow between scape and pedicel, which differentiates it from Baissidae. Also its propleuron is extending forward to form a pronounced 'neck' (not present in †Bassidae). However, the current family rank of Baissidae might be challenged since lacking clear apomorphies (Rasnitsyn, 2013) and retaining several plesiomorphies (listed in Engel, 2017). With 11 preserved antennomeres, and a reduced hind wing venation (viz. without 1r cell closed by tubular vein). Paleoaulacus gen. nov. vs. cannot be assigned to the †Praeaulacidae that have more than 20 flagellomeres. The family †Praeaulacidae is currently considered as the 'ancestral' group of Evanioidea and is characterized by a complete wing venation, except for some species of the subfamily †Cretocleistogastrinae (Rasnitsyn, 1988; Li et al., 2013). Hence, Paleoaulacus gen. nov. cannot be assigned to this family since it has not such a wing venation. Paleoaulacus gen. nov. has not the typical horn with toruli, the long scape and the high number of antennomeres (20-21) proper to the †Othniodellithidae. It cannot be

assigned to the †Anomopterellidae because it has only 12(?) antennomeres vs. at least 18 for the latter family.

Following Goulet & Huber (1993), the extant Aulacidae differ from the Gasteruptiidae *sensu* Engel (2017) (= †Kotujellitinae + Gasteruptiinae + †Hypselogastriinae + Hyptiogastrinae) in having an anterior margin of metasomal segment 1 distinctly separated from metanotum, and metatibia at most slightly widened toward apex (vs. metasomal segment 1 with anterior margin almost touching metanotum, and metatibia gradually enlarged or markedly swollen toward apex, appearing clubshaped in Gasteruptiidae). *Paleoaulacus* gen. nov., thereby, keys out as an aulacid wasp. Following the key to suprageneric groups of Aulaciformes of Engel (2017), *Paleoaulacus* gen. nov. keys out near the tribes Aulcini and Electrofoenini.

Engel (2017) proposed to use the relative positions of the eyes and antennae to separate Gasteruptiidae (see. Engel, 2017: Appendix: 'toruli never at or bordering ventral tangent of compound eyes') and Aulacidae (see. Engel, 2017: Appendix: 'toruli usually at or near ventral tangent of compound eyes'). Following this character *Paleoaulacus* gen. nov. also keys out as an Aulacidae. *Paleoaulacus* gen. nov. has a forewing venation developed as in some other Aulacidae, with the veins 1rs-m, 2rs-m and 2m-cu present (absent in Hyptiogastritinae). In view of all these characters, we assigned *Paleoaulacus* gen. nov. to the subfamily Aulacinae.

Following the key proposed by Engel (2017), *Paleoaulacus* gen. nov. would key out in the Aulacini because of the forewing with 2Rs+M shorter than 1Rs+M, and shorter than 2Rs *plus* short mandibles. However, *Paleoaulacus* gen. nov. clearly lacks the mesoscutum transversely carinate, challenging a confident placement within the tribe Aulacini. Moreover, *Paleoaulacus* gen. nov. has a metasoma with narrow tube-like petiole, sharply segregated from a cone-like base of the gaster. This character is unknown in Aulacini but occurs in some early diverged Aulacidae (e.g., *Archeofoenus* Engel, 2017 and some †Electrofoenini).

Similarly, *Paleoaulacus* gen. nov. could fall in the †Electrofoenini because of the mesoscutum not transversely carinate. It also shares with the recently described *Electrofoenia jehani* Jouault

et al., 2020 (in †Electrofoenini) a forewing with 2Rs+M shorter than 1Rs+M, and similar shape of the metasoma, unlike the remaining †Electrofoenini. Additionally, *Paleoaulacus* gen. nov. shares with most †Electrofoenini (except *Electrofoenia jehani* Jouault et al., 2020) subequal veins 2Rs and 1Rs but *Paleoaulacus* gen. nov. lacks the projected mandibles typical of the †Electrofoenini (see Engel, 2017; Jouault et al., 2020). The complete wing venation with veins 1rs-m, 2rs-m and 2m-cu present is recorded in the Cretaceous tribe †Electrofoenini (e.g. Engel, 2017; Jouault et al., 2020) but also in the tribe Aulacini (e.g. Chen et al., 2016; Ramage and Jouault, 2020).

Based on these comparisons, it appears that *Paleoaulacus* gen. nov. shares with both tribes Aulacini and †Electrofoenini different wing venation and body characters, challenging a confident placement within one of the two tribes. Therefore, we choose to not place our new genus within the Aulacini nor in the †Electrofoenini until the discovery of additional specimens.

Among Aulacinae, it seems that *Paleoaulacus* gen. nov., shares with the extant genus *Aulacus* similar habitus and wing venation (*e.g.* cell and vein disposition really similar, see Fig. 2C). However, it has a long and cylindrical metasoma while the two extant aulacid genera, *Pristaulacus* Kieffer, 1900 and *Aulacus*, usually have a trapezoidal one (see Goulet and Hubert, 1993; Chen et al., 2016; Ramage and Jouault, 2020). *Paleoaulacus* gen. nov. shares this putative plesiomorphic character state (see Engel and Wang, 2016) with the Gasteruptiidae from which it can be differentiated in its metasomal insertion clearly separated from metanotum. The Gasteruptiinae can also be distinguished by peculiar connected U-shaped notauli (synapomorphic character) (Grimaldi & Engel, 2005: 419), present since Early Cretaceous (Rasnitsyn, 1991), while the notauli are not connected in *Paleoaulacus* gen. nov. Basibuyuk and Quicke (1997) proposed to characterize the extant Gasteruptiidae by the presence of three hamuli. Unfortunately this character is not visible in the holotype of *Paleoaulacus* gen. nov. Moreover, *Paleoaulacus* gen. nov. does not share the synapomorphies of the aulacid sister group (†Kotujellitinae + Gasteruptiinae), viz. forewing vein 2rs-m spectral or conspicuous (absent or spectral in *Paleoaulacus* gen. nov.), without any indication of a meeting between Rs and M (while present in *Paleoaulacus* gen. nov.), and vein 2m-cu absent (pre-

sent in *Paleoaulacus* gen. nov.). *Paleoaulacus* gen. nov. would rather display the 'groundplan' venation features of Evanioidea (Basibuyuk et al., 2002).

Chen et al. (2016) proposed to separate *Aulacus* and *Pristaulacus* on the basis of the following characters: occipital carina absent in *Aulacus* vs. present in *Pristaulacus*; forewing with crossvein 2r-m, with 2Rs+M long in *Aulacus* vs. short in *Pristaulacus*; hind coxa of female without groove or notch on inner lateral surface in *Aulacus* vs. present in *Pristaulacus*; pretarsal claw not pectinate, without tooth-like processes along inner margin in *Aulacus* vs. pectinate, with tooth-like processes along inner margin in *Pristaulacus*. Even if we cannot discern the presence of a 'groove or notch on inner lateral surface of hind coxa', all the others characters allow us to assume that *Paleoaulacus* gen. nov. shares more characters with *Aulacus* than with *Pristaulacus*. However, *Paleoaulacus* gen. nov. differs from *Aulacus* in having more developed eyes (completely occupying the sides of head); cylindrical metasoma; cell 1M narrow with parallel to sub-parallel sides; and 2Rs+M short.

5. Conclusion.

The present description of *Paleoaulacus* gen. nov. suggests that the diversity of aulacid wasps in mid-Cretaceous Burmese amber is underestimated. *Paleoaulacus* gen. nov. possesses characters of the †Electrofoenini and of the Aulacini, challenging a clear placement in one of these two tribes. *Paleoaulacus* gen. nov. could belong to a new tribe, maybe 'intermediate' between extant and extinct tribes. By sharing with the extant genus *Aulacus* a similar wing venation, *Paleaoaulacus* gen. nov. shows an atypical stability of the wing venation though time; while the evolutionary tendency in the Aulacidae, and Hymenoptera as a whole, seems to have been towards a reduction and a simplification of the wing venation during geological time.

Acknowledgment

- We sincerely appreciate critical reviews and comments from the editor (Dr. Eduardo Koutsoukos),
- an anonymous reviewers, and Prof. Alexandr Rasnitsyn in improving our manuscript. We are grate-
- ful to Dr. Valérie Ngô-Muller (ISYEB, MNHN) who has kindly imaged the specimen.

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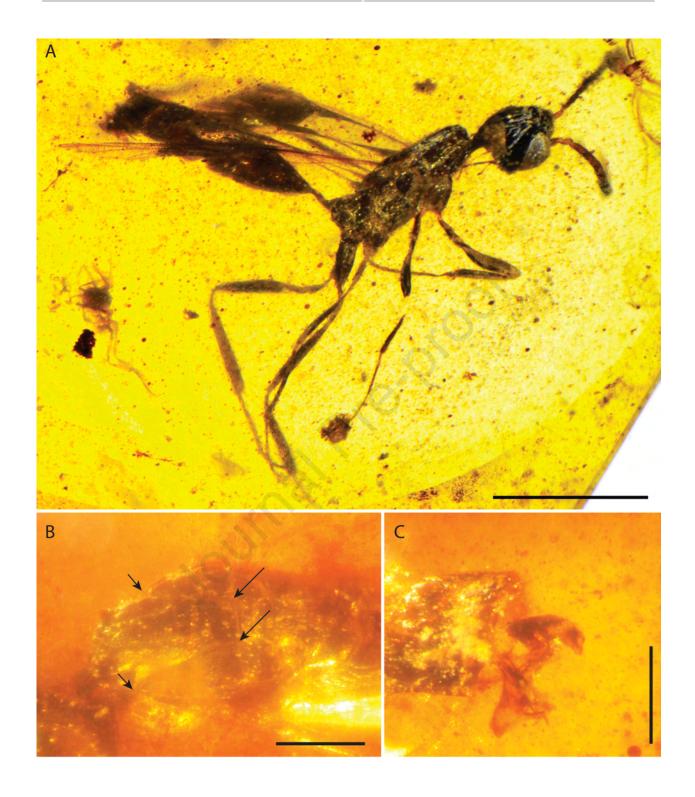
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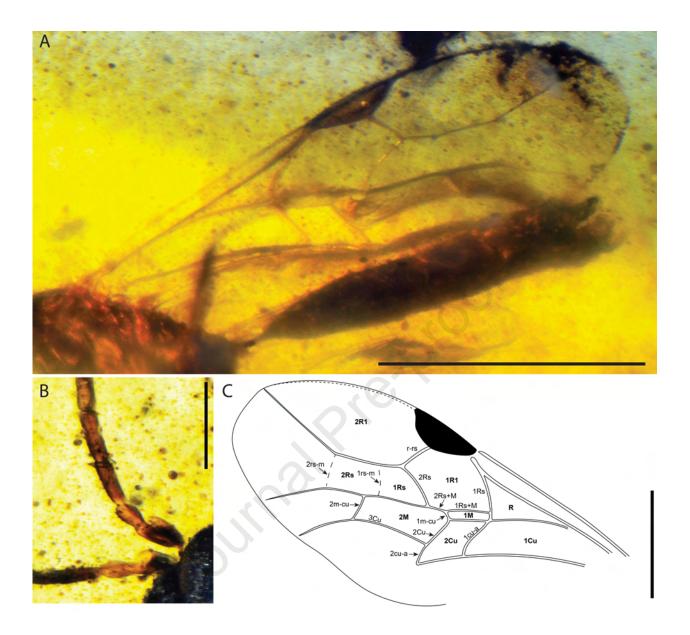
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335								
336	Figure caption							
337	Table 1. List of	fossil Aula	cidae with t	heir distribution.				
338	Figure 1. Paleoa	ulacus mir	<i>utus</i> sp. n	ov. Jouault & Ne	l, holoty _l	pe IGR.BU-014	4 A: Hal	oitus in side
339	view. B: Mes	sosoma in d	orso-lateral	l view (short arrov	vs pointi	ng bases of not	auli and	long arrows
340	pointing apic	es). C: Gen	italia. Scale	e bars: 0.5 mm (A); 0.25 m	m (B,C).		
341	Figure 2. Paleo	aulacus mi	<i>nutus</i> sp. r	ov. Jouault & Ne	el, holoty	pe IGR.BU-01	4. A: Ph	otograph of
342	wings. B: Ph	otograph of	antennae b	pasal part. C: Line	drawing	of forewing ve	enation (cell name in
343	bold). Scale l	bars 0.5 mm	n (A,C); 0.1	25 (B).				

	Species	Distribution	Period	References
Aulacidae				
Vectevania Cockerell	1	United Kingdom	Eocene	Cockerell, 1922
Hyptiogastritinae				
Hyptiogastrites Cockerell	1	Myanmar	Late Cretaceous	Cockerell, 1917
Archeofoenus Engel	2	Myanmar	Late Cretaceous	Engel, 2017; Turrisi and Ellenberger, 2019
Protofoenus Cockerell	1	Myanmar	Late Cretaceous	Cockerell, 1917
Aulacinae				
Electrofoenops Engel	3	Myanmar	Late Cretaceous	Engel, 2017; Turrisi and Ellenberger, 2019
Electrofoenus Cockerell	1	Myanmar	Late Cretaceous	Cockerell, 1917
Aulacus Jurine	1	France	Eocene	Nel et al., 2004
	2	Baltic amber	Eocene	Brues, 1933;
	1	USA	Oligocene	Bruce, 1910
Exilaulacus Li et al.		Myanmar	Late Cretaceous	Li et al., 2018
Pristaulacus Kieffer		USA	Eocene	Brues, 1910; Cockerell, 1916
riistaulacus Niellei		Baltic amber	Eocene	Brues, 1923 ;Brues, 1933; Jennings and Krogmann, 2009
Electrofeonia Jouault et al. Paleoaulacus Jouault & Nel		Myanmar Myanmar	Late Cretaceous	Jouault et al., 2020 Jouault and Nei, 2021





Jouault C: Conceptualization, writing original draft, visualization, analysis, revise manuscript. Nel A: Supervision, writing original draft, revise manuscript.

We declare that we have no conflict of interest

Yours sincerely

Corentin