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# Phylogeny and classification of the shrimp genera *Acetes, Peisos*, and *Sicyonella* (Sergestidae: Crustacea: Decapoda)

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Despite their role in marine systems, Sergestidae remain one of the most poorly understood families amongst planktonic shrimps with regard to phylogeny. Recent morphological and phylogenetic revisions of a number of sergestid genera have disentangled classificatory problems and emphasized the importance of reproductive structures for the taxonomy and phylogeny of the Sergestidae. Only three genera, *Acetes*, *Peisos*, and *Sicyonella*, remain unrevised phylogenetically. We undertook a phylogenetic analysis of these groups based on 124 morphological characters (120 binary, four multistate). Eighteen new characters were based on scanning electron microscopy studies of the clasping organ and petasma. The phylogenetic analysis revealed statistically supported monophyly of the clades *Sicyonella* and *Acetes* + *Peisos*. We combine *Peisos* and *Acetes* into a monophyletic genus *Acetes*, give emended diagnoses and keys to all species of *Sicyonella* and *Acetes*, and discuss morphological trends within these genera. We present maps of geographical distribution for all valid species of *Acetes*.

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ADDITIONAL KEYWORDS: anatomy – cladistic analysis – identification key – morphological phylogenetics – morphology – phylogenetics – taxonomy.

## INTRODUCTION

Sergestid shrimps are important components of marine systems. The family Sergestidae comprises 16 pelagic genera widely distributed in the Atlantic, Indian, and Pacific Oceans (Appendix 1). These genera encompass 74 valid species, which have been recently revised (Vereshchaka, 2000, 2009; Vereshchaka, Olesen & Lunina, 2014; Vereshchaka & Lunina, 2015). In addition to these genera, the Sergestidae comprises three minor genera: Sicyonella with three species, Acetes with 14 species, and the monotypic Peisos. Economically, Acetes is one of the most important organisms in Asian and East African waters (Fig. 1D); during certain parts of the year species of Acetes form conspicuous

aggregations near the shore, leading to an extensive fishing activity (Omori, 1975). Emended diagnoses and keys to species of *Acetes* and a review of their geographical distribution may be important for fishery planning. The genera *Sicyonella* and *Peisos* are not significant for fisheries.

All three species of Sicyonella were recently redescribed with the use of scanning electron microscopy (SEM; Fukuoka, Tamaki & Kikuchi, 2005), whereas the latest revision of Acetes was more than 40 years ago (Omori, 1975). The phylogenetic status of the monotypic Peisos has not been tested since the description of its only species, Peisos petrunkevichi (Burkenroad, 1945). The genus Acetes (Fig. 1A–C) was established by H. Milne Edwards (1830) for Acetes indicus (type by original designation). Thirteen additional species were described between 1893 (Acetes americanus Ortmann, 1893) and 1975 (Acetes intermedius Omori, 1975, and Acetes marinus Omori,

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**Figure 1.** Acetes japonicus in situ, lateral (A) and upper (B) view; Acetes paraguayensis, allotype, ZMUC CRU-09812 in collection of the Natural History Museum of Denmark (C); Acetes sp. at a Indonesian fish market (D).

1975). During this period, the genera *Sicyonella* Borradaile, 1910 and *Peisos* Burkenroad, 1945 were also described.

Recent phylogenetic studies of Sergestidae using cladistic methods resulted in significant taxonomic changes both below and above the genus level (Vereshchaka *et al.*, 2014; Vereshchaka & Lunina, 2015). At this stage, it is appropriate to revise the remaining minor sergestid genera, *Sicyonella*, *Acetes*, and *Peisos*, to bring the phylogenetic studies of the Sergestidae near completion.

In contrast to the other genera of Sergestidae, *Acetes*, *Peisos*, and *Sicyonella* occur in coastal and even fresh

waters and have a unique set of characters compared with other sergestids. Relative to other sergestids, *Acetes* and *Peisos* have simplified mouthparts, rudimentary or absent pereopods, and a simplified petasma, etc. Conversely, *Sicyonella* is characterized by well-developed pereopods with full-sized chelae and a complete set of segments in the natatory pereopods (all pereopods are significantly reduced in the other sergestids), and an elaborated petasma, etc. It remains unclear as to which of these genera represent early branching lineages or rather terminal clades. All three genera are important for understanding the phylogeny of Sergestidae, as has been noted before (Burkenroad, 1945).

The ultrastructure of clasping and copulatory organs has been shown to be important both taxonomically and phylogenetically (Vereshchaka et al., 2014; Vereshchaka & Lunina, 2015). In the primitive state, these organs are simple finger-like, undivided, and lacking specialized additional structures, whereas in the derived state they demonstrate various branching, change in shape, and the presence of numerous minute structures, amongst other specializations. (Vereshchaka et al., 2014; Vereshchaka & Lunina, 2015). In Acetes and Peisos, the clasping and copulatory organs have been studied with the use of light microscopy only (Hansen, 1919; Omori, 1975). No comparative analysis of the morphology of the clasping and copulatory organs has been carried out between Acetes and Peisos, and Sicyonella.

In this paper we provide information on the ultrastructure of the clasping and reproductive organs of *Acetes* and *Peisos* with the use of SEM as the basis for a morphology-based phylogenetic analysis of these groups. Our overall goals were to test the monophyly of all three genera, to analyse their status and position within Sergestidae, to discuss morphological trends within these genera, to revise the classification of the family, and to provide a key to valid species as well as maps of their geographical distribution.

#### MATERIAL AND METHODS

All 14 recognized species of *Acetes*, three species of *Sicyonella*, *Peisos petrunkevichi*, and 16 type species representing all other genera of Sergestidae were included as terminals. Character state scoring for each species was derived from examination of specimens (see Appendix 1). Characters were polarized using three species representing three families of Penaeoidea, each occurring in three different types of oceanic habitats (benthopelagic, pelagic, and benthic) and therefore representing different types of morphology. The three outgroup species – all of which are type species for their genera – were *Aristeomorpha foliacea* (Aristeidae), which is benthopelagic; *Gennadas parvus* (Benthesicymidae), which is pelagic; and *Penaeus monodon* Fabricius, 1798 (Penaeidae), which is benthic.

Character state scoring for each outgroup species was derived from examination of specimens (see Appendix 1).

Prior to treatment for SEM, the relevant parts (clasping organs, petasma, etc.) of selected specimens were dissected in order to expose important structures for detailed study. The material was dehydrated in an ethanol series, critical point dried, mounted, and coated by a mixture of platinum and palladium following standard procedures (e.g. Olesen, Richter & Scholtz, 2003). The scanning electron microscope used was a JEOL JSM-6335F (with a field emission gun). The images were processed and photo plates were created in standard graphical software such as CorelDraw X7 and various Adobe programs.

We used the data matrix of Vereshchaka & Lunina (2015) as a basis for this work. The character states are figured in Vereshchaka (2000, 2009), Vereshchaka *et al.* (2014), and Vereshchaka & Lunina (2015). We used 124 modified characters (120 binary, four multistate, amongst which 18 are new – see Appendix 2). The data matrix is presented in Appendix 3.

Data were handled and analysed using maximum parsimony in a combination of programs: WINCLADA/NONA, Nexus Data Editor (NDE), TNT, and MES-QUITE (Nixon, 1999; Goloboff, Farris & Nixon, 2000).

All characters were unordered (non-additive) and equally weighted; missing data were scored as unknown. Trees were generated in TNT under the 'implicit enumeration' procedure. Relative stability of clades was assessed by standard bootstrapping (sampling with replacement) with 10 000 pseudoreplicates and by Bremer support (tree bisection-reconnection algorithm, saving up to 10 000 trees up to three steps longer). We considered the clades statistically significant if they were supported either by Bremer support  $\geq 3$  or bootstrap values  $\geq 80$ .

#### RESULTS AND DISCUSSION

ULTRASTRUCTURE OF SEXUAL CHARACTERS AND POSSIBLE HOMOLOGIES

As for other sergestids, the sexual structures of *Sicyonella* and *Acetes* (the clasping organ and the petasma) are well known to provide reliable information for classification (Hansen, 1919; Vereshchaka & Lunina, 2015). As not all information was available under light microscopy, SEM was used here for more detailed examination.

Serrated setae are present on the clasping organs and differ from the ordinary setae in being more robust and possessing distal serrations as seen by light microscopy (Vereshchaka, 2000, 2009). However, under SEM it can be seen that the serrated setae exhibit a complicated ultrastructure (Fig. 2). In *Acetes* and *Peisos* 

(except A. americanus and Acetes binghami), the serrated setae have reticulate distal parts and resemble morel mushrooms (Fig. 2A–C), whereas in A. americanus and A. binghami the setae bear longitudinal ribs (Fig. 2D). The function of these setae is uncertain but owing to their robustness they may assist in holding the female during mating and/or have chemo-/mechanoreceptory functions.

The clasping tubercle is very similar in ultrastructure to the serrated setae (Fig. 3A) and we assume that they are homologous. The clasping tubercle is most likely derived from a serrated setae of the outer male antennular flagellum. It is remarkable that *Acetes* has two tubercles; all other sergestid genera have a single one. The second tubercle may be either rudimentary (Figs 4A, C, E, 5C, 6A, 7A, C, 8C, E) or developed (Figs 5A, E, 6C, E, 7E, 8A).

Scales on the clasping organs were reported by Fukuoka et al. (2005) for Sicyonella inermis (Fig. 9A). We found similar structures in Acetes paraguayensis and A. marinus (Fig. 7A, C). Further examination of clasping organs showed that fine scales generally are present (Fig. 3A–F); they are absent only in a few species with rudimentary clasping organs (Fig. 3F). The fine scales most likely assist in clasping of the female during mating. They are absent in two species of Sicyonella (Fig. 9C, E) and in 13 species of Acetes, (Figs 4–6, 7E, 8); in these species they are replaced by two characteristic rows of serrated setae (Sicyonella) or by scattered pairs of robust claw-like setae (Acetes) that suggest an alternative clasping mechanism.

The pars media of the petasma is present in all species of Sicyonella (Fig. 8B, D, F) and Acetes except A. marinus (Fig. 7B). Both genera show much variation in the shape of the pars media, from being entire in S. inermis (Fig. 9B) and vestigial in A. paraguayensis (Fig. 7D) to branched in the rest of Sicvonella (Fig. 9D, F) and in A. americanus (Fig. 4D). The capitulum of the pars media in all Sergestidae, except Acetes, is armed with hooks visible under light microscopy (Vereshchaka, 2000, 2009; Vereshchaka et al., 2014). SEM shows that these hooks are squamose (Fig. 10D-F) and may form pincers in Sicyonella (figs 4, 10, 15 in Fukuoka et al., 2005) and in other sergestids (Fig. 10F). Homologies between lobes and processi in these genera have been convincingly established previously (Hansen, 1919, 1922; Vereshchaka, 2000, 2009; Fukuoka et al., 2005; Vereshchaka et al., 2014).

In *Acetes*, the capitulum of the pars media is armed with true claws (Fig. 10A–D), which are different from the squamose hooks of other sergestids (Fig. 10E–G). In *A. binghami*, *A. americanus*, and *A. petrunkevichi* (Fig. 4B, D, F), the pars media is divided and the longer branch bears tubular apical claws with serrated tips (Fig. 4D, F). These apical claws may represent a transitional state between the entire claws of *Acetes* and

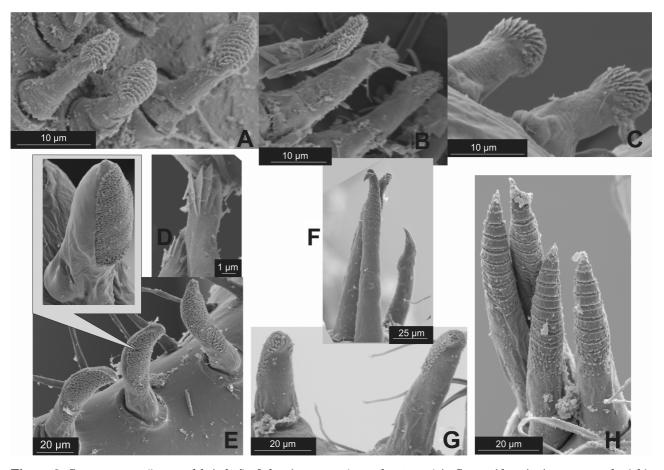


Figure 2. Sensory setae ('serrated bristles') of clasping organ (second antenna) in Sergestidae: A, Acetes petrunkevichi; B, Acetes intermedius; C, Acetes marinus; D, Acetes americanus; E, Petalidium foliaceum; F, Robustosergia robusta; G, Challengerosergia challenger; H, Lucensosergia lucens.

*Peisos* and the squamose hooks of other sergestids. We assume that the branches of the petasma in *A. binghami*, *A. americanus*, and *A. petrunkevichi* are nonhomologous to the lobi and processi of other sergestids. Only the processus ventralis, when present, can be recognized with certainty, owing to its characteristic position.

The pars astringens and pars externa in *Sicyonella* and *Acetes* are certainly homologous to those in other sergestid genera. In *Acetes*, the pars astringens is extraordinarily variable in morphology: It may be large, with a hook on the processus uncifer (Fig. 4B), large without a hook (Figs 5D, 6B, 7B, D, 8D, F), vestigial (Figs 5B, 7F), or absent (Figs 4D, F, 5F, 6D, F).

### SUPPORTED CLADES

Each of analyses with *Ar. foliacea* (Analysis 1), *G. parvus* (Analysis 2), and *Pen. monodon* (Analysis 3) as outgroups retrieved nine minimal length trees, with 231, 237, and 232 steps, respectively. The topology of all three

trees was identical; all supported clades are shown in Figure 11. The clade Sicyonella is sister to the rest of Sergestidae and received high Bremer support (4–5). Within the clade Sicyonella, a terminal clade Sicyonella antennata + Sicyonella received strong statistical support (8 Bremer and 98 bootstrap support). Within the rest of Sergestidae, Acetes is sister to the remaining genera, and received extraordinarily high Bremer (13) and bootstrap (100) support. Within the clade Acetes, the terminal clade Acetes vulgaris + Acetes sibogae + Acetes intermedius + Acetes erythraeus is supported statistically (4 Bremer and 79–80 bootstrap support).

# THE MONOPHYLY OF SICYONELLA AND ACETES AND THE STATUS OF PEISOS

Analyses 1–3, each with a different outgroup, revealed statistical support for the clade *Sicyonella*, thus suggesting its monophyletic origin. The genus is supported by the following synapomorphies, which are common for analyses 1–3 (Fig. 12): maxillula in adults

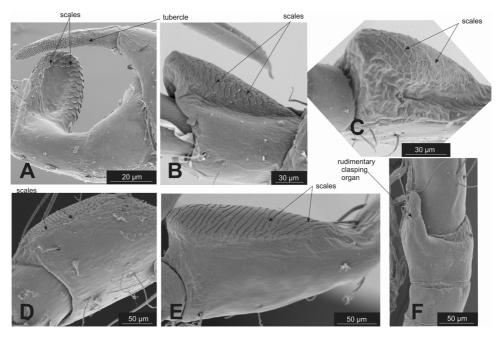


Figure 3. Tubercle and scales of the clasping organ in Sergestidae: A, Acetes marinus; B, Challengerisergia challenger; C, Lucensosergia lucens; D, Robustosergia robusta; E, Deosergestes corniculum; F, Challengerosergia talismani.

with two endites (character 29 – see Appendix 2), enlarged third maxilliped (31), and slightly reduced chela of second and third pereopods (43, 50).

Both the number of synapomorphies and the statistical robustness of the clade confirm the status of *Sicyonella* as being both monophyletic and significantly derived, justifying its taxonomic status as a separate genus.

Analyses 1–3 also revealed a very high level of statistical support for the clade Acetes, suggesting that this clade is also monophyletic. The clade Acetes without Acetes petrunkevitchi (= former Peisos petrunkevitchi) never gained robust support. We therefore combine all species of Acetes and the former Peisos petrunkevitchi into a single genus and consider Peisos as a junior synonym of Acetes. The genus is supported by the following synapomorphies, which are common for analyses 1–3: arthrobranchs on somites IX–XII absent (10, 12, 14, 15, 18), very elongated third antennular segment (24), single endite on the maxillula in adults present (29), endopod on the first maxilliped reduced or absent (30), much reduced chela on first pereopod (40), fifth pereopod in males much reduced (70), two clasping tubercles (79–80), and presence of a strong distal tooth on the fourth segment of the clasping organ (89).

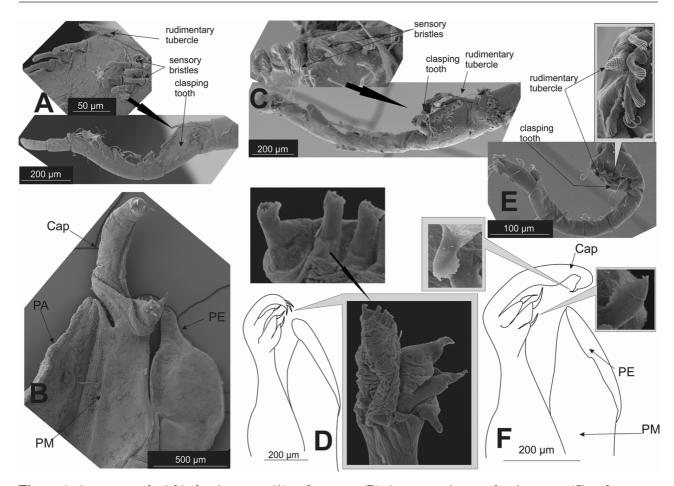
#### TAXONOMY OF SICYONELLA AND ACETES

A key to the genera of Sergestidae is given in Vereshchaka *et al.* (2014). The results of the phylogenetic

analyses provided here indicate the need for an emended diagnosis of the genera *Sicyonella* and *Acetes*.

# SICYONELLA BORRADAILE, 1910

Diagnosis: Carapace and abdomen smooth, firm; labrum not much separated from antennae and eyes; rostrum acute with oblique frontal margin and two dorsal teeth; supraorbital, pterygostomial, and hepatic teeth present; sixth abdominal somite and telson in males without ventral processes; telson with four pairs of lateral spines; eyestalks not elongated, cornea well pigmented; third antennular segment without ventral processes, shorter than first segment; stylocerite present, mobile; mandible with palp; maxillula in adults with two endites; first maxilliped with three-segmented exopod and epipod; second maxilliped with epipod; third maxilliped > two times as long as first pereopod, not dimorphic sexually, dactyl four-segmented; first pereopod with ischium lacking strong movable spines and normally developed chela, fingers subequal; second pereopod with merus lacking proximal protrusion and reduced chela lacking elongated setae, fingers subequal; third pereopod with reduced chela lacking strong curved spines and elongated setae, fingers subequal; pereopods IV-V progressively decreasing in length, flat, seven-segmented, carpi and meri setose only on one margin; uropodal exopod setose at distal part 1/5 of outer margin, with small tooth by far not reaching distal end of exopod. Male clasping organ: well developed, without



**Figure 4.** Acetes petrunkevichi: clasping organ (A) and petasma (B); Acetes americanus: clasping organ (C) and petasma (D); Acetes binghami: clasping organ (E) and petasma (F). Cap, capitulum; PA, pars astringens; PE, pars externa; PM, pars media.

clasping tubercle, bearing two contiguous rows of strong sensory setae or numerous scales. Petasma: pars astringens present, pars media developed, armed with pincers; processus uncifer developed, laminar, lacking hook. Photophores: dermal organs and organ of Pesta absent. Branchiae: podobranch on somite VIII, dendritic anterior arthrobranchs on somites IX–XIII, rudimentary posterior arthrobranchs on somites IX–XIII, and dendritic posterior arthrobranchs on somite XIII.

Type species: By monotypy, Sicyonella maldivensis Borradaile, 1910.

Species: Sicyonella antennata Hansen, 1919, Sicyonella inermis (Paul'son, 1875), Sicyonella maldivensis Borradaile, 1910.

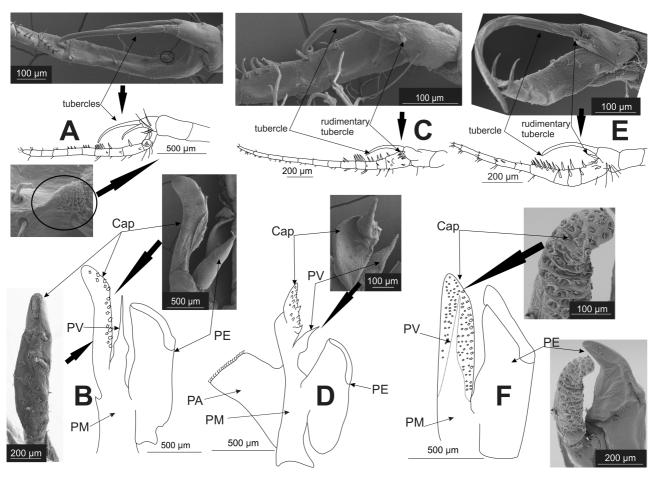
Remarks: Paul'son (1875) described a new species Aphareus inermis from the Red Sea in Russian and referred it to the family Penaeidae. Later Borradaile (1910) established a new genus Sicyonella for a new species, S. maldivensis, and, being ignorant of the

Russian paper by Paul'son, was unaware that Sicyonella was identical to Aphareus Paul'son, 1875. Borradaile (1910) established a new subfamily Sicyoninae of the family Penaeidae. Calman (1913) discovered that the generic name Aphareus had been preoccupied by Cuvier (1830) for a fish and so proposed a new name Aphareocaris, also being unaware that this genus was identical to Sicyonella Borradaile, 1910. Hansen (1919) synonymized Aphareocaris with Sicyonella and described the latest valid species, S. antennata.

The excellent redescription of all valid species of *Sicyonella* by Fukuoka *et al.* (2005) with the use of SEM makes morphological comments redundant. Here we present a key to species as one was not provided in Fukuoka *et al.* (2005). Citations to illustrations that supplement the key are also given.

#### ACETES H. MILNE EDWARDS, 1830

*Diagnosis:* Carapace and abdomen smooth, firm; labrum not much separated from antennae and eyes; rostrum acute, with oblique frontal margin; supraorbital and

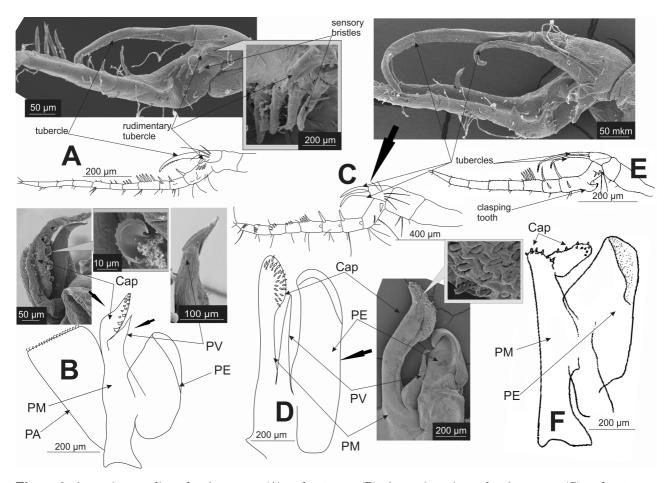


**Figure 5.** Acetes chinensis: clasping organ (A) and petasma (B); Acetes erythraeus: clasping organ (C) and petasma (D); Acetes indicus: clasping organ (E) and petasma (F). Cap, capitulum; PA, pars astringens; PE, pars externa; PM, pars media; PV, processus ventralis.

hepatic teeth present; sixth abdominal somite and telson in males without ventral processes; telson without lateral spines; eyestalks not elongated, cornea well pigmented; third antennular segment without ventral processes, longer than first segment; stylocerite present, mobile; mandible with palp; maxillula in adults with a single endite; first maxilliped with endopod and epipod; second maxilliped with epipod; third maxilliped < two times as long as first pereopod, not dimorphic sexually, dactyl subdivided; first pereopod with ischium lacking strong movable spines and reduced chela, fingers subequal; second pereopod with merus lacking proximal protrusion and much reduced chela lacking elongated setae, fingers subequal; third pereopod with much reduced chela lacking strong curved spines and elongated setae, fingers subequal; fourth pereopod in female absent or five-segmented, in male absent or threesegmented; fifth pereopod absent or three-segmented in female, absent or one-segmented (only coxa present) in male; uropodal exopod setose for distal outer margin, with small tooth by far not reaching distal end of exopod. Male clasping organ: well developed, with two developed or rudimentary clasping tubercles, no sensory bristles adjacent to clasping tubercle. Petasma: pars astringens absent or present, pars media if present armed with strong claws; pars externa developed. Photophores: dermal organs and organ of Pesta absent. Branchiae: podobranch on somite VIII, anterior arthrobranchs on somites VIII–XIII, posterior arthrobranchs absent.

Type species: By original designation, Acetes indicus H. Milne Edwards, 1830.

Species: Acetes americanus Ortmann, 1893, Acetes binghami Burkenroad, 1934, Acetes chinensis Hansen, 1919, Acetes erythraeus Nobili, 1905, Acetes indicus H. Milne Edwards, 1830, Acetes intermedius Omori, 1975, Acetes japonicus Kishinouye, 1905, Acetes johni Nataraj, 1949, Acetes marinus Omori, 1975, Acetes natalensis Barnard, 1955, Acetes paraguayensis Hansen, 1919, Acetes petrunkevitchi (Burkenroad, 1945), Acetes serrulatus (Krøyer, 1859), Acetes sibogae Hansen, 1919, Acetes vulgaris Hansen, 1919.



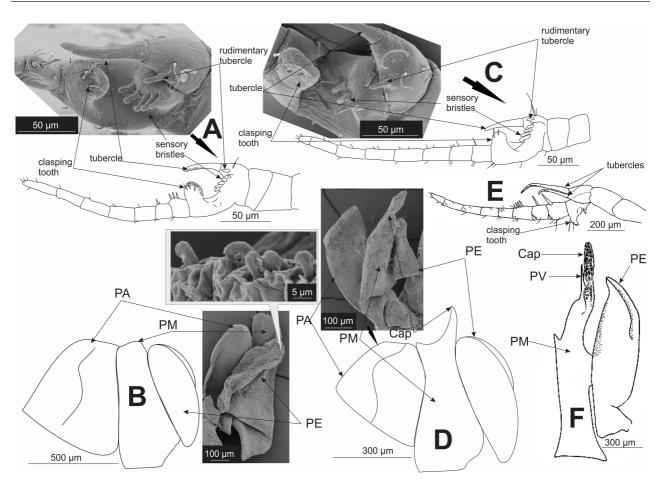
**Figure 6.** Acetes intermedius: clasping organ (A) and petasma (B); Acetes japonicus: clasping organ (C) and petasma (D); Acetes johni: clasping organ (E) and petasma (F). E and F were modified from Omori (1975). Cap, capitulum; PA, pars astringens; PE, pars externa; PM, pars media; PV, processus ventralis.

Remarks: The genus Acetes was established by H. Milne Edwards (1830) who described a new species, A. indicus, from the Ganges estuary. A comprehensive study by Omori (1975) of all valid species of Acetes included diagnoses and figures of all known species of the genus. This paper disentangled most of the taxonomic questions within the genus and even made it possible to identify all females, which is unique for Sergestoidea as usually only the adult males can be identified with certainty. Our SEM data on the ultrastructure of the petasma have provided hitherto-unknown morphological details and formed a basis for establishing homologies to other taxa, which is a necessity for phylogenetic studies. In the earliest available identification keys to species of Acetes, males and females were treated separately (e.g. Hansen, 1919), a tradition continued by Omori (1975). Below we provide a combined key for both males and females with citations for the illustrations necessary for reliable identification.

# MORPHOLOGICAL TRENDS IN SICYONELLA AND ACETES

Morphological analyses show that Sicyonella differs most clearly from the rest of Sergestidae in characters that may be associated with its occurrence at the benthopelagic shelf. These characteristics may provide additional protection, manoeuvrability, and feeding opportunities near the water–bottom interface (Vereshchaka, 1990, 1995). Some of these characters are synapomorphic (Fig. 12): two endites at the maxillulae (character 29 – see Appendix 2), enlarged third maxillipeds (31), and well-developed chelae of second and third pereopods (43, 50). Other characters (two dorsal teeth on the rostrum, pterygostomial tooth, four moveable lateral spines on the telson, a complete set of segments in fourth and fifth pereopods) may have been inherited from a common penaeid ancestor.

Most other characters within *Sicyonella* are associated with mating (male petasma and coupling struc-



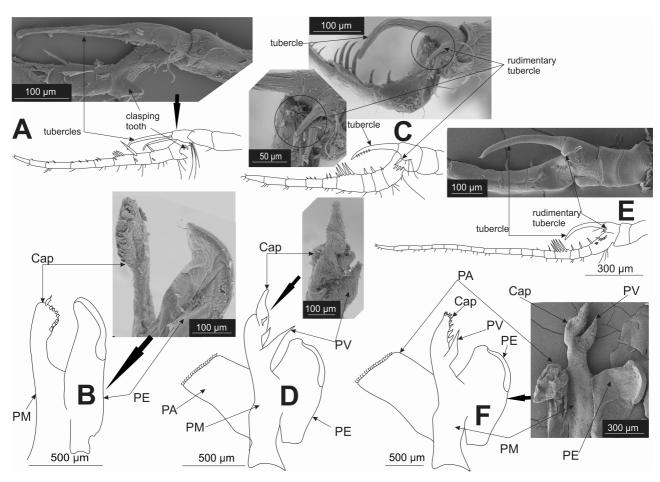
**Figure 7.** Acetes marinus: clasping organ (A) and petasma (B); Acetes paraguayensis: clasping organ (C) and petasma (D); Acetes natalensis: clasping organ (E) and petasma (F). E and F were modified from Omori (1975). Cap, capitulum; PA, pars astringens; PE, pars externa; PM, pars media; PV, processus ventralis.

tures). Hence, the species S. inermis is characterized by the presence of fine scales opposite the tubercle in the clasping organ (87) and by the presence of squamose hooks (100) the entire capitulum of the petasma (Fig. 12). In addition, another clade, S. antennata + S. maldivensis, is supported by a set of synapomorphies related to mating and coupling (Fig. 12): claw-like setae at the clasping organs opposite the tubercle (83), a divided capitulum of the petasma armed with squamose hooks and pincers (101, 104), well-developed lobi armatus, connectens, and terminalis (104, 107, 108-109), well-developed, elongated, twice-branched processus ventralis (110,111, 113, 114, 117, 118). The claw-like setae of the clasping organs may have a holding function. The petasmas of S. antennata and S. maldivensis are probably the most elaborate within Dendrobranchiata, with all known processi and lobi and the branched procesus ventralis being present at the same time.

The clade *Acetes* is supported by synapomorphies related to the reduction of the branches (10, 12, 14, 15, 18), of the mouthparts (29–30), of the first chelae (40), and of the pereopods (adapted for swimming) (70).

These characters, along with the general reduction in size and body compression, are probably adaptations to estuarine and freshwater shallow habitats, and to the fast reproductive cycles observed in the genus (Omori, 1975). These habitats have permanently high concentrations of oxygen, provide no possibility for extensive vertical migrations (too shallow), are productive, and in general favour short life cycles; some species have two generations in a season (Omori, 1975). Yet another set of synapomorphies is related to the presumed coupling procedure: the elongated third antennular segment (24), the presence of two clasping tubercles (79–80), and the strong distal tooth in the clasping organ (89).

All species of *Acetes*, except *A. petrunkevitchi*, show further reduction and loss of structures related to movement (fourth and fifth pereopods – Fig. 12) that may be further adaptations to shallow habitats where significant vertical migrations are impossible. Within *Acetes*, the phylogeny is based only on sexual characters, e.g. the clasping organ and the petasma (Fig. 12).



**Figure 8.** Acetes serrulatus: clasping organ (A) and petasma (B); Acetes sibogae: clasping organ (C) and petasma (D); Acetes vulgaris: clasping organ (E) and petasma (F). Cap, capitulum; PA, pars astringens; PE, pars externa; PM, pars media; PV, processus ventralis.

Omori (1975: fig. 3) manually depictured six clades of Acetes: (1) A. marinus + A. paraguayensis, (2) A. erythraeus + A. intermedius + A. sibogae + A. vulgaris, (3) A. indicus, (4) A. serrulatus + A. johni + natalensis, (5) A. japonicus + A. chinensis, and (6) A. americanus + A. binghami

However, our analysis supports only three of Omori's clades (Fig. 12):

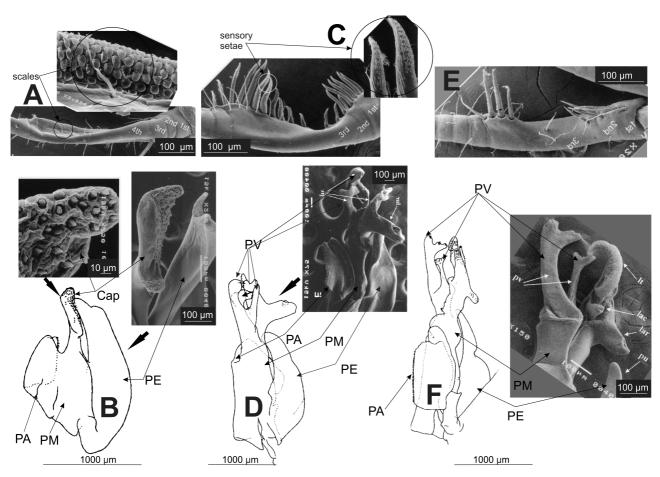
- 1. The clade *A. marinus* + *A. paraguayensis*, which is not statistically robust but supported by the presence of fine scales on the clasping organ opposite the tubercle (87) and by the absence/strong reduction of the capitulum of the petasma (94).
- 2. The clade *A. americanus* + *A. binghami*, which is also not statistically robust but supported by the presence of specialized serrated setae with longitudinal ribs in the clasping organ (86).
- 3. The terminal clade A. erythraeus + A. intermedius + A. sibogae + A. vulgaris, which is statistically robust (Fig. 11) and supported by the following synapomorphies of the petasma: the presence

of well-developed pars astringens (90, 92), additional enlarged claws on the capitulum (98), and an elongated processus ventralis (115, 117).

Most of the morphological variation within *Acetes* is seen in sexual characters, but because of substantial homoplasy, the phylogeny within the genus is hard to resolve. Like *Sicyonella*, *Acetes* is supported by synapomorphies that may be regarded as adaptations to estuarine habitats. Once adapted to such a habitat, the subsequent speciation within both genera has seemingly only been related to mating mechanisms (clasping and copulation).

# GEOGRAPHICAL REMARKS ON THE DISTRIBUTION OF Acetes

The distributions of all recognized species of *Acetes* are summarized in Figure 13. The clades *A. marinus* + *A. paraguayensis* and *A. americanus* + *A. binghami* are geographically isolated from the rest of *Acetes* and occur in Central, South and North America. These clades

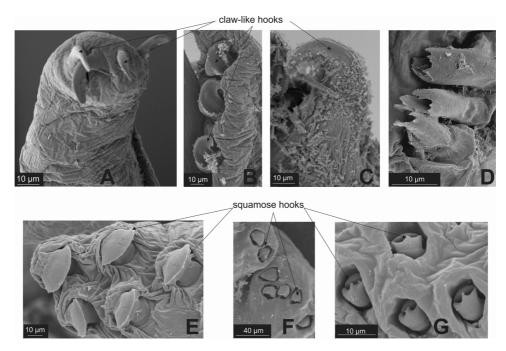


**Figure 9.** Sicyonella inermis: clasping organ (A) and petasma (B); Sicyonella maldivensis: clasping organ (C) and petasma (D); Sicyonella antennata: clasping organ (E) and petasma (F). All photos were modified from Fukuoka et al. (2005). Cap, capitulum; PA, pars astringens; PE, pars externa; PM, pars media; PV, processus ventralis.

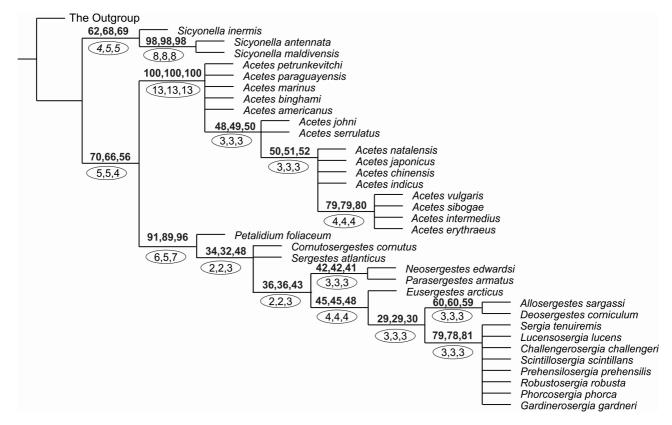
originated through allopatric divergence, whereby new species arise from geographically isolated populations of the same ancestral species. One of these clades, A. marinus + A. paraguayensis, has adapted to low-salinity environments in South America and speciation within this clade reflects the degree of this adaptation: A. marinus lives in brackish waters, whereas A. paraguayensis is the only freshwater sergestid (Hansen, 1919; Omori, 1975). The second American clade, A. americanus + A. binghami, occurs in the coastal waters of North, Central, and South America and shows a parapatric distribution. The two species are geographically isolated from each other by the Isthmus of Panama, which prevents gene exchange between them.

The terminal clade of *Acetes* occurs in the Indo-West Pacific. Speciation within this clade took place allopatrically for *A. johni* and *A. erythraeus*, which are geographically isolated from the other species of *Acetes* and inhabit coastal waters of the Western Indian Ocean. The rest of the species occur along a coastal line between West India and Japan and have undergone sympatric speciation. We assume that this sympatric speciation is a likely outcome of competition for resources, when the evolution of assortative mating leads to reproductive isolation between ecologically diverging subpopulations (Dieckmann & Doebeli, 1999). Cross-breeding between sympatric species of *Acetes* is prevented by highly specialized copulatory structures, which differ greatly even between closely related species.

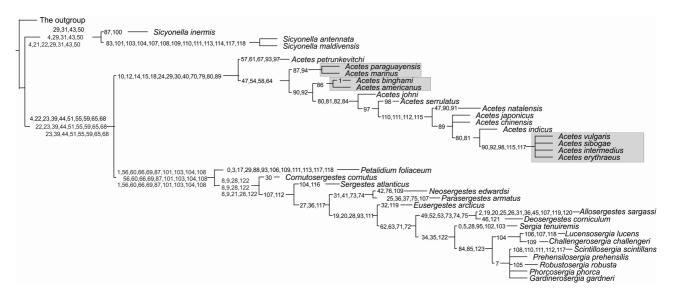
Figure 13 shows a number of blank areas from where *Acetes* has not yet been reported. This is probably because of a lack of sampling as the genus occurs in all well-explored coastal areas of tropical and subtropical areas. This is especially true in regard to the west coast of Africa, which contains the estuaries of big rivers such as Congo, Ebola, Gambia, Niger, Orange, and Senegal. Estuaries of these rivers would definitely be expected to harbour various species of *Acetes*.



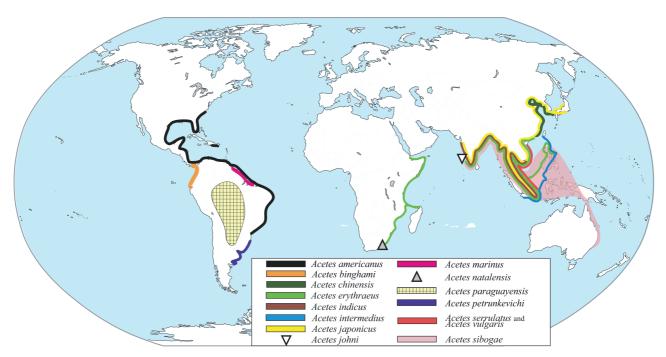
**Figure 10.** Armature of the capitulum of petasma in Sergestidae: A, Acetes petrunkevichi; B, Acetes intermedius; C, Acetes sibogae; D, Acetes americanus; E, Petalidium foliaceum; F, Robustosergia robusta; G, Lucensosergia lucens.



**Figure 11.** Supported clades of Sergestidae using three different outgroups: *Aristeomorpha foliacea* (Risso, 1827), *Gennadas parvus* Bate, 1881, and *Penaeus monodon* Fabricius, 1798. Tree topologies were identical for each analysis and so only one tree is shown. Bootstrap values for each of three analyses are separated by commas and presented above the nodes; Bremer support values for each of three analyses are separated by commas and presented below the nodes in ovals.



**Figure 12.** Strict consensus trees of Sergestidae with various outgroups: *Aristeomorpha foliacea* (Risso, 1827), *Gennadas parvus* Bate, 1881, and *Penaeus monodon* Fabricius, 1798. Tree topologies were identical for each analysis and so only one tree is shown; the nodes are marked with synapomorphies (see character list in Appendix 2). For the four cases in which synapomorphies differed amongst the analyses, the character numbers are shown in three rows, each representing one analysis. Omori's (1975) clades are marked in grey.



**Figure 13.** Provisional geographical distribution of *Acetes*. Coloured lines indicate position of estuarine habitats for each of the known species. Semitransparent coloured areas indicate species ranges of freshwater *Acetes paraguayensis* and estuarine *Acetes sibogae*.

	Key to species of the genus Sicyonella
1.	Clasping organ with fine scales, without robust setae; pars media of petasma entire
-	Clasping organ with robust setae (Fig. 9C, E), fine scales absent; pars media of petasma divided (Fig. 9D, F)
2.	Clasping organ with gap between contiguous rows of strong setae < 1/8 as long as third segment of outer antennular
	flagellum; lobus armatus of petasma nearly as large as the part including lobus inermis, lobus connectens, and
	lobus terminalis
-	Clasping organ with gap between contiguous rows of strong setae nearly half as long as third segment of outer
	antennular flagellum; lobus armatus of petasma half as large as the part including lobus inermis, lobus connectens,
	and lobus terminalis

Key to species of the genus <i>Acetes</i>
1. Rostrum without dorsal teeth
- Rostrum with one to two dorsal teeth
2. Rostrum with a single dorsal tooth
- Rostrum with two dorsal teeth
3. Fourth pereopod (three or five segments) present (but reduced)
- Fourth pereopod absent
4. No tooth on distal inner margin of coxa of third pereopod in female; pars astringens of petasma vestigial
- Tooth present on distal inner margin of coxa of third pereopod in female; pars astringens of petasma absent (Fig. 5E
or well developed (Fig. 5D)
5. Apex of telson triangular. In male, clasping organ with a single developed tubercle opposed to scattered pairs of
strong, specialized setae, which are positioned without large prominence (Fig. 5E)
- Apex of telson rounded or truncated. In male, clasping organ with two developed tubercles (Fig. 5B) or with on
tubercle opposed to a large prominence which is lacking scattered pairs of strong, specialized setae (Fig. 7A)
1
6. In female, basis of third pereopod with acute projection on inner margin. In male, pars astringens absent, no en
larged claws in addition to ordinary claws
- In female, no acute projection on inner margin of third pereopod. In male, pars astringens present, enlarged claw
in addition to ordinary claws present (Fig. 6B)
7. Bases of first pleopods with strong, curved tooth between them
- No strong, curved tooth between bases of first pleopods
8. First segment of antennular peduncle no longer than second and third segments combined. In female, basis of
third pereopod with blunt projection on distal inner margin. In male, three to five enlarged claws along oute
margin
- First segment of antennular peduncle longer than second and third segments combined. In female, no projection
on distal inner margin of basis of third pereopod. In male, one to two enlarged claws along outer margin
9. In female, basis of third pereopod with small projection on distal inner margin. In male, one to two enlarged claw
along outer margin
- In female, no projection on distal inner margin of basis of third pereopod. In male, three enlarged claws alon
outer margin
10. In female, third thoracic segment with a pair of barbs (visible in lateral view) just behind posterior teeth of coxa
In male, clasping organ with a single developed tubercle (Fig. 7A); petasma with pars astringens (Fig. 7A)
- In female, no barb (or not visible in lateral view) just behind posterior teeth of coxa on third thoracic segment. I
male, clasping organ with two developed tubercles (Fig. 5A); petasma without pars astringens (Fig. 5B)1
11. In female, coxa of third pereopod with small blunt tooth. In male, petasma without capitulum
- In female, coxa of third pereopod with large acute tooth. In male, petasma with rudimentary capitulum

# **ACKNOWLEDGEMENTS**

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# APPENDIX 1

TERMINAL TAXA (IN ALPHABETICAL ORDER) AND SOURCES OF CHARACTER SCORING

BMNH, British Museum (Natural History); MNHN, Museum National d'Histoire Naturelle, Paris; SMNH, Naturhistoriska Riksmuseet, Sweden; USNM, United States National Museum; YPM, Yale Peabody Museum; ZMUC, Zoological Museum, University of Copenhagen.

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	C CRU-04533 (syntypes)
	C CRU-06077 (syntypes)
	C CRU-05590 (holotype, dissected), ZMUC CRU-04528
2 2	C CRU-03726
* '	C-CRU-04419
	C CRU-04425
	C CRU-04526
	C CRU-5626 (postlarva), ZMUC CRU-04507
	H Type 2338 (holotype)
· · · · · · · · · · · · · · · · · · ·	C-CRU-004445
	1888.22, 1888.22, 1903.6.6.16 (syntypes, damaged), UC CRU-20546
40 Phorcosergia phorca (Faxon, 1893) ZMU	C CRU-04434
8 1	H 1888.2
	M 7316
	C CRU-03613
	N, NA 331 (syntypes, bad condition), ZMUC CRU-04542
	C CRU-08362
	C ALV-1
	M 1026370
	C CRU-04443

# APPENDIX 2

# LIST OF CHARACTERS USED

New characters are marked with asterisks (\*)

Character no.	Character state	State no.	Reference to figure and source
Body			
0	Integument firm	0	
	Integument membranous	1	
1	Rostrum bears 2 or more dorsal teeth behind the orbital margin	0	
_	Rostrum bears 0–1 dorsal teeth behind the orbital margin	1	
2	Frontal margin of rostrum oblique	0	2B, D – Vereshchaka <i>et al.</i> (2014)
	Frontal margin of rostrum vertical	1	2A – Vereshchaka et al. (2014)
3	Supraorbital tooth absent Supraorbital tooth present	0 1	2A-C - Vereshchaka et al. (2014)
4	Pterygostomial tooth absent	0	2D – Vereshchaka <i>et al.</i> (2014)
4	Pterygostomial tooth present	1	
5	Hepatic protrusion prominent	0	
o .	Hepatic protrusion inconspicuous	1	
6	Hepatic spine absent	0	
· ·	Hepatic spine present	1	2D – Vereshchaka et al. (2014)
7	Hepatic barb absent	0	, , , , , , , , , , , , , , , , , ,
	Hepatic barb present	1	2A-C – Vereshchaka et al. (2014)
8	Somite VIII, arthrobranch developed	0	
	Somite VIII, arthrobranch rudimentary or absent	1	
9	Somite VIII, podobranch absent	0	
	Somite VIII, podobranch present	1	
10	Somite IX, posterior arthrobranch present	0	
	Somite IX, posterior arthrobranch absent	1	
11	Somite IX, posterior arthrobranch dendritic	0	
	Somite IX, posterior arthrobranch lamellar	1	
12	Somite X, posterior arthrobranch present	0	
	Somite X, posterior arthrobranch absent	1	
13	Somite X, posterior arthrobranch dendritic	0	
	Somite X, posterior arthrobranch lamellar	1	
14	Somite XI, anterior arthrobranch present	0	
	Somite XI, anterior arthrobranch absent	1	
15	Somite XI, posterior arthrobranch developed	0	
10	Somite XI, posterior arthrobranch reduced	1	
16	Somite XI, posterior arthrobranch dendritic	0 1	
17	Somite XI, posterior arthrobranch lamellar Somite XII, posterior arthrobranch developed	0	
11	Somite XII, posterior arthrobranch reduced	1	
18	Somite XII, posterior arthrobranch present	0	
10	Somite XII, posterior arthrobranch absent	1	
19	Somite XII, posterior arthrobranch developed	0	
	Somite XII, posterior arthrobranch reduced	1	
20	Somite XII, posterior arthrobranch dendritic	0	
	Somite XII, posterior arthrobranch lamellar	1	
21	Somite XIII, posterior arthrobranch present	0	
	Somite XIII, posterior arthrobranch absent	1	
22	Telson, movable lateral spines present	0	
	Telson, movable lateral spines absent	1	
Appendages			
23	Antennule, very elongated first segment (by half or more than	0	2E – Vereshchaka et al. (2014)
	third segment) present		
	Antennule, very elongated first segment (by half or more than	1	
0.4	third segment) absent	0	OF V
24	Antennule, very elongated third segment (by half or more than first segment) present	0	2F – Vereshchaka <i>et al.</i> (2014)
	Antennule, very elongated third segment (by half or more than	1	
	first segment) absent	1	
25	Stylocerite absent	0	
· <del></del>	Stylocerite present	1	
26	Fixed stylocerite absent	0	
-	Fixed stylocerite present	1	
27	Mobile stylocerite absent	0	
	Mobile stylocerite present	1	
28	Distal tooth of scaphocerite not reaching distal end of blade	0	
	Distal tooth of scaphocerite reaching distal end of blade	1	7G – Vereshchaka $et\ al.\ (2014)$
	Distal tooth of scaphocerite overreaching distal end of blade	2	7E – Vereshchaka et al. (2014)

Character no.	Character state	State no.	Reference to figure and source
29	Maxillula in adults with 4 endites	0	
	Maxillula in adults with 3 endites	1	
	Maxillula in adults with 2 endites	2	
	Maxillula in adults with a single endite	3	
30	First maxilliped with 3-4-segmented endopod	0	
	First maxilliped with 2 segmented endopod	1	
24	First maxilliped with endopod rudimentary or absent	2	OT 11 1 1 1 1 (001 ()
31	Third maxilliped moderately developed, < 2.0 times as long as first pereopod	0	2E – Vereshchaka et al. (2014)
20	Third maxilliped enlarged, > 2.0 times as long as first pereopod	1	2F – Vereshchaka et al. (2014)
32	Third maxilliped not sexually dimorphic, dactyl not modified	0	3D – Vereshchaka et al. (2014)
33	Third maxilliped sexually dimorphic, dactyl modified in males	1 0	$3E$ – Vereshchaka $et\ al.\ (2014)$
99	Third maxilliped with entire dactyl Third maxilliped with dactyl subdivided	1	
34	Third maximped with dactyl subdivided Third maxilliped, dactyl subdivided into ordinary subsegments	0	
94	absent		OE W
0.5	Third maxilliped, dactyl subdivided into ordinary subsegments present	1	2E – Vereshchaka <i>et al.</i> (2014)
35	Third, with dactyl subdivided into specialized subsegments absent	0	
	Third maxilliped, dactyl subdivided into specialized subsegments present	1	2F – Vereshchaka <i>et al.</i> (2014)
36	Third maxilliped, dactyl consisting of 4 specialized subsegments	0	6 – Vereshchaka (2009)
	Third maxilliped, dactyl consisting of 5 specialized subsegments	1	6 – Vereshchaka (2009)
	Third maxilliped, dactyl consisting of 6 specialized subsegments	2	6 – Vereshchaka (2009)
	Third maxilliped, dactyl consisting of 7 specialized subsegments	3	6 – Vereshchaka (2009)
37	First pereopod, ischium lacking strong movable spines	0	
0.0	First pereopod, ischium bearing strong movable spines	1	3F – Vereshchaka <i>et al.</i> (2014)
38	First pereopod with chela	0	
39	First pereopod without chela First pereopod, normal chela (palm nearly as long as fingers)	1 0	
	absent First pereopod, normal chela (palm nearly as long as fingers)	1	
40	present First pereopod, reduced chela (palm > 10 times as long as	0	
	fingers) absent First pereopod, reduced chela (palm > 10 times as long as fingers) present	1	
41	Second pereopod, ischium lacking strong distally curved tooth	0	
	Second pereopod, ischium bearing strong distally curved tooth	1	3G - Vereshchaka et al. (2014)
42	Second pereopod, merus lacking distal protrusion	0	
	Second pereopod, merus bearing strong distally curved tooth	1	3G - Vereshchaka et al. (2014)
43	Second pereopod, slightly reduced chela (palm twice as long as fingers) absent	0	
	Second pereopod, slightly reduced chela (palm twice as long as fingers) present	1	
44	Second percopod, much reduced chela (palm > 10 times as long as fingers) absent	0	
	Second percopod, much reduced chela (palm > 10 times as long as fingers) present	1	
45	Second percopod, fixed finger in chela rudimentary, shorter than dactyl	0	3H – Vereshchaka et al. (2014)
	Second pereopod, fixed finger developed, as long as dactyl	1	3I – Vereshchaka et al. (2014)
46	Second percopod, the la lacking very long setae overreaching	0	oi – veresiteliaka et at. (2014)
10	setae in tufts	ŭ	
	Second pereopod, chela bearing very long setae overreaching setae in tufts	1	3I – Vereshchaka <i>et al.</i> (2014)
47	Third pereopod, coxa lacking mesial tooth Third pereopod, coxa bearing mesial tooth	0 1	
48	Third percopod in female, rounded basis	0	
10	Third percopod in female, basis with small projection or tooth	1	
49	Third pereopod, propodus lacking strong, curved spines proximal to tufts of setae	0	
	Third percopod, propodus bearing strong, curved spines proximal to tufts of setae	1	3J – Vereshchaka et al. (2014)
50	Third percopod, slightly reduced chela (palm three times as long as fingers) absent	0	
	Third percopod, slightly reduced chela (palm three times as long as fingers) present	1	

Character no.	Character state	State no.	Reference to figure and source
51	Third percopod, much reduced chela (palm > 10 times as long	0	
	as fingers) absent Third percopod, much reduced chela (palm > 10 times as long as fingers) present	1	
52	Third pereopod, fixed finger shorter than dactyl	0	3J – Vereshchaka et al. (2014)
	Third pereopod, fixed finger as long as dactyl	1	3K– Vereshchaka et al. (2014)
53	Third pereopod, chela lacking very long setae overreaching setae in tufts	0	
	Third pereopod, chela bearing very long setae overreaching setae in tufts	1	3J – Vereshchaka <i>et al.</i> (2014)
54	Fourth pereopod present in female	0	
	Fourth percopod absent in female	1	
55	Fourth percent in female not 7-segmented	$0 \\ 1$	
56	Fourth percopod in female 7-segmented Fourth percopod in female not 6-segmented	0	
50	Fourth percopod in female 6-segmented	1	
57	Fourth pereopod in female not 5-segmented	0	
	Fourth pereopod in female 5-segmented	1	
58	Fourth pereopod present in male	0	
	Fourth pereopod absent in male	1	
59	Fourth pereopod in male not 7-segmented	0	
00	Fourth pereopod in male 7-segmented	1	
60	Fourth pereopod in male not 6-segmented Fourth pereopod in male 6-segmented	0 1	
61	Fourth percopod in male o-segmented  Fourth percopod in male not 5-segmented	0	
01	Fourth percopod in male 5-segmented	1	
62	Fourth percopod, no carpus and propodus setose along both margins	0	
	Fourth percopod, carpus and propodus setose along both margins	1	
63	Fourth pereopod, carpus and propodus setose along one margin only	0	
	Fourth pereopod, no carpus and propodus setose along one margin only	1	
64	Fifth percopod present in female	0	
65	Fifth pereopod absent in female Fifth pereopod in female not 7-segmented	1 0	
00	Fifth percopod in female 7-segmented Fifth percopod in female 7-segmented	1	
66	Fifth percopod in female 1-segmented Fifth percopod in female not 6-segmented	0	
	Fifth pereopod in female 6-segmented	1	
67	Fifth pereopod in female not 3-segmented	0	
	Fifth pereopod in female 3-segmented	1	
68	Fifth pereopod in male not 7-segmented	0	
	Fifth pereopod in male 7-segmented	1	
69	Fifth pereopod in male not 6-segmented	0	
=-	Fifth pereopod in male 6-segmented	1	
70	Fifth percopod in male not consisting only of coxa	0 1	
71	Fifth pereopod in male consisting only of coxa Fifth pereopod, carpus and propodus setose along both margins absent	0	
	Fifth percopod, carpus and propodus setose along both margins present	1	
72	Fifth pereopod, carpus and propodus setose along one margin only present	0	
	Fifth pereopod, carpus and propodus setose along one margin only absent	1	
73	Uropodal exopod without outer spine	0	10 – Vereshchaka (2009)
T.4	Uropodal exopod with outer spine	1	10 – Vereshchaka (2009)
74	Uropodal exopod, proximal segment not setose along outer margin Uropodal exopod, proximal segment setose along outer margin	0	10 – Vereshchaka (2009) 10 – Vereshchaka (2009)
75	Uropodal exopod, no proximal segment partly setose along	0	10 – Vereshchaka (2009) 10 – Vereshchaka (2009)
10	outer margin Uropodal exopod, proximal segment partly setose along outer	1	10 – Vereshchaka (2009)
76	margin Uropodal exopod, no proximal segment entirely setose along	0	10 – Vereshchaka (2009)
	outer margin Uropodal exopod, proximal segment entirely setose along uropodal exopod, proximal segment entirely setose along outer	1	10 – Vereshchaka (2009)
	margin	-	(2000)

Character no.	Character state	State no.	Reference to figure and source
Male clasping organ			
77	Clasping tubercle absent/rudimentary	0	4B – Vereshchaka et al. (2014)
	Clasping tubercle present	1	4C–E – Vereshchaka et al. (2014
78*	A single clasping tubercle absent	0	4C–E – Vereshchaka et al. (2014
	A single clasping tubercle present	1	
79*	Two clasping tubercles (may be rudimentary) absent	0	4C–E – Vereshchaka <i>et al.</i> (2014
	Two clasping tubercles (may be rudimentary) present	1	Figure 5A,C – present paper
80*	One rudimentary and one well-developed tubercle absent	0	
	One rudimentary and one well-developed tubercle present	1	Figure 5C – present paper
81*	Two well-developed clasping tubercles absent	0	
o o di	Two well-developed clasping tubercles present	1	Figure 5A – present paper
82*	Claw-like setae positioned in scattered groups opposite the tubercle absent	0	71 70
	Claw-like setae positioned in scattered groups opposite the tubercle present	1	Figure 5C – present paper
83*	Claw-like setae positioned in two contiguous rows opposite the tubercle absent	0	
	Claw-like setae positioned in two contiguous rows opposite the tubercle present	1	Figure 9C – present paper
84*	A set of serrated bristles opposite the tubercle absent	0	
	A set of serrated bristles opposite the tubercle present	1	Figure 4C – present paper
85*	Serrated bristles with reticulate distal part absent	0	
	Serrated bristles with reticulate distal part present	1	Figure 2A–C – present paper
86*	Serrated bristles with longitudinal ribs in distal part absent	0	
	Serrated bristles with longitudinal ribs in distal part present	1	Figure 2D – present paper
87*	Fine scales opposite the tubercle absent	0	
	Fine scales opposite the tubercle present	1	Figure 3A – present paper
88	A row of serrated bristles adjacent to the tubercle absent	0	
	A row of serrated bristles adjacent to the tubercle present	1	
89	Strong distal tooth or projection on the fourth segment absent	0	71 71 7
	Strong distal tooth or projection on the fourth segment present	1	Figure 7A, C – present paper
Petasma		0	F: 5D
90	Pars astringens absent	0	Figure 5B – present paper
01*	Pars astringens present	1	Figure 5D – present paper
91*	Vestigial pars astringens absent	0	E: 7E
92*	Vestigial pars astringens present	1 0	Figure 7F – present paper
92.	Well-developed pars astringens absent	1	Figure 7P D progent namer
93	Well-developed pars astringens present Processus uncifer without hook	0	Figure 7B,D – present paper 5C, 6D – Vereshchaka <i>et al</i> .
30	Processus uncifer with a hook	1	(2014) 5A,B,D – Vereshchaka <i>et al</i> .
			(2014)
94	Capitulum absent/vestigial	0	Figure 7B,D – present paper
0.54	Capitulum present	1	Figure 7F – present paper
95*	No capitulum armed with strong claws	0	E: 104 C
96	Capitulum armed with strong claws	1 0	Figure 10A–C – present paper
90	Divided capitulum with strong claws absent Divided capitulum with strong claws present	1	Figure 4B,D,F – present paper
97	Entire capitulum with strong claws present	0	Figure 4B,D,F – present paper
91	Entire capitulum with strong claws absent  Entire capitulum with strong claws present	1	Figure 9B – present paper
98*	Capitulum, enlarged claws in addition to ordinary claws absent	0	Figure 6D – present paper Figure 6D – present paper
	Capitulum, enlarged claws in addition to ordinary claws present	1	Figures 8D,F, 10C – present paper
99*	Capitulum armed with squamose hooks and pincers absent	0	paper
	Capitulum armed with squamose hooks and pincers present	1	Figure 10E-G - present paper
100*	Entire capitulum armed with squamose hooks and pincers absent	0	
	Entire capitulum armed with squamose hooks and pincers present	1	Figure 9B – present paper
101*	Divided capitulum armed with squamose hooks and pincers absent	0	
	Divided capitulum armed with squamose hooks and pincers present	1	Figure 9C,E – present paper
102	Capitulum with pincers, complete set of undivided lobi or	0	
102	processi absent		

Character no.	Character state	State no.	Reference to figure and source
103	Capitulum with pincers and significantly modified (divided or reduced) lobi/processi absent	0	
	Capitulum with pincers and significantly modified (divided or reduced) lobi/processi present	1	
104	Lobus armatus rudimentary	0	5E – Vereshchaka et al. (2014)
	Lobus armatus developed	1	5A-D – Vereshchaka et al. (2014)
105	Twisted lobus connectens and lobus terminalis absent	0	5A-E - Vereshchaka et al. (2014)
	Twisted lobus connectens and lobus terminalis present	1	6D – Vereshchaka et al. (2014)
106	Rudimentary lobus connectens absent	0	
	Rudimentary lobus connectens present	1	5C – Vereshchaka et al. (2014)
107	Well-developed lobus connectens absent	0	
	Well-developed lobus connectens present	1	5A,D,E – Vereshchaka <i>et al</i> . (2014)
108	Lobus terminalis absent or rudimentary	0	(====/
	Lobus terminalis developed	1	5A-D – Vereshchaka et al. (2014)
109	Divided lobus terminalis absent	0	5A-D – Vereshchaka et al. (2014)
100	Divided lobus terminalis present	1	6A – Vereshchaka et al. (2014)
110	Processus ventralis absent	0	5C – Vereshchaka et al. (2014)
110	Processus ventralis present	1	5A,B,D – Vereshchaka <i>et al.</i> (2014)
111	Processus ventralis rudimentary	0	5E – Vereshchaka <i>et al.</i> (2014)
	Processus ventralis developed	1	5A,B,D – Vereshchaka <i>et al</i> .
112	Entire processus ventralis absent	0	(=011)
	Entire processus ventralis present	1	5A,B,D – Vereshchaka <i>et al</i> . (2014)
113	Divided processus ventralis absent	0	
	Divided processus ventralis present	1	9C,E – present paper
114	Twice-divided processus ventralis absent	0	
	Twice-divided processus ventralis present	1	9C,E – present paper
115	Needle-like processus ventralis absent	0	respectively.
	Needle-like processus ventralis present	1	5F – present paper
116	Triangular processus ventralis absent	0	or prosent paper
110	Triangular processus ventralis present	1	5E – Vereshchaka et al. (2014)
117	Elongate processus ventralis absent	0	01
111	Elongate processus ventralis present	1	5B,D present paper
118	Processus ventralis, hooks and suckers absent	0	5A-E – Vereshchaka <i>et al.</i> (2014)
110	Processus ventralis, hooks and suckers present	1	6B – Vereshchaka et al. (2014)
119	Processus ventralis, simple spines absent	0	5A,C,E – Vereshchaka <i>et al</i> . (2014)
	Processus ventralis, simple spines present	1	5B.D – Vereshchaka et al. (2014)
120	Processus ventralis, stellate spines absent	0	5A,D,E – Vereshchaka <i>et al</i> . (2014)
	Processus ventralis, stellate spines present	1	5B – Vereshchaka <i>et al.</i> (2014)
121	Processus ventralis, apical lashes absent	0	5B-E – Vereshchaka et al. (2014)
	Processus ventralis, apical lashes present	1	5A– Vereshchaka et al. (2014)
Photophores	г		
122	The organ of Pesta absent	0	2F – Vereshchaka et al. (2014)
	The organ of Pesta present	1	2E – Vereshchaka et al. (2014)
123	Dermal photophores absent	0	
-	Dermal photophores present	1	

The character numbers also refer Figs. 12–13 and (in brackets) to the subchapters: THE MONOPHYLY OF SICYONELLA AND ACETES AND THE STATUS OF PEISOS and MORPHOLOGICAL TRENDS IN SICYONELLA AND ACETES.

0 5 10 15 20 25 30 35 40

# APPENDIX 3

#### DATA MATRIX

Missing data indicated by question marks (?); inapplicable data by hyphens (-)

Characters 0-40

Aristaeomorpha foliacea Gennadas parvus Penaeus monodon Acetes americanus Acetes binghami Acetes chinensis Acetes erythraeus Acetes indicus Acetes intermedius Acetes japonicus Acetes johni Acetes natalensis Acetes serrulatus Acetes sibogae Acetes vulgaris Acetes marinus Acetes paraguayensis Acetes petrunkevitchi Sicyonella inermis Sicyonella maldivensis Sicyonella antennata Petalidium foliaceum Sergia tenuiremis Gardinerosergia gardneri Phorcosergia phorca Robustosergia robusta Prehensilosergia prehensilis Scintillosergia scintillans Challengerosergia challengeri Lucensosergia lucens Deosergestes corniculum

Eusergestes arcticus Sergestes atlanticus

Cornutosergestes cornutus

Allosergestes sargassi

Parasergestes armatus

Neosergestes edwardsi

00-01010000000000000001000000000000--0010 1100010001000000000000100001000000--0010 00-000100100000000001110000000000--0010 00010010101-1-11-01--11011010320000--0001 01010010101-1-11-01--110110103200000-0001 00010010101-1-11-01--110110103200000-0001 00010010101-1-11-01--110110103200000-0001 00010010101-1-11-01--110110103200000-0001 00010010101-1-11-01--110110103200000-0001 00010010101-1-11-01--110110103200000-0001 00010010101-1-11-01--110110103200000-0001 00010010101-1-11-01--110110103200000-0001 00010010101-1-11-01--110110103200000-000100010010101-1-11-01--110110103200000-0001 00010010101-1-11-01--110110103200000-0001 00010010101-1-11-01--110110103200000-0001 00010010101-1-11-01--110110103200000-0001 00010010101-1-11-01--110110103200000-0001 0001101010000000000100010101020101010001000011010100000000001000101010201010100010 00011010100000000001000101010201010100010 110000101001010011011110010101000?00????? 110001000101010010000011011000000110-0100 010000010101010010000011011020000110-0100 010000010101010010000011011000000110-0100 010000010101010010000011011000000110-0100 010000010101010010000011011010000110-010001000001010101010010000011011010000110-0100010000100101010010000011011010000110-0100 010000100101010010000011011010000110-0100 01000010010101001000001001101000010120100 010100100101010010000011011010000110120100 01010010010101001001101001012000010140100 01010010010101001001101001012010010140100 01000010010101001001101000002001010101100 01000010010101001001101001102001010120100

#### APPENDIX 3 Continued

Characters 41-81

Aristaeomorpha foliacea Gennadas parvus Penaeus monodon Acetes americanus Acetes binghami Acetes chinensis Acetes erythraeus Acetes indicus Acetes intermedius Acetes japonicus Acetes johni Acetes natalensis Acetes serrulatus Acetes sibogae Acetes vulgaris Acetes marinus Acetes paraguayensis Acetes petrunkevitchi Sicyonella inermis Sicyonella maldivensis Sicyonella antennata Petalidium foliaceum Sergia tenuiremis Gardinerosergia gardneri Phorcosergia phorca Robustosergia robusta Prehensilosergia prehensilis Scintillosergia scintillans Challengerosergia challengeri Lucensosergia lucens Deosergestes corniculum Eusergestes arcticus Sergestes atlanticus Cornutosergestes cornutus Allosergestes sargassi Parasergestes armatus

Neosergestes edwardsi

41 46 51 56 61 66 71 76 81

#### APPENDIX 3 Continued

Characters 82-123

82 87 92 97 102 107 112 117 122 Aristaeomorpha foliacea Gennadas parvus Penaeus monodon Acetes americanus Acetes binghami Acetes chinensis Acetes erythraeus Acetes indicus Acetes intermedius Acetes japonicus Acetes johni Acetes natalensis Acetes serrulatus Acetes sibogae Acetes vulgaris Acetes marinus Acetes paraguayensis Acetes petrunkevitchi Sicyonella inermis Sicyonella maldivensis Sicyonella antennata Petalidium foliaceum Sergia tenuiremis Gardinerosergia gardneri Phorcosergia phorca Robustosergia robusta Prehensilosergia prehensilis Scintillosergia scintillans Challengerosergia challengeri Lucensosergia lucens Deosergestes corniculum 000001001011100001010110011011100001000110 000001001011100001010110011011100001010010 Eusergestes arcticus Sergestes atlanticus 00000100101010000101010001101010001000010 Cornutosergestes cornutus Allosergestes sargassi Parasergestes armatus Neosergestes edwardsi