

Research Paper

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Biological responses of *Tetranychus urticae* (Acari: Tetranychidae) to sub-lethal concentrations of the entomopathogenic fungus *Beauveria bassiana*

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Abstract

Tetranychus urticae (Acari: Tetranychidae) is one of the most important pests of agricultural crops with worldwide distribution causing considerable damage to different products. Application of chemical acaricides is one of the most important strategies used for the control of this pest. Entomopathogenic fungi, however, have been proposed as alternative control agents. In this study, sub-lethal effects ($LC_{10} = 6.76 \times 10^2$, $LC_{20} = 8.74 \times 10^3$, and $LC_{30} = 55.38 \times 10^3$ conidia ml^{-1}) of *Beauveria bassiana* strain TV on the life table parameters of *T. urticae* were evaluated under laboratory conditions. The results demonstrated that by increasing the concentration, a significant decline was observed in adult longevity of both male and female individuals. Total fecundity of *T. urticae* was calculated as 45.16, 36.28, 23.98, and 18.21 eggs in control, LC_{10} , LC_{20} , and LC_{30} treatments, respectively. Sub-lethal concentrations drastically affected the population parameters of this mite pest. The intrinsic rate of increase (r) ranged from 0.1983 to 0.1688 day^{-1} for the mites treated with distilled water and LC_{20} treatments, respectively. The net reproductive rate (R_0) was affected by the sub-lethal concentrations (lower value at LC_{30} concentration: 11.19 offspring/individual). Considering the detrimental effects of *B. bassiana* on some biological parameters of *T. urticae*, it can be concluded that this product can be used to develop targeted interventions aimed at integrated pest management of this pest.

Introduction

The two-spotted spider mite, *Tetranychus urticae* Koch (Acari: Tetranychidae), is one of the most destructive mite pests with worldwide distribution (Migeon and Dorkeld, 2006–2017) on different crops such as maize, soybean, cotton, bean, eggplant, tomato, cucumber, potato, strawberry, currants (Sedaratian *et al.*, 2009, 2011; van Leeuwen *et al.*, 2010; Khanamani *et al.*, 2013; Maleknia *et al.*, 2016; Mollaloo *et al.*, 2017; Azadi-Qoort *et al.*, 2019). Direct feeding of *T. urticae* causes a loss of leaf chlorophyll content reducing the net photosynthetic rate, resulting yield losses and finally decline and death of the host plants (Tomczyk and Kropczynska, 1985; Campbell *et al.*, 1990; Park and Lee, 2002; Meck *et al.*, 2012, 2013; Abou El-Ela, 2014).

Applications of synthetic acaricides are noted as the most important strategy for combating the population of this pest (Chen *et al.*, 2019). Unfortunately, high reliance on these chemicals leads to some undesirable consequences such as pest resurgence, development of resistance, environmental pollutions, and negative effects on non-target organisms especially natural enemies (Croft, 1990; Shi *et al.*, 2005; Maniania *et al.*, 2008).

Biocontrol agents are generally discussed as safe alternatives for the management of pest mites (Wekesa *et al.*, 2015), as they are compared to selective chemical pesticides and can be the main component of integrated mite management programs (Ullah and Lim, 2017). Among different biocontrol agents, entomopathogenic fungi (EPF) have been considered as an inseparable part of integrated strategies for suppressing different mites and insect species (Wekesa *et al.*, 2006). *Beauveria bassiana* (Bals.) Vuill. play a critical role in management programs of numerous pest species (Irigaray *et al.*, 2003; Wekesa *et al.*, 2006; Maniania *et al.*, 2008; Seiedy *et al.*, 2010). Indeed, a possible effect on different mite species has been documented by several researchers (Maniania *et al.*, 2008; Geroh *et al.*, 2015). The usage of EPF in biological control is increasing mostly because of greater environmental awareness, food safety concerns, and the failure of conventional chemicals due to an increasing number of insecticide-resistant species (Shahid *et al.*, 2012; Yuan *et al.*, 2018). Keeping these advantages in view, several limitations (I: need specific environmental conditions; II: short shelf life; III:

Table 1. Toxicity of *Beauveria bassiana* strain TV on female individuals of *Tetranychus urticae*

No.*	df	Sub-lethal concentrations (conidia ml ⁻¹)*				Slope ± SE	χ ²	P
		LC ₁₀ (confidence limits)	LC ₂₀ (confidence limits)	LC ₃₀ (confidence limits)	LC ₅₀ (confidence limits)			
640	5	6.76 × 10 ² (2.29 × 10 ² –16.14 × 10 ²)	8.74 × 10 ³ (4.01 × 10 ³ –16.88 × 10 ³)	55.38 × 10 ³ (29.84 × 10 ³ –97.11 × 10 ³)	11.70 × 10 ⁵ (6.71 × 10 ⁵ –21.44 × 10 ⁵)	2.69 ± 0.30	3.90	0.007

often slow acting and require high application rate (Maina *et al.*, 2018)) could drastically affect the biological performance of these natural enemies, and among them, reception of sub-lethal concentrations has considerable potential (He *et al.*, 2013; Song *et al.*, 2013).

Little or non-toxic to non-target organisms; the narrow area of toxic action; residues have no known adverse effects on the environment; reduce chemical insecticide use; self-perpetuating under ideal environmental conditions are some of the benefits of EPF (Maina *et al.*, 2018). Sub-lethal effects of fungal pathogens can have important conceptions on the population density of related hosts, which finally contributes to the status of the target organisms as a pest (Arthurs and Thomas, 2000; Blanford and Thomas, 2001; Hornbostel *et al.*, 2004; Quesada-Moraga *et al.*, 2004; Seyed-Talebi *et al.*, 2012).

Demographic toxicology has been suggested as the best way to evaluate total effects (lethal and sub-lethal) of entomopathogens, because it is based on both survivorship and fecundity parameters (Stark and Banks, 2003; Sedaratian *et al.*, 2013, 2014; Liu *et al.*, 2019). Considering the potential effects of sub-lethal exposure to EPF (*B. bassiana*) on pest populations, this study's objective is to improve the management of *T. urticae*.

Materials and methods

Host plant and mites

Host plant (*Cucumis sativus* L. cv. Veolla F₁ (Cucurbitaceae)) was grown in plastic pots (10 cm height, 5 cm diameter) under controlled greenhouse condition (25 ± 5°C, 60 ± 10% RH, and a photoperiod of 16: 8 (L: D) hours). To prevent unwanted infestations, all plants were maintained in mesh cages (1.5 × 1 × 1 m). The initial population of *T. urticae* was collected from infested greenhouses of Pakdasht (South-Eastern part of Tehran, Iran); which had been never exposed to pesticides. The spider mites were transferred to the greenhouse conditions and released on cucumber plants after being identified in the laboratory.

Fungus

In the current study, a native strain (TV) of *B. bassiana* (soil origin) was obtained from the College of Agriculture and Natural Resource, University of Tehran (Tehran, Iran). This strain was grown on Potato Dextrose Agar (PDA) and maintained at 25 ± 2°C, 70 ± 5% RH, and darkness photoperiod for 2 weeks to reach the sporulation stage. Cultures were scrapped after sporulation (≈14 days) and conidia were used for experimentations (Goettel and Inglis, 1997). To prepare a homogenous suspension of conidia, the conidia harvested from the surface of the Petri dishes vortexed with distilled water in a 100 ml tube for 20 min. The mixture is passed through Whatman No. 1 filter paper to separate the spores. The spore concentration was then determined using a hemocytometer slide under a microscope (ZIESS®) with 40× magnification (Sumikarsih *et al.*, 2019).

Bioassay

In bioassays, the following were used: seven fungal concentrations (10², 10³, 10⁴, 10⁵, 10⁶, 10⁷, and 10⁸ conidia ml⁻¹) that the mortality covering the range of 10–90%, and control treatment, where adult was treated with distilled water +0.02% Tween-80. Additionally, 20 same-aged adult mites (24 h-old ten males and ten females) were placed on the treated leaf discs (4 cm diameter) for each concentration by using a soft pointed brush. Then, the leaf discs of cucumber were sprayed (Posada *et al.*, 2007) with one of the different concentrations (10², 10³, 10⁴, 10⁵, 10⁶, 10⁷, and 10⁸ conidia ml⁻¹ and control treatment) of *B. bassiana* and then, were dried for 30 min at room conditions. After this period of time, the Petri dishes lid was closed and the leaf discs were transferred to the incubator. The experiments were conducted at the laboratory conditions of 25 ± 2°C, 70 ± 5% RH and 16:8 (L:D) hours photoperiod.

Twenty-four hours later, the lids were replaced with new ones which had a hole in their center (4 cm diameter). These holes were covered with fine net mesh. After the leaf discs were inoculated with different concentrations of *B. bassiana*, initial mortality was counted after 5–7 days. Mites were considered dead when they did not move after stimulation. Each bioassay was replicated four times for each of the seven concentrations and control.

Life-table assay

To assess the sub-lethal effects of *B. bassiana* (LC₁₀, LC₂₀, and LC₃₀) on the biological parameters of *T. urticae*, 50 adult mites were transferred into fresh cucumber leaf discs (4 cm diameter), each of which was placed on a wet cotton in a Petri dish. The leaf discs were treated with distilled water and sub-lethal concentrations of *B. bassiana* (table 1). After 72 h, dead mites were removed and surviving females were separately transferred onto untreated discs (4 cm diameter) and allowed to oviposit for 24 h. After this period of time, one egg was selected randomly and transferred into a new leaf disc. In this way, 70 eggs were used for the experiments in each sublethal concentration. All experiments were conducted in a growth chamber at 25 ± 2°C, 70 ± 5% RH, and a photoperiod of 16:8 (L:D) h, while the development and survival time were checked daily. After adult emergence, female individuals were coupled with males from the same treatment. Adult longevity and their survivorship were documented every day. Daily fecundity of female mites was also recorded. Observations were performed until the death of the last individual.

Data analysis

To calculate the sub-lethal concentrations of *B. bassiana* and their 95% fiducial limits, Probit analysis was performed (SPSS ver. 19.0). The original raw data for all individuals were analyzed according to the age-stage, two-sex life table procedure (Chi and Liu, 1985; Chi, 1988), and computer program, TWSEX

Table 2. The effect of different concentrations of *Beauveria bassiana* on the duration of different life stages (mean \pm SE) of *Tetranychus urticae*

Life stages and fecundity	Control	LC ₁₀	LC ₂₀	LC ₃₀
Male				
Egg (day)	4.1 \pm 0.1b	4.6 \pm 0.11a	4.6 \pm 0.12a	4.2 \pm 0.1b
Larva (day)	2.0 \pm 0.0a	2.0 \pm 0.08a	1.8 \pm 0.08a	1.8 \pm 0.0a
Protonymph (day)	2.0 \pm 0.1a	2.0 \pm 0.08a	1.8 \pm 0.09 a	1.7 \pm 0.1a
Deutonymph (day)	1.8 \pm 0.1a	1.7 \pm 0.10a	1.6 \pm 0.12a	1.5 \pm 0.1a
Adult lifespan (day)	11.0 \pm 0.2a	9.2 \pm 0.21b	7.8 \pm 0.29c	7.0 \pm 0.2c
Total life span (day)	20.9 \pm 0.3a	19.4 \pm 0.43b	17.8 \pm 0.38c	16.3 \pm 0.3d
Female				
Egg (day)	4.4 \pm 0.0a	4.2 \pm 0.0ab	4.1 \pm 0.0ab	4.0 \pm 0.0b
Larva (day)	2.3 \pm 0.0a	2.1 \pm 0.0ab	2.0 \pm 0.0b	1.8 \pm 0.0b
Protonymph (day)	2.1 \pm 0.0a	2.0 \pm 0.0a	1.8 \pm 0.0ab	1.7 \pm 0.0b
Deutonymph (day)	2.0 \pm 0.0a	1.9 \pm 0.0ab	1.7 \pm 0.0b	1.7 \pm 0.0b
Adult lifespan (day)	12.3 \pm 0.1a	11.8 \pm 0.1a	10.4 \pm 0.1b	9.6 \pm 0.1c
Total life span (day)	23.3 \pm 0.2a	22.0 \pm 0.1b	20.2 \pm 0.1c	19.0 \pm 0.2d

Means within a row followed by the same letter are not significantly different (paired bootstrap test, $P < 0.05$).

MSChart (Chi, 2019). All population parameters including gross (GRR) and net (R_0) reproductive rates, intrinsic (r) and finite (λ) rates of increase as well as mean generation time (T) were estimated by bootstrap procedure using $\times 100,000$ samples. Furthermore, the paired bootstrap test was used to evaluate statistical differences among different biological parameters of *T. urticae* (Efron and Tibshirani, 1993; Huang and Chi, 2012). The paired bootstrap test based on the confidence interval of differences was used to assess the differences between treatments (Wei et al., 2020; Yang et al., 2020).

Results

Concentration-response bioassay

The estimated LC₅₀ for the *B. bassiana* was 11.70×10^5 conidia ml⁻¹; while no mortality was recorded at control treatment. In addition, the values of LC₁₀, LC₂₀, and LC₃₀ were 6.77×10^2 , 8.75×10^3 , and 55.38×10^3 conidia ml⁻¹, respectively (table 1).

Sub-lethal effects on different developmental stages and fecundity

Table 2 presents the sub-lethal effects of *B. bassiana* on different developmental stages of both female and male individuals of *T. urticae*. Based on the results, a significant difference was observed between male (egg stage) and female individuals (in the duration of egg [$F = 7.15$, $P = 0.0003$, $df = 3,68$; female: $F = 4.76$, $P = 0.0033$, $df = 3,168$], larvae [male: $F = 1.35$, $P = 0.26$, $df = 3,68$; female: $F = 9.65$, $P < 0.0001$, $df = 3,168$] and protonymph [male: $F = 1.87$, $P = 0.14$, $df = 3,68$; female: $F = 7.34$, $P < 0.0001$, $df = 3168$] as well as deutonymph [male: $F = 1.99$, $P = 0.12$, $df = 3,68$; female: $F = 6.55$, $P < 0.0001$, $df = 3,168$] stages). The higher values for developmental stages at male and female individuals were observed in control and LC₁₀ concentration. When the individuals were treated with LC₃₀ of *B. bassiana*, males and adult lifespan of females, as well as total life span, were significantly affected. Adult lifespan and total life span of female individuals

had the lowest values when LC₃₀ treatment was applied (table 2).

Ovipositional period and total fecundity

The females treated by the LC₂₀ and LC₃₀ concentrations of *B. bassiana* had significant shorter total pre-oviposition periods (APOP: the duration from female emergence to first oviposition; TPOP: duration from egg to first oviposition) compared to the other treatments. The value of TPOP ranged from 10.3 to 11.9 days at LC₃₀ and Control treatments, respectively.

The adult pre-ovipositional period (APOP) was not significantly affected by different concentrations of *B. bassiana* (table 3). The lowest duration of ovipositional period was 7.5 days at LC₃₀. *Beauveria bassiana* had significant sub-lethal effects on the total fecundity of *T. urticae* (table 3). Fecundity of female individuals was lowest at LC₃₀ (18.2 eggs); whereas at Control treatment it was markedly higher (45.1 eggs).

Age-specific survivorship, age-specific and age-stage specific fecundity

The age-stage specific survival rates (s_{xj}) of *T. urticae* in different treatments are presented in fig. 1. Overlap between different developmental stages of *T. urticae* at different treatments was related to the variation of development rate of these biological stages (figs 1a–d). Age-specific survivorship (l_x), age-specific fecundity (m_x), and age-stage specific fecundity (f_{xj}) of *T. urticae* at different concentrations of *B. bassiana* are plotted in fig. 2. Total lifetime for *T. urticae* was 26, 25, 23, and 22 days in Control, LC₁₀, LC₂₀, and LC₃₀ treatments, respectively. The highest value of m_x for control mites was 3.7 eggs/female which was observed on day 17. However, maximum values of m_x for LC₁₀, LC₂₀, and LC₃₀ treatments were 3.1, 2.6, and 2.0 eggs/female which occurred on days 16, 15, and 13, respectively (fig. 2).

Table 3. Mean (\pm SE) reproductive period and total fecundity of offspring from females of *Tetranychus urticae* for control and different concentrations of *Beauveria bassiana*

Parameters	Control	LC ₁₀	LC ₂₀	LC ₃₀
APOP (day) ^a	1.0 \pm 0.0a	1.0 \pm 0.0a	1.0 \pm 0.0a	1.0 \pm 0.1a
TPOP (day) ^b	11.9 \pm 0.2a	11.4 \pm 0.1a	10.7 \pm 0.1b	10.3 \pm 0.1b
Ovipositional period (day)	10.3 \pm 0.1a	9.8 \pm 0.1b	8.3 \pm 0.1b	7.5 \pm 0.1c
Fecundity (egg)	45.1 \pm 0.7a	36.2 \pm 0.7b	23.9 \pm 0.5c	18.2 \pm 0.4d

Means within a row followed by the same letter are not significantly different (paired bootstrap test, $P < 0.05$).

^aAPOP = adult pre-ovipositional period (the duration from adult emergence to the first oviposition).

^bTPOP = total pre-ovipositional period (the time between egg to the first oviposition).

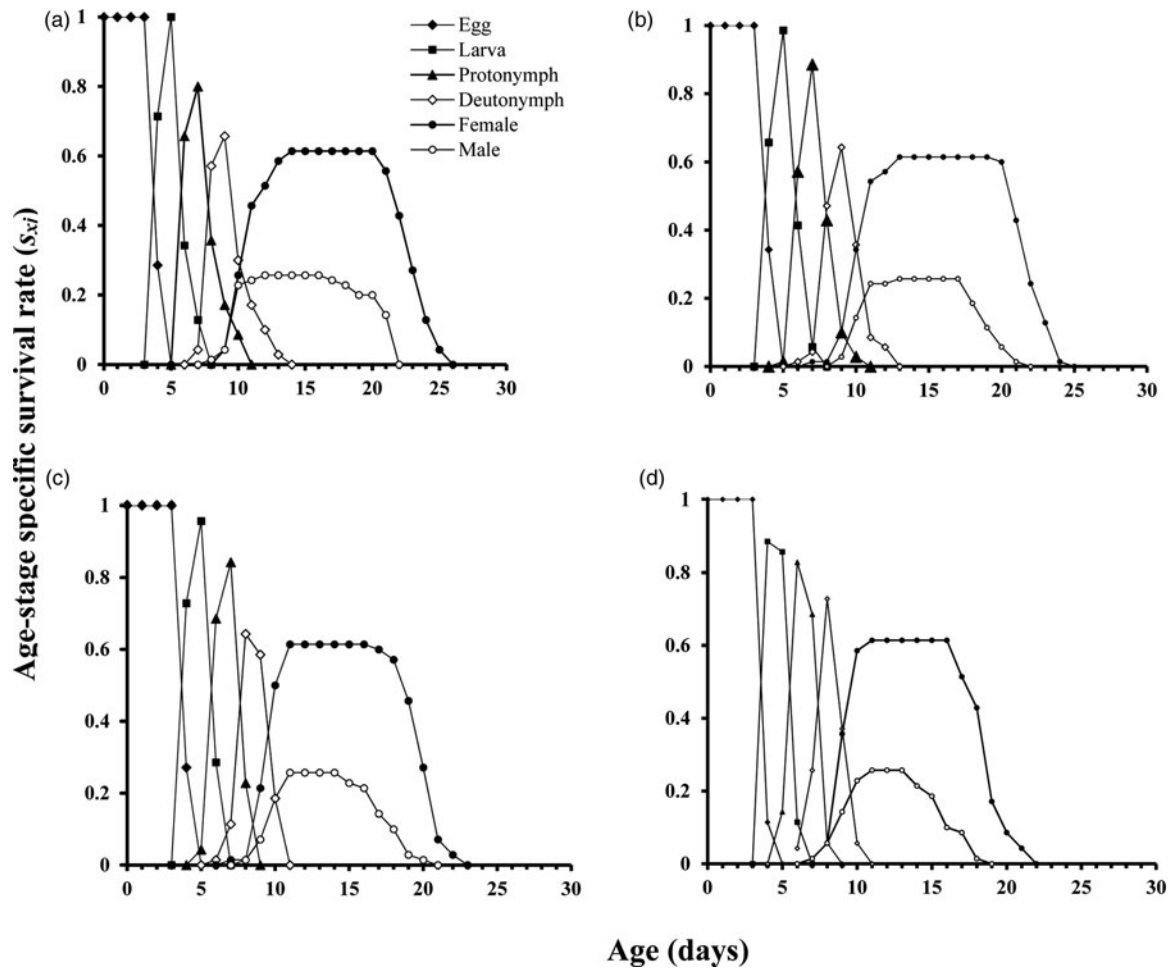


Fig. 1. The age-stage specific survival rate (s_{xj}) of *Tetranychus urticae* at different sub-lethal concentrations (a = Control, b = LC₁₀, c = LC₂₀, and d = LC₃₀) of *Beauveria bassiana*.

Population parameters

As displayed in table 4, different treatments drastically affected the population parameters of *T. urticae*. The gross reproductive rate (GRR) varied from 15.3 to 35.8 offspring/individual at LC₃₀ and control treatments, respectively. Also, the lowest value of net reproductive rate (R_0) was obtained for the mites exposed to the LC₃₀ treatment. The assessed intrinsic rate of increase (r) for mites influenced by sub-lethal concentrations ranged from 0.168 (day^{-1}) at LC₃₀ to 0.198 (day^{-1}) in the control, respectively. The finite rate of increase (λ) indicated a significant difference

with increasing concentration from control to LC₃₀. The mean generation time (T) had the highest value at control (16.7 days); followed by LC₁₀, LC₂₀, and LC₃₀ (table 4).

Discussion

Although pesticides are considered as an economic, labor-saving, and efficient tool of pest management with great popularity in most sectors of the agricultural production, several undesirable effects have restricted their applications in modern agricultural systems (Kaplan *et al.*, 2012). Accordingly, there is a critical

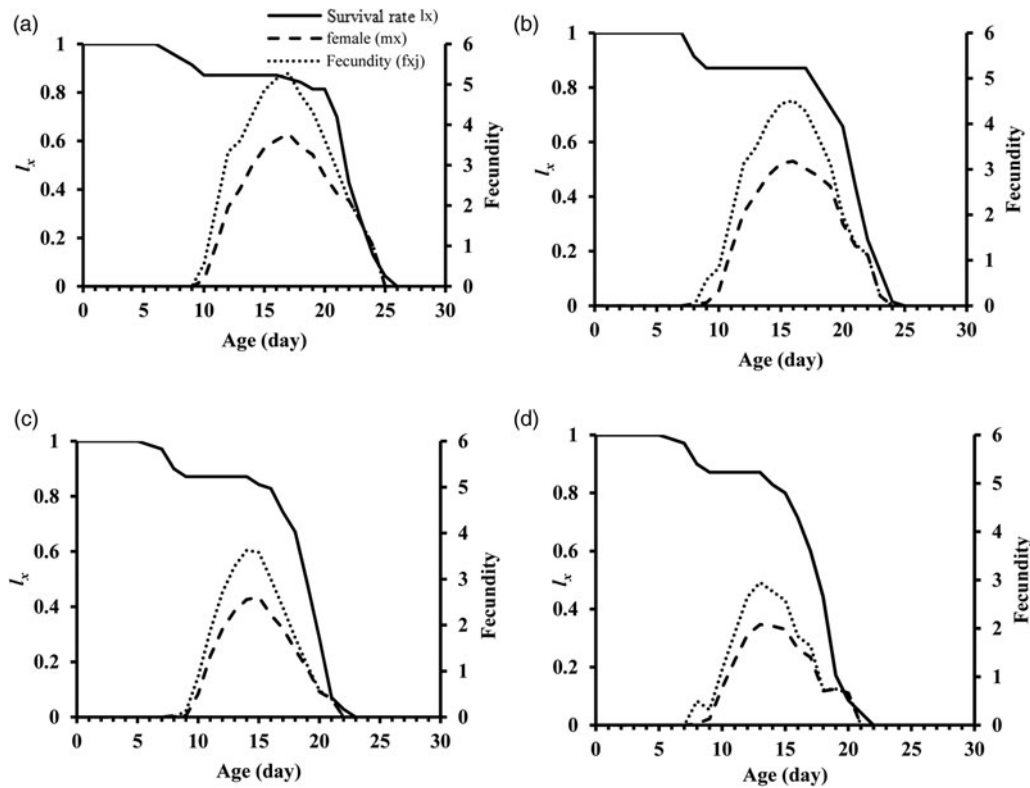


Fig. 2. Age-specific survivorship (l_x), age-specific fecundity (m_x), and age-stage specific fecundity (f_{xj}) of *Tetranychus urticae* at different sub-lethal treatments (a = Control, b = LC₁₀, c = LC₂₀, and d = LC₃₀) of *Beauveria bassiana*.

Table 4. The effects of different treatments of *Beauveria bassiana* on the population parameters (mean \pm SE) of *Tetranychus urticae*

Treatments	Gross reproduction rate (GRR)	Net reproductive rate (R_0)	Intrinsic rate of increase (r)	Finite rate of increase (λ)	Mean generation time (T)
Control	35.8 \pm 2.5a	27.7 \pm 2.6a	0.198 \pm 0.003a	1.219 \pm 0.01a	16.7 \pm 0.1a
LC ₁₀	28.3 \pm 1.9b	22.2 \pm 2.1a	0.194 \pm 0.005a	1.214 \pm 0.04a	15.9 \pm 0.1b
LC ₂₀	18.9 \pm 1.4c	14.7 \pm 1.4b	0.179 \pm 0.004b	1.196 \pm 0.05b	14.9 \pm 0.1c
LC ₃₀	15.3 \pm 1.1c	11.1 \pm 1.0b	0.168 \pm 0.007b	1.183 \pm 0.02b	14.2 \pm 0.1d
Unit	Offspring/individual	Offspring/individual	Day ⁻¹	Day ⁻¹	Day ⁻¹

Means within a column followed by the same letter are not significantly different (paired bootstrap test, $P < 0.05$).

demand to find reliable alternatives for the sustainable management of phytophagous pests. Chandler *et al.* (2000) reviewed the opportunities of exploiting entomopathogens for biological control of phytophagous mites. However, consideration of EPF effects should be assessed further to specify its influence on offsprings' life history, including growth, development, and reproduction (Latifian *et al.*, 2010). Numerous studies have been conducted for evaluating the lethal and sub-lethal effects of different EPF such as *B. bassiana* and *Metarhizium anisopliae* Metchinkoff on *Tetranychus* species (Wekesa *et al.*, 2006; Shi and Feng, 2009; Seyed-Talebi *et al.*, 2014; Wu *et al.*, 2016). However, no evidence is available regarding the sub-lethal effects of *B. bassiana* on the biological attributes (such as survival, longevity, fecundity, and demographic parameters) of *T. urticae*.

Our findings revealed an obvious variation in susceptibility at different life stages of *T. urticae* to *B. bassiana*. In contrast with our results, Zhou *et al.* (2010) concluded that sub-lethal doses

of *Isaria fumosorosea* Wize had no significant effects on developmental stages of *Axinoscymnus cardilobus* Ren and Pang (Coleoptera: Coccinellidae).

The adult longevity and total lifespan for both sexes were significantly lower in fungus-treated individuals than Control. Our results are in agreement with those reported by Irigaray *et al.* (2003) and Gatarayiha *et al.* (2010) which reported that *B. bassiana* had greater deleterious effects on adult females of *T. urticae* than immature stages. Afifi *et al.* (2010) reported that mite mortality was increased with increasing conidia concentration and exposure time to *B. bassiana*. Their results indicated that 14 days after treatment with 2×10^6 and 2×10^8 conidia ml⁻¹, an average mortality of 62.5 and 83.3% for *Panonychus ulmi* (Koch) and 82.6 and 91.7% for *Phyllocoptura oleivora* Ashmed were recorded. On the other hand, *B. bassiana* strain 447 had no significant effects on *T. urticae* under laboratory conditions, even when applied at a concentration of

1×10^9 conidia ml^{-1} (Antonio *et al.*, 1999). Several factors may be responsible for this variation in the efficacy of *B. bassiana* against spider mites, including strain identity, concentrations used, experimental conditions, host species and life stages, exposure time, etc.

Our study showed that sub-lethal concentrations of *B. bassiana* drastically reduced reproductive parameters of *T. urticae* in comparison with untreated individuals. The present study, however, exhibited that *B. bassiana* had no detectable effects on pre-ovipositional period of *T. urticae*. Our findings indicated negative effects of sub-lethal treatments on ovipositional period and total fecundity of *T. urticae*. These observations are in line with those reported by Shi and Feng (2009) in the case of *B. bassiana* strain Bb2860, *Paecilomyces fumosoroseus* and strain Pfr116 and *M. anisopliae* strain Ma759 on the *T. urticae*. Similarly, Ullah and Lim (2017) illustrated that *B. bassiana* strain GHA significantly decreased the fecundity of *T. urticae*. This evidence might be derived from a decrease in the female physiological state related to: (i) fungal colonization of tissues such as fat body (source of vitellogenins) and ovaries (Blay and Yuval, 1999) and (ii) fungal toxin production to transcend insect cellular and humeral immune reactions (Inglis *et al.*, 2001; Quesada-Moraga *et al.*, 2004). It has been shown that applying the sub-lethal concentrations of *B. bassiana* strain GZGY-1-3 had no significant effects on the fecundity of *Frankliniella occidentalis* (Pergande) (Zhang *et al.*, 2015).

The life table technique has been applied as a reliable procedure for assessing the population dynamics of both phytophagous organisms and their natural enemies (Biondi *et al.*, 2013; Cira *et al.*, 2017; Nawaz *et al.*, 2017). Regarding the curves of the age-specific survivorship (l_x) and age-specific fecundity (m_x) of *T. urticae* at different treatments, increasing sub-lethal concentrations had detectable effects on the mortality rate and fecundity of this pest. Similar reports were published for the survivorship schedules of *T. urticae*, *F. occidentalis*, *Encarsia formosa* Gahan, and *Planococcus citri* Risso when treated by *B. bassiana* strain 432.99, *B. bassiana* strain SZ-26, and *Lecanicillium longisporum* strain LRC190 and *L. longisporum*, respectively (Chandler *et al.*, 2000; Wu *et al.*, 2014; Fazeli-Dinan *et al.*, 2016; Ghaffari *et al.*, 2017). Among different population parameters, the intrinsic rate of increase (r) has been considered as the most applicable index that takes both fecundity and survivorship of individuals into account. The results obtained herein revealed that higher concentrations of *B. bassiana* obviously affected the r value of *T. urticae*. Likewise, Seyed-Talebi *et al.* (2012) and Rashki and Shirvani (2013) reported that the r and λ values were significantly lower in *T. urticae* and *Aphis gossypii* Glover which were treated with strains EUT105 and DEBI008 of *B. bassiana*, respectively. Despite our results, Baverstock *et al.* (2006) showed that *B. bassiana* has no significant effects on the r value of pea aphid, *Acyrtosiphon pisum* (Harris). It can also be hypothesized that feeding deficiency caused by fungal infection may drastically affect the reproductive potential of female individuals which have great energetic demands (Yuan *et al.*, 2018). In addition to these parameters, the values of net (R_0) and gross (GRR) reproduction rates and mean generation time (T) in treated individuals were also inferior to Control treatment. Similar to our findings, Huang *et al.* (2010) and Yuan *et al.* (2018) documented significant effects of different concentrations of *I. fumosoroseus* and *B. bassiana* strain JLGZL-14 on the R_0 and T values of *Bemisia tabaci* (Gennadius) and *Phthorimaea operculella* Zeller, respectively.

In conclusion, the present study displayed that *B. bassiana* not only has considerable pathogenicity to *T. urticae*, but also causes different sub-lethal effects such as reducing the total life-span as well as total fecundity of female individuals. Furthermore, our results clarified the considerable potential of *B. bassiana* strain TV, as an efficient biocontrol agent for sustainable management of *T. urticae*. More attention, however, should be devoted to investigate these effects under semi-field and field conditions.

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