

# The Larva of *Drusus nigrescens* Meyer-Dür, 1875 (Trichoptera: Limnephilidae: Drusinae) with notes on its ecology, genetic differentiation and systematic position

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The paper presents a description of the hitherto unknown larva of *Drusus nigrescens* Meyer-Dür, 1875. Information on the morphological and genetic identification of this species is given, and the most important diagnostic features are illustrated. Its systematic position within the genus *Drusus* is affirmed and some zoogeographical and ecological notes are added.

Keywords: Trichoptera, *Drusus nigrescens*, fifth instar larva, description, identification, distribution, ecology, mitochondrial DNA.

## Introduction

So far, 24 Drusinae species are reported from Austria, Germany and Switzerland (Lubini & Vicentini 2005; Malicky 1999, 2004; Robert 2001, 2004). Seven of them (*Drusus alpinus* (Meyer-Dür), *D. chapmani* McLachlan, *D. franzi* Schmid, *D. improvisus* McLachlan, *D. melanochaetes* McLachlan, *D. nigrescens* Meyer-Dür and *D. noricus* Malicky) are still unknown in the larval stage.

In the meantime, however, we managed to get larval specimens of *D. nigrescens* from Switzerland whose identity with the collected adults was confirmed by genetic analyses. This material enabled us to work out reliable diagnostic characters permitting integration of *D. nigrescens* in the key by Waringer & Graf (1997, 2004).

## Material and methods

Larvae which obviously were not yet included in existing Drusinae keys were collected from a spring brook in Switzerland where the dominant adult was *D. nigrescens*. To support species affiliation, larval and adult specimens were genetically analysed by means of DNA nucleotide sequence analysis following the methods outlined in Pauls (2004) and Pauls et al. (2006). A 498 base pair (bp) long section of the mtCOI region

revealed identical haplotypes between three previously unknown larvae with five adults of *D. nigrescens* (3 males, 1 female) from the same locality. One further male carried a haplotype which differed by one bp change. In Pauls (2004) a very close relationship between *D. nigrescens* and *D. monticola* was observed. We thus also generated mtCOI sequence data for 2 adults and 4 larvae of *D. monticola* and one adult of *D. melanochaetes*, a closely related species (Pauls 2004). We generated uncorrected pairwise distances between individuals using the DNADist function as implemented in BioEdit 7.0.5.3 (Hall 1999) (Table 1). No haplotypes were shared between species (Table 1). Haplotypes of *D. nigrescens* and *D. monticola* were, however, very close and interspecific uncorrected pairwise distance between haplotypes ( $p_{\text{nig/mon}}$ : 0.4-1.21%) was not much higher than within species variability ( $p_{\text{nig}}$ : =0.4%;  $p_{\text{mon}}$ : =0.82%) (Table 1). To further elucidate the close relationship between haplotypes of *D. nigrescens* and *D. monticola*, a median joining (MJ) network (Bandelt et al. 1999) was computed using the default settings in Network 4.2 (Fluxus Technologies 2005) (Fig. 11). The network shows that, although the relationship between *D. monticola* (light grey) and *D. nigrescens* (black) is very close, the two species are delimited. The species also remain differentiated and monophyletic in a three gene phylogeny of the Drusinae (Pauls, unpublished data).

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### Description of the fifth instar larva of *Drusus nigrescens*

Material examined: Five fifth and five fourth instar larvae and 14 males and 15 females from the Furka-Paß, Switzerland (46° 35,3' N, 08° 25,8' E, 2348 m asl), collected by W.G; 2 males, 1 female and 2 larvae collected by S.P. (Table 1). Two females and 4 larvae of *D. monticola*, and the female of *D. melanchaetes* were collected from various sites by W.G. and S.P. (Table 1).

Body length of final instar larvae: 8.7 - 10.5 mm; head width: 1.30 - 1.40 mm (head width of fourth instar larvae: 0.88 - 1.02). Length of larval case: 7.5 - 8.2 mm. The case is distinctly curved, tapering posteriorly (the mean width at anterior opening is 2.45 mm and at the posterior opening 1.45 mm) and consists completely of mineral particles with grain sizes increasing distinctly in anterior direction.

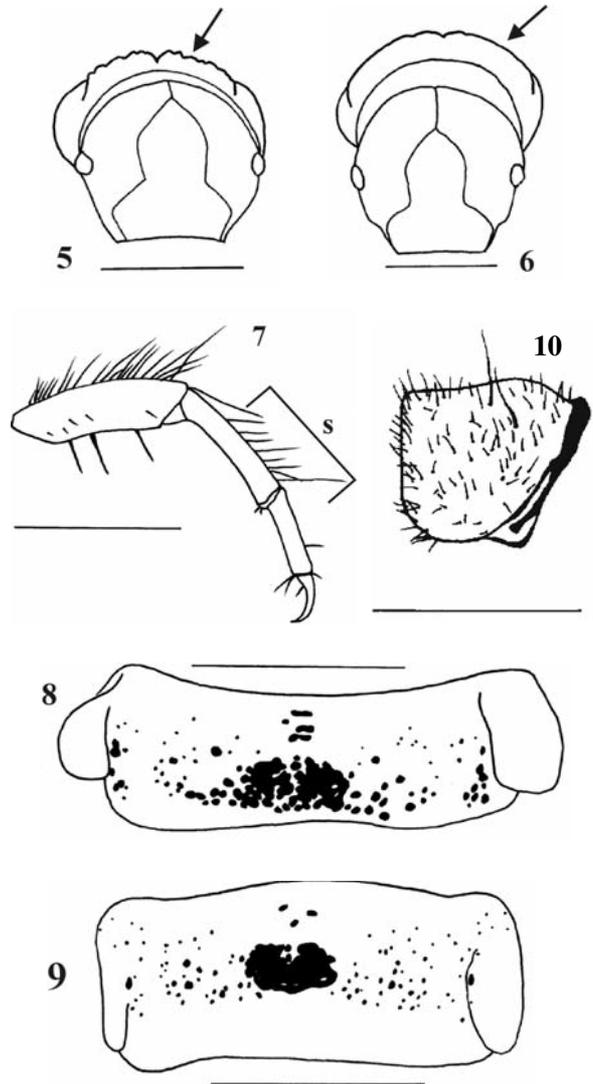
Head capsule and all body sclerites dark brown to blackish brown. The head capsule (Fig. 2) lacks additional setae or spines that are typical in some other Drusinae larvae (e.g. *Ecclisopteryx* spp., *Drusus trifidus*). Mandibles lacking terminal teeth along edges as well as ridges in the central concavity (Fig. 2).

In profile, dorsal line of pronotum with a distinct ridge in its posterior third (Fig. 2, 3) extending anterolaterally with its height decreasing (Fig. 3). The dark-brown pronotal surface is covered by black setae along the lateral borders, leaving the central pronotal area bare. Prosternite inconspicuous, prosternal horn present. Mesonotum completely covered by two chestnut brown sclerites. Metanotum partially covered by three pairs of sclerites; anterior metanotal sclerites large, ovoidal, their median separation being distinctly smaller than their maximum extension along the body axis (Fig. 1). Setal bases at the central section of the first abdominal sternum large and heavily concentrated, with a marked tendency of fusing, thereby creating a multi-lobed sclerotized pattern (Fig. 8). Overall setal counts at the sternum are up to 75, excluding the setal cover of the lateral protuberances. The sclerotized pattern is close to the structure observed in *Metanoea rhaetica* and *M. flavipennis*; however, the sclerotized plate in *Metanoea* is much less patchy and has a more clearly-defined outer border (Fig. 9) than in *D. nigrescens*.

Dorsal gills are present from the second (postsegmental position) to the fifth (postsegmental position). Ventral gills range from second (presegmental) to seventh segment (postsegmental). Lateral gills are present on the second and third segment (both in postseg-

mental position). Lateral fringe present from the posterior third of third to the beginning of eighth abdominal segment.

Setae are present at anterior and posterior faces of all femora. The row of dorsal setae at the mid and hind legs extend, proximally, well over the half of the tibia (Fig. 7).



Figs 5-10. 5: *Drusus nigrescens*, head and pronotum, frontal view (arrow: serrated pronotal ridge); 6: *Drusus monticola*, head and pronotum, frontal view (arrow: smooth pronotal ridge); 7: *Drusus nigrescens*, right midleg, posterior face (s= dorsal tibial setae); 8: *Drusus nigrescens*, first abdominal sternum (black: sclerotized setal bases); Fig. 9 - 10: *Metanoea flavipennis*, fifth instar larva; 9: first abdominal sternum (black: sclerotized plate and setal bases); 10: pronotum, left lateral view. Scale bars: 1mm.

## Morphological separation of *Drusus nigrescens* from other European Trichoptera

A summary of morphological features for the identification of limnephilid and Drusinae larvae is given in Waringer (1985). Within the framework of the limnephilid key by Waringer & Graf (1997, 2004), *Drusus nigrescens* is separated from other species by the following features:

- metanotum covered by three pairs of small sclerites (Fig. 1, m);
- head and pronotum without a thick layer of woolly hairs (Fig 2);
- head capsule without groups of additional spines, without central concavity and rims surrounding the frontoclypeus (Fig. 2);
- first abdominal sternum with a large median sclerotized patch (Fig. 8);
- Mandibles without teeth around edges, spoonshaped (Fig. 2, arrow).

At this position *D. nigrescens* keys out together with *Metanoea rhaetica* and *M. flavipennis*, where *D. nigrescens* is easily separated from the *Metanoea* species by the presence of a distinct sharp pronotal rim (Fig. 3); in *Metanoea*, the dorsal line of the pronotum in profile is evenly rounded in its posterior third, thereby creating a small dorsal hump (Fig. 10; Waringer & Graf 1997, 2004; Waringer, Graf & Maier, 2000).

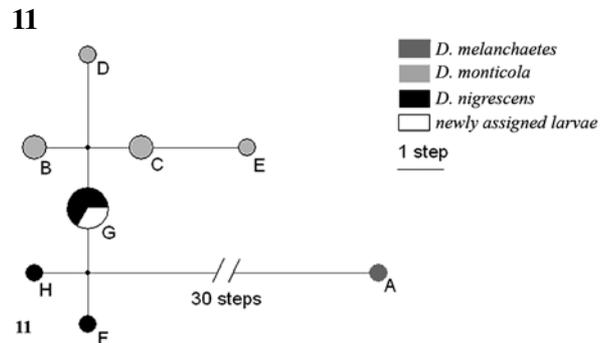
In case the sclerotized patch has been overlooked or interpreted as just an agglomeration of isolated setal bases, *D. nigrescens* keys out together with *Drusus monticola* (limnephilid key in Waringer & Graf (1997, 2004). Here, both species can be easily identified by the structure of the pronotal ridge in lateral (Figs. 3 & 4) and anterior view: in *D. nigrescens*, the ridge is serrate (Fig. 5), whereas in *D. monticola* the dorsal outline is smooth (Fig. 6). In addition, the dorsal setal fringes on the mid and hind tibiae are present within the proximal half of the tibiae (Fig. 7), whereas in *D. monticola* the setal fringes are restricted to the distal thirds of the tibiae. Finally, the agglomeration of sclerotized setal bases (Fig. 8) clearly identifies *D. nigrescens*; in *D. monticola*, sclerotized setal bases are few and widely separated.

## Habitat, phenology and distribution

Last and penultimate instar larvae of *D. nigrescens* were collected on 21 July 2006 at the Furka-Paß, Switzerland (46° 35,3' N, 08° 25,8' E, 2348 m asl). The small, spring-fed, 50 m long, first order tributary is part of the Mutt watershed; it is a clean, fast-flowing, summer-cold, mountain brook bordered by meadows

(Fig. 12). At this location, *D. nigrescens* was sympatric with *D. muelleri*, *Lithax niger* and Plecoptera such as *Dyctyogenus fontium*, *Protonemura lateralis*, *Leuctra ravizai*, *L. rosinae*, *Nemoura mortoni* and *N. sinuata*.

According to Malicky (2004), *D. nigrescens* is a western alpine species; records exist from Switzerland (Lubini-Ferlin & Vicentini 2005) and Italy (Piemonte and Lombardia; Cianficconi 2002), but the species is lacking in Germany (Robert 2001, 2004). In addition, doubtful records of *D. nigrescens* are known from Poland as well as the Ukraine (Czachorowski, personal communication). In Austria, it is reported from the Tyrol only (Malicky 1999). *D. nigrescens* is on the wing from July to September.



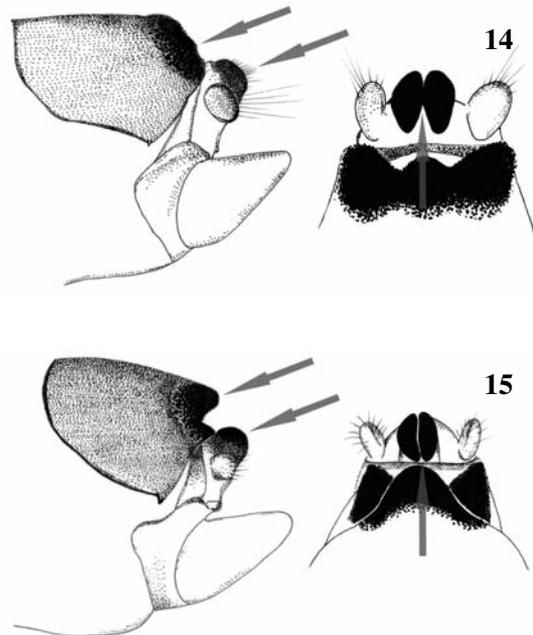
Figs 11-13. 11: Median-joining network of mtCOI haplotypes. Shaded circles represent haplotypes, haplotype names refer to Table 1. Shading coding indicates the species of specimens carrying a particular haplotype; diameter is relative to the number of specimens carrying a particular haplotype. Lines between haplotypes represent genetic distances between haplotypes; branch length is relative to distance, very long branches are shown with breaks. Small black circles represent median vectors. 12: View of the sampling site of *D. nigrescens* at the Furka-Paß, Switzerland, showing the small, spring-fed, first order tributary of the Mutt. 13: *D. nigrescens*, habitus of the coal-black male.

### Discussion

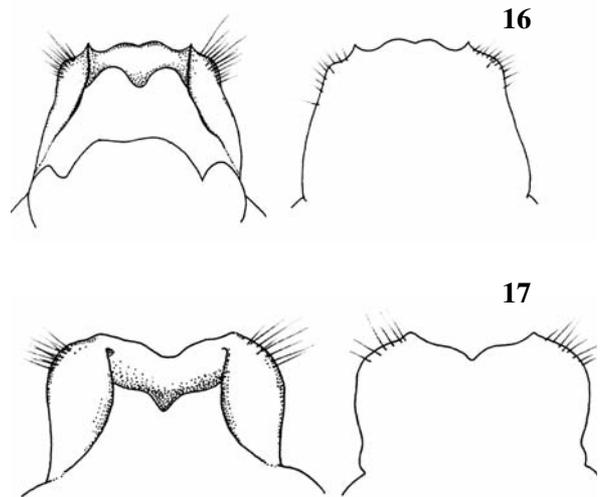
In his seminal work on the Drusinae, Schmid (1956) placed *D. nigrescens* within the *bosnicus*-group. Members of this group are dark brown in colour and have a large spinulated dorsal plate on the 8<sup>th</sup> tergite. The superior appendices are relatively small. The intermediate appendices are relatively large, prominent and are turned toward one another, but remain distinct. The inferior appendices are prominent and very divergent. Schmid (1956) considers *D. nigrescens* a very close neighbour to *D. monticola*, which was confirmed in our analysis of mtCOI in this study and in a preliminary phylogeny on the group (Pauls 2004). In fact our results show that, although both the larval and adult stages allow for clear morphological distinction, that divergence between the two species is minimal. The two species and their distinct morphologies are presumably the result of a very recent speciation event. More detailed molecular and morphological phylogenetic analyses are necessary to elucidate this divergence in more detail.

Schmid (1956) and Malicky (1983) illustrated both species, and greatly improved drawings of *D. monticola* and *D. nigrescens* are included, side by side, in Malicky (2004) together with the remark that identification is difficult. Although minute differences in both sexes are quite clearly shown in the latter publication, we add figures of the male and female genitalia, redrawn from own material (Figs. 14-17), to facilitate separation. Besides genitalia the most striking character is the general appearance, which is not shown anywhere. *D. nigrescens* is black as coal (Fig. 13) whereas *D. monticola* is brownish and much paler.

The current study presents the second example of a larval description based on molecular matching with adult caddisflies of the Drusinae (e.g. Graf et al. 2005). Molecular associations between sexes or life stages are becoming more commonplace in caddisflies (Shan et al. 2004) and other insect groups (e.g. Miller et al. 2005, Willassen 2005), exemplifying how nucleotide sequence markers can facilitate and provide supportive evidence in taxonomic and systematic research. However our study also shows the potential limits of using only DNA based-taxonomy between extremely closely related species, especially if lineage sorting is still incomplete (Pamilo & Nei 1988, Morando et al. 2004). The use of a single mitochondrial gene region as proposed for “DNA barcoding” may not be sufficient to resolve such situations, and the choice of the marker of utmost importance (Mueller 2006).



Figs. 14-15. Male genitalia (left lateral and dorsal view): Fig. 14: *Drusus monticola*; Fig. 15: *D. nigrescens*. The most important diagnostic characters are marked by arrows.



Figs 16-17. Female genitalia (ventral and dorsal view): Fig. 16: *Drusus monticola*; Fig. 17: *D. nigrescens*.

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