

# AMERICAN MUSEUM *Novitates*

PUBLISHED BY THE AMERICAN MUSEUM OF NATURAL HISTORY  
CENTRAL PARK WEST AT 79TH STREET, NEW YORK, NY 10024  
Number 3572, 39 pp., 14 figures, 4 tables  
May 16, 2007

## How Time Flies for Flies: Diverse Diptera from the Triassic of Virginia and Early Radiation of the Order

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### ABSTRACT

The most diverse and best-preserved early fauna of flies (order Diptera) is described from the Late Carnian (Late Triassic, ca. 220 Ma) of Virginia, USA. Complete flies are preserved as aluminosilicate films on very fine-grained shales from the Cow Branch Formation, which is part of the Newark Supergroup of Early Mesozoic rift basins from eastern North America. The dipteran fauna consists of eight families (one new), 11 genera (five new), and 16 species (11 new), and includes the following taxa (Blagoderov and Grimaldi are the authors of all new names): *Architipula youngi* Krzemiński, *Metarchilimonia krzeminskorum* n.gen., n.sp., and *M. solita* n.sp. (Limoniidae); *Triassopsychoda olseni* n.gen., n.sp. (Psychodidae); Culicomorpha indet.; *Yalea argentata* (Krzemiński), *Y. rectimedia* n.sp., *Alinka cara* Krzemiński (Procramptonomyiidae); *Veriplecia rugosa* n.sp., *Virginiptera certa* n.gen., n.sp., *V. similis* n.sp., *V. lativentra* n.sp. (Paraxymyiidae); *Brachyrhyphus distortus* n.gen. n.sp. (Protorhyphidae); ?*Crosaphis virginiensis* n.sp. (Crosaphididae); and *Prosechamyia trimedia* n.gen., n.sp., *P. dimedia* n.sp. (Prosechamyiidae, new family). Particularly significant is a culicomorphan with a long proboscis, which is the earliest fossil record of a structure specialized apparently for blood feeding. Also, *Prosechamyia* appears to be a stem group to the very diverse infraorder Brachycera, the earliest definitive members of which appear in the Early Jurassic. Phylogenetic relationships of major clades of living and extinct nematoceros Diptera are analyzed, indicating that infraordinal-level diversification was complete by the Late Triassic. Flies did not reach modern levels of ecological abundance until the mid-Jurassic, apparently due to diversification within most infraorders by that time.

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## INTRODUCTION

The Diptera, or true flies, comprise one of the oldest lineages of holometabolous insects, with the earliest definitive flies known from the mid-Triassic of France, approximately 230 Ma (Krzemiński and Krzemińska, 2003; Krzemiński et al., 1994). Since the Triassic, flies have evolved into one of the most diverse orders of living insects, with approximately 120,000 described species, and estimates place the actual numbers near one million or more (Gaston, 1991). Ecologically, flies are arguably the most diverse of all orders of insects, including species that are vertebrate ectoparasites, arthropod parasitoids, phytophages, pollenivores, predators, and, of course, saprophages (Grimaldi and Engel, 2005). In this study we make a substantial contribution to knowledge of the evolutionary history of the Diptera by describing the most diverse dipteran fauna now known from the Triassic.

Relationships among the major clades, or infraorders, of Diptera is not settled (Hennig, 1973; Oosterbroek and Courtney, 1995; Wood and Borkent, 1989), which is due in part to vague concepts for some of the infraorders, such as Psychodomorpha and Bibionomorpha. Vague infraordinal concepts themselves may be a result of morphological work that is insufficient to date or in its entirety. This is doubtful, though, since the basal relationships of all other major holometabolous orders have been well resolved using morphological characters (Grimaldi and Engel, 2005), and there is no reason to believe Diptera should be any different. Unlike the other holometabolous orders, though, basal relationships of Diptera have not been examined using DNA sequences. If indeed the morphology of Recent Diptera is insufficient to resolve basal relationships, it is possible that early fossils could possess intermediate features that better reveal homologies, such as is the case with Tanyderidae and Psychodidae (Ansorge, 1994) and Valesegyidae and Scatopsoidea (Amorim and Grimaldi, 2006). In taxa with extensive extinction of stem groups, obscure relationships among crown groups would be expected. Alternatively, relationships among crown groups can also be obscured if they diverged very close in time or simultaneously.

**EARLY FOSSIL RECORD:** The earliest and closest known relatives of flies appeared in the Permian. These include *Permotipula particia* Tillyard, 1929 and *Robinjohnia tillyardi* Martynova, 1948, from the Late Permian (Tatarian: 255 Ma) of Belmont and Warner's Bay, New South Wales, Australia (Willmann 1989a, b; Shcherbakov et al., 1995). Also related to Diptera, but more distantly, are the extinct families Liassophilidae (Triassic to mid-Jurassic of Eurasia) and the Permotanyderidae (Permian of Australia). *Permotipula* is known only from a single wing; specimens of the other groups were four-winged insects, though the hind wings of Liassophilidae were reduced in size. Deposits of insects from the Early Triassic are very rare (reviewed in Rasnitsyn and Quicke, 2002), and so there are no Diptera known from this period. In the mid- to Late Triassic, though, some 230–210 Ma, a sudden diversity of true (i.e., two-winged) Diptera occurred, representing all but one of the six living infraorders, 14 families (all but one extinct), and 28 species, not including the present study (table 1). Triassic Diptera are known from seven deposits in Asia, Australia, Europe, and North America (reviewed by Krzemiński 1992; Krzemiński and Evenhuis, 2000; Krzemiński and Krzemińska, 2003; Shcherbakov et al., 1995, fig. 4).<sup>4</sup> To date, no Diptera are known from the insect-bearing Triassic deposits in Argentina (Martins-Neto, 2003), and only one is known from the prolific Molteno Formation of South Africa (Anderson and Anderson, 1993), which is of indeterminate identity (Blagoderov and Grimaldi, unpubl.).

**THE VIRGINIA TRIASSIC:** All of the Diptera specimens based on the current study are from the quarries of the Solite Corporation in Cascade, Virginia (hereafter, the "Solite quarries") on the border of Virginia and North Carolina. These quarries lie in the Danville–

<sup>4</sup>*Triassochoeristites jinsuoguanensis* (Hong and Guo 2003) was recently described in the Mesopanorpididae (Mecoptera) but probably belongs to the dipteran family Vladipteridae. This conclusion is based on the structure of the cubital complex, with a convex CuA and associated concave *iCu* vein, a distinct kink at the base of the radius, three long ( $R_3$ – $R_5$ ) veins and a short, oblique vein  $R_2$  in the radial sector (this last vein was misinterpreted on the drawing, but is clearly seen on the published photograph). These are all apomorphies of Diptera.



TABLE 1  
(Continued)

<i>Amne triassica</i> Krzemiński et Jarzembowski, 1999	INFRAORDER CULICOMORPHA Family Chironomidae Stransham, UK		Rhaetian
Indet. sp. 1, 2	Family indet. Cow Branch Formation Solite, Virginia, USA		Carnian-Norian
<i>Yalea argentata</i> (Krzemiński) 1992	INFRAORDER BIBIONOMORPHA Family Procramp-tonomyiidae Solite, Virginia, USA		Carnian-Norian
<i>Yalea rectimedita</i> , n.sp.	Cow Branch Formation Solite, Virginia, USA		Carnian-Norian
<i>Alinka cara</i> Krzemiński, 1992	Cow Branch Formation Solite, Virginia, USA		Carnian-Norian
<i>Austrocramptonomyia minuta</i> Blagoderov, 1999	Lower Ipswich Series Mt. Crosby, Queensland, Australia		Carnian
<i>Vymrhyphus tuomikoskii</i> Blagoderov, 1995	Family Prothorhyphidae Madygen Formation Dzhailoucho, Kyrgyzstan, Central Asia		Ladinian-Carnian
<i>Vymrhyphus triassicus</i> Blagoderov, 1995	Madygen Formation Dzhailoucho, Kyrgyzstan, Central Asia		Ladinian-Carnian
<i>Vymrhyphus blagoderovi</i> Krzemiński, 2003	Grès a Voltzia Vosges, France		Anisian
<i>Brachyrhyphus distortus</i> , n.sp.	Cow Branch Formation Solite, Virginia, USA		Carnian-Norian
<i>Veriplecia handlirschi</i> Blagoderov, 1999	Family Paraxymyiidae Lower Ipswich Series Mt. Crosby, Queensland, Australia		Carnian
<i>Veriplecia rugosa</i> , n.sp.	Cow Branch Formation Solite, Virginia, USA		Carnian-Norian
<i>Virginiptera certa</i> , n.sp.	Cow Branch Formation Solite, Virginia, USA		Carnian-Norian
<i>Virginiptera similis</i> , n.sp.	Cow Branch Formation Solite, Virginia, USA		Carnian-Norian
<i>Virginiptera lativentra</i> , n.sp.	Cow Branch Formation Solite, Virginia, USA		Carnian-Norian
<i>Crosaphis anomala</i> Evans, 1971	Family Crosaphididae Lower Ipswich Series (Upper?) Mt. Crosby, Queensland, Australia		Carnian
? <i>Crosaphis virginiensis</i> , n.sp.	Cow Branch Formation Solite, Virginia, USA		Carnian-Norian
<i>Gallia aelsatica</i> Krzemiński, 2003	SUBORDER BRACHYCERA/STEM GROUPS Vosges, France		Anisian
<i>Prosechanyia trimedita</i> , n.sp.	Cow Branch Formation Solite, Virginia, USA		Carnian-Norian
<i>Prosechanyia dimedita</i> , n.sp.	Cow Branch Formation Solite, Virginia, USA		Carnian-Norian



Fig. 1. Nick Fraser standing on an excavated section of the insect layer in the Cow Branch Formation from the Solite Quarry. The layer is a fine-grained black shale that dips northeast at  $34^\circ$ .

Dan River Jurassic-Triassic rift lake basin of the Eastern U.S. (LeTourneau and Olsen, 2003). Fossil insects from this deposit were first reported by Olsen et al. (1978), then by Fraser et al. (1996) based on abundant new material. Olsen's material is housed in the Peabody Museum of Natural History at Yale University, New Haven, CT (hereafter, "YPM"), and all other specimens are housed in the Virginia Museum of Natural History, Martinsville, VA (hereafter, "VMNH"). In the quarries there are outcrops of the upper part of the Carnian-aged (ca. 235–223 Ma) Cow Branch Formation (figs. 1, 2). This formation originated in one of a series of rift basin lakes created during the formation of Pangaea. Some of these ancient lakes reveal striking 21,000-year periods in sedimentation, caused by Van Houten cycles (Olsen, 1986; Witte et al., 1991). Van Houten cycles are cyclical changes in precipitation caused by Milankovitch-type orbitally driven changes in solar insolation (Olsen, 1986), whose periodicity is affected by the 21,000-year cycles in the

precession of the equinoxes. Over 30 Van Houten cycles are preserved in the two main Solite quarries, and one cycle has proven to be particularly fossiliferous, including many insects (fig. 2).

The richest strata in the Solite quarries contain insects and the best-preserved plants and vertebrates, which also have the least bioturbation and the finest microlamination. Beard et al. (submitted) suggest the insect layer may reflect an environment in transition from ephemeral lake or bacterially matted lake margin to standing water marked by periodic carbonate precipitation. High  $\text{CO}_2$  in the lake sediment may have rendered it toxic to scavengers and other benthic organisms, either directly or by promoting the growth of toxic algae. A modern analog may be the saline, alkaline dolomite lakes of South Australia. Insects are found in very fine-grained, black shales, preserved entirely as two-dimensional, silvery films composed of aluminosilicates with essentially no relief. Preservation of the insects is of microscopic scale, including microtrichia about  $1\ \mu\text{m}$  in thickness. The insects are preserved with a significant diversity of vascular land plants, including ferns, ginkgophytes, cycads, and abundant conifers and Bennettitales, which appear to be close relatives of conifers (Fraser et al., 1996; Axsmith et al., 1997). Insect diversity includes at least 11 orders of pterygote insects: Blattodea, Coleoptera, Diptera, Mecopterida, Odonata, Orthoptera, Plecoptera, Sternorrhyncha, Auchenorrhyncha, Heteroptera, and Thysanoptera, comprising perhaps 50 species known thus far (Fraser and Grimaldi, 1999, 2003). The Cow Branch Formation is the only significant deposit of Triassic insects from North America, and contains taxa that are among the oldest known records of Thysanoptera (Grimaldi et al., 2004), and certain families in several orders (e.g., Belostomatidae [Heteroptera], Staphylinidae [Coleoptera]).

The suggestion that the Cow Branch Formation "may be younger than judged from paleobotanical data" (Shcherbakov et al., 1995: 78) has little foundation, since the stratigraphy of the formation is well documented and correlated with other Late Triassic and Early Jurassic deposits of the

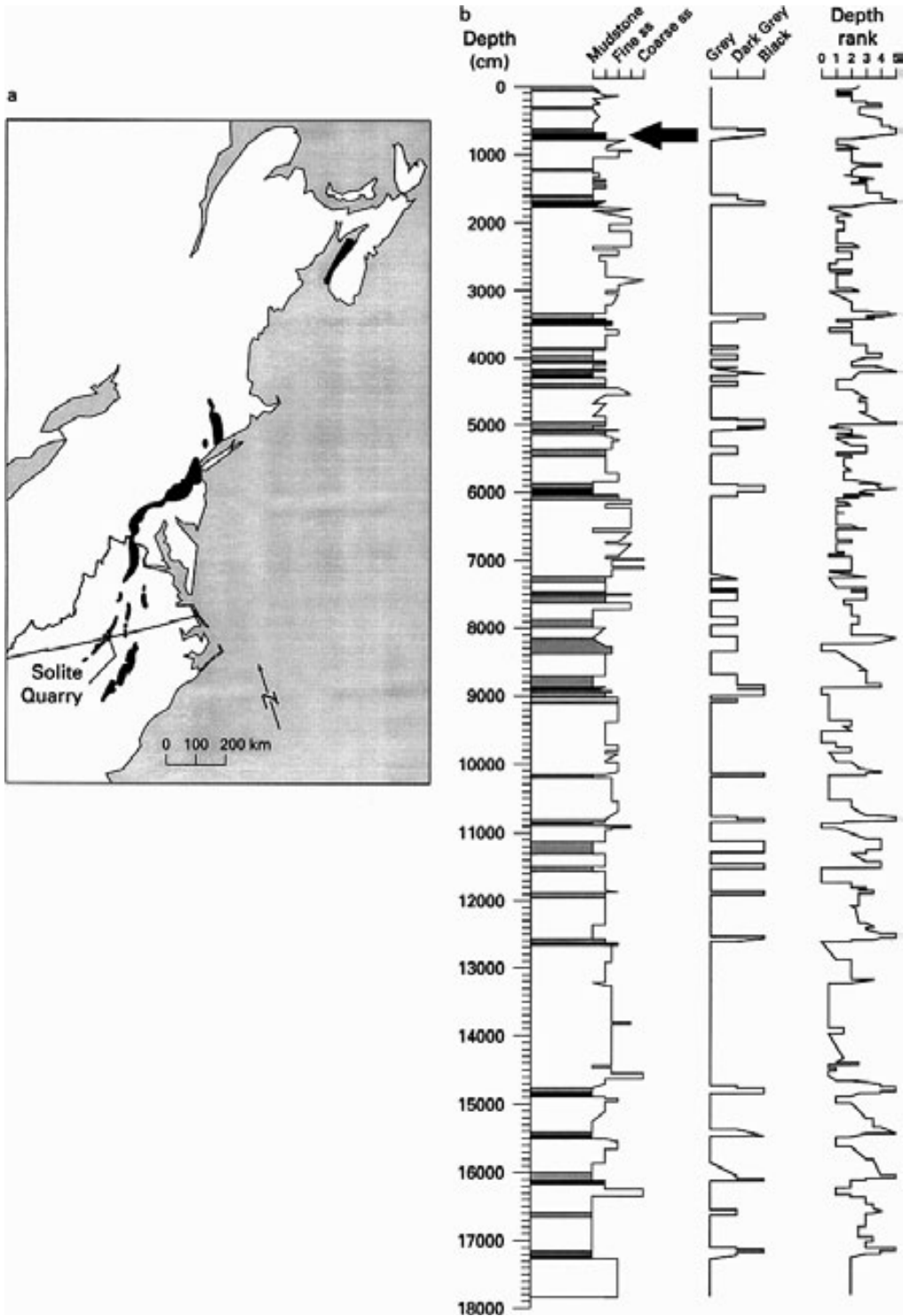


Fig. 2. Map and stratigraphy of the Triassic-Jurassic Newark Basin deposits. **a.** Map showing location of deposits, and of the Solite quarries. **b.** Stratigraphic sequence, with an arrow indicating the insect layer.

eastern North American rift valleys and with the Chinle Formation of the southwestern U.S., including the use of paleomagnetic dating (Olsen et al., 1978; Kent et al., 1995). The vertebrates in the assemblage also correlate with Carnian-aged vertebrates (Lucas and Huber, 2003), including the abundant amphibious protosaurian reptile *Tanytrachelos*. The age of the formation is Late Carnian, about 220 Ma.

## MATERIALS AND METHODS

Perhaps the most significant feature of the Virginia Triassic deposits is their preservation. Where most other Triassic flies are known only as isolated and dissociated wings, the Virginia Triassic flies are mostly completely articulated. Having features of the legs (e.g., size of coxae, presence and size of tibial spurs) and head (e.g., structure of antennae) greatly improves phylogenetic inference. Proper study of the insects from the Solite quarries, however, requires wetting of the rock surface to improve contrast and an intense but diffuse light source. Specimens were wet with an evaporative solvent, generally 70% ethanol, then carefully blotted dry after study using a soft tissue. A fiberoptic ring light is absolutely necessary to illuminate the dim silvery images; directional light from fiber optic light guides do not adequately reveal details. Photography was done by attaching a ring light to a 1000 W output fiberoptic flash unit (MicroOptics, Inc.: ML-1000), and photographing with an Infinity K-2 long distance microscope and a Nikon D-1 digital camera (see [www.microoptics-usa.com](http://www.microoptics-usa.com)). Wetted specimens were illustrated using a drawing tube attached to a stereoscope using ring-light illumination; use of polarizing film also helps.

## SYSTEMATICS

### INFRAORDER TIPULOMORPHA

Known as crane flies, the infraorder includes about 15,000 species in four living families: Tipulidae, Limoniidae, Cylindrotomidae, and Trichoceridae (the latter is sometimes placed near Psychodidae based on larval morphology). The Triassic family

Vladipteridae, which has the most generalized venation among Diptera, also belongs here. Larvae of tipulomorphs are aquatic, semi-aquatic or terrestrial, but usually associated with moist habitats, often found in decomposing plant material.

### FAMILY LIMONIIDAE RONDANI, 1856

This is one of the largest families of Diptera with more than 11,000 Recent and 300 fossil species described (Evenhuis, 1994). The majority of Mesozoic limoniids belong to the basal subfamily Architipulinae Handlirsch.

#### *Architipula youngi* Krzemiński, 1992 figures 3a, 8a

*Architipula youngi* Krzemiński, 1992: 43.

In addition to the description of Krzemiński (1992) a faint pterostigma is observed at the apex of  $R_1$  and  $R_2$ . Krzemiński indicated holotype collection number as 1103, but actually the specimen is registered under number YPM 35968. Three other paratypes are very poorly preserved. Specimen YPM 30077 can be identified as *Tipulomorpha incertae sedis*; specimen YPM 35973 (indicated as # 100000) is not conspecific with *Architipula youngi* since the fork  $R_{2+3}$  and  $R_4$  is shorter than its stem; specimen YPM 35969 lacks wings and could belong to *Culicomorpha* or *Bibionomorpha* as well as *Tipulomorpha*.

#### *Metarchilimonia* Blagoderov and Grimaldi, new genus

**DIAGNOSIS:** Small tipuloids with relatively wide wings; Sc short, approximately  $\frac{1}{2}$  the wing length.  $R_2$  meets  $R_1$ .  $R_4$  forms a stalk with  $R_5$ ,  $r-m$  connects  $M_{1+2}$  and  $R_5$ . Crossvein  $m-cu$  connected to the base of fork of  $M_{3+4}$ .

**TYPE SPECIES:** *Metarchilimonia krzeminskorum* n.gen., n.sp.

**DISCUSSION:** The new genus is close to *Archilimonia* Krzemiński and Krzemińska, 2003, by virtue of a short Sc, a similar system of vein forks,  $m-cu$  close to the base of the fork of  $M_{3+4}$ . *Mabelisia* Shcherbakov and *Gnomusca* Shcherbakov, both from the Madygen Formation in Kyrgyzstan, also have

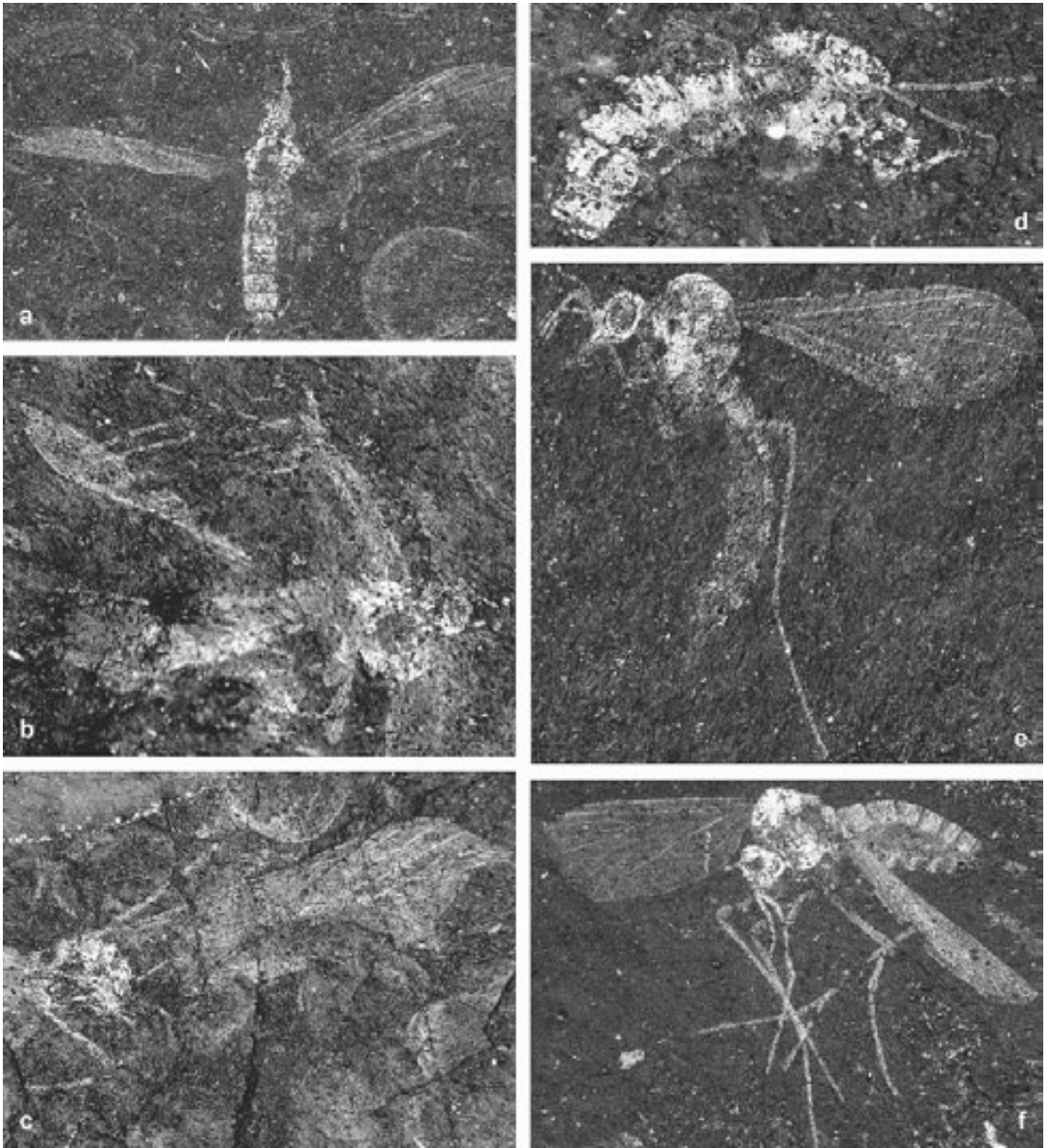


Fig. 3. Photomicrographs of tipulomorph Diptera from the Late Triassic of the Cow Branch Formation, Solite Quarry. **a.** *Architipula youngi* Krzemiński, YPM 35986. **b.** *Tipulomorpha* indet., YPM 30077. **c.** *Tipulomorpha* indet., YPM 35973. **d.** *Diptera* indet., YPM 35969. **e.** *Metarchilimonia krzeminskorum* Blagoderov and Grimaldi, holotype VMNH 732. **f.** *Metarchilimonia krzeminskorum* Blagoderov and Grimaldi, paratype VMNH 3671. For size scales see fig. 8.

vein Sc rather short and have a similar structure of RS (at least in *G. renixa*), but they are significantly different in many other aspects of venation. *Metarchilimonia* differs from *Architipula youngi* Krzemiński, also from

the Solite quarries, by the former having a wider wing, more proximal position of  $R_2$ , more distal position of *r-m* with respect to discal cell and the forking pattern of the radial veins.

ETYMOLOGY: From the Greek word  $\mu\epsilon\tau$ , meaning "after" or "next to", and *Archilimonia*. The name is feminine.

*Metarchilimonia krzeminskorum*

Blagoderov and Grimaldi, new species  
figures 3e, f, 8c, d

DIAGNOSIS:  $Sc_2$  preapical, discal cell shorter than fork of  $M_{1+2}$ .

DESCRIPTION: Female. Measurements: body length 4.7 mm, wing length 3.2 mm. Head round, antennae with 14 flagellomeres, gradually tapered in width toward apex. Scape short, smaller than basal flagellomere. Mouthparts forming rostrum, approximately  $1.3\times$  greatest length of eye; palpi short (segmentation not preserved). Scutum arched. Wing moderately wide, width  $0.39\times$  the length. Sc short,  $0.53\times$  the wing length, ends slightly proximal to level of base of  $R_{2+3}$  stem.  $Sc_2$  present, preapical. Base of stem of RS at  $0.37\times$  the wing length. Five branches of RS, four reaching wing margin.  $R_2$  meeting  $R_1$  near wing margin. Stem of  $R_{2+3}$  parallel to  $R_1$ , slightly bent at base.  $R_{4+5}$  very short,  $0.5\times$  the length of section of  $R_5$  anterior to *r-m*. Base of  $R_5$  transverse. Length of  $R_5$  nearly  $2\times$  longer than stem of RS. Basal fork of M at level of Sc apex.  $M_3$  connected to  $M_{1+2}$  near the middle of stem of latter. Anterior margin of discal cell  $1.3\times$  longer than stem of  $M_{1+2}$  and  $0.8\times$  the length of  $M_1$ . Posterior margin of discal cell  $0.7\times$  length of  $M_4$ .  $M_4$   $1.6\times$  longer than stem of  $M_{3+4}$ . Crossvein *m-cu* in line with base of  $M_{3+4}$  fork. CuP and A veins not observed. Tibial spurs absent, legs with short setae. Oviscape as long as tergite 10, sclerotized, tapered at apex; cerci and valves not apparent.

MATERIAL: Holotype VMNH 732, sex unknown; paratype VMNH 3671, ♀.

ETYMOLOGY: The specific epithet is a patronym honoring Drs. Wiesław Krzemiński and Ewa Krzemińska, for their significant contributions to the study of early Diptera.

*Metarchilimonia solita*

Blagoderov and Grimaldi, new species  
figures 4a, 8b

DIAGNOSIS: Differs from *M. krzeminskorum* by being nearly half the size, and by

various venation features: stem of  $M_{1+2}$  longer than the fork of  $M_{1+2}$  (vs. longer fork); apical  $Sc_2$ ; discal cell longer.

DESCRIPTION: Measurements: body length 2.3–2.5 mm (holotype, 2.5), wing length 2–2.2 mm (holotype, 2.2). Head round, rostrum not observed. Antenna moniliform, 14-segmented, flagellomeres very slightly tapered toward apex, each with single apical seta. Only basal 8 flagellomeres well preserved. Legs densely setose, without tibial spurs. Sc short,  $0.46\times$  the wing length.  $Sc_2$  apical, its length about equal to short, distal segment of  $Sc_1$ . Base of  $R_5$  oblique. Anterior margin of discal cell  $1.2\times$  longer than stem of  $M_{1+2}$  fork and  $1.2\times$  longer than  $M_1$ . Posterior margin of discal cell  $1.2\times$  longer than  $M_4$ .

MATERIAL: Holotype VMNH 3013.

ETYMOLOGY: The specific epithet is for the Solite Corporation, which has quarried the locality.

INFRAORDER PSYCHODOMORPHA

This infraorder includes the living families Tanyderidae and Psychodidae, and sometimes Ptychopteridae, the Blephariceromorpha, and Scatopsoidea. The first two families are known from the Early Jurassic (Ansorge, 1994). Psychodidae are widely distributed and their larvae breed in wet to foul substrates. Phlebotominae feed on vertebrate blood and insect hemolymph. In the widest sense this infraorder may include the following extinct taxa: Hennigmatidae, Tillyardipteridae, Rhaetaniidae, and Eoptychopteridae.

FAMILY PSYCHODIDAE NEWMAN, 1834,  
OR NEAR

*Triassopsychoda* Blagoderov and Grimaldi,  
new genus

DIAGNOSIS: Wing drop-shaped, with highly reduced anal lobe; veins with long setae. Vein Sc meets C, ends slightly distad of basalmost fork of R. Vein R with four branches, M with three branches, *m-m* absent; bases of  $R_{4+5}$ ,  $R_5$ , *r-m* in apical half of wing; base of M faint.

TYPE SPECIES: *Triassopsychoda olseni* n.sp.

ETYMOLOGY: The genus name is from the Triassic; the name is feminine.

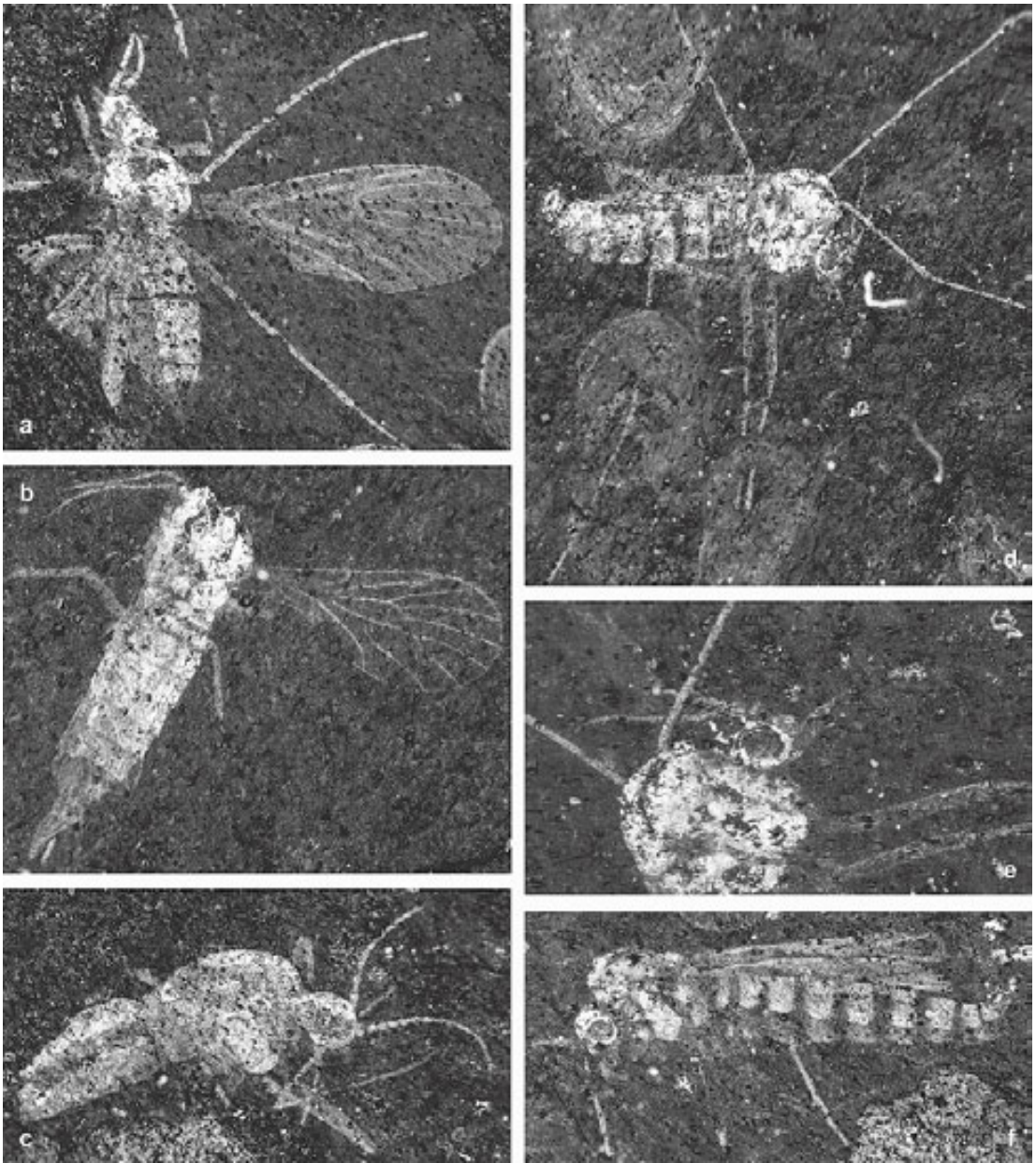


Fig. 4. Photomicrographs of tipulomorphan (a), psychodomorphan (b), and culicomorphan (c-f) Diptera from the Cow Branch Formation. **a.** *Metarchilimonia solita* Blagoderov and Grimaldi, holotype VMNH 3013. **b.** *Triassopsychoda olseni* Blagoderov and Grimaldi, holotype VMNH 733. **c.** Culicomorpha indet., VMNH 808. **d.** Culicomorpha indet., VMNH 951. **e.** Culicomorpha indet., VMNH 951, detail of head, showing proboscis. **f.** Culicomorpha indet., VMNH 932. For size scales see fig. 9.

*Triassopsychoda olseni*

Blagoderov and Grimaldi, new species  
figures 4b, 9a

DIAGNOSIS: As for genus.

DESCRIPTION: Female. Measurements: body length = 2.1 mm, wing length = 1.4 mm. Head round, eyes reniform, with large facets. Length of mouthparts, including palps, approximately  $\frac{1}{4}$  height of head. Antennae with 14 flagellomeres, each with whorl of long setae; length of each flagellomere approximately  $2\times$  its width. Apex of wing broadly rounded, anal lobe absent, wing base petiolate. Wing veins with long setae including C. Costal cell very wide; Sc meets C, its length approximately  $0.4\times$  the wing length. Base of RS stem at  $0.35\times$  wing length, base of  $R_{3+4+5}$  fork at  $0.53\times$  the wing length, so radial-medial cell is long and wide.  $R_2$  lost, base of  $R_{4+5}$  fork distal to base of  $R_3$ . Lengths of stem of  $R_{4+5}$  fork, basal section of  $R_5$ , *r-m*, and stem of  $M_{1+2}$  fork are subequal. Base of  $M_{3+4}$  is at  $0.47\times$  the wing length. Veins  $M_3$  and *im* absent; discal cell open and very wide. Nine segments of abdomen apparent, subequal in length. Cerci/oviscapt long, narrow, triangular, length equal that of segments 8 and 9 combined.

MATERIAL: Holotype VMNH 733.

ETYMOLOGY: The species epithet is in honour of Dr. Paul Olsen of Lamont-Doherty Earth Observatory, who has done the original work on the Solite quarry fossils, and who has brought tremendous insight to the Mesozoic rift basins of eastern North America.

DISCUSSION: Despite the reduced venation (absence of  $R_2$ ,  $M_3$ , base of M) this fly possesses some very plesiomorphic features: the distal positions of the base of RS, and radial and medial forks, and crossveins *r-m* and *m-cu*, and a very wide discal cell. The structure of the radial veins of the new genus resembles the living genus *Horaiella* Tonnoir from India and especially *Eatonisca* Meunier from Baltic amber (both Psychodidae), but differs from those genera by the distal position of the radial and medial forks and the absence of  $M_3$ . It differs from the Early Jurassic genera *Tanypsycha* Ansoerge, *Liassopsychodina* Ansoerge, and *Libanophlebotomus* Azar et al.,

1999, by the absence of veins  $R_2$  and  $M_3$ , from the latter genus also by having Sc merging with C instead of  $R_1$ .

The position of *Triassopsychoda* is uncertain. The reniform eye shape, setal whorls on the flagellomeres, long cerci/oviscapt, broad wing with narrow stem, reduced anal lobe, and setose wing veins are, in combination, diagnostic of Psychodidae. The striking feature of *Triassopsychoda*, distinguishing it from other Psychodidae is a complete loss of vein  $M_3$  while preserving crossvein *tb* (the basal-most section of  $M_4$ ). Many authors, following McAlpine (1981) interpret the posterior longitudinal vein in the hind medial fork as  $CuA_1$  (Azar et al., 1999; Duckhouse, 2000). As we discuss below (see Phylogeny),  $CuA$  in Diptera is always a nonbranched convex vein. In Psychodidae, however, vein relief is unapparent, veins tend to be equally sclerotized and the membrane weakly corrugated; longitudinal veins are long and parallel; and crossveins and fork bases are shifted proximad. *Triassopsychoda*, conversely, has  $CuA$  long and thick, just like in other Diptera, and the base of  $M_4$  is at the middle of the wing. Another objection against including *Triassopsychoda* in Psychodidae is the structure of vein C, which is evanescent slightly apical to the apex of  $R_5$  (vs. circumambient in Psychodidae, although this state may be plesiomorphic because some Tipulomorpha share it). Thus, despite the highly reduced venation, *Triassopsychoda* possesses several plesiomorphies not found in living Psychodidae.

## INFRAORDER CULICOMORPHA

Culicomorpha include more than 12,000 Recent species in eight families: Ceratopogonidae, Chaoboridae, Chironomidae, Corethrellidae, Culicidae, Dixidae, Simuliidae, and Thaumaleidae. The Culicidae, Corethrellidae, Simuliidae and basal Ceratopogonidae are essentially hematophagous. Larvae of almost all culicomorphs are aquatic or live in moist habitats. About 300 fossil species are described, about 60% of which are Ceratopogonidae. The earliest known fossil of the infraorder is *Aenne* from the uppermost Triassic of England, which has been assigned

to the Chironomidae (Krzemiński and Jarzembowski, 1999).

*Culicomorpha indet.*  
figures 4c–f, 5a, b, 9b, c

Several specimens from the Solite quarries have their venation obscured, so they can not be described and named formally. Some of these specimens, however, are otherwise exquisitely preserved, retaining some distinctive features that allow placing them within the infraorder Culicomorpha. Despite differences in sizes and leg lengths, specimens VMNH 808, 873, 932, 951, 2956, and 3056 possess: (1) slender moniliform antennae with long setae on each flagellomere; (2) a round antennal pedicel that is larger than the scape; (3) round (vs. reniform or emarginate) eyes with large facets; (4) a broad katapisternum that is rounded ventrally; (5) a short, arched mesonotum that nearly overhangs the pronotum; (6) small, unprotruding scutellum; (7) wide insertion of the abdomen; and (8) male genitalia curved dorsally. Features 2, 5, and 8 are particularly distinctive to Culicomorpha, but venation is needed to confirm this. Most significantly, VMNH 951 bears a long, slender proboscis that is nearly twice as long as the head height (figs. 4d, e; 9b, c).

Body and leg proportions of VMNH 951 are very similar to that of other culicomorphans (VMNH 808, 873, 932, 2956, 3056), but it is doubtful that the specimens are conspecific, since there are no known instances in any living species of Diptera where a proboscis is so dramatically dimorphic. In various Recent species of biting midges the styletlike mandibles of females are well developed and used to puncture the skin of their host in order to feed on blood; males feed on nectar and have vestigial mandibles, but the proboscis size barely differs with that of females. Since angiosperms did not appear until nearly 100 Ma later, it is possible that the proboscis was used to probe Bennettitalean or Gnetalean reproductive structures, and bennettitaleans occurred in the Triassic of Virginia. Recent Gnetales produce pollination droplets, on which insects feed, but these are external, making a long proboscis unnecessary. VMNH 951 is the earliest fossil evidence

of a structure in Diptera specialized for blood feeding, in a group in which the ground-plan adult diet is hematophagy (Grimaldi and Engel, 2005; Lukashevich and Mostovski, 2003). If this is the case, VMNH 951 is the earliest known blood-feeding insect. It is not until the Jurassic that definitive insect ectoparasites appeared, and not until the Cretaceous that definitive hematophagous insect groups appeared (Grimaldi and Engel, 2005).

INFRAORDER BIBIONOMORPHA

The composition of the infraorder is in flux, partly because it is paraphyletic with respect to the higher flies, the Brachycera. It includes diverse Mesozoic taxa, such as Eliidae and various families of sciaroids, in addition to the families discussed below. The infraorder also includes very diverse Recent families, such as Mycetophilidae and Cecidomyiidae, totaling 8,000 species in some 25 families. In contrast to the other major infraorders of nematoceros flies, larvae of Bibionomorpha and those of their stem group (the Brachycera) are almost exclusively terrestrial, being saprophagous, fungivorous, phytophagous, and sometimes predatory.

FAMILY PROCRAMPTONOMYIIDAE  
KOVALEV, 1985

A small family of basal bibionomorphs known from Late Triassic to Early Cretaceous of North America, Australia, Asia, and Europe. Kovalev (in Kalugina and Kovalev, 1995) referred the family to Anisopodoidea based on the presence of a discal cell. However, a more accurate placement would be at the base of all Bibionomorpha s.l. (Shcherbakov et al., 1995).

Genus *Yalea* Krzemiński, 2004

*Yalea* Krzemiński, 1992: 44. Type species: *Yalea argentata* Krzemiński, 1992, monotypic, by original designation (preoccupied by *Yalea* Chu, 1979 [Geometridae]).  
*Yalea* Krzemiński, 2004: 229. Replacement name.

DIAGNOSIS: R<sub>4</sub> base proximal to *r-m*, section of M<sub>2</sub> proximal to *m-m* and section of M<sub>3</sub> proximal to *m-m* are nearly equal, with length 0.3–0.5× that of discal cell.

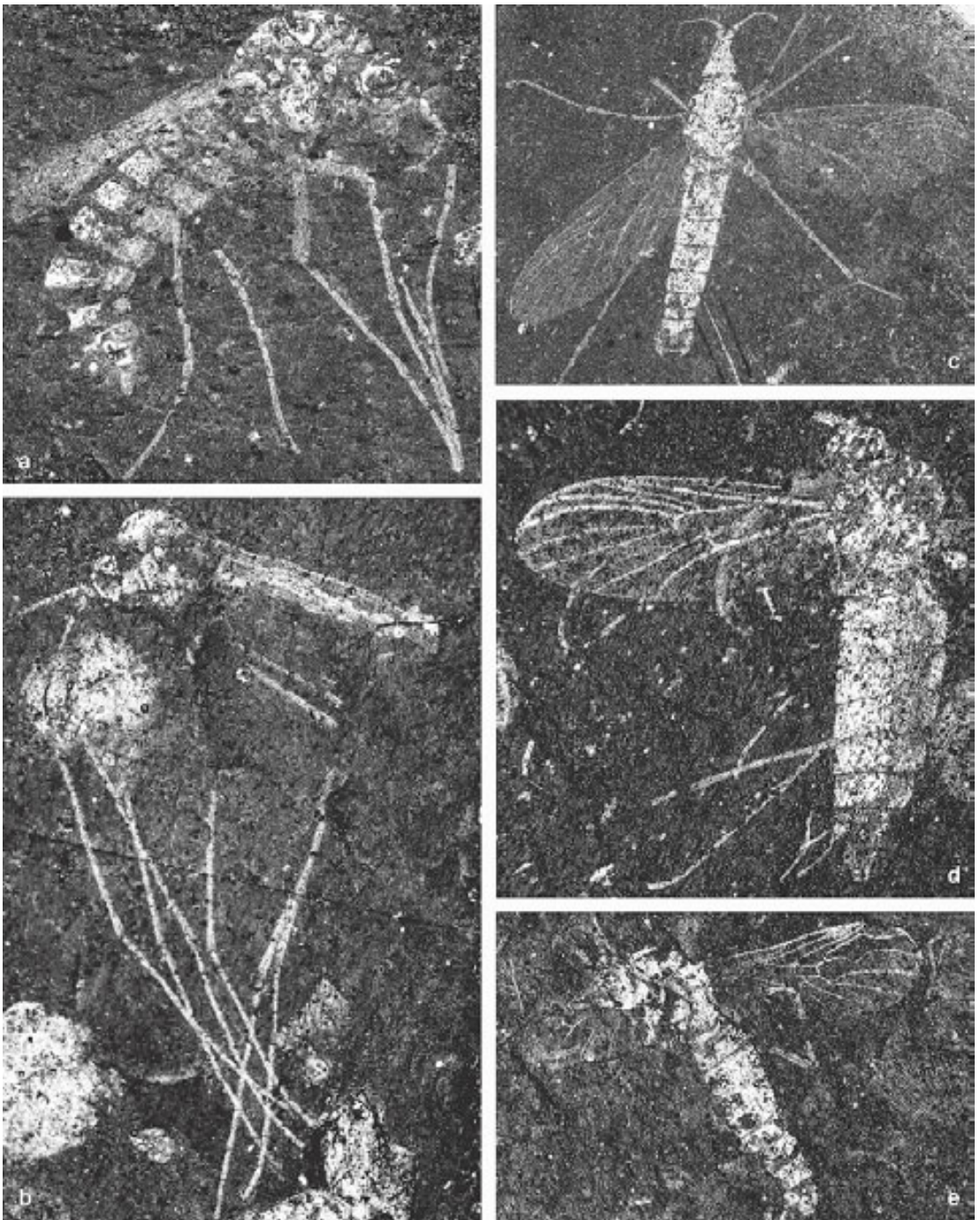


Fig. 5. Photomicrographs of culicomorphan (a, b) and bibionomorph (c–e) Diptera from the Cow Branch Formation. **a.** *Culicomorpha* indet., VMNH 2956. **b.** *Culicomorpha* indet., VMNH 3056. **c.** *Yalea argentata* (Krzemiński). **d.** *Yalea rectimedia* Blagoderov and Grimaldi, VMNH 1041. **e.** *Veriplecia rugosa* Blagoderov and Grimaldi, YPM 35989. For size scales see fig. 10.

INCLUDED SPECIES: *Yalea argentata* (Krzemiński) (figs. 5c, 10b) and *Y. rectimedia* Blagoderov and Grimaldi, new species.

*Yalea rectimedia* Blagoderov and Grimaldi,  
new species  
figures 5d, 10a

DIAGNOSIS: Nearly 0.7× smaller than *Yalea argentata*. Sc shorter, ends at the level of  $R_{2+3}$  base (vs. slightly distad).  $M_3$  and  $M_4$  parallel (vs. slightly divergent and curved).  $R_{2+3}$  2× the length of stem of RS (vs. 1.5× the length). Length of  $M_{1+2}$  stem equal to section of  $M_2$  proximal to *m-m* (vs. 1.5× the length).

DESCRIPTION: Female. Measurements: body = 3.8 mm, wing = 2.2 mm. Scape and pedicel wider than flagellum, hemispheric; flagellomeres cylindrical, with lengths about equal to widths. Sc ends at C at level of base of  $R_{2+3}$ .  $R_1$  almost straight,  $R_{2+3}$  sigmoid, curved anteriorly at apex.  $R_4$  and  $R_5$  curved posteriorly.  $R_{2+3}$  2× the length of the basal section of RS. Length of basal section of RS 2.5× that of section of RS between bases of  $R_{2+3}$  and *r-m*. M stem weak. Section of  $M_{1+2}$  stem proximal to *r-m* 2.3× the length of stem section distal to *r-m*; length of  $M_1$  and  $M_2$  fork 3.5× that of  $M_{1+2}$  stem. Apex of CuA distal to level of *r-m* crossvein. Cerci apparently one-segmented.

MATERIAL: Holotype VMNH 1041.

ETYMOLOGY: The species epithet is from the Latin *rectus*, meaning “straight”, in reference to details of wing venation.

#### Genus *Alinka* Krzemiński, 1992

*Alinka* Krzemiński, 1992: 46. Type species: *Alinka cara* Krzemiński, 1992, monotypic, by original designation.

Krzemiński (1992: fig. 16) described *Alinka cara* as the earliest representative of Brachycera known at the time. This opinion was later critiqued by Shcherbakov et al. (1995), Grimaldi and Cumming (1999), and others. Our re-examination of the holotype (YPM 16826 [not 100 000 as originally cited]) supports the placement of *Alinka* in the family Procrampptomomyiidae. Eleven flagellomeres are preserved in the antenna, although the actual number could be greater because the apical flagellomere is incomplete and without

the apical tuft of setulae originally reported (Krzemiński, 1992: fig. 6a). The only synapomorphy of *Alinka* with Brachycera was a short  $R_4$ , but this character state appears several times in Diptera, such as in Paraxemyiidae, Eoditomyiidae, and Protorhyphidae (see below). The paratype of *A. cara* is too poorly preserved to be identified as a conspecific with certainty.

SUPERFAMILY SCIAROIDEA BILLBERG, 1820  
FAMILY PARAXEMYIIDAE ROHDENDORF, 1946

This family includes basal sciaroids (“fungus gnats”) with three branches of RS and a distinct, though sometimes faint base of M. We are including in this family the Eomycetophilidae Ansoerge, 1996, which is apomorphic in having  $R_4$  very short, apical, and transverse. We consider this family to be a subfamily of Paraxemyiidae. Four genera of Paraxemyiidae are known from the Late Triassic to the Early Cretaceous: *Paraxemyia* Rohdendorf, 1946; *Veriplecia* Blagoderov, 1999; *Eomycetophila* Kovalev, 1990; and *Complecia* Blagoderov, 1999.

#### Genus *Veriplecia* Blagoderov, 1999

*Veriplecia* Blagoderov, 1999: 12. Type species: *V. clara* Blagoderov, 1999, monotypic.

*Veriplecia rugosa* Blagoderov and Grimaldi,  
new species  
figures 5e, 10d

DIAGNOSIS: Differs from *Veriplecia clara* from the Late Triassic of Australia by the more basal position of the base of  $R_{2+3}$ , which is slightly basal to the level of crossvein *tb* (basal section of  $M_{3+4}$  stem proximal to crossvein *m-cu*).

DESCRIPTION: Male. Measurements: body = 3.1 mm, wing = 2.2 mm. Head round, eyes large, genae setulose. Antennae moniliform, with scape slightly larger than individual flagellomeres. Scutum with fine setulae; no large, bristlelike setae. Costa setulose. Sc short, 0.41× the length of wing, ends at C slightly proximal to level of base of  $R_{2+3}$ .  $Sc_2$  not apparent or absent. Ratios of length of stem of RS: stem of  $R_{4+5}$ :  $R_5$  are 1:1:3.

Crossvein *r-m* located between bases of  $R_{2+3}$  and  $R_4$ ; length of section of RS between *r-m* and base of  $R_4$  is  $0.5\times$  the length of section between bases of  $R_{2+3}$  and *r-m*. Distinct pterostigma at apex of  $R_1$ . Ratios of lengths of M stem, and sections of  $M_{1+2}$  proximal and distal of *r-m* when straightened out are 5:1:1. Basal stem of M evanescent toward wing base. Length of  $M_1$  and  $M_2$  fork  $5.7\times$  the length of stem of  $M_{1+2}$  distal to *r-m*. Length of  $M_{1+2}$  stem proximal to *r-m* is  $2.6\times$  the *tb* length. Crossvein *m-cu* is  $1.7\times$  the length of *tb*. CuA very slightly curved back. 10 visible segments of abdomen; segment 3 largest, 8 smallest, 10 apparently large, covering ventrally situated genitalia. Abdomen and genitalia setose. Pair of short, protruding appendages present (probably gonostyli).

**MATERIAL:** Holotype YPM 35989.

**ETYMOLOGY:** The specific epithet is from the Latin *rugosus*, meaning "wrinkled", in reference to the deformed wing of the unique specimen.

*Virginiptera* Blagoderov and Grimaldi,  
new genus

**DIAGNOSIS:** Costa ends slightly beyond tip of  $R_5$ . Sc ends at C distal to the base of RS and at the level of base of fork  $R_{2+3}$ – $R_{4+5}$ . R and CuA considerably thicker than M. Base of  $R_{2+3}$  proximal to crossvein *r-m*.  $R_4$  very short,  $0.5\times$  the length of RS. Stem of M weak, incomplete. Crossvein *tb* very short. CuA evenly curved. Tibial spurs (1:1:2) present, length ca.  $1.3\times$  width of tibia. Female cerci 2-segmented. Male genitalia held ventrally.

The new genus is close to *Complecia clara* Blagoderov, 1999, from the Late Triassic of Australia, which differs from the new genus in having Sc shorter and ending free, base of  $R_{2+3}$  very close to the base of *r-m*, and a more regular spacing of veins in the radial sector (ratios of the distances along C between tips of  $R_1$ ,  $R_{2+3}$ , and  $R_5$  are 1.3:1.1:1.0) and in the medial sector (ratios of the distances along wing margin between apices of  $R_5$ ,  $M_1$ ,  $M_2$ ,  $M_{3+4}$ , and CuA are 1:1.3:1.6:1.9).

**TYPE SPECIES:** *Virginiptera certa*, new species.

**ETYMOLOGY:** The genus name derives from Virginia, the U.S. state of origin of the fossil. The genus name is feminine.

**COMMENTS:** The new genus has a venation similar to that of Paraxymiidae (Eomycetophilinae) and Eoditomyiidae: three branches in RS, with  $R_4$  extremely short, the base of  $R_{2+3}$  proximal to *r-m*, three branches of M, and the discal cell reduced. *Virginiptera* differs from *Eoditomyia* Ansoerge (Early Jurassic, Germany) by having a complete Sc, and in *Eoditomyia* crossveins *r-m*, *tb* and *m-cu* are nearly in line, and the base of M is completely reduced. It differs from *Crosaphis* (Anisopodoidea) by the presence of a short  $R_4$  vein.

The new genus is closely related to or within Sciaroidea based on the following features:

- Length of coxae approximately equal to thoracic height. Anisopodoidea in general have much shorter coxae (length ca.  $0.3\times$  the depth of thorax) than in Sciaroidea, the latter of which have a coxal length that may even exceed the thoracic height.
- Tibial spurs as long as the tibial diameter, and protruding at approximately  $45^\circ$ . Sciaroidea have short to extremely long tibial spurs, but the position is always protruding away from the tibia. In contrast, Anisopodoidea (including fossil Anisopodidae and Protorhynchidae) have short tibial spurs in line with the tibia.
- CuA is evenly curved. In most Anisopodoidea the section of CuA distal to *m-cu* is apomorphically straight or sinuous.
- Abdominal tergite I lacks laterodorsal tubercles or protrusions. Sciaroidea never possess these structures.
- Anepisternum is bare. In those fossil Sciaroidea for which preservation allows observation, as well as in many basal Recent sciaroids, the pleural sclerites are bare. Anisopodoidea very often have the setose pleural sclerites.
- Segment 8 is short in females. Size of the eighth segment varies in Sciaroidea and Anisopodoidea, but it is generally reduced in the former and well developed in the latter group.
- Tibiae with rows of thick dorsal setae. Anisopodoidea usually lack thick setae on the tibiae.
- MA or arculus not seen at the wing base. Anisopodoidea have a strong veinlike arculus; in Sciaroidea MA is usually

represented only by a kink in the wing membrane at the base of R.

*Virginiptera certa* Blagoderov and Grimaldi,  
new species  
figures 6b, 11c

DIAGNOSIS: Sc ends at C, very slightly proximal to base of  $R_{2+3}$  and distal to crossvein *tb*. Crossvein *r-m* as long as section of RS between bases of  $R_{2+3}$  and *r-m*. Stem of M slightly kinked at *r-m* and *tb*. Length of section of  $M_{1+2}$  proximal to *r-m*  $1.6\times$  that of *r-m*. Section of  $M_{1+2}$  stem distal to *r-m* with length  $1.8\times$  that of *r-m*.

DESCRIPTION: Measurements: body length = 2.8–3.1 mm (holotype 3.1 mm), wing length = 1.9–2 mm (holotype 1.9 mm), antennal length = 0.6 mm. Antennae short, approximately  $0.7\times$  thorax length; flagellum 14-segmented (13-segmented in VMNH 2998); flagellomeres cylindrical, length of each ca.  $0.5\times$  the width. Scutum with short irregular setulae, no bristlelike setae, several longer setae caudally. Scutellum small, not protruding. Katepisternum wider than deep. Abdomen setulose, attachment slightly narrowed; widest at segments 3–4. Length of segment 8  $0.5\times$  that of segment 7. Cerci 2-segmented. **Wing:** Costa ends slightly beyond tip of  $R_5$ ,  $\frac{1}{2}$  the length between tips of  $R_5$  and  $M_1$  on holotype's left wing. Sc ends at C, slightly proximal to base of  $R_{2+3}$  and distal to crossvein *tb*. Sc length  $0.4\times$  the wing length. R and CuA veins distinctly thicker and more sclerotized than medial veins.  $R_1$  length  $0.67\times$  the wing length.  $R_{2+3}$  nearly parallel to  $R_1$ , both veins very slightly curved toward anterior margin of wing.  $R_4$  very short, straight,  $0.5\times$  the length of  $R_5$ . Ratios of distances between tips of  $R_1$ ,  $R_{2+3}$ , and  $R_5$  on C are 2:2.5:1.  $R_{4+5}$  stem straight basally, in apical quarter curved slightly backward, ending at wing tip. Crossvein *r-m* as long as section of RS between bases of  $R_{2+3}$  and *r-m*. Stem of M slightly kinked at *r-m* and *tb*;  $M_{1+2}$  stem proximal to *r-m* is very weak. Section of  $M_{1+2}$  stem distal to *r-m*  $1.8\times$  that of section proximal to *r-m*. Length of section of  $M_{1+2}$  proximal to *r-m*  $1.6\times$  that of *r-m*. Length of  $M_1$ – $M_2$  fork  $4.2\times$  the length of section of  $M_{1+2}$  distal to *r-m*. Ratios of distances at wing margin between apices of  $R_5$ ,  $M_1$ ,  $M_2$ ,  $M_{3+4}$ ,

and CuA are 1:1.5:2.5:2.5. Crossvein *tb* very short, but distinct. CuA evenly curved. Legs setulose, hind tibiae with distinct row of dorsal setae. Tibial spurs  $1.5\times$  the tibial diameter, protruding  $45^\circ$  from axis of tibia.

MATERIAL: Holotype VMNH 731, ♀; paratype VMNH 2998, ♂.

ETYMOLOGY: The species epithet is from the Latin *certus*, meaning "certain".

*Virginiptera similis* Blagoderov and Grimaldi,  
new species  
figures 6d, 11b

DIAGNOSIS: Size slightly less than half that of *V. certa*. Sc ends at C, at the level of crossvein *tb* and significantly proximal to level of *r-m*. Crossvein *r-m*  $2\times$  longer than section of RS between bases of  $R_{2+3}$  and *r-m*. Length of section of  $M_{1+2}$  proximal to *r-m* is  $1.3\times$  that of *r-m*. Length of section of  $M_{1+2}$  distal to *r-m* is  $2\times$  that of  $M_{1+2}$  section proximal to *r-m*.

DESCRIPTION: Measurements: body = 1.3 mm, wing = 1.3 mm, antenna = 0.7 mm. Flagellum 14-segmented, flagellomeres cylindrical. Scutum arched. Abdomen with male terminalia large and folded under apex of abdomen. **Wing:** Costa ends slightly beyond tip of  $R_5$ . Sc ends at C, at level of crossvein *tb*. Length of Sc  $0.42\times$  the wing length.  $Sc_2$  preserved. Length of  $R_1$   $0.6\times$  the wing length.  $R_{2+3}$  nearly straight, curved at apex toward anterior margin of wing. Ratios of distances between tips of  $R_1$ ,  $R_{2+3}$ , and  $R_5$  on C are 1.5:2.5:1.  $R_5$  short, straight;  $R_4$  slightly curved,  $0.5\times$  length of  $R_5$ . Crossvein *r-m*  $2\times$  longer than section of RS between bases of  $R_{2+3}$  and *r-m*. Stem of M nearly straight, base evanescent. Length of  $M_{1+2}$  section proximal to *r-m*  $1.3\times$  that of *r-m*. Length of section of  $M_{1+2}$  distal to *r-m* is  $2\times$  that of section of  $M_{1+2}$  proximal to *r-m*. Length of  $M_1$  and  $M_2$  fork is  $4\times$  the length of  $M_{1+2}$  section distal to *r-m*. Crossvein *tb* very short, but distinct. Ratios of distances between apices of  $R_5$ ,  $M_1$ ,  $M_2$ ,  $M_{3+4}$ , and CuA at wing margin are 1:1.5:2:2. CuA evenly curved.

MATERIAL: Holotype VMNH 825, ♂; paratype VMNH 1042, ♂.

ETYMOLOGY: The species epithet is from Latin *similis*, meaning "similar to" or "resembling".

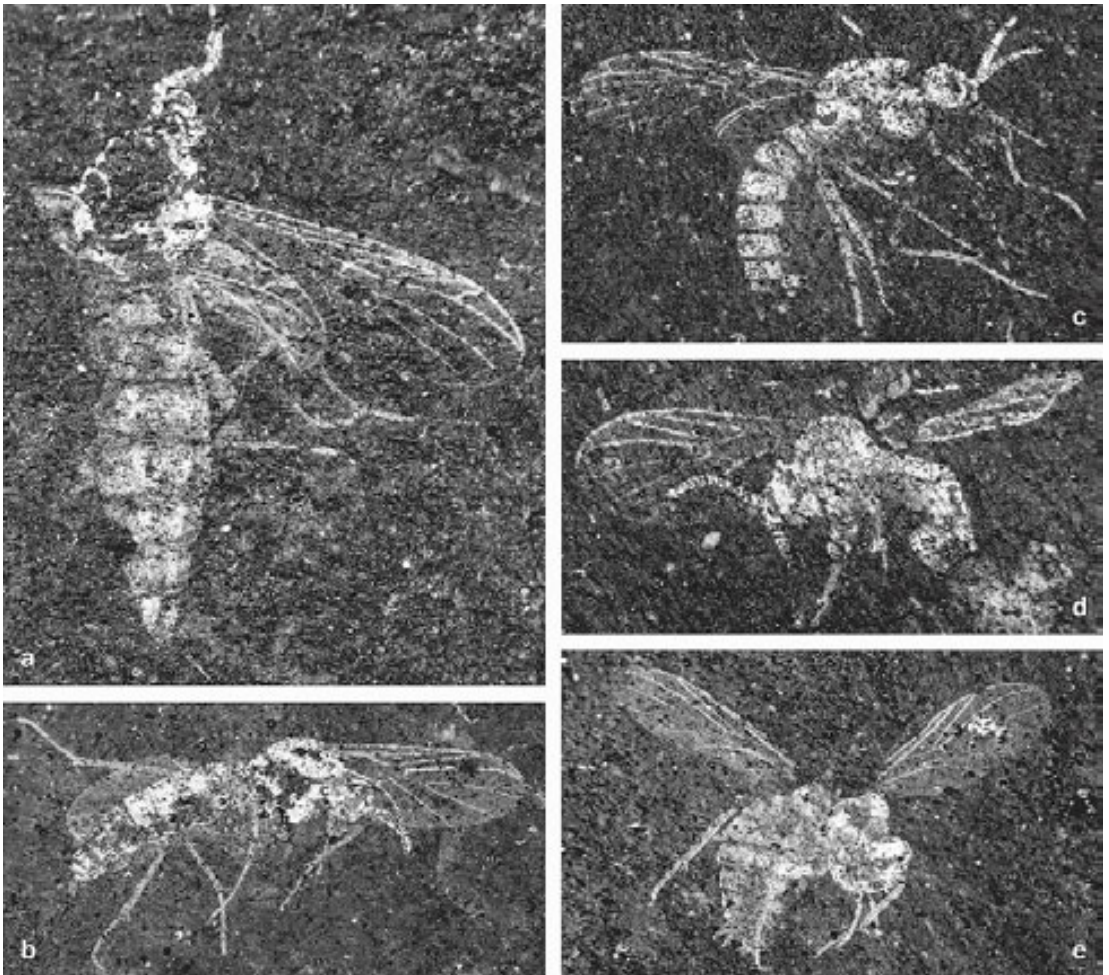


Fig. 6. Photomicrographs of bibionomorphan Diptera from the Cow Branch Formation. **a.** *Alinka cara* Krzemiński, YPM. **b.** *Virginiptera certa* Blagoderov and Grimaldi, holotype VMNH 731. **c.** *Virginiptera certa* Blagoderov and Grimaldi, paratype VMNH 2998. **d.** *Virginiptera similis* Blagoderov and Grimaldi, VMNH 825. **e.** *Virginiptera lativentra* Blagoderov and Grimaldi, holotype VMNH 2923. For size scales see fig. 11.

***Virginiptera lativentra***

Blagoderov and Grimaldi, new species  
figures 6e; 11d

DIAGNOSIS: Sc very short, ends at C slightly proximal to base of *m-cu*. C ends well beyond tip of  $R_5$ , almost to tip of  $M_1$ . Crossvein *r-m* as long as section of RS between bases of  $R_{2+3}$  and *r-m*. Length of section of  $M_{1+2}$  proximal to *r-m*  $1.4\times$  that of *r-m*. Length of section of  $M_{1+2}$  distal to *r-m*

$1.8\times$  the length of section proximal to *r-m*;  $M_{3+4}$  connects directly to stem of M.

DESCRIPTION: Measurements: body = 2.2 mm, wing = 1.6 mm, antennae = 0.8 mm. Palpi 5-segmented, long, length nearly equal to length of head. Flagellum 14-segmented, flagellomeres cylindrical. Tibial spurs  $1.5\times$  the tibial diameter, hind tibia with a row of dorsal bristles, fore tibia with one apical spur, hind tibia with two, midtibial spurs obscured. Abdominal tergite 4 longest

and widest. Seventh segment short,  $0.5\times$  the sixth; eighth segment very short, retracted. Ninth tergite with two triangular dorsal processes. **Wing:** Costa ends beyond tip of  $R_5$  at  $2/3$  the length between tips of  $R_5$  and  $M_1$ . Length of Sc  $0.31\times$  the wing length. Sc ends at C proximal to *m-cu*.  $R_1$   $0.54\times$  the wing length.  $R_1$  and  $R_{2+3}$  nearly parallel, straight. Ratios of distances between tips of  $R_1$ ,  $R_{2+3}$ , and  $R_5$  along C are 1.2:1.6:1.  $R_{4+5}$  stem and  $R_{2+3}$  divergent for entire length.  $R_4$  short, straight,  $0.5\times$  length of  $R_5$ . Crossvein *r-m* as long as section of RS between bases of  $R_{2+3}$  and *r-m*. Stem of M weak, but can be traced nearly to base of wing. Length of section of  $M_{1+2}$  stem proximal to *r-m*  $1.4\times$  that of *r-m*; section distal to *r-m*  $3.4\times$  that of *r-m*. Fork of  $M_1$  and  $M_2$   $3.7\times$  longer than its stem. Crossvein *tb* absent, *m-cu* touches M stem. Ratios of distances between apices of  $R_5$ ,  $M_1$ ,  $M_2$ ,  $M_{3+4}$ , and CuA at wing margin are 1:1.3:1.4:2.5. CuA evenly curved.

**MATERIAL:** Holotype VMNH 2923, VMNH 914, ♂ (part and counterpart).

**ETYMOLOGY:** The species epithet is from Latin *latus*, meaning "wide". and *venter*, meaning "abdomen".

#### SUPERFAMILY ANISOPODOIDEA KNAB, 1912

The superfamily includes about 200 species in three small families: Mesozoic Protoryphidae, Jurassic-Recent Anisopodidae sensu lato, and Triassic Crosaphididae. It is often believed to be the living sister group to the Brachycera (Woodley, 1989; Oosterbroek and Courtney, 1996).

#### FAMILY PROTORHYPHIDAE HANDLIRSCH, 1906

#### *Brachyrhynchus* Blagoderov and Grimaldi, new genus

**DIAGNOSIS:** Sc  $0.45\times$  the wing length, ends at C slightly beyond the level of the basal fork of M, level with base of  $R_{2+3}$ .  $R_{2+3}$  slightly converging with  $R_1$ ;  $R_4$   $0.25\times$  the length of stem of  $R_{4+5}$ ,  $0.28\times$  length of  $R_5$ , almost straight. M stem faint, crossvein *m-m* connects  $M_2$  and  $M_3$ . CuA evenly curved posteriad. The new genus differs from all known Protoryphidae by the very short vein  $R_4$ .

**TYPE SPECIES:** *Brachyrhynchus distortus* n.sp.

**ETYMOLOGY:** The genus name is from the Greek βραχης, meaning "short", and *Rhynchus* (a generic name in the family), in reference to the short  $R_4$ . The name is masculine.

**COMMENTS:** Five genera and 10 species of Protoryphidae are known from the Triassic to Late Jurassic-Early Cretaceous deposits of Eurasia. *Vymrhynchus blagoderovi* Krzemiński and Krzemińska, 2003, from the mid-Triassic of France differs from the other two species of the genus by  $R_{2+3}$  converging with  $R_1$  and diverging with  $R_4$ , *r-m* situated distal to  $R_4$  base, and by the very distal position of the discal cell. Thus, it could be considered as a separate genus.

#### *Brachyrhynchus distortus*

Blagoderov and Grimaldi, new species  
figures 7a, 11e

**DIAGNOSIS:** As for genus.

**DESCRIPTION:** Measurements: body length (partial) = 3.1 mm, wing length = 1.7 mm. **Wing:**  $R_1$  straight,  $R_{2+3}$  slightly sigmoid, converging but not fusing with  $R_1$ .  $R_4$  almost straight, very short; length of  $R_5$   $3.5\times$  that of  $R_4$ . Section of RS between  $R_{2+3}$  and *r-m* about equal to length of *r-m*. Costa continues beyond apex of  $R_5$  at least  $1/2$  distance between the apices of  $R_5$  and  $M_1$  (this portion of both wings not completely preserved). Crossveins *r-m* and *tb* near the middle of the discal cell. Crossvein *m-m* connects  $M_2$  and  $M_3$ . M stem short, evanescent. Apices of  $M_1$ ,  $M_2$ ,  $M_3$ , and  $M_4$  almost evenly spaced at wing margin. Distance at wing margin between apices of  $M_4$  and CuA  $2\times$  the distance between apices of CuA and CuP.

**MATERIAL:** Holotype VMNH 2927, ♀.

**ETYMOLOGY:** The species epithet is from the Latin *distortus*, meaning "distorted", in reference to the distorted proportions of the specimen as a result of fossilization.

**COMMENTS:** The new genus from the Virginia Triassic is unique among Protoryphidae in its possession of a very short  $R_4$ . A distal shift and reduction of  $R_4$  is common among Bibionomorpha s.l. (*Alinka*, Paraxymyiidae, Eoditomyiidae) and Brachycera, and probably is a result of costalization (see below

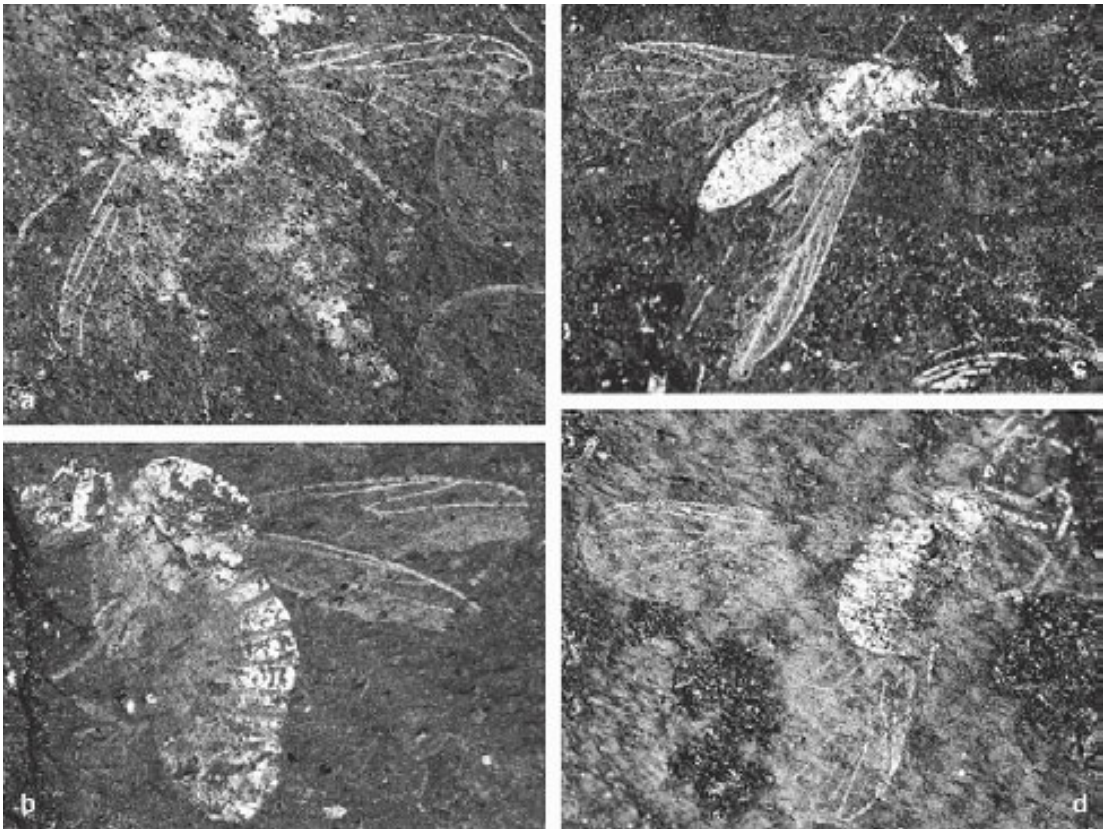


Fig. 7. Photomicrographs of bibionomorphan (a, b) and possible stem-group brachyceran (c, d) Diptera from the Cow Branch Formation. **a.** *Brachyrhyphus distortus* Blagoderov and Grimaldi, holotype VMNH 2927. **b.** *?Crosaphis virginiensis* Blagoderov and Grimaldi, holotype VMNH 797. **c.** *Prosechamyia trimedia* Blagoderov and Grimaldi, holotype VMNH 1371. **d.** *Prosechamyia dimedia* Blagoderov and Grimaldi, VMNH 957. For size scales see fig. 12.

“Phylogeny”). Anisopodidae, which appears to be a sister group or crown-group relative of Protorhyphidae, have lost  $R_4$ , thus the new genus demonstrates an intermediate condition between the two families.

FAMILY CROSAPHIDIDAE KOVALEV, 1983

*?Crosaphis virginiensis*

Blagoderov and Grimaldi, new species  
figures 7b, 12a

**DIAGNOSIS:** Sc short, ends considerably proximal to base of RS; tip of  $R_1$  at level of base of  $M_{1+2}$  fork; base of  $R_{2+3}$  distal to base of  $M_{3+4}$ .

**DESCRIPTION:** Male. Measurements: body length = 1.78 mm, wing = 1.36 mm.

Antennae moniliform with 12–13 flagellomeres, flagellomere length nearly equal to the width; scape and pedicel rounded, slightly wider than flagellum. Anterodorsal tubercles may be present on first abdominal tergite. **Wing:** Sc short, its length  $0.28\times$  the wing length, meets C well before the base of RS.  $Sc_2$  not apparent.  $R_4$  absent. Base of  $R_{2+3}$  distal to the base of  $M_{3+4}$ . Stem of  $M_{1+2}$  fork at least  $2\times$  the length of section of  $M_{1+2}$  proximal to  $r-m$ . Very slight angle in CuA kinked at  $m-cu$ , apical part of CuA straight. Legs either without or with very short tibial spurs. **Abdomen** broad and short, with hemispheric apical segment, presumably  $\delta$  genital capsule including gonocoxites.

**MATERIAL:** Holotype VMNH 797,  $\delta$ .

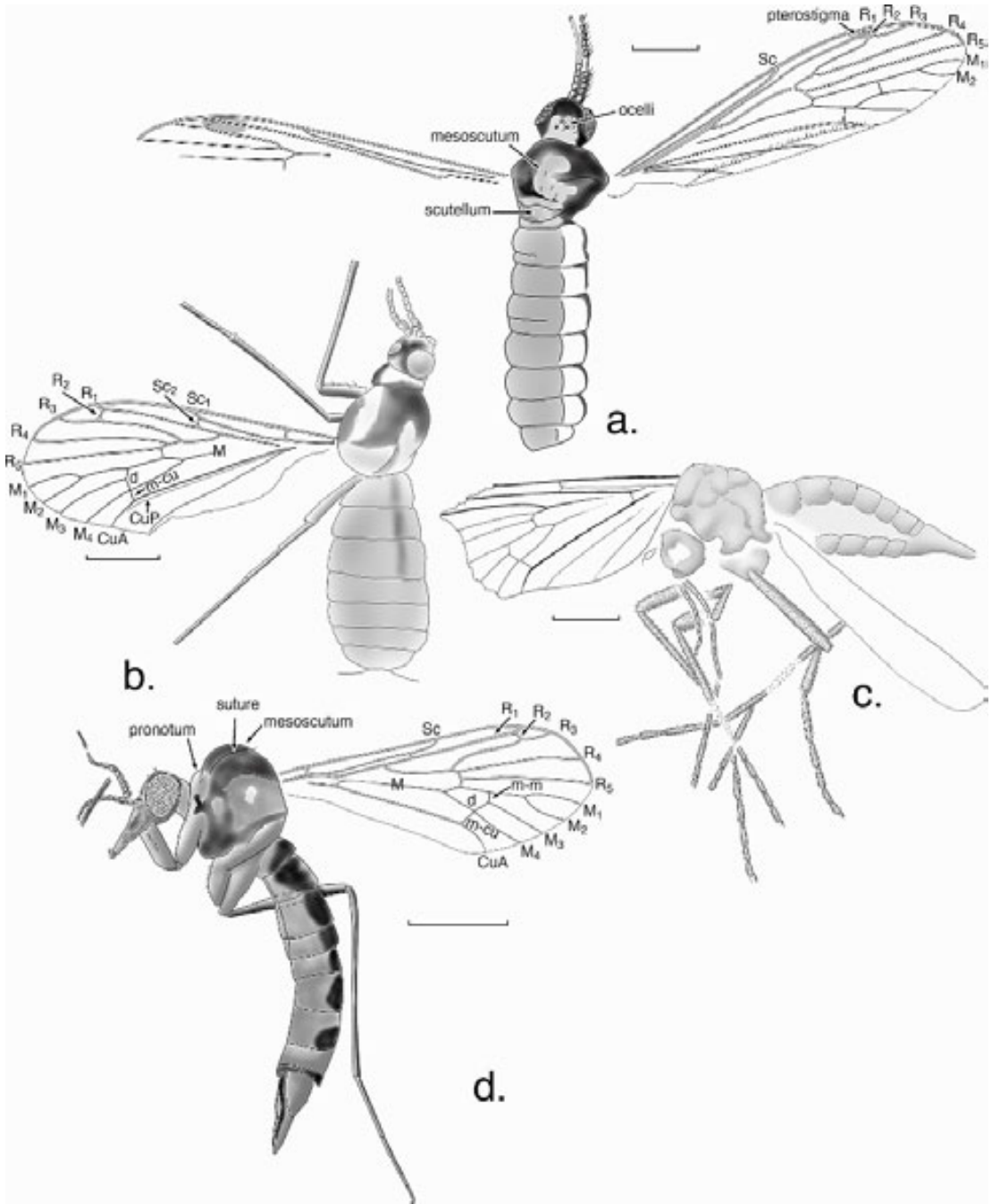


Fig. 8. Tipulomorpha from the Solite quarries. **a.** *Architipula youngi* Krzemiński YPM 35986 (Architipulinae) **b.** *Metarchilimonia solita* Blagoderov and Grimaldi, VMNH 3013. **c.** *Metarchilimonia krzeminskorum*, VMNH 3672. **d.** *Metarchilimonia krzeminskorum* Blagoderov and Grimaldi, VMNH 732. Scales = 0.5 mm.

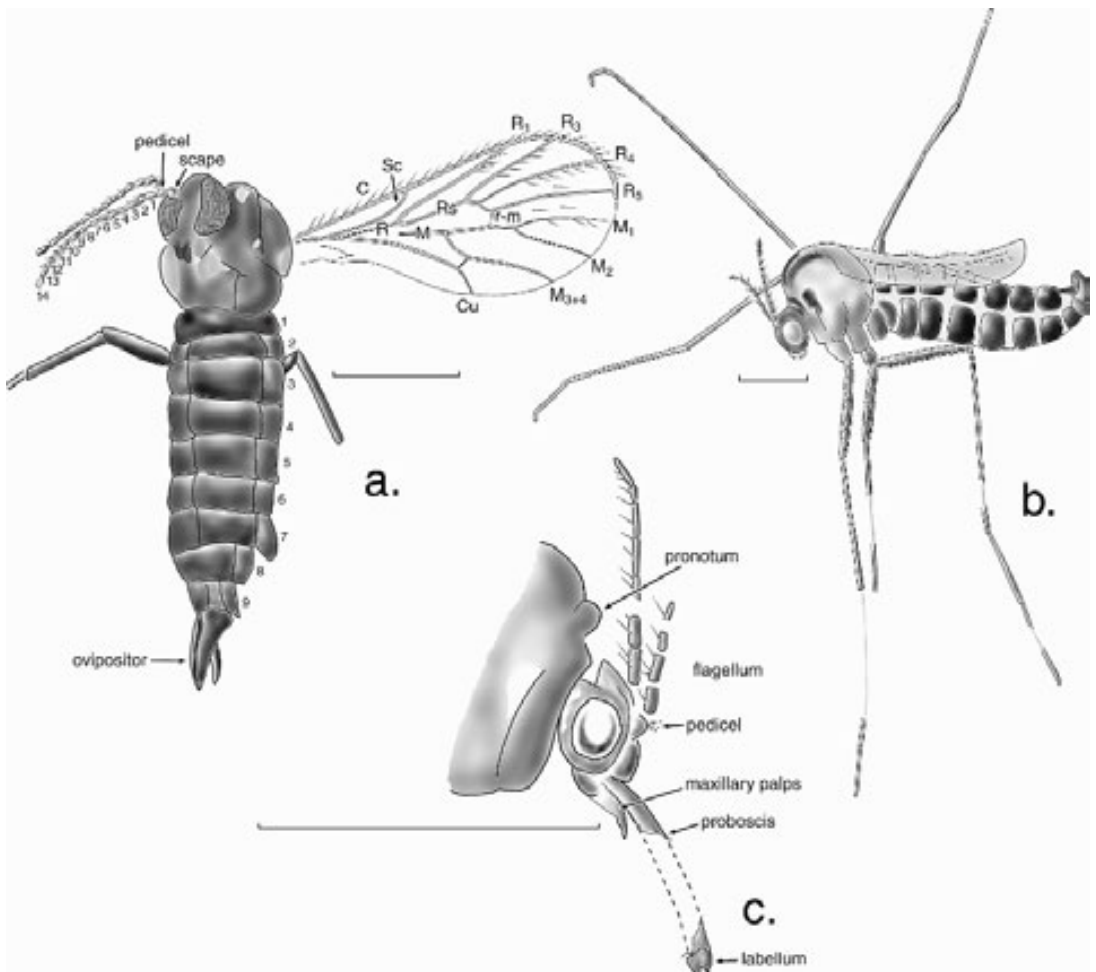


Fig. 9. Psychodomorpha and Culicomorpha from the Solite quarries. **a.** *Triassopsychoda olseni* Blagoderov and Grimaldi (?Psychodidae), VMNH 733. **b.** Culicomorpha (Chironomoidea?), VMNH 951. **c.** Detail of head of culicomorphan VMNH 951 (obverse), showing a long proboscis. Long proboscides in Culicomorpha are generally associated with blood feeding, so this may be the earliest instance of specialization for blood feeding in the fossil record. Scales = 0.5 mm.

**ETYMOLOGY:** From Virginia, the state of origin of the fossil.

**DISCUSSION:** *?Crosaphis virginiensis* is smaller than the other described Bibionomorpha from the Solite quarries. It differs from Paraxymyidae by the longer stem of  $M_{1+2}$ , which is at least  $2\times$  longer than the section of  $M_{1+2}$  proximal to *r-m*.  $R_4$  is either absent or not apparent because the apex of the wing is not preserved. The apical part of CuA is almost straight. These features ally this species with *Crosaphis*, and the short coxae, which are smaller than in Sciarioidea, together

with short or no apparent tibial spurs, suggest the species belongs in Anisopodoidea.

Incomplete preservation of the type species indicates that this species can only tentatively be assigned to the genus *Crosaphis*. *Crosaphis anomala* Evans, 1956, was described from a single wing impression from the Late Triassic of Australia. After restudying the fossil, V. Kovalev (1983) concluded that it did not belong to Hemiptera, as Evans supposed, but rather was close to mycetobiine Anisopodidae. Restudy of the holotype of *Crosaphis anomala* in the Queensland Museum

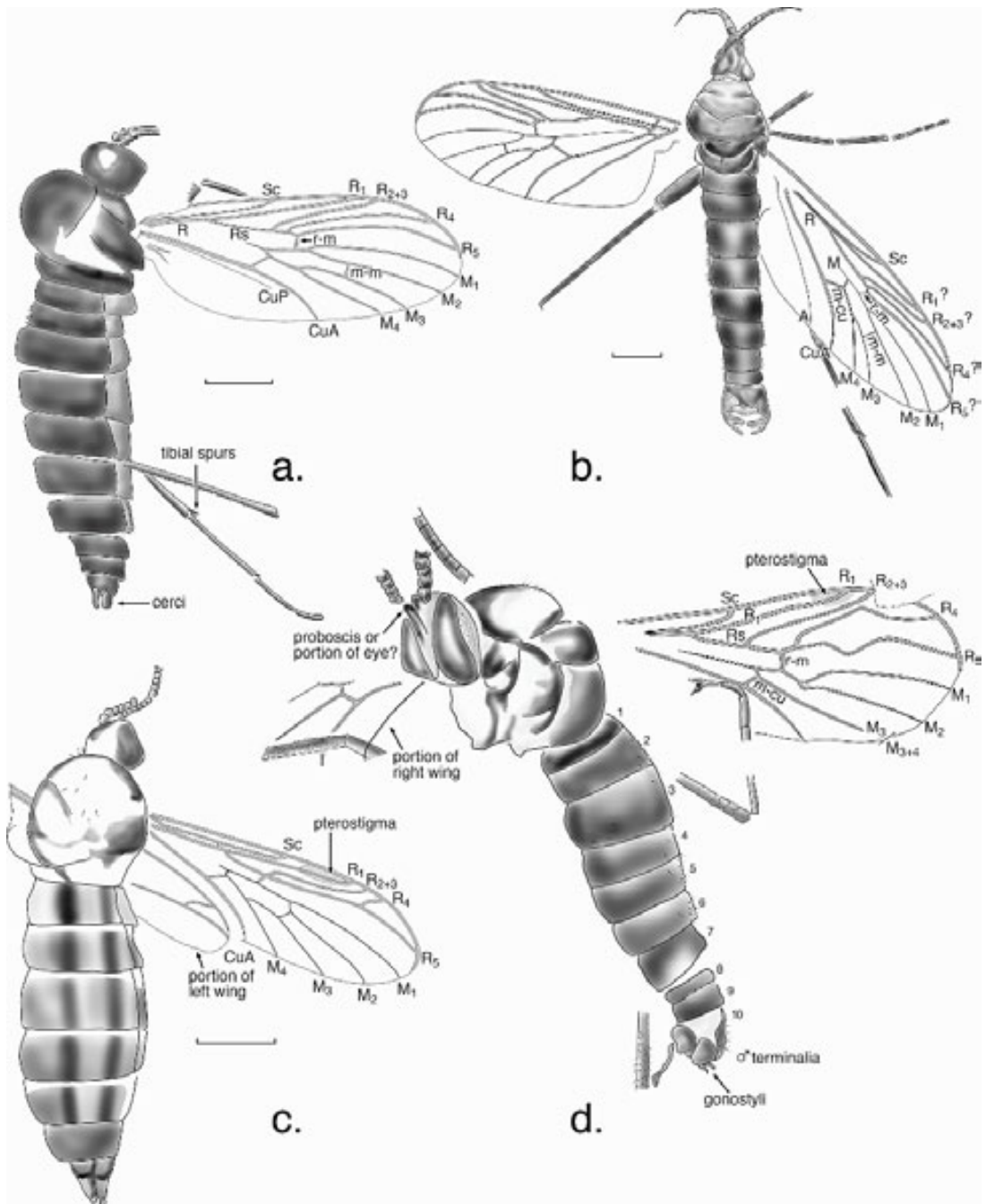


Fig. 10. Bibionomorpha from the Solite quarries. **a.** *Yalea rectimedia* Blagoderov and Grimaldi (Procramptonomyiidae), VMNH 1041. **b.** *Yalea argentata* (Krzemiński) (Procramptonomyiidae), VMNH 3673. **c.** *Alinka cara* Krzemiński (Procramptonomyiidae), YPM. **d.** *Veriplecia rugosa* Blagoderov and Grimaldi (Paraxymyiidae), VMNH 35989.

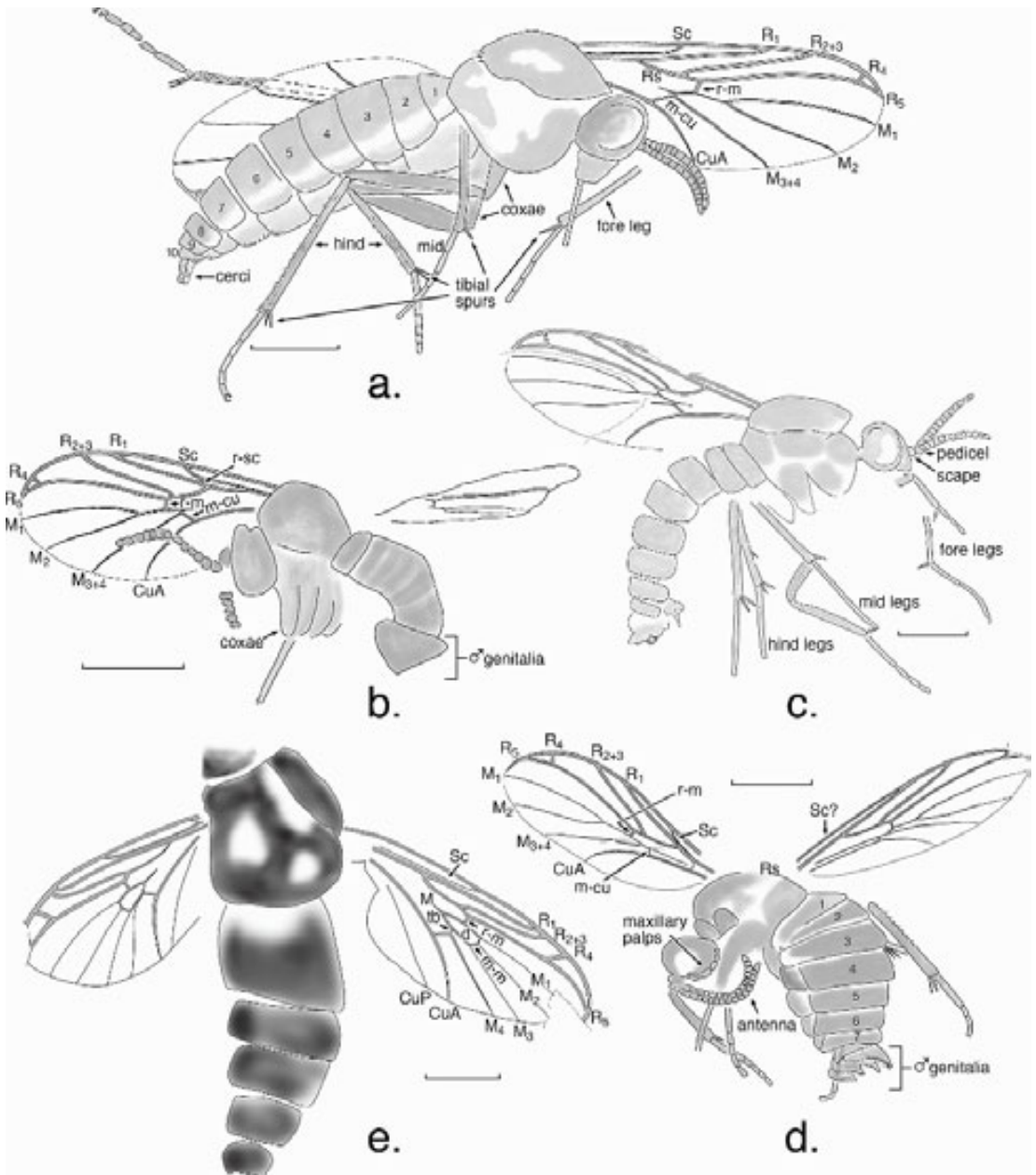


Fig. 11. Bibionomorpha from the Solite quarries. **a.** *Virginiptera certa* Blagoderov and Grimaldi, female, VMNH 731. **b.** *Virginiptera similis* Blagoderov and Grimaldi, VMNH 825. **c.** *Virginiptera certa*, male, VMNH 2998. **d.** *Virginiptera lativentra* Blagoderov and Grimaldi, holotype male, VMNH 2923. **e.** *Brachyrhynchus distortus* Blagoderov and Grimaldi (Protorhynchidae), VMNH 2927. Not to the same scale (scales = 0.5 mm).

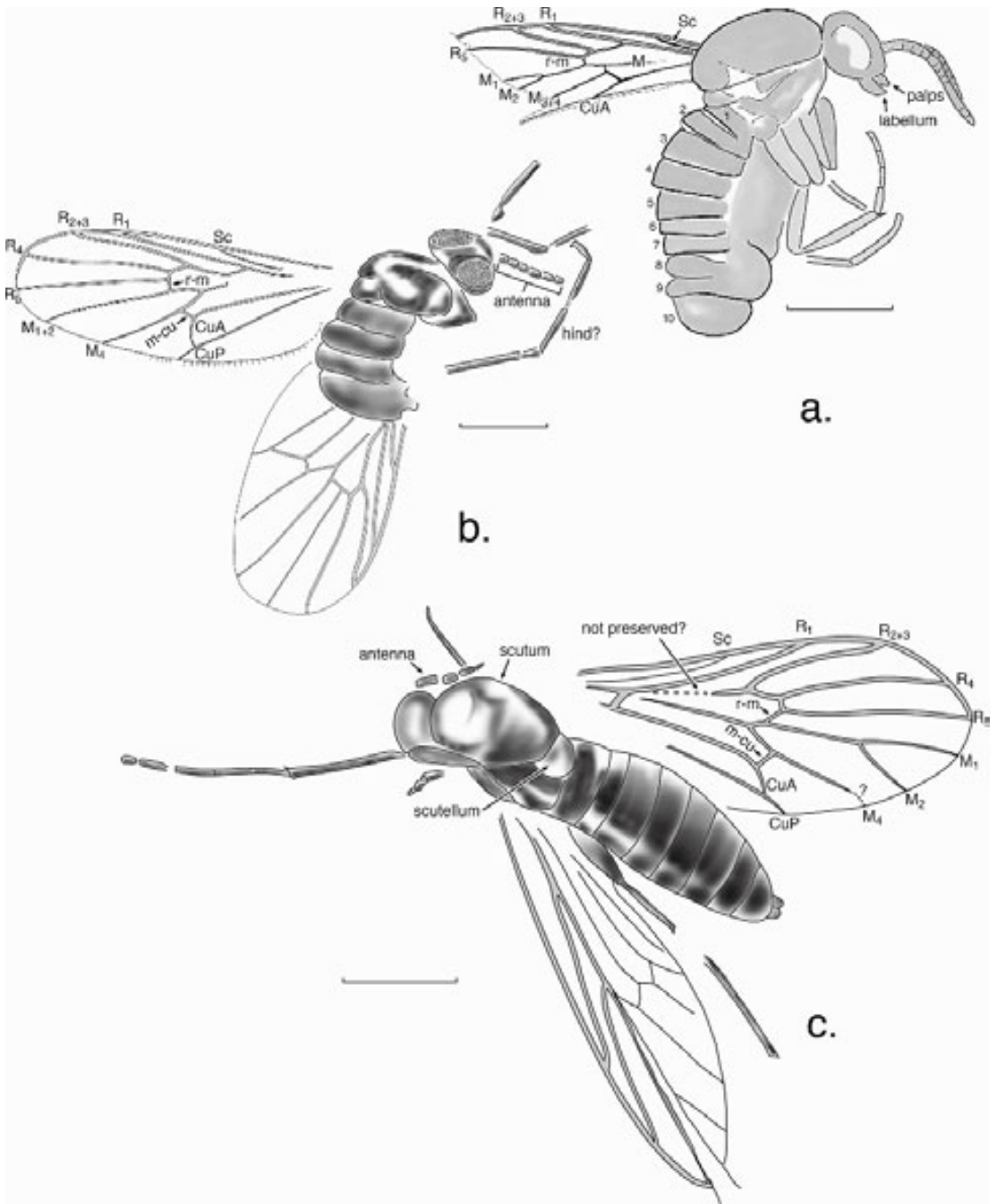


Fig. 12. Assorted Diptera from the Solite quarries. **a.** *?Crosaphis virginiensis* Blagoderov and Grimaldi (Crosaphididae: Anisopodoidea). VMNH 797. **b.** *Prosechamyia dimedia* Blagoderov and Grimaldi (Prosechamyiidae), VMNH 957. **c.** *Prosechamyia trimedia* Blagoderov and Grimaldi, VMNH 1371. *Prosechamyia* appears to be an extinct stem group to the large, diverse lineage of flies, the Brachycera. Scales = 0.5 mm.

by one of us (DG) found it to be poorly preserved. Although the presence of parts of the venation were consistent with the venation drawn by Kovalev (1983), much venation was not observable against the rough matrix. It is possible that there may have been some deterioration since Kovalev studied it. Evenhuis (Evenhuis, 1994; Krzemiński and Evenhuis, 2000) doubted the position of *Crosaphis* in Anisopodoidea, on account of the venation that is supposedly similar to that of Bombyliidae mythicomiine. Not only is Late Triassic far too early for a group as recently derived (phylogenetically) as Bombyliidae, but a nearly complete crosaphid specimen indicates it is not even a brachyceran. Amorim and Tozoni (1994) placed *Crosaphis* in the Mycetobiidae, a living family to which crosaphids are very closely related.

STEM-GROUP BRACHYCERA  
FAMILY PROSECHAMYIIDAE, NEW FAMILY

DIAGNOSIS: CuA meets CuP well before wing margin and making a closed cell, as in most Brachycera; M<sub>4</sub> lost, as in many Bibionomorpha and some living and extinct "Rhagionidae" s.l. (e.g., *Bolbomyia*, *Palaeobolbomyia*, *Mesobolbomyia*, *Litoleptis*), *m-m* crossvein absent, so there is no discal cell; *r-m* meeting R<sub>5</sub>; R<sub>4</sub> and R<sub>5</sub> very long and connected to basal cell; antennae plesiomorphically moniliform.

TYPE GENUS: *Prosechamyia* n.gen.

***Prosechamyia*** Blagoderov and Grimaldi,  
new genus

DIAGNOSIS: Same as for family.

TYPE SPECIES: *Prosechamyia trimedia* n.sp.

ETYMOLOGY: The genus name is a compound one from the Greek words προσεχής, meaning "bordering" or "adjacent to", and μύια, meaning "fly". The name is feminine.

DISCUSSION: The new genus has a unique combination of characters unknown for any Diptera, living or extinct. The petiolate anal cell is a synapomorphy of virtually all Brachycera (some species, such as *Pseudoernia*, some Rhagionidae and Bombyliidae have CuA and CuP converging but not meeting).

On the other hand, all earliest known Brachycera (Protobranchyceridae, Rhagionidae, Oligophryneidae, and others) have very generalized venation with a discal cell, four branches of vein M, and rather short R<sub>4</sub> and R<sub>5</sub> veins. The new genus has no closed discal cell, no fork of M<sub>3+4</sub>, very long branches of R<sub>4</sub> and R<sub>5</sub>, and moniliform antennae. All living Brachycera, and fossil ones with excellent preservation, have moderately to highly reduced antennae with differentiated antennal segments. Antennae of *Prosechamyia trimedia* are not preserved well enough to determine the number of flagellomeres with certainty, but apical segments are tapered, suggestive of basal Brachycera and resembling the condition observed in *Litoleptis* Chillcott, *Arthroceras* Williston (Rhagionidae), Xylophagidae, and Xylomyiidae. Differentiation of flagellomeres in *P. dimedia* is not preserved. Other synapomorphies of living Brachycera, like posterior elongation of the larval head into the prothorax, vertical movements of larval mandibles, or the reduced number of maxillary palpomeres (from five to two) in adults unfortunately could not be observed in the fossil material.

*Prosechamyia* definitely is not in Bibionomorpha s.l., some of which have a reduced discal cell and three branches of the radius (e.g., Protopleciidae, Crosaphididae). In those bibionomorph groups R<sub>2+3</sub> is usually curved anteriorly and the tips of R<sub>1</sub> and R<sub>2+3</sub> are close to each other; the base of M<sub>3+4</sub> is strongly kinked at *m-cu*, suggesting an ancestral state of discal cell structure similar to that of Procramptonomyiidae. The new genus has radial branches almost evenly spaced as in many Brachycera, and M forks symmetrically, similar to Protorhaphidae, Anisopodidae, and many basal Brachycera. The similarity in the wing venation of *Prosechamyia* to the Recent genus *Litoleptis* is especially striking: it differs from the latter by the longer fork of R<sub>4</sub> and R<sub>5</sub>, crossvein *r-m* connected directly to R<sub>5</sub> and by the much longer basal cell. Thus, the only synapomorphy of the new genus with true Brachycera is a convergence of CuA and CuP forming a petiolate cell. It is probable that the new genus constitutes an extinct stem group to true Brachycera, having acquired only one of numerous brachyceran synapomorphies.

Caution must be exercised when interpreting relationships—particularly one as dramatic as a sister group to all Brachycera—on the basis of one character. An interesting example of independent evolution of brachyceran characters occurs in some Archizelmiridae, a family of basal Sciaroidea from the Late Jurassic to Late Cretaceous. In one species of this family the antenna is very similar to that in Brachycera, complete with apical flagellomeres modified into a hairlike arista (Grimaldi et al., 2003). If such striking convergence can occur in a relatively complex structure, it seems at least equally probable for there to be convergence in the structure of vein Cu as well.

*Prosechamyia trimedia*

Blagoderov and Grimaldi, new species  
figures 7c, 12c

DIAGNOSIS: Three medial veins,  $M_1$  and  $M_2$  strongly divergent; base of  $M_{1+2}$ – $M_{3+4}$  fork at the level of  $R_{2+3}$  base;  $R_4$  and  $R_5$  connected directly to basal cell, without a stem; length of  $r-m$  nearly equal to basal section of  $M_{1+2}$ ; basal section of  $M_{3+4}$  twice the length of basal sections of  $M_{1+2}$ .

DESCRIPTION: Measurements: body length 1.9 mm, wing length 1.7 mm. Head round, eyes large. Only four flagellomeres seen, but these are widely separated and presumably some were lost; flagellomeres tapered in width apicad, presumably apical one 1/4 size of most basal preserved one. Sc ends at C at level of base of  $R_{2+3}$ . Sc is  $0.42\times$  length of wing.  $R_1$  short,  $0.58\times$  the wing length. Base of RS apparently not preserved. Crossvein  $r-m$  oblique, connecting to  $R_5$  near its base. Length of  $R_{4+5}$   $2\times$  the length of basal section of  $R_5$  and  $1.5\times$  the length of  $r-m$ . Base of  $M_{1+2}$ – $M_{3+4}$  fork at the level of the base of  $R_{2+3}$ . Stem of  $M_{1+2}$   $1.2\times$  that of  $M_1$ , the section distal to  $r-m$  equal in length to  $M_1$ .  $M_2$  widely divergent from  $M_1$ . Basal section of  $M_{3+4}$   $2\times$  the length of basal section of  $M_{1+2}$ . Crossvein  $m-m$  and vein  $M_3$  absent.  $M_4$  incompletely preserved in unique specimen. CuA forms closed anal cell with CuP, with stem approximately equal to length of  $m-cu$ . Legs rather long, slender, no tibial spurs visible. Eight visible segments of

abdomen, terminalia not apparent, sex unknown.

MATERIAL: Holotype VMNH 1371.

ETYMOLOGY: In reference to the number of medial veins.

*Prosechamyia dimedia*

Blagoderov and Grimaldi, new species  
figures 7d, 12b

DIAGNOSIS: Differs from *P. trimedia* by: vein  $M_{1+2}$  simple, not forked;  $M_{1+2}$ – $M_{3+4}$  stem proximal to level of  $R_{2+3}$  base (vs. at the same level);  $r-m$   $0.5\times$  the length of basal section of  $M_{1+2}$  (vs. nearly equal), and  $r-m$  transverse (vs. oblique); basal sections of  $M_{1+2}$  and  $M_4$  (proximal to  $r-m$  and  $m-cu$ ) equal in length (vs. basal section of  $M_4$  twice the length).

DESCRIPTION: Measurements: body length 1.7 mm, wing length 1.8 mm. Head and eyes large. Six segments of antenna visible, only slightly tapered to most apically preserved flagellomere. Sc ends at C; its length  $0.4\times$  wing length. Length of  $R_{4+5}$   $2\times$  that of basal section of  $R_5$  and  $1.2\times$  the length of  $r-m$ . Crossvein  $r-m$  transverse, veins  $M_{1+2}$  and  $M_{3+4}$  simple (apparently  $M_2$  and  $M_3$  lost),  $m-cu$  absent. Basal sections of  $M_{1+2}$  and  $M_{3+4}$ , proximal to crossveins  $r-m$  and  $m-cu$  respectively, are subequal. Tibia (hind) with at least one tibial spur, slightly longer than tibial diameter. End of abdomen curled ventrally; sex unknown, but probably male.

MATERIAL: Holotype VMNH 957.

ETYMOLOGY: In reference to the number of medial veins.

DISCUSSION

PHYLOGENY

This study is an attempt to determine the phylogenetic placement of early fossils of the order Diptera; it is not intended to be the definitive solution to the poorly understood relationships among nematoceros flies. Nematoceran phylogeny has proven problematic thus far, and there is significant disagreement among the major studies. Disagreement may be due to the fact that most of these studies have been based on certain character systems; alternatively, the major lineages may have arisen contemporaneously or nearly so

(and so relationships would be difficult to decipher). Many studies, for example, hypothesize that Tipulomorpha is the sister group to the rest of the Diptera (Hennig, 1954, 1973; Wood and Borkent, 1989; Michelsen, 1996), but one major study hypothesizes that the Tipulidae is much more recently derived and very closely related to the Brachycera (Oosterbroek and Courtney, 1995). Several studies have been largely or wholly relegated to characters of immatures (Wood and Borkent, 1989; Sinclair, 1992); or to particular areas of the adult body, such as the cervical region (Michelsen, 1996) or venation, particularly of fossils (Krzemiński and Krzemińska, 2003; Shcherbakov et al., 1995). Phylogenetic hypotheses by Hennig (1954, 1973) and Oosterbroek and Courtney (1995) were based on immatures and adults. Oddly, there is no comprehensive study of nematoceran relationships based on DNA sequences, and the only one that has been done was based on one gene and minimal taxon sampling (Friedrich and Tautz, 1997). Other DNA studies, mentioned below, have focused on particular infraorders or families. A serious caveat of DNA studies, of course, is that they cannot incorporate fossil DNA, and so they must incorporate morphological structures of living and fossil species to understand the placement of extinct taxa. No study has incorporated many morphological characters of living and extinct Diptera.

We examined the phylogeny of living and extinct Diptera using the morphology of 75 exemplar taxa, representing 45 Mesozoic genera and 30 basal genera of living families of nematoceran Diptera, and two basal families of living Brachycera (Xylomyidae and Xylophagidae). Eighty morphological characters were studied, which represented most major body structures but also included external and internal genitalia of both sexes. Of the 80 morphological characters, 37 (46%) of them were based on wing structure, shape, and venation, which is essential for interpreting Triassic fossil Diptera since most of these taxa are wholly or largely defined by venation. Of the 6,000 cells in the data matrix, 1,891 (31%) of them were not scored or were considered ambiguous because of sexual dimorphism or incomplete preservation of

fossils. Most of these unscored cells were characters 30–74 and pertain to minute structures on the body that are generally obscured in compression fossil insects. Most characters were coded for just two states (0, plesiomorphic; 1, apomorphic), but 11 characters with two or more apomorphic states were also scored. With the exception of character 80, all states were nonadditive. The phylogenetic program NONA (Goloboff, 1999) was used, employing the parsimony ratchet with 200 iterations (10 trees per iteration), which generated 264 most-parsimonious trees (MPTs) of length 324, CI 30, RI 72. Figure 13 is the strict consensus of these MPTs, which is some 51 steps longer (L 375, CI 26, RI 66). As expected with a consensus tree, there is less resolution than in the MPTs, in this case primarily among the basal Bibionomorpha. Otherwise, the relationships within the other infraorders and major taxa are well resolved in the consensus tree.

**Tipulomorpha** in the present study is the sister group to the remaining Diptera, as is traditionally hypothesized (e.g., Hennig, 1954, 1973; Wood and Borkent, 1989), and there is a significant grade of extinct taxa in this infraorder from the Triassic to the present. The infraorder typically includes just the Tipulidae s.l. (or Tipulidae s.s., *Cylindrotomidae* s.s., and *Limoniidae* s.s.), though sometimes the winter crane flies, family Trichoceridae, are also included (e.g., Oosterbroek and Courtney, 1995). Typical breeding sites of tipulids are the sediments of shallow streams and ponds, and organically rich mud and wet soils; trichocerids breed in decayed wood. Tipulidae s.l. is the largest family of Diptera with approximately 14,000 described species.

**Blephariceromorpha** in this study is represented by only one Recent taxon (*Edwardsina*: Blephariceridae) and a primitive Jurassic genus, *Ansoargius*. Deuterophlebiidae is another Recent family in the group, and monophyly of the infraorder is essentially certain. The odd, neotenic family Nymphomyiidae is related either to these (Oosterbroek and Courtney, 1995; Wood and Borkent, 1989) or it belongs to the Culicomorpha (Courtney, 1994). This infraorder, which is the smallest one (approximately 100 species worldwide), is

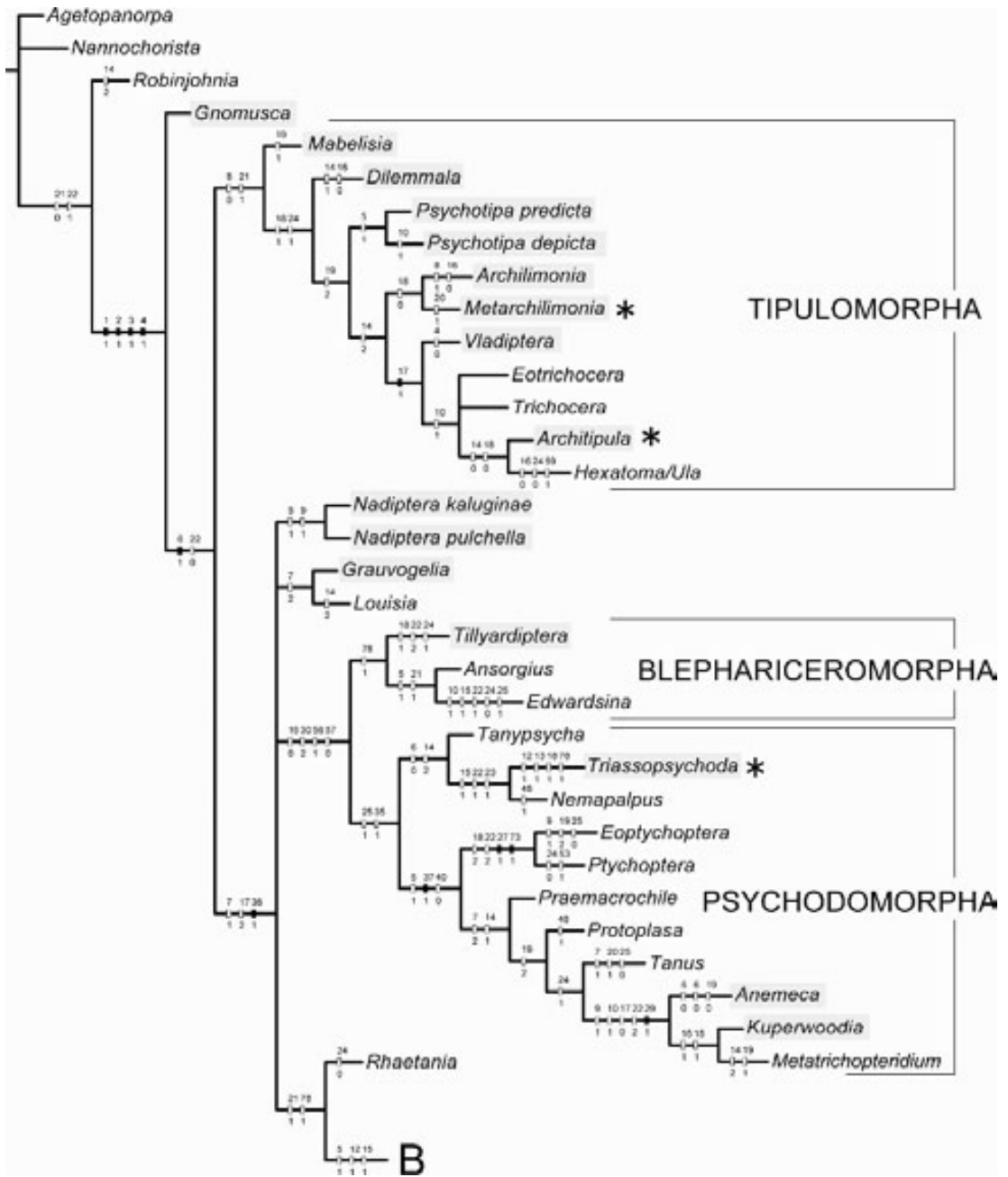


Fig. 13. Strict consensus tree (L 375, CI 26, RI 66) of 264 most-parsimonious trees, based on 80 morphological characters of 75 exemplar genera of Diptera (45 of them Mesozoic; see tables 2, 3). Shaded taxa are from the Triassic (others are later Mesozoic and Cenozoic); those with an asterisk are from the Solite quarries. Major infraorders of the Diptera appeared by the Late Triassic.

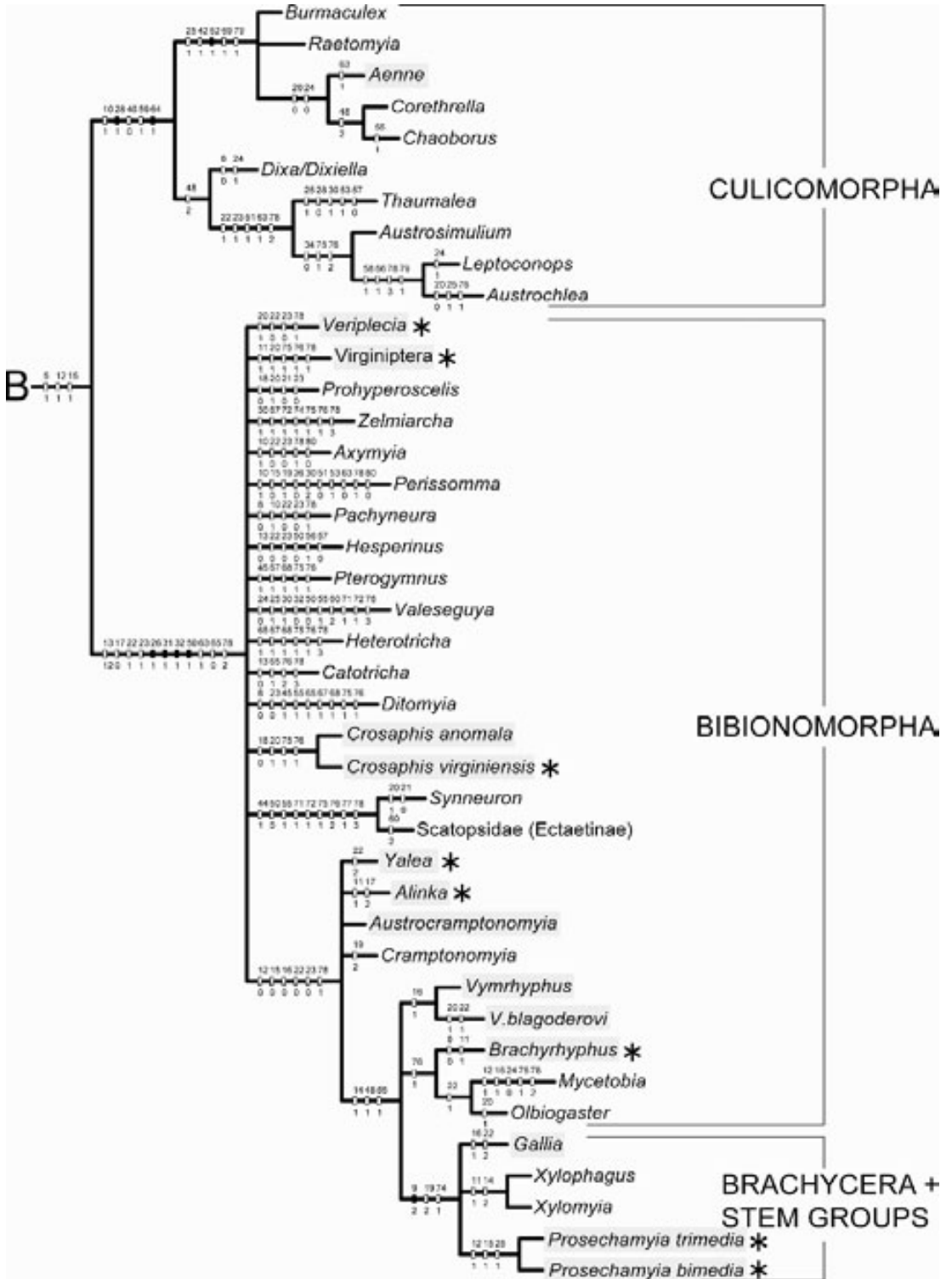


Fig. 13. Continued

TABLE 2  
Character Matrix of Select Mesozoic and Recent Basal Diptera (A = 01, B = 02; see table 3 for characters)

	1	1111111	112	22	222	2222	333333	3333	444444	444	4	5555555555	6666666666	7777777778
	1234567890	1234567	890	12	345	6789	012345	6789	012345	678	9	0123456789	0123456789	0123456789
Agetopanorpa	0000000100	0000000	000	12	120	0000	-----	--00	-00-0?	000	-	-----?	--0	-----00000-
Robinjohnia	0000000100	0002000	000	01	000	0000	-----	---0	-----	-00	-	-----?	--0	-----0000-
Nannochorista	0000000100	0000010	000	12	020	0000	000000	0000	000000	000	0	020011?100	0000-11000	00000000000
Vladiptera	1110010000	0002011	120	10	010	0000	-----	---	-----	--0	-	-----?	--0	-----0000-
Dilemmala	11110--000	0001000	100	10	010	0000	-----	---	-----	--0	-	-----?	--0	-----0000-
Psychotipia	1111110000	0000010	120	10	010	0000	-----	---	-----	--0	-	-----?	--0	-----0000-
Psychotipadepicta	1111110001	-000010	120	10	010	0000	-----	---	-----	--0	-	-----?	--0	-----0000-
Archilimonia	1111010100	0002000	020	10	010	0000	-----	---	-----	--0	-	-----?	--0	-----0000-
Mabelisia	1111010000	0000010	010	10	020	0000	-----	---	-----	--0	-	-----?	--0	-----0000-
Architipula	1111010001	-000011	020	10	010	0000	-----	--00	-00--	000	-	-----?00	--0	-----00000-
Metarchilimonia	1111010000	0002010	021	10	01A	0000	-----	--00	-00--	000	-	-----?0	--0	-----00000-
Gnomusca	1111000100	0000010	001	01	000	0000	-----	---	-----	--0	-	-----?	--0	-----0000-
Grauvogelia	1111012100	0000012	001	00	020	0000	-----	---	-----	--0	-	-----?	--0	-----0000-
Lousia	1111012100	0002012	001	00	020	0000	-----	---	-----	--0	-	-----?	--0	-----0000-
Nadipterakaluginae	1111111110	0000012	001	00	020	0000	-----	---	-----	--0	-	-----?	--0	-----0000-
Nadipterapulchella	1111111110	0000012	000	00	020	0000	-----	---	-----	--0	-	-----?	--0	-----0000-
Tanus	1111111100	0001002	021	00	010	0000	-----	-1--	-----	--0	-	-----?	--0	-----0000-
Kuperwoodia	1111112111	-001010	120	02	011	0001	-----	---	-----	--0	-	-----?	--0	-----0000-
Anemeca	1111002111	-001000	000	02	011	0001	-----	---	-----	--0	-	-----?	--0	-----0000-
Tillyardiptera	1111011100	0000002	100	02	010	0000	-----	---	-----	--0	-	-----?	--0	-----0001-
Rhaetania	1111011100	0000012	001	10	000	0000	-----	---	-----	--0	-	-----?	--0	-----0001-
Triassopsychoda	111100-1-0	011-1-2	1-1	01	121	0000	-----1	--00	-1?-	-00	-	-----?0	--0	-----0001-
Aenne	1111111101	-10-1-2	0-0	10	001	0010	-----	---	-----	--0	-	-----?	--1	-----0001-
Yalea	1111111100	0010000	100	12	020	1000	-----	--00	-0?-	-00	-	-----?0	--1	-----00010-
Alinka	1111111100	1010002	100	10	020	1000	-----	--00	-0?-	-00	-	-----?0	--1	-----00010-
Austrocramptonomyia	1111111100	0010000	100	10	020	1000	-----	---	-----	--0	-	-----?	--1	-----0001-
Vymrhypus	1111111100	0001010	000	10	020	1000	-----	---	-----	--1	-	-----?	--1	-----0001-
Vblagoderovi	1111111100	0001010	001	11	020	1000	-----	---	-----	--1	-	-----?	--1	-----0001-
Brachyrhynchus	111111-000	1011000	000	10	020	1000	-----	---	-0?-	--1	-	-----?0	--1	-----01010-
Veriplecta	111111-100	011-1-0	1-1	10	020	1000	-----	---	-0?-	--0	-	-----?0	--1	-----00010-
Virginiptera	111111-1-0	112-1-0	1-1	11	120	1000	-----	--00	-0?-	--0	1	-----?0	--1	-----11010-
Crosaphisanomala	111111-100	-11-1-0	0-1	11	120	1000	-----	---	-----	--?	-	-----?	--1	-----1102-
Crosaphisvirgin	111111-1-0	-11-1-0	0-1	11	120	1000	-----	--00	-0?-	--?	1	-----?0	--1	-----11020-
Gallia	111111-120	0001010	020	12	000	1000	-----	---	-?-?	--1	-	-----?0	--1	-----0001?-
Prosechamiatrimedia	111111-120	010-1-0	1-0	10	001	1000	-----	--00?	-0?-	--01	-	-----?0	--1	-----100010-

TABLE 2  
(Continued)

1	111111	112	22	222	2222	333333	33333	444444	444	4	5555555555	6666666666	7777777777	
1234567890	1234567	890	12	345	6789	012345	6789	012345	678	9	0123456789	0123456789	0123456789	
Prosechmyadimedia	11111-120	010-1-0	1-0	10	001	1000	---	--0?	-01	-	-----?0-	---1-----	----100010-	
Eoptychoptera	1111111100	0000002	221	02	0100	-----	-100	-00---	000	1	--00---	0000?1???	?001000000-	
Ansorgus	1111111100	0000002	000	10	020	0000	-----0	-10?0-	100	-	-----?00	---0-----	-----00010-	
Praemacrocchile	1111112100	0001002	000	0B	021	0000	-----	-?0-	--0	1	--0-----?	---0-----	-----0000--	
Eotrichocera	1111010001	-002011	120	12	010	0000	-----	---	--0	-	-----?	---0-----	-----0000--	
Rhaetomyia	111111-101	-10-1-2	0-1	10	021	0010	-----	---	--0	-	-----?	---0-----	-----0001--	
Metatrichopteridium	1111112111	-002010	110	02	011	0001	-----	---	--0	-	-----?	---0-----	-----0000--	
Tanypsyscha	111100-100	0002002	001	00	021	0000	-----	---	--0	-	-----?	---0-----	-----0000--	
Prohyperoscelis	1111111100	-12-1-0	0-1	01	020	1000	-----	---	--0	-	-----?	---0-----	-----0002--	
Zelmaicera	1111111100	-12-1-0	1-0	11	120	--00	1110-0	1000	000	1	1?001---	0001000100	00101110302	
Ptychoptera	1111111100	0000002	200	02	001	0100	200011	1100	000000	000	1	0001112000	0000011000	000100000000
Edwardsina	1111111101	-0001-2	0-0	11	001	0000	200010	?000	110000	100	0	0100011000	0100?11000	00000000100
Protoplasa	1111112100	0001002	020	00	021	0000	200111	?100	000000	001	0	0000111001	0000011000	00000000000
Hexatoma Uja	1111010001	-00000A	020	12	000	0000	000111	0000	100000	000	0	0200110101	0000011000	10000000000
Axymyia	1111111101	-11-1-0	1-0	10	020	1000	011010	1000	100000	000	1	120010?100	0001000000	00000000100
Trichocera	1111010001	-002011	120	12	010	0000	100110	0000	100000	000	1	020010?100	0000010000	10000000000
Perissomma	1111111101	-12-010	110	11	120	0000	211010	1000	100000	000	1	1001100100	0000000000	00000000100
Synneuron	1111111100	-12-1-0	0-1	01	120	1000	211010	1000	100010	000	1	020011?100	0001000000	01100121302
Scatopsidae Ectatinae	1111111100	-12-1-0	2-0	11	120	-000	111010	100A	100010	000	1	020011?100	2001000000	01100121302
Nemapalpus	1111001100	00021-2	0-1	01	121	0000	200011	1000	101110	001	1	0?0011?001	0000011001	00000000000
Dixa Dixiella	1111111001	-10?1-2	121	10	010	0010	000011	1010	000000	002	1	0200100101	0000?10000	00000000101
Corethrella	1111111101	-10-1-2	0-0	10	001	0010	000000	1010	001000	002	1	0210000111	00001?1001	00000000111
Chaoborus	1111111101	-10-1-2	0-0	10	001	0010	000000	1010	001000	002	1	0210001111	0000101001	00000000111
Thaumalea	1111111101	-10?1-2	2-1	11	121	0000	100011	1010	000000	002	1	0101100001	0001110000	00000000201
Austrosimulium	1111111101	-10?1-2	2-1	11	120	0-10	000001	1010	000000	002	1	010010-101	0001110000	00000012101
Leptoconops	11111111-1	-10?1-2	1-1	11	110	0-10	00000A	1010	001000	002	1	0100101111	0001111000	00000121311
Pachyneura	1111111001	-11?1-0	1-0	10	020	1000	011010	1000	100000	000	1	120010?100	00010?0000	00000000102
Cramptonomyia	1111111100	0010000	120	10	020	1000	011010	1000	100000	000	1	120010?100	00010?0000	00000000102
Hesperimus	1111111100	-10?1-B	1-0	10	02?	1000	011010	1000	10000A	000	1	0200101000	00010?0000	00000000202
Mycetobia	1111111100	-11-1-0	2-0	11	000	1000	111010	1000	100010	011	1	1200110100	0001000000	00000110202
Olbigaster	1111111100	0001000	201	11	020	1000	111010	1000	100010	011	1	1200110100	0001000000	00000010102
Pterogymnus	1111111100	-12-1-0	1-0	11	120	1000	011010	1000	100001	000	1	120010?100	00010?0110	00000110202
Burmaculex	1111111101	-10-1-2	0-1	10	021	0010	?-11	1000	?01000	00? 1	0?101?10-	0000111001	0000000011?	
Valesguya	1111111100	-12-1-0	1-0	11	101	-000	110010	1000	100000	000	1	020011?100	2001000000	0110000030?
Austrochlea	111111110?	?10-1-2	0-0	11	121	-010	??001	1010	?0?000	002	1	0?00101111	0001111000	00000110311

TABLE 2  
(Continued)

1	1111111	112	22	222	222	333333	3333	444444	444	4	5555555555	6666666666	7777777777
	1234567890	890	12	345	6789	012345	6789	012345	678	9	0123456789	0123456789	0123456789
	111111100	?11-1-0	1-0	11	120	-000	0?1010	1000	10000?	000	1	1?0010?100	0001010110
Heterotricha	111111100	?10-1-0	1-0	11	120	-000	0?1010	1000	?0000?	000	1	1?0010?100	0001010000
Catotricha	111111000	-11-1-B	1-0	11	020	1000	011010	1000	100001	000	1	1200110100	0001010110
Ditomylia	111111120	1002000	120	10	020	1000	311010	1001	100000	001	1	020011?100	1001000000
Xylophagus	111111120	1002000	120	10	020	1000	311010	1001	100000	001	1	020011?100	1001000000
Xylomyia													

closely related to the Psychodomorpha in our analysis. Larvae live on rocks in shallow, cold, swift streams, especially near waterfalls.

The infraorder **Psychodomorpha** has been one of the most problematic of all major lineages in the Diptera. It typically contains the family of moth flies, Psychodidae (Michelsen, 1996; Oosterbroek and Courtney, 1995), but some authors also include in this infraorder the phantom crane flies (Ptychopteridae) and Tanyderidae (Hennig, 1973; this study), as well as the blephariceromorphs (Hennig, 1973). Wood and Borkent (1989) included in this infraorder the Psychodidae, Trichoceridae, Perissomatidae, Anisopodidae, and Scatopsioidea. Psychodidae and Scatopsioidea breed in wet, foul substrates.

**Culicomorpha** is the least problematic of the major infraorders, since there is good consensus on the monophyly, composition, and relationships (Hennig, 1954, 1973; Wood and Borkent, 1989; Oosterbroek and Courtney, 1995; Miller et al., 1997; Pawlowski et al., 1997). Our results likewise indicate a monophyletic Culicomorpha, but relationships differ slightly with the other major studies, perhaps because of minimal sampling in this study. *Aenne*, which is known from the Triassic of Britain (Krzemiński and Jarzembowski, 1999) to the Cretaceous of Lebanon (in amber), belongs to the Chironomidae, so in the cladogram in figure 13 it should be the sister group to the other chironomid, which is the very basal living genus *Austrochlea*. Culicomorphan larvae are essentially aquatic, and the ancestral diet of adult females is insect hemolymph and/or vertebrate blood (Grimaldi and Engel, 2005). An undetermined culicomorphan from the Solite quarries reported here has a long proboscis, which is suggestive of its use in sucking blood. As such, it would be the oldest known example among insects of a structure specialized for blood feeding.

**Bibionomorpha** is a large, complex infraorder whose monophyly is well supported in this study, though internal relationships are poorly resolved. Better resolution is indicated by some studies, such as the close relation of *Valeseguya* (Valeseguyidae) to the Scatopsioidea (Scatopsidae + Synneuridae; Amorim and Grimaldi, 2006). Relationships

TABLE 3

Characters Used in Phylogenetic Analysis of Select Mesozoic and Recent Basal Diptera<sup>a</sup>

1. Branches of vein Sc: (0) two or more, (1) one.
2. Base of vein R: (0) straight, (1) kinked.
3. Veins M<sub>5</sub> and CuA: (0) fused, (1) separated.
4. Vein A<sub>2</sub>: (0) visible, (1) reduced or lost.
5. Allular incision on wing: (0) absent, (1) present.
6. Anal lobe of wing: (0) absent, (1) present.
7. Shape of vein 1A: (0) reaching wing margin distal to cu-a, (1) reduced, not reaching wing margin, (2) reaching wing margin proximally to cu-a (anal loop).
8. Apex of vein CuP: (0) in distal half, (1) in basal half.
9. Veins CuP and CuA: (0) parallel or slightly divergent or convergent, (1) strongly divergent, (2) convergent, forms anal cell.
10. Branching of vein RS: (0) R<sub>2+3</sub>+R<sub>4+5</sub>, (1) R<sub>2+3+4</sub>+R<sub>5</sub>.
11. Fork of vein R<sub>4+5</sub>: (0) long, (1) shorter than its stem.
12. Vein M<sub>3+4</sub>: (0) forked, (1) M<sub>3</sub> lost.
13. Base of vein MP (M<sub>1</sub> section): (0) developed, (1) vestigial, (2) lost.
14. Vein *m-cu* joins: (0) M<sub>3+4</sub>, (1) M<sub>4</sub>, short stalk between discal cell and *m-cu*, (2) M<sub>3+4</sub> fork base.
15. Vein *im*: (0) present, (1) absent.
16. Vein *im* joining: (0) M<sub>2</sub>, (1) M<sub>1+2</sub> or base of fork.
17. M forking: (0) symmetrically, (1) discal cell anterior of M stem, (2) discal cell posterior of M stem.
18. CuA distad of *m-cu*: (0) rather straight, (1) kinked and strongly curved posteriorly, (2) straight or curved anteriorly.
19. Vein *r-m* joining discal cell: (0) at the middle of discal cell, (1) distally, (2) proximally.
20. Base of discal cell: (0) proximal to CuP apex or at the same level, (1) distal.
21. Section RS<sub>1</sub> length: (0) at least 3× the length of R base, (1) less than 2× the R base.
22. Sc length: (0) about 0.5× the wing length, (1) shorter than 0.4×, (2) longer than 0.6×.
23. CuP: (0) complete, (1) reduced, not meeting wing margin.
24. Bases of veins *r-m* and *m-cu*: (0) at same level, line connecting them almost vertically, (1) *r-m* proximad, (2) *m-cu* proximad.
25. Base of vein R<sub>5</sub>: (0) oblique or horizontal, (1) vertical, aligned with *r-m*.
26. Tip of vein R<sub>3</sub>: (0) tips of R<sub>1</sub>, R<sub>3</sub>, R<sub>4</sub> and R<sub>5</sub> evenly spaced, (1) R<sub>3</sub> closer to R<sub>1</sub> rather than R<sub>4</sub>.
27. Relief of vein CuP: (0) convex for entire length, (1) strongly convex distally.
28. Radial cells: (0) same width, (1) cell R<sub>5</sub>/M<sub>1</sub> wider than R<sub>4</sub>/R<sub>5</sub>, parallel-sided.
29. Vein R<sub>5</sub>: (0) normal, (1) continues to the base of wing as a fold.
30. Meron: (0) contiguous with coxa 2, (1) associated with epimeron 2, (2) fused with epm 2 to meropleurite, (3) inconspicuous.
31. Precervical sclerite: (0) absent, (1) present.
32. Laterocervical sclerite: (0) contacts proepisternum, (1) contacts episternal lobe.
33. Laterocervical sclerite: (0) without orifice, (1) with orifice.
34. Mandibles: (0) present and usually better developed or only functional in female, (1) absent.
35. Ocelli: (0) present, (1) absent.
36. Arolium: (0) present, (1) replaced with empodium.
37. Last tarsomere of male: (1) normal, not prehensile, (1) capable of folding against penultimate tarsomere.
38. Pedicel: (0) normal, (1) enlarged.
39. Flagellomeres: (0) 10-14, (1) fewer than 8.
40. Parameres: (0) separate, (1) fused to each other.
41. Claws: (0) normal, not folding, (1) fold against distitarsus.
42. Length of setae on flagellomeres: (0) short, (1) as long as flagellomere.
43. Antennal scoids: (0) absent, (1) present.
44. Movement of gonostyli: (0) horizontal, (1) vertical.
45. Aedeagus: (0) tubular, (1) flattened.
46. Middle legs: (0) normal, (1) shortest of the three pairs.
47. Dorsolateral tubercles on tergite 1: (0) absent, (1) present.
48. MA (arculus): (0) slightly sclerotized, mostly as a crease, (1) as a strong vein.
49. Antepnotum and proepisternum: (0) antepnotum much larger than proepisternum, (1) antepnotum about the size of proepisternum.
50. Basalar pit: (0) absent, (1) present.
51. Postcervical sclerite: (0) normal, (1) produced mesoventrally or absent.

TABLE 3  
(Continued)

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52.	Posterior margin of wing: (0) with setae, (1) with scales.
53.	Transepimeral suture: (0) present, (1) absent.
54.	Coronal suture: (0) present, (1) absent.
55.	Ninth sternite of male: (0) well developed, (1) reduced to apodemes.
56.	Spermathecal ducts: (0) open separately, (1) with common openings, (2) partly fused.
57.	Subalifer: (0) present, (1) absent.
58.	Distribution of setae on flagellomeres: (0) irregular, (1) in whorls.
59.	Ocelli: (0) present, (1) vestigial or absent.
60.	Maxillary palp: (0) 3–5 segments, (1) 2 segments, (2) 1 segment, (3) lost.
61.	Wings: (0) normal, (1) dehiscent.
62.	Wing position at rest: (0) along the body, (1) over and away from body.
63.	Costa: (0) circumambient, (1) abbreviated between tips of veins R <sub>5</sub> and M <sub>1</sub> .
64.	Empodium: (0) pulvilliform, (1) setiform.
65.	Pulvilli: (0) present, (1) absent.
66.	Clypeus: (0) small, anterior margin flat or rounded, (1) long and pointed.
67.	Coxae: (0) short, (1) as long as thorax height or longer.
68.	Tibial spurs: (0) short or absent, (1) several times longer than tibial diameter.
69.	Setulae on wing veins: (0) short, (1) long.
70.	Male cerci: (0) present, (1) absent.
71.	Mesothorax vestiture setae: (0) present, (1) mesothorax devoid of macrosetae.
72.	Katepisternum: (0) vertical, (1) extended posteriorly, mesopleural suture kinked.
73.	Prehalter: (0) absent, (1) present.
74.	Flagellomere specialization: (0) flagellomeres moniliform or clavate, (1) flagellum shortened and apical part forms arista or style.
75.	Vein distribution: (0) even through membrane, (1) radial veins condensed with costa.
76.	Vein thickness: (0) even, (1) medial veins weakened, (2) medial and cubital veins weakened.
77.	Length of radial veins: (0) continue to the tip of the wing, (1) not approaching the tip of the wing.
78.	Number of radial branches: (0) 5, (1) 4, (2) 3, (3) 2.
79.	Setae on flagellomeres: (0) short, (1) male antennae plumose.
80.	Hair plates: (0) absent, (1) radial and axillary, (2) radial and anal (Rohdendorf, 1946).

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<sup>a</sup>Character 80 is additive; the rest are nonadditive.

among the Sciaroidea (fungus gnats and allies) have been well treated (Chandler, 2002; Jaschhof, 2001; Jaschhof and Didham, 2002; Blagoderov and Grimaldi, 2004; Hipa and Vilkamaa, 2005; Vilkamaa and Hipa, 2004), and *Veriplecia* and *Virginiptera* (Paraxymyiidae: Late Triassic to Cretaceous) appear to be the basal members of the infraorder. Interestingly, the larval biology of Bibionomorpha is unique among nematoceros flies, in that they are largely terrestrial like the large, monophyletic group, the Brachycera (bibionomorphans, however, feed primarily on fungal matter; most basal brachyceran larvae are predaceous). Several studies (i.e., Hennig, 1973), including this one, break with Wood and Borkent (1989) and Oosterbroek and Courtney (1995) in including Anisopodidae in the Bibionomorpha. In fact, it

appears that Anisopodidae is probably the sister group to the Brachycera, thus rendering the Bibionomorpha paraphyletic. Anisopodidae is represented in the Triassic by *Vymrhyphus*, and fossils of the family become more diverse through the later Mesozoic.

**Brachycera** is the largest and most diverse infraorder in the Diptera, with 110 families and probably over 100,000 described species. It is a definitively monophyletic group as based on, among other features: (1) larval features (obviously not preserved in fossils); (2) reduction of the antennal flagellum to eight or fewer segments (commonly four or fewer), and significant differentiation of the flagellomeres; (3) reduction of the maxillary palp to 1–2 segments; (4) veins CuA and CuP (often interpreted as CuA<sub>2</sub> and A<sub>1</sub>) with their apices meeting before the wing margin (forming

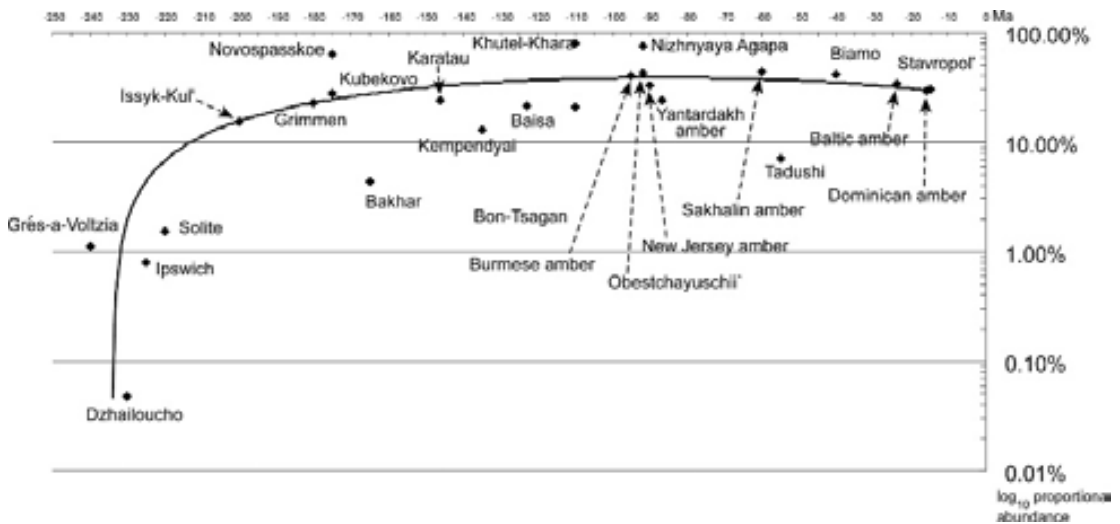


Fig. 14. Relative abundance of Diptera through time, as measured by  $\log_{10}$  proportion of Diptera specimens among total number of fossil insect specimens from representative, significant deposits (see table 4). Dipteran abundance asymptotes toward modern levels in the Jurassic because of diversification within infraorders.

a closed cell) or nearly so. Fusion of veins  $R_2$  and  $R_3$  (Hennig, 1973) has been suggested as a fourth defining character, but this is widespread among nematocerous flies. *Gallia*, from the mid-Triassic of France, is known only from the wing, and it has been classified as a brachyceran (Krzemiński and Krzemińska, 2003). This is based, however, just on wing venation. Our discovery of *Prosechamyia* from the Solite quarries reveals an additional Triassic fly with convergent Cu veins, but the antennae (albeit incomplete) are filiform, indicating *Prosechamyia* is a stem group to true Brachycera. Thus, in lieu of antennal and other structures of *Gallia*, it is quite possible that it too is a stem group to true brachycerans.

#### DIPTERA ABUNDANCE AND DIVERSITY IN THE TRIASSIC

Our results indicate that stem-group and even some derived members of most major dipteran infraorders existed by the Late Triassic. Diptera, however, were rare in the Triassic, which contrasts starkly with their abundance later in the Mesozoic (fig. 14). For example, in the Triassic deposits at Dzhailoucho (Kyrgyzstan, Central Asia) only

11 fly specimens were found among approximately 20,000 insects; in the Ipswich Series of Australia only four out of 500 (Anderson and Anderson, 1993); and at Grés-a-Voltzia in France only 59 adults out of 5,300 other insects (Krzemiński and Krzemińska, 2003; Marchal-Papier et al., 2000). In the Early Jurassic of Germany—some 40 million years later—Diptera comprised approximately 23% of all insect specimens, second only to Hemiptera (26%) (Ansorge, 1996); and in the Late Jurassic of the spectacular insect Concentrat Lagerstätte of Karatau, Kazakhstan, Diptera comprised 24% (4,791 of 19,879) of all insect specimens ([www.palaeontolog.ru](http://www.palaeontolog.ru)). By the Early Cretaceous, Diptera comprised approximately 21% of all insect specimens, such as in the diverse deposits at Baissa, Siberia. These proportions of course reflect biases in the preservation of lacustrine and marine littoral deposits, but the Cretaceous and Tertiary abundance of Diptera is also apparent in amber from those periods, with Diptera (particularly Chironomidae and Ceratopogonidae) typically being the most abundant insect order in amber (e.g., Rasnitsyn and Quicke, 2002; Grimaldi, 2000; Grimaldi et al., 2002).

TABLE 4  
Abundance of Diptera in Major Deposits of Fossil Insects from the Triassic to Neogene  
(C, compression; A, amber)

Locality	Age	%	Insects (N)	Diptera (N)	Taphonomy	Reference
Grés-a-Voltzia	-240	1.11%	5300	59	C	Krzemiński, 2003, Papier et al., 2000
Dzhailoucho	-230	0.05%	23332	11	C	www.palaeontomolog.ru
Ipswich	-225	0.80%	500	4	C	www.palaeontomolog.ru
Solite	-220	1.52%	3300	50	C	
Issyk-Kul'	-200	15.23%	2836	432	C	www.palaeontomolog.ru
Grimmen	-180	22.67%	1200	272	C	Ansorge, 1996
Novospasskoe	-175	63.26%	3100	1961	C	Kalugina, Kovalev, 1985
Kubekovo	-175	28.18%	1022	288	C	www.palaeontomolog.ru
Bakhar	-165	4.35%	5421	236	C	www.palaeontomolog.ru
Karatau	-146	24.10%	19879	4791	C	www.palaeontomolog.ru
Kempendyai	-135	12.94%	1182	153	C	www.palaeontomolog.ru
Baissa	-123	21.25%	24476	5200	C	www.palaeontomolog.ru
Bon-Tsagan	-110	21.12%	9959	2103	C	www.palaeontomolog.ru
Khutel-Khara	-110	80.01%	3331	2665	C	Sontag, 2003
Myanmar	-95	40.17%	3012	1210	A	Grimaldi et al., 2002
Obestchayuschii	-92	42.85%	1202	515	C	www.palaeontomolog.ru
Nizhnyaya Agapa	-92	75.10%	257	193	A	www.palaeontomolog.ru
New Jersey	-90	33.46%	1052	352	A	Grimaldi et al., 2000
Yantardakh	-87	23.89%	854	204	A	www.palaeontomolog.ru
Sakhalin	-60	43.41%	940	408	A	Zherikhin, 1978
Tadushi	-55	7.16%	3843	275	C	www.palaeontomolog.ru
Baltic	-40	41.44%	7111	2947	A	www.palaeontomolog.ru
Biamo	-24	34.17%	1718	587	C	www.palaeontomolog.ru
Dominican Rep.	-16	29.65%	11814	3503	A	D.Grimaldi, unpubl.
Stavropol'	-15	30.53%	3449	1053	C	www.palaeontomolog.ru

Diptera from the Triassic of Virginia comprise some 1.5% of all insect specimens, which is comparable in abundance to other Triassic deposits. Overall, the trend indicates that Diptera were rare in the Triassic, comprising approximately 1% of all insects and dramatically increasing in abundance in the latest Triassic to earliest Jurassic. By the Late Jurassic ~150 Ma the abundance of Diptera nearly reached that of the Tertiary level, and remained there. By the end of the Triassic flies were global. All Diptera from the locality that has yielded the oldest such specimens, the Grés-a-Voltzia sandstones, belong to extinct and the most basal lineages. Indeed, most basal families (Vladipteridae, Archilimoniidae, Grauvogeliidae, Nadipteridae) occur in Europe and Central Asia. A virtual absence of Diptera in the Australian Triassic is possibly a sampling problem, but insects from the Newcastle Formation in Belmont, Australia are very distinctive among what are otherwise globally uniform Permian insect faunas, so the

Australian Triassic Diptera could reflect regional differences (Ponomarenko and Sukacheva, 1998). There has been little change in the tectonic position of Australia during those 30 Ma between the Triassic and Permian deposits.

In conclusion, the Diptera evolved from a Late Permian *Permotipula* or *Robinjohnia*-like ancestor that had saprophagous immatures and adults. The exact timing is difficult to discern since the Early Triassic is virtually unknown for insects, but we hypothesize that true, two-winged Diptera originated in the earliest Triassic ca. 245 Ma. By 20 Ma after the Permo-Triassic boundary, stem groups and some derived members of the major infraorders had appeared. This would suggest a very rapid diversification or radiation of Diptera in the Early Triassic, which could be the basis for why relationships among infraorders and other higher lineages of living nematoceros flies appear to be so obscure.

## ACKNOWLEDGMENTS

We thank Tam Nguyen and Steve Thurston for their help in composing the plates; to the former Solite Corporation, particularly John Roberts and C. H. Gover, Richard Familia and Giant Cement, and to Roy Larimer for their help with the excavations; to Tim White for loans of specimens from the Yale Peabody Museum; and to Michael Engel and an anonymous reviewer for their helpful commentary. Lastly, we are indebted to the National Geographic Society and the National Science Foundation (NSF EAR 0106309 to NF and DG) for funding this project.

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