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ABSTRACT

Role of eye-stalk of *Macrobrachium gangeticum* Bate, 1868 in its reproductive behaviour has been examined by conducting deletion and addition experiments. Eye-stalk ablation induced gonadal maturity in both sexes, leading to change in colour and size of ovaries and increase in GSI and oocyte diameter in females and increased length of testes and diameters of seminiferous tubules in males. Injection of eye-stalk extracts tended to at least partly restrict the effects in both sexes. The experiments thus suggested that the eye-stalk of *M.gangeticum* released some gonad inhibiting factors.

Key words : Macrobrachium gangeticum, Eye-stalk, Gonads.

INTRODUCTION

The eye-stalk of crustanceans possesses X-organ-sinus gland complex which is an important neuroendocrine organ of crustaceans, (Cooke, 1982; Bliss, 1971 and Passano, 1951.) The histoarchitecture and seasonal variations of the X-organ sinus gland complex of *Macrobrachium gangeticum* Bate, 1868, have earlier been reported, (Gupta, 2007 a & b) and the present communication deals with the role of this organ in regulation of the reproductive behaviour of the species.

Twenty immature specimen each of male and female *M. gangeticum* (ranging in size 13-15 cm) were collected from Ganga river near Buxar and Ballia in the month of March (2004-2005) and brought to the laboratory. Morphological characteristics of gonads were noted and they were then fixed

in Bouin's solution and sections (5-6 μ m thick) were stained with eosin/hematoxylin.

The animals were grouped under 8 groups (I-VIII) with five prawns in each group (groups I-IV include females and V-VIII include males)

Groups I & V - Control (female and male)

Groups II & VI - Eye-stalk ablated (female and male)

Groups III & VII - Normal females and males injected with eye-stalk extract.

Groups IV & VIII - Eye-stalk ablated females and males injected with the eye-stalk extract.

Eye stalk ablation was done in animals kept under ice cold water. The homogenate of the eye-stalk was prepared according to the methods described by Nagabhushanam and Diwan, (1974), Twenty microlitres of the eye-stalk extract, prepared by homogenizing 2 eye-stalks in 20 μ l distilled water, (equivalent to two eye-stalk/prawn) were injected on the 3rd, 6th and 9th days in the abdominal musculature of prawns of groups III, IV, VII and VIII with the help of hypodermic microsyringe. Prawns were sacrificed on day 15 and examined.

OBSERVATIONS

Effects of eye-stalk on ovarian maturation:

The eye-stalk ablated females showed a noticeable change in colour of their ovaries-from pale greenish to green, a significant increase in mean GSI and also in oocyte diameter at the termination of the experiment as compared to the corresponding values of controls. The females of group III showed the values of mean GSI and oocyte diameter to be somewhat lower than those of group II (eye-stalk ablated) counterparts. Similarly, when the eye-stalk ablated animals were injected with eye-stalk extract (Group IV), the above values were reduced as compared to those of group II females and were nearly touching the corresponding values of control animals (Table-1).

Effects of eye-stalk on testicular maturation:

The testes of eye-stalk ablated animals showed somewhat advanced stage of maturity both in physical appearance as well as in histology at the termination of the experiment. The colour was white with pink spots, the average length measured 3.21 + 0.59 cm. against the control value of 2.41 ± 0.54 cm and the average seminiferous tubule diameter was 44.31 \pm 4.55 μ m against the control value of $30.54 \pm 3.52 \mu m$. While the seminiferous tubules of control animals had spermatocytes and few spermatogonia as their dominant cell types, those of eye-stalk ablated animals had spermatids and spermatozoa. The animals of group VII, exhibited opposite results. Length of testes as well as tubule diameter showed much reduced values, nearly approaching the control values. Normal/Eye-stalk ablated animals, when supplemented with eye-stalk injections (Group VII & VIII), exhibited reduced testicular maturity. Length of testes and tubule diameter were only a little more than the corresponding values of control animals. Spermatocytes were dominant in their seminiferous tubules (Table-2).

Table-1

Animal groups (size range 13-15 cm)	GSI mean \pm SD	Oocyte diameter ±SD(μm)	Colour of the ovary at the end of experiment
Group-l	2.87±0.24	24.91±2.41	Pale Greenish
Group-II	3.32±0.31	35.95±0.23	Green
Group-III	2.95±0.25	28.51±3.54	Pale Greenish
Group-IV	2.90±0.29	26.33±2.51	Pale Greenish

(All the data represent values at the termination of experiment)

Table-2

Effects of eye-stalk on testicular maturation of *M.gangeticum*

Animal groups (size range 13-15 cm)	Length of testes (cm)	Diameter of seminiferous tubules (µm)	Dominant cell types
Group-V	2.41±0.54	30.54±3.52	spermatocytes, few spermatogonia
Group-VI	3.21±0.59	44.31±4.55	spermatids and spermatozoa
Group-VII	2.53±0.41	33.45±3.45	spermatocytes and few spermatogonia
Group-VIII	2.82±0.34	35.24±4.21	spermatocytes

(All the data represent values at the termination of experiment)

DISCUSSION

Deletion and addition experiments have revealed the physiological role of the eye-stalk of *M. gangeticum*. Eye-stalk ablation of females led to change in the colour of ovaries from light green to dark green and increase in GSI and oocyte diameter. All these changes indicated accelaration of the ovarian maturity. However, when normal or eye-stalk ablated animals were injected with eye-stalk extracts, the above changes were, more or less, restricted.

Almost similar effects were produced in males post eye-stalk ablation. Length of testes as well as diameters of seminiferous tubules got increased. Presence of spermatids with few spermatozoa in the testes of eyestalk ablated animals indicated precocious maturity. As in case of females, the effects of eye-stalk ablation in males too were more or less restricted following injection of eye-stalk extract. Thus, in case of both males and females, eye-stalk ablation leads to advancement of gonadal maturity. However, specimen of groups III and VII, which include normal females and males injected with eyestalk extracts, show values somewhat higher than that of groups I and V (controls). However, the differences are not very significant. But the values of groups II and VI (eye-stalk ablated females and males) are significantly higher than those of the other groups of males and females respectively.

These observations, clearly indicate that the x-organ sinus gland complex located inside the eye-stalk exerts some gonad inhibiting effects. This conclusion is in agreement with the findings of Rangnekar and Deshmukh, 1968. John and Sivadas, 1979 noted early maturity of ovary in eye-stalk ablated estuarine crab, *Scylla serrata*. Similar reports were made by Panouse, 1943 in *Palaemon serratus*. Mirajkar *et al.*, 1985 reported high ovary inhibiting activity of the

eye-stalk in spawned prawn Macrobrachium kistensis. While Adiyodi and Adiyodi, (1974) were of the opinion that the ablation effect on reproduction of crustaceans depends on the maturation stage of the ovary, Bomirski and Klek, (1974) suggested that the ovary inhibiting hormone of eye-stalk is responsible for the resting phase of the ovary and controls the role of vitellogenesis. Han and Kim, (1993) also found the gonad inhibiting effect of xorgan of Macrobrachium nipponense to be stronger in resting period. Other studies showing the gonad inhibiting effects of eyestalk include those by Mirajkar, (1980) on Macrobrachium kistensis, Kumar and Pandian on Macrobrachium nobilii, Rajender Rao, (1987) on *Macrobrachium lammarrei*, (1987) Biswas, (1991) on Macrobrachium dayanum, Wilder et al., (1994), Perez-Cruz et al., (1995) and Kumar and Pandey, (2003) on Macrobrachium rosenbergii and (Upadhyaya, 2000) on Macrobrachium malcomsonii.

In the present study, the precocious ovarian maturity following eye-stalk ablation did not lead to oviposition. This is in accordance with the finding of Sarojini *et al.*, (1982 & 1983), Sambhasiva Rao (1986), Biswas, (1991) *etc*. Eye-stalk is also not reported to affect the process of hatching, (Saigusa, 2000).

Several studies also confirm the present observation that the factors released from eye-stalk bring about precocious maturity of testes as well. Mirajkar, (1980), Sarojini *et al.*, (1982), Rajender Rao, (1989), Biswas, (1991) and Upadhyaya (2000) have reported early maturity of testes of various *Macrobrachium* species after eye-stalk ablation. However, the findings of Carlisle, (1954) and Aoto and Nishidha, (1956) do not correspond to the present view. Eye-stalk ablation is also reported to increase the intermoult period as well as the growth rate.

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