On the first *Silis* Charpentier, 1825 from Baltic amber (Coleoptera, Cantharidae)

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Abstract. Only in recent years have new genera and species of the subfamily Silininae Mulsant, 1862 been described as inclusions in amber. However, no representative of the genus *Silis* Charpentier, 1825 had been described from Baltic amber, even if few specimens were already known at the generic level. *Silis lombardii* sp. nov. is entirely dark brown and shows (as usual for the genus) the two characteristic lobes in the sides of pronotum, elongated elytra, and a basal small tooth only on the anterior claws. The Eocene findings show that the subfamily is of ancient origin and that at least in the Eocene it was much more abundant than today in the same territories, where only two species are known. (urn:lsid:zoobank.org:act:76887127-3D24-41DA-86CC-7DCAB40DC3B7)

1 Introduction

The subfamily Silininae Mulsant, 1862 is widespread with a very high number of species in the eastern Palaeartic, Africa, South America and South-East Asia (Delkeskamp, 1939, 1977; Kazantsev and Brancucci, 2007; Constantin, 2010), but it is very poor in genera and species diversity in the western Palaeartic and Europe. Biodiversity hotspots appear to be former Indo-China, Indonesia and South America, even if the systematic knowledge at the generic level is still to be studied in depth (Ramsdale, 2002) for many regions and in particular for the Oriental region and North America. Also noteworthy is the fact that no species of the subfamily has reached southern Australia, Tasmania and New Zealand (Delkeskamp, 1977; Ramsdale, 2002). Recently, three extinct genera probably related to *Silis* and a species of the genus *Autosilis* Kazantsev, 2011 have been described from Baltic amber (Kazantsev, 2013; Alekseev and Kazantsev, 2014; Fanti and Damgaard, 2018; Fanti and Pankowski, 2018). Concerning the genus *Silis* Charpentier, 1825 in fossil records, the species *Silis chiapasensis* has been described from Miocene Mexican amber (Wittmer, 1963), and few specimens are known at the generic level for Eocene Baltic amber (Klebs, 1910; Bachofen-Echt, 1949; Fanti, 2017). These findings denote and suggest an ancient presence in Europe (Fanti and Damgaard, 2018) starting at least from the Eocene (no representative of the Silinae subfamily is currently known from the Cretaceous), and with a strong current rarefaction due to causes unknown to date. With the exception of *S. mingrelica* (Kazantsev, 1994) from Georgia, only another species, i.e. *S. ruficollis* (Fabricius, 1775), is currently present and widespread in Europe including the Baltic region (Kazantsev and Brancucci, 2007).

2 Material and methods

The amber piece was cleaned, polished and comes from a Yantarny mine in the Sambian Peninsula, Russia. The inclusion was photographed with a Zeiss Axiocam ICc 3 digital camera mounted on a Zeiss Stemi 2000c-stereomicroscope, with the addition of focus stacking software Helicon Focus, and with a Nikon D3100 digital camera equipped with AF-S DX Micro-Nikkor lens (40 mm f/2.8G). Figures were then processed and produced using PhotoImpact Viewer SE. Measurements were taken with an ocular micrometer mounted on a Leica S8APO microscope. The material is deposited.
in the University of Molise (Unimol), Italy, under accession code Unimol AAA003FP (AAA denotes Department of Agricoltura (agriculture), Ambiente (environment) and Alimenti (foods); 003 denotes the sequential number; FP denotes the Francesco Parisi collection).

3 Systematic paleontology

Family **Cantharidae** Imhoff, 1856
Subfamily **Silinae** Mulsant, 1862
Tribe **Silini** Mulsant, 1862
Genus **Silis** Charpentier, 1825
Subgenus **Silis** Charpentier, 1825

**Silis (Silis) lombardii** sp. nov.
(Figs. 1–3)

**Description**

Adult, winged, male-defined on the basis of the long antennae and elongate lateral lobes of the pronotum. Body length: 5.2 mm, elytra: 3.9 mm, antennae: around 3.2 mm. Entirely dark brown.

Head large, as wide as pronotum, slightly rugose, little exposed being partially covered by pronotum. Eyes elliptical and protruded, inserted in the upper and lateral part of the head. Maxillary palpi 4-segmented, unequal in length with the last palptomere very elongate and secundiform. Labial palps 3-segmented with the last palptomere secundiform. Antennae inserted near the eyes, just reaching the metafemora, 11-segmented, filiform, all antennomeres covered with small setae; scape elongate and enlarged, club-shaped; antennomere II (pedicel) extremely short and globular; antennomeres III–IV longer than antennomere II, robust; antennomere V slightly longer than antennomeres III–IV; antennomeres VI–IX slender and longer than antennomere V; antennomere X slightly shorter than antennomeres VI–IX; antennomere XI filiform, oblong with rounded apex. Pronotum slightly elongate, surface undulating with concavities and small setae and punctuation; apical margin convex; sides posteriorly expanding, with two elongated and apically pointed long processes: the posterior processes are sinuous and with a small basal acute tooth and the anterior ones are slender and straight. Scutellum triangular. Elytra wider than pronotum with slightly rugose microsculpture and some setae; elongate and covering the last abdominal segments; enlarged at humeri and narrow in the middle; apex strongly rounded. Posterior wings almost completely covered by elytra. Metasternum very elongate with rounded posterior margin, abdominal segments transverse and rugose. Legs long, slender and pubescent; coxae short and robust; trochanters elongate and triangular-shaped with rounded apex; femora enlarged and slightly curved; tibiae cylindrical, shorter than femora, with an apical spur. Tarsal formula 5-5-5; first tarsomere elongate and slightly enlarged apically; second shorter than first; third tarsomere triangular and shorter than tarsomere II; fourth tarsomere deeply bilobed at sides; fifth tarsomere very elongate, slender, flat and curved; proclaws with an obtuse basal tooth, meso- and metaclaws simple without basal tooth. Female unknown, sexual dimorphism is supposed.
Figure 2. *Silis (Silis) lombardii* sp. nov. in Baltic amber, holotype. 
(a) Detail of pronotum (arrow), scale bar = 0.4 mm. 
(b) Detail of right antenna, scale bar = 0.4 mm.

**Etymology**

Named in honour of Fabio Lombardi (Department of Agricultural Science, Mediterranea University of Reggio Calabria, Italy), dear friend and colleague of the first author.

**Holotype**

Male, in Baltic amber, deposited at the University of Molise (Unimol) with accession no. Unimol AAA003FP.

Figure 3. *Silis (Silis) lombardii* sp. nov. in Baltic amber, holotype. 
(a) Detail of pronotum (ventral view), elytra, legs and antennae with numbered antennomeres, scale bar = 0.5 mm. 
(b) Detail of last abdominal segments (lateral view), scale bar = 0.5 mm.

**Type locality**

Yantarny mine, Sambian Peninsula, Kaliningrad region, Russia.

**Type horizon**

Middle Eocene (Lutetian) (47.8–41.2 Myr) to Late Eocene (Priabonian) (37.8–33.9 Myr).
Syninclusions

Air bubbles and debris (botanical remains).

Remarks

The amber piece is flat and elongate and measures approximately 18 mm × 6–8 mm. The inclusion is complete and well visible. The specimen is crumpled but not damaged.

Systematic placement

Maxillary palpi sub-equal in length with last palpomere sec-uriform, elytra elongated, abdomen with only eight visible urites, protarsal claws with small tooth, meso- and metatarsal claws simple, and pronotum laterally with two long lobes in each part clearly make this new species belonging to the sub-family Silinae and to the genus Silis (Brancucci, 1980; Constantin, 2010, 2017).

Differential diagnosis

Silis lombardii sp. nov. differs from S. chiapasensis Wittmer, 1963 (Mexican Chiapas amber) in a slightly larger size, darker colour, and different shape of pronotum and lateral processes (Wittmer, 1963). Silis lombardii sp. nov. differs from the extant S. ruficollis in the body colour (S. ruficollis is black with red pronotum), the smaller size (5.2 instead of 6–7.5 mm in S. ruficollis) and in the more elongate lateral lobes of the pronotum.

4 Discussion

In temperate and northern Europe the genus Silis com-poses species of plains, rarely of higher altitudes, particularly present on shores of lakes or on marshy meadows (Moscardini, 1972). These habitats were typical in the Baltic Eocene, where no evident orogenetic events were reported (Sadowski, 2017). Thus, Silis is considered herein a hygrophilous genus and of temperate climate. Elsewhere, the genus Silis appears more thermophilic, since it is very frequent in the tropics (Constantin, 2010, 2017) suggesting that Silis can be paraphyletic. The elevated temperatures and the extreme envi-ronmental variety evidently favour the diversification and the evolution of the Silinae; for this reason, in Eocene amber many more species could be found again in the future. The thermal gradient and the subtropical environment present during the Eocene (Weitschat and Wichard, 2010; Sadowski et al., 2017) may, however, have played an important role in their spread in the Baltic area. The decrease in temperatures and annual precipitation at the end of the Eocene and contin-ued in the Oligocene (Weitschat and Wichard, 2010) reduced the number of taxa in northern temperate zones of Europe.

However, for example in the temperate zones of North America there is a higher number of species than in Eu-

tope (Arnett Jr., 1963; Green, 1966; Ramsdale, 2002). In contrast with Europe the biodiversity centre is in montane regions, particularly in California (Ramsdale, 2002). This fact suggests, rather, a Trans-Beringian Pliocene spread of temperate-cold species, as for example the fossil Opsimini (Coleoptera Cerambycidae) from Baltic amber (Vitali and Damgaard, 2016). The oldest finds of the Silinae subfam-
ily are the species included in Baltic amber to which an Eocene species, as adpression, from fossiliferous deposits of Florissant, Colorado, USA (Wickham, 1914; Fanti, 2017), and an undescribed specimen from the argillaceous lime-
stone of the Tertiary strata (Eocene or Oligocene) of Aix-
en-Provence, France (de Serres, 1843; Pictet, 1854; Fanti, 2017), are added. However, a much older evolution of the subfamily, referable to the Cretaceous as it is for the subfam-
ily Cantharinae (Fanti, 2017), is hypothesised.

Data availability. The material included in the paper is accessible in the Department of Agricultural, Environmental and Food Sciences of the University of Molise (Unimol), Italy, and all data are included in the description.

Author contributions. Both authors contributed equally to the article.

Competing interests. The authors declare that they have no conflict of interest.

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