



Observations on the laboratory culture of the harpacticoid copepod *Euterpina acutifrons* (Dana, 1847) using different diets

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ABSTRACT

Hatching success, nauplii production, maturation and survival of the harpacticoid copepod *Euterpina acutifrons* (Dana, 1847) fed on mono/mixed culture of *Isochrysis galbana*, *Tetraselmis gracilis*, *Chaetoceros calcitrans* and *Chlorella marina* were studied. The time taken for *E. acutifrons* to mature was the lowest with a diet of *I. galbana* followed by *C. calcitrans*. Percentage of survival was highest with a mixed diet of 75% *I. galbana* and 25% *C. calcitrans* (84%) followed by *T. gracilis* (68%). The study also revealed that both feed type and concentration played significant role in production and survival of *E. acutifrons*. Mixed diet of *I. galbana* and *C. calcitrans* showed maximum adult production followed by a diet of *T. gracilis*. The mean values of all the three stages of the copepod viz., adults, ovigerous females and copepodite-nauplii were found much greater in the treatment with replacement of the culture medium and feed. Mass culture using different organic feeds was also carried out and it was found that mean growth in terms of adult stage production was found to be higher when fed with rice bran followed by groundnut oil cake (GOC) and the least in cow dung treatment. In rice bran treatment, copepodites showed higher production rate when compared to the other two feeds. Mean growth in terms of nauplii production was also higher in rice bran fed group followed by GOC fed group and the least in cow dung treatment.

Keywords: Algal feeds, Artificial feed, Copepod culture, Harpacticoid copepod, *Euterpina acutifrons*, Mass culture, Population density

Introduction

Production of marine copepods in the hatchery is important for solving feed related problems in the rearing of marine fin fish larvae. Copepods have been used for marine fish species, which normally do not accept artificial feeds. Therefore in recent years, much attention has been focused on mass culture of copepods under laboratory and hatchery conditions. Attempts on mass culture of different species of copepods have been carried out worldwide (Stottrup and Norsker, 1997; Nanton and Castell, 1998; Schipp *et al.*, 1999; Payne and Rippingale, 2000; Stottrup, 2000) in view of their worldwide distribution, abundance, high rate of reproduction, short generation time, high nutritive value and smaller naupliar stages. Gopakumar *et al.* (2008) gave an account on mixed culture of the harpacticoid species *Euterpina acutifrons* (Dana, 1847) and the calanoid copepod *Psuedodiaptomus sericaudatus* as live feed for ornamental fish larval rearing.

There is great interest in culturing harpacticoid copepods since they have smaller naupliar stages, high reproductive potential, fast population growth, flexible diet and tolerance to wide ranges of environmental factors

such as temperature and salinity (Sun and Fleeger, 1995; Stottrup and Norsker, 1997). Moreover, harpacticoid copepods provide a broad spectrum of prey sizes suitable for developing fish larvae, help to keep the walls of the fish larval rearing tanks clean by grazing on algae and debris (Delbare *et al.*, 1996) and are also rich in essential fatty acids that are extremely important for larval fish survival and growth (Stottrup and Norsker, 1997). Nassogne (1970) and Kraul (1990) have reported on culture of *E. acutifrons*. The culture conditions for harpacticoids are less demanding than those for calanoids. Filtered seawater may be used and a whole range of inert food is acceptable to many harpacticoid species. This simplifies the culture method and eliminates the need for culture of phytoplankton.

E. acutifrons is a holoplanktonic suspension feeding copepod which is a common species in coastal waters off Vizhinjam. It is a hardy species and easy to maintain in laboratory conditions. In the present study, an attempt was made to investigate the diets that will improve production of *E. acutifrons* under laboratory conditions using different algal feeds as well as selected inert feeds.

Materials and methods

The marine harpacticoid copepod *E. acutifrons* was collected from Vizhinjam Bay. Copepods were cultured on marine algae with continuous aeration at room temperature, at 35 ppt salinity. Four algal species each with specific nutritional characteristics, *viz.*, *Isochrysis galbana*, *Tetraselmis gracilis*, *Chaetoceros calcitrans* and *Chlorella marina* in monoculture as well as in combinations *i.e.*, *I. galbana* plus *C. calcitrans*, *T. gracilis* plus *I. galbana* and *C. calcitrans* plus *I. galbana* were fed to copepods to elucidate the ideal algal feed for culture. Algal cultures at the required density were prepared by counting samples using a haemocytometer, followed by dilution using enriched seawater. Initial trials were carried out to evaluate development and survival using different algae to find out the most suitable feed. Ten millilitre capacity glass vials were used for the experiment to test different algal concentrations *viz.*, 1×10^6 , 2×10^6 , 3×10^6 , 5×10^6 and 6×10^6 cells ml^{-1} . Newly hatched nauplii were daily picked out of the experimental vials with a Pasteur pipette and counted. The animals were transferred to new vials with appropriate algal concentration on every third day. Experiments were carried out until next generation of ovigerous copepods were obtained.

Mass culture experiments of copepods were done employing the ideal concentration arrived at during the initial vial experiment for each microalga. Ovigerous females were stocked in 1000 ml glass beakers containing the following algae: Group I - *Chlorella marina*; Group II - *Chaetoceros calcitrans*; Group III - *Isochrysis galbana*; Group IV - *Tetraselmis gracilis* and Group V - Mixed culture of *I. galbana* and *C. calcitrans*.

Twenty ovigerous females were stocked in glass beakers containing 1 litre of the respective microalgal culture. Three beakers were assigned to each diet group. Algal culture was separately maintained for feeding the copepods and desired concentration was obtained through centrifugation and dilution. Gentle aeration was maintained *via* an open glass capillary tube. Two sets of experiments were conducted, in the first set media and algae were not changed during the experimental period. In the second set, 50% of medium was replaced and filled with water containing fresh algae on alternate days. Counts were taken on alternate days from 5 ml sample collected from the well stirred beaker. Copepod population density (no. per 1000 ml) was noted and counts were taken separately for nauplii, copepodites, adults and ovigerous females.

Another set of experiment was conducted in 5 litre transparent plastic buckets (in triplicates) using organic material as source of feed. Cow dung, groundnut oil

cake (GOC) and rice bran in combination with urea and superphosphate were used to prepare culture media. For each 100 g of prime component, 0.5 g urea and 2 g superphosphate were added. Dry cow dung was powdered and again dried in oven for 24 h. This was used for preparing the medium in filtered seawater. Rice bran and GOC were collected from the local market and dried in the oven and a suspension was prepared by soaking in filtered seawater. The culture media were inoculated with known numbers of *E. acutifrons*. The culture was maintained for a period of four weeks with continuous aeration. The rearing water quality parameters were maintained at optimal levels, *i.e.*, temperature - $28 \pm 0.5^\circ\text{C}$, salinity - 35‰ and pH - 8 to 8.2. In all the culture groups, copepod count (no. per 1000 ml) was taken at weekly intervals.

Results

Results of the study indicated that the feed type and concentration played a significant role in the survival of *E. acutifrons* (Table 1). The time taken for the animals to mature was significantly shorter ($p < 0.01$) with a diet of *I. galbana* (6 days) at a concentration of 1×10^6 cells ml^{-1} and with mixed diet of *I. galbana* and *C. calcitrans* (7 days). Time to maturity was comparatively higher with *C. calcitrans* (9 days) at an optimum cell concentration of 2×10^6 cells ml^{-1} and *C. marina* (9 days) at a cell concentration of 6×10^6 cells ml^{-1} . Copepods fed with *T. gracilis* took the longest time to mature (10 days) at a concentration of 1×10^6 cells ml^{-1} , but the best survival rate was observed in this feed. The highest percentage of survival was observed with mixed diet of 75% *I. galbana* and 25% *C. calcitrans* (84%) on the 7th day followed by *T. gracilis* (68%) on 10th day, a mixed diet of 75% *C. calcitrans* and *I. galbana* 25% (61%) on 7th day, *C. calcitrans* (50%) on 9th day, *I. galbana* (41%) on 6th day and *C. marina* (27%) on 9th day (Table. 1).

The mass culture experiments conducted in 1 litre glass beakers showed that the food for copepod growth is stage-specific *i.e.*, requirements of feed differ with developmental stage. In the first set, without replacement of culture medium and algae, group V with mixed diet of *I. galbana* and *C. calcitrans* showed maximum adult production followed by group IV diet of *T. gracilis* (Table 1). Group II showed least performance with respect to adult production and growth. Analysis of variance along with *post-hoc* analysis showed highly significant difference among different treatments ($p < 0.01$). Performance of group IV and V did not differ significantly. Similarly group I and II also performed equally with respect to adult production and growth. In the mass culture experiment without replacement of culture medium, maximum density for adults was observed in group IV (119.67 nos. l^{-1}) and minimum in group II (39.33 nos. l^{-1}).

Table 1. Hatching success, nauplii production, time to maturity and survival of *Euterpina acutifrons* fed on different algal diets

| Algal species | Algal concentration (x 10 ⁶ cells ml ⁻¹) | Number of nauplii (Mean±SD) | Number of adults with eggs (Mean± SD) | Time to maturity (days) | % Survival |
|--|---|-----------------------------|---------------------------------------|-------------------------|------------|
| <i>Chaetoceros calcitrans</i> | 1 | 18 ± 2 | 3 ± 2 | 14 | 17 ± 0 |
| | 2 | 32 ± 4 | 16 ± 5 | 09 | 50 ± 1 |
| | 3 | 18 ± 3 | 2 ± 1 | 12 | 11 ± 1 |
| <i>Isochrysis galbana</i> | 0.5 | 10 ± 1 | 0 | - | 0 |
| | 1 | 34 ± 1 | 14 ± 2 | 06 | 41 ± 1 |
| | 2 | 26 ± 4 | 0 | - | 0 |
| <i>Tetraselmis gracilis</i> | 3 | 13 ± 2 | 0 | - | 0 |
| | 0.5 | 10 ± 1 | 0 | - | 0 |
| | 1 | 25 ± 2 | 17 ± 3 | 10 | 68 ± 1 |
| <i>Chlorella marina</i> | 2 | 17 ± 3 | 0 | - | 0 |
| | 3 | 13 ± 2 | 0 | - | 0 |
| | 5 | 11 ± 1 | 0 | - | 0 |
| <i>C. calcitrans</i> (25 %) + <i>I. galbana</i> (75 %) | 6 | 11 ± 2 | 3 ± 1 | 09 | 27 ± 1 |
| | 1 + 2 | 37 ± 3 | 31 ± 3 | 07 | 84 ± 0 |
| <i>C. calcitrans</i> (75 %) + <i>I. galbana</i> (25 %) | 1 + 2 | 23 ± 3 | 14 ± 4 | 07 | 61 ± 1 |
| <i>C. calcitrans</i> (50 %) + <i>I. galbana</i> (50 %) | 1 + 2 | 33 ± 3 | 0 | - | 0 |

Table 2. The maximum mean density of ovigerous females was obtained in group II (30.33 nos. l⁻¹) and minimum in group III. Maximum density of the early stages such as nauplii and copepodites were recorded in Group III (136.33 nos. l⁻¹) and minimum in group I (49.33 nos. l⁻¹).

In the mass culture experiment with replacement of culture medium, group V with mixed diet of *I. galbana* and *C. calcitrans* showed maximum production followed closely by group IV and III (Table 3) as observed without replacement of culture medium. Lowest density

Table 2. Production of different stages of *E. acutifrons* in mass culture experiment using different algal diets, without replacement of culture medium

| Stage | Group | Density (no. l ⁻¹) Mean± SD |
|---------------------|-----------------------------|--|
| Adults | I (<i>C. marina</i>) | 40.33 ± 5.50 ^a |
| | II (<i>C. calcitrans</i>) | 39.33 ± 7.00 ^a |
| | III (<i>I. galbana</i>) | 79.67 ± 21.50 ^b |
| | IV (<i>T. gracilis</i>) | 119.67 ± 12.38 ^c |
| | V (Mixed) | 131.33 ± 14.76 ^c |
| Ovigerous females | I (<i>C. marina</i>) | 13.00 ± 6.25 ^a |
| | II (<i>C. calcitrans</i>) | 30.33 ± 13.00 ^b |
| | III (<i>I. galbana</i>) | 13.33 ± 2.18 ^a |
| | IV (<i>T. gracilis</i>) | 14.00 ± 1.50 ^a |
| | V (Mixed) | 24.67 ± 2.50 ^b |
| Copepodites+Nauplii | I (<i>C. marina</i>) | 49.33 ± 31.43 ^a |
| | II (<i>C. calcitrans</i>) | 62.00 ± 42.00 ^a |
| | III (<i>I. galbana</i>) | 136.33 ± 15.98 ^b |
| | IV (<i>T. gracilis</i>) | 122.33 ± 10.58 ^b |
| | V (Mixed) | 129.00 ± 15.87 ^b |

Values bearing same superscripts within each stage do not differ significantly (p<0.01)

Table 3. Production of different stages of *E. acutifrons* in mass culture experiment using different algal diets, with replacement of culture medium

| Stage | Group | Density (no. l ⁻¹) Mean±SD |
|---------------------|-----------------------------|---|
| Adults | I (<i>C. marina</i>) | 68.00 ± 6.54 ^a |
| | II (<i>C. calcitrans</i>) | 74.67 ± 5.89 ^a |
| | III (<i>I. galbana</i>) | 159.67 ± 21.50 ^b |
| | IV (<i>T. gracilis</i>) | 164.00 ± 15.32 ^b |
| | V (Mixed) | 187.00 ± 4.33 ^c |
| Ovigerous females | I (<i>C. marina</i>) | 19.33 ± 2.18 ^a |
| | II (<i>C. calcitrans</i>) | 45.00 ± 4.82 ^c |
| | III (<i>I. galbana</i>) | 28.67 ± 7.86 ^b |
| | IV (<i>T. gracilis</i>) | 18.33 ± 2.78 ^a |
| | V (Mixed) | 57.00 ± 3.12 ^d |
| Copepodites+Nauplii | I (<i>C. marina</i>) | 73.67 ± 23.52 ^a |
| | II (<i>C. calcitrans</i>) | 98.67 ± 20.35 ^b |
| | III (<i>I. galbana</i>) | 221.00 ± 30.01 ^d |
| | IV (<i>T. gracilis</i>) | 154.67 ± 7.86 ^c |
| | V (Mixed) | 158.00 ± 12.22 ^c |

Values bearing same superscripts within each stage do not differ significantly (p<0.01)

was encountered in group I and II. Analysis of variance along with *post-hoc* analysis showed highly significant difference among different treatments (p<0.01). Group III and IV performed almost equally with not much difference. Similarly group I and II also performed equally with respect to adult production. Group V as well as group II showed maximum production and growth with respect to ovigerous stage in the first set without replacement of feed. Among the others, group III gave better output than I and IV. In the second set with replacement of feed, production of copepodites and nauplii was maximum in group III

followed by group V and IV. All these three groups did not differ significantly as indicated by Duncan's multiple range *post-hoc* analysis. Group I and II showed poor performance compared to other three groups with much low values (Table 2 and 3).

The paired samples analysis was done group-wise in both treatments. In all the groups, mean density of all the three stages of animals *viz.*, the adults, ovigerous females and copepodites+nauplii were found to be significantly higher ($p < 0.01$) in treatment with replacement of culture media and feed. The algal concentration was found to have more significant effect ($p < 0.01$) on the mean density of different stages of copepods.

In the mass culture experiment employing organic inert feeds, adult stage production was found to be significantly higher ($p < 0.01$) in treatment III (rice bran fed copepods) followed by treatment II (GOC fed copepods) and least in cow dung treatment (Table 4). The adults registered a similar growth pattern in all the treatment groups during the first week. During the second week, a sharp decline was noticed in treatment II and III. In treatment III, further growth was observed in the second week which then started to decline. Ovigerous female production was found higher in rice bran fed population followed by GOC fed copepods in treatment II and least in cow dung treatment (Table 4). Production of copepodites also registered almost the same production pattern as of adults. In rice bran treatment, copepodites showed a higher production rate when compared to the other two feeds (Table 4). But unlike in adults, a sharp decline was noticed initially in all the three diets which thereafter registered a steady growth. Nauplii production was also found to be higher in treatment III fed with rice bran followed by GOC fed treatment II, with least production in cow dung treatment (Table 4). The nauplii registered

similar growth pattern during the first week except in cow dung treatment, which showed a steady stage. Then it declined and again after third week registered a slight increase in number. In treatment II and III, after first week, a steady decline was noticed.

Discussion

Feed type and concentration play vital role in the reproductive responses of copepods. The amount of food supplied directly influences their reproductive rate. It is evident that food type, size and concentration can independently as well as in combination influence the reproductive potential and hatching, growth rate, development and survival time to maturity of the copepod population. In the first experiment in vials (10 ml) hatching success, nauplii production, time to maturity and percentage of survival of *E. acutifrons* mainly depended on diet. The hatching success and percentage survival were highest with a mixed diet of 75% *I. galbana* and 25% *C. calcitrans*. The observations by Shrivastava *et al.* (1999) is in agreement with our findings, wherein they concluded that the combined feed of *I. galbana* and *Chaetoceros* sp. to be most satisfactory for the estuarine calanoid copepod *Heliodiaptomus cinctus*. Nassogne (1970) reported that the best results were obtained when *E. acutifrons* was fed with a mixture of algal species. In the present study, the time to maturity for *E. acutifrons*, was shortest with a diet of *I. galbana*, followed by *C. calcitrans*, *C. marina* and *T. gracilis*. Similarly Payne and Rippingale (2000) recorded highest egg production in the calanoid copepod *Gladioferens imparipes* on a diet of *I. galbana* followed by *C. calcitrans*. As in the present work, Stottrup and Jensen (1990) recorded higher egg production in *Acartia tonsa* on a diet of *I. galbana*. The fatty acid profile of algae used in copepod diets (Payne and Rippingale, 2000) showed that *I. galbana*

Table 4. Production (Mean \pm SD) of different stages of *E. acutifrons* in mass culture experiment using different organic feed inputs

| Stage | Treatment | Day 7 | Day 14 | Day 21 | Day 28 |
|-------------------|-----------|-------------------------------|-------------------------------|-------------------------------|------------------------------|
| Ovigerous females | I | 1.00 \pm 0 ^b | 1.50 \pm 0.55 ^a | 1.00 \pm 0 ^a | 1.00 \pm 0 ^b |
| | II | 0.50 \pm 0.55 ^a | 1.00 \pm 0 ^a | 1.00 \pm 0 ^a | 0.50 \pm 0.55 ^a |
| | III | 1.00 \pm 0 ^b | 1.50 \pm 0.55 ^a | 1.50 \pm 0.55 ^b | 1.00 \pm 0 ^b |
| Adults | I | 6.50 \pm 1.64 ^a | 8.00 \pm 1.1 ^a | 4.50 \pm 0.55 ^a | 2.50 \pm 0.55 ^a |
| | II | 6.50 \pm 0.55 ^a | 8.50 \pm 0.55 ^a | 6.50 \pm 1.64 ^a | 3.50 \pm 0.55 ^b |
| | III | 8.50 \pm 0.55 ^b | 8.50 \pm 1.64 ^a | 11.50 \pm 2.74 ^b | 7.00 \pm 1.1 ^c |
| Copepodites | I | 7.50 \pm 0.55 ^a | 22.50 \pm 4.93 ^a | 14.50 \pm 2.74 ^a | 10.00 \pm 1.1 ^a |
| | II | 10.00 \pm 1.1 ^b | 26.50 \pm 1.64 ^b | 13.50 \pm 4.93 ^a | 11.00 \pm 1.1 ^a |
| | III | 15.00 \pm 1.1 ^c | 47.00 \pm 0 ^c | 15.50 \pm 4.93 ^a | 14.00 \pm 1.1 ^b |
| Nauplii | I | 14.00 \pm 2.19 ^b | 14.00 \pm 3.29 ^a | 5.50 \pm 0.55 ^a | 6.50 \pm 1.64 ^b |
| | II | 7.50 \pm 3.83 ^a | 18.00 \pm 4.38 ^a | 7.50 \pm 1.64 ^a | 4.50 \pm 0.55 ^a |
| | III | 13.00 \pm 2.19 ^b | 26.00 \pm 2.19 ^b | 11.50 \pm 2.74 ^b | 9.00 \pm 2.19 ^c |

Treatment I - Cowdung; Treatment II - GOC; Treatment III - Rice bran. Values bearing same superscripts within each stage do not differ significantly ($p < 0.01$)

had a high DHA:EPA ratio and this may be the reason for the use of *I. galbana* for sustained viable egg production through multiple generations of copepod species. After *I. galbana*, *C. calcitrans* was found to be the next beneficial diet for culture of *E. acutifrons* in the present study. Copepods fed on *I. galbana* tended to have shorter time for maturity than those fed on *T. gracilis*. The present study is in conformity with the studies made by Payne and Ripplingale (2000) wherein they found *I. galbana* as the second best diet for *G. imparipes*. *C. marina*, resulted in poor hatching and survival to maturity. However, a mixture of the best two diets *I. galbana* and *C. calcitrans* showed good hatching, survival and egg production within 7 days. Miralto *et al.* (2005) studied the effect of food type (two diatoms and two dinoflagellates) on the egg production and hatching success of *Centropages typicus*. Reproductive responses were strongly affected by food type and fecundity was twice high with larger cells and hatching success was 2-3 times higher with dinoflagellate diets.

In the mass culture experiments with algal diets, adult production and survival were higher with a mixed diet of 75% *I. galbana* and 25% *C. calcitrans*, in both set of experiments with and without replacement of culture medium and algae. This is in agreement with earlier referred works of Shrivastava *et al.* (1999) and Nassogne (1970). Robin *et al.* (2005) reported that *Chaetoceros-Isochrysis* combination enabled favourable production of *Paracalanus* sp. Schipp *et al.* (1999) similarly reported benefit of using mixed algal diet for mass production of *Acartia* sp. Corkett and Urry (1968) reported that *I. galbana* and the diatom *Lauderia borealis* were the most satisfactory food organisms for *Paracalanus elongates*. The second best feed which produced more adults was *T. gracilis* in place of *I. galbana*. The survival rate was in agreement with the initial glass vial experiments. Similar observation was reported by Huei-Meei Su *et al.* (2005) in the production of *Apocyclops royi*. When fed with *Tetraselmis* sp., *A. royi* matured earlier and produced more offsprings, than copepods fed with other 10 diets. Copepods fed on *Tetraselmis* sp., which contains low amount of eicosapentaenoic acid (EPA) and no docosahexaenoic acid (DHA) grew faster than *I. galbana* which is rich in DHA indicating that the highly unsaturated fatty acid content may not be critical in the growth of *A. royi*. Same may be true in the case of *E. acutifrons* also. The growth pattern may change in mass culture depending on the population size, concentration and size of feed. The positive effect of *I. galbana* was also observed in several studies worldwide. Similarly, in copepodite and nauplii production also, *I. galbana* was observed to be the best feed than the mixed diet and *T. gracilis*. The findings illustrate a stage specific

microalgal requirement of the copepod, with smaller developmental stages of *E. acutifrons* benefiting from *I. galbana* diet and adults favouring *T. gracilis* diet. It is in line with the findings of David *et al.* (2005) where *Chaetoceros* proved to be an inferior diet to *Isochrysis* for nauplii. Several studies have reported high survival rates and fecundity among adult copepods offered with large celled microalgae (Murray and Marcus, 2002; McKinnon *et al.*, 2003). Adults seemed to prefer comparatively large celled *T. gracilis* to *I. galbana* and mixed diet. Microalgal size is an important factor affecting feeding efficiency of different copepod developmental stages (Berggreen *et al.*, 1988). In the present study, it was found that the size of food organisms also influenced the growth and biomass of *E. acutifrons*. Similar results were earlier recorded by Nassogne (1970) in *E. acutifrons* and Corkett (1970) in *Acartia tonsa*.

With respect to ovigerous females, comparatively good production was noticed with *C. calcitrans* and mixed algal diet. In all other diet treatments, almost same rate of production was noticed. Razouls *et al.* (1986) reported that cell size is an important factor affecting production of *Temora stylifera*. In all the treatment groups, production of all the three stages *viz.*, adults, ovigerous females and copepodites-nauplii was found to be much higher with replacement of culture medium. Comparatively low production in the second set without replacement of culture medium, could be attributed to the inadequate feed in the medium. Food is an important limiting factor to support the growth and density of copepod in laboratory culture. The abundance of nauplii, copepodites and adults of *E. acutifrons* increased in proportion to the increased food concentration throughout the culture. Guisande *et al.* (1996) observed that if the food concentration in the food column diminished, mean number of eggs also decreased, resulting in lower production of next generation.

In the mass culture trials using organic feed medium, almost similar pattern was observed in the production of adults, ovigerous females, copepodites and nauplii. Rice bran proved to be the best for copepod production, followed by groundnut oil cake with cow dung showing the lowest production. Turk *et al.* (1982) described rice bran culture as a simple, inexpensive and successful method for production of *A. tonsa* and successfully cultured the population for four months in seawater with rice bran. He opined that it provides all the nutrients for the copepods. Vilela (1992) reported that the caloric content and nutritional quality was better in rice bran-*Platymonas* mix for culture of *Tigriopus brevicornis*. Barber and Barber (1974) established that rice bran contained many important amino acids and vitamins which are of high nutritive value. Joebert *et al.* (2005)

reported that the decreasing trend in copepod production in the second week onwards could be due to predation or cannibalism. In the present study, 14 days of culture was found to be optimal under laboratory conditions. The candidate species *E. acutifrons* had a generation time of 10-14 days, which depend on the temperature and availability of food. The maximum production of nauplii was attained on 14th day of culture and the production decreased from 15th day onwards during most of the experimental trials.

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