A key to the West Palaearctic genera of stoneflies (Plecoptera) in the larval stage

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Introduction

Plecoptera constitute a numerically and ecologically significant component in freshwater ecosystems, mainly in running waters of all sizes, all over the world. The fauna of the vast Holarctic Region is relatively uniform, and its stonefly families are essentially endemic to it, only some extend into the Oriental Region. The East Palaearctic and the Nearctic subregions are particularly similar, sharing the families Pteronarcyidae, Styloperlidae, and Peltoperlidae, and a number of genera and even single species that all lack from the West Palaearctic subregion. The wingless Scopuridae are endemic to Japan and Korea. Europe and the immediately adjacent parts of Asia and Palaearctic North Africa together are relatively distinct from the other subregions, not only by the absence of the beformentioned taxa but also by shared endemic genera.

Adult stonefly taxonomy is well advanced, the terrestrial imagines can reliably be identified, although in certain regions of the world and in some genera limitations remain. However, larvae are generally less well known than adults. Given the fact that many adults can be identified to species only by genital characters, a lastingly lesser taxonomic resolution must be anticipated in larvae, even after further study. In North America, excellent comprehensive literature on larval Plecoptera exists (STEWART & STARK 2002). The genera of the Russian Far East and Siberia have been treated synoptically, although in a less detailed way (ZHILTZOVA & TESLENKO 1997). There is a preliminary key to the genera of the incompletely studied Chinese fauna (HARPER 1994).

In Europe, there are several good regional generic keys to larvae in the relatively impover-ished faunas in the North and in the center of the continent. Central Europe holds an outstanding position through the work of the late JAROSLAV RAUSER (1980), who published a key to the species from former Czechoslovakia. However, all these keys are of limited use in many parts of South Europe with much more diverse faunas. These include a number of endemic genera whose larvae have been described, but not all in a comparative way, nor were they distinguished in keys.

The present paper keys all Plecoptera genera of the West Palaearctic Region. Brief comments on each genus are included. Where appropriate, suggestions or critical remarks on specific identifications of central European species are added, especially for species included in official lists of indicator organisms. There are also 2 appendices dealing with the German species of genus *Nemoura* and with selected species level taxa in genus *Leuctra*, respectively.

Geographic Range

West Palaearctic Region (i.e., Europe eastwards to Ural Mts and western shore of Caspian Sea; Anatolia, Caucasus, Lebanon and areas to the South, including Israel and Africa North of Sahara). Plecoptera are unknown from Libya and areas to the East, as well as from the Arabian Peninsula.

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Included Taxa

All genera in the above range. In genera occurring also outside the West Palaearctic Region the key is designed for West Palaearctic species, exotic species of the same genera cannot necessarily be identified with the present key. All of the West-Palaearctic families of Plecoptera belong in the suborder Arctoperlaria.

For monotypic genera, the single species is always named. For other genera, the attempt is occasionally made to identify selected species by characters listed in the text accompanying the genus.

Identifiable life stages

Characteristic character expressions of larvae develop stepwise, last instar larvae with fully formed wing-pads are optimal study objects. However, much younger larvae can also be identified with the present key but no larvae of less than *ca* 3 mm bodylength, or even first instars.

Counting of body segments

Segments of the trunk are counted from front to rear, parts of appendages from base to tip. Mind that of the 1st abdominal segment only the tergite is visible, the 1st sternite being fused to the metasternum. The first free, visible abdominal sternite is actually of segment 2. The female gonopore develops on sternite 8; its attachment scar moves backward at molts until it reaches the rear margin, interrupting the terminal hair fringe in its middle. The male gonopore is located at the end of sternite 9 but barely noticeable.

Anal gills

A collective term for gill filaments arising from the soft medial faces of paraprocts. Fine whitish filaments protruding out of the anus are no gills but gut fungi of the group Harpellales, which are common in Plecoptera. If one seizes them with forceps they can easily be pulled out of the rectum

Nomenclature and systematic arrangement

Nomenclature follows the 4th edition of the International Code for Zoological Nomenclature (1999). The world catalogues by ILLIES (1966) and ZWICK (1973) were used as base, Recent literature, year 2002 included, was added. The system is that of ZWICK (2000).

Illustrations

Illustrations are originals by P. ZWICK, except when differently indicated. Illustrations were drawn at a variety of maginifications, from whole specimens or mounted parts, in direct or transmitted light. Illustrations are not generally to scale, absolute measurements are given where needed. Overall body length (BL) of fully grown larvae is indicated under each genus.

"Hot-keys"

A few taxa of various taxonomic rank can immediately be recognized, by reference to single key characters:

- Gill tufts on thoracic segments: Perlidae.
- 4 branched cervical gills: *Amphinemura*.
- 6 tubuliform or sausage-shaped cervical gills: *Protonemura*.
- 1 three-segmented telescoping gill filament on the inner face of each coxa; abdominal tergites with unpaired projections: *Taeniopteryx*.
- Maxillary palpus with asymmetrically inserted tiny needle-like terminal segment: Chloroperlidae.
- Head trapezoidal, extremely setose: *Isoptena*.
- Thoracic dorsum shield-like, with wide parallel wing-pads, setose; very small: *Capnopsis*.

Superfamilies

- 1' No transverse suture crossing upper face of head between antennae and labrum (L), i.e. frontoclypeus (F+C) undivided (Fig. 1b). Labrum transverse, more than twice as wide as long. Mandibles flat and slender, without mola (Fig. 2b); in lateral view, mandibles are several times longer than high at the basal insertion; maxillary palpus long and slender (Fig. 1b). Glossae much shorter than paraglossae (Figs 6b, 7b). Coxae wide apart, thorax sternites between coxae extended, the area between furcal pits and spina about the size of a coxa (Figs 21b, c). Last tarsal segment about twice as long as two basal segments together; the latter are of equal size and very short (Fig. 3a) . Perloidea

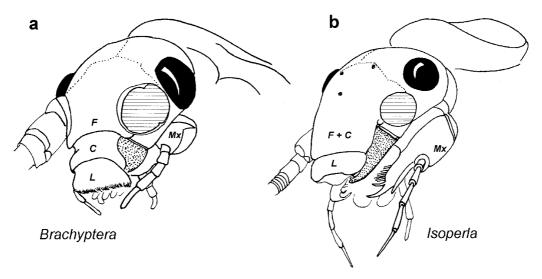


Fig. 1. Head, oblique anterior view. Left antenna removed, antennal foramen striate, left mandible stippled: Brachyptera sp. (a), Isoperla sp. (b). C = Clypeus, F = frons, L = labrum, Mx = maxilla.

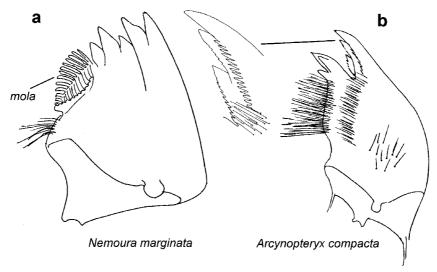


Fig. 2. Ventral view of left mandible of *Nemoura marginata* PICTET, 1836 and *Arcynopteryx compacta* (MCLACHLAN, 1872) (with enlarged apex of mandible).

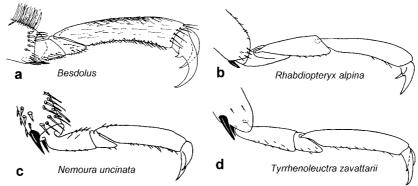


Fig. 3. Hind tarsus of *Besdolus imhoffi* (PICTET, 1841) (a), *Rhabdiopteryx alpina* (KÜHTREIBER, 1934) (b), *Nemoura uncinata* (DESPAX, 1934) (c) and *Tyrrhenoleuctra zavattarii* (CONSIGLIO, 1956) (d).

Families of Nemouroidea

Conventional keys rely mainly on the orientation of wing-pads. In Plecoptera, first small rudiments of wing-pads appear in the antepenultimate instar (ZWICK 2003), the definite shape is attained at subsequent molts. Using wing-pad position, reliable identifications are possible only for specimens in the two last instars. However, much younger larvae can be identified by less obvious characters; the present key uses position of wing-pads only as an auxiliary character.

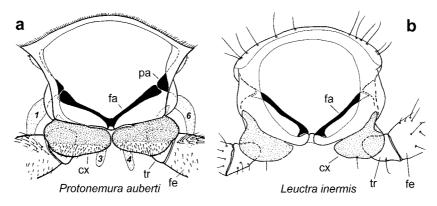


Fig. 4. Cross section through prothorax behind fore coxae, seen from behind (diagrammatic), only sceletal elements shown, internal ones exposed by the cut are in black. Left, *Protonemura auberti* ILLIES, 1954, only 4 of the 6 sausage-shaped gills (numbered) partly visible in background. Right, *Leuctra inermis* KEMPNY, 1899. cx, coxa; fa, furcal arm; fe, femur; pa, pleural arm; tr, trochanter.

- 2' More slender, extended hind legs do not attain end of abdomen. Wing-pads variably positioned, mostly parallel to body axis (Abb. 5c). Thoracic sternites distinctly sclerotized, furca- and spinasternum of mesothorax together forming a well-delimited Y-shaped area between the coxae. Furcal pits well visible. Coxae of normal shape, medially not extended and not strikingly projecting from sternal surface. Never with gills . 3
- Mentum plate-like, extending far sideward and forward, largely covering bases of maxillae (Fig. 5d). Wing-pads always parallel to body axis (Fig. 5c) Leuctridae
- 3' Mentum small, base of labium elongate, mouthparts completely exposed (resembling Fig. 5b). Wing-pads parallel to body axis except in one single rare genus whose wing-pads diverge (like in Nemouridae); however, the present genus lacks hairs ... Capniidae

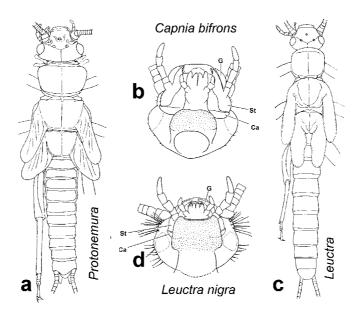


Fig. 5. Habitus of *Protonemura* sp. (a) and *Leuctra* sp. (c; from RAUŠER 1980). Head in ventral view, mentum stippled: *Capnia bifrons* (NEWMAN, 1839) (b) and *Leuctra nigra* (OLIVIER, 1811) (d). Ca = Cardo, G = Galea, St = Stipes.

Families of Perloidea

Larvae of this family have a distinctive habitus: they appear flat, with laterally extended flat legs with long setal fringes. Wing-pads little apparent even in the last instar, rounded on the outside and medially, separated by a wide shallow V-shaped notch.

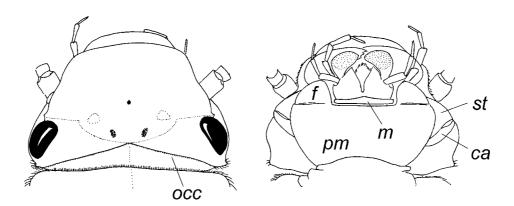


Fig. 6: *Dinocras cephalotes* (CURTIS, 1827), head in dorsal and ventral views; the cushion-like paraglossae stippled.

ca = cardo; f = front corner of postmentum; m = mentum; pm = postmentum; occ = occipital fold; st = stipes.

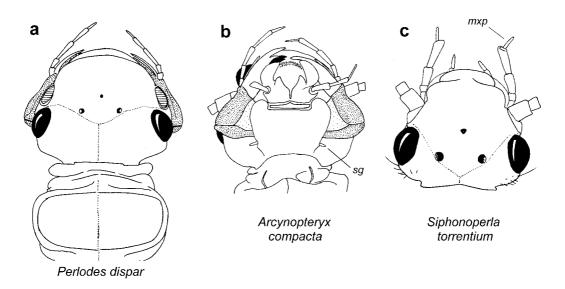


Fig. 7. a, *Perlodes dispar* RAMBUR, 1842, head and thorax in dorsal views; cardo, stipes and lacinia of maxillae stippled; antennae removed, foramina striate. b, *Arcynopteryx compacta* (MCLACHLAN, 1872), head in slightly oblique ventral view, only right eye and mandible (black) partly visible, only basal segment of labial palpus shown; cardo and stipes of maxillae stippled. c, *Siphonoperla torrentium*(PICTET, 1841), head in dorsal view; mxp = reduced, asymmetrically inserted terminal segment at tip of maxillary palpus. sg = submental gill.

Genera of Taeniopterygidae

Genus *Oemopteryx* KLAPÁLEK is not included. A single species, *O. loewii* (ALBARDA, 1899) once occurred in large European streams; it is missing for about 100 years. A few females are preserved in museums, but no single male. The larva was never found. Larvae of the 4 North American species of *Oemopteryx* differ between themselves, inferences on *O. loewii* are therefore impossible.

A telescoping 3-segmented retractable gill protruding from inner face of each coxa (Fig. 8a). Long-legged larvae lacking hair fringes and resembling Nemouridae except that they possess unpaired processes on adominal tergites (Figs 8b, c). In most species, entire abdominal dorsum appears serrate, sometimes also thorax, especially pronotum, with prominent processes. Sternite 9 normal, paraprocts freely visible

BL up to 13 mm. The only genus of subfamily Taeniopteryginae, unmistakable by the coxal gills. *Taeniopteryx* has a disjunct North Atlantic distribution. The numerous North American species lack tergal processes. Most of the approximately 10 Palaearctic species occur in South Europe, several of them also in central European mountains. In potamal streams of central European lowlands only *T. nebulosa* (LINNÉ, 1758) is found. It ranges from South to North Europe and East to central Asia. *T. nebulosa* can be identified by missing tergal processes on segments 8 and 9 (Fig. 8c) which the other Palaearctic species have (Fig. 8b). *T. araneoides* KLAPÁLEK, 1902 (wingless; Elbe, Danube) not found for over 100 years, larva unknown.

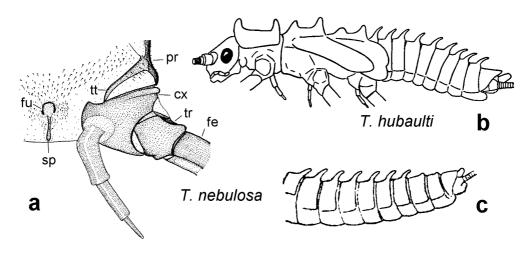


Fig. 8. a, *Taeniopteryx nebulosa* (LINNÉ, 1758), oblique ventro-anterior view of middle thorax, with 3-segmented telescoping gill originating from coxa. b, *T. hubaulti* AUBERT, 1946, lateral view, contour (after AUBERT, 1950). c, *T. nebulosa*, contour of abdomen, lateral view (after ILLIES, 1955). cx, coxa; fe, femur; fu, furcal pit; pr, pleural ridge; sp, spina; tr, trochanter; tt, trochantinus.

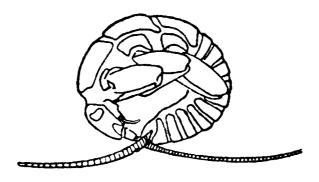


Fig. 9. Rolled larva of Brachyptera sp. (from ILLIES, 1955).

- - BL up to 12 mm. Endemic West Palaearctic genus. *B. risi* (MORTON, 1896) occurs in all of Europe and is the only Scandinavian species. In central European mountain streams *B. seticornis* (KLAPÁLEK, 1902) is the commonest species. Several species from potamal stream sections that had disappeared for decades were recently again found in a few places. Many species are endemic in the Mediterranean Region. The literature suggests specific colour patterns but there are examples of striking variation. Last instars can reliably be identified by structural details of sternite 9 and paraprocts illustrated by ILLIES (1955; Fig. 10c) and AUBERT (1959); allegedly distinctive colour patterns to be viewed with caution!

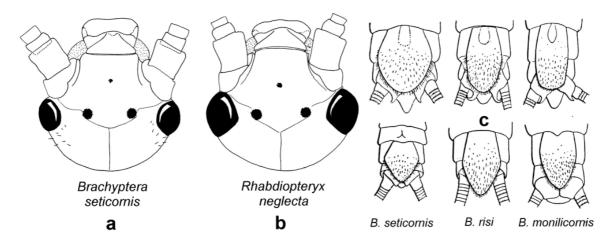


Fig. 10. Dorsal view of heads of *Brachyptera seticornis* (KLAPÁLEK, 1902) (a) and *Rhabdiopteryx neglecta* (ALBARDA, 1889) (b); note angle at which branches of occipital suture meet. *Brachyptera* spp., ventral view of abdominal tip with postgenital plate of males (top) and females (bottom) (c; from ILLIES, 1955)

Genera of Nemouridae

The present key works for larvae of near 2 mm body length or larger. The two genera with cervical gills do not yet have these in the first instar and the characteristic number and shape of gills develop stepwise, at early molts. *Nemurella* is in the literature normally diagnosed by a characteristic length relation of tarsal segments; however, this relation develops only late in larval life.

1	With cervical gills (Amphinemurinae)	 2
1'	No gills (Nemouringe)	3

² The larvae of some recently described Southern European species of *Rhabdiopteryx* are unknown, it cannot presently be tested whether the difference holds for all. However, according to AUBERT (1959), the Alpine *B. trifasciata* (PICTET, 1832) resembles genus *Rhabdiopteryx*. *B. trifasciata* occurred abundantly in Alpine rivers, However, it was entirely missing during the last 50 years but was recently again found, at few sites and in low numbers. I have seen no larvae of *B. trifasciata*.

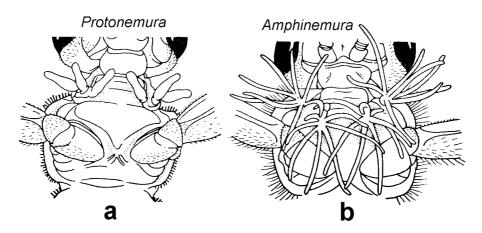


Fig. 11. *Protonemura* sp. (a) and *Amphinemura* sp. (b), ventral view of prothorax and neck showing sausage-shaped and branched gills, respectively (from RAUŠER, 1980).

3 Very long-legged: end of hind tibia lies clearly on coxa when folded back. Ground pilosity of femora weak, but several spreading setae in distal third exceed femur width. In late instars 1st and 3rd tarsal segments of hind legs about equal (Fig. 12c). Body normally held in a characteristic posture, with upcurved abomen and arched long antennae and cerci (Fig. 12a). Late instar male unmistakable, with enlarged upcurved paraprocts (Fig. 12b). Marginal setation of tergites consisting of only 4 large setae, others not larger than weak intercalary pilosity³ (Fig. 12d). Lateral setae longer and stronger than paramedian ones, setae on tergite 9 strongest. Setae along tibiae long, several longer than tibial width, exceptionally long erect sensory hairs, especially in small larvae. Occiput often strikingly lighter than dark purplish central portion of head. Setation of cerci sparse but strong and spreading, most developed and distinctly longer than segments around segments 5-7; distal setae fine and increasingly short. Small larvae (Fig. 12e) best identified by cerci attaining body length, unique long setae on several basal flagellar segments (Fig. 12g; disappear in larger specimens), long and strong pilosity of legs, and details of tergal and cercal setation described above

Nemurella KLAPÁLEK Only Nemurella pictetii KLAPÁLEK, 1900. BL up to 11 mm. From northern Spain to central Siberia. Common at sites with few other Nemouridae, especially in swampy springs, upper courses with diffusely entering groundwater, outlets of drain water, spring basins etc., but also in strongly acid waters like outlets of peat bogs. Occasionally said to be ubiquituous, but not resistant against pollution. Voltinism varies; most central European populations are bi- to trivoltine, univoltine in cool mountain sites, North European mountain populations semivoltine.

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³ The term "intercalary" designates pilosity on the surface of a sclerite, not at its edge, for example on the surface of tergites or on the surface of a cercus segment, between base and apical setal ring.

3'	Body shape, length of legs, shape of cerci and setation very different between the nu-
	merous species; 1st tarsal segment always shorter than last. Occiput of central Euro-
	pean species not distinctly paler than middle of head. If legs are long and slender, they
	lack very large spreading femoral spines. Tibiae usually with dense and strong setae
	along outer edges, but setae not longer than tibial width. Large setae different from
	intercalary tergal setation along tergal margins more numerous. Male paraprocts short,
	unmodified. Antennae never with long setae near base of flagellum
	BI mostly 8-9 mm, some species up to 11 mm. Many species stout and more plump than Namuralla

BL mostly 8-9 mm, some species up to 11 mm. Many species stout and more plump than *Nemurella*, cerci usually straight and shorter. See Appendix I for a preliminary key to German species.

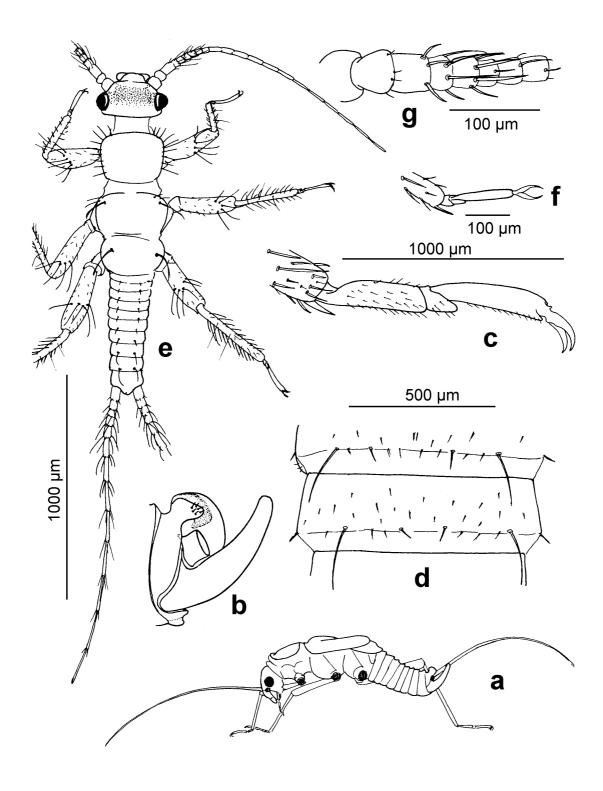


Fig. 12. *Nemurella pictetii* KLAPÁLEK, 1900. Characteristic curved posture (a; only right appendages shown), paraprocts of last instar male (b; lateral view), hind tarsus (c) and tergite 5 (d) of last instar specimen. Small larva, habitus (e), hind tarsus (f) and base of antennal flagellum (g). Figures without scale are not to scale.

Genera of Leuctridae

- 1' Pleural membranes divide also abdominal segments 5-8 into tergites and sternites 2
- 2' Head trapezoidal, widest posteriorly, narrowed in almost straight line towards front (Fig. 13a). Clypeus with prominent anterior corners. Pronotum transverse, ca. 1.5 times wider than long. Long erect hairs along sides of body and legs. Antennae with normal base, rapidly thinning apically, distally very fine and slender. Cerci multi-segmented but thin, thread-like, at base thinner than antennae (Fig. 13), miserable in comparison with large (up to 11,5 mm long) massive body Pachyleuctra DESPAX BL up to 11 mm. P. benllochi (NAVÁS, 1917) and two closely related species in the Pyrenees. Through their pilosity, half grown specimens greatly resemble Leuctra nigra (OLIVIER, 1811) and L. braueri KEMPNY, 1898; the latter has also angular front corners of clypeus. Both Leuctra species have normally shaped roundish heads (Appendix II, Fig. 34), and in both, pronotum, antennae and cerci are average.

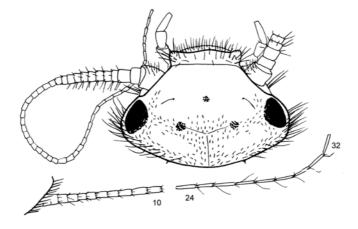


Fig. 13. Pachyleuctra sp., head and cercus (with segment numbers) of last instar larva,; all to same scale.

Genera of Capniidae

- 1' Pilosity inconspicuous, at most individual long hairs, larvae sometimes appearing completely bare. Abdominal segments 1-8 divided by pleural membrane. Pale larvae 3

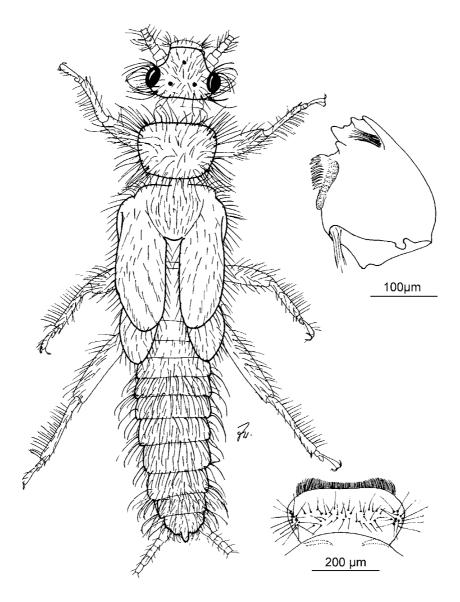


Fig. 14. Capnopsis schilleri (ROSTOCK, 1892), last instar larva, habitus (not to scale) and labrum in dorsal, left mandible in ventral view.

Genera of Perlidae

Among the West Palaearctic Plecoptera, the tufts of highly branched thoracic gills are characteristic. Each consists of 1 to 3 trunks which may fork into several branches each of which carries many straight fine filaments. These gills occur in characteristic positions to which the traditional gill nomenclature refers. It is followed here because it underlines homologies with similar gills observed also in other related families (within the infra-order Systellognatha). However, such examples do not exist among West Palaearctic representatives. The following gills are observed in Perlidae:

- a substigmatic gill below each of the 3 first spiracles;
- an anterior supracoxal gill above and in front of the fore coxa;
- a posterior supracoxal gill above the middle and hind coxa, behind the pleural fold.

1	Postmentum with transverse lines separating each front corner from main plate-like part (Fig. 6b); marginal setation of sternite 7 complete
1'	Front corners continuous with main part of postmentum; marginal setation of sternite 7 medially incomplete, setal fringe divided 4
2	Transverse fold across occiput regularly curved, crossing unpaired median branch of occipital suture distinctly behind forking point of suture. Long, club-shaped setae stand in a single row along transverse fold (Fig. 15b), also laterally where the fold is curved down regularly, merging into the sharp lateral edge of head below the eyes. Abdominal tergites only with fine pilosity contrasting markedly with strong marginal spines (Fig. 15c). Small; 2 pale longitudinal bands on abdomen (Fig. 15a)
	The only species is <i>Eoperla ochracea</i> (KOLBE, 1885), BL up to 17 mm. Strictly Mediterranean: North Africa, Spain, Southern France, Makedonia, Greece, Asia Minor. Absent from the Italian region.
2'	Transverse occipital fold angled forward, medially almost touching the forking point of occipital suture (Fig. 6a). Setae along transverse fold short, barely projecting over fold when viewing intact specimens. Well visible setae occur only laterally behind eyes where fold enters into sharp lateral edge of head, and are irregularly placed instead of in an orderly row. In slide preparations of exuviae, the setae are better seen; they are stouter than in <i>Eoperla</i> , almost spatulate (Fig. 15d). Abdominal tergites with dense cover of short fairly strong spines that differ less strikingly from marginal spines (Fig. 15e). No pale longitudinal bands on abdomen
3	not known. Transverse occipital fold bent angularly forward behind inner edge of eye and meeting
J	the fold running around eye and to side of head at a distinct, forward-directed angle ⁵ . Sidewards from this point, setation includes much longer pointed setae (Fig 16a, inset); cercus base never with seam of erect soft hairs
3'	Transverse occipital fold and fold running around eye merging in a regularly curved or only slighty undulating line (Fig. 18); no striking change in setation at fusion point. Cercus base with (Fig. 17b) or without seam of erect fine hairs

⁴ Much used in the world generic key to larvae of Perlinae (SIVEC et al. 1988); however, may be difficult to assess when there are fine and much smaller setae in the centre of setal fringe VII than on the sides which is not exceptional.

⁵ The same is true of the closely related East Asian genus *Oyamia* KLAPÁLEK.

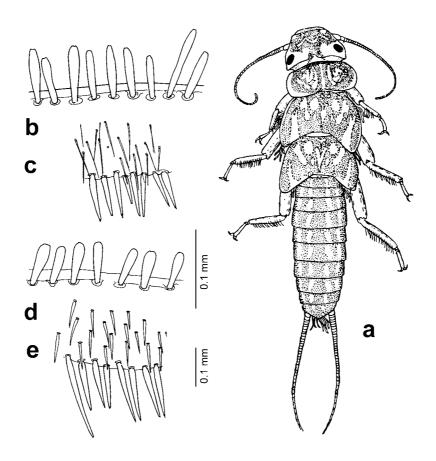


Fig. 15. Eoperla ochracea (KOLBE, 1885), habitus (a; modified from DESPAX, 1931). Setae along occipital fold (b) and detail of setation of tergite 5 (c). Dinocras cephalotes (CURTIS, 1827), setae along occipital fold (d) and detail of setation of tergite 5 (e). Figures b and d, and c and e, respectively, are to the same scale, respectively; mind that Eoperla is not even half the size of Dinocras!

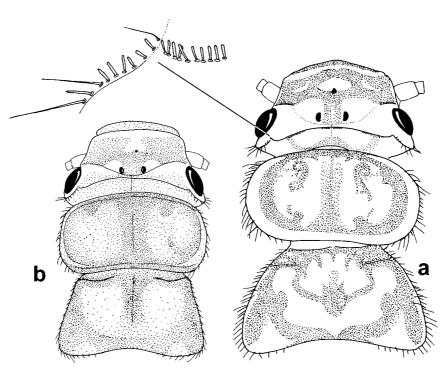


Fig. 16. Agnetina elegantula KLAPÁLEK, 1905, habitus of fore body (a) and detail of occipital crest and its pilosity at bend behind left eye (specimen from Austria, Burgenland, Lafnitz); *Marthamea selysii* (PICTET, 1841), fore body of brachypterous last instar male from River Moselle (b).

- Silky fine hairs on cercus form a dorsal fringe, especially on basal segments. Rings of dark spines striking already on basal cercus segments, not much longer on distal segments (Fig. 17a). Fine pilosity about half as long as segment, closely appressed to segment surface. Most species bright yellow with contrasting dark brown pattern 5

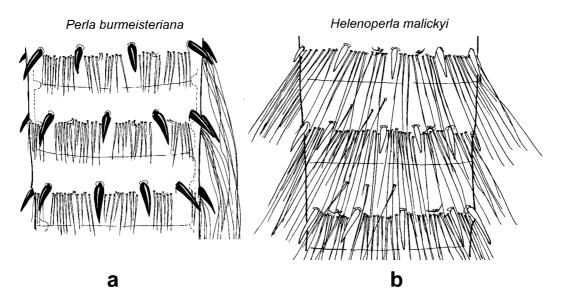


Fig. 17: Right cercus (dorsal side to the right), segments 6-8, of *Perla burmeisteriana* CLAASSEN, 1936 (a) and *Helenoperla malickyi* SIVEC, 1997 (b); not to scale.

BL up to 40 mm. *Perla* is endemic to the West Palaearctic Region. Species occur from North Africa and Great Britain to Iran; Cyprus is the only Mediterranean island with a *Perla* species (*P. caucasica* GUÉRIN, 1838). The genus does not inhabit North Europe. Species identities seemed to be well sorted out until SIVEC & STARK (2002) showed convincingly that species taxonomy is chaotic. There seem to be many more species than presently recognized. The authors studied only the eggs, other life stages can presently not be identified to the same degree. Therefore, specific identifications in *Perla* are presently preliminary and as of *status quo*.

Provided the study area is rigidly restricted to Germany ⁶, *P. marginata* (PANZER, 1799) can be recognized by lack of gills on paraprocts (so-called anal gills); head pattern distinctive (Fig. 18b). *P. burmeisteriana* CLAASSEN, 1936 has stout femora (hind femora about 3 times as long as wide), head pattern also distinctive (Fig. 18c). The long-legged (hind femur ca. 4 times as long as wide) Alpine larvae would (according to SIVEC & STARK 2002) all belong to a species presently called *P. grandis* RAMBUR, 1842.

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⁶ The polytypic *P. pallida* Guérin, 1838 (Slovenia and eastward) and *P. madritensis* RAMBUR, 1842 (Iberian Peninsula) are siblings of *P. marginata* and easily confused with it (SIVEC & STARK, 2002). *Perla carantana* SIVEC & GRAF, 2003 from Austria and Slovenia is indistinguishable from *P. burmeisteriana*, except in the characteristic egg stage (SIVEC & GRAF, 2003)!

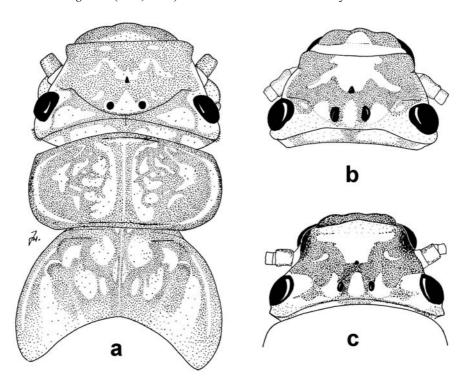


Fig. 18. Paragnetina spinulifera ZHILTZOVA, 1967, fore body (a; from ZWICK, 1971); dorsal view of head, Perla marginata (PANZER, 1799) (b) and P. burmeisteriana CLAASSEN, 1936 (c); not to scale.

Agnetina KLAPÁLEK BL up to 25 mm. The genus was for a long time known by the name *Phasganophora* KLAPÁLEK; see ZWICK (1984) for the confused history and the identity of the 3 West-Palaearctic species. Unfortunately, upon discovery (SCHOENEMUND 1925) the larva of *A. elegantula* was mistaken for *Marthamea vitripennis*; for a long time this severely hindered distinction of the two genera. Only *A. elegantula* KLAPÁLEK, 1905 occurs in central Europe. It is very rare and known from very few central European localities (GRAF, 1997). *A. senilis* KLAPÁLEK, 1921 and *K. werneri* (KEMPNY, 1908) are poorly known Caucasian and Anatolian species, respectively. The dark brown dorsal pattern on yellow ground of *A. elegantula* resembles the American *A. capitata* (PICTET, 1841) much. Figures of the habitus of *A. elegantula* are in SCHOENEMUND (1925; as *Marthamea vitripennis*) and GRAF (1997; photography).

Genera of Perlodidae

- 1' Paraprocts blunt, their sclerotised portion apically surrounded by soft, membraneous strip; dorsal cercal hair fringe long, distinct, also at base (Figs 19c, d). Body surface bare or with different kinds of pilosity; fine procumbent hairs as seen in *Isoperla* occur only in a single Spanish species with narrow unidentate lacinia. Occipital setation variable. Variable number of abdominal segments divided into tergite and sternite....... 2

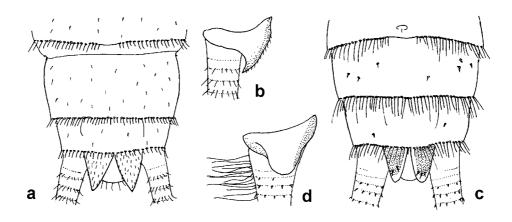


Fig. 19. *Isoperla* sp. (a, b) and *Dictyogenus alpinum* (PICTET, 1841) (c, d), abdominal tip of female larvae in ventral view, and lateral view of detached left paraproct and cercus. In the figure of *Dictyogenus*, stippling identifies sclerotized parts of paraprocts; the strongly bulging anal membrane is concealing both edge and setal fringe of tergite 10.

in intermittent streams; not in the Pyrenees. BL up to 14 mm.

- 4' An (often irregular or incomplete) row of setae across occiput, between rear margins of eyes (Fig. 21a). Body surface more or less densely pilose or setose. Lacinia variable, apical teeth diverge, setation along inner edge distinct (for example, Figs 20c, 22a, e) 6

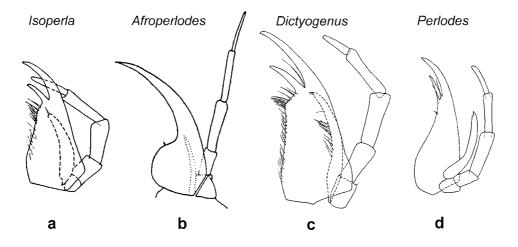


Fig. 20. Left maxillae of several genera of Perlodidae.

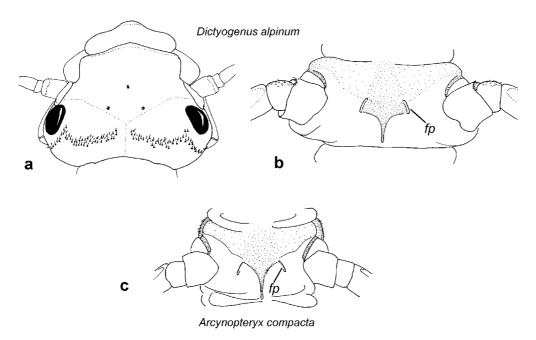


Fig. 21. Head and ventral view of mesothorax of *Dictyogenus alpinum* (PICTET, 1841) (a, b) and ventral view of mesothorax of *Arcynopteryx compacta* (MCLACHLAN, 1872) (c). Density of stippling varies to indicate degree of sclerotisation on the largely soft ventral face of thorax. Fp = furcal pits.

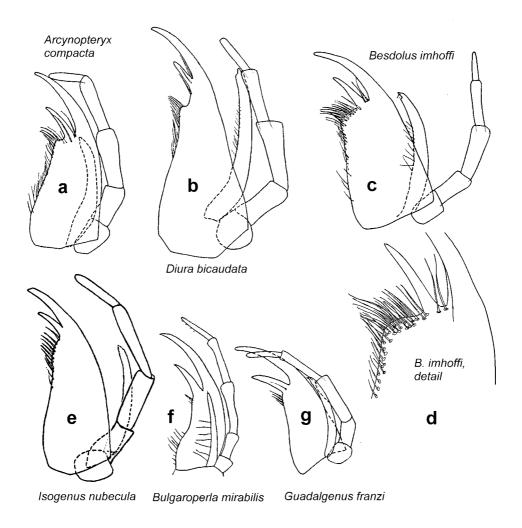


Fig. 22. Maxillae of several genera of Perlodidae; Bulgaroperla after BRAASCH & JOOST (1971).

Genera of Chloroperlidae

Family Chloroperlidae comprises two subfamilies. The 3 genera constituting the Paraperlinae are restricted to the East Asian mainland and to North America, while Chloroperlinae are of Holarctic distribution, with 5 genera in the West Palaearctic Region. Chloroperlinae are structurally very homogenous. While adults of the European species can easily and reliably be assigned to the various genera by genital characters, distinction of larvae (except *Isoptena*, which is globally unique and cannot be mistaken for any other stonefly larva, from the first instar on) is tricky, in all faunal regions.

The various European species within a given genus differ a little between them which renders generic distinctions difficult; for example. *S. montana* (PICTET, 1841) has resemblances with *Chloroperla*-species. At the same time, the various species are not sufficiently characteristic to be recognized at the specific level. In a regularly sampled population of *S. torrentium* (PICTET, 1841), ontogenetic change in all characters was found to be important, creating great problems. For example, not only segment number but also general shape and relative length of cerci change importantly during growth, from short with few segments and a strongly concical contour in early instars to relatively longer, with up to 16 segments and a much more slender overall shape in the last instar. Similarly, eyes of early instars are much smaller in relation to the head capsule than they are in later and especially in the last instar. Early instars

tend to be unicolorous and generally pale, but patterns may develop and be distinct in late instars; however, no comparative study of the patterning of the European *Siphonoperla* has so far been made. *Chloroperla* larvae tend to be generally pale.

The present key is based on study of last instar specimens. Users are warned to attempt to identify single young specimens. Only if much material collected at the same site over a longer period of time is available it may be possible to follow ontogenetic change and assign also middle sized larvae to a particular taxon with confidence.

Present illustrations are of mounted whole last instar specimens observed in transmitted light and only this way (or at best when specimens are viewed from all sides in strongly oblique illumination) can the length of the sometimes exceedingly thin and usually colourless setae be precisely recognized. In slide mounts, and consequently also in the present illustrations, not all setae are in their natural positions. Therefore the direction in which they point in the figures is not decisive. The present figures never show the very fine procumbent ground pilosity. Four pore-like sensilla on the head are visible only in transmitted light; they provide no means for distinctions.

Proportions of pronotum mentioned in the key refer to measured (not estimated!) distances between insertion points of setal rows surrounding the notum. The notal sclerite itself is frequently only partly delimited by a fine edge resembling a delicate line.

- - BL up to 11 mm. The single species, *I. serricornis* (PICTET, 1841), occurs in North and central Europe, mainly in areas with Pleistocene sands where larvae live in shifting sand at the bottom of streams, sometimes at considerable depth.
- 1' Shape normal, pilosity sparse and usually soft (for example, Figs 24, 25) 2

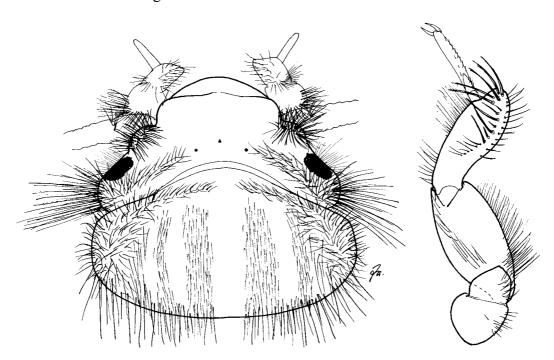


Fig. 23. Isoptena serricornis (PICTET, 1841), dorsal view of head and prothorax; ventral view of right foreleg.

- 3' Various patterns, no longitudinal abdominal bands ⁷. At least some cercal setae as long as or longer than the respective cercus segment (Fig. 25a) Siphonoperla ZWICK BL up to 12 mm, most species slightly smaller. Endemic in the West Palaearctic Region, about 9 species. Whether the North African S. lepineyi (NAVÁS, 1935) belongs here is not entirely clear. S. torrentium (PICTET, 1841) is the most common and most widely distributed species; in Scandinavia occurs only S. burmeisteri (PICTET, 1841).
- 4' Elongate pale larvae whose pronotal sclerite is distinctly delimited in front and posteriorly (Fig. 24c). Pronotal hair fringes much denser than in preceding species (Fig. 24c), hair fringe on tergite 10 much more slender. Setation of cerci along inner face near base short, not as long as segments. However, distally individual setae along dorsal and ventral faces of cerci are very long and fine (Fig. 25d) *Chloroperla* NEWMAN BL up to 8 mm. Endemic in central and South Europe and Anatolia. The typical species, *C. tripunctata* (SCOPOLI, 1763), is very widely distributed, preferably at moderate elevations. Regionally endemic additional species occur in the major European mountains, at higher elevations. Larvae apparently live relatively deep in stream bottom substrata, probably in the hyporheic zone. *C. tripunctata* has 5-7 very long dorsal and ventral hairs on each distal cercus segment, while *C. breviata* NAVÁS, 1918 (Pyrenees) and *C. susemicheli* ZWICK, 1967 (Alps) have only about 3 on each side (Fig. 25d).

⁷ Xanthoperla may have paramedian sections of the tergal antecostae darker than the rest, which may resemble banding; however, the tergite itself behind the darker portion is not pigmented!

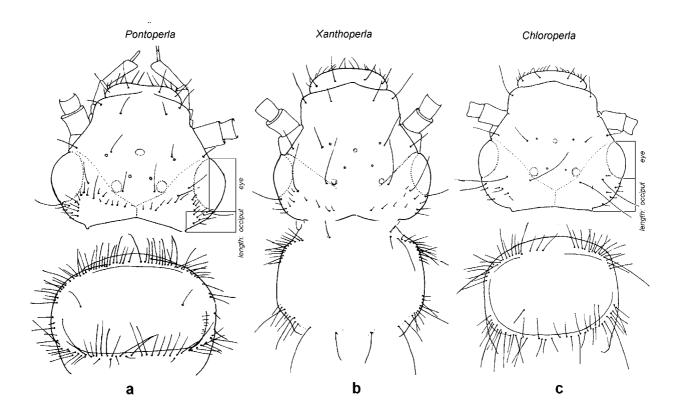


Fig. 24. Head and pronotum of *Pontoperla teberdinica* (BALINSKY, 1950) (a), *Xanthoperla apicalis* (NEWMAN, 1836) (b) and *Chloroperla susemicheli* ZWICK, 1967; whole mounts, in transmitted light.

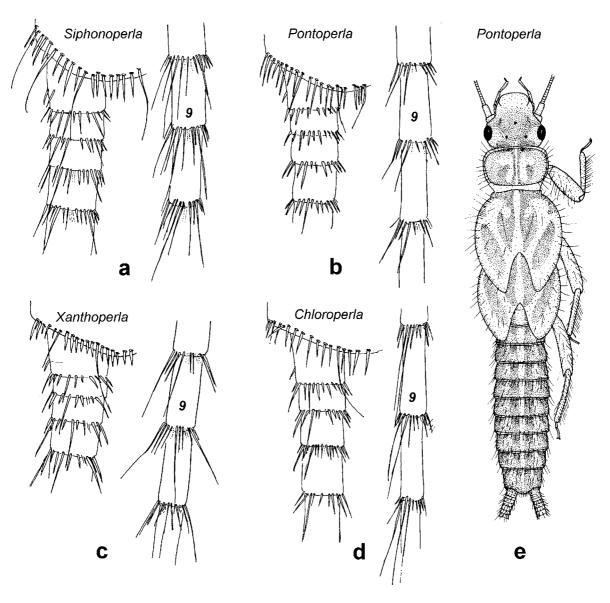


Fig. 25. Dorsal view of left cercus (base and segments 9 & 10) of *Siphonoperla torrentium* (PICTET, 1841), *Pontoperla teberdinica* (BALINSKY, 1950), *Xanthoperla apicalis* (NEWMAN, 1836) and *Chloroperla susemicheli* ZWICK, 1967; whole mounts in transmitted light. To the right, last instar larva of *Pontoperla teberdinica* (from ZWICK, 1971).

Appendix I:

Preliminary key to the German species of genus *Nemoura*, with notes on a few South European species

RAUŠER, in his masterly pioneer work (1980) provided a key to the Czechoslovakian species of *Nemoura* in which the relative length of setation on particular cercus segments is of prime importance. This approach requires that segments be accurately counted because the diagnostic proportions change along the cercus. Generally, the relative length of setae increases from the base to about the basal third of the segment and then decreases again. Although these characters are basically valid, there are two problems. First, the cercus grows during development by addition of segments near the base, similar to the base of the antennal flagellum where new segments are also added. Even in last instar larvae, this basal cercal area is usually incompletely segmented, the number of dividing sulci on the outside may exceed the number on the inside. Similarly, the most basal setal fringes at the end of segments are often only developed on the outer side. Therefore, the "first" segment after the cercus base is hard to identify. Second, the angle at which setae diverge from the segment varies, which influences estimates of relative length. These details may also be perceived differently in slide mounts than in intact specimens viewed in direct light under a dissecting microscope.

An alternative preliminary key was proposed by ZWICK in a lecture handout, in 1993. Although not formally published this document was nevertheless circulated by users. The present key is a re–arranged and extended version of this earlier key, a definite key to all species can still not be presented. Because of the need for future work and also because of the wider geographic frame of the present key at generic level, a few southern European species (whose larvae were previously inadequately described, or completely unknown) are also included.

Exuviae of reared males or mature males in their larval skin of all German species were available. Intact specimens were studied under direct light and slide mounts of entire larvae (or of exuviae) in transmitted light. Magnifications ~100x are sometimes required.

- Setae along caudal margin of abdominal tergites short, the longest barely half as long as the corresponding tergite, of normal spine shape. Setae around pronotum and along wing margins pointed. Setae on cerci inconspicuous, the longest not quite as long as the respective segment. Pedicellus concolorous with scapus and antennal flagellum
 - Ubiquituous stream species, tolerates some pollution, found also in acid peat bogs. Rare in typical Plecoptera habitats, but may be very common in marginal habitats that other Plecoptera fail to colonize.
- 2' Setation along caudal margin of abdominal tergites heterogenous: most setae very short and apically truncate and frazzled (Fig. 28), while several others are extremely thin, pointed and attain segment length; they are apically thinned and turn into a very fine, thin, flexible hair that is easily overlooked. Setae along pronotal and wing margins

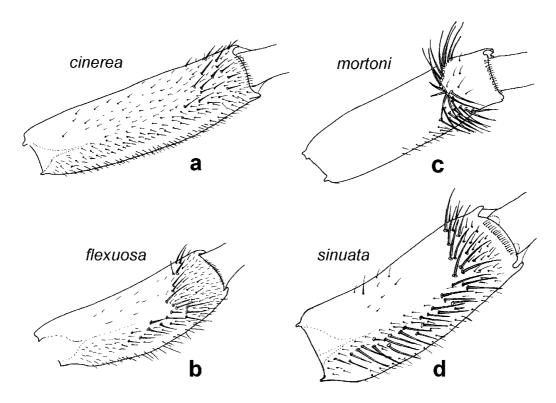


Fig. 26. Nemoura spp., right front femur in dorsal view; not to scale.

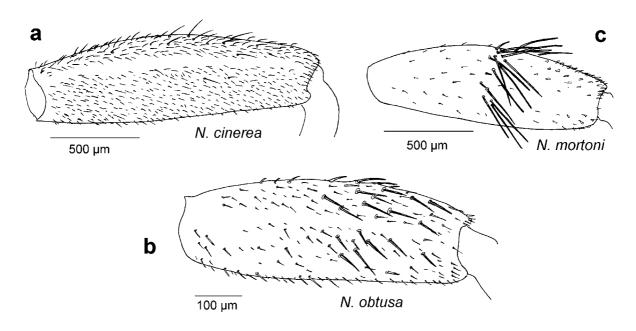


Fig. 27. Nemoura spp., examples of shape and setation of hind femur.

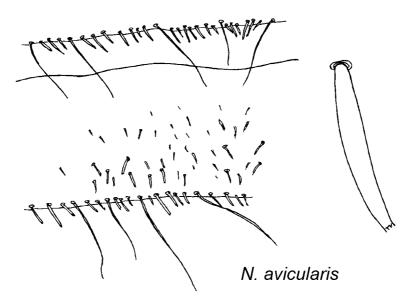


Fig. 28. *Nemoura avicularis*, left half of tergites 4 and 5, and detail of a marginal seta with frazzled apex; not to scale.

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⁸ LILLEHAMMER (1988) says the first tarsal segment to be curved which is not the case in German specimens.

6' Tergal marginal setation consisting exclusively of pointed and rather long sparse spines, the longest about as long as the corresponding tergite. Pronotal setation a little wavy...

palliventris AUBERT, 1953
Widespread rhithral Italian species, from the Maritime Alps to Sicily.

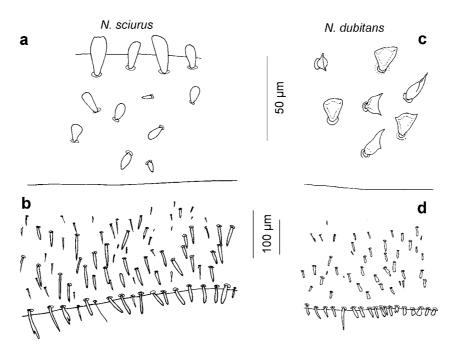


Fig. 29. *Nemoura sciurus* AUBERT, 1949 (a, b) and *N. dubitans* MORTON, 1894 (c, d). Setae on disc of pronotum, near midline (and along front margin, for *N. sciurus*; a, c) and left half of tergite 5 (b, d).

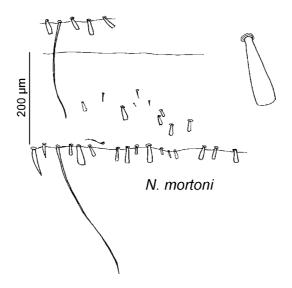


Fig. 30. Nemoura mortoni RIS, 1902, left half of tergites 4 and 5; detail of a spatulate marginal seta not to scale.

- 7' Wing-pads with fine, erect long hairs in linear arrangement......lacustris E. PICTET, 1865 Along tiny, slow-flowing streams in southern France and on the Iberian Peninsula.

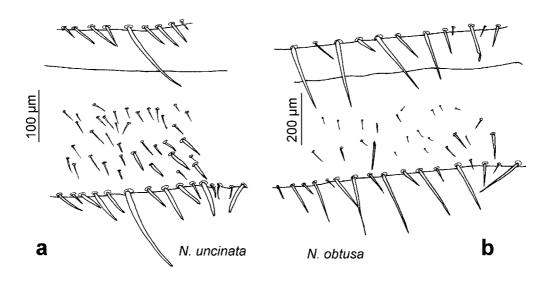


Fig. 31. *Nemoura uncinata* DESPAX, 1934 and *N. obtusa* RIS, 1902, left half of tergites 4 and 5; to different scales.

- 9 Setae on antero-lateral margin of pronotum acute. Tiny setae in lines on wing-pads are small pointed spines and occasional slender hairs (Fig. 32a, b)............ obtusa RIS, 1902 Springs and little streamlets in the Alps. A single record from the Polish Tatra.
- 9' Setae on antero-lateral margin of pronotum almost cylindrical, only apex narrowed to a short point. Tiny setae in lines on wing-pads ovoid (Figs 32c, d) sinuata RIS, 1902 Springs and little streamlets in the Alps.

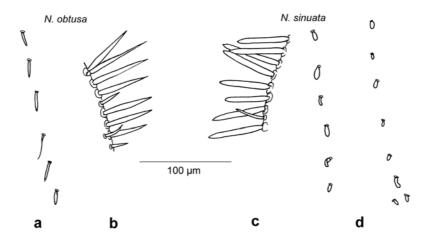


Fig. 32. *Nemoura obtusa* RIS, 1902 (a, b) and *N. sinuata* RIS, 1902 (c, d), rows of tiny setae on wing-pads (a, d) and lateral marginal setae of pronotum, near anterior corner (b, c).

Appendix II:

Diagnoses of selected characteristic species and species groups of Leuctra

RAUŠER (1980) offered a key to the Czechoslovak species of *Leuctra*. However, in this very large and difficult genus specific identifications remain problematic, with few exceptions. Below, new diagnioses are presented for a few individual species that can be reliably identified, and for two major species groups that are proposed as Operational Taxonomic Units (OTU) until more precise identifications become possible.

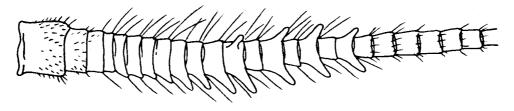


Fig. 33. Leuctra geniculata (STEPHENS, 1835), base of antenna (from RAUŠER, 1980)

- Very hairy, especially head (Fig. 34) and sides of pronotum:
 - Clypeus with prominent pointed corners (Fig. 34a); basal part of antennal flagellum with striking long setae. Lower side of head below antennal insertion normal. Found mainly in deep aerated deposits of fine detritus L. braueri KEMPNY, 1898

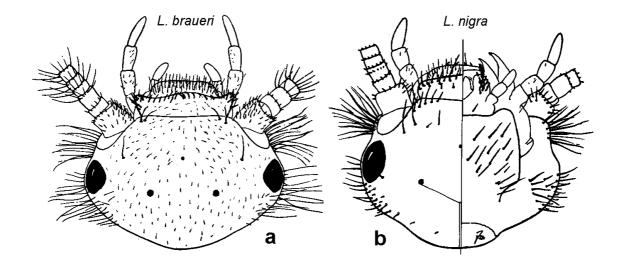


Fig. 34. Leuctra braueri KEMPNY, 1898 (a), head in dorsal view, and L. nigra (OLIVIER, 1811) (b) head in combined dorsal and ventral views.

- Macroscopic pilosity of normal density, or sparse, sometimes almost absent:

Notes on central European members of the *prima-hippopus-inermis*-group:

- Some members of the *prima*–subgroup (for example, *prima* KEMPNY, 1899, *autumnalis* AUBERT, 1948, *signifera* KEMPNY 1899, *pseudosignifera* AUBERT, 1954, probably also others) appear relatively hairy because hairs are stiff and tend to be well pigmented.
- *L. hippopus* KEMPNY, 1899 and the species of the *inermis*—subgroup (for example, *L. inermis* KEMPNY, 1899, *L. rauscheri* AUBERT, 1957, *L. handlirschi* KEMPNY, 1898, *L. teriolensis* KEMPNY, 1900) have paler, less easily noticed hairs but those on the abdominal segments are nevertheless about half as long as the segment length and well visible in profile.

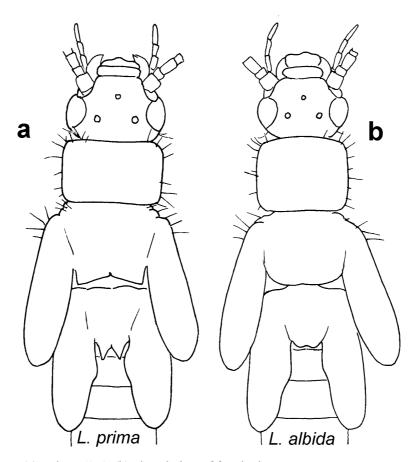


Fig.35: Leuctra prima (a) and L. albida (b), dorsal view of fore body.

Notes on central European members of the *fusca*–group: large group with many autumnal species.

- Some are normally shaped and normally pilose like *L. albida* KEMPNY, 1899 (Fig. 36b), *L. fusca* (L., 1758) and *L. digitata* KEMPNY, 1899 (Fig. 36a; the distinction of the latter two species after RAUŠER 1980 fails!);
- Many have short hairs and are slender (like L. aurita NAVÁS, 1919; Fig. 36c),
- Some are almost hairless, very slender and pale. The subterranean *L. major* BRINCK, 1949 and *L. leptogaster* AUBERT, 1949 are extremes.

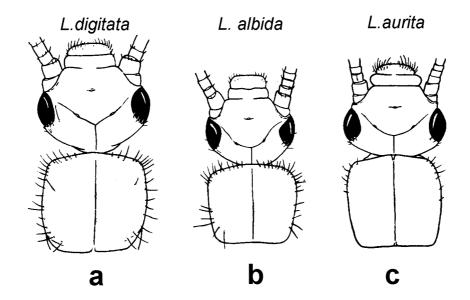


Fig. 36. Head and thorax of *Leuctra* spp. (from RAUŠER, 1980).

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