Effect of Prey Density on Biology and Foraging Potential of Mallada boninensis (Okamato) (Neuroptera: Chrysopidae)

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ABSTRACT

The present study aimed to evaluate the effect of prey density on the biology and functional response of green lacewing, Mallada boninensis (Okamato) (Neuroptera: Chrysopidae). Newly emerged larvae of M. boninensis were fed 20, 30, 40, 50, 60, 70, 80, 90 and 100 fresh eggs of Corcyra cephalonica (Lepidoptera: Gelechiidae) in plastic vials. It was observed that the prey density had a significant effect on the positive consumption rate, development, and fecundity of M. boninensis. In general, maximum consumption with shortest developmental time, maximum fecundity, and most extended adult longevity was observed as prey density increased. The predatory potential was high when the prey density was raised. The daily predation rate of M. boninensis increased slowly during the first two instars and reached its peak in the third larval instar. The results indicated that M. boninensis feeding potential and developmental period might vary from 6.00 to 11.33 days based on food density and having a difference in per day consumption. The 100 Corcyra eggs/day treatment had the highest egg intake of 87.88 eggs per day, followed by 90 eggs/day (79.33 eggs) and 80 egg day−1 (69.75 eggs).

Keywords: Corcyra cephalonica; Chrysopidae; Prey density; Lacewing; Mallada boninensis

INTRODUCTION

There are several natural enemies of insect pests which co-exist with them in the different ecosystems. Amongst a very complex network of bio agents, the Chrysopid is known to be the most effective predator. Chrysopids are commonly termed green lacewings. Chrysopids have a tremendous potential to consume pests of crops. Because of their large geographical distribution, broad habitats with a high relative frequency of occurrence, strong seeking ability, and ease of laboratory rearing, C. zastrowi sillemi (Esben-Peterson) and Mallada boninensis (Okamoto) (Neuroptera: Chrysopidae) are the most thoroughly researched species of Chrysopids. Green lacewings are one of the most efficient generalist predators employed in biological control (McEwen et al., 2001). Adults are generally not predatory and feed on nectar, pollen or honeydew while a few of them are predatory (Coppel and Mertins, 1977). Amongst the Mallada spp., M. boninensis, M. basalis, M. aster, and M. desjardinsi are essential as these are found to be potential predators of aphids, leaf miners, psylla, blackfly, and whitefly (Syed et al., 2008; Riddick, 2009). In recent years use of green lacewing species has been recommended for the IPM programme (Nehare et al., 2004). They can be successfully reared on eggs of Corcyra cephalonica Stainton in the laboratory.

The predation phenomena of the M. boninensis, are sometimes not as simple as mentioned above but change with varying prey densities. It has been observed with many insects and small animal predators that when prey population increased, prey consumption also increased and consumption rate is the function of food density (Elango and Sridharan, 2018). Such a changing behavior ultimately affects the predator’s release pattern in a bio-control program and needs to be studied for better understanding under different ecosystems. Therefore, a study was designed to evaluate the predation rate of laboratory-reared M. boninensis, on Corcyra eggs with the following objectives: (i) to determine the predation rate of all larval instars under nine prey density levels and (ii) to study the effect of prey density on the biology of M. boninensis.

MATERIAL AND METHODS

Culture of Rice moth, Corcyra cephalonica (Stainton)

Rice moth, C. cephalonica has been widely used as an efficient alternative host for the mass rearing of
many biocontrol agents. Bajra (Pennisetum glaucum L.) grains were coarsely milled and broken into 2-3 pieces in a milling machine. The broken grains are heat sterilized at 100°C for 1 hour to eliminate the residual population of stored product insects viz., Rhizopertha dominica, Sitotroga cerealella, Tribolium castaneum and fungal contaminants. Upon sterilization, the grains were cooled under the fan in a clean area. The grains were then transferred to plastic basins @ 2.5 kg/basin. Groundnut kernel in required quantity was broken using a pounding machine, a mechanical blender, or a machine (domestic mixer). After that, 100 g of the broken kernel was placed in each basin and well mixed by hand. The materials were well combined after adding dry yeast (Bakers) and wettable sulphur at a rate of 5g/basin. This was followed by a spray of 10 mL of 0.01-0.05 percent streptomycin sulphate and mixing the contents. Corcyra larvae are reared in the bajra medium.

The eggs used to start a Corcyra colony must be devoid of pollutants such as moth scales and broken limbs, and they must not be exposed to UV radiation. The number of trays that can be infected with eggs is estimated by measuring the volume of overnight deposited eggs collected. A cc of eggs is estimated to contain between 16000 and 18000 eggs. Cumbu medium with optimal amounts of Corcyra eggs Corcyra eggs infest 0.5 cc of water each basin. After that, the basins were covered with a clean khada fabric and secured with rubber fasteners. After the eggs have been infested for 28-30 days, the adults emerge. On the inside of the khada fabric, the growups may be seen. They are either aspirated or collected using a mechanical moth catcher with specimen tubes. The entire procedure takes place under a mosquito net tent.

Along with the eggs, scales and shattered limbs were discovered in higher proportions. The eggs were carefully rolled on filter paper to another container to clean the collecting. The eggs were quantified in measuring cylinders and used to build up the stocks and natural enemy production.

**Mass culturing of Mallada boninensis**

Grubs were reared in G.I. round basins (28 cm dia) @250 larvae/basin covered with khada cloth. The eggs of Corcyra cephalonica were given as feeding material for the larvae in the laboratory. In ten days, the M. boninensis larvae puate into round white-colored silken cocoons. The cocoons were collected with fine brush and transferred into 1 litre plastic container with a wire mesh window for emergence of adults. The adults were collected daily and transferred to pneumatic glass troughs or G.I. round troughs (30 cm × 12 cm). Before allowing the adults, the rearing troughs were wrapped inside with a brown sheet that acted as egg receiving card. About 250 adults (60 % females) were allowed into each trough and covered with white nylon or georgette cloth secured by a rubber band. Three bits of foam sponge (2 sq.in) dripped in water were kept above the nylon cloth cover. Besides, an artificial protein rich diet (yeast, fructose, honey, Proteinex® and water in the ratio 1:1:1:1) was provided in semisolid paste form in three spots on the cloth outside. The adults were collected daily and allowed into fresh rearing troughs with fresh food. From the old troughs, the brown paper sheets and M. boninensis eggs were removed and used to maintain a culture of green lacewing.

**Prey density on biology of M. boninensis**

The experiment was designed in a completely randomized (CRD) design. There were nine (food densities) treatments, viz., 20, 30, 40, 50, 60, 70, 80, 90, 100 Corcyra eggs/grub/day. Each treatment was replicated thrice. Each replication has three samples (Mallada grubs). Freshly harvested M. boninensis eggs were placed in 9 cm Petri dish sealed with parafilm to avoid desiccation and were observed daily. Newly emerged larvae of M. boninensis were transferred into plastic vials with counted number of fresh Corcyra eggs for each food density/ replication. The larvae of M. boninensis were transferred carefully to the capsules with the help of fine camel hair brush. Vials contents were observed under stereo microscope every day to determine the number of unconsumed eggs and any change in larval biology. The numbers of unconsumed eggs were subtracted from the total number of offered eggs (prey density) and data were recorded on a daily basis. Biological parameters like the duration of development of each larval instar, pupation, adult emergence, and mortality occurring in each treatment was recorded daily in all prey densities. After pupation, each pupa was observed for adult emergence and recorded. The emerging adults were transferred to G.I. round troughs (30 cm × 12 cm). The adults were fed daily on a thick viscous solution of water + honey + yeast. The adult’s basins were observed every 24 h for egg-laying
Statistical analysis

The data collected under laboratory experiments in completely randomized design were analyzed using analysis of variance (ANOVA) using AGRES 3.01 and AGDATA software. Data in the form of numbers were transformed to square root values and those in numbers were transformed to and analyzed. The mean values of the treatments were compared using DMRT at 5 per cent level of significance.

RESULTS AND DISCUSSION

Effect of food density on Mallada boninensis consumption

The larvae of *M. boninensis* responded to increasing prey densities with increasing food consumption and older larval stages displayed a higher rate of predation than younger ones (Fig 1). The consumption rate increased progressively during each day. Maximum the *M. boninensis* fed 703.00 ±0.18 *Corcyra* eggs within minimum developmental period of 6.00±0.01 days in the 100 eggs/day treatment. When the intake rate was reduced (173.67±0.07 *Corcyra* eggs) in the low number of eggs per treatment (20 eggs/day), the developmental time was increased by 11.33±0.02 days. The results indicated that *M. boninensis* feeding potential and developmental period may vary (6.00±0.01 to 11.33±0.02 days) based on food density and the difference in per day consumption. Maximum consumption 87.88 ±0.01 eggs/ day were observed in 100 *Corcyra* eggs/day treatment followed by 90 eggs/day (79.33±0.05 eggs) and 80 egg day⁻¹ (69.75 ±0.03 eggs).

Table 1. Effect of food density on duration of grub, pupal and adult periods of *Mallada boninensis*

| Treatments (Eggs) | First instar (Days) | Second instar (Days) | Third instar (Days) | Pupal period (Days) | Developmental period (Days) | adult longevity (Days) |
|------------------|---------------------|----------------------|---------------------|---------------------|----------------------------|------------------------|
| 20               | 3.67                | 3.00                 | 4.67                | 9.33                | 20.67                      | 2.33                   |
|                  | (1.92)              | (1.73)               | (2.16)              | (3.05)              | (4.55)                     | (1.53)                 |
| 30               | 3.67                | 3.00                 | 4.33                | 8.33                | 19.33                      | 4.67                   |
|                  | (1.92)              | (1.73)               | (2.08)              | (2.89)              | (4.40)                     | (2.16)                 |
| 40               | 3.33                | 2.33                 | 4.00                | 8.67                | 18.33                      | 5.67                   |
|                  | (1.82)              | (1.53)               | (2.00)              | (2.94)              | (4.28)                     | (2.38)                 |
| 50               | 3.33                | 2.33                 | 3.67                | 8.00                | 17.33                      | 7.67                   |
|                  | (1.82)              | (1.53)               | (1.92)              | (2.83)              | (4.16)                     | (2.77)                 |
| 60               | 3.00                | 2.33                 | 3.33                | 7.67                | 16.33                      | 8.00                   |
|                  | (1.73)              | (1.53)               | (1.82)              | (2.77)              | (4.04)                     | (2.83)                 |
| 70               | 3.00                | 2.33                 | 3.33                | 6.33                | 15.00                      | 8.33                   |
|                  | (1.73)              | (1.53)               | (1.82)              | (2.52)              | (3.87)                     | (2.89)                 |
| 80               | 3.00                | 2.33                 | 2.67                | 6.00                | 14.00                      | 9.00                   |
|                  | (1.73)              | (1.53)               | (1.63)              | (2.45)              | (3.74)                     | (3.00)                 |
| 90               | 2.67                | 2.00                 | 2.33                | 5.00                | 12.00                      | 9.67                   |
|                  | (1.63)              | (1.41)               | (1.53)              | (2.24)              | (3.46)                     | (3.11)                 |
| 100              | 2.33                | 1.67                 | 2.00                | 4.67                | 10.67                      | 10.67                  |
|                  | (1.53)              | (1.29)               | (1.41)              | (2.16)              | (3.27)                     | (3.27)                 |

*Mean of three replications. Values in the parentheses are square root transformed values. In a column, means followed by the common letter(s) are not significant in DMRT @ 5% level of significance.*
Food density had pronounced effects on the biological parameters of *M. boninensis*. The *M. boninensis* have three instar in grub period and developmental periods also change based on the food density. The first instar grub has duration from 2.33 days (100 eggs/day) to 3.67 days (20 eggs/day). The first instar duration was the same in the treatment 60, 70 and 80 eggs/day with 3.00 days. Similar results were also found in the duration of first instar grub was maximum on *A. gossypii* with 3.54 days and minimum on *C. cephalonica* with 2.48 days (Guntupalli and Kalyanasundram, 2016). The duration wise there was not much wide variance in high and low density prey in the first instar. The second instar *M. boninensis* having duration from 1.67 days to 3.00 days. Averagely in all treatments other than 90 and 100 eggs/day, the duration of the second instar was 2.33 days. The third instar duration range from 2.00 days (100 eggs/day) to 4.67 days (20 eggs/day)(Table.1) In the third instar *M. boninensis* was voraciously feeding in *Corcyra* eggs and there may be variance in the third instar duration of *M. boninensis*. Earlier research indicated the larval duration of a related species *Mallada astur* (Banks) as 11.6 days (Venkatesan et al., 2002) and *Mallada basalis*, 11.8 days (Chang, 2000) which are close to our observations. In the present trials, it was noticed that although *M. boninensis* larvae completed development in each of the nine prey densities, an increase in prey density reduced development time and mortality rate.

**Duration of developmental stages**

Food density directly affected the 3rd instar larval duration, pupal period, and adult longevity of *M. boninensis*. Increased prey densities reduced the developmental time and mortality rate of *M. boninensis*. (Fig. 2). Maximum duration 20.67 days of grub period was recorded in 20 eggs/day treatment whereas; minimum duration 10.67 days was recorded in 100 eggs/day treatment. In the case of pupal period, it was shortest in 70 and 80 eggs/day treatment (6.63 and 6.00 days) whereas, the longest pupation period (9.33 days) was recorded in 20 eggs/day treatment. The longest adult longevity was observed in 80, 90 and 100 eggs/day treatments, while shortest adult longevity was noticed in 20, 30 and 40 eggs/day treatments. The results were on par with the results of Nagamallikadevi et al. (2013).

**CONCLUSION**

It was noticed from the data that younger *M. boninensis* grub consumed less food in all prey densities than older grub, probably due to smaller in size larvae having less mobility and prey handling efficiency than a larger sized grub or older grub.

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**Ethics statement**

No specific permits were required for the described field studies because no human or animal subjects were involved in this research.

**Consent for publication**

All the authors agreed to publish the content.

**Competing interests**

There was no conflict of interest in the publication of this content.

**Data availability**

All the data of this manuscript are included in the M.S. No separate external data source is required. If anything is required from the M.S., certainly, this will be extended by communicating with the corresponding author through corresponding official mail: elaento@gmail.com

**AUTHOR CONTRIBUTIONS**

Research grant - NA; Idea conceptualization - SJN & SS; Experiments - KE; Guidance - SJN; Writing- original draft - K.E.; Writing- reviewing & editing - KE&SJN

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