Growth and survival of post-larval giant tiger shrimp *Penaeus monodon* feeding on mangrove leaf litter biofilms

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ABSTRACT: Biofilm associated with decomposing mangrove leaf litter is a food source for shrimp post-larvae (PL). PL of giant tiger shrimp *Penaeus monodon* foraging on leaf litter of *Rhizophora mucronata* with developing biofilm at 1, 3, 4, 6, and 8 wk of decomposition were tested for specific growth rate (SGR, %) and percentage survival (SR, %). Biofilm was analyzed for species composition, abundance, and biomass of microalgae and epifauna. Microalgal biomass increased with the progress of litter decomposition. Diatoms, especially *Navicula* sp. and *Nitzschia* sp., dominated the first 6 wk of litter decomposition with a percentage cover of 88 and 99% during the third and fourth weeks, respectively. Cyanobacteria dominated in the 8 wk old biofilm with 61% cover. Copepods dominated the epifauna during the first 3 wk of litter decomposition. In the 4 to 5 wk old litter, polychaetes were most abundant whereas nematodes were dominant in litter that had decomposed for 8 wk. PL foraging on 4 wk old litter had a higher SGR (1.6 ± 0.5%) and SR (39.8 ± 4.8%) coincident with the maximum abundance of microalgae and epifauna. The study illustrated that nutritionally rich biofilm for PL shrimp is: (1) developed during the fourth week of mangrove leaf litter decomposition and is dominated by diatoms, polychaetes, harpacticoid copepods, and oligochaetes; and (2) limited by the collapse of the epifauna and subsequent colonization by cyanobacteria in mangrove leaf litter decomposed beyond 4 wk.

KEY WORDS: Mangrove · Decomposition · Biofilm · Microalgae · Epifauna · Shrimp

INTRODUCTION

The natural diet of penaeid post-larval (PL) shrimp has been characterized at different stages of the life cycle from nauplius, protozoa, mysis to PL (Dall et al. 1990, Rothlisberg 1998). Protozoa are typically herbivorous, while the mysis and PL become increasingly carnivorous. However, the PL are opportunistic feeders; for example, if diatoms dominate the environment, they will also dominate the diet of the shrimp (Preston et al. 1992). The composition and the diversity of microalgae may both directly and indirectly determine the nutritional quality of the natural food for PL. For instance, diatoms have a positive effect on growth and survival (Brown & Farmer 1994, Burford 1997, Ying et al. 2000) as they are rich in polyunsaturated fatty acids (PUFAs) (Parrish et al. 2000), which positively affect growth and metamorphosis in crustacean larvae, and survival and resilience to stressful conditions in juveniles (Read 1981,
Bell et al. 1984, Lavens & Sorgeloos 2000). Differences in the microalgal taxonomic composition rather than size or shape appear to be the critical factor (Rothlisberg 1998). *Penaeus semisulcatus* larvae achieve better growth when fed Chlorophyta of the species *Tetraselmis suecica* rich in proteins and essential fatty acids (D’Souza & Kelly 2000). The nontoxic blue-green alga *Trichodesmium* sp. was ingested by larvae of *P. merguiensis*, although it did not support growth or survival (Preston et al. 1998). Burford (1997) and Primavera (1998) suggested a variety of microalgal indicators, such as abundance and species diversity, as proxies for the actual food conditions and water quality (Casé et al. 2008) in shrimp culture ponds.

Penaeid PL also eat a wide variety of small invertebrates and plant material (Chen & Chen 1992, Rothlisberg 1998, Nunes & Parsons 2000). Prey in the natural diet of juvenile *P. esculentus* in seagrass beds include gastropod species, bivalves, crustaceans, and polychaetes. The epifauna is high in protein content and PUFA, and relatively low in carbohydrate (Dall et al. 1991). Polychaetes of the families Spionidae, Capitellidae, Eunicidae, Nereidae, Pilargidae, and Sebaliidae have been used in penaeid shrimp predation experiments, especially with *P. subtilis* (Nunes & Parsons 2000).

The importance of mangrove leaf litter as food for shrimp larvae may be linked to the associated periphytic biofilm. The present study defined the periphytic biofilm developing on the decomposing mangrove litter as a network of primary (microalgae) and secondary (epiflora) trophic levels mediated by a microbial loop (bacteria) (Azim & Wahab 2005). Stable isotope studies have demonstrated that giant tiger shrimp *P. monodon* feed on phytoplankton and epiphytic microalgae in a riverine mangrove forest in Guimaras (central Philippines; Primavera 1996). In the Ilamaraca estuary, Brazil, Schwamborn et al. (2002) found penaeid PL to have the highest carbon isotope delta values ($\delta^{13}$C) among the decapod larvae, indicating a negligible contribution of mangrove carbon to their nutrition.

PL feeding response may be influenced by the litter and biofilm palatability, which may be influenced negatively by the increasing level of tannins in the decomposing mangrove leaves (Rajendran & Kathiresan 2000). Direct observations on penaeid PL (Primavera 1996) and other benthic invertebrates (Bouillon et al. 2002) showed a reduced preference for mangrove-derived carbon compared to local and imported algal sources. Ronnback et al. (2002) observed that the distribution of juvenile *P. indicus* did not vary in abundance between sediments of low and high organic content. However, juvenile penaeid shrimp aggregate around mangrove litter colonized by bacteria (Rajendran & Kathiresan 2004, 2007). On average, 60 to 75% of nitrogen and 20 to 40% of carbon in highly decomposed mangrove leaf litter derive from heterotrophic bacteria and not from the remaining plant tissues (Tremblay & Benner 2006). In natural systems, bacteria may not occur as a separate functional or ecological entity but are present in combination with other microbiota forming benthic and epiphytic biofilms or as a flocculated mass in suspension (Burford et al. 2003, Azim & Wahab 2005).

The biological components of the biofilm and their nutritive value to PL in culture ponds have not been studied, although it is pivotal information to manage the mangrove litter in shrimp ponds. We hypothesized that the food value of mangrove leaf litter and the associated biofilm to PL depends on the assemblage and biomass of microalgae and epifauna as the decomposition of the mangrove litter progresses. We therefore investigated the major taxa of microalgae and invertebrate fauna colonizing the decomposing mangrove litter of *Rhizophora mucronata* and their application as diet for PL of *P. monodon*.

**MATERIALS AND METHODS**

**Study site**

The study was carried out in a mangrove forest and shrimp ponds at the mangrove conservation farm (Majaoni Silvofishery) located in Mtwapa Creek on the northern coast of Kenya (3° 57’S, 39° 42’ E). This site was used to incubate mangrove litter in ponds of 75 to 100 cm water depth. Mtwapa Creek is characterized by a reforested mangrove forest dominated by *Rhizophora mucronata*. PL of *Penaeus indicus* and *P. monodon* are commonly fished within this creek.

**Test organism**

PL of *P. monodon*, aged between 15 and 25 d from the time of hatching (PL 15–25) and hatched from the same brooder (cohort), were obtained from the Alphakrust shrimp hatchery at Mafia Island, Tanzania (www.alphaafrica.com). They were transported in plastic bags with a pure oxygen headspace for not more than 6 h and acclimatized to local conditions in a culture tank for 1 wk before the start of the ex-
periment. During the acclimatization period, the PL were fed on a shrimp larvae compound feed (CP) imported from India (Higashimaru zoea to PL 20 feed; crude protein over 52%, see www.aquafeed.com/documents/1254938830_1.pdf).

**Litter incubation and biofilm biomass**

Senescent mangrove leaves (hereafter referred to as litter), which had just turned yellow-brown and dropped from the trees, were dried in the shade to a constant weight and incubated in water hanging on 3 ropes (representing 3 replicates) in a shrimp pond, not stocked with shrimps, for a period of 8 wk. The incubation of the leaves was staggered by 1 wk to provide litter of the same age to the different treatments throughout the experiment. Given that the litter ropes were set up in triplicates, the set-up ultimately used 24 ropes to accommodate the 8 wk duration of litter incubation. However, to ensure that the biofilm fed to the shrimp was replenished every 4 d, the litter was harvested, in replicates and pooled, every 4 d within the same week to ensure that the week factor was maintained. The associated biofilm was also sampled by carefully scraping 3 leaves from each rope replicate. To estimate the biomass, the biofilm was dried in an oven at 70°C for 48 h and quantified as weight per unit leaf surface area.

**Microalgae and epifauna**

Litter was sampled weekly in triplicate by pooling 3 decomposing leaves per replicate. The biofilm was gently washed from the surface of the decomposing litter with a known volume of filtered sea water and preserved in 2% Lugol’s iodine solution (1:2 iodine: iodide: glacial acetic acid solution). Microalgae were classified and counted after first diluting each replicate sample 5 to 10 times; 5 sub-replicates of 0.02 ml were then sub-sampled and examined under an inverted microscope to determine microalgal abundance (cells l⁻¹).

To assess the microalgal biomass, mangrove leaf litter was sampled weekly in triplicate by pooling 3 leaves per sample. The periphytic biofilm was gently scraped from the surface of the litter with a known volume of filtered sea water and filtered through a glass fiber filter (GF/F, 0.45 µm mesh, 47 mm diameter). The surface area of both sides of the leaf was determined in order to convert the algal biomass from chlorophyll a (chl a) μg l⁻¹ to mg cm⁻² of the total leaf surface. Phytopigments were extracted from the biofilm after adding 10 ml 90% acetone to the lyophylized GF/F at 4°C in the dark, and the supernatant was analyzed for chl a according to the modified protocol of Granger & Lizumi (2001).

We examined the epifauna, which included both the meiofauna (metazoans that pass through a 1 mm sieve and are retained on a 38 µm sieve) and macrofauna (organisms retained on a 1 mm sieve). Litter was sampled weekly in triplicate by pooling 9 decomposing leaves per replicate in a plastic bag. Each group of decomposing leaves was immediately mixed with 8% MgCl₂ to shock the attached epifauna, thoroughly agitated, and subsequently sieved through 1 mm and 38 µm sieves. The sieved fauna was gently washed from the 38 µm sieve with a soft spray of filtered fresh water, preserved in 4% formalin, and stained with a few drops of 1% Rose Bengal solution. Epifauna was identified and counted at the highest possible taxon level using a binocular microscope and recorded as number of individuals per cm² of leaf surface.

**Species diversity and evenness**

The Shannon-Wiener index \( (H') \) and equitability index \( (EH) \) were used to estimate the microalgal and epifaunal community diversity and evenness based on the natural log (Shannon 1948), according to the following formulas:

\[
H' = - \sum_{i=1}^{n} p_i \ln p_i \quad (1)
\]

\[
EH = \frac{H'}{\ln S} \quad (2)
\]

where \( S \) is the total number of species in the community; \( p_i \) is the proportion of \( S \) made up by the \( i \)th species. Species equitability or evenness \( (EH) \) was interpreted within the range of 0 (dominance by a single species) to 1 (many species present in equal numbers).

**Feeding experiment**

PL 15–25 of \( P. \) monodon were starved for 24 h before the start of the experiment and then stocked for 16 d into 70 l circular laboratory tanks in triplicate at a density of 2 PL l⁻¹. The culture water was obtained from a reservoir tank and treated by first passing through a sand filter, then through a cartilage filter and disinfected by UV exposure. The cul-
ture water was replenished by adjusting the flow-through system to achieve a water exchange rate of 50% d⁻¹. Dissolved oxygen was maintained by continuously aerating water in both reservoir and experimental tanks through airstones. The following food treatments were tested: (1) mangrove leaf litter with biofilm at 1, 3, 4, 6, and 8 wk of decomposition; (2) commercial compound feed (CP); and (3) no food, as a control treatment. After removing the old batch of leaf litter from the tank, the PL were supplied every 4 d with new mangrove leaf litter, originating from the stock of leaves in the incubation pond, to maintain a litter density not exceeding 0.5 g l⁻¹ (Hai & Yakupitiyage 2005). The litter density was translated to supply an amount of dry biofilm biomass which would provide a feeding rate of 10% dry body weight. To estimate the dry weight, prior to the feeding experiment, 10 PL were sampled at random and dried in an oven at 70°C for 48 h and later weighed. This dry weight was extrapolated to the total PL stocked in the experimental tanks and used to calculate the daily proportion of the dry food. As all food sources in different treatments were provided in the same biomass, the same feeding rate of 10% dry body weight was used for the CP feed. Food items were offered twice a day (morning and afternoon) after removing the remaining food, and doubled every 3.5 d. PL were sampled weekly for specific growth rate (SGR, %) and at the end of the experiment for percentage survival (SR, %).

The growth and survival indices were calculated using the following formulae (Busacker et al. 1990),

\[ SGR\% = \frac{\ln(BW_t) - \ln(BW_0)}{T} \times 100 \]  

\[ SR\% = \frac{N_t}{N_0} \times 100 \]  

where SGR is the specific growth rate (% BW d⁻¹); BWₜ is the final body wet weight (g); BW₀ is the initial body wet weight (g); T is duration of the experiment (days); SR is the survival (in %); Nₜ is the number of shrimp collected at sampling time t; N₀ is the number of shrimp initially stocked. The wet weight was based on sampling groups of 70 individuals. The following water quality parameters were measured weekly: temperature, dissolved oxygen, pH, salinity, and total ammonium nitrogen (TAN). Temperature, dissolved oxygen, and pH were measured using meters, salinity was measured using a refractometer, and TAN was analyzed in the laboratory according to Eaton et al. (2005).

Data analysis

Univariate statistical analyses (ANOVA) were conducted with Statistica 7.0 software. All data were checked for normality and variance homogeneity requirements for parametric analysis. Data which did not meet normality requirements after being transformed were analyzed non-parametrically (Kruskal-Wallis ANOVA and median test). Multidimensional scaling (MDS) and analysis of similarity (ANOSIM) were used to compare similarity in the distribution of target parameters using Primer 6.0 software (Clarke & Gorley 2006).

RESULTS

Biofilm and microalgal biomass

Biofilm increased in both quantitative and chl a estimated biomass throughout the period of litter decomposition. The highest biofilm biomass was in the litter decomposed for 8 wk, which was significantly different from the biomass on leaf litter decomposed for less than 6 wk (1-way ANOVA: F₆,₁₄ = 145.43, p < 0.05 and Tukey post hoc p < 0.05). Microalgal biomass on the litter decomposed for 1 and 2 wk (Tukey post hoc p < 0.05). Biofilm on litter had significantly higher biomass of microalgae (294.07 ± 0.79 µg l⁻¹ chl a) than the pond water (0.79 ± 0.83 µg l⁻¹ chl a) throughout the incubation period (p < 0.05; see Fig. 2a).

Taxonomic composition and species diversity of microalgae

Among the microalgae, 32 species were identified including 14 diatoms, 8 cyanobacteria, 7 dinoflagellates, 1 coccolith, and 1 flagellate species. Compared to diatoms and cyanobacteria, dinoflagellates, coccoliths, and flagellates had a low cover of <10% during the entire period of litter decomposition. Diatoms dominated the biofilm during the first 6 wk of litter decomposition, with a relative cover of 88 and 99% during Weeks 3 and 4, respectively. Among the diatom species, Navicula sp. dominated the first 4 wk of decomposition, reaching a maximum proportional cover of 90.8% during Week 3, whereas Nitzschia sp. dominated the later period of decomposition with a maximum cover between 64.2 and 78.7% during Weeks 5 and 6. Cyanobacteria exhibited lower abun-
dance during the early period of decomposition but recorded a sharp increase during Week 8, with a percentage cover of 61%, which exceeded the cover of diatoms (38%). Among the cyanobacteria, *Microcystis* sp. dominated the 8 wk old litter with a proportional cover of 55.3%. The lowest microalgal diversity was recorded during Weeks 3, 4, and 5 (Table 1). The microalgal community differed significantly in the different weeks of litter decomposition (ANOSIM: R = 0.602; p < 0.05; Fig. 1a).

### Microalgal species abundance

Diatoms varied significantly in abundance during the 8 wk of litter decomposition (Kruskal-Wallis test, $H_{6, 21} = 13.26$, $p < 0.05$), with a maximum abundance in Week 3 due to the high densities of *Navicula* sp. and in Week 6 due to *Nitzschia* sp. (Fig. 2c). Cyanobacteria exhibited lower abundance during the early period of litter decomposition but recorded a sharp increase during Week 8, exceeding the abundance of diatoms. *Microcystis* sp. dominated the cyanobacteria (Fig. 2d). *Peridinium* sp. dominated the dinoflagellates during Week 6 (Fig. 2e).

### Epifauna diversity and abundance

We identified 19 major taxa of epifauna on the epiphytic biofilm (Table 2). The epifaunal species colonizing the biofilm differed significantly in the different weeks of litter decomposition (ANOSIM: R = 0.887; $p < 0.05$; Fig. 1b). Copepoda dominated the epifauna, colonizing the biofilm during the first 3 wk of litter decomposition with a proportional percentage cover

<table>
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<tr>
<th>Taxon</th>
<th>Week 1</th>
<th>Week 2</th>
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<td>78.7</td>
<td>64.2</td>
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<td>89.2</td>
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<td>51.8</td>
<td>73.3</td>
<td>90.8</td>
<td>7.2</td>
<td>23.7</td>
<td>5.7</td>
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<td><em>Thallasionema nitzchoides</em></td>
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<td><em>Halsea</em> sp.</td>
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<td><em>Choanoflagellate</em> sp.</td>
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<td>Diversity index</td>
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<td>1.054</td>
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<td>0.445</td>
<td>0.861</td>
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<td>Species evenness</td>
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<td>0.108</td>
<td>0.036</td>
<td>0.041</td>
<td>0.079</td>
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Fig. 1. Multidimensional scale plot of similarity in (a) microalgae and (b) epifauna colonizing biofilm at the different stages of mangrove leaf litter decomposition. Symbols refer to duration of litter decomposition in weeks.

Fig. 2. Biomass and assemblage of microalgae in biofilm developing on *Rhizophora mucronata* litter at different stages of decomposition. (a) Biomass of microalgae in the pond water and in biofilm. Absolute abundance of (b) microalgal communities, (c) diatoms, (d) cyanobacteria, (e) dinoflagellates, and (f) rare microalgal communities.
between 59 and 68%, while Polychaeta dominated during Weeks 4 and 5, with a proportional percentage cover of 90 to 93%; at this stage, the development of biofilm on the litter had the lowest diversity of epifauna and species evenness (Table 2). Concomitantly, the maximum abundance of epifauna was recorded. However, abundance declined as the litter decomposed further towards Weeks 6 and 8 (ANOVA, \( F_{6,14} = 93.26, p < 0.05; \) Tukey post hoc \( p < 0.05 \)). Epifauna in the 4 and 5 wk old biofilm was dominated by meiofauna of the lower size class (collected in 38–250 µm sieves) compared to the larger meiofauna (retained in 250 µm – 1 mm sieves) (Fig. 3a). Nematoda were dominant (representing 40% of the community) in the biofilm developing on litter that had decomposed for 8 wk (Fig. 3b).

**Shrimp performance**

**Growth rate (SGR)**

PL foraging on the biofilm at the different stages of litter decomposition recorded significantly lower SGR, compared to PL that were fed with compound feed (CP; ANOVA \( F_{6,14} = 30.423, p < 0.05; \) Tukey post hoc \( p < 0.05 \)). However, among the treatments receiving the litter, PL foraging on the biofilm on 4 wk old decomposed litter showed overall better performance (Tukey post hoc \( p < 0.05 \)). PL foraging on the biofilm growing on litter that had decomposed for 1, 3, and 6 wk showed reduced growth and in some cases negative SGR. Although PL foraging on the litter that had decomposed for 8 wk showed positive growth, their SGR was not significantly different from the treatments recording negative growth rates and PL in the starvation treatments (Tukey post hoc \( p < 0.05; \) Fig. 4).

**Survival (SR)**

PL fed compound feed (CP) recorded the highest SR (ANOVA, \( F_{6,14} = 73.85, p < 0.05; \) Tukey post hoc \( p < 0.05 \)). Among the PL foraging on the biofilm on the decomposing litter, PL foraging on the 4 wk old litter recorded the highest mean SR, although they did not differ statistically from PL foraging on the litter decomposed for 1, 3, and 6 wk (Tukey post hoc: \( p > 0.05 \)). However, pairwise comparisons revealed significantly lower SR in PL foraging on the litter decomposed for 8 wk (Tukey post hoc \( p < 0.05 \)). PL in the starvation treatments (NF) recorded the lowest SR of 8.4 ± 0.9% (Tukey post hoc \( p < 0.05 \)), although they did not differ significantly from the PL foraging on the litter decomposed for 8 wk (Tukey post hoc: \( p > 0.05; \) Fig. 4).

**Table 2. Proportional assemblage (\( P_i \) in Eq. 1) of epifauna on biofilm developing on *Rhizophora mucronata* litter at different stages of decomposition. All values, except diversity index and species evenness, are given in percentages**

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DISCUSSION

Microalgae

In the first 6 wk of the experiment, there was a clear dominance of diatoms but also a shift in the community composition, with *Navicula* sp. dominating the leaf litter until Week 4 of decomposition. After Week 4, *Navicula* sp. was replaced by *Nitzschia* sp. at Weeks 5, 6, and 8. The reason for this shift in diatom species is not understood, although its implication to the diet of PL shrimp could be considerable. *Nitzschia* sp. has been reported to be richer in proteins (Brown & Jeffrey 1995) and therefore could present a better source of proteins to the immediate grazers.

Microalgal succession from diatoms to cyanobacteria was prominent at the later stages of the leaf litter decomposition. Cyanobacterial growth can be promoted by high nutrient input (Amado & Monserrat 2010). The pond water showed a gradual slight increase in TAN after Week 4 of litter incubation. Hence the success of cyanobacteria on the litter may have been a micro-scale bloom triggered by an increase in nutrients leached from the decomposed litter such as the ammonified particulate and dissolved organic nitrogen (Alongi et al. 2000) and phosphorus (Lin & Sternberg 2007). In addition, the first weeks of biofilm development were characterized by a strong growth of copepods which could feed on the diatoms. Diatoms have a C:N ratio of about 5, and this could influence carbon and nitrogen contents of copepod fecal pellets (Morales 1987, Siuda & Dam 2010). Copepods that were grazing on diatoms could have released a lot of nutrients, especially ammonium. This may have stimulated the cyanobacterial blooms.

Apart from the increase in cyanobacteria, there was a general decline in the other microalgal species belonging to the diatoms, dinoflagellates, and flagellates during the advanced stages of litter decomposition. Microalgae reached a stationary phase between Weeks 4 and 6 of decomposition and progressed to the death phase during Week 8. The decline of the microalgae may result from grazing by epifauna (Coman et al. 2003). However, the similar decline in epifauna after Week 5 (Fig. 3a) suggests that grazing may not have been the only reason for the decline.
Decomposing mangrove litter leaches a phenolic compound, viz. tannin (Rajendran & Kathiresan 2000, Hernes et al. 2001, Lin et al. 2007). Tannin concentrations above 18 mg l⁻¹ inhibit the growth of the microalgae *Skeletonema* sp. and *Dunaliella* sp. in the Gulf of Mexico (Herrera-Silveira & Ramirez-Ramirez 1996). Tannins leached from the decomposing litter could have had a cumulative negative effect on certain species of microalgae and in this way may have provided space and reduced competition for the proliferation of cyanobacteria.

**Epifauna**

Epifauna identified on the biofilm included several juvenile stages of macroinvertebrates such as gastropods, bivalves, crustaceans, polychaetes, insects, and annelid worms which are potential food for PL (Tacon 1996, Rothlisberg 1998). In our study, adult copepods were the major component of the biofilm during the first 3 wk of litter decomposition and mainly consisted of representatives of the order Harpacticoida, which occur in sediments but are also very diverse in epiphytic communities such as on seagrass leaves (De Troch et al. 2003). Juvenile stages of copepods (nauplii larvae), egg cocoons of polychaetes and oligochaetes, and bivalve trochophore and veliger larvae, although enumerated together with the adult species, were possible contributors to the increase in epifauna in the size range of 38 to 250 µm which dominated the 4 and 5 wk old biofilm. The increase in juvenile stages suggests recruitment at this phase of the decomposition process. The increase in polychaetes after Week 3 suggests that at this stage of litter decomposition, polychaetes increasingly contribute to the shrimp food (Fig. 3b). According to our previous study (Gatune et al. 2012), litter decomposed for 3 and 4 wk supported a climax microbial abundance and essential fatty acids which may nutritionally influence a healthy assemblage of epifauna. At this stage of litter decomposition, there is also a large pool of nitrogen, amino sugars, and amino acids (Rajendran & Kathiresan 2000) which would nourish the primary trophic levels occupied by bacteria and microalgae. These microbti are potential food sources for the epifauna (Woitchik et al. 1997). According to our previous study (Gatune et al. 2012), PL that were stocked in litter were isotopically closer to periphytic biofilm associated with mangrove litter in comparison to PL from bare mangrove zones and open creek habitats. Although the present experiment did not include an isotopic assessment to prove the actual uptake of the prey items by PL, the observation from Gatune et al. (2012) justifies this assumption.

The decline in the abundance of epifauna after the recruitment stage (Weeks 4 and 5) could be due to the reduced survival of juveniles advancing into the adult stages. Reduced supply of quality food because of the decline and aging of diatoms concomitant with an increase of non-palatable cyanobacteria could be the main reasons. Litter decomposed for more than 4 wk could be characterized by low levels of reduced amino sugars and amino acids (Rajendran & Kathiresan 2000, Tremblay & Benner 2006). On the other hand, tannins leached from decomposing litter can form complexes with proteins and enzymes leading to antimicrobial properties (Lin et al. 2007) and immobilize nutrients such as nitrogen and phosphorus (Lin & Sternberg 2007). Such chemical modifications can make decomposing mangrove litter insufficient to support a climax community of microbiota which could serve as an important food source for the epifauna (De Mesel et al. 2004, Pascal et al. 2008a,b, Wieltschnig et al. 2008). Tannins can also impart a direct negative effect on epifauna such as meiofauna (Alongi 1987) and macrofauna (Dittmann 2001).

**Performance of post-larval shrimp foraging on biofilm**

PL foraging on biofilm of litter decomposed for a period of less than 3 wk appeared to experience retarded growth. This scenario was also repeated in the PL foraging on 6 wk old litter. A notable improvement in growth and survival was observed in PL foraging on 4 wk old litter. Growth performance of PL can be predicted from quality and/or palatability of food which may be influenced by artificial formulation or favorable environmental conditions. For instance, PL fed CP feed had much improved growth and survival. In the present case, the results of the PL fed with biofilm could not have been affected by underfeeding, since remnant biofilm food was observed. Furthermore, the differences observed especially in SGRs in PL fed with biofilm were clearly separated statistically.

The CP feed, which acted as a reference feed, was specially formulated to provide target nutrients which are essential for improved growth performance (Hertrampf & Piedad-Pascual 2000, Lavens & Sorgeloos 2000). The use of natural diets in this study did not necessarily provide a straightforward solution towards improving the commercial diet. However, in
Potential effect of microalgae and epifauna on post-larval shrimp growth and survival

Apart from the diatoms Chaetoceros sp. and Thalassiosira sp. which occurred in small proportions in the biofilm developed on the litter decomposed for 1 and 5 wk (Table 1), the species of microalgae constituting the biofilm grown in these experiments differed from the microalgal species commonly used in the commercial culture of shrimp (e.g. diatoms Chaetoceros sp., Skeletonema sp., and Thalassiosira sp.). However, non-diatom microalgae such as prymnesiophytes (Isochrysis sp., Pavlova sp.), chlorophytes (Tetraselmis sp., Dunaliella sp., Nannochloris sp.), eustigmatophytes (Nannochloropsis sp.), and cryptophytes (Chroomonas sp.) are also commonly used in aquaculture (Brown & Miller 1992, Brown & Farmer 1994, Borowitzka 1997). In a microcosm study by Brown & Jeffrey (1995), Nitzschia closterium was found to be richest in proteins (38% content) compared to 6 other diatoms of the genera Navicula, Skeletonema, Lauderia, and Cylindrotheca. The ecological role of the dominating diatoms on the less decomposed litter cannot be ignored given the importance of diatoms in the energy transfer to the consumer trophic levels. For instance, the characteristic fatty acid composition of diatoms is readily distinguishable from those of other microalgal groups (Ying et al. 2000). Diatoms are typically rich in PUFAs such as eicosapentaenoic acid (EPA, 20:5ω3) (Parrish et al. 2000). Certain fatty acids such as the PUFAs, highly unsaturated fatty acids, phospholipids, and sterols affect reproduction, growth, metamorphosis of crustacean larvae to juveniles, survival, and resilience to stressful conditions (Read 1981, Bell et al. 1984, Lavens & Sorgeloos 2000). The nutritional quality of diatoms may deteriorate at the advanced age. While comparing 3 growth phases in 4 marine diatoms, among them N. closterium and Chaetoceros gracilis, Liang & Mai (2005) observed that saturated and monounsaturated fatty acids increased while the PUFAs decreased with culture age.

PL foraging on litter decomposed for 4 wk may also have benefited from the ample food supply from the increased recruitment of epifauna. The low species diversity and evenness of epifauna and the high abundance of copepods and polychaetes observed at this stage (Fig. 3b) emphasize the ecological importance. Zooplankton and epibenthos contribute to the nutrition of PL in aquaculture installations (Chen & Chen 1992). The establishment of an abundant assemblage of epifauna is therefore an important prerequisite to stocking (Tacon 1996, Coman et al. 2003) and can be part of the supplementary feeds that can directly be consumed by cultured shrimp. Such natural food items include insects, annelid worms, crustaceans, and mollusks (Tacon 1996). Meiofauna and small macrofauna can constitute a major food source for juvenile fish and shrimp (Dittmann 2001). Rothlisberg (1998) and Hill & Wassenberg (1987) observed that the diet of juvenile and adult shrimp consisted of a wide variety of zoobenthos and macroinvertebrates (gastropods, bivalves, crustaceans, and polychaetes).

PL foraging on the 6 and 8 wk old biofilm may also have been adversely affected by the replacement of diatoms by cyanobacteria. In the present study, the cyanobacteria Microcystis sp. was dominant; it produces a potent hepatotoxin, a microcystin that is also produced by a number of planktonic cyanobacterial genera such as Anabeana, Anabaenopsis, Nostoc, and Planktothrix (Oscillatoria) (Amado & Monserrat 2010) and other bioactive metabolites with potential to degrade the nutritional status of aquaculture species (inhibitors of proteases and grazer deterrements) (Smith et al. 2008). The reduced epifaunal abundance could also have caused reduced growth and survival of PL foraging in the litter decomposed for more than 4 wk.

CONCLUSION

The present study underlines the ecological importance of the periphytic biofilm developed on the mangrove litter decomposed for 4 wk as an important
food and stage of development in supporting PL. Biofilm of high food quality is fully developed and is dominated by diatoms, polychaetes, harpacticoid copepods, and oligochaetes. The ecological function of decomposing litter in providing quality natural food to PL is limited if retained in the pond water beyond a period of 4 wk. This is illustrated by the collapse of the epifauna and subsequent colonization by cyanobacteria of low nutritional quality. This needs to be considered when controlling the residence time of decomposing mangrove litter in shrimp ponds.

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LITERATURE CITED

> Chen YLL, Chen HY (1992) Juvenile Penaeus monodon as effective zooplankton predators. Aquaculture 103:35−44


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