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NEW KEYS FOR THE IDENTIFICATION OF INDO-WEST PACIFIC CORAL ASSOCIATED PONTONIINE SHRIMPS, WITH OBSERVATIONS ON THEIR ECOLOGY

(Crustacea : Decapoda: Palaemonidae)

A. J. Bruce

Crustacea Section, Queensland Museum, P.O. Box 3300, South Brisbane, Queensland, 4101, Australia

ABSTRACT

Correct identification is the keystone for investigating the ecology of coral reef animals. New keys for the identification of coral-associated Indo-West Pacific pontoniine shrimps are provided to replace those in Bruce (1972). Information on their geographical distribution and host preferences is up-dated and their ecology discussed. The key includes 48 species of 16 genera of obligatory associates, together with 6 species of two genera of less certain ecology.

Key words: Crustacea, Decapoda, Natantia, Palaemonidae, coral associates, Indo-West Pacific, identification keys, hosts, ecology, distributions.

INTRODUCTION

Pontoniine shrimps are the dominant caridean associates of Indo-West Pacific scleractinian corals. The first reported associations were those of *Oedipus graminea* and *O. superba* (now *Coralliocaris graminea* and *C. superba*) by Dana (1852), but as was usual with earlier workers, the host was not even generically identified. Even a century after Dana, there were few records of the coral hosts of these shrimps. In more recent years the intimate relationship, often obligatory and highly specific, has become increasingly recognized, and the host's identities more often reported.

The obligatory association of caridean shrimps in general, and those of the sub-family Pontoniinae in particular, with scleractinian corals seems to be a phenomenon particularly noteworthy in the Indo-West Pacific region. In the eastern Pacific region, both of the pontoniine genera associated with corals, *Harpiliopsis* (2 species) and *Fennera* (1 species), with *Pocillopora* and *Porites* (Hol-thuis 1951), are probably derived from the Indo-West Pacific by the transportation of planktonic larvae by the Equatorial Counter Current and "island-hopping". In the Atlantic-Caribbean region, they are even less conspicuous and

there appear to be no pontoniine associates of western Atlantic scleractinian corals (P. Castro pers. comm.). Even the large genus *Acropora* appears to be without dedicated pontoniine commensals in that region, in marked contrast to the Indo-West Pacific region, where it has at least 20 species of 6 genera. The absence of *Harpiliopsis* and *Fennera* from the Caribbean suggests that they may have colonised the eastern American seaboard after the development of the Panamanian isthmus.

The information available in 1969 was reviewed in Bruce (1972). General information on the biology of Indo-West Pacific pontoniine and other shrimps can be found in Bruce (1974b, 1976a, 1977b, 1979b). The host corals were reviewed by Bruce (1977a). Further details concerning the taxonomy and distribution of many species are now available but little more is known of their life cycles or trophic relationships with their hosts. Their longevity is particularly obscure. Post-larval specimens may often be found on the host corals but as these are very small and usually nearly transparent, with relatively few chromatophores, they are easily overlooked (Bruce 1985). These have presumably arrived and settled from planktonic larvae. Abbreviated larval development is not known to occur in coral associated pontoniine species. It may be noted that in many pontoniine species, including several from coral hosts, observed under aquarium conditions in Tanzania, Zanzibar), (inclusive of Kenya and the Seychelle Islands, the ova hatched on the night of each spring tide, generally between midnight and 2 am, and a new batch was already in place on the pleopods by the following morning. Field collections made at the time of spring tides throughout the year almost always yielded adult females bearing a full complement of ova, either on the point of hatching or freshly laid. In two weeks these ova were ready to hatch, for the next spring tide, throughout the year. In less equable environments these shrimps may well be expected to show distinct seasonal patterns in their reproductive activities. The first larval stages of the various genera can generally be readily separated on morphological grounds (although the larvae of many genera still remain to be described). Identification of species is much more difficult, although in many cases each species appears to have a distinctive pattern of chromatophores. None have as yet been reared to post-larval stages and host selection by post-larval shrimps remains unstudied.

The identification of pontoniine shrimps from coral hosts presents a number of practical problems, many in common with other carideans. While many are abundant on appropriate hosts, many are also apparently rare, or at least rarely caught by collectors, and consequently inadequately known and often poorly described by modern standards. Thus, *Coralliocaris macrophthalma*, described by Henri Milne Edwards in 1837, from an indefinite locality, has only been reported on two occasions in the subsequent literature. Some specimens of *C. graminea* may show a close resemblance to the descriptions of *C. macrophthalma* but detailed comparison is not possible as the type material of the latter is no longer extant. Many species have characteristic colour patterns but the colouration of Milne Edwards' species was not recorded. Further information on these species would be most useful. It should also be kept in mind that abnormal rostra and chelae, through regeneration after injury, are not uncommon in coral associated shrimps (Bruce 1976b), probably due often to intraspecific conflict, and that species that normally have well developed similar subequal chelae may have small dissimilar unequal chelae, for example in *Coralliocaris*. In other species, such as *Jocaste* spp., the chelae are normally asymmetrical.

In making collections of coral associated shrimps, many of which can be quite active, it is fundamentally important that there is no cross contamination between different hosts. Segregation of intertidal coral hosts presents no major problems. From deeper water, smaller corals can be collected by divers and brought to the surface in clip-seal polythene bags. Large corals can be raised in buckets, preferably lidded, and still larger ones in garbage bins also fitted with lids and with small mesh-covered windows cut into the lower sides to allow the outflow of the contained water without losing the catch. Different coral species should not be mixed in the same container. A few drops of rotenone-type chemicals will then dislodge the shrimps, except for *Paratypton siebenrockii*, the only gall-inhabiting species. These shrimps will survive well if transferred quickly to fresh sea-water. The freshly caught shrimps should be quickly sorted by colour pattern. They can be allowed to die, in small containers, by hyperthermia or hypothermia (in tropical sunlight or in containers of iced water or by refrigeration) before preservation to avoid the excessive autotomy of their appendages. Overcrowding should be avoided as this may also result in unnecessary damage to the shrimps. Short term preservation with formaldehyde solution often allows some temporary preservation of the colour pattern, although the shades of colour may change. Subsequent conservation in 50% ethylene glycol also preserves the colour pattern for a further period but final preservation should be in 70% ethanol. A few undamaged, well preserved specimens are preferable to a larger number of poorly preserved bodies with a mass of detached appendages, especially if a mixture of species is represented. Most useful of all are specimens accompanied an authoritative host identification and a description of the colour pattern, or a photograph of the live shrimp.

The importance of colour pattern in the identification of species has only recently become fully apparent. This has been greatly facilitated by the use of scuba diving techniques and underwater photography, although, unfortunately, many photographers do not collect specimens of their subjects. Some of the species that have been photographed on coral and other hosts can not as yet be referred to any known species. Many species of pontoniine shrimp, particularly the commensal species, have striking species-specific colour patterns that are not conspicuous in the free-living species, which tend to be semi-transparent or cryptically coloured. These patterns are generally consistent over wide areas of geographical distribution, although the colour tones may show some variation.

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Distinctive colour patterns within a morphological "species" may be the first indication of the existence of a group of sibling species or, at least, species with minimal morphological differences. These may also be associated with different host animals. Once preserved in alcohol, this valuable clue is rapidly lost, often for ever. In East Africa, some specimens of "*Periclimenes lutescens*" were noticed to consist of two distinct colour patterns, one associated with *Acropora* and the other with *Stylophora* (Bruce 1972). The former proved eventually to be *P. lutescens* (Dana) s.str., while the other was resurrected as *P. consobrinus* De Man, a very closely related species separable by only small morphological differences, whose author (De Man 1902) had himself originally (De Man 1888) identified his specimens as *P. lutescens*, the hosts and colour patterns of both being then unknown. Holthuis (1952) synonymized the two species, although Kemp (1922) had treated them as distinct, although for inadequate reasons.

The importance of sibling species in marine animals has been reviewed by Knowlton (1986, 1993). It appears probable that the frequency of sibling species in the marine environment has been much underestimated until recently and that in many groups of coral reef animals, including both corals and decapod crustaceans, the species diversity may be underestimated by a factor of three to five times (Knowlton & Jackson 1994). Duffey (1996) has shown that the sponge-associated alpheid shrimp "Synalpheus rathbunae" Coutiére, 1909 consists of three sympatric species, separable by their allozyme genotypes and with minimal host overlap. It is likely that some of the "species" dealt with in this report consist of similar sibling species. This is well exemplified by the Coralliocaris venusta taxa, where shrimps with two characteristic colour patterns of conspicuous white patches live and behave like distinct species (Bruce 1974b), but without any consistent morphological differences having as yet been detected. Both are associated with Acropora hosts, but it is not clear if they associate with different species. On any particular coral host the shrimp population always consists of examples sharing the same colour pattern. The original description of C. venusta, by Kemp (1922), did not provide any indication of the colour pattern of his specimens, which are not available for study, so that it cannot be determined which of the two taxa is C. venusta Kemp s str. However, a third pattern, without conspicuous white patches also occurs (Bruce 1977b) and may represent a further sibling species.

Similarly, atypical colour patterns have been noted in *C. superba* (Dana), which may also represent sibling species, but these are relatively rare, whereas both colour patterns of *C. venusta* seem quite common. Unfortunately, detailed information on the host preferences of these taxa, if any, is not readily available. *Coralliocaris viridis* has recently been separated from the closely related *C. graminea* (Dana) by Bruce (1974a, 1976b). It may be noted that while these two species are very easily and consistently separated by their colour patterns and morphological features in the western Indian Ocean, some specimens are much less easily distinguished in Great Barrier Reef material. Similar situations

may be expected to occur in other species or species groups. The group represented by *Periclimenes holthuisi* and related species, some of which also associate with actinians and scyphozoans, show particularly characteristic colour patterns on the dorsum of the third abdominal segment (Bruce 1990), but the number of colour patterns as reflected in the underwater photographic literature seems to greatly exceed the number of presently recognized taxa. The genus *Hamopontonia* contains at present only two recognized species, one of which, *H. corallicola*, has been reported to associate with actinarians as well as corals in Japanese waters (Suzuki & Hayashi 1977; Nomura 1989). Nomura concludes that "the colour patterns may be affected by the host species". The actinarian associates would be likely candidates for a sibling species. It also seems probable that the various monospecific pontoniine genera that are associated with a diversity of host genera may dissolve into complexes of sibling species.

Although pontoniine shrimps have been reported in association with a wide variety of coral hosts, they are most frequently found on close-branching coral species, such as many Acropora, Pocillopora, Stylophora, etc., or in species in which the tentacles of the polyps are fully expanded by day, such as Galaxea, Euphyllia, etc. Many coral species that appear to be suitable as potential hosts, such as Aorehelia, have not so far provided the expected pontoniine or other associates. Coral hosts in this report are identified only to genus. Most pontoniine shrimps are associated with hosts at a generic level and many of the species level host identifications, particularly the more ancient ones, should be verified in terms of modern scleractinian taxonomy. This may be particularly important in view of the recent recognition of sibling species in some corals such as Stylophora pistillata (Esper 1797) (Gattuso et al. 1991) and indications that Pocillopora damicornis (L., 1758) also not a single species (Knowlton & Jackson 1994). It seems likely that many of the other "well known" common and widespread coral species may also be polyspecific. These two coral genera, Stylophora and Pocillopora, have a long history of being studied in the Indo-West Pacific for their decapod commensals but are referred to at species level in numerous papers only from 1966, when Patton reported the association of several shrimp species with a variety of coral hosts, including species of these two genera. The situation of the numerous Acropora "species" appears even more complex, possibly due to the existance of hybrids as well as "pseudo-species" (Wallace & Willis 1994). There are numerous records of pontoniine shrimps (Anapontonia, Ischnopontonia and Platycaris) from Galaxea fascicularis (L. 1758), some of which should perhaps be referred to the closely related G. alta, until recently overlooked on the Great Barrier Reef (Harrison 1988), and possibly elsewhere also.

The distributions of many coral-associated pontoniine shrimp species are still not well known, due largely to the haphazard nature of collecting activities. It is apparent that some common species are very widely distributed, ranging, for example, from the Red Sea or East Africa to Mexico or Panama (*Harpiliopsis depressa* and *H. spinigera, Fennera chacei*, the latter first described from the eastern

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Pacific region). However, some species do appear to have genuine rarity or very restricted distributions. The recently discovered Yemenicaris trullicauda, found in Yemen in Stylophora and Pocillopora spp, would surely have been found elsewhere in these hosts if it occurred commonly in the western Indian Ocean or Red Sea. Similarly, *Tectopontonia maziwiae* is still known only from the holotype specimen, from Maziwi Island, Tanzania, collected in 1970, although its host is a common Acropora species, reported as A. surculosa (Dana 1846), now A. hyacinthus (Dana 1846), ranging from the Indian Ocean east to Tahiti. However, rarity may be more apparent than real, as is illustrated by the example of the alpheid shrimp Racilius compressus, first described from the Red Sea by Paulson (1875). It was not reported again until 1958, 70 years after its discovery, when a further, but incomplete, specimen was found in Moçambique (Barnard 1958). The species was apparently of great rarity. Once its host, the oculinid coral Galaxea fascicularis, where it is often found with pontoniine shrimps of the genera Anapontonia, Ischnopontonia and Platycaris, had been identified, the shrimp was found to be abundant and widespread throughout most of the Indo-West Pacific region, whereever the host coral occurred.. Some of the "rare" pontoniines may be rare for similar reasons. There are still many genera and species of corals, particularly those from greater depths, that appear potentially suitable as hosts from which no shrimp associates have yet been reported.

Almost all the shrimps included in the following key are obligate associates of scleractinian corals. Other pontoniine shrimps may also often be found in collections of coral associates. These include particularly Periclimenella petitthouarsi and P. spinifera, which are probably free-living browsers and frequently found in coral free intertidal pools. This genus is therefore included in the key, together with the similar genus Exoclimenella. The associations of the latter genus must be considered as tentative only as they are based on very limited information and the various species may be also free-living browsers. Further information on these two genera can be found in Duris & Bruce (1995). Not included are the species of the free-living, micropredatory *Periclimenes grandis* (Stimpson, 1860) species-group or the ubiquitous Palaemonella rotumana (Borradaile, 1898). Keys for the identification of the species of these taxa can be found in Bruce (1987, 1991). Some of the already described but rarer pontoniine species, such as Eupontonia noctalbata Bruce, not as yet known to be of commensal life-styles, may eventually prove to be commensally associated with coral hosts. Some other species may be accidentally associated with corals due to the vagaries of collection. Others that have been attributed to coral hosts, such as some Periclimenaeus species, are probably always associated with encrusting sponges or colonial tunicates attached to the coral. Such species have been omitted from the present key. The key is intended for use with adult specimens, preferably ovigerous females. Further details for most species are to be found in Kemp (1922), Holthuis (1952, 1993), Chace & Bruce (1993). Müller (1993) also gives full details of their coral hosts.

Abbreviations used: R., rostral dentition, dorsal teeth/ventral teeth; pocl, postorbital carapace length. Further details of the species marked with an asterisk in the keys are to be found in the similarly marked items listed in the references.

KEYTO GENERA

1.	First pereiopod chelae with strongly spatulate, pectinate fingers 2
_	First pereipod chelae with simple, non-pectinate fingers
2.	Major second pereiopod with phonogenic fossae on opposing cutting
	edges of fingers Periclimenella
-	Phonogenic fossae absent from second pereiopod fingers Exoclimenella
3.	Mandibles with palp Vir
_	Mandibles without palp 4
4.	Dactyls of ambulatory perciopods with hoof-shaped basal process
_	Dactyls of ambulatory pereiopods without hoof-shaped basal process 6
5.	Hepatic spine present, second pereiopod chelae markedly unequal, dissim-
	ilarJocaste
_	Hepatic spine absent, second pereiopod chelae subequal, similar
6.	Dactyls of ambulatory pereiopods simple
	Dactyls of ambulatory pereiopods twisted laterally, carinate Harpiliopsis
7.	Posterior margin of telson normally armed with three pairs of spines 10
_	Posterior border of telson not normally armed with three pairs of spines 8
8.	Posterior border of telson without marginal spines in adults
_	Posterior telson expanded scoop-shaped, margin with lateral and interme-
	diate spines only
9.	Posterior margin of telson narrow, with paired of curved hook-like process-
	es, rostrum well developed Hamopontonia
_	Posterior margin of telson broadly rounded, unarmed; rostrum obsolete,
	gall inhabiting
10.	Body form depressed 11
_	Body subcylindrical or compressed 13
11.	Body strongly depressed; rostrum short, broadly expanded laterally, un-
	armed
_	Body moderately depressed only, rostrum compressed, dentate
12.	Carapace with hepatic, antennal and post-antennal spines Fennera
	Carapace with hepatic and antennal spines only Tectopontonia
13.	Body subcylindrical; rostrum well developed, dorsally and ventrally den-
	tate; second pereiopods with large subequal similar chelae; caudal fan with-
	out special armament14
-	Body compressed, rostrum feebly developed, ventrally unarmed 15
14.	Hepatic spine presentPericlimenes
-	Hepatic spine absent Philarius

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15.	Body strongly compressed; rostrum dorsally dentate, caudal fan strongly
	armed
_	Body moderately compressed, rostrum unarmed, caudal fan normal 17
16.	Distolateral margin of uropodal exopod strongly multidentate, without ar-
	ticulated spine Anapontonia
	Distolateral margin of uropodal exopod not multidentate, unarmed, with
	strong hamate spine distally Ischnopontonia
17.	Orbital margin non-pectinate Metapontonia
_	Orbital margin strongly pectinate, inhabiting grooves on host's surface

KEYS TO SPECIES

Anapontonia Bruce, 1966

Coralliocaris Stimpson, 1860

- 4. Dactyl of second pereiopod laterally carinate or with proximal swelling... 5
- 5. Fixed finger of second pereiopod with distinct oval fossa on proximal cutting edge; R. 4-5/0...... *C. pavonae* Bruce, 1972

*See addendum

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[Host genus: *Pavona*. Distribution: Type locality: Sigatoka, Viti Levu, Fijian Islands. Also known from Taiwan].

- 7. Colour pattern with large white patches C. venusta a

lands, Kiribati and Society.Islands].

 Proximal segment of antennular peduncle wider than long, with poorly developed distolateral lobe bearing obsolescent distolateral tooth; rostrum reaching only to middle of proximal peduncular segment; R. 0/0.....

Ctenopontonia Bruce, 1979

Exoclimenella Bruce, 1994

- 1. Supraorbital spines present; R. 7/4...... *E. sudanensis* Duris & Bruce, 1995 [Host genus: *Stylophora*. **Distribution:** Type locality: Port Sudan, Sudan. Known only from the type material].

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_	Carpus of second pereiopod with three distal teeth
3.	Merus of second pereiopods with several distoventral spines; mandibular
	palp present; R. 1+6-7/3 <i>E. denticulata</i> (Nobili, 1906)
	[Host genus: Acropora. Distribution: Type locality: Gatavake, Mangareva Atoll, Gambier Is-
	lands. Also known from Great Barrier Reef, South China Sea, Marshall Islands and Tuamotu
	Islands].
-	Merus of second perciopod unarmed; mandibular palp absent; R. 6/2
	<i>E. sibogae</i> (Holthuis, 1952)
	[Host genera: ? Distribution: Type locality: Banda, Indonesia, 9-36m. Known from type speci- men only].

Fennera Holthuis, 1951

Hamopontonia Bruce, 1970

Harpiliopsis Borradaile, 1915

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Island, Palmyra Island, Johnson Atoll and Hawaiian Islands. Also Galapagos Islands, Mexico, Costa Rica, Panama, Colombia and Ecuador].

Ischnopontonia Bruce, 1966

Jocaste Holthuis, 1952

Metapontonia Bruce, 1967

Paratypton Balss, 1914

One species only..... P. siebenrocki Balss, 1914

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[Host genus: *Acropora*. Distribution: Type localities: Koseir; Mersa Sheik, Egypt; Jaluit, Marshall Islands; Samoa. Also known from Kenya, Zanzibar, Tanganyika, La Réunion, Seychelle Islands, Indonesia, Japan, Great Barrier Reed, Marshall Islands and Fijian Islands].

Periclimenella Bruce, 1994

- 1. Supraorbital spines absent; R. 1+5-8/3-5 .. *P. petitthouarsii* (Audouin, 1826) [Host genera: various. Distribution: Type locality: Egypt. Also known from Israel, Sudan, Eritrea, Saudi Arabia, Jibuti, Yemen, Persian Gulf, Kenya, Zanzibar, Tanganyika, Comoro Islands and Madagascar].
- Supraorbital spines present; R. 1+5-8/3-4...... *P. spinifera* (De Man, 1902) [Host genera: various. Distribution: Type localities: Ternate, Ambon, Indonesia; Tahiti, Society Islands. Also known from Madagascar to Tahiti and Wake Island, to southern Great Barrier Reef, north to the Ryukyu Islands, but absent from the north west Indian Ocean and Red Sca].

Periclimenes Costa, 1844

- Supraorbital spines absent 3

- 4. Medial margins of terminal and preterminal segments of second maxilliped endopod forming straight medial border; R.1+ 6-7/2-3

P. lutescens (Dana, 1852) [Host genera: *Acropora*, rarely *Pocillopora*, *Seriatopora*. Distribution: Type locality: Tongatabu, Tonga Islands. Also known from Egypt, Israel, Eritrea, Sa'udi Arabia, Kenya, Zanzibar, Tanganyika, Moçambique, Comoro Islands, Madagascar, Seychelle Islands, Maldive Islands, Nicobar Islands, Singapore. Indonesia, Nansha Islands, Japan, Great Barrier Reef, Coral Sea, Solomon Islands, Samoan Islands, Society Islands (?) and Marquesas Islands (?)].

Kenya, Tanganyika, Comoro Islands, La Réunion, Thailand, Philippines and Great Barrier Reef].

- Fingers of second perciopods with few (1-3) larger, acute, proximal teeth...8

- Fourth thoracic sternite with transverse linguiform median plate; ambulatory propods with well developed spines, R. 5-6/1... *P. difficilis* Bruce, 1976 [Host genus: *Porites*. Distribution: Type locality: Baie Ste Anne, Praslin, Seychelle Islands. Known only from the type locality, and Bird Island, also some on alcyonaria].

- 11. Major second pereiopod dactyl with single small acute tooth proximally,

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opposing two small teeth on fixed finger, without distinct distal diastema, major chela robust; ambulatory perciopods without ventral spines; ambulatory dactyl more than 0.25 of propod length; R. 1+6-7/1

P. madreporae Bruce, 1969 [Host genera: *Acropora, Pocillopora, Seriatopora, Stylophora, Turbinaria*. Distribution: Type locality: Erskine Island, Queensland, Australia. Also known from La Réunion, Indonesia, Philippines, Coral Sea, Caroline Islands (?) and Solomon Islands].

- Major second pereiopod with fixed finger tooth at 0.8 of length, separated by deep notch from tip; dactylar tooth obsolete; R. 5-7/0-2.....

P. diversipes Kemp, 1922 [Host genera: Acropora, Galaxea, Goniopora, Montipora, Pavona, Pocillopora, Porites, Psammocora, Seriatopora, Stylophora.. Distribution: Type locality: Kilakarai, Gulf of Manaar, India. Also known from Egypt, Yemen, Kenya, Zanzibar, Tanganyika, Comoro Islands, Madagascar, La Réunion, Seychelle Islands, Singapore, Thailand, Northern Territory, Great Barrier Reef and Coral Sea].

- Major second pereiopod with fixed finger tooth large, acute, at about 0.6 of finger length, with large gap proximally; dactylar tooth obsolete, distal cutting edge more or less straight; R. 5-7/0-1..... *P. gonioporae* Bruce, 1989* [Host genera: *Galaxea, Goniopora Lobophyllia, Montipora, Porites.* Distribution: Type locality: Ras Iwatine, Kenya. Also known from La Réunion and Great Barrier Reef].

Philarius Holthuis, 1952

- 1. Supraorbital spines present; R 7-10/1 *P. lifuensis* (Borradaile, 1898) [Host genus: *Acropora*. Distribution: Type locality: Lifu, Loyalty Islands, New Caledonia. Also known from the Great Barrier Reef].
- Carpus of second perciopod with strong medial spine distally; postrostral teeth present on carapace; R. 7-9/1 *P. imperialis* (Kubo, 1940) [Host genus: *Acropora*. Distribution: Type locality: Haha-jima, Ogasawara-Islands. Also known from Israel, Kenya, Zanzibar, Tanganyika, Comoro Islands, La Réunion, Singapore, Nansha Islands, Northern Territory, Great Barrier Reef, Coral Sea, Marshall Islands and Caroline Islands].

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donesia, Nansha Islands, Thailand, Japan, Philippines, Northern Territory, Great Barrier Reef, Coral Sea, Solomon Islands, Fijian Islands, Samoan Islands, Marshall Islands and Kiribati J.

Platycaris Holthius, 1952

Pontonides Borradaile, 1917

- Margin of lateral rostral carina rectangular; proximal lateral aspect of antepenultimate segment of third maxillipeds with blunt flattened setae; R. 0/0
 Pontonides sp. α* [Host genus: *Dendrophyllia*. Distribution: Tomioka, Amakusa Islands, Japan only, Fujino and Miyake, 1969, *as P. unciger* Calman].

Tectopontonia Bruce, 1973

Vir Holthuis, 1952

Yemenicaris Bruce, 1997

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Addendum: It has recently been shown by Mitsuhashi & Takeda (1998) that *C. pavonae* Bruce, 1972 is a junior synonym of *C. taiwanensis* Fujimo & Miyake, 1972 which is therefore the correct name for this species.

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