Comparative Development, Survival and Fecundity of *Helicoverpa armigera* (Hubner) (Lepidoptera: Noctuidae) on Different Chickpea Cultivars

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ABSTRACT

The development, body weight, survivorship, and reproduction of the gram pod borer, *Helicoverpa armigera* (Hûbner), were evaluated in the laboratory at $25\pm5^{\circ}$ C, $65\pm5^{\circ}$ R. H. and a light : dark cycle of 16:8 h on six different cultivars of chickpea. The larval period ranged from 16.41 to 22.71 days on cultivars CMC-211s and Pb-2008 respectively. Similarly the larval period ranged 9.50 to 14.01 days on CMC-211s and Pb-2008 respectively. The average number of eggs oviposited by adults reared on Pb-2008, Bittal-98, Parbat, CM-2000, Dasht and CMC-211s were 299.68, 323.66, 353.36, 462.84, 492.28 and 522.54 respectively. The survival percentage survival of all larval instars of *H. armigera* ranged from 73% on comparatively resistant cultivars to 93% on susceptible cultivars of chickpea. Dasht, CM-2000, Parbat and Bittle-98 showed intermediate trend. Cultivar Pb-2008 was resistant as compared to other cultivars.

INTRODUCTION

Chickpea (*Cicer arietinum*) is the third most important grain legume of the World (Sharma *et al.*, 2005; Sarwar *et al.*, 2011). It is a major source of protein for the poor people of Pakistan. Its protein value ranges from 25.3 to 28.9% (Hulse, 1991; Shrestha *et al.*, 2011; Sarwar, 2013). In Pakistan, it is mainly grown in rainfed and irrigated areas of the Punjab and covers an area of 1.11 m ha with a grain yield of 475 thousand tons (Anonymous, 2008). Production constraints in chickpea primarily include insect pests and diseases (Nadeem *et al.*, 2011). Among the insect pests, *Helicoverpa armigera* (Hübner) (Lepidoptera: Noctuidae) is the major constraint in the production of crop worldwide (Sharma *et al.*, 2005; Shafique *et al.*, 2009).

H. armigera is highly polyphagous, cosmopolitan, devastating and worldwide distributed pest (Sharma *et al.*, 2005; Sarwar *et al.*, 2009). It causes yield losses in cotton, okra, tomato and in few other crops and vegetables (Saleem and Yunus, 1982). Yield losses due to this pest in chickpea may range from 70 to 95% (Prakash *et al.*, 2007). Its larvae causes serious damage to this crop during the fruiting stage. A single larva can consume many pods before reaching the pupal stage (Nadeem *et al.*, 2010). In different areas of Pakistan, 3-34% pod damage to important cultivars of chickpea was recorded (Ahmed and Hashmi, 1976).



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Key words

Helicoverpa armigera, host plant resistance, chickpea cultivars, gram pod borer.

Chemical control is a primary management tactic to control the *H. armigera*. However, the reliance on insecticides had resulted into many problems like pest resurgence, outbreak of secondary pests, environmental pollution and insecticide resistance (Kranthi *et al.*, 2002). It is reported that this pest has developed resistance to pyrethroids, organophosphates, organochlorines, and carbamates (Ahmed *et al.*, 1997, 2001). Resistance of *H. armigera* to the major classes of insecticides is threat to grain legume production in Asia (King and Sawicki, 1990). Therefore, an integrated approach for managing this pest is needed.

Host plant resistance (HPR) as one of the important component of integrated pest management, can play major role in management of *H. armigera* (Sharma *et al.*, 1999; Sarwar *et al.*, 2011; Sarwar and Sattar, 2013). It is economically reliable, ecologically safe and compatible with other IPM strategies (Sharma *et al.*, 1999, Li *et al.*, 2004; Nadeem *et al.*, 2010). HPR helps in developing cultivars that give stability to host plants against different insects. This stability is important in terms of growth, development and behavior of herbivorous insects (Umbanhowar and Hastings, 2002; Awmack and Leather, 2002).

Antixenosis and antibiosis are the mechanisms that confer resistance in plants. Antibiosis mechanism of plants may cause reduction in insect size, weight, survival, longevity, reproduction and may be resulted in long developmental time (Sarfraz *et al.*, 2006). Resistant chickpea plants were reported to show non-preference for oviposition and larval feeding by *H. armigera* (Lateef, 1985). Host plant resistance in Chicpea against *H*.

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armigera was studied by many researchers (Wakil *et al.*, 2005; Shafique *et al.*, 2008; Sarwar *et al.*, 2009; 2011; Nadeem *et al.*, 2010, 2011).

The objectives of this research were: (1) to investigate the preference/non preference of *H. armigera* larvae towards different chickpea cultivars. (2) to study the effect of different cultivars on the development, survival and fecundity of *H. armigera*.

MATERIALS AND METHODS

Collection and rearing of H. armigera

A field population of *H. armigera* (third to fifth instar larvae) was collected from tomato fields of National Agriculture Research Center Islamabad and cultured at $25\pm2^{\circ}$ C, $60\pm5\%$ RH with 16:8 (light :dark) cycle. The larvae were reared on artificial diet until pupation and then transferred to a plastic box lined with tissue paper. On adult emergence, individuals were transferred to transparent rearing jars and fed with 10% sugar solution. Nappy liner strips were provided for egg laying and eggs were collected on daily basis. The insects were continuously fed on the artificial diet for three generations before testing to reduce the possible influence of the host source of the tested insect population.

Development H. armigera fed on different chickpea cultivars

The seeds of following six different chickpea cultivars *viz.*, Bittal-98, Pb-2008, Parbat, CMC-211s, Dasht, CM-2000, were obtained from National Agriculture Research Center Islamabad. These seeds were sown in plastic pots of 16 cm diameter having sand, soil and farm yard manure in 1:1:1 ratio.

The neonates hatched on the same day (< 24 h old) were divided into six groups (as six cultivars) and randomly transferred to Petri dishes (one larval per Petri dish to prevent cannibalism). All treatments were reared at 25°C with a photoperiod of L:D 16:8 and provided with pods of one of the six cultivars. The pods were provided to larvae until pupation. Larvae were examined daily and the development time and weight of each larval instar was recorded. Total developmental time from neonate to pupation was observed. Pupal weight was recorded one day after pupation with sex determination of each insect. The pupae were placed individually in 250 ml plastic cups covered with white netting secured by a plastic band. The number of adults emerged was recorded daily and the proportion that emerged successfully was also recorded together with the total developmental time from neonate to adult.

Reproductive capacity of H. armigera

Adults that emerged from larvae reared on each cultivar were allowed to mate and then transferred to the 250 ml cups for egg laying. (One pair per cup, 10 pairs) supplied with 10% honey solution as food source. When a female started egg laying, the insect pair was transferred to a new cup every 48 h until they died. Data regarding number of eggs laid per female, pre oviposition, oviposition, and post oviposition period was recorded.

Feeding preference of H. armigera

Pods of different cultivars were placed in each Petri dish (diameter 16 cm). Fifteen Larvae were released in centre of each Petri dish. The experiment was replicated five times. After 24 hours, the feeding preference of these larvae was recorded by observing the number of larvae attracted to different pods.

Statistical analysis

Developmental time, body weight, pre-oviposition time, oviposition time, post oviposition time, fecundity and feeding preference of *H. armigera* reared on different host plants were analyzed with analysis of variance (ANOVA) by using M-STAT, and means were separated least significance difference at p<0.05.

RESULTS

Development of H. armigera on different chickpea cultivars

The mean larval duration (neonate to pupation) of H. armigera was significantly differ (F_{5.29}=1452.07, P<0.01) on the six test cultivars of chickpea (Table I). The mean development time of *H. armigera* larvae was longest on Pb-2008 with 22.71 days and shortest on CMC-211s with 16.41 days. Test cultivars had significant effect on the development time of 1^{st} instar larvae of H. armigera (F_{5.29}=169.54, P< 0.01). Maximum duration of 1st instar larvae was 2.75 days on Pb-2008. While minimum duration was 2.17days on CM-211s (Table I). There was significant differences among the development time of 2nd instar larvae on selected cultivars (F_{5.29}=226.35, P< 0.01). 2nd instar larvae reared on CMC-211s developed slowly (2.31 days) as compare to other cultivars. Maximum larval duration of 3rd instar was observed on Pb-2008 (3.53 days) followed by Bittle-98 (3.1 days) and Parbat (3.08 days) while Bittle-98 and Parbat were statistically similar to each other. Minimum larval duration of 3rd instar was observed on CMC-211s (2.62 days) followed by Dasht (2.74 days) and CM-2000 (2.84 days) (F_{5.29}=346.09, P<0.01) (Table I). Cultivars

CMC_211e	Dasht	CM-2000	Parbat	Bittal-98	Pb-2008	Cultivar	Table II Mean larv:	The means followed by di	LSD	CMC-211s 2.1	Dasht 2.2	CM-2000 2.3	Parbat 2.4	Bittal-98 2.5	РЬ-2008 2.7	Cultivar 1 ^s	Table I Mean deve
	0.28 ± 0.00^{b}	$0.25 \pm 0.00^{\circ}$	0.19±0.01 ^d	0.18±0.00°	$0.13 \pm 0.00^{\text{f}}$	1 st instar	al (1st to 6 th inst	ifferent letters in	0.05	7±0.05 ^t	3±0.03 °	3 ± 0.07^{d}	7±0.05 °	4±0.03 ^b	5±0.01 ^a	^t instar	lopment time of
	6.65 ± 0.10^{b}	$6.12 \pm 0.04^{\circ}$	4.34 ± 0.03^{d}	3.36±0.02°	$2.52\pm0.10^{\text{f}}$	2 nd instar	ar) and pupal w	the same column	0.04	2.31±0.02 °	2.52 ± 0.02^{d}	2.63±0.05 °	2.66±0.04 °	2.73 ± 0.04^{b}	2.96 ± 0.02^{a}	2 nd instar	f larvae and pup
	20.56±0.	19.52±0.	$18.22 \pm 0.$	17.88±0.	17.31±0.	3 rd inst	eight (mg ± SE) (is are significantly	0.05	2.62±0.07 °	2.74±0.04 ^d	$2.84\pm0.04^{\circ}$	3.08 ± 0.05^{b}	3.1 ± 0.03^{b}	3.53 ^a ±0.04 ^a	3 rd instar	ae (days ± SE) of
	.03 ^a 75.2t	.13 ^b 73.5	.03° 66.4	.06 ^d 64.19	.07 ^e 58.3	lar 4 th	of Helicoverpa a	y different ($P < 0$.	0.04	$2.81 \pm 0.04^{\text{f}}$	$2.92 \pm 0.03^{\circ}$	2.97 ± 0.06^{d}	$3.46\pm0.03^{\circ}$	3.52 ± 0.03^{b}	3.91 ± 0.03^{a}	4 th instar	f <i>H. armigera</i> on
	5 ± 0.08^{b}	2±0.14 °	8 ± 0.18^{d}	9±0.51 °	4 ± 0.20^{f}	instar	<i>rmigera</i> on diff	05, least signifi	0.05	$3.17 \pm 0.05^{+1}$	3.43 ± 0.03	$3.50\pm0.05^{\rm d}$	3.73 ± 0.03	3.92 ± 0.02^{b}	4.50 ± 0.03^{a}	5 th instar	different chick
	303.26±0.41 ^b	298.64±0.34 °	295.05±0.24 ^d	293.29±0.56°	290.47±0.62 ^f	5 th instar	erent chickpea cult	cant difference).	0.04	$3.32\pm0.01^{\text{f}}$	² 3.74±0.03 ^e	3.92 ± 0.08^{d}	² 4.22±0.04 ^c	4.33±0.03 ^b	5.05±0.04 ^a	6 th instar	pea cultivars.
510 00.0 01 a	515.71 ± 0.03^{b}	513.66±0.08 °	507.04±0.03 ^d	505.29±0.02 °	503.85 ± 0.01 ^f	6 th instar	ivars		0.21	16.41 ± 0.05^{t}	17.59±0.04 °	$18.08 \pm 0.34^{\text{d}}$	19.66±0.11°	20.24±0.21 ^b	22.71±0.05 ^a	Total time	
רו 112 חג+1 כו	198.74 ± 0.23 ^{ab}	157.32 ± 3.21^{b}	180.83 ± 0.45 ab	175.85±0.74 ^{ab}	163.53±0.63 ^b	Pupa				$09.50\pm0.02^{\circ}$	10.29 ± 0.15^{d}	11.52 ± 0.03 °	13.03 ± 0.50^{b}	13.25 ± 0.05 ^b	14.01±0.01 ^b	Pupa	ŝ

The means followed by different letters in the same columns are significantly different (P < 0.05, least significant difference).

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differ significantly with respect to development time of 4th instar larvae (F_{5.29}=1119.64, P<0.01). Larvae reared on cultivar Pb-2008 had longest development time with 3.91 days (Table I). Minimum duration of 5th instar was recorded on CMC-211s with 3.17 days, followed by Dasht and CM-2000 with 3.43 and 3.50 days respectively (F_{5.29}=857.18, P< 0.01). Chickpea cultivars differed significantly in larval duration of 6^{th} instar of H. armigera larvae (F_{5.29}=1452.07, P<0.01). The mean pupal development time of H. armigera was significantly differ $(F_{5,29}=464.86, P< 0.01)$ on chickpea cultivars (Table I). The percentage survival (Table III) of larvae of H. armigera was minimum 73% on cultivar Pb-2008 and was maximum 93% on cultivar CMC-211s and differ significantly in all the test cultivars (F_{5,29}=30.62, P< 0.01). The percent adult emergence of H. armigera was significantly different in all the test cultivar (F_{5.29}=79.41, P< 0.01).

Table III.-Percentage survival (Mean ± SE) of larvae and
pupae of *Helicoverpa armigera* on different
chickpea cultivars

Cultivar	Larva	Pupa		
Pb-2008	73.00 ±0.02 ^d	78.06±3.17 ^d		
Bittal-98	78.00±0.71 °	81.00±0.65 °		
Parbat	79.00±1.07 °	80.96±0.95 °		
CM-2000	87.00±0.08 ^b	87.37±2.05 ^b		
Dasht	88.00±0.52 ^b	88.63±0.35 ^b		
CMC-211s	93.00±0.64 a	94.52± 0.15 ^a		
LSD	1.01	1.03		

The means followed by different letters in the same columns are significantly different (P < 0.05, least significant difference).

Body weight

The results of weight of *H. armigera* on different chickpea cultivars are given in Table II. The mean larval weight of 1st instar (F_{5.29}=3978.06, P<0.01), 2nd instar (F_{5,29}=2823.83, P<0.01), 3rd instar (F_{5,29}=1903.71, P<0.01), 4th instar (F_{5.29}=5560.56, P<0.01), 5th instar (F_{5,29}=660.62, P< 0.01) and 6th instar (F_{5,29}=589.08, P<0.01) of *H. armigera* were significantly different in all the test cultivar. Maximum weight of 1st instar larvae was 0.33 mg on CMC-211s (Table II). The 2nd instar larvae reared on CMC-211s had maximum weight (7.32 mg), followed by Dasht and CM-2000. Larvae of 3rd instar reared on CMC-211s were heavier (20.64 mg) and lighter on Pb-2008 (17.31 mg) as compared to other cultivar. The 4th instar larvae had maximum weight (78.09 mg) and when reared on CMC-211s minimum weight (58.34 mg) cultivar Pb-2008. The 5th instar larvae maximum weight (303.02 mg on CMC-211s, followed by Dasht and CM-2000. The 6th instar larvae was 518.20 mg on CMC-211s and was 503.85 mg on Pb-2008.

The mean pupal weight of *H. armigera* did not differ significantly in all the test cultivars ($F_{5,29}$ =2.20, P> 0.05). Maximum pupal weight was 212.08 mg on CMC-211s which was statistically similar to pupal weight on Dasht (198.74 mg), Parbat (180.83 mg) and Bittle-98 (175.85 mg).

Fecundity of H. armigera on different chickpea cultivars

There was significant difference in fecundity of *H. armigera* on different chickpea cultivars ($F_{5,29}$ =589.26, P< 0.01). The total number of eggs laid were highest by the females of *H. armigera* larvae reared on CMC-211s (522.54 eggs) while there was significant reduction in number of eggs laid by the females of the larvae developed on PB-2008 (299.68 eggs) (Table IV).





Effect of different chickpea cultivars on feeding behaviour of H. armigera

The feeding preference of *H. armigera* were significantly different among all the treatments ($F_{5,29}$ =8.66, P<0.01). The maximum larvae were recorded on CMC-211s (3.8) while minimum larvae were recorded on Pb-2008 (1.2) (Table IV).

DISCUSSION

Plant species differ greatly in suitability as hosts for specific insects when measured in terms of survival, development and reproductive rates. Prolonged

Cultivar	Pre oviposition period	Oviposition period	Post oviposition period	Fecundity	
Pb-2008	3.12±0.08 ^a	6.14±0.04 ^a	2.06±0.04 ^a	299.68±1.73 ^f	
Bittal-98	2.93±0.03 ^b	5.81±0.02 ^b	1.92±0.01 ^b	323.66±1.69 °	
Parbat	2.88±0.05 ^b	5.33±0.10°	1.73±0.00 °	353.36±1.15 ^d	
CM-2000	2.63±0.02 °	4.93±0.03 ^d	$1.42\pm0.02^{\text{ d}}$	462.84±2.77 °	
Dasht	2.42 ± 0.02^{d}	4.71±0.06 ^e	1.31±0.01 ^e	492.28±2.27 ^b	
CMC-211s	2.18±0.05 °	4.25±0.11 ^f	$1.22\pm0.03^{\text{ f}}$	522.54±2.31 ^a	
LSD	0.05	0.17	0.03	2.57	

 Table IV. The mean pre, post-oviposition , oviposition periods (Days ± SE) and fecundity (Number of eggs/ female) of *Helicoverpa armigera* emerging from larvae reared on different chickpea cultivars.

The means followed by different letters in the same columns are significantly different (P < 0.05, least significant difference).

development time on particular species of host means longer life cycle and slower population growth (Singh and Parihar, 1988). Larval stages of *H. armigera* when prolonged may augment the efficacy of its management tactics by using insecticides and natural enemies (Du *et al.*, 2004). In the present study *H. armigera* completed its larval development in six instars which have been reported previously by Goyal and Rathore (1988) and Borah and Duttta (2002).

Development time of 1^{st} instar larvae was longest on Pb-2008 (2.75 days) and shortest on CM-211s (2.17days). These results are in contradiction with Fathipour and Naseri (2011), who reported the development time of 1^{st} instar of *H. armigera* larvae from 17.30 to 26.20 days on soybean cultivars. This variation in results might be associated with food quality of the host species.

The larval periods ranged from 19.66 days on Parbat to 22.71 days on Pb-2008. Whereas Naseri et al. (2009) reported total development time of H. armigera larvae from 17.30 to 26.20 days on different soybean cultivars. A possible explanation of variations in results may be due to difference in the nutritional value of the host species tested. Body weight is an important indicator of fitness of an insect, which can be measured easily (Liu et al., 2004). Host plants have great influence on the body weight of an insect species. This is evident from Sharma et al. (1999) that larvae reared on resistant cultivars had considerably lower weight than reared on susceptible cultivars. This is in agreement with our finding. Pupal weight of H. armigera was maximum on CMC-211s (212.08 milligram) while minimum on cultivar CM-2000 (157.32milligram). Whereas, Srivastava and Srivastava (1990) reported pupal weight from 231 to 310 milligram on different chickpea genotypes. The difference in the results is might be due to the physiological differences present in host plants.

The survival of H. armigera larvae differed

significantly on different cultivars of chickpea. The survival percentage of all larval instars of *H. armigera* ranged from 73% on comparatively resistant cultivars to 93% on comparatively susceptible cultivars of chickpea. These results are in partial agreement to those of Srivastava and Srivastava (1990) who reported 77-90 % survival of larvae of *H. armigera* on different chickpea cultivar.

Many factors affect host suitability, including nutrient content and secondary substances of the host (Liu et al., 2004). The exact cause of the differences found among host plants in larval growth rates, mortality, adult fecundity and survival remains unknown (Liu et al., 2004). However, it is reported that the presence of secondary plant substances or poor food quality in wild varieties while malic acid and oxalic acid are the principal components of resistance to H. armigera in the cultivated chickpea, which result in oviposition nonpreference and antifeedant effects on H. armigera (Yoshida et al., 1995). It is also reported that antibiosis seems to be the major component of resistance in the wild relatives of chickpea (Sharma et al., 2005). There is evidence of substances (flavonoids) present in some chickpea cultivars responsible for antifeedant activity against the H. armigera larvae (Simmonds and Stevenson, 2001). In the present study, preference of larval feeding of H. armigera revealed significant difference. The maximum larvae were recorded on CMC-211s (3.8) while minimum larvae were recorded on Pb-2008 (1.2). The differences in larval attraction towards cultivars might be attributed to differences in primary or secondary compounds or physiological characteristics present in different cultivar pods (Sharma et al., 2005).

There was significant difference in fecundity of *H. armigera* on different chickpea cultivar. The total number of eggs laid were highest by the females developed from larvae reared on CMC-211s (522.54 eggs) while there was significant reduction in number of eggs laid by the

females developed on PB-2008 (299.68 eggs). These results are in accordance with the Narayanamma *et al.* (2007), who reported lower fecundity on resistant chickpea parent hybrids as compared to susceptible parent hybrids. In conclusion, the more body weight, faster development, highest survival percentage and highest fecundity of *H. armigera* suggested that cultivar CMC-211s was susceptible as compared with the other cultivars. Cultivar Pb-2008 was resistant as compare to other cultivars. Dasht, CM-2000, Parbat and Bittle-98 showed intermediate trend.

The development of new chickpea resistant varieties is being recommended to enhance crop protection and ultimately crop production. The development of cultivars with resistance would provide an effective complementary approach in integrated pest management to minimize the extent of losses due to this pest (Sharma *et al.*, 2005).

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