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Study on the Growth Performance and Survival of Hatchery Produced *Macrobrachium rosenbergii* (De Man) Larvae in Natural Brackishwater of Different Grades

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Abstract: A study was undertaken to evaluate the possibility of utilizing natural brackishwater (from Shibs River) in the hatching operation and larvae culture of *Macrobrachium rosenbergii*. Three treatments each with three replications were tested. Natural brackish water diluting by rainwater (T₁), natural brackishwater diluting by under ground tap water (T₂) and natural brackish water with salinity range from 12 to 20 ppt. (T₃) were considered. In case of dilution (in T₁ and T₂) salinity concentration range was maintain from 4-18 ppt as per requirement of the experiment and stage of brood and larvae. Significant higher survival (23.38%) for T₁ was observed followed by T₃ with 13.63% and T₂ with 4.38% up to post larvae (PL) stage. Significant ($p < 0.05$) difference in survival rate of larvae between T₁ and T₂ and T₂ and T₃ was observed. Study reveals that natural brackishwater diluting by rain water can be effectively used in the hatching operation and larvae culture of the freshwater giant prawn, *M. rosenbergii*.

Key words: *M. rosenbergii* larvae, metamorphosis, growth performance, survival rate and natural water

Introduction

Macrobrachium rosenbergii is a freshwater prawn and in Bangladesh most commonly known as Golda chingri found in fresh water rivers with tidal influence. They are also found in ponds, canals, irrigation ditches having connection with tidal water. It is indigenous to the south-east Asia including Bangladesh. But in our country the availability of this species in nature is decreasing day by day demanding its culture practice. In recent time, the heavy demand of this prawn in both for national and international markets due to its good nutritional value as well as deliciousness, expand the culture system of the animal. To date, in our country monoculture of this species are being practiced at Khulna, Bagerhat, Piruzpur, Gopalgong, Madaripur and Narail districts covering a total area of about 9000 hector. Polyculture of prawn with carp in ponds, rice-cum-prawn culture is also being practices in our country (DOF, 1994, 1996). Moreover there is enormous potentiality of mixed culture of freshwater prawn *M. rosenbergii* with brackishwater shrimp *P. monodon* in coastal region of the country particularly in Cox's Bazar and greater Khulna districts (FRI, 1994). But main obstacle in flourishing culture practice of *M. rosenbergii* is the scarcity of seeds from nature as well as from hatcheries due to various difficulties. One of the most practical difficulties of providing prawn seed from natural source is that, it is almost impossible to identify prawn larvae from other freshwater prawn species which prohibits the monoculture system of *M. rosenbergii* directly. Though in the recent year a considerable number of backyard prawn hatcheries started to produce prawn larvae in the country, but depending upon the location of the hatchery, source of saline water/brine and water management in hatchery operation, the production rate of prawn post larvae for most of this hatcheries are found below the standard rate (not more than 20%) which is again another important drawback for prawn culture development. So effort should be directed towards artificial sources with proper water management. Nowadays, giant fresh water prawn's larvae are reared with the conventional techniques where artificial brackishwater of about 12 ppt. Salinity is prepared from seawater. The utilization of natural brackishwater in the hatching and larvae culture of giant prawn has so far received very little attention from aquaculturists. In present study the attempt was taken to utilize the natural brackishwater for hatching and larval rearing of giant prawn.

Materials and Methods

The study was conducted from mid April to late September 1998 at the experimental hatchery unit of Brackishwater Station, Bangladesh Fisheries Research Institute, Paikgacha, Khulna. For dilution of brackishwater into different required concentrations, natural brackishwater from the nearby Shibs River was used. For

removing dirt, suspended particle and microbial cells, collected natural water was at first filtered by primary filtration unit (filter formed by gravel) and stored into under ground water tank from where it was lifted up to overhead tank and passed through sand filtration unit and ultra violet (UV) unit and stored into ground tank then again it was lifted up to overhead tank and finally stored in cistern. Prior to use for the hatching activity and larvae culture, 60% chlorinated bleaching powder at the rate of 12 ppm was added to this water for further purification and kept for 2 days for settling followed by treatment with sodium thiosulphate at the rate of 10 ppt. and allowed another 2 days for settle which was followed by agitation using air blower for 1 days. After agitation it was kept 1 days for settle down all waste material. Then supernatant was used for the experimental work.

All necessary equipment's i.e. siphon pipe, bucket, bowl, net pieces etc were washed twice daily prior to use for siphoning and water exchanging by chlorinated water (60% chlorinated bleaching powder) at the rate of 12 ppm. Larvae produced were siphoned off and the number of larvae was estimated by the Aliquot method prior to stock in larval rearing tanks.

Three different water type (considered as treatments) each with three replications were used. The treatments were as follows: T₁ for natural brackishwater mixed with rainwater; T₂ for natural brackishwater mixed with underground tap water and T₃ for natural brackishwater only.

The study was divided into three batches and the experimental treatments were set in randomized block design. In each batch 3 berried females were used as mother prawn. Three rectangular fiberglass tanks each of 152 L holding capacity were filled with the rearing medium three days prior to stocking.

Hatchlings were reared with varying densities as 44.70-66.82 larvae/L and fed with *Artemia* nauplii. *Artemia* nauplii were supplied after 48 hours of hatch at a rate of 5 naupli/ml of water and *Artemia* nauplii as feed were used up to PL stage. During rearing of larvae in the tanks aeration was provided at a rate of 10 L/min with 30 minutes break after continuous one-hour operation. To keep the tanks bottom clear, water exchange was done every morning and evening of the day at the rate of 90 and 50% respectively by siphoning. The rearing process continued till the attainment of about 95% of the stock as post larvae (PL) (Table 1).

Water quality parameters of each rearing medium such as temperature, dissolved oxygen, pH and ammonia were recorded twice daily. The total hardness of each rearing medium was also determined. The salinity of the rearing medium was checked once a week with a hand refractometer.

Table 1: Replication wise dates of hatching, stocking dates and density of hatchlings

Replication	Date		No. of larvae/tank	No. of larval/L
	Hatching	Stocking		
T ₁ R ₁	April 15	April 17	6711	44.70
T ₁ R ₂	April 18	April 20	6923	46.20
T ₁ R ₃	April 23	April 25	6812	45.40
T ₂ R ₁	June 5	June 7	8913	59.40
T ₂ R ₂	June 9	June 11	8125	54.20
T ₂ R ₃	June 13	June 15	9045	60.30
T ₃ R ₁	August 2	August 4	10,023	66.82
T ₃ R ₂	August 6	August 8	8837	58.90
T ₃ R ₃	August 9	August 11	9759	65.00

Table 2: Treatment wise average survival rate of *M. rosenbergii* up to the attainment of post larval stage

Treatment	Survivability (%)			Treatment mean (%)
	R ₁	R ₂	R ₃	
T ₁	24.03	22.25	23.87	23.38(±0.985)
T ₂	3.68	5.07	4.37	4.38 (±0.695)
T ₃	14.03	14.47	12.34	13.63 (±1.125)

Table 3: Duration of metamorphosis of *M. rosenbergii* hatchlings during this larval rearing under different treatments.

Treatment	Metamorphosis (days)			Mean of metamorphosis (days)		Average (days)
	R ₁	R ₂	R ₃	First	Last	
T ₁	24-45	29-47	31-42	28	44.66	36.33
T ₂	31-47	29-46	33-48	31	47.00	39.00
T ₃	34-47	30-47	31-46	31.66	46.66	39.17

Results and Discussion

Significant higher survival (23.38%) of the post larvae was observed in T₁ followed by T₃ (13.63%) and T₂ (4.38%) with significant variation among the treatment average ($p < 0.05$). The findings clearly represent the impact of water quality on larval survivability. Because, the maximum survivability in T₁ might due to the better quality of rain water added during dilution than the under ground water, that contains higher amount of iron as well as hardness of water is maximum, those directly influenced larval metabolism and development. Due to variation in water quality and due to presence of some undesirable compound, Ong (1975) and Aniello and Singh (1980) made similar observation on survivability. Different in composition of the rearing media may consider as possible cause of difference in mortality rate among the treatments (Table 2).

The other cause of low survival on identified were cannibalism among the members, improper acclimatization of the larvae before stocking in the rearing tank, high iron content of underground water and adhesion of the larvae along the inside wall of rearing tank due to jumping. Larval growth period to first day of metamorphosis was also found variable (Table 3). Data reveals that T₁ had the earliest mean at day 28, followed by T₂ and T₃ at 31th and 31.66th respectively. The table also shows that T₁ had the shortest mean larval cycle of 44.66 days, followed by T₃ (46.66 days) and T₂ (47 days).

The variation in length of larval cycle might due to the impact of temperature of culture environment that has direct effect on metabolism and growth performance of the animal. The course of study as per water parameter record, the T₁ part was performed in a warmer environment than T₂ and T₃. During the operation of T₂ and T₃, due to occasional rain during the period, temperature went down and retards animal's metabolism and growth. Similar observation on larval growth due to variation in temperature was reported by Gibson (1975), Suharto *et al.* (1980) and New and Singholka (1982). Authors also reported that temperature lower than optimum (i.e. below 24-26°C) required more larval rearing time.

In Bangladesh, generally proto type recirculatory *Macrobrachium* sp. hatchery with biofiltration unit does not acceded survivability 25% up to its post larval development and time length required for attaining post larvae 25-30 days (DOF, 1996) which is in both the cases more or less similar with the findings of T₁ where clear rain water was used for alteration of the natural brackishwater up to the required level. So the findings of the experiment indicates that natural brackishwater diluted with rainwater can be effectively used for larvae culture of giant freshwater prawn.

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