

# Crustacean fish parasites from Segara Anakan Lagoon, Java, Indonesia

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Received: 21 August 2006 / Accepted: 9 November 2006 / Published online: 12 January 2007  
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**Abstract** The present study is the first investigation on ectoparasites of commercial important fish from Segara Anakan, a brackish water lagoon located at the southern coast of Java, Indonesia. Eight economically important marine fish species (*Mugil cephalus*, *Siganus javus*, *Scatophagus argus*, *Caranx sexfasciatus*, *Lutjanus johnii*, *Eleutheronema tetradactylum*, *Johnius coitor*, and *Epinephelus coioides*) were examined for crustacean parasites. Prevalence and intensity data for each parasite species are given, together with an analysis of the origin and possible transmission pathways. A highly diverse copepod fauna consisting of 23 different species and two isopods was found. All fish species were at least infested with two copepod species, with the exception of *L. johnii*, *S. argus*, and *M. cephalus*. With seven and six species, respectively, they harboured the most species-rich ectoparasite fauna. The copepods *Ergasilus* sp. 3 and *Caligus acanthopagri* on *S. argus* showed the highest prevalence (78.6) and intensity

[17.8 (1–233) and 5.3 (1–22)] of infestation. The recorded parasite fauna is represented by marine, brackish water, and probably also freshwater components. The brackish water environment of Segara Anakan does not prevent disease outbreaks due to parasitic copepods by preventing pathogenic marine or freshwater species to enter the lagoon. This might cause fish health problems if the Segara Anakan Lagoon would be developed for finfish mariculture in future.

## Introduction

Indonesia is the largest archipelago in the world, consisting of more than 17,000 islands spread on 5.8 million km<sup>2</sup> of the sea (Harris 2000). With a coastline of about 81,000 km, Indonesia has the second largest coastal zone in the world. Approximately 830,000 ha of this area are potentially useful for brackish water aquaculture development and more than 140,000 ha are suitable for mariculture cages (Pollunin 1983; Harris 2000). The Segara Anakan area is a brackish water lagoon of approximately 4,000 ha, which is located on the western side of Cilacap (southern coast of Java) and is surrounded by about 14,000 ha of mangrove forests (Naamin 1991). This lagoon plays an important role as a nursery ground, and also has a major ecological function to support a large and productive mangrove ecosystem (Romimohtarto et al. 1991). Around 45 fish species were reported to occur in the Segara Anakan Lagoon (White et al. 1989), but the real species number is probably much higher (Dudley 2000). From 45 species, 10 are commonly caught and reported as economic important species in this area (Naamin 1991).

Indonesia can be considered as the center of marine biodiversity. For example, 23% of the so far described 254

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fish parasitic trypanorhynch cestodes have been recorded from Indonesian waters (Palm 2004). Jakob and Palm (2006) recorded 38 fish parasite species from mainly deep-water fish species from Pelabuhan Ratu, southern Java coast. The knowledge on the Indonesian marine fish parasites, however, is still scarce (Palm 2000). In Indonesia, more than 400 different parasite species have been recorded from around 240 marine fish species so far (also see Jakob and Palm 2006). This is only a fraction of the expected fish parasite biodiversity in approximately 3,000 marine fish species in Indonesia, underlining the poor exploration with respect to the occurrence of marine fish parasites (Hutomo 1986). Information from brackish water environments such as the Segara Anakan Lagoon is completely missing. This, nowadays, has become a major problem in identification and treatment of parasites and diseases in the rapidly developing mariculture industry (Sugama 1999; Zafran et al. 2000; Hartono et al. 2001; Roza et al. 2002; Dewi et al. 2003). There is an urgent need to better describe and identify the Indonesian fish parasite fauna to prevent and overcome future fish disease and parasite outbreaks.

The establishment of future mariculture facilities within and in the surrounding of the Segara Anakan Lagoon needs information on potential threats by fish parasites. The present study is the first investigation on ectoparasites from commercial important fish species from Segara Anakan. It contributes to our knowledge on fish parasitic crustaceans on free-living marine fish species from the southern Java coast, which might also be cultivated in Indonesia in future. Because fish parasites are one of the large health problems within mariculture facilities, parasite identification is

provided to facilitate the best treatment and more detailed investigation in the future.

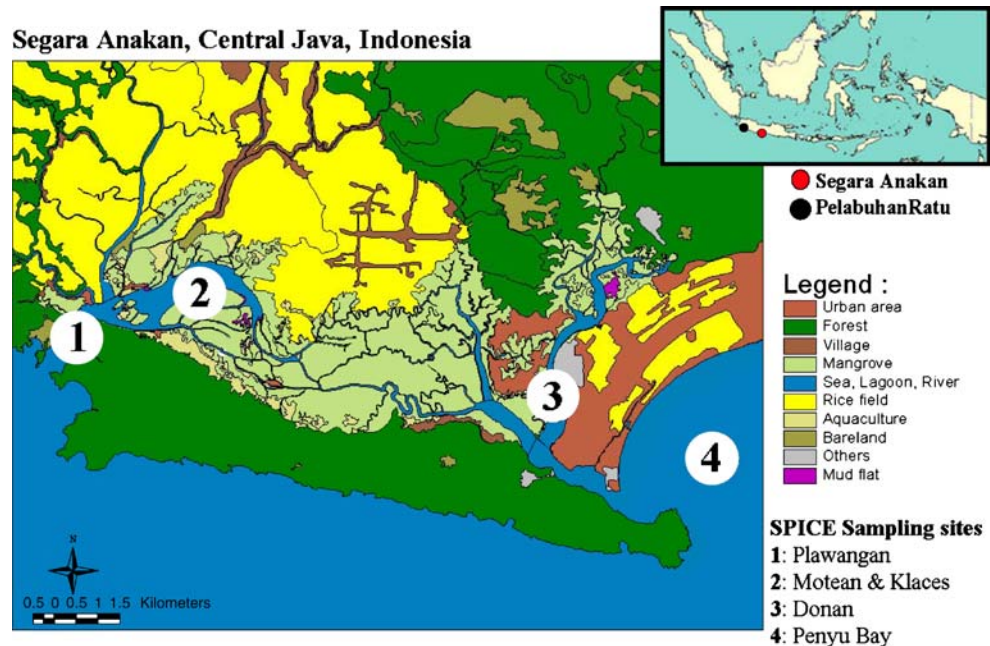
## Materials and methods

Samples were taken from August to November 2004 at two different localities in the Segara Anakan Lagoon (Central Java). Both selected sampling sites (Motean and Klaces/area 2 and Donan/area 3) are part of the sampling areas of the project “Science for the Protection of Indonesian Coastal Ecosystems” (SPICE) within the German–Indonesian cooperation in Marine Sciences (Fig. 1). They are differentiated by its environmental characteristics. Area 2 is located in the center of the lagoon and is influenced by several rivers that carry freshwater runoff and sedimentation. The salinity is low (19.7–28.0). Area 3 is located near to an outlet to the Indian Ocean. Therefore, the salinity is higher (29.3–31.2).

The fish was obtained directly from local fishermen. The following marine and brackish water fish species were studied from the two sampling sites within Segara Anakan Lagoon: *Mugil cephalus* L. (70), *Scatophagus argus* (L., 1766) (70), *Siganus javus* (L., 1766) (5), *Caranx sexfasciatus* Quoy and Gaimard, 1825 (8), *Lutjanus johnii* (Bloch, 1792) (8), *Eleutheronema tetradactylum* (Shaw, 1804) (8), *Johnius coitor* (Hamilton, 1822) (20), and *Epinephelus coioides* (Hamilton, 1822) (21).

Fish dissection was done in the Parasitology and Entomology Laboratory of the Biology Faculty, Jenderal Soedirman University (UNSOED), Purwokerto. Before examination for parasites, the unfrozen fish were measured for length and

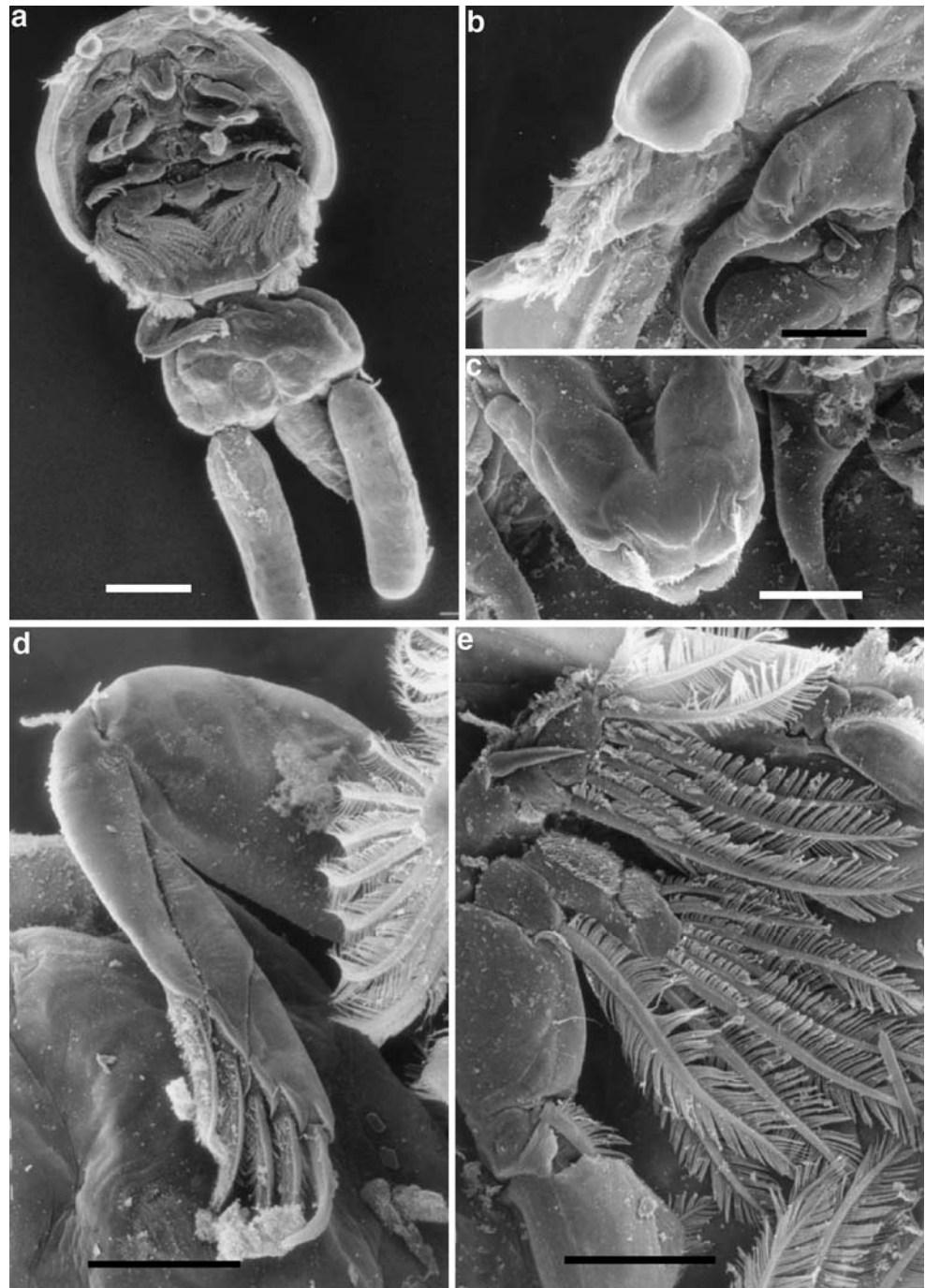
**Fig. 1** Segara Anakan, Cilacap, Indonesia and the sampling localities according to the project sites in “Science for the Protection of Indonesian Coastal Ecosystems (SPICE 2003–2007)”. (Map source: Segara Anakan map, SPICE Project)



weight. Each fish was examined microscopically for the presence of parasitic crustaceans based on a method described by Kabata (1985). The collection and preservation methodology for crustacean parasites followed Pritchard and Kruse (1982). Pictures were taken by using a digital camera, Canon PC 1015, attached to the Axioskop 40, and in some cases to a stereomicroscope (STEMI SV 11 Zeiss, Germany). The ecological terms in parasitology, prevalence, intensity, and mean intensity, follow Bush et al. (1997).

Selected specimens were prepared following Robinson et al. (1985) for scanning electron microscopical studies (Fig. 2). A LEITZ-AMR 1000 and International Scientific Instruments ISI-100 B were used for scanning the samples at 30 and 15 kV. The scanning electron microscopy photomicrographs were made with the help of a Leitz, LEICA MD-2, Canada and reflex camera with AGFA APX 25 professional 135, Ilford FP4 plus 125 and Ilford Panf plus 50.

**Fig. 2** Scanning electron microscopy of *Caligus acanthopagri* (♀) collected from *Scatophagus argus*. **a** Complete specimen, ventral view; **b** lunulae and first antenna; **c** mouth cone; **d** second leg; **e** third leg; Scale bars: **a** 500  $\mu\text{m}$ ; **b** 70  $\mu\text{m}$ ; **c** 50  $\mu\text{m}$ ; **d** and **e** 100  $\mu\text{m}$



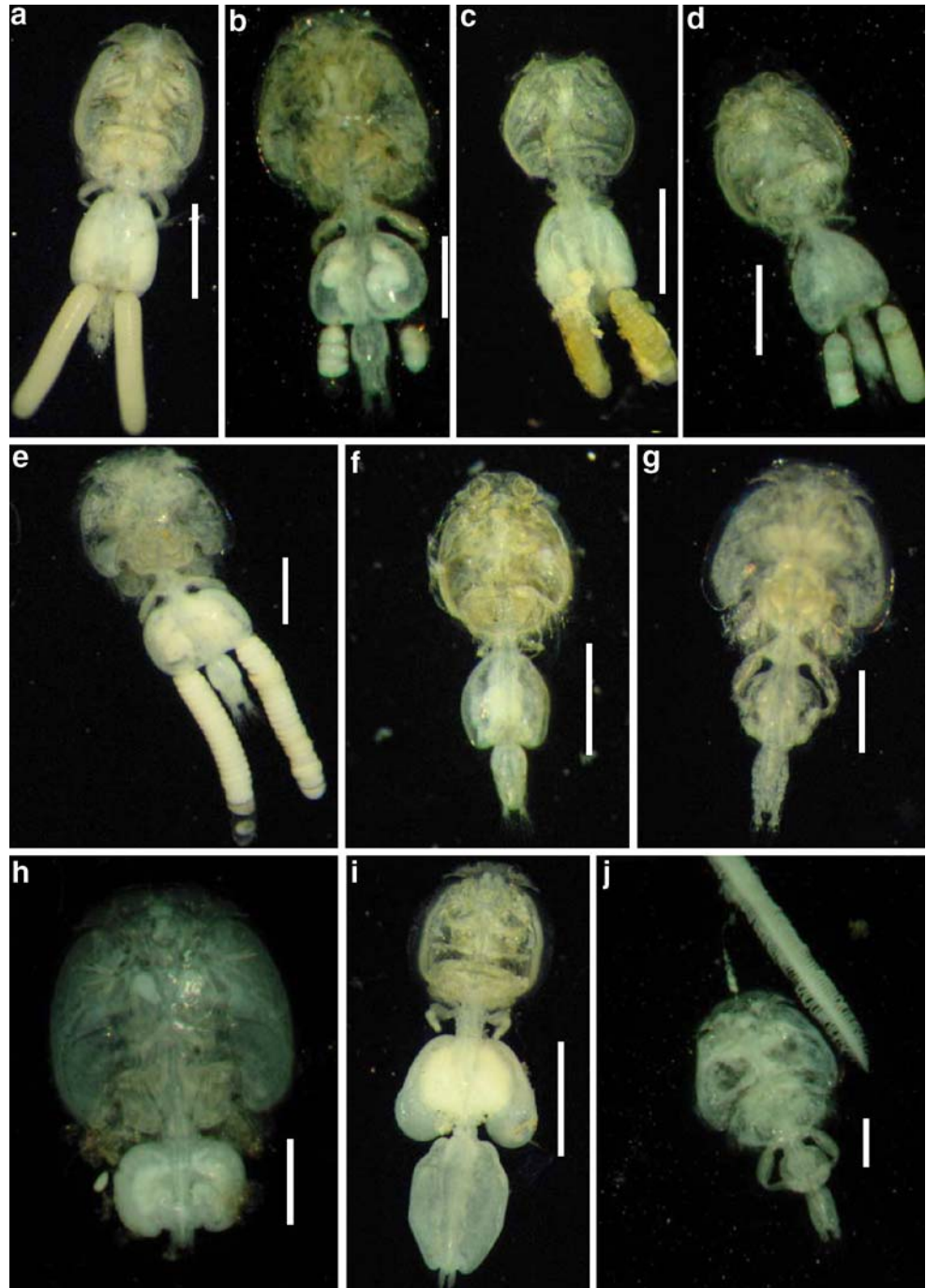
## Results

During the present study, 210 fishes (eight species belonging to eight families) from the Segara Anakan Lagoon, central Java, were investigated for the presence of parasitic crustaceans. The isolated parasites belonged to various copepod and two isopod families. All fish species, with the exception of *L. johnii*, were at least infested with two crustacean species.

Out of all studied fish specimens, only 10 were found uninfested. Copepods were the most common ectoparasites and occurred in nearly all fish samples. In contrast, only two fish species were infested with Isopoda.

A total of 23 parasitic copepod (Figs. 2, 3, and 4) and two isopod species were found, 23 species occurred as adult and three species were found in the larval stage (Table 1). The copepods of the families Ergasilidae and Caligidae were the most common parasites on the examined fish. *M.*

**Fig. 3** Caligid copepod species from Segara Anakan. **a** *Caligus phipsoni* (♀) with egg sacs from *Eleutheronema tetradactylum*; **b** *Caligus* sp. (♀) with egg sacs from *Johnius coitor*; **c** *Caligus* cf. *confusus* (♀) with egg sacs from *Caranx sexfasciatus*; **d** *Caligus* cf. *quadratus* (♀) with egg sacs from *Siganus javus*; **e** *Caligus acanthopagri* (♀) with egg sacs from *Scatophagus argus*; **f** *Caligus* cf. *epinepheli* (♀) from *Epinephelus coioides*; **g** *Caligus rotundigenitalis* (♂) from *Mugil cephalus*; **h** *Caligus epidemicus* (♀) from *S. argus*; **i** *Parapetalus hirsutus* (♀) from *E. tetradactylum*; **j** Chalimus stage. Scale bars: **a** and **d** 1.2 mm; **b**, **e**, and **g** 0.55 mm; **c** 0.8 mm; **f** 0.65 mm; **h** 0.5 mm; **i** 1.5 mm; **j** 0.3 mm



**Fig. 4** Copepod species from Segara Anakan. **a** *Nothobomolochus* sp. (♀) with egg sacs from *Mugil cephalus*; **b** *Ergasilus* sp. 1 (♀) with egg sacs from *M. cephalus*; **c** Ergasilidae gen. et sp. indet. (♀) from *M. cephalus*; **d** *Ergasilus* sp. 4 (♀) with egg sacs from *Siganus javus*; **e** *Thysanote* sp. (♀) with egg sacs from *Scatophagus argus*; **f** *Peniculus* cf. *scomberi* (♀) from *Caranx sexfasciatus*; **g** Pennellidae gen. et sp. indet. from *Epinephelus coioides*; **h** *Naobranchia* cf. *polynemi* (♀) from *Eleutheronema tetradactylum*. Scale bars: **a–c** 300 µm; **d** 70 µm; **e** 1.5 µm; **f** 1 mm; **g** 200 µm; **h** 50 µm



*cephalus* and *S. argus* harbored the most diverse crustacean parasite fauna. A brief description of the collected crustaceans from this unique environment together with notes on the species identification is given below.

Subclass: Copepoda Milne Edwards, 1830

Order: Cyclopoida Sars, 1886

Family: Bomolochidae Sumpf, 1871

Only a single bomolochid copepod species was collected from the gill mucus of *M. cephalus* (Fig. 4a). *Nothobomolochus* sp. is characterized by the structure of the antenna

(three modified setae on the proximal segment of antennule) and the number of setae in caudal rami (one major setae). The species was identified after the genus description of Kabata (1979). Many of the 34 species of *Nothobomolochus* appear remarkably similar, and these similarities have resulted in great taxonomic difficulties. None of these tiny bomolochids is easy to identify. Body shapes and even proportions vary with swelling due to the state of maturity.

Family: Ergasilidae von Nordmann, 1832

**Table 1** Fish parasitic Crustacea from Segara Anakan Lagoon (A=adult; L=larval stage)

| Fish species                             | Parasite species (stage)                      | Site  | Prevalence (%) | Mean intensity (intensity) |
|--|---|---|----------------|----------------------------|
| <i>Mugil cephalus</i> (n=70)             |   |   |                |                            |
| Copepoda                                 | <i>Nothobomolochus</i> sp. (A)**              | Gill mucus                                      | 17.1           | 1.8 (1–5)                  |
|  | <i>Ergasilus</i> sp. 1 (A)**                  | Gill filaments                                  | 51             | 5.7 (1–37)                 |
|  | <i>Ergasilus</i> sp. 2 (A)*, **               | Gill filaments                                  | 1.4            | 3 (3)                      |
|  | Ergasilidae gen. et sp. indet. (A)*, **       | Gill cavity                                     | 24.3           | 4.1 (1–15)                 |
|  | <i>Caligus rotundigenitalis</i> (A)*          | Inner operculum                                 | 24.3           | 2.2 (1–15)                 |
|  | Chalimus stage (L)                            | Gill filaments                                  | 24.3           | 2.3 (1–8)                  |
| <i>Siganus javus</i> (n=5)               |   |   |                |                            |
| Copepoda                                 | <i>Ergasilus</i> sp. 4 (A)*, **               | Gill filament                                   | 20             | 1 (1)                      |
|  | <i>Caligus epidemicus</i> (A)*                | Skin  | 20             | 1 (1)                      |
|  | <i>Caligus</i> cf. <i>quadratus</i> (A)**     | Gill filaments, gill cavity and inner operculum | 100            | 1 (1)                      |
| <i>Scatophagus argus</i> (n=70)          |   |   |                |                            |
| Copepoda                                 | <i>Ergasilus</i> sp. 2 (A)*, **               | Inner operculum and gill mucus                  | 7.1            | 21.8 (1–65)                |
|  | <i>Ergasilus</i> sp. 3 (A)*, **               | Gill filaments                                  | 78.6           | 17.8 (1–233)               |
|  | <i>Caligus acanthopagri</i> (A)*              | Inner operculum and gill filaments              | 78.6           | 5.3 (1–22)                 |
|  | <i>Caligus epidemicus</i> (A)*                | Skin  | 4.3            | 1.33 (1–2)                 |
|  | <i>Pseudocaligus</i> sp. (A)*, **             | Skin  | 5.7            | 1.75 (1–4)                 |
|  | Chalimus stage (L)                            | Gill filaments                                  | 75.7           | 6.2 (1–44)                 |
|  | <i>Thysanote</i> sp. (A)*, **                 | Nasal cavity                                    | 65.7           | 1.8 (1–2)                  |
| Isopoda                                  | <i>Cymothoa</i> sp. (A)*, **                  | Gill rakers and mouth cavity                    | 14.3           | 1.4 (1–2)                  |
|  | Gnathiidae gen. et sp. Indet. (L)*            | Gill filaments                                  | 2.8            | 1 (1)                      |
| <i>Caranx sexfasciatus</i> (n=8)         |   |   |                |                            |
| Copepoda                                 | <i>Caligus</i> cf. <i>confusus</i> (A)*       | Gill filaments                                  | 100            | 8 (4–17)                   |
|  | Chalimus stage (L)                            | Gill filaments and mouth cavity                 | 37.5           | 2 (1–3)                    |
|  | <i>Peniculus</i> cf. <i>scomberi</i> (A)*, ** | Dorsal fin                                      | 12.5           | 1 (1)                      |
| Isopoda                                  | Gnathiidae gen. et sp. indet. (L)*            | Gill filaments                                  | 12.5           | 1 (1)                      |
| <i>Lutjanus johnii</i> (n=8)             |   |   |                |                            |
| Copepoda                                 |   |   |                |                            |
| <i>Eleutheronema tetradactylum</i> (n=8) |   |   |                |                            |
| Copepoda                                 | <i>Caligus phipsoni</i> (A)*                  | Inner operculum and gill filaments              | 75             | 2.8 (1–6)                  |
|  | <i>Parapetalus hirsutus</i> (A)*              | Inner operculum                                 | 75.5           | 1.6 (1–3)                  |
|  | Chalimus stage (L)                            | Gill filaments                                  | 25             | 1 (1)                      |
|  | <i>Naobranchia</i> cf. <i>polynemi</i> (A)*   | Gill filaments                                  | 12.5           | 1 (1)                      |
|  | <i>Lernanthropus polynemi</i> (A)*            | Gill filaments                                  | 88             | 3.7 (2–8)                  |
| <i>Johnius coitor</i> (n=20)             |   |   |                |                            |
| Copepoda                                 | <i>Caligus</i> sp. (A)                        | Gill filaments                                  | 25             | 2 (1–4)                    |
|  | Chalimus stage (L)                            | Gill rakers                                     | 30             | 1.3 (1–2)                  |
|  | <i>Lernanthropus</i> sp. (A)*                 | Gill filaments                                  | 5              | 1 (1)                      |
|  | <i>Peniculus</i> cf. <i>scomberi</i> (A)*     | Dorsal fin                                      | 5              | 1 (1)                      |
| <i>Epinephelus coioides</i> (n=21)       |   |   |                |                            |
| Copepoda                                 | <i>Caligus</i> cf. <i>epinepheli</i> (A)*     | Gill filaments                                  | 4.8            | 1 (1)                      |
|  | Pennellidae gen. et sp. indet. (L)*, **       | Gill filaments and gill rakers                  | 71.4           | 30.1 (1–233)               |

\* new locality record, \*\* new host record

A total of five ergasilid species were collected (Fig. 4b–d). The species were identified by using the identification key to ergasilid genera by Boxhall and Halsey (2004). *Ergasilus* sp. 1 was collected from *M. cephalus* (Fig. 4b). It is characterized by a typical cyclopiform body shape with clear external segmentation, the presence of biramous fourth legs, antenna with one claw, and the first swimming legs without modified endopod. *Ergasilus* sp. 2 was collected from *S. argus* at high

prevalence and intensity and from *M. cephalus* at low prevalence and intensity of infestation. This species differs from *Ergasilus* sp. 1 in several characters, such as the rounded cephalothorax, the structure and smaller size of the second antenna, and differences in the morphology of the caudal rami, which are not biramous in the present species. *Ergasilus* sp. 3 was collected from *S. argus*. It differs from the previous ergasilid species by having a

more rounded body shape, differences in caudal rami, and the fine structure of the second antenna. *Ergasilus* sp. 4 was found at a low prevalence and intensity on *S. javus* (Fig. 4d). This species differs from the other ergasilids in its body shape and size (shape of thoracic segments, caudal branches, and number of seta.), the morphology of caudal rami and extremely long egg sacs. The morphology of all four species did not correspond to any other known *Ergasilus* species; the species descriptions will be a matter of another communication.

Ergasilidae gen. et sp. indet. (Fig. 4c)

The genus *Ergasilus* contains 135 tentatively valid species. Some additional species are treated as species inquirendae or incertae sedis. This species-rich genus lacks revision, making taxonomic work very difficult. In addition, several species have not been well described or not seen again since the original description. In many cases, the type specimens are either inaccessible or not longer extant (Lin and Ho 1998).

Order: Siphonostomatoida Latreille, 1829

Family: Caligidae Burmeister, 1835

A total of 10 caligid species were collected (Fig. 3). Out of these, eight species belong to the genus *Caligus*, one species to *Parapetalus*, and one species to *Pseudocaligus*. *Caligus acanthopagri* Lin, Ho, Shen, 1994 is characterized by a two-segmented antennule, 3-segmented antenna, a small abdomen, and long egg sacs that are longer than one-half of the body length. *C. acanthopagri* was collected from the gills and opercula of *S. argus* at a high prevalence and intensity (Fig. 2). The species was identified by using the original description of Lin et al. (1994). *Caligus* cf. *confusus* Pillai, 1961 was collected from *C. sexfasciatus*. It is characterized by a suborbicular cephalothorax shield, short and 1-segmented abdomen, 3-segmented antenna, and egg sacs shorter than the body. The species is most similar to *C. confusus* described by Pillai (1961), but showed slight differences in the armature of the antenna and the swimming legs. *Caligus epidemicus* Hewitt, 1971 was collected from the skin of *S. argus* and *S. javus*. This species is characterized by a subcircular cephalothorax shield, a small and 1-segmented abdomen, short caudal rami, 3-segmented antenna, and a subquadrate sternal furca. *C. epidemicus* was identified by using Ho and Lin (2004). *Caligus* cf. *epinepheli* Yamaguti, 1936 was found on the gill filaments of *Epinephelus coiodes*. This species shared most common features with *C. epinepheli*, e.g., a subcircular cephalothorax shield, a long abdomen, and 3-segmented antenna. However, the specimens from Segara Anakan showed slight differences in the antenna and swimming legs. *Caligus phipsoni* Bassett-Smith, 1898 was isolated from the gills of *E. tetradactylum*. The structure of the first antenna and swimming legs of the present specimen are similar to those in the original description of *C. phipsoni*. As another important character,

*C. phipsoni* has bigger lunulae than any other *Caligus* species. *Caligus* cf. *quadratus* Shiino, 1954 was found on *S. javus*. It is characterized by a suborbicular cephalothorax shield, a long abdomen, caudal rami ration 1.6 times longer than wide, and a 3-segmented antenna, but showed slight differences in the armature of the antenna and the swimming legs. The species was identified after Ho and Lin (2004). *Caligus rotundigenitalis* Yü, 1933 was found on the inner operculum of *M. cephalus*. This species is characterized by a subcircular cephalothorax shield, 2-segmented abdomen with smaller proximal segment, and a 3-segmented antenna. This caligid copepod was also identified after Ho and Lin (2004).

*Caligus* sp. was collected from the gill filaments of *J. coitor*. The specimens are characterized by a subcircular cephalothorax shield, the presence of a sterna furca, small abdomen, and a well-developed fourth leg. This species could not be identified to the species level and might represent a currently undescribed species. *Parapetalus hirsutus* (Bassett-Smith, 1898) was found on the inner operculum of *E. tetradactylum*. It is characterized by a subcircular cephalothorax shield, 1-segmented large abdomen, oval caudal rami, and a 3-segmented antenna (Ho and Lin 2004).

*Pseudocaligus* sp. was obtained on the skin of *S. argus*. The characters of *Pseudocaligus* are similar to those of *Caligus*, except for a vestigial fourth leg or the complete absence of this leg (Ho and Lin 2004). This species is characterized by a round cephalothorax, short abdomen, and the presence of a sternal furca. The morphology of the present specimens was identical with the diagnosis of *Pseudocaligus* given by Ho and Lin (2004).

*P. hirsutus* was collected from the inner operculum of *E. tetradactylum*. The shape of this species is different with *Caligus* and it is characterized by a subcircular cephalothorax shield, 1-segmented large abdomen, caudal rami oval, and a 3-segmented antenna.

Family: Lernaepodidae Milne Edwards, 1840

Two species of the family Lernaepodidae were isolated. *Naobranhia* cf. *polynemi* Tripathi, 1962 occurred on the gill filament of *E. tetradactylum* (Fig. 4h). In the present study, only a single non gravid female was found, and was identified by using the original species description by Tripathi (1962). However, the description is brief and *N. polynemi* has not been redescribed. *Thysanote* sp. was isolated from the nasal cavity of *S. argus*. It is characterized by the short cephalothorax, a large trunk, and unbranched processes at the hind end of the trunk. The species was identified to the genus level by using the key of lernaepodid genera by Boxhall and Halsey (2004).

Family: Lernanthropidae Kabata, 1979

Two species of lernanthropid copepods were found. *Lernanthropus polynemi* Richiardi, 1881 was found on the gill filaments of *E. tetradactylum*. This species can be

identified by its cephalothorax shape, the subtriangular head, the shape of the urosome, and the straight egg sacs. The morphological features of the specimen correspond to the redescription by Piasecki and Hayward (2002). Another species of the Lernanthropidae, *Lernanthropus* sp., was collected from the gill filaments of *J. coitor*. According to Boxshall and Halsey (2004), the genus *Lernanthropus* consists of 119 species, but an identification key is not available.

Family: Pennellidae Burmeister, 1835

One adult and one larval stage of the family Pennellidae were found. *Peniculus* cf. *scomberi* Gnanamuthu, 1951 was collected from the dorsal fin of *C. sexfasciatus* and *J. coitor* (Fig. 4f). This species is characterized by an oval, short cephalothorax with a prominent proboscis-like mouth cone. Two narrow free thoracic segments are interposed between the cephalothorax and the genital complex, forming a neck-like structure. The body shape corresponds to the original description of *P. scomberi* by Gnanamuthu (1951). However, the original description is insufficient (Alexander 1983) and needs redescription. The larval pennellids attached on the gill filaments and gill rakers of *E. coioides* could not be further identified (Fig. 4g). They are characterized by attachment organs (frontal filament), the presence of a mouth cone and curious structures around the mouth cone. These characters allow assignment to the family Pennellidae; however, the life cycle and developmental stages of this family are widely unknown.

Class: Malacostraca Latreille, 1806

Order: Isopoda Latreille, 1817

Family: Cymothoidae Leach, 1818

One adult Isopoda, *Cymothoa* sp., was isolated from the mouth cavity of *S. argus*. The genus *Cymothoa* is mainly characterized by the general body shape (cephalon sunken into preonite), and the morphology of the pleon (distinct from pereon) and the cephalon. The actual species composition of the genus *Cymothoa* is unclear and a revision is needed (Bunkley-Williams and Williams 1998).

Family: Gnathiidae Harger, 1880

The pranzia stages of gnathiid isopods were found on the gill filaments of *S. argus* and *C. sexfasciatus*. The pranzia stages could be identified to the family level only due to the lack of species-specific characters. Gnathiid larvae can be recognized by its characteristic body shape, resembling a thin female with large eyes. The living larvae are characterized by a red color due to the uptake of large quantities of blood.

## Discussion

According to Froese et al. (1996) and Tomascik et al. (1997), Indonesia is part of the world's center of marine aquatic biodiversity. The exceptionally high biodiversity of

the marine fauna in the Indonesian Archipelago is a result of its geographical location and geological history. Though less than 10% of the Indonesian marine and brackish water fish fauna have yet been studied, the parasite fauna appears to be highly diverse. Parasites are an important but often underestimated part of marine biodiversity (Marcogliese and Price 1997). Palm et al. (1999) estimated that more than three different metazoan parasites occur in each marine fish species, leading to the assumption that more than 9,000 metazoan marine fish parasites occur in Indonesia. To date, 242 marine and brackish water fish species have been studied for their parasite fauna in Indonesian waters, revealing more than 400 species. However, the Indonesian fish species are far from being fully studied and in some regions, nobody has ever worked on marine fish parasites.

This is the first study on fish ectoparasites from Segara Anakan Lagoon. A species-rich ectoparasite fauna was recorded. Twenty-two species or genera were recorded for the first time from Indonesian waters or from the southern coast of Java. In addition, eight new host records could be established. The collected crustaceans consist of typical shallow water species. According to Boxshall and Halsey (2004), caligid, ergasilid, and lernanthropid copepods are known as common parasites of shallow water fish. Copepods of the family Ergasilidae are mostly known as freshwater parasites, and only few species are known from the brackish water or marine environment (Boxshall and Halsey 2004). The other collected copepod families (Bomolochidae, Caligidae, Lernanthropidae, Lernaepodiidae, and Pennellidae) are mainly or exclusively known as marine fish parasites (Kabata 1979; Boxshall and Halsey 2004), and also the isopods *Cymothoa* sp. and gnathiids are known as ectoparasites of marine fish (Möller and Anders 1986).

Within the present study, *M. cephalus*, *S. argus*, *E. tetradactylum*, and *J. coitor* had a species-rich copepod fauna. Six parasitic copepods were recorded from *M. cephalus*. Even though this fish species has a wide distribution and has been well studied for copepod parasites (e.g., Paperna and Overstreet 1981; El-Rashidy and Boxshall 1999), several copepods from the present study represent new host records. According to El-Rashidy and Boxshall (1999), at least nine different ergasilid genera have been recorded from mugilid fish, and *M. cephalus* alone harbors six different genera (*Dermoergasilus*, *Ergasilus*, *Mugilicola*, *Nipergasilus*, *Paraergasilus*, and *Thersitina*). Three species of ergasilid copepods were recorded from Segara Anakan; two of them belonging to *Ergasilus* and one species probably represents an undescribed genus. *S. argus* was infested with seven copepod species, and showed the highest number of species. Beside copepods, the parasitic isopod *Cymothoa* sp. was found to infest *S.*



*argus*. Isopods belonging to *Cymothoa* are common in the mouth cavity of their hosts (Bunkley-Williams and Williams 1998). Mladineo and Valic (2002) observed the mechanism of infection in cymothoid isopods. The larvae actively search their fish hosts after leaving the female, and *S. argus* is a suitable host for *Cymothoa* sp.

In contrast to the fish species mentioned above, *E. coioides*, *J. coitor*, *S. javus*, and *C. sexfasciatus* harbored only a low number of host specific copepods. According to Boxshall and Halsey (2004), most parasitic copepods are known to have high host specificity. For example, *L. polynemi* and *P. hirsutus* occur exclusively on polynemid fish (Pillai 1962; Ho and Lin 2001; Piasecki and Hayward 2002). Others such as some *Caligus* spp. are known to have low hosts specificity (Ho and Lin 2004). For example, *Caligus elongatus* Nordmann, 1832 has been recorded from more than 100 host species, both teleosts and even elasmobranchs, belonging to 47 families (Williams and Bunkley-Williams 1996). Within the present study, seven out of eight fish species were infested with *Caligus* spp. Similarly, the gnathiid isopods on *S. argus* and *C. sexfasciatus* show low host specificity. According to Möller and Anders (1986), pranzia stages were recorded to infest a high number of different fish species. Most copepods from Segara Anakan were host-specific, with 19 species infesting only a single host fish species. However, some species such as *C. epidemicus* and *C. rotundigenitalis* are known to infest several fish species, thus having a low level of host specificity such as the isolated isopods.

Segara Anakan is a brackish water environment and seems to exclude some typical marine helminths such as the trypanorhynch cestodes (Yuniar 2005). Remane (1934) developed a diagram to describe the faunistic distribution along a marine freshwater salinity gradient. According to him, the true brackish water species are less diverse than those living in marine or freshwater environments. According to Nybakken (2001), three faunistic components occur within estuaries, marine, freshwater, and brackish water or estuarine species. The species composition depends on the actual salinity. Beside true brackish water species, additionally, some freshwater and marine species with a high salinity tolerance can be found in such environments (Nybakken 2001). Zander (1998) published a review on parasites in brackish water environments based on research in the Baltic Sea, the largest brackish water environment in Europe. The author stated that the parasite fauna in brackish water regions is poor relative to truly marine habitats, host specificity is low, new hosts have been acquired, the number of hosts in life cycles is reduced, and there are adaptations to brackish water hosts. Studies on fish parasites in brackish water habitats in tropical areas are scarce and have been mainly carried out on copepods in India and Brazil (Babu and Raj 1985; Boxshall and Montú 1997).

Data from Indonesian waters are lacking. Within the present study, the parasites collected from Segara Anakan Lagoon show a wide salinity range. According to Zander (1997), within a salinity of 6–8 ppt, the species diversity is minimal. Twenty-four parasitic crustacean species were recorded in the studied brackish water Segara Anakan, showing high ectoparasite richness. This might be explained by a still high salinity ranging from 19–31 ppt within the lagoon, close to the high salinity in marine environments. This contrasts, however, the observation that some marine helminth species do not enter the lagoon, though being common in the marine system along the southern Java coast (Yuniar 2005, Palm 2004, Jakob and Palm 2006).

### Importance of fish parasites for future mariculture development in Segara Anakan Lagoon

Marine finfish is often cultivated in coastal fish ponds. The floating net cage-culture system is mainly extensive and expanding rapidly throughout Southeast Asia. These systems nowadays serve as an important source of fish for the local and international markets (Leong 1992). The impact of parasites on marine finfish culture is well documented, with catastrophic losses reported for disease outbreaks that result in high mortality levels. Prerequisite for establishing effective control measures in fish cultures is a quick and exact diagnosis of the causative agent, and the knowledge on the complex host–parasite–environment relationships (Moravec 1994). The correct parasite identification is of great significance, and may decide on the survival or death of the infested fish. Today, more than 60 parasite species are known that directly affect the Indonesian mariculture. Unfortunately, identification of many of these parasites is still unclear.

According to Ho (2000) and Johnson et al. (2004), caligid copepods are economically important parasites in mariculture facilities. Within Segara Anakan, 10 species of caligid copepods were collected from eight fish species. *C. acanthopagri*, *C. epidemicus*, and *C. rotundigenitalis* already caused severe problems within the Asian mariculture (Ho and Lin 2004). In Indonesia, several cases of fish mortalities related to the infestation with *Caligus* spp. have been recorded, such as in grouper culture in Gondol Research Station, Bali (Yuasa et al. 1998; Zafran et al. 2000). Lin and Ho (1998) stated that two species belonging to the family Ergasilidae were found on fish cultured in brackish water in Taiwan. Within the present study, five species of ergasilid copepods were recorded from Segara Anakan Lagoon. *Lernanthropus* sp. was recorded as a parasite of snapper (*L. johnii*) cultured in floating net cages in Malaysia (Leong and Wong 1989). *Lernanthropid*

copepods were also collected in Segara Anakan, infesting *J. coitor* and *E. tetradactylum*.

Isopods are also common parasites in mariculture facilities. Papapanagiotou and Trilles (2001) listed several isopods of the family Cymothoidae, which are known to infest cultivated marine fish, also causing great financial loss. For example, *Ceratothoa parallela* (Otto, 1828) caused over 50% mortality of *Sparus aurata* L., 1758 cultured in Greece (Papapanagiotou and Trilles 2001). Isopods on fish cultured along the Peruvian coast were published by Williams and Bunkley-Williams (2000). The authors recorded that *Ceratothoa gaudichaudii* (Milne Edwards, 1840) caused a 15% loss in body weight, costing approximately 1.3 billion kilograms of annual loss to the fishermen. *Cymothoa oestrum* (L., 1793) was reported to cause superf infections in caged fishes in the Caribbean (Williams and Bunkley-Williams 2000). In Indonesia, problems in mariculture facilities in some localities were caused by Isopoda. For example cymothoid isopods were recorded in the nasal and gill cavity of groupers cultured in Bali (Koesharyani et al. 2001). In Segara Anakan Lagoon, one species of Isopoda of the same family was found in the mouth cavity of *S. argus*. This species might also become a problem in future mariculture activities within the region.

## Conclusions

The present study of ectoparasites from commercial fish species in Segara Anakan Lagoon demonstrates a high parasite richness within this tropical brackish water environment. All recorded species are typical shallow water species, and some of them are known as economically important pathogens within the Asian mariculture. Consequently, the brackish water environment does not prevent disease outbreaks due to parasitic copepods by preventing pathogenic marine or freshwater species to enter the lagoon. This might cause disease problems if the Segara Anakan Lagoon would be developed for commercial finfish mariculture in the future. The studied fish species varied in the number of copepod species. These differences depend on the host specificity of the parasites and the habitat preference of the host. Most of the isolated copepods appear to have high host specificity within the lagoon, occurring on a single host fish species. However, with *C. epidemicus* and *C. rotundigenitalis*, also species with lower host specificity and wider host range were found. Salinity is considered one of the main factors influencing the parasite infestation with ectoparasitic copepods. Some euryhalin metazoan ectoparasites can tolerate a wider salinity range (e.g., ergasilid copepods), but most species found are stenohalin and prefer the marine

environment. The parasitic copepod fauna of Segara Anakan Lagoon is dominated by marine species, all of them with obviously a high degree of salinity tolerance, thus being euryhalin. The present study demonstrates a high ectoparasitic crustacean richness also in a brackish water tropical ecosystem. This and further studies will determine the actual contribution of the Indonesian parasite fauna to marine biodiversity. Larger samples and studies from other Indonesian regions are needed to describe and analyze the real parasite diversity and to verify the potential threats of fish parasites for the Indonesian mariculture industry.

**Acknowledgements** We would like to thank Prof. H. Mehlhorn (Institut für Zoomorphologie, Zellbiologie und Parasitologie, Heinrich-Heine-Universität Düsseldorf) for the possibility to carry out parts of scanning electron microscopy. We would also like to thank Prof. G.A. Boxshall (National Museum of Natural History, London) and Prof. J.-S. Ho (Department of Biological Sciences, California State University, California) for confirmation of some copepod identification and literature support. We also thank SPICE project members Dr. T. Jennerjahn (ZMT Bremen), E. Ardli, M.Sc. (Jenderal Soedirman University and ZMT Bremen), and Dr. B. Heru, Arthadi, M.S. and S. Subadrah, S.U. of the Parasitology and Entomology Laboratory, Jenderal Soedirman University (UNSOED), Purwokerto, for their support during data collection. Financial support was provided by the German Academic Exchange Service (DAAD), the German Federal Ministry of Education and Research (BMBF), and the German Research Council PA 664/4-1.

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