

## BIOLOGY OF LADYBIRD BEETLE *Micraspis discolors* (Fab.) (COCCINELLIDAE: COLEOPTERA)

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### ABSTRACT

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The experiment was conducted at the Department of Entomology, Hajee Mohammad Danesh Science and Technology University, Dinajpur, during January to April 2007 at 21.02°C ± 4.5 °C (Room temperature) and 66.05 ± 0.95 % RH to observe the biology of ladybird beetle. The mean pre-oviposition period of the ladybird beetle *Micraspis discolor* was 5.1±0.48 days and the oviposition period was 40.10 ±0.46 days. The female beetles laid on an average 235.50 ±7.96 eggs and the hatching % was 83.93 ±2.11. The average incubation period was 3.1± 0.22 days and the duration of total the larval period from egg to adult was 10.76 ±1.35 days. The pre-pupal and pupal periods were 1.50 ±0.20 and 2.60± 0.21days, respectively. The mean longevity of the male and the female beetles were 40.20 ±1.0 and 47.50 ±0.82 days, respectively.

**Keywords:** Ladybird beetle, incubation, larval stage

### INTRODUCTION

The ladybird beetles have been associated with good fortune in many myths and legends. They have been honoured through the centuries as they vernacular indicate that the term 'Lady' is in reference to biblical Mother Mary (Roache, 1960). The ladybird beetles have been known worldwide as a predator of a number of insects. They are distributed in many countries of Asia, including Bangladesh (Islam and Nasiruddin, 1978). This beetles, often called ladybug or coccinellids. They are the most commonly known of all beneficial insects. In Europe these beetles are called ladybirds (William, 2002). They are of great economic important as predaceous both in their larval and adult stages on various important crop pests such as aphids, coccids and other soft bodied insects including aphids (Hippa *et al*, 1978; Kring *et al*, 1985), while the species *M. discolor* feed on many insect pests such as aphids, brown plant hopper, corn borer, Lepidopteron insects, mealybug, white flies (Rao *et al*, 1989; Mani, 1995). This predaceous coccinellids is also found in association with those insects infesting bean, wheat, chilli, sorghum, tobacco, cotton, maize, potato, lathyrus, soyabean, sweet potato, lentil, mustard, brinjal, groundnut, sunflower and cabbage. (Gautam *et al*, 1995; Duffied, 1995).

Country bean (*Lablab purpureus* Linn.) is a common and popular vegetable of Bangladesh. Locally it is well known as "Sheem" and cultivated widely in winter season in our country. It contains 4.5% protein in pod, 25% protein in dry seed (Rashid, 1976). It also content with appreciable amount of vitamin, phosphate, calcium and sodium (Gopalan *et al*, 1982). Generally it is also known as income generating crop in our country. This crop is also important for its atmospheric nitrogen fixation (Kalra, 1979).

The aphid is one of the most destructive pests and its distribution is world wide. Bean aphid, *Aphis craccivora* Koch. attack the bean plant and other leguminous crops attacked by the nymphs and the adults of aphid cause damage by sucking the sap from the flowers, buds, pods, tender shoots and reduce the market value (Srivastava and Singh, 1986). At the time of infestation plants fail to give flowering and pods setting resulting in 20-40% yield loss (Islam, 2007).

In Bangladesh the bean growers use various insecticides to control bean aphid. Insecticidal control is not only expensive but also its residues left over the sprayed surface of the crops or in the soil and have become a matter of concern of environmental pollution. The indiscriminate use of pesticides causes phytotoxicity and destruction of beneficial organisms such as predators, parasitoids microorganisms and pollinators (Luckman and Metcalf, 1978). Global warning has cautioned us and the adverse consequences of insecticide use are always alarming and also inducing pest out break because of pest resistance. These entomological backlashes have compelled the scientists to be concerned with entomologically compatible pest management programs (Hodek, 1970).

Now a day, Integrated Pest Management (IPM) is well known to entomologists, where all suitable pest control techniques are being used to find ecologically sound and environmentally safe ways of pest control. Biological control should be regarded as the backbone of any IPM program and about 90% of all potential pests are already under biological control (Debach and Rosen, 1991). The biological control is one of the most effective means of achieving insect control (Pedigo, 2004). The vegetable growers use less insecticide or avoid insecticides in

developed countries. In recent years, pest control particularly for aphids has been revolutionized by the application of predators and parasitoids in those countries (Bari and Sardar, 1998). The coccinellid beetles are considered to be a great economic importance in agro-ecosystem through their successful employed in the biological control of many injurious insect (Agarwala *et al*, 1988).

In our country most of the farmers are not well acquainted with grubs and pupae of the lady birds. They are accustomed to use insecticide indiscriminately without monitoring the pest population above the Economic Threshold Level (ETL) as well as consulting experts of this line. Its adverse side effects have already caused threat to a great extent to human being, beneficial predators. Our poor farmers are becoming poorer and crops vulnerable to attack and disease. In this aspect exploration of predators like *M. discolor* in Bangladesh may play a vital role as part of IPM Programme.

The study of the biology of *M. discolor* would help to use this insect of proper biological control. But reviews of biology of *M. discolor* is limited and environmental conditions of Dinajpur are different from other areas of the country. So, the present study was undertaken to observe the biology of *M. discolor*

## MATERIALS AND METHODS

The experiment was conducted at the laboratory of the Department of Entomology, Hajee Mohammad Danesh Science and Technology University (HSTU), Dinajpur, during the period of January to April 2007. All the experiment were completed at  $21.02^{\circ}\text{C} \pm 4.5^{\circ}\text{C}$  (Room temperature) and  $66.05 \pm 0.95$  % RH.

### *Collection and mass culture*

A culture of large number of larvae and adult predator of *Micrapis discolor* was established in the laboratory in order to supply necessary insects for the experiment. For this reason, some males and females of the *Micrapis discolor* were collected by sweep net from the unsprayed horticulture field and were confined in petridishes. Bean aphids were also collected daily with infested bean leaves, stems, twigs and inflorescences from the same unsprayed field and supplied as food. These beetles were sexed and paired in petridishes (6.0 x1.0 cm). The bottom of the petridishes was covered with blotting paper (Whitman filter paper no 1). After hatching of eggs, the grubs were transferred to several medium sized petridishes (11cm diameter) and reared on bean aphids. The emerged adults were sexed and confined in pairs in the petridishes and fed on bean aphids. Egg masses were collected and reared as above and continued for several times for obtaining large number of larvae and adult predators.

### *Study of Biology*

Egg laid by each female during 24 hours were counted and kept in separate petridish to determine the total number of eggs laid per female and also hatching period including hatching percentage (%) of eggs were recorded. After hatching of eggs young larvae were then transferred individually in petridishes (6.0 x 1.0 cm), containing blotting paper at the bottom. Fresh bean shoots infested with aphids were placed in each petridish to provide food for the larvae every morning. The larvae were observed twice daily at 12 hrs interval until pupation. The number of instars and period of each instar were recorded. The pupae were kept undisturbed in the respective petridishes until the emergence of adult. At the same time the pupal period was recorded. Ten replications were used in this case.

### *Statistical analysis*

The Data were analyzed by Analysis of Variance (ANOVA) and the mean values were separated by Duncan's Multiple Range Test (DMRT). All analyses were performed using SPSS, version 12.01 (SPSS, Chicago, IL, USA).

## RESULTS AND DISCUSSIONS

### *Pre-oviposition*

The time between the date of adult emergence and the first egg deposition was considered as pre-oviposition period. The pre-oviposition period of *M. discolor* was 3 to 7 days with an average of  $5.10 \pm 0.48$  days (Table 1)). Agarwala *et al* (1988) observed that the pre-oviposition period was 6 to 10.33 days on *A. craccivora* at  $16-26^{\circ}\text{C}$ . Prodhan *et al* (1995) studied that the pre-oviposition period of *M. discolor* was 3 to 7 days, which is similar to the present findings.

### *Oviposition*

Oviposition period was the duration between first and last egg laying. The oviposition period in the present study was 38 to 43 days. (Table 1) and the mean oviposition period was  $40.10 \pm 0.46$  days. Prodhan *et al* (1995)

reported that the period of *M. discolor* lasted from 35 to 42 days using *A. craccivora* as food, which is similar to the present observation.

**Incubation**

Incubation period was the duration between the dates of egg laying and hatching. The incubation period was 2 to 4 days with an average of  $3.10 \pm 0.22$  days (Table 1). Islam and Nasiruddin (1978) observed that the incubation period of *M. discolor* was 2 days on cotton aphid. However, Ngammuang (1987) and Prodhan *et al* (1995) reported that it was 2 and 3 days respectively using bean aphids as host. These results seem to be close with the present observation.

Table 1. Duration of different stages of *M. discolor* as noted in the study

Different stages	Duration of days		Mean $\pm$ SE
	Minimum	Maximum	
Pre-oviposition period	3	7	$5.10 \pm 0.48$
Oviposition Period	38	43	$40.10 \pm 0.46$
Incubation period	2	4	$3.10 \pm 0.22$
Total larval period	8.8	14	$10.76 \pm 0.47$
Pre-pupal period	1	2	$1.50 \pm 0.20$
Pupal period	2	4	$2.60 \pm 0.21$
Total developmental period from egg to adult	12.5	25	$17.96 \pm 1.35$
<b>Longevity of adult</b>			
Male	35	45	$40.20 \pm 1.00$
Female	45	53	$47.50 \pm 0.82$
Average longevity of the beetle			$43.85 \pm 0.91$

**Duration of larval stages**

The total larval period (1<sup>st</sup> instar to 4<sup>th</sup> instar) was  $10.76 \pm 0.47$  days (Table 1). Nasiruddin and Islam (1979) observed that the total period of *M. discolor* was 11.8 to 12.5 days, which is higher than the present findings. Prodhan *et al* (1995) observed that the total larval period of *M. discolor* varied from 7 to 9 days on bean aphid. This result was lower than the present study. However, Sakurai *et al* (1991) reported that the quality of food and environmental factors like temperature, humidity also play an important role on different aspects of the biology of coccinellid beetles. So, this variation may be due to the quality of food and environmental factors like temperature and humidity.

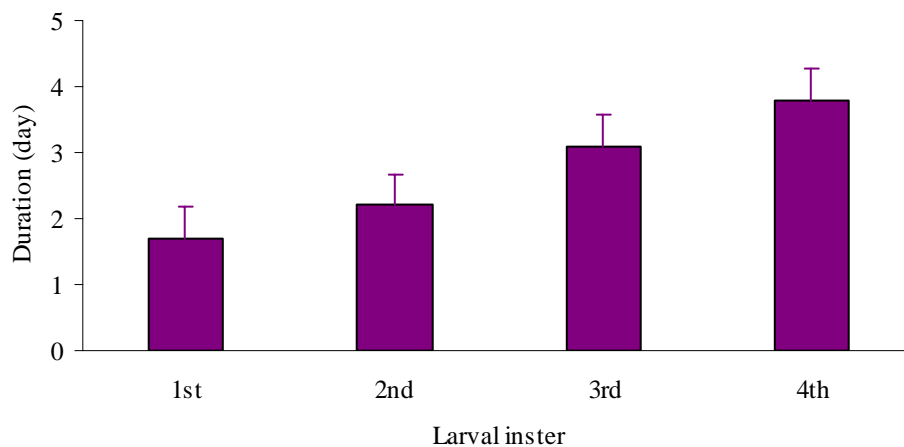


Figure 1. Duration (mean  $\pm$  SE) of different larval instars of *M. discolor* reared on bean aphid.

**First instar**

The newly hatched larval period was 1 to 3 days and on an average of  $1.71 \pm 0.20$  days (Figure 1). Prodhan *et al* (1995) reported that this period of *M. discolor* was 2 to 3 days using bean aphid as a host. This result supported to the present observation.

**Second instar**

The results of the duration of the 2<sup>nd</sup> instar varied from 1.50 to 3 days and the mean duration was  $2.20 \pm 0.16$  days (Figure 1). Nasiruddin and Islam (1979) found that the duration of the 2<sup>nd</sup> instar larvae of *M. discolor* was 2.4 to 3.1 days on different aphid, which is comparatively similar to the results of the present findings. Prodhan *et al* (1995) found that the duration of 2<sup>nd</sup> instar of *M. discolor* varied from 1 to 2 days using bean aphid using cabbage aphid as a host.

**Third instar**

The result indicated that the duration of the 3<sup>rd</sup> instar larvae lasted from 2 to 4 days. The mean duration of 3<sup>rd</sup> instar larvae was  $3.10 \pm 0.17$  (Fig. 1). Nasiruddin and Islam (1979) found that the duration of the 3<sup>rd</sup> instar larvae of *M. discolor* varied from 3.1 to 3.8 days on maize, bean and chilli aphids as host. This result supported to the present observation.

**Fourth instar**

Observation made on the larval duration of the 4<sup>th</sup> instar larvae on an average  $3.75 \pm 0.19$  days with minimum of 3 days and maximum of 5 days (Figure 1). Prodhan *et al* (1995) reported that the duration of final instar larvae of *M. discolor* was 3 days. Nasiruddin and Islam (1979) found that the duration of the 4<sup>th</sup> instar larvae of *M. discolor* varied from 3.8 to 4.2 days on maize, bean, and chilli aphids. This duration was within the range of present findings.

**Pupal period**

The pupal period was 2 to 4 days with an average  $2.60 \pm 0.21$  days (Table 1). Nagammuang (1987) recorded that the mean pupal duration of *M. discolor* was  $3.43 \pm 0.57$  days when larvae reared on *A. craccivora*. Different findings revealed that the pupal period of coccinellid beetles varied with the differences of food and it was correlated with the temperature (Sakurai *et al* 1991).

**Facundity and hatching rate**

The number of eggs laid per female were 190 to 270 with an average  $235.50 \pm 7.96$ . The mean hatching percentage were  $83.93 \pm 2.11$  (Table 2). Ngammuang (1987) reported that the number of eggs deposited per female of *M. discolor* was  $181.07 \pm 66.37$  on *A. craccivora*, and 70.15% eggs were hatched. Prodhan *et al* (1995) observed that the fecundity of female varied from 200-300 eggs with mean of 270.5 and with average 70.15% eggs were hatched. These results seem to be close with the present findings.

Table 2: Fecundity and hatching rate of *M. discolor* as observed in the study

No of observation	No. of egg laid	No. of egg hatched	% of egg hatching
1	228	205	71.18
2	196	181	92.35
3	242	210	86.78
4	250	208	83.20
5	242	220	90.91
6	222	193	86.94
7	255	211	82.75
8	260	191	73.46
9	190	170	89.47
10	270	222	82.22
Mean $\pm$ SE	$235.50 \pm 7.96$	$201.10 \pm 5.08$	$83.93 \pm 2.11$

**Adult Longevity**

The longevity of adult beetles was counted from the emergence of the adult to its death. The longevity of the male beetles varied from 35 to 45 days with an average of  $40.20 \pm 1.00$  days. Whereas, the longevity of the female beetles varied from 45 to 53 days with an average of  $47.50 \pm 0.82$  days. The average longevity of the beetle (male and female) were  $43.85 \pm 0.91$  days (Table 1). It showed that the longevity of the male beetle was shorter than the female. But there were no significant differences. Samal and Misra (1985) reported that the adult of *M. discolor* lived for 24 to 40 days in September- November. This result was near similar to the present study. Ngammuang (1987) found that the longevity of male and female were  $37.8 \pm 15.24$  and  $59.53 \pm 23.53$

days when fed on *A. craccivora*, in the laboratory at temperature of  $28 \pm 2^{\circ}\text{C}$  with 74% RH. These results are also close to the present observation.

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