EFFECT OF DIETS AND TEMPERATURE DEGREES ON THE BIOLOGICAL ASPECTS OF THE LAELAPID MITE, OLOLAELAPS USSURIENSIS BREGETOVA & KOROLEVA, 1964

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ABSTRACT: The biological aspects of the laelapid mite, Ololaelaps ussuriensis (Laelapidae: Mesostigmata) was investigated at laboratory conditions under different diets and two temperature degrees. The obtained results showed that the diets as acarid mite, Tyrophagus putrescentiae (Schrank) and free living nematodes, Rhabditella muscicola (Andrassy) and temperature degrees (15 and 25 °C) were significantly influenced the periods of incubation period, longevity and life span of the predatory mite, O. ussuriensis, but there was no significant differences in the case of combination or interaction between these factors .Data , also, indicated that the adult female of O. ussuriensis laid more eggs (41.8 eggs) when fed on the free living nematode at 25 °C compared with the acarid mite diet . The lowest number of laid eggs (33 eggs) was observed when the females fed on T. putrescentiae at 15 °C.

Key words: Ololaelaps ussuriensis, diets, free live nematodes, biological aspects.

INTRODUCTION

Mites represent the majority of the soil and organic manure arthropods fauna, and playing different important roles in nature which vary from harmful and beneficial. Soil mites play an important role in increasing soil fertility by its effect on organic matter decomposition. Owing to their numerical importance, the soil mites have received more attention than other soil Acari. Actinedida and Gamasida represent the major groups of soil mites which found in many soil habitats (Usher, 1971; Wallwork, 1976; Mostafa, 1980 and Convey et al., 2000). Although the Acarina communities of Egyptian soil have not been widely studied, some information is available on the prostigmatid and mesostigmatid mites' fauna. The majority of these species appear to predators associated with small and immature stages of insects, mites, and nematodes in the soil surface (Karg 1961; Sardar and Murphy 1987; Bilgrami (1994). Some of them feed on fungi and helps in control soil born fungi diseases (Ragusa and Zedan 1985 and Ahmed 1998). Smaller

deep-litter and soil forms are predominantly nematophagous and are the most important predators of nematodes in many habitats. Interaction within a community and therefore predator mites have the ability to keep free living nematodes and mite pests under the threshold level. In Egypt, Ibrahim 1982, Metwally et al., 1983, Ahmed 1998, Kaid1998 and Ezz El-Dein 2003 investigated the behaviour of the different ground mites. The mite family Laelapidae (Mesostigmata) is ecologically diverse, including obligate and facultative parasites of vertebrates, insect paraphages, and free-living predators that inhabit soil litter habitats and the nests of vertebrates and arthropods (Evans and Till 1966; Strong and Halliday 1994; Lindquist et al., 2009).

The present work aims to study the effect of different diets and temperatures on the biological aspects of the laelapid mite, *O. ussuriensis* at laboratory conditions.

MATERIALS AND METHODS

Collection and preparation of the commonest predacous soil mite:-Individuals

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belonging to the laelapid mite, Ololaelaps bregetova (Laelapidae: Mesostigmata) Shereef and Soliman used in biological studies were collected from normal untreated soil of maize, wheat and soybean at Qaha region, Qaluobia Governorate. For individual rearing, newly deposited eggs were transferred individually to a new rearing plastic cell. Each newly hatched larva was supplied with prey. Devoured prey was replaced daily until maturity. Emerged females were allowed to mate with males and monitored for oviposition. All biological aspects were recorded twice daily during the mite development. Other necessary data dealing with the predator's biology, fecundity and other biological aspects were recorded.

Rearing procedures:

To rear the predacious mite, О. bregetova, there are two types of cages were used. The first for culturing mites (large glass Petri dishes filled up to 0.5 cm with a mixture of plaster of Paris and Charcoal (9:1) and the second for individual rearing (plastic ring 2.5 cm in diameter and 1.5 cm in depth) filled with 0.7 cm with a mixture of plaster of Paris and Charcoal (7: 3) according to Metwally et al., (1983). For culturing of mites, several adult females and males of O. bregetova were placed in Petridishes. The bottom was kept moist, thus the relative humidity was suitable by adding one or two drops of water every two days. A camel hair brush was used to transfer newly deposited eggs to the plastic rings. After hatching, each larva was supplied with free living nematodes and acarid mite as prevs. Observations were made twice daily using stereomicroscope to determine different biological aspects. Copulated female were kept for determining pre-oviposition, oviposition and postoviposition periods. Observation was taken at 15 and 25+2 °C. and relative humidity 75+5 %.

Source of food:

a. Free-living nematode:

Soybean and wheat soil samples were

put in Baermann funnel for 24 hours for extracting free living nematodes *R. muscicola*, Abou-EI-Sood (1992). The extraction of free living nematode was conducted in Petri-dishes contains slides of Potatoes. Petri-dishes were kept at 25 °C. Camel hair brushes were used to add drops of food in rearing cells of the predatory mites as the main source of food.

b. Acarid mite:

The culture of acarid mite, *T. putrescentiae* was maintained and already obtained form Acarology Dep., Plant Prot. Res. Inst.,. The collected individuals of both predators and prey species were transferred to a covered glass Petri-dishes which covered with a layer of Plaster of Paris and charcoal mixture. Moisture is kept by adding drops of water every two days.

Statistical analysis:

Suitable statistical analysis was used to clearify the results of the work. All biological studies data were subjected to one way analysis of variance (ANOVA) and means were separated by Duncan's multiple range tests, Duncan (1955).

RESULTS AND DISCUSSION Incubation period:

It was clearly obvious from Tables (1&2) that there were highly significant differences between the incubation period which give rise to_males of *O. bregetova* when fed on different diets. This period was 6.5, 5.5 & 3.1 and 2.4 days at 15 and 25 °C, when feeding of females and males, respectively on acarid mites. However, these periods changed to 7.9, 7.3 days at 15 °C and 2.85 and 2.3 days at 25 °C for females and males, respectively when fed on nematode.

However, statistical analysis of obtained data indicated that there were highly significant differences between the individuals fed on different diets at different temperatures. Generally, the effect of interaction between the temperature and food type on the incubation period of *O*. *bregetova* male individuals was observed not affected, Table (4).

Larval stage:

The obtained results in Tables (1&2) indicates that the highest recorded larval stage of *O. bregetova* was noticed when females fed on the acarid mite, *T. putrescentiae* at 15 $^{\circ}$ C (6.0 days) and the

lowest larval stage was noticed when the male individuals reared on the free nematodes at 25 °C (1. 3 days). On the other hand, the quiescent larval stage of *O. bregetova* durated the highest period 1.6 days when the female fed on the acarid mite at 15 °C, decreased to recorded 0.33 days on the free nematodes at 25 °C

 Table (1): Biological aspects of the laelapid mite, Ololaelaps ussuriensis when fed on acarid mite, Tyrophagus putrescentiae at different temperatures

Biological aspect		15 °C		25 ℃	
		Ŷ	8	Q +	8
Incubation period		6.5 <u>+</u> 0.53 (6-7)	5.5 <u>+</u> 0.53 (5-6)	3.1 <u>+</u> 0.74 (2-4)	2.4 <u>+</u> 0.52 (2-3)
Larva	а	6.0 <u>+</u> 0.67 (5-7)	5.0 <u>+</u> 0.82 (4-6)	2.43 <u>+</u> 0.5 (2-3)	1.73 <u>+</u> 0.52 (1.25-2.25)
	q	1.6 <u>+</u> 0.52 (1-2)	0.93 <u>+</u> 0.37 (0.25-1.25)	1.25 <u>+</u> 0.4 (1-2)	0.7 <u>+</u> 0.39 (0.25-1)
Protonymph	а	6.2 <u>+</u> 0.79 (5-7)	5.2 <u>+</u> 0.79 (4-6)	2.7 <u>+</u> 0.52 (2-3.25)	2.28 <u>+</u> 0.4 (2-3)
	q	1.95 <u>+</u> 0.4 (1-2.25)	1.34 <u>+</u> 0.54 (0.25-2)	0.98 <u>+</u> 0.23 (0.25-1.25)	0.8 <u>+</u> 0.56 (0.25-2)
Deutonymph	а	7.68 <u>+</u> 0.43 (7-8)	6.88 <u>+</u> 0.47 (6-7.25)	3.58 <u>+</u> 0.51 (3-4.25)	2.83 <u>+</u> 0.41 (2.25-3.25)
	q	1.63 <u>+</u> 0.39 1-2.25)	1.23 <u>+</u> 0.32 (1-2)	1.0 <u>+</u> 0.29 (0.25-1.25)	0.63 <u>+</u> 0.39 (0.25-1)
Immature stages		25.06 <u>+</u> 1.25 (24.5-25.6)	20.58 <u>+</u> 1.27 (20-21.1)	11.94 <u>+</u> 0.68 (11.3-12.6)	8.97 <u>+</u> 0.59 (8.6-9.3)
Life cycle		31.55 <u>+</u> 1.39 (29.25-33.25)	25.78 <u>+</u> 1.78 (23.5-28.5)	14.93 <u>+</u> 1.18 (13.5-16.75)	11.25 <u>+</u> 1.23 (9-12.5)
Longevity		18.5 <u>+</u> 0.85 (17-20)	16.83 <u>+</u> 1.04 (15-18)	12.5 <u>+</u> 1.33 (11-15)	10.93 <u>+</u> 1.0 (10-13)
Life span		50.25 <u>+</u> 1.24 (48.25-51.5)	42.58 <u>+</u> 1.74 (39.5-44.5)	(27.43 <u>+</u> 1.92 (24.5-30.75)	22.18 <u>+</u> 1.19 (20.75-23.75)

<u>+</u> Standard deviation a = active q = quiescent

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Biological aspect		15 °C		25 °C	
		Ŷ	0,	9	6
Incubation period		7.9 <u>+</u> 0.88 (7-9)	7.3 <u>+</u> 0.67 (6-8)	2.85 <u>+</u> 0.61 (2-4)	2.3 <u>+</u> 0.56 (1.25-3)
Larva -	а	7.3 <u>+</u> 0.82 (6-8)	6.4 <u>+</u> 0.7 (6-8)	1.53 <u>+</u> 0.56 (1-2.25)	1.3 <u>+</u> 0.39 (1-2)
	q	1.5 <u>+</u> 0.53 (1-2)	0.75 <u>+</u> 0.44 (0.25-1.25)	0.7 <u>+</u> 0.39 (0.25-1)	0.33 <u>+</u> 0.24 (0.25-1)
Protonymph	а	6.9 <u>+</u> 0.74 (6-8)	5.8 <u>+</u> 0.63 (5-7)	2.25 <u>+</u> 0.42 (2-3)	1.42 <u>+</u> 0.41 (1-2)
	q	1.35 <u>+</u> 0.61 (0.25-2)	0.75 <u>+</u> 0.44 (0.25-1.25)	0.7 <u>+</u> 0.39 (0.25-1)	0.48 <u>+</u> 0.36 (0.25-1)
Deutonymph -	а	7.5 <u>+</u> 0.71 (7-9)	6.6 <u>+</u> 0.7 (6-8)	3.33 <u>+</u> 0.47 (2-3)	1.78 <u>+</u> 0.46 (1.25-2.25)
	q	1.08 <u>+</u> 0.59 (0.25-2)	1.23 <u>+</u> 0.13 (1-1.25)	0.7 <u>+</u> 0.39 (0.25-1)	0.48 <u>+</u> 0.3 (0.25-1)
Immature stages		26.63 <u>+</u> 2.25 (26.0-17.3)	21.53 <u>+</u> 1.08 (21.1-22.2)	9.21 <u>+</u> 1.0 (9-10.3)	5.79 <u>+</u> 0.45 (5-6.4)
Life cycle		33.53 <u>+</u> 1.67 (31-36)	28.73 <u>+</u> 2.09 (24.5-31.25)	11.03 <u>+</u> 1.85 (7.75-14.5)	8.08 <u>+</u> 1.54 (5.5-11.5)
Longevity		14.8 <u>+</u> 1.69 (13-19)	13.93 <u>+</u> 1.65 (12-18)	9.2 <u>+</u> 13.2 (7-11)	8.93 <u>+</u> 1.63 (7-12)
Life span		48.38 <u>+</u> 1.51 (45.5-50.5)	42.56 <u>+</u> 1.40 (40.5-45.25)	20.23 <u>+</u> 2.59 (17.5-24.5)	17.0 <u>+</u> 2.36 (14.5-20.75)

Table (2): Biological aspects of	the laelapid mite,	, Ololaelaps ussuriensis	s when fed on
free living nematodes	, Rhabditella musci	cola at different tempera	atures

+ Standard deviation a = active q = quiescent

Protonymphal stage:

Results in Tables (1&2) indicated that the protonymphal active stage of the predatory mite, *O. bregetova* took the longest period 6.2 days when fed as female individuals on *T. putrescentiae* at 15 °C, and took the shortest period 1.42 days when male fed on the free nematodes at 25 °C. On the other hand, the quiescent protonymphs had the same trend of active stage but the periods differed, 1.95 days for females and 0.48 days for males.

Deutonymphal stage: As shown in Tables (1&2) the deutonymphal stage of *O*.

bregetova was high when the active mite females and males fed on *T. putrescentiae* 7.68 and 6.88 days at 15 °C, respectively, but the quiescent period of male when fed on free nematode was the lowest period 0.48 days at 25 °C.

Immature stages:

It is clear from the current study that the immature stages of the predatory mite *O. bregetova* took 25.06, 20.58; 11.94 and 8.97 days when the individuals fed on the acarid mite, *T. putrescentiae* at 15 and 25 °C for females and males, respectively, Tables (1&2). However, this period averaged 26.63,

21.53; 9.21 and 5.79 days when the same individuals fed on the free living nematodes, respectively.

Life cycle:

Concerning the life cycle, Tables (1, 2 &4), statistical analysis using L.S. D. at 0.05 pointed out that the life cycle of *O. bregetova* was highly significantly differed at different temperatures for (different sexes) males and females. The obtained results indicated that the longest duration period of the life cycle was recorded when the females fed at 15 °C (31.55 days) on the acarid mite, while the shortest period was recorded when the male individuals fed on the free living nematode at 25 °C (8.08 days).

Longevity:

Statistical analysis of the obtained results in Tables (1, 2 and 4) revealed the occurrence of significances differences of longevity period of O. bregetova when fed on the free living nematodes and the acarid mite. Longevity took 18.5, 16.83 & 12.5 and 10.93 days at 15 °C and 25 °C for females and males, respectively when the mites fed on the acarid mite, T. putrescentiae as a food. However, these periods changed to recorded 14.8, 13.93 & 9.2 and 8.93 days when the mites fed on the nematode at the conditions the same of experiment. Generally, the effect of temperature and food on the longevity of O. bregetova individuals was observed highly significant, Table (4).

Pre-oviposition, oviposition and post-oviposition periods:

The respective durations of pre-oviposition, oviposition and post-oviposition of 0 bregetova on T.putrescentiae and free nematodes are summarized in Table (3). The obtained data showed that there were obvious differences between the O. bregetova female individuals fed on both the acarid mite and the nematode at different temperature in case of pre-oviposition period. This period

lasted 4.5 and 2.4 days when the predatory mite fed on the acarid mite at 15 and 25 °C, respectively. On the other hand, this period took 3.25 and 1.7 days when the mite fed on nematode. On the other hand, the longest oviposition period of *O. bregetova* was noticed when the female fed on the acarid mite at 15 °C (8.9 days), but the shortest period was noticed when the mites fed on nematodes at 25 °C, (5.0 days). However, the longest postoviposition period was 5.1 days when the mites fed on the acarid mite at 15 °C, while the shortest period took 2.5 days at 25 °C on the nematode, Table (3).

Fecundity:

The eggs of *O. bregetova* females laid individually and the number of deposited eggs are represented in Table (3). The study shows the influence of different temperatures and hosts on the fecundity of the predatory adult females of *O. bregetova*. Data clearly indicated that the feeding at 25 °C on nematode gave the highest level of egg (41.8 eggs) while the lowest number was 33.0 eggs, when the mites fed at 15 °C on the acarid mite.

Life span:

Accordingly, the life span of *O. bregetova* differed at different temperatures, Tables (1&2). However, this period lasted the highest level (50.25 days) when the females fed on the acarid mite at 15 °C, changed to the lowest level when the males reared on nematodes at 25 °C (17.0 days).

The statistical analysis of data, Table (4) showed that the sex (male and female), diets (acarid mites and nematode) and temperature (15 and 25 °C) had affected on the incubation period, longevity and life span of *O.bregetova* with highly significant effect, but there was no any significant difference in the case of combination or interaction between these factors collectively on these periods. On the other hand, there was no any obviously effect for interaction of these factors on the mite life cycle.

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Biological aspect	Diet	15 ºC	25 ⁰C
Preoviposition period	Acarid mite	4.5 <u>+</u> 0.53 (4-5)	2.4 <u>+</u> 0.46 (2-3)
	Nematode	3.25 <u>+</u> 0.72 (2.5-5)	1.7 <u>+</u> 0.67 (1-3)
Oviposition period	Acarid mite	8.9 <u>+</u> 0.88 (8-10)	7.5 <u>+</u> 0.97 (6-9)
	Nematode	7.1 <u>+</u> 0.99 (6-9)	5.0 <u>+</u> 0.82 (4-6)
Postoviposition period	Acarid mite	5.1 <u>+</u> 0.88 (4-6)	2.6 <u>+</u> 0.7 (2-4)
	Nematode	4.45 <u>+</u> 0.50 (4-5)	2.5 <u>+</u> 0.53 (2-3)
Fecundity	Acarid mite	33.0 <u>+</u> 1.83 (30.36	39.6 <u>+</u> 1.43 (37-42)
	Nematode	38.8 <u>+</u> 1.23 (37-41)	41.8 <u>+</u> 1.87 (40-5)

Table (3): Longevity (in days) and fecundity (No. of eggs) of the laelapid mite adult female, Ololaelaps bregetova fed on different diets at different temperatures

+ Standard deviation

Table (4): Effect of different diets and temperatures on the biological aspects of the laelapid mite, *Ololaelaps ussuriensis*

Biological aspect	Source	F.	Ρ.	L.S.D. at 0.05 level
Incubation period	Sex	24.74	0.000***	
	Diet	24.74	0.000***	0.2856
	Temp.	834.22	0.000***	
	Int. sex x diet x temp.	0.1903	0.6639 ns	
	Sex	141.18	0.000***	
Life evelo	Diet	2.206	0.1418 ns	0.7214
Life cycle	Temp.	2634.55	0.000***	
	Int. sex x diet x temp	0.0298	0.6847 ns	
	Sex	13.30	0.000***	0.6012
Longevity	Diet	97.307	0.000***	
	Temp.	347.871	0.000***	
	Int. sex x diet x temp	0.1717	0.6798 ns	
Life span	Sex	182.77	0.000***	
	Diet	76.74	0.000***	0.8064
	Temp.	3595.65	0.000***	
	Int. sex x diet x temp	0.0021	0.9632 ns	

*** highly significant ns = non-significant

Imbriani and Mankau (1983) observed voracious feeding by Lasioseius sculpatus on Aphelenchus avenae and Cephalobous sp. increased population of mite resulted in a significant decline of A. avenae. The most definite association between mites and nematodes came from the work of Rodriguez et al., (1972) who cultured Macrocheles muscaedomesticae on Rhabditis sp., and found it to prefer house fly eggs over nematodes. Its prot-and deutero-nymphs under same conditions, however, preferred nematodes.

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تاثير الغذاء ودرجات الحرارة على المظاهر البيولوجية للاكاروس الليلابيدى Ololaelaps ussuriensis عادل محمود مصطفى^(۱) ، هناء ابراهيم محمود^(۲) ، عصام محمد عبدالسلام ياسين^(۱) ، **حسنية عبد الفتاح عفيفى علوان^(۱) ، عابدين محمود خليل^(۱)** ^(۱) معهد بحوث وقاية النباتات- مركز البحوث الزراعية - الدقى - جيزة - مصر ^(۲) قسم علم الحيوان - كلية العلوم (بنات)- جامعة الازهر - مدينة نصر – القاهرة - مصر.

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