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Thermal Effect on the Biology and Life Tables of Bemisia tabaci Gennadius (Homoptera: Aleyrodidae)

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Abstract: The life history and life table of *Bemisia tabaci* Gennadius on cucumber was studied under laboratory conditions at 20, 25 and 30°C. The overall developmental time varied from 34.8 days at 20°C to 14.1 days at 30°C. Immature mortality decreased from 45.8 to 17.3% with increasing temperature. The threshold temperatures of egg, 1st, 2nd, 3rd and 4th nymphal stage and a generation were 14.72, 14.36, 10.18, 11.40, 14.36 and 13.07°C whereas the degree-day requirement at each stage was 64.44, 42.39, 49.19, 33.19, 35.46 and 229.52 DD, respectively. Female longevity ranged from 16.8-34.1 days. Mean total fecundity ranged from 150-263 eggs/female. Mean daily fecundity ranged from 4.2-12.7 eggs/female, increasing with increasing temperature. Values for r_m varied from 0.066 to 0.191 being least at 20°C and greatest at 30°C. Generation times decreased from 43 to 19 days with increasing temperature. The results indicate that *B. tabaci* is well adapted to high temperatures and may extend its distribution if mean world temperatures increase as a result of global warming.

Key words: Bemisia tabaci, cotton whitefly, biology, life table, thermal, cucumber

INTRODUCTION

The cotton whitefly, Bemisia tabaci Gennadius (Homoptera: Aleyrodidae) is an important pest on various crops. Its first record as a pest was reported in cotton fields of Greece in 1889 (Cock, 1993). It is a polyphagous species with a world wide distribution (Mound and Halsey, 1978; Cock, 1993; Mckenzie et al., 2004). Damage caused includes reduction in yield and fruit quality as well as virus transmission (Byrne et al., 1990; Oetting and Buntin, 1996; Schuster et al., 1996). In addition, B. tabaci is a polymorphic species. It is variable not only in its morphology (size and form of nymphs) (Mound and Halsey, 1978; Bethke et al., 1991) but also in its ecological characteristics (environmental requirements, development, fecundity, insecticide resistance, virus transmission, natural enemy complexes and endosymbiont complement (Xu et al., 2003; Al-Zeyoud and Sengonca, 2004; Horowitz and Ishaya, 1996; Markham et al., 1996; Rowland et al., 1991; Costa et al., 1993a, b; Bedford et al., 1994; Costa et al., 1995; Kirk et al., 2000). Therefore, for each pest management program, an exact determination of the crucial population parameters is required.

In Khouzestan province (Southwestern of Iran) B. tabaci utilizes several key plant species, particularly cucumber and other cucurbitaceous plants for feeding and reproduction. Very little information is available

regarding life history traits of *B. tabaci* population infesting cucurbitaceous host plants in this area (Kocheili *et al.*, 2005). In the present study, the effect of temperature on development, mortality, longevity, fecundity, sex ratio and life table characteristics of *B. tabaci* on cucumber was investigated.

MATERIALS AND METHODS

Stock culture maintenance: Adult cotton whitefly, *Bemisia tabaci* used in these studies were collected by an aspirator in September 2006 from a cucumber (*Cucumis sativus* L.) field near Ahwaz, Iran and reared on the foliage of cucumber plants (cultivar Superdominus) grown from seeds in plastic pots (10 cm diameter). Infested plants were kept in wooden-framed rearing cages (120×60×60 cm) covered with white nylon mesh of 210 μm apertures. They were maintained in a laboratory with seasonal temperature ranging from 16-25°C and relative humidity 40-50%. The photoperiod was 16:8 (L:D) h, with illumination (4000 lux) provided from fluorscent lamps. Plant were kept in the cages until they were severely damaged by the whiteflies, new plants being added when needed.

Development and mortality: Forty to fifty adult whiteflies of both sexes (roughly 50% male: 50% female) were placed

each clip cages attached to the underside of expanding cucumber leaves. The clip cage were similar those described by Lewis (1973) with some modifications. The clip cage was made of a plastic vial (1.5) cm diameter, 1 cm length). The bottom of the cage was covered with nylon organdy for ventilation. A 0.6 cm hole in the body of the cage served as an entrance for the whiteflies. The hole was accommodated by a small cork stopper. The test plants were maintained at 25°C for 24 h to allow deposition of eggs, after which the clip cages and adult whiteflies were removed. The plants were then placed in temperature-controlled cabinets at three constant temperatures: 20, 25 and 30±1°C. Relative humidity was kept at a minimum of 50% and a photoperiod of 16:8 (L:D) was maintained with light intensity set at 1800 lux. The plants were watered as needed. The development of each life stage, determined by measuring the size of the insect (Nechols and Tauber, 1977), was monitored with a dissection microscope until adult eclosion. The duration of immature stages and mortality were recorded daily.

Longevity, fecundity, sex ratio and population parameters: The effect of different temperatures on longevity, fecundity and other population parameters was studied by confining one adult female and one adult male immediately on emergence on the undersurface of a cucumber leaf by means of a plastic clip cage. Every other day, whiteflies were transferred to a new leaf and the eggs laid were counted. Any male insect that was accidentally lost, or that died before the female, was replaced. Each experiment was terminated with the natural death of the female. All eggs laid in fecundity experiment were observed daily until adult eclosion and the numbers of male and female whiteflies were recorded to determine the sex ratio.

Statistical analysis: Analysis of variance (ANOVA) and Duncan multiple range tests was used to examine differences in developmental times, longevity and fecundity across temperatures (SAS Institute, 1997). A series of Chi-square tests were conducted to determine if there were any significant differences in stage mortality for insects reared at different temperatures.

The reciprocals of the observed developmental times, in days, provided developmental rates for each stage at each temperature. Lower developmental threshold temperatures were estimated by the X-intercept method of Arnold (1959). The mean number of degree-days (DD) required for development of each life stage was calculated using the equation:

DD = D(T-t)

where:

D = The developmental time (days)

 Γ = The temperature (°C) during development

t = The lower developmental threshold (°C) (Price, 1984).

A life table using time-specific survival rates (l_x) and fecundity (m_x) for each 24 h period was constructed for calculating the following life table parameters (Southwood, 1978).

Net reproduction rate of increase: $R_o = \sum l_x m_x$; Mean generation time (in days): $T = \sum l_x m_x / R_o$; Intrinsic rate of increase: $r_m = \ln R_o / T$; Finite rate of increase: $\lambda = e^{rm}$; Population doubling time: $DT = \ln 2/r_m$. Data on the different treatments were compared with the Jacknife program developed by Maia *et al.* (2000).

RESULTS AND DISCUSSION

Development and mortality: The developmental times for all immature stages were inversely proportional to temperature. The duration of the egg stage varied from 11.86 days at 20°C to 4.18 days at 30°C. Mean development times for first-to third-instar B. tabaci were usually slightly shorter than for its egg and fourth instar (pupal) development times. Some authors have reported that the egg and fourth-instar are the longest stages for other aleyrodids (Powell and Bellows, 1992a; Roermond and van Lenteren, 1992; Shishehbor and Brennan, 1995). It is of interest to know which life stages are longest when making pest management decisions, such as which biological control agents to use (e.g., egg parasitoid versus nymphal parasitoid) or which pesticide is most appropriate (e.g., one with ovicidal versus insecticidal properties) (Leddy et al., 1995).

The mean total developmental period for *B. tabaci* varied from 34.84 days at 20°C to 14.10 days at 30°C (Table 1). Analysis of variance indicated significant differences in development time between the temperature examined (F = 1500; df = 2, 57; p = 0.001). In a comparable study Powell and Bellows (1992a) obtained a total development time of 38.20, 20.22 and 17.36 days on cucumber at 20, 25.5 and 29°C, respectively, which is longer than the results in our study. The discrepancy between the two studies may be due to differences in whitefly populations and experimental conditions (host plant cultivar). Bethke *at al.* (1991) examined biology, morphometrics and development of two populations of *B. tabaci* on cotton and poinsettia. Their study indicated

Table 1: Developmental times of the immature stages of *Bemisia tabaci* in days (Mean±SE) on cucumber

	Temperature (°C)				
Stages	20	25	30		
Egg	$11.86\pm0.19a$	6.50±0.06b	$4.18\pm0.09c$		
First instar	$6.18\pm0.13a$	5.00±0.13b	$2.50\pm0.09c$		
Second instar	$5.75\pm0.13a$	$2.75\pm0.11b$	$2.63\pm0.09b$		
Third instar	$3.03\pm0.10a$	2.30±0.08b	$2.27\pm0.07b$		
Fourth instar	7.16±0.23a	$2.77\pm0.08b$	$2.32\pm0.09c$		
Total	34.84±0.25a	19.23±0.13b	$14.10\pm0.11c$		
N^a	(59)	(62)	(52)		

Means in each row followed by the same letter(s) were not statistically significantly different (p>0.01); *: Sample size (N) in parenthesis

Table 2: Developmental rates (Y; 1/day) for immature instars regressed on constant temperatures (X), stimated lower developmental thresholds (t) and mean numbers of Degree-Days (DD) required for development of the immature stages of *B. tabaci*

	Rate (Y) regressed on temperature (X) (°C) $Y = -a+bX$				
Stages	a	b	\mathbb{R}^2	t (°C)	DD±SE
Egg	0.228	0.015	0.99	14.72	64.44±1.24
First instar	0.342	0.024	0.87	14.36	42.39±5.54
Second instar	0.210	0.021	0.80	10.18	49.19±4.28
Third instar	0.130	0.011	0.77	11.40	33.19±4.80
Fourth instar	0.418	0.029	0.92	14.36	35.46±3.10
Total	0.055	0.004	0.99	13.07	229.52±13.8

that, based on both morphometric and fecundity differences, there are distinct populations exploiting both cotton and poinsttia.

The lower threshold temperatures differed among the developmental stages (Table 2). Overall, the lower threshold for complete development of *B. tabaci* was apparently higher than 13.07°C. The predicted minimum thresholds of *B. tabaci* is lower than other reports for this aleyrodid. Powell and Bellows (1992 a) observed the lower thresholds for development of *B. tabaci* to be 14.65°C on cotton and 16.71°C on cucumber. However, Gerling *et al.* (1986) reported that the lower developmental threshold for *B. tabaci* is 11°C.

The mean number of degree-days required by *B. tabaci* to complete its development was 229 DD (Table 2). This is higher than that of Powell and Bellows (1992 a) for *B. tabaci* (195 DD) on the same host plant. The differences could be partly due to the lower minimum threshold temperature of *B. tabaci* in this study.

There were significant differences in mortality for different temperature regimes for eggs ($\chi^2 = 52$; df = 3; p = 0.04) and the first instar ($\chi^2 = 7.8$; df = 2; p = 0.04) but not for the second ($\chi^2 = 0.193$; df = 2; p = 0.97), third ($\chi^2 = 1.29$; df = 2; p = 0.73) and fourth ($\chi^2 = 0.29$; df = 2; p = 0.96) instars (Table 3). Total mortality was highest at 20°C (45.8 %), declined as temperature increased from 20 to 30°C (17.3 %). Present result agree with those of Powell and Bellows (1992a) who stated that total pre-adult mortality of *B. tabaci* decreased as temperature increased (from 20 to 32°C).

Table 3: Percentage mortality within immature stages of *Bemisia tabaci* reared on cucumber at different temperatures

Stages	Temperature (°C)			
	20	25	30	
Egg	25.4 (15)	6.5 (4)	3.8 (2)	
First instar	13.6(6)	3.4(2)	0.0(0)	
Second instar	5.3(2)	7.1 (4)	4.0(2)	
Third instar	5.6(2)	1.9(1)	6.3 (3)	
Fourth instar	5.9(2)	3.9(2)	4.4 (2)	
Total	45.8 (59)	20.9 (62)	17.3 (52)	

Sample size (n) in parenthesis is number dying in each stage except for the total which is the initial number entering the egg stage $\frac{1}{2} \int_{-\infty}^{\infty} \frac{1}{2} \left(\frac{1}{2} \int_{-\infty}^{\infty} \frac{$

Table 4: Longevity in days (Mean±SE, range, (N)) of adult female and male Bemisia tabaci on cucumber at three different temperature

Sex	Temperature (°C)		
	20	25	30
Female	34.14±3.13a	26.75±1.95b	16.88±1.20c
Range	15-51	9-39	2-23
(N) ^a	(14)	(16)	(17)
Male	29.50±2.99a	17.50±2.24b	11.23±0.79c
Range	7-42	7-37	5-15
(N) a	(14)	(16)	(17)

Means in each row followed by the same letter(s) were not statistically significantly different (p>0.01); *: Sample size (N) in parenthesis

Longevity: An inverse relationship exists between temperature and mean adult longevity of B. tabacia across the full temperature range investigated (Table 4). ANOVA indicated that temperature was a highly significant factor affecting the longevity of both females (F = 81.16; df = 2, 74; p = 0.0001) and males (F = 383.35; df = 2, 36; p = 0.0001). The maximum longevity observed for an individual whitefly was 51 days for a female at 20° C.

The longevities of adult B. tabaci determined in the present study were greater than those reported in other studies conducted at similar constant temperatures. Three studies (Butler et al., 1983; Powell and Bellows, 1992b; Fekrat and Shishelbor, 2004) conducted on cotton, cucumber and aubergine, respectively, at constant temperature reported adult longevity shorter than those obtained in the present study. Butler et al. (1983) reported that at 26.7 and 32.2°C females lived 8.0 and 10.4 days and males 7.6 and 11.7 days, respectively. Powell and Bellows (1992b) reported that females lived 24.6, 15.5 and 9.64 days and males 18.6, 12.23 and 7.03 days at 20, 25.5 and 29°C, respectively. Fekrat and Shishehbor (2004) found that at 20, 25 and 30°C females lived 18.14, 13.14 and 8.0 days and males 12.71, 9.78 and 5.92 days, respectively. As in the present study, most researchers reported that female insects lived longer than males (Butler et al., 1983; Hendi et al. 1984; Powell and Bellows, 1992b; Fekrat and Shishebor, 2004). Differences in population of whiteflies and host plants may account for greater longevity of B. tabaci in the present study.

Table 5: Total fecundity, daily fecundity (Mean±SE, range) and sex ratio of B. tabaci females on cucumber at three different temperatures

	Temperature (°C)			
Parameters	20	25	30	
Total egg	150.29±22.8b	263.75±24.4a	204.71±16.9ab	
range (N)a	49-324 (14)	204-466 (16)	44-291 (17)	
Average eggs/day	4.24±0.44c	9.92±0.69b	12.75±0.44a	
Range	3-7	7-13	8-16	
Sex ratio (female)	$0.502\pm0.012c$	0.560±0.008b	0.599±0.120a	
Female:Male	1.04:1	1.27:1	1.49:1	

Means in each row followed by the same letter(s) were not statistically significantly different (p>0.01); "Sample size (N) in parenthesis

Fecundity and sex ratio: ANOVA indicated significant overall temperature effects on mean total fecundity (F = 22.19; df = 2, 74; p = 0.0001) and mean daily fecundity (F = 373.19; df = 2, 74; p = 0.0001). Peak egg production occurred at 30°C (204.7 eggs) (Table 5). At this temperature, the maximum number of eggs produced in a single day by an individual female was 16.

At 30°C Hendi et al. (1984) reported means of 203.1 eggs laid by B. tabaci on tomato which compares favourably with the results obtained in the present study. However, other laboratory studies have reported a variety of fecundity values for this species. Horowitz (1983) reported that B. tabaci deposited a mean total number of 95.5 eggs on cotton at 30°C. At 20, 25.5 and 29°C Powell and Bellows (1992b) found that B. tabaci laid 196.5, 175.3 and 208.6 eggs, respectively, on cucumber. At 26.7 and 32.2°C Butler et al. (1983) found that B. tabaci laid a mean of 81 and 72 eggs, respectively, on cotton. Fekrat and Shishehbor (2004) found that B. tabaci laid a mean of 78.6, 71.3 and 51.8 eggs at 20, 25 and 30°C, respectively, on aubergine. By comparison in the present study on cucumber at 20, 25 and 30°C, the mean total eggs were 150.29, 263.75 and 204.71, respectively, generally higher than most other reports. The differences may be explained by disparities in host plant suitability to B. tabaci and population differences of whiteflies used in these studies.

Daily oviposition rates of *B. tabaci* obtained in the present study were higher than the values cited by Hendi *et al.* (1984), Powell and Bellows (1992b) and Fekrat and Shishehbor (2004). Hendi *et al.* (1984) reported that *B. tabaci* female laid 8 eggs daily at 30°C on tomato. In a laboratory experiment, Powell and Bellows (1992b) reported daily egg production rates of 4.95, 5.97 and 9.71 for *B. tabaci* female on cucumber at 20, 25.5 and 29°C, respectively. Fekrat and Shishehbor (2004) reported that *B. tabaci* laid 3.9, 5.0 and 5.8 eggs daily at 20, 25 and 30°C, respectively, on aubergine.

The sex ratio varied from 1:1 (male:female) at 20°C to 1:1.49 at 30°C (Table 5). Gameel (1978) reported that the sex ratio of *B. tabaci* is usually 1:1 on cotton. Coudriet *et al.* (1986) stated that he found approximately a 1:1 sex ratio during the summer.

Table 6: Life table parameters (Mean±SD) of *B. tabaci* on cucumber at three different temperatures

	Temperature (°C)			
Parameters	20	25	30	
$r_{\rm m}$	0.066±0.003a	0.136±0.002b	0.191±0.003c	
R_0	18.000±2.660a	38.430±3.550b	40.480±3.330c	
λ	$1.068\pm0.003a$	1.145±0.003b	1.210±0.004c	
T	43.860±0.940a	26.890±0.660b	19.340±0.410c	
DT	10.460±0.440a	5.100±0.110b	3.620±0.050c	

Means in each row followed by the same letter(s) were not statistically significantly different (p>0.01)

Life table parameters: Calculated daily intrinsic rates of natural increase (r_m) ranged from 0.066 for whiteflies held at 20°C to a maximum peak rate of 0.191 at 30°C (Table 6). The finite rate of increase (λ) ranged from 1.068 times per individual per day at 20°C to 1.210 times per individual per day at 30°C (Table 6). Mean generation time (T) decreased consistently with rising temperature across the entire temperature range observed. The time required to double population number reached a minimum of only 3.62 days at 30°C.

The values of r_m of B. tabaci on cucumber at 20 and 25.5°C were found by Powell and Bellows (1992b) to be 0.062 and 0.142 day⁻¹, respectively, which corresponds well with the results at the same temperatures in the present study. However, Fekrat and Shishehbor (2004) reported that at 20, 25 and 30°C the intrinsic rate of increase and net reproductive rate of B. tabaci on aubergine were 0.081, 0.092, 0.141 and 23.80, 21.67 and 18.12 day⁻¹, respectively. The values for both rates are lower than found in the present study (Table 6), reflecting higher juvenile mortality, lower fecundity and shorter adult life span in the study of Fekrat and Shishehbor (2004). Differences in the ecological factors, viz., population of whiteflies, host plants as well as measurement methods may provide and explanation for higher r_m and R_n values for B. tabaci on cucumber than on aubergine.

The optimum temperature for development and reproduction of *B. tabaci* was 30°C, within the temperature range examined. Present data indicate that *B. tabaci* is adapted to warm areas. In the context of global warming, it would appear that *B. tabaci* has a greater chance of survival, consequently it may widen its geographical distribution and pose a new threat to agroecosystems, as a result of a greater access to new host plants.

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