Results and Discussion

IV. RESULTS AND DISCUSSION

4.1-Initial and bioresidual toxicity of different pesticides on the cotton leafworm larvae:

a- Laboratory reference strain:

Results in Table (1) demonstrate the percentages of initial and residual larval mortalities occurring after feeding for 2 days on cotton leaves treated with the tested insecticides at different time intervals, following application under field conditions, among the four groups of 4th instar larvae of S. littoralis laboratory strain.

Data representing initial toxicity for the 1st group of larvae which were exposed for 24 h. feeding period on treated cotton leaves collected at zero day and the survivors larvae continued their feeding for another 24 h. on treated cotton leaves collected at 1 day following application are presented in Table (1). The percentage initial mortalities after 48h. feeding on treated cotton leaves and 72h. feeding on untreated cotton leaves reached 100 % for Chromafenozide (400 and 200 ml./fed.) and Ethoxazole (25 and 12.5 ml./fed.) while reached 80 and 70% for Pyriproxyfen (300 and 150 ml./fed.) and 74 and 62% for Azadirachtin (100 and 50 ml./fed.).

The bioresidual toxicity of the tested insecticides on the second group of larvae fed for 48h. on treated cotton leaves collected at 7 and 8 days following application and followed by feeding 3 days on untreated leaves are also presented in Table (1). The larval mortalities reached 82, 76, 74, 73, 73, 68, 63 and 62 % for Chromafenozide 400 ml./fed.), Pyriproxyfen 300 ml/fed., Ethoxazole 25 ml/fed., Chromafenozide 200 ml/fed., Ethoxazole 12.5 ml/fed., Azadirachtin 100 m./fed., Pyriproxyfen 150 ml/fed. And Azadirachtin 50 ml/fed., respectively.

Results of the 3rd group of larvae fed for 48h. on treated cotton leaves collected from the field at 14 and 15 days after application and

(Laboratory reference strain) fed on treated cotton leaves for 48h. followed by 3 days feeding on Table (1). Initial and bioresidual toxicity of different insecticides against the 4th instar larvae of S. littoralis untreated cotton leaves.

		% Larval mortal	lity for groups of la	% Larval mortality for groups of larvae fed on treated leaves collected at the indicated days	eaves collected at th	e indicated days	
Insecticides	Rate ml/	Initial toxicity		Bioresidual toxicity	al toxicity		Overall mean*
	feddan	157	2^{nd}	3rd	414	Mean	±S.d.
		(0 - 1 day)	(7-8 day)	(14-15 day)	(21-22 day)	меап	
Pyriproxyfen	300	80	92	62	40	59.33	64.5 ± 18.06 ab
(10% EC)	150	70	63	54	39	52.00	56.5 ± 13.37 ab
Chromafenozide	400	100	82	92	70	76.00	82.0 ± 12.96 a
(5% EC)	200	100	73	72	99	70.33	77.75 ± 15.15 ab
Azadirachtin	100	74	89	53	40	53.66	58.75 ± 15.3 ab
(4.5% EC)	50	62	62	51	36	49.66	52.75 ± 12.31 b
Ethoxazole	25	100	74	56	50	00.09	70.00 ± 22.44 ab
(10% SC)	12.5	100	73	48	48	56.33	67.25 ± 24.81 ab
Control	1	2	3	2	4	3.0	2.75 ± c

* Means followed by the same letter are not significantly different according to Duncan's multiple range test. L.S.D. at 0.05 = 15.9

then fed for 3 days on untreated cotton leaves are presented in Table (1). The residual percentage mortalities reached 76, 72, 62, 56, 54, 53, 51 and 48% for Chromafenozide (400 and 200 ml/fed.,), Pyriproxyfen (300 ml/fed.), Ethoxazole (25 ml/fed.), Pyriproxyfen (150 ml/fed.,), Azadirachtin (100 and 50 ml/fed.) and Ethoxazole (12.5 ml/fed.), respectively.

Following the bioresidual toxicity of the tested insecticides over 21 days, the data obtained for the 4th group of larvae (Table 1) demonstrate that the efficacy reached 70, 66, 50, 48, 40, 40, 39 and 36 % for Chromafenozide (400 and 200 ml/fed.), Ethoxazole (25 and 12.5 ml/fed.), Pyriproxyfen (300 ml/fed.), Azadirachtin (100 ml/fed.), Pyriproxyfen (150 ml/fed.) and Azadirachtin (50 ml/fed.), respectively.

The mean residual toxicity of the 2nd, 3rd and 4th groups of larvae are also presented in Table (1). It demonstrated 76.0, 70.3, 60.0, 59.3, 56.3, 53.6, 52.0 and 49.6 % larval mortality for Chromafenozide (400 and 200 ml/fed.), Ethoxazole (25 ml/fed.), Pyriproxyfen (300 ml/fed), Ethoxazole (12.5 m./fed.), Azadirachtin (100 ml/fed.), Pryiproxyfen (150 ml/fed.), Azadirachtin (50 ml/fed.), respectively.

The overall mean larval mortality percentages were 82.0, 77.75, 70.0, 67.25, 64.5, 56.5, 58.75 and 52.75 % for Chromafenozide (400 and 200 ml/fed.), Ethoxazole (25 and 12.5 ml/fed.), Pyriproxyfen (300 and 150 ml/fed.) and Azadirachtin (100 and 50 ml/fed.), respectively.

The statistical analysis of the results showed that Chroma-fenozide was the most effective compound against the larval population of *S. littoralis* at both rates (400 and 200 ml/fed.), followed by Ethoxazole at the rates of (25 and 12.5 ml/fed.) and Pyriproxyfen at the rates of (300 and 150 ml/fed.) The least toxic compound was Azadirachtin at the rates of (100 and 50 ml/fed.) (Table 1).

b- Early-season field strain (before spraying program):

Data in Table (2) show the initial and residual percent mortalities of the 4th instar larvae of *S. littoralis* occurring within 2 days feeding on cotton leaves treated with different insecticides and followed by 3 days feeding on untreated cotton leaves. For the 1st group of larvae, the initial mortality percentages after 48 h. feeding period on treated cotton leaves collected at zero and 1 day after application, followed by feeding on untreated cotton leaves for 3 days were 100, 77, 67, 66, 63, 61, 57 and 55 % for Chromafenozide 400 ml/fed., Chromafenozide (200 ml/fed.), Ethoxazole (25 ml/fed.), Pyriproxyfen (300 ml/fed.), Azadirachtin (100 ml/fed.), Ethoxazole (12.5 ml/fed.), Pyriproxyfen (150 ml/fed.) and Azadirachtin (50 ml/fed.), respectively.

As for the 2nd group of larvae (Table 2), the mortality percentages reached 78, 72, 62, 61, 53, 53, 45 and 32 % for Chromafenozide (400 and 200 ml/fed.), Pyriproxyfen (300 ml/fed.), Ethoxazole (25 ml/fed.), Pyriproxyfen (150 ml/fed.), Ethoxazole (12.5 ml/fed.), and Azadirachtin (100 and 50 ml/fed.), respectively.

Results of the 3rd group of larvae fed for 48h. on treated cotton leaves collected from the field at 14 and 15 days after applications are also presented in Table (2). The data revealed that the accumulative latent toxicity (after 5 days) reached 48, 47, 43, 42, 41, 39, 34 and 20% for Pyriproxyfen (300 ml/fed.), Chromafenozide (400 ml/fed.), Pyriproxyfen (150 ml/fed.), Ethoxazole (25 ml/fed.), Chromafenozide (200 ml/fed.), Ethoxazole (12.5 ml/fed.), and Azadirachtin (100 and 50 ml/fed.), respectively.

For the 4th group of larvae (Table 2), the resulted percentage mortalities were 45, 36, 35, 34, 33, 30, 21, and 15% for Chromafenozide (400 and 200 ml/fed.), Ethoxazole (25 ml/fed.), Azadirachtin (100 ml/fed.), Ethoxazole (12.5 ml/fed.), Pyriproxyfen (300 and 150 ml/fed.) and Azadirachtin (50 ml/fed.), respectively.

Initial and bioresidual toxicity of different insecticides against the 4th instar larvae of S. littoralis early-season field strain (before spraying program) fed on treated cotton leaves for 48h. followed by 3 days feeding on untreated cotton leaves. Table (2).

		% Larval mortali	ity for groups of lar	% Larval mortality for groups of larvae fed on treated leaves collected at the indicated days	aves collected at the	indicated days	
Insecticides	Rate	Initial		Bioresidual toxicity	toxicity		Overall mean*
	feddan	18	2nd	3rd	4"	Mean	+ S.d.
		(0 - 1 day)	(7-8 day)	(14-15 day)	(21-22 day)		
	000	99	29	48	30	46.66	51.5 ± 16.2 ab
Pyriproxyten	300	25	53	43	21	39.00	43.5 ± 16.11 ab
(10% EC)	001	6	36	47	45	56.66	$67.5 \pm 26.41 a$
Chromafenozide	400	IOO	0 6	=======================================	3,6	49.66	56.5 ± 20.98 ab
(5% EC)	200	7.7	7/	T.	2		44 0 ± 12 60 ab
Azadirachtin	100	63	45	34	34	37.66	44.0 ± 15.08 au
Azaun azınını	05	35	32	20	15	22.33	30.5 ± 7.82 b
(4.5% EC)	200	29	19	42	35	46.00	51.2 ± 15.19 ab
Ethoxazole	57	6	53	39	33	41.66	46.5 ± 12.79 ab
(10% SC)	7.71	,	6	2	4	3.0	2.75 ± c
Control		7	,				

* Means followed by the same letter are not significantly different according to Duncan's multiple range test.

L.S.D. at 0.05 = 26.11

The mean of bioresidual activity percentages after the insecticidal applications of the 2nd, 3rd and 4th groups of S. littoralis larvae were 56.66, 49.66, 46.66, 46.0, 41.66, 39.0, 37.66 and 22.33% for Chromafenozide (400 and 200 ml/fed.), Pyriproxyfen (300 ml/fed.), Ethoxazole (25 ml/fed.), Ethoxazole (12.5 ml/fed., Pyriproxyfen (150 ml/fed.), and Azadirachtin (100 and 50 ml/fed.), respectively.

The overall mean larval mortality percentages of early-season field strain (before spraying program) were 67.5, 56.5, 51.5, 51.2, 46.5, 44.0, 43.5 and 30.5% for Chromafenozide (400 and 200 ml/fed.), Pyriproxyfen (300 ml/fed.), Ethoxazole (25 and 12.5 ml/fed.), Azadirachtin (100 ml/fed.), Pyriproxyfen (150 ml/fed.), and Azadirachtin (50 ml/fed.), respectively.

From the statistical analysis of the results it can be said again that Chromafenozide was the most toxic compound and was superior to the other tested insecticides at both rates 400 and 200 cm³/fed. It was followed by Ethoxazole and Pyriproxyfen especially at the high rates. The least toxic compound was Azadirachtin.

c- Late-season field strain (after spraying program):

Data demonstrated in Table (3) indicate the mortality percentages of the 4th instar larvae of late-season field strain (after spraying program) when fed for 48h. on cotton leaves treated with different insecticides at 0 and 1 day following application. The insecticides can be arranged according to their efficacy in the following descending order, Chromafenozide 400 ml/fed. (81 %), Ethoxazole 25 ml/fed. (80%), Chromafenozide 200 ml/fed. (79 %), Ethoxazole 12.5 ml/fed. (61%), Azadirachtin 100 and 50 ml/fed. (59 and 55 %) and Pyriproxyfen 300 and 150 ml/fed. (52 and 43 %), respectively. As for the 2nd group of larvae, the percent mortalities demonstrated were 65 % for Chromafenozide (400 ml/fed.) 59 % for Ethoxazole (25 ml/fed.), 57 % for Azadirachtin (100 ml/fed.), 50 % for Chromafenozide (200 ml/fed.), 46 % for Pyriproxyfen (300 ml/fed.), 45 % for Ethoxazole

season field strain (after spraying program) fed on treated cotton leaves for 48h. followed by 3 days Initial and bioresidual toxicity of different insecticides against the 4th instar larvae of S. littoralis latefeeding on untreated cotton leaves. Table (3).

		% Larval mortali	ity for groups of lar	vae fed on treated l	% Larval mortality for groups of larvae fed on treated leaves collected at the indicated days	indicated days	
Insecticides	Rate ml/	Initial toxicity		Bioresidual toxicity	l toxicity		Overall mean*
	feddan	181	2 nd	3 rd	4"	Mean	± S.d.
		(0 - 1 day)	(7-8 day)	(14-15 day)	(21-22 day)		
Pyrinroxyfen	300	52	46	33	28	35.66	39.57 ± 11.14 ab
(10% EC)	150	43	28	11	5	14.66	$21.75 \pm 17.19 \mathrm{b}$
Chromafenozide	400	81	65	58	42	55.00	61.5 ± 16.17 a
(5% FC)	200	79	50	45	40	45.00	53.5 ± 17.48 a
Azadirachtin	100	59	57	43	26	42.00	46.25 ± 15.26 ab
(4.5% EC)	50	55	40	32	23	31.66	37.5 ± 13.57 ab
Ethoxazole	25	80	59	45	32	45.33	54.0 ± 20.54 a
(10% SC)	12.5	61	45	40	30	38.33	44.0 ± 12.93 ab
Control		2	3	2	4	3.0	2.75± c

* Means followed by the same letter are not significantly different according to Duncan's multiple range test.

L.S.D. at 0.05 = 11.5

(12.5 ml/fed.), 40 % for Azadirachtin (50 ml/fed.) and 28 % for Pyriproxyfen (150 ml/fed.), respectively. Data in Table (3) also show the mortality percentages of the 3rd group of larvae which were fed for 48 h. on treated cotton leaves collected at 14th and 15th days following insecticidal applications. The mortality percentages were 58, 45, 45, 43, 40, 33, 32 and 11% for Chromafenozide (400 and 200 ml/fed.), Ethoxazole (25 ml/fed.), Azadirachtin (100 ml/fed.), Ethoxazole (12.5 ml/fed.), Pyriproxyfen (300 ml/fed.), Azadirachtin (50 ml/fed.) and Pyriproxyfen (150 ml/fed.), respectively.

The mortality percentages of the 4th group of S. littoralis lateseason field strain (after spraying program) were 42 and 40 % for Chromafenozide (400 and 200 ml/fed.), 32 and 30 % for Ethoxazole (25 and 12.5 ml/fed.), 28% for Pyriproxyfen (300 ml/fed.), 26 and 23% for Azadirachtin (100 and 50 ml/fed.), and 5 % for Pyriproxyfen (150 ml/fed.), respectively.

The mean residual larval mortality percentages were 55.0, 45.0, 45.3, 42.0, 38.3, 35.6, 31.6 and 14.6 % for Chromafenozide (400 and 200 ml/fed.), Ethoxazole (25 ml/fed.), Azadirachtin (100 ml/fed.), Ethoxazole (12.5 ml/fed.), Pyriproxyfen (300 ml/fed.), Azadirachtin (50 ml/fed.), and Pyriproxyfen (150 ml/fed.), respectively.

The statistical analysis of the overall mean larval mortality percentages revealed that Chromafenozide at both rates of 400 and 200 ml/fed. was significantly superior compared with the other tested insecticides. The least toxic compound was Pyriproxyfen and both Ethoxazole and Azadirachtin were moderately toxic in their effects on the larvae of *S. littoralis* (Table 3).

In general, it can be concluded from the prementioned results that feeding period on untreated leaves play an important role in the performance of the toxicity later on, particularly for compounds with new mode of action such as insect growth regulators. However, data of previous work carried out early by **Rizk and Radwan** (1975) indicated that both new chitin biosynthesis inhibitors, PH-6040 and PH-6038

induced justified performance when the tested instar larvae of the cotton leafworm *S. littoralis* were fed continuously for 0-5 days after application. Likewise **Watson** (1981) found that the highest efficacy was achieved when mortality counts were assessed after 48h. exposure or/and feeding on treated leaves.

The lack of food intake by *Schistocerca* nymphs fed on *Melia volkensii*, lead to very poor relative growth and high mortality (Mwangi, 1982). Azadirachtin disrupted growth of young instars of *S. littoralis* than old ones and also affected ecdysis (Koul, 1985).

From the results of the present work, the feeding period on treated leaves was significantly effective factor in larval toxicity particularly insect growth regulators having low initial acute toxicity. These results confirm those found by Ishaaya and Klein (1990) who found that *S. littoralis* larvae collected from cotton fields in Israel that have been heavily sprayed with conventional insecticides showed strong resistance to organophosphates and pyrethroids and mild tolerance to benzoylphenyl ureas.

Recently, Smagghe et al. (1995) indicated that variations in toxicity of tebufenozide to laboratory and field strains of S. littoralis may be attributed to differences in compound stability and excretion. Further studies carried out by Smagghe et al. (2000) revealed that methoxyfenozide was the most toxic compound against S. littoralis larvae and RH-5849 was the least toxic.

4.2- Latent effect of different pesticides on certain developmental aspects in the cotton leafworm biology:

1- Effect on accumulated larval mortality:

Under laboratory conditions, surviving larvae after feeding on and exposure to residues of certain insecticides on cotton leaves, collected after various periods of spraying, were allowed to complete their larval stage duration. Results of overall accumulated larval mortality are given in Tables (4-6). The tested time intervals at which treated leaves were collected are as follows:

- a- Zero and 1st day time interval for the 1st group of larvae.
- b-7th and 8th day time interval for the 2nd group of larvae.
- c- 14th and 15the days time interval for the 3rd group of larvae.
- d- 21st and 22nd day time interval for the 4th group of larvae.

a- Laboratory reference strain:

Data in Table (4) elucidate the overall mortality of the larval stage of S. littoralis till pupation. The results of the 1st group of larvae which were fed on treated cotton leaves collected at 0-1 days after spraying revealed that both Chromafenozide and Ethoxazole caused 100% mortality, Pyriproxyfen caused 81-86% and Azadirachtin exhibited 67-80 % mortality. Regarding the 2nd group of larvae, the resulted mortalities were 76-87 %, 76-81 %, 77-79 % and 73-77 % for Chromafenozide, Pyriproxyfen, Ethoxazole and Azadirachtin, respectively. In the 3rd group of larvae which were fed on treated cotton leaves collected at 14-15 days after spraying, the accumulated larval mortalities reached 78-81 %, 60-69 %, 54-62 % and 57-61 % for Chromafenozide, Pyriproxyfen, Ethoxazole and Azadirachtin, respectively. In the 4th period of exposure the larvae fed on cotton leaves collected after 21-22 days from spraying showed 74-77 %, 55-55 %, 41-50 % and 37-48 % mortalities for Chromafenozide, Ethoxazole, Pyriproxyfen and Azadirachtin, respectively (Table 4). The

Accumalated larval mortality till pupation for groups of 4th instar larvae of S. littoralis (Laboratory strain) fed for 48h. on treated leaves collected at different time intervals after applying certain insecticides under field conditions. Table (4).

	Data	Overall % lar	val mortality till	Overall % larval mortality till pupation for groups of larvae fed	ups of larvae fed	Mean*
	ml/	on tre	ated leaves colle	on treated leaves collected at the indicated days	ted days	Larval Mortality
Insactivides	feddan	151	2 nd	3rd	4"	± S.d.
Inscrince		(0 - 1 dav)	(7-8 day)	(14-15 day)	(21-22 day)	
	000	98	81	69	50	71.5 ± 16.01 ab
Pyriproxyfen	2000	20 20	92	09	41	$64.5 \pm 18.04 \text{ ab}$
(10% EC)	061	1001	87	81	77	86.25 ± 10.04 a
Chromafenozide	400	100	76	78	74	82.0 ± 12.11 ab
(5% EC)	700	100	2 !		48	66 5 + 14 89 ab
Azadirachtin	100	80	77	10	0+	10111
Acada acada	05	29	73	57	37	58.5 ± 15.78 b
(4.3% EU)	20	100	79	62	55	78.5 ± 23.30 ab
Ethoxazole	50.	100	77	54	55	71.5 ± 21.76 ab
(10% SC)	12.3	100		٧.	7	$5.25 \pm 1.26 c$
Control	•	4	٥			44

* Means followed by the same letter are not significantly different according to Duncan's multiple range test.

L.S.D. at 0.05 = 23.29

mean percent accumulated larval mortalities were 82.0-86.2 %, 71.5-78.5 %, 64.5-71.5 % and 58.5-66.5 % for Chromafenozide, Ethoxazole, Chromafenozide at 400 ml/fed. caused a significant increase in larval and Azadirachtin, mortality (86.2 %) when compared with all other treatments while Azadirachtin at 50 ml/fed. demonstrated the least significant effect on larval mortality (58.5 %).

b- Early-season field strain (before spraying program):

In the 1st group of larvae (Table 5) it was obvious that larvae suffered high percentage of mortality after feeding on leaves treated with insecticides reached 83-100 % for Chromafenozide, 64-73 % for Ethoxazole, 63-71 % for Pyriproxyfen and 63-68 % for Azadirachtin.

In the 2nd period of exposure (7-8 days after spraying), the accumulated larval mortalities reached 78-83 % for Chromafenozide, 60-67 % for Pyriproxyfen, 58-67 % for Ethoxazole and 40-45 % for Azadirachtin.

As for the 3rd group of larvae, the obtained results of mortalities showed 54-68 %, 48-58 %, 46-50 % and 32-38 % for Pyriproxyfen, Chromafenozide, Ethoxazole and Azadirachtin, respectively.

results of the 4th group of larvae indicated that Chromafenozide, Ethoxazole, Pyriproxyfen and Azadirachtin caused 45-51 %, 41-42 %, 27-39 % and 18-38 % mortalities, respectively.

The statistical analysis of the mean accumulated larval mortality percentages demonstrated that Chromafenozide (400 ml/fed.) was the most effective (73.0 %) treatment while Azadirachtin (50 ml/fed.) was the least effective (38.25 %) treatment.

Late season field strain (after spraying program):

The overall larval mortality percentages till pupation for groups of late-season field strain S. littoralis larvae fed on treated cotton

field strain (before spraying program) fed for 48h. on treated leaves collected at different time Accumalated larval mortality till pupation for groups of 4th instar larvae of S. littoralis early-season intervals after applying certain insecticids under field conditions. Table (5).

	Rate	Overall % lar	val mortality till	Overall % larval mortality till pupation for groups of larvae fed	ips of larvae fed	Mean*
	ml/	on tre	ated leaves collec	on treated leaves collected at the indicated days	ted days	Larval Mortality
Insecticides	feddan	1 st	2^{nd}	3^{rd}	4"4	% ± S.d.
		(0 - 1 day)	(7-8 day)	(14-15 day)	(21-22 day)	
Pyriproxyfen	300	71	67	54	39	57.75 ± 14.45 ab
(10% EC)	150	63	99	89	27	54.5 ± 18.62 ab
Chromafenozide	400	100	83	58	51	73.0 ± 22.64 a
(5% EC)	200	83	78	48	45	63.5 ± 19.77 ab
Azadirachtin	100	89	54	38	38	49.5 ± 14.46 ab
(4.5% EC)	50	63	40	32	18	38.25 ± 18.84 b
Ethoxazole	25	73	67	50	42	58.0 ± 14.45 ab
(10% SC)	12.5	64	58	46	41	52.25 ± 10.59 ab
Control		4	5	5	7	$5.25 \pm 1.26 c$

* Means followed by the same letter are not significantly different according to Duncan's multiple range test.

L.S.D. at 0.05 = 23.42

leaves collected at different time intervals after applying insecticides in the field are presented in Table (6). In the 1st groups of larvae, the mortality percentages reached 93, 91, 87-89 and 87-90 % for Chromafenozide, Azadirachtin, Ethoxazole and Pyriproxyfen, respectively. In the 2nd tested period, the accumulated larval mortalities were 89-91 % for Chromafenozide, 82-85% for Ethoxazole, 80-89% for Azadirachtin and 84-85% for Pyriproxyfen. In the 3rd tested interval, the accumulated larval mortality percentages demonstrated 81-84, 85-88, 81-84 and 82 % for Chromafenozide, Ethoxazole, Azadirachtin and Pyriproxyfen, respectively. The 4th group of larvae fed on treated cotton leaves collected at (21-22 days) after spraying insecticides in the field exhibited 79-82, 70-72, 67-76 and 76-78 % mortalities for Chromafenozide, Ethoxazole, Azadirachtin and Pyriproxyfen, respectively, However, the statistical analysis of the mean accumulated larval mortality percentages revealed that while there was significant increase in larval mortality in treatments than control, there was no significant variations between different treatments itself and that Chromafenozide at (400 ml/fed.) caused the highest accumulated larval mortality (86.75 %) while Ethoxazole at 12.5 ml/fed. showed the least mean accumulated larval mortality (81.0

The remarkable high overall larval mortalities 58.5-86.25 % in laboratory strain (Table 4), 38.25-73.0 % in early-season field strain before spraying program (Table 5) and 79.75-86.75 % in late-season field strain after spraying program (Table 6) agreed with the previous findings of **Bayoumi** et al. (1998) found that percentage accumulative mortality varied according to the compound, concentration, larval instar and/or strain studied.

Accumalated larval mortality till pupation for groups of 4th instar larvae of S. littoralis (late-season field strain) fed for 48h, on treated leaves collected at different time intervals after applying certain insecticids under field conditions. Table (6).

	Rafe	Overall % lar	val mortality till	Overall % larval mortality till pupation for groups of larvae fed	ups of larvae fed	Mean*
	ml/	on tre	ated leaves colled	on treated leaves collected at the indicated days	ted days	Larval Mortality
Insecticides	feddan	18	2 nd	3rd	4"	% ± S.D.
		(0 - 1 day)	(7-8 day)	(14-15 day)	(21-22 day)	
Pvriproxvfen	300	87	84	82	78	82.75 ± 3.77 a
(10% EC)	150	06	85	82	76	83.25 ± 5.85 a
Chromafenozide	400	93	91	84	79	86.75 ± 6.45 a
(5% EC)	200	93	68	81	82	86.25 ± 5.74 a
Azadirachtin	100	91	68	84	76	$85.0 \pm 6.68 a$
(4.5% EC)	50	91	80	81	. 67	79.75 ± 9.84 a
Ethoxazole	25	68	85	88	72	83.5 ± 7.85 a
(10% SC)	12.5	87	82	85	70	81.0 ± 7.62 a
Control		4	5	5	7	$5.25 \pm 1.26 \text{ b}$

* Means followed by the same letter are not significantly different according to Duncan's multiple range test.

L.S.D. at 0.05 = 9.50

2- Effect on larval duration:

Tables (7-9) elucidate the duration of *S. littoralis* larval stage till pupation when 4th instar larvae were allowed to feed for 48 hr. on treated cotton leaves collected at different time intervals after applying the tested insecticides in the field and continue feeding on untreated leaves till death or pupation

a- Laboratory reference strain:

In the 1st group of larvae which were fed on treated cotton leaves collected at 0-1 days after spraying insecticides, Chromafenozide and Ethoxazole at both rates of application caused 100 % larval mortality while an increase in the larval duration reached 67.3-81.3 % and 60.0-62.0 % was recorded for Pyriproxyfen and Azadirachtin, respectively (Table 7).

Regarding the 2nd group of larvae fed on treated cotton leaves collected at 7-8 days after spraying insecticides, a pronounced increase in the larval period was recorded, 49.3-68.7 % for Pyriproxyfen, 42.0-48.0 % for Azadirachtin, 32.7-36.7 % for Chromafenozide and 18.0-34.0 % for Ethoxazole.

The same trend was also observed in the 3rd group of S. littoralis larvae. The increases in larval periods were 34.7-56.7 % for Pyriproxyfen, 28.7-34.7 % for Azadirachtin, 23.3-36.0 % for Chromafenozide and 23.3-24.0 % for Ethoxazole.

A relatively lower increase in the larval duration of the 4th group of larvae was observed, 20.6-21.3 % for Pyriproxyfen, 6.9-11.9 % for Chromafenozide, 0.6-1.3 % for Azadirachtin and 0.6 % for Ethoxazole.

b- Early-season field strain (before spraying program):

Results in Table (8) represent the percent changes in larval durations for groups of early-season larvae of *S. littoralis* fed on treated cotton leaves collected at different time intervals after applying

Larval duration for groups of 4th instar larvae of S. littoralis (laboratory strain) fed for 48h. on treated leaves collected at different time intervals after applying insecticides under field conditions. **Table** (7).

		Mea	ın larval du	ration (in da	ays) for gro	Mean larval duration (in days) for groups of larvae survived after feeding on	e survived	after feeding	c on
	Rate			treated leav	ves collected	treated leaves collected at the indicated days.	cated days.		
Insecticides	(m)/	$1^{st} (0 - 1^{st} day)$	1^{st} day)	$2^{nd} (7 - 8^{th} day)$	8''' day)	3^{rd} (14 - 15" day)	15" day)	4" (21 - 22th day)	22th day)
	feddan		%	12.	%		%		%
		Duration	Change	Duration	Change	Duration	Change	Duration	Change
Pvrinroxvfen	300	27.1	+81.3	25.3	+ 68.7	23.5	+ 56.7	19.4	+21.3
(10% EC)	150	25.1	+ 67.3	22.4	+ 49.3	20.2	+34.7	19.3	+ 20.6
Chromafenozide	400	0.0	1	20.5	+36.7	20.4	+36.0	17.9	+11.9
(5% EC)	200	0.0	ŧ	19.9	+ 32.7	18.5	+ 23.3	17.1	+ 6.9
Azadirachtin	100	24.3	+ 62.0	22.2	+ 48.0	20.2	+34.7	16.2	+1.3
(4.5% EC)	50	24.0	+ 60.0	21.3	+ 42.0	19.3	+ 28.7	16.1	+ 0.6
Ethoxazole	25	0.0	1	20.1	+34.0	18.6	+ 24.0	16.1	+ 0.6
(10% SC)	12.5	0.0		17.7	+ 18.0	18.5	+ 23.3	16.1	+ 0.6
Control		15.0		15.0		15.0		16.0	t

spraying program) fed for 48h. on treatd leaves collected at different time intervals after applying Larval duration for groups of 4th instar larvae of S. littoralis early-season field strain (before insecticides under field conditions. Table (8).

	Rate	Mea	an larval du	ration (in d treated lea	lays) for gro	Mean larval duration (in days) for groups of larvae survived after feeding on treated leaves collected at the indicated.	le survived	after feedin	g on
Insecticides	/Im	$1^{57}(0 -$	$1^{57} (0 - 1^{57} day)$	2nd (7 -	2"d (7 - 8th dav)	3rd (14 15th Jell	cated days.	th in	
	feddan		%		/0	- +1	(Kan cr	4" (21 - 22" day)	22" day)
		Duration	Change	Duration	Change	Duration	%	Duration	%
Pyriproxyfen	300	0 90			Sump		Change		Change
	200	70.07	+ /3.3	24.1	+ 60.7	20.2	+ 34 7	16.0	
(10% EC)	150	23.3	+55.3	21.5	+ 43 3	101	0 0	10.7	0.0+
Chromafenozide	400	00		t	0.01	17.1	+ 21.3	16.4	+2.7
(C. F. 704)		0.0		17.1	+ 14.0	16.2	+8.0	153	77
(5% EC)	200	21.8	+45.3	163	+87	15.4		200	t t
Azadirachtin	100	3 66	0	2.5	0.0	13.4	+2.7	11.5	- 28.1
	201	6.77	+ 20.0	20.4	+ 36.0	15.5	+3.3	13.2	17.5
(4.5% EC)	50	21.2	+41.3	18.4	+22.7	15.3	-	1.0.	C./.I
Ethoxazole	25	20.1	+ 34 0	101		0.01	1.2.0	13.2	- 17.5
(30% 501)	4 01		2	10./	+ 74.7	17.0	+13.3	16.4	+27
(36)/01)	17.3	18.8	+ 25.3	18.0	+ 20.0	15.2	+ 12	16.0	i
Control	1	15.0	1	15.0		2001	Ci	7.01	+1.3
				0.61		15.0		16.0	1

insecticides in the field. The treatment of Chromafenozide (400 ml/fed.) demonstrated 100% larval mortality. Larval durations were increased (+25.3-73.3 %) in treatments of all other tested insecticides when the 1st group of larvae were exposed and fed on treated leaves collected at 0-1 days after applying insecticides.

In the 2nd and 3rd groups of larvae an increase of 14.0-60. 7% and 1.3-34.7 %, respectively in larval duration was observed by the use of all insecticides.

Regarding the 4th group of larvae which were fed on treated cotton leaves collected at 21-22 days after spraying insecticides in the field, different reactions in the larval periods were achieved. Pyriproxyfen and Ethoxazole resulted in slight increase in the larval periods, 2.7-6.0 % and 1.3-2.7 % respectively, while Chromafenozide and Azadirachtin exhibited a decrease in the larval periods by 4.4-28.1 % and 17.5 % respectively.

c- Late-season field strain (after spraying program):

Data demonstrated in Table (9) illustrate the larval duration till pupation when larvae were fed on cotton leaves treated with insecticides for 48h. and allowed to continue feeding till pupation on untreated leaves.

In the 1st group of larvae which were fed at the first testing interval (0-1 days after spraying), the larval durations in Pyriproxyfen, Chromafenozide and Ethoxazole treatments where remarkably longer than control, the percent changes were +6.7-20.0 %, while in Azadirachtin treatment no changes were observed in larval period.

In the 2nd group of larvae, there were no changes in the larval duration except for Chromafenozide treatment which caused an increase of 6.7 %.

In the 3rd group of larvae which were fed on treated cotton leaves collected 14-15 days after spraying insecticides, there were no changes in the larval period in all treatments.

Larval duration for groups of 4th instar larvae of S. littoralis Late-season field strain (after spraying program) fed for 48h, on treated leaves collected at different time intervals after applying insecticides under field conditions. Table (9).

		Rate	Mea	l du	ration (in di	ays) for gro	ration (in days) for groups of larvae survived	ae survived	after feedir	no Si	
	Insecticides	ml/	$1^{st}(0 -$	$1^{st} (0 - 1^{st} day)$	2"d (7 - 8"h dav)	8th dav)	3 rd (14 - 15 th dow)	15th days	- 1		_
-		feddan		%		%		o day)	4 (21 - 22" day)	77, day)	_
			Duration	Change	Duration	Change	Duration	Change	Duration	%	
-	Pyriproxyfen	300	18	+ 20	15		15	Cuange		Change	
	(10% FC)	150	10		2	>	CI	0	16	0	-
1	(270,02)	001	10	+ 20	15	0	15	0	16	0	
	Chromafenozide	400	18	+ 20	16	+67	15		2		
	(5% EC)	200	16	174	1.		CT		CI	- 6.3	
~	zadiroohtin	100		10.7	10	+ 6.7	15	0	15	-63	
Ç	Azaun acııtın	100	15	0	15	0	15	0	15		
	(4.5% EC)	50	15	0	15	c	15		CI	- 0.3	
E	Ethoxazole	25	18	+ 20	15		11		CI	- 6.3	
	(10% SC)	12.5	16	177	2 .		CI	0	16	0	
	Control			10.7	CI	0	15	0	16	0	
	Control		15	1	15		15	9	16		
							-		_	-	

In the 4th group of larvae, a decrease in the larval duration (6.3 %) was observed in the treatments of both Chromafenozide and Azadirachtin at both rates of application.

In this respect, the increase in larval duration which was observed in the treatments of Pyriproxyfen and Chromafenozide agreed with the previous findings of Moustafa and Attal (1985) who found that the chitin inhibitor triflumuron at low concentration produced significant prolongation in *S. littoralis* larval period.

3- Effect on larval weight:

a- Laboratory reference strain:

Results illustrated in Table (10) show that all insecticides at both rates caused a decrease in the larval weight of *S. littoralis* for the four groups of larvae by 1.2-57.8 %, 28.7-85.9 %, 29.5-76.8 % and 27.2-76.8 % for Pyriproxyfen, Chromafenozide, Azadirachtin and Ethoxazole respectively.

b- Early-season field strain (before spraying program):

Data in Table (11) revealed slight increases in larval weights compared with that of the control by 4.3, 2.3 and 0.4 % for Pyriproxyfen at 150 ml/fed. for 0-1 days, 7-8 days and 14-15 days after treatment, whereas a pronounced decrease (-50.4 %) in the larval weight was recorded in the 4th group, fed on treated leaves at 21-22 days after treatment (Table 11). A considerable reductions in the larval were demonstrated in treatments of Chromafenozide, weights Azadirachtin and Ethoxazole, the reduction percentages were between 17.5-78.2 %, 28.7-42.6 % and 25.1-48.9 %, respectively during the four testing intervals. The results showed in general that all insecticides at both rates decreased the larval weight of S. littoralis fed on treated cotton leaves collected at different time intervals, the decrease was remarkable in the treatments of Chromafenozide, Azadirachtin and Ethoxazole, while it was of low magnitude in the Pyriproxyfen treatment.

Table (10). Mean larval weight (till 8 days) of S. littoralis (Laboratory reference strain) for groups of 4th instar larvae fed for 48h, on treated cotton leaves collected at different time intervals after applying insecticides under field conditions.

Insecticides ml/ 1st (0 - 1st day) 2nd (7 - 8th day) 3rd (14 - 11st day) 406		Rate	Mear	Mean weight of larvae (mg.) till 8 days for groups of larvae fed on treated leaves collected at the indicated days	arvae (mg.)	ng.) till 8 days for groups of lar collected at the indicated days	or groups o	f larvae fed	on treated	leaves
feddan Weight % Weight % Weight en 300 192.9 -13.2 128.7 -42.0 112.3 !% EC) 150 220.3 -9.4 219.2 -1.2 208.9 ozide 400 31.3 -85.9 82.4 -62.9 112.4 % EC) 200 41.7 -81.2 109.8 -50.5 160.7 in 100 51.5 -76.8 81.3 -62.9 150.1 % EC) 50 59.0 -73.5 104.6 -52.9 150.1 % EC) 50 59.0 -73.5 104.6 -52.9 150.1 % SC) 12.5 52.3 -76.8 110.5 -47.3 155.5 ol - 222.4 - -73.6 116.7 -47.3 155.5	Insecticides	/lm	$1^{st}(0 -$	1^{st} day)	2 nd (7 -	8th day)	3rd (14 -	15th day)	4th (21 -	4th (71 - 27th day)
Pyriproxyfen 300 192.9 -13.2 128.7 -42.0 112.3 Chromafenozide (10% EC) 150 220.3 -9.4 219.2 -1.2 208.9 Chromafenozide (5% EC) 400 31.3 -85.9 82.4 -62.9 112.4 Azadirachtin (4.5% EC) 200 41.7 -81.2 109.8 -50.5 160.7 Ethoxazole (10% SC) 50 59.0 -73.5 104.6 -52.9 150.1 Ethoxazole (10% SC) 12.5 52.3 -76.8 110.5 -50.2 111.1 Control - 222.4 - 76.5 116.7 -47.3 155.5		feddan	Weight	% Change	Weight	Change	Weight	%	Weight	%
% EC) 150 220.3 - 9.4 219.2 - 1.2 208.9 ozide 400 31.3 - 85.9 82.4 - 62.9 112.4 % EC) 200 41.7 - 81.2 109.8 - 50.5 160.7 in 100 51.5 - 76.8 81.3 - 63.4 91.4 % EC) 50 59.0 - 73.5 104.6 - 52.9 150.1 % SC) 12.5 52.3 - 76.8 110.5 - 50.2 111.1 % SC) 12.5 52.3 - 76.5 116.7 - 47.3 155.5 ol - 222.4 - 73.6 273.5		300	192.9	- 13.2	128.7	- 42 0	1123	Change 50 1	0 30	Change
ozide 400 31.3 -85.9 82.4 -62.9 112.4 %EC) 200 41.7 -81.2 109.8 -50.5 160.7 in 100 51.5 -76.8 81.3 -63.4 91.4 %EC) 50 59.0 -73.5 104.6 -52.9 150.1 %EC) 50 59.0 -76.8 110.5 -52.9 150.1 %SC) 12.5 52.3 -76.5 116.7 -47.3 155.5 ol - 222.4 - -221.0 -225.9	(10% EC)	150	220.3	- 9.4	219.2	-12	208.0	7.2	9.00	8./6-
%EC) 200 41.7 -81.2 109.8 -50.5 160.7 in 100 51.5 -76.8 81.3 -63.4 91.4 %EC) 50 59.0 -73.5 104.6 -52.9 150.1 25 51.6 -76.8 110.5 -50.2 111.1 %SC) 12.5 52.3 -76.5 116.7 -47.3 155.5 51 - 222.4 - 73.5	Chromafenozide	400	31.3	- 85.9	82.4	0 69 -	112.4	0.7	11117	4.6-
in 100 51.5 -76.8 81.3 -63.4 91.4 % EC) 50 59.0 -73.5 104.6 -52.9 150.1 25 51.6 -76.8 110.5 -50.2 111.1 % SC) 12.5 52.3 -76.5 116.7 -47.3 155.5 51 - 222.4 - 221.0	(5% EC)	200	41.7	- 81.2	100 0	50.5	112.4	2.00.2	114./	- 49.6
(ii) 100 51.5 - 76.8 81.3 - 63.4 91.4 % EC) 50 - 73.5 104.6 - 52.9 150.1 25 51.6 - 76.8 110.5 - 50.2 111.1 % SC) 12.5 52.3 - 76.5 116.7 - 47.3 155.5 31 - 222.4 - 76.5 176.9 - 77.5	(2000)	2	177.7	7.10	109.0	- 20.3	160.7	- 28.7	161.9	- 28.8
% EC) 50 59.0 -73.5 104.6 -52.9 150.1 25 51.6 -76.8 110.5 -50.2 111.1 % SC) 12.5 52.3 -76.5 116.7 -47.3 155.5 51 - 222.4 - 221.0 -225.5	Azadirachtin	100	51.5	- 76.8	81.3	- 63.4	91.4	- 59.5	136.8	- 39 0
% SC) 12.5 52.3 - 76.8 110.5 - 50.2 111.1 1.0	(4.5% EC)	50	59.0	- 73.5	104.6	- 52.9	150.1	- 33.4	160 5	20.5
6 SC) 12.5 52.3 - 76.5 116.7 - 47.3 155.5 - 222.4 - 221.0 - 225.5	Ethoxazole	25	51.6	- 76.8	110.5	- 50.2	111	- 50 7	140.7	20.7
- 222.4 - 221.9	(10% SC)	12.5	52.3	- 76.5	116.7	- 47.3	155.5	-310	1657	230.7
1111.7	Control	1	222.4	ĭ	221.9		225.5	0.10	2775	7.17-

Mean larval weight (till 8 days) of S. littoralis early-season field strain (before spraying program) for groups of 4^{th} instar larvae fed for 48h, on treated cotton leaves collected at different time intervals after applying insecticides under field conditions. Table (11).

		Mean	weight of la	rvae (mg.)	till 8 days fo	Mean weight of larvae (mg.) till 8 days for groups of larvae fed on treated leaves	larvae fed	on treated I	eaves
	Rate			ollo	scted at the	collected at the mulcarcu units:	Eth down	4th (21 - 22th day)	22th day)
	\(\frac{1}{2}\)	1st (0 - 1st day)	(adav)	$2^{na} (7 - 8^{na} day)$	8" day)	3 (14-	(S) uay)	-	%
Insecticides	(mil)/ feddan		%	Woight	%	Weight	%	Weight	Change
		Weight	Change	weight	Change	0	Change		62.0
		, 104	11	204 4	- 7.8	201.9	- 10.4	109.7	- 34.0
Pvriproxyfen	300	205.1	/-/-			2363	+ 0.4	112.8	- 50.4
7100 EC	150	231.9	+ 4.3	227.2	+ 7.3	2.027		1030	28.4
(10% EC)	201	7.07	70.7	121.4	- 45.3	158.8	- 29.6	103.0	1.07
Chromafenozide	400	48.0	7.0/ -		7 00	1653	- 26.6	187.6	- 17.5
	000	105.5	- 52.6	136.8	- 38.4	100.0			27.0
(5% EC)	707			1203	- 41 3	135.3	- 40.0	141.2	-3/.7
Azadirachtin	100	127.6	- 42.6	130.7	C.T.L	1000	202	162.2	- 28.7
	9	136.3	- 38.7	142.1	- 36.0	7.761	2,70	, ,,,	36.1
(4.5% EC)	25		10.01	120.5	- 45.7	141.3	- 37.3	143.4	1.00-
Ethoxazole	25	113.6	- 40.7	2000	10.6	158.0	- 29.9	170.4	- 25.1
(J8 78017	12.5	120.0	- 46.0	127.6	- 47.3	13000		2 100	
(10%95)				2219	•	225.5	•	C-177	
Control		222.4		*******				D	

c- Late-season field strain (after spraying program) :

The results concerning the effect of the tested insecticides on the larval weight of S. littoralis, late-season field strain (after spraying program) are presented in Table (12). A decrease in the larval weight was obtained in all treatments and rates of applications. The highest decrease was demonstrated by Ethoxazole at 25 ml/fed. (68.6 %) while the least one (29.4 %) was obtained by Ethoxazole at 12.5 ml/fed. for the 1st group of larvae. The highest decrease in larval weight during the 2nd group of larvae was 46.6% by the treatment of Azadirachtin at 100 ml/fed., while the least one (12.2 %) was found by the treatment of Ethoxazole at 12.5 ml/fed. The highest decrease in larval weight was recorded for Pyriproxyfen (30.8 %) at 300 ml/fed. during the 3rd testing interval while it was 27.7 % for Azadirachtin at 100 ml/fed. during the 4th testing interval, however, the lowest decrease was obtained by the treatment of Ethoxazole (10.1 %) at 12.5 ml/fed. for the 3rd group of larvae and the treatment of Ethoxazole at 300 ml/fed. (9.1 %) for the 4th group of larvae.

The obtained data are in agreement with **Ascher and Nemney** (1976a) who found that *S. littoralis* larvae weighting 150 mg when fed on diflubenzuron-treated alfalfa, while it reached 222.4 mg in control. **Asher et al.** (1984) revealed that larvae of *S. littoralis* taken from field-treated plants weighting 30-50 mg but the untreated of the same age (weighting 100-200 mg).

Podoler *et al.* (1985) indicated a remarkable increase in the final weight of the last instar larvae of *S. littoralis* with the use of the hormone analogue (ZR-619).

Dimetry and Abdalla (1988) showed that larval weights of *S. littoralis* were reduced by extracts of seeds of *Abrus precatorius*.

4- Effect on pupation and pupal malformation :

Data in Tables (13-15) demonstrate pupation percentage and pupal abnormality resulted after feeding groups of the 4th instar larvae

Mean larval weight (till 8 days) of S. littoralis late-season field strain (after spraying program) of 4^{th} instar larvae fed for 48h, on treated cotton leaves collected at different time intervals after applying insecticides under field conditions. Table (12).

Rate Insecticides (ml)/ feddan		Mean we	ight of lary	ae (mg.) till	Mean weight of larvae (mg.) till 8 days for groups of larvae fed	groups of la	rvae fed	
		0	n treated le	aves collect	on treated leaves collected at the indicated days.	licated days		
feddan	$1^{st} (0 - 1^{st} day)$	(st day)	$2^{nd} (7 - 8^{th} \text{ day})$	8" day)	3^{rd} (14 - 15 th day)	15" day)	4" (21 - 22th day)	22th day)
	Weight	%	Weight	%	Weight	%	Weight	%
	weight	Change	Weight	Change	1119111	Change	200	Change
Pyriproxyfen 300	116.8	- 47.4	135.1	- 39.1	155.9	- 30.8	171.0	- 24.8
(10% EC) 150	152.9	- 31.3	162.9	- 26.6	182.2	- 19.2	186.3	- 18.1
Chromafenozide 400	71.4	- 67.9	139.9	- 36.9	161.4	- 28.4	178.1	- 21.7
(5% EC) 200	91.3	- 58.9	155.7	- 29.8	166.6	- 26.1	180.6	- 20.6
Azadirachtin 100	96.4	- 56.7	118.4	- 46.6	163.8	- 27.4	164.5	- 27.7
(4.5% EC) 50	115.5	- 48.1	135.9	- 38.8	189.7	- 15.9	196.3	- 13.7
Ethoxazole 25	8.69	- 68.6	184.8	- 16.7	198.0	- 12.2	205.5	- 9.7
(10% SC) 12.5	157.1	- 29.4	194.8	- 12.2	202.8	- 10.1	206.7	- 9.1
Control -	222.4		221.9	·	225.5		227.5	ı

on insecticide-treated leaves collected at different time intervals after application of insecticides under field conditions.

a- Laboratory reference strain:

Survivors after feeding or exposure of the 4th instar larvae for 48h. to insecticide-treated leaves were allowed to pupate as shown in Table (13). Pupation percentages revealed that at the 1st testing interval (0-1 day) no pupation percent was recorded in Chromafenozide and Ethoxazole treatments where accumulated larval mortality reached 100 %. Also, it was obvious that the highest pupation percentage (33 %) in the 1st interval (0-1 day) was recorded in Azadirachtin (50 ml/200l) and that 27 % out of them were abnormal (malformed) while 6 % were healthy. The normal pupation ranged between 4 % and 19 % in the 2nd interval (7-8 day) and between 10 % and 20 % in the 3rd interval (14-15 days) while it ranged between 10 % and 19 % in the 4th testing interval (21-22 day).

Generally, it was obvious that all insecticidal treatments resulted in remarkable reduction percentages in pupation in compare with control. The highest reduction in pupation was recorded by Chromafenozide treatments.

Regarding the latent effects on percent pupal abnormality (malformed) it was obvious that it ranged between 7 and 27% in the 1st testing interval (0-1 day), between 2 and 20 % in the 2nd testing interval (7-8 day), and between 3 and 30 % in the 3rd testing interval (14-15 day) while it was between 13 and 44 % in the 4th testing interval (21-22 day).

However, it was obvious in general that the highest abnormality (malformation) percentages were mostly resulted in the low rates of Azadirachtin (50 ml/200 L) followed by Pyriproxyfen (150 ml/200 L) in most testing intervals, respectively.

Pupational and pupal abnormality percentages of S. littoralis (laboratory strain) for groups of larvae fed for 48h. on treated cotton leaves collected at different time intervals after applying certain insecticides under field conditions. Table (13).

	24.0		% Pupati	on for grou	% Pupation for groups of larvae survived after feeding on treated cotton leaves collected at the indicated days.	survived af	ter feeding cated days.	on treated	
	Kale	187 (0)	1st (0 - 1st day)	$2^{nd} (7 - 8^{th} \operatorname{day})$	8th day)	3^{rd} (14 - 15 th day)	(5th day)	4th (21 - 22th day)	22th day)
Insecticides	foddon	Normal	Abnormal	Normal	Abnormal	Normal	Abnormal	Normal	Abnormal
	300	7	7	17	2	20	11	12	38
Pyriproxyten	150	·	14	14	10	17	23	16	43
(10% EC)	007			9	7	16	3	10	13
Chromatenozide	004			4	20	18	4	11	15
(5% EC)	7007	. -	10	12	-	14	25	13	39
Azadirachtin	100	OI V	27	1 2	15	13	30	19	44
(4.5% EC)	00 40	0	14	15	9	10	28	14	31
Ethoxazole (100% SC)	12.5			19	4	17	29	12	33
Control		96		95	1	95		93	1
COURS									

b- Early-season field strain (before spraying program):

Data in Table (14) revealed that the highest pupation percentage (37%) in the 1st testing interval (0-1 day) was recorded in Azadirachtin at 50 ml/fed. and that 19 % out of them were abnormal while 18 % were healthy. The normal pupation ranged between 14 and 29 % in the 1st testing interval (0-1 day, between 7 and 30 % in the 2nd testing interval (7-8 day), and between 13 and 29 % in the 3rd testing interval (14-15 day) while it was between 4 and 25 % in the 4th testing interval (21-22 day) versus 93-96 % normal pupation in control. Comparison between mean normal pupation in different treatments within the four testing intervals and control, it was obvious that all insecticides resulted in remarkable reduction in normal pupation, and the highest reduction or/and the least mean normal pupation was recorded mostly in Chromafenozide treatments.

As for the abnormality in pupation, it was obvious that it ranged between 5 and 20 % in the 1st testing interval, between 6 and 35 % in the 2nd testing interval and between 15 and 45 % in the 3rd testing interval while it ranged between 39 and 60 % in the 4th testing interval (21-22 day). However, it was obvious that percent abnormal pupation increased remarkably at the later testing intervals where insecticide residues were at its lower level (sublethal). Also, it was obvious that the highest abnormality percentage 16-19 %, 22-35 %, 39-45 % and 58-60 % were recorded in Azadirachtin treatments at 1st, 2nd, 3rd and 4th testing intervals, respectively.

c- Late-season field strain (after spraying program):

Data in Table (15) elucidate pupation percentages either normal (healthy) or abnormal (malformed) for individuals that survived and tolerated feeding and exposure of the 4th instar larvae to treated cotton leaves collected at different time intervals after spray.

Healthy pupation percentages reached 3-8 %, 4-12 %, 5-13 % and 6-18 % for the 1st, 2nd, 3rd and 4th testing intervals versus 93-96

Pupation and pupal abnormality percentages of S. littoralis early-season field strain (before spraying program) for group of larvae fed for 48h. on treated cotton leaves collected at different time intervals after applying insecticides under field conditions. Table (14).

			% Pupati	on for grou	% Pupation for groups of larvae survived after feeding on treated	survived af	ter feeding	on treated	
	Rate			cotton lear	cotton leaves collected at the indicated days.	l at the indi	cated days.		
Insecticides	(ml)/	$1^{37}(0 -$	$1^{st} (0 - 1^{st} day)$	$2^{nd} (7 - 8^{th} day)$	8" day)	3^{rd} (14 - 15 th day)	15 th day)	4^{th} (21 - 22 th day)	22th day)
	feddan	Normal	Abnormal	Normal	Abnormal	Normal	Abnormal	Normal	Abnormal
Pyriproxyfen	300	14	15	20	13	28	18	22	39
(10% EC)	150	17	20	22	18	17	15	25	48
Chromafenozide	400	1	ı	8	6	13	29	4	45
(5% EC)	200	17	1	7	15	27	25	7	48
Azadirachtin	100	16	16	24	22	23	39	2	09
(4.5% EC)	. 50	18	19	25	35	23	45	- 24	58
Ethoxazole	25	22	5	27	9	22	28	18	40
(10% SC)	12.5	29	7	30	12	29	25	6	50
Control		96	•	95		95	•	93	•

Pupation and pupal abnormality percentages of S. littoralis late-season field strain (after spraying program) for groups of larvae fed for 48h. on treated cotton leaves collected at different time intervals after applying insecticides under field conditions. Table (15).

			% Dung	for for					
	Rate		70 rupar	on tor grou cotton lea	cotton leaves collected at the indicated described	survived at	fter feeding	on treated	
Insecticides	(ml)/	$1^{st}(0 -$	$1^{57} (0 - 1^{57} day)$	2nd (7 - 8th	8th day)	3rd (14 15th 3	teth 3	th in	
	feddan	Normal	Abnormal	Normal	Ahnomol	- +T	(yab)	4" (21 - 22th day)	22th day)
Perincoration	000			TAGEMIAL	ADHOLINA	Normal	Abnormal	Normal	Abnormal
A Jarbi Oayten	200	0	∞	7	6	10	0	Ç	
(10% EC)	150	000	2	0	9	2	0	71	10
Chuomoforeil	001		1		0	13	2	13	=
Can omalenozide	400	4	n	2	4	v	11		
(5% EC)	200	m	4	-		,	TT	0	15
Azadinoohti-	100		-	t	\	9	13	6	6
AZZAUH ACHUN	100	9	Э	7	4	7	o	Ç	
(4.5% EC)	50	4	٧	12			7	13	11
Ethoxazole	25	v	,	77	0	177	7	18	15
	61	0	0	∞	7	10	2	12	16
(10% SC)	12.5	8	S	6	6	11		77	10
Control	,	96		90		11	4	12	18
				22		5		00	

% for control while the pupal abnormality percent reached 2-8 %, 4-9 %, 2-13 % and 9-18 % for the same testing intervals, respectively versus no abnormality percent in control. It was worthy mentioning that the abnormality recorded ca. 40-50 % of the overall pupation percentages in all treatments and that the highest abnormality percent was recorded remarkably in Pyriproxyfen, Chromafenozide and Ethoxazole during the 1st, 2nd, 3rd and 4th testing intervals, respectively.

Reviewing the literature, similar low level of normal pupation was recorded when studying the latent effect of feeding *S. littoralis* larvae on Pyriproxyfen (Moawad et al. 1996) or on IKI-7899 (Abdel Mageed et al., 1986). Likewise, El-Badawy (1979) indicated that percentages of pupation and adult eclosion decrease significantly with the increase of diflubenzuron (IGR) doses as well as the treated instar of *S. littoralis*. Also, a positive correlation was recorded between the age of the residues of emulsion of seed kernels of *Azadirachtin indica* and the pupation rate (Meisner et al., 1983).

Recently, Antonious and Rizk (1994) found that larval treatment with a high concentration of neem seed oil resulted in larvae being highly impaired at ecdysis as exhibited by various deformities (malformation) in the formed pupae.

5- Effect on pupal duration:

Data in Tables (16-18) demonstrate the pupal duration of *S. littoralis* after feeding the 4th instar larvae for 48 h. on treated cotton leaves, collected at four time intervals (0-1 days, 7-8 days, 14-15 days and 21-22 days) after spraying different pesticides, followed by feeding three days on untreated leaves.

a- Laboratory reference strain:

The effect of the tested insecticides on the pupal duration of *S. littoralis* (laboratory strain) are presented in Table (16). A pronounced

decrease in the pupal duration was recorded in all treatments in pupae resulted from the four groups of larvae, ranged between 33.3 and 72.2 %, 18.2 and 60.0 %, 5.6 and 44.4 % and 8.3 and 33.3 % in the 1st (0-1 day), 2nd (7-8 day), 3rd (14-15 day) and 4th (21-22 day) testing intervals, respectively.

However, the highest reduction in pupal duration was recorded in Pyriproxyfen treatments whereas the least was recorded in Ethoxazole treatments.

b- Early-season field strain (before spraying program):

The results illustrated in Table (17) show that the pupal durations in control were 5.4, 5.5, 5.5 and 6.0 days for the 1st, 2nd, 3rd and 4th groups of larvae, the mean pupal duration was 5.6 days. All insecticidal treatments caused considerable decrease in the pupal durations, of Pyriproxyfen (2.0-6.0 days), Chromafenozide (3.0-5.5 days), Azadirachtin (3.0-6.0 days) and Ethoxazole (3.3-5.5 days) during all testing intervals.

The calculated reduction in pupal duration relative to untreated control reached 22.2-62.9 %, 27.3-45.9 %, 14.6-45.5 % and 8.3 % for treatment of the 1st, 2nd, 3rd and 4th testing intervals, respectively. However, the highest reduction in pupal duration was recorded for Pyriproxyfen treatments while the least was recorded for Ethoxazole.

c- Late-season field strain (after spraying program):

The data representing the pupal duration of *S. littoralis* after the feeding of 4th instar larvae on treated cotton leaves collected at different time intervals are presented in Table (18). These data revealed that the 1st group of larvae which were fed on treated cotton leaves collected at 0 and 1 days from application was remarkably affected by the treatments of insecticides, the pupal durations were 3.0-3.5, 0, 4.5-5.0 and 4.8-5.2 days for Pyriproxyfen, Chromafenozide, Azadirachtin

Table (16). Pupal duration of S. littoralis (Laboratory strain) for groups of larvae fed for 48 h. on treated cotton leaves collected at different time intervals after applying insecticides under field conditions.

	Date	M	ean pupal d	uration (in c	lays) resulte	Mean pupal duration (in days) resulted from groups of larvae survived after feeding on treated cotton leaves collected at the indicated days.	ups of larvae	survived af	ter
Insecticides	(ml)	13 (0 -	$1^{st} (0 - 1^{st} \operatorname{day})$	2 nd (7 - 8 th day)	8th day)	$3^{rd}(14-15^{th} \text{ day})$	15th day)	$4^{th}(21 - 22^{th} \text{ day})$	22th day)
	feddan	Princetion	%	Duration	%	Duration	%	Duration	% ;
		Duration	Change	Duramon	Change	Toma and	Change		Change
Pyriproxyfen	300	1.5	- 72.2	2.2	- 60.0	3.0	- 44.4	5.5	- 8.3
(10% EC)	150	1.8	- 66.7	2.4	- 56.4	3.0	- 44.4	5.5	- 8.3
Chromafenozide	400	0		4.0	- 27.3	4.1	- 24.1	4.0	- 33.3
(5% EC)	200	0		4.4	- 20.0	5.0	- 7.4	4.0	- 33.3
Azadirachtin	100	3.5	- 35.2	4.1	- 25.5	4.8	-11.1	0.9	0
(4.5% EC)	20	3.6	- 33.3	4.5	- 18.2	5.1	- 5.6	0.9	0
Ethoxazole	25	0		3.7	- 32.7	4.5	- 16.7	5.5	-8.3
(10% SC)	12.5	0		4.5	- 18.2	4.6	- 14.8	5.5	- 8.3
Control		5.4		5.5		5.5	•	0.9	

Pupal duration of S. littoralis early-season field strain (before spraying program) for groups of larvae fed for 48h, on treated cotton leaves collected at different time intervals after applying insecticides under field conditions. Table (17).

	Rate	N	lean pupal feedin	Mean pupal duration (in days) resulted from groups of larvae survived after feeding on treated cotton leaves collected at the state of	days) result	ed from grou	ups of larvae	e survived at	ter
Insecticides	(ml)/		$1^{st}(0-1^{st}day)$	2 nd (7 -	2"d (7 - 8th dav)	3rd (14	3rd (14 15th Jan)	ed days.	
	feddan		%		70	111	(KRD CT	4 (21 - 22" day)	22''' day)
		Duration	Change	Duration	Change	Duration	Change	Duration	%
Pyriproxyfen	300	2.0	- 62.9	3.0	45.0		Cuange		Change
(C)	400			200	43.7	6.5	- 18.2	0.9	0
(10% EC)	061	2.5	- 53.7	3.2	- 41.8	4.7	-146	0.9	
Chromafenozide	400	0	1	3.0	15 5		7.4.0	0.0	0
(F07 EC.)	000	(200	- 43.3	3.0	- 45.5	5.5	- 8.3
(370 EC)	200	0		3.5	- 36.4	3.4	20 7		
Azadirachtin	100	3.0	- 44 4	3.6			7.06 -	0.0	-8.3
70 Z 70 Z 70	9			2.0	- 34.0	3.5	- 36.4	0.9	0
(4.3% EC)	200	3.5	- 35.2	4.0	- 27.3	4.0	- 773	03	
Ethoxazole	25	3.3	- 38.9	3.5	- 36.4	4.	500	0.0	0
(10% SC)	12.5	4.2	- 22.2			C.	7.81 -	5.5	- 8.3
			7.77	7.4	- 2/.3	4.5	- 18.2	5.5	200
Control		5.4		5.5	ï	5.5		60	

Pupal duration of S. littoralis late-season field strain (after spraying program) for groups of larvae fed for 48h, on treated cotton leaves collected at different time intervals after applying insecticides under field conditions. Table (18).

		M	ean pupal d	Mean pupal duration (in days) resulted from groups of larvae survived after	lays) resulte	ed from grou	ips of larvae	survived at	fter
	Kate		reeding	reeding on treated cotton leaves conected at the indicated days.	cotton leave	s collected a	t the marcat	eu uays.	
Insecticides	(ml)/	$1^{st}(0 -$	$1^{st} (0 - 1^{st} day)$	$2^{na} (7 - 8^m \text{ day})$	8''' day)	3^{ra} (14 - 15 ^m day)	15''' day)	4" (21 - 22th day)	22th day)
	feddan		%	A.	%		%		%
		Duration	Change	Duration	Change	Duration	Change	Duration	Change
Pyriproxyfen	300	3.0	- 44.4	5.5	0	5.5	0	0.9	0
(10% EC)	150	3.5	- 35.2	5.5	0	5.5	0	0.9	0
Chromafenozide	400	0	•	5.4	- 1.8	5.5	0	0.9	0
(5% EC)	200	0	Ĩ	5.4	- 1.8	5.5	0	0.9	0
Azadirachtin	100	4.5	- 16.7	5.5	0	5.5	0	0.9	0
(4.5% EC)	90	5.0	- 7.4	5.5	0	5.5	0	0.9	0
Ethoxazole	25	4.8	- 11.1	5.4	- 1.8	5.5	0	0.9	0
(10% SC)	12.5	5.2	- 3.7	5.4	- 1.8	5.5	0	0.9	0
Control	1	5.4	•	5.5	•	5.5	•	0.9	1

and Ethoxazole, respectively, whereas pupal period in control was 5.4 days. It is obvious from Table (18) that the feeding of the 2nd, 3rd and 4th groups of larvae on treated cotton leaves collected at 7-8, 14-15 and 21-22 days, respectively did not affect the pupal duration of S. littoralis.

In conclusion, it could be said that all treatments decreased the pupal duration of *S. littoralis* within the three tested strains, the highest decrease was recorded in laboratory and early-season strain particularly in Chromafenozide (400 ml/fed. followed by Pyriproxyfen (300 ml/fed.). In agreement, **Moawad** *et al.* (1996) found that pyriproxyfen at 10 ppm cause lifetime of larvae being longer than control and reduced pupal duration.

6- Effect on pupal weight:

Data presented in Tables (19-21) the effect of tested insecticides on the pupal weight of the three different *S. littoralis* strains after feeding four groups of 4th instar larvae on treated cotton leaves collected at different time intervals after spray.

a- Laboratory reference strain:

The obtained data in Table (19) expressed as percent changes in pupal weight showed that the highest decrease was recorded in the 1st interval by the higher rate of Pyriproxyfen (-62.8 %), in the 2nd interval by the higher rate of Azadirachtin (-49.8 %), in the 3rd interval by Chromafenozide (-37.6 %) while it was (-29.9 %) for the same treatment at the 4th testing interval. In general it was obvious that Pyriproxyfen at 300 ml/fed. resulted in remarkably the least mean pupal weight (85.7 mg/pupa) when compared with the control (230.7 mg/pupa) which was obviously higher than other treatments.

Table (19). Pupal weight (mg.) of S. littoralis (laboratory strain) for groups of larvae fed for 48 h. on treated cotton leaves collected at different time intervals after applying insecticides under field conditions.

		M	on woight	ot numbe (m)	r) resulted	Mass waight of number (mg.) resulted from oronns of larvae survived after	of larvae s	urvived afte	
	Rate	II.	feedin	or pupae (mg	cotton leav	feeding on treated cotton leaves collected at indicated days.	it indicated	days.	
Insecticides	(m)	1st (0 - 1st day)	r dav)	2"d (7 - 8" day)	th day)	3^{rd} (14 - 15 th day)	5th day)	$4^{th} (21 - 22^{th} day)$	2" day)
	feddan	Weight	%	Weight	%	Weight	%	Weight	%
		(mg)	Change	(mg)	Change	(mg.)	Change	(mg.)	Change
Pyrinroxyfen	300	85.7	- 62.8	135.4	- 43.5	182.3	- 24.1	195.3	- 21.9
(10% EC)	150	90.06	- 60.9	148.9	- 37.9	201.6	- 16.1	225.2	- 9.9
Chromafenozide	400	,	,	143.1	- 40.3	150.0	- 37.6	175.2	- 29.9
(5% EC)	200		,	186.3	- 22.3	200.1	- 16.7	233.3	- 6.7
Azadirachtin	100	100.0	- 56.7	120.4	- 49.8	170.8	- 28.9	201.2	- 19.5
(4.5% EC)	50	140.1	- 39.3	152.5	- 36.4	182.9	- 23.9	202.3	- 19.1
Ethoxazole	25		•	150.0	- 37.4	177.3	- 26.2	242.2	3.1
(10% SC)	12.5			188.1	- 21.5	195.0	- 18.9	242.3	- 3.1
Control		230.7	•	239.7		240.3	1	250.0	

b- Early-season field strain (before spraying program):

As shown in Table (20) it was obvious that all tested insecticides exhibited considerable reduction in pupal weight ranged between-16.0 and -38.8 % in the 1st testing interval, between -16.6 and -41.4 % in the 2nd testing interval and between -4.3 and -24.3 % in the 3rd testing interval while the reduction was between -2.0 and -7.2% in the 4th testing interval. The highest reduction in pupal weight was recorded for Pyriproxyfen (38.8 %) at the rate of 300 ml/fed. in the 1st testing interval., and for Chromafenozide (41.4 %) at the rate of 400 ml/fed. at the 2nd testing interval and for Azadirachtin (24.3 %) at rate of 100 ml/fed. in the 3rd testing interval. However, based on comparison between all testing intervals it was obvious that the least mean pupal weight (140.4 mg/pupa) was recorded for Chromafenozide treatment (400 ml/fed.) which was remarkably lower than other treatments including control.

c- Late-season field strain (after spraying program):

The obtained data in Table (21) expressed a percent change in pupal weight revealed that all insecticidal treatments during the 3 first testing intervals resulted in remarkable reduction in pupal weight ranged between 56.6 and 30.3 % in the 1st testing interval (0-1 day), 55.3 and 19.9 % in the 2nd testing interval (7-8 day) and 54.1 and 16.7 % in the 3rd testing interval (14-15 day). Chromafenozide at 400 ml/fed. recorded the highest reduction in pupal weight in the first three testing intervals. As for the 4th testing interval similar trend was achieved but of less magnitude except in case of both treatments of Pyriproxyfen and Ethoxazole where considerable increase of +20.2 and 24.1 % was recorded for the former and +32.2 and 40.0 % was recorded for the later.

In agreement, **Zidan** *et al.* (1996) recorded a high reduction in pupal weight of *S. littoralis* when larvae were treated with Pyriproxyfen, Dipel 2X and mineral oil. It is suggested by the authors

Table (20). Pupal weight (mg.) of S. littoralis early-season field strain (before spraying program) for groups of larvae fed for 48 h. on treated cotton leaves collected at different time intervals after applying insecticides under field conditions.

		Mean weig	ht of pupa	ie (mg.) resi	ulted from	Mean weight of pupae (mg.) resulted from groups of larvae survived after feeding on	arvae sur	rived after	feeding on
	Rate	treated cott	on leaves c	treated cotton leaves collected at indicated days.	dicated da	vs.			
Insecticides	(m)/	$1^{37} (0 - 1^{37} \text{ day})$	day)	$2^{nd} (7 - 8^{th} day)$	" day)	3^{rd} (14 - 15 th day)	5th day)	4''' (21 - 22'' day)	2''' day)
	feddan	Weight	%	Weight	%	Weight	%	Weight	%
		(mg)	Change	(mg)	Change	(mg•)	Change	(mg·)	Change
Pvrinroxvfen	300	141.2	- 38.8	146.0	- 39.1	199.7	- 16.9	232.1	- 7.2
(10% EC)	150	156.7	- 32.1	169.6	- 29.3	220.0	- 8.5	233.3	- 6.7
Chromafenozide	400	3	•	140.4	- 41.4	195.0	- 18.9	233.0	- 6.8
(5% EC)	200	193.8	- 16.0	200.0	- 16.6	220.0	- 8.5	234.2	- 6.3
Azadirachtin	100	161.2	- 30.1	178.9	- 25.4	182.0	- 24.3	233.3	- 6.7
(4.5% EC)	50	182.5	- 20.9	184.0	- 23.2	211.0	- 12.2	245.0	- 2.0
Ethoxazole	25	180.0	- 21.9	186.1	- 22.4	199.5	- 17.0	250.0	0
(10% SC)	12.5	182.7	- 20.8	189.5	- 20.9	230.0	- 4.3	250.0	0
Control	t	230.6	1	239.7		240.3	•	250.0	•

Table (21). Pupal weight (mg.) of S. littoralis late-season field strain (after spraying program) for groups of larvae fed for 48 h. on treated cotton leaves collected at different time intervals after applying insecticides under field conditions.

,	Rate	2	lean weigh feedin	Mean weight of pupae (mg.) resulted from groups of larvae survived after feeding on treated cotton leaves collected at indicated decided at the contract of th	g.) resulted	from group	s of larvae	survived aft	er	T-
Insecticides	(m)/	$1^{57} (0 - 1^{57} day)$	1st day)	2nd (7 - 8th dav)	dav)	3rd (14 15th Jan)	Eth dan	u days.	- 1/4	
	feddan	Weight	%	Waight	/0	1 - 47 6	c nay)	4 (21 -	4 (21 - 22" day)	_
		(mg)	Change	(ma)	0,70	Weight	%	Weight	%	_
Demission	000		9	(48)	Спапве	(mg.)	Change	(mg.)	Change	
Lyriproxylen	300	112.7	- 51.1	115.4	- 51.9	120.7	40.0	2000		
(10% EC)	150	1403	300	1000		1.07	- 47.0	300.3	+ 20.2	
:		0.01	- 37.4	190.0	- 20.5	200.3	- 16.7	310.3	+ 24 1	
Chromatenozide	400	100.1	- 56.6	107.2	- 553	1103	51.1	0000		
(5% FC)	200	160.9	, ,	, 00,		270.5	- 24.1	200.8	- 19.6	
(27.00)	2007	100.0	- 30.3	192.1	- 19.9	120.3	- 49 9	2006	10.7	
Azadirachtin	100	119.6	- 48.1	129.7	- 45 9	140.2	41.7	0.004	- 17./	
(4.5% EC)	50	122.3	- 46 0	1336	1	7.0.1	- 41./	200.4	- 19.8	
Toroxout I	40		7.05	132.0	- 44./	160.3	- 33.3	240.1	- 4.0	
Ethoxazole	67	132.7	- 42.5	143.1	- 40.3	1502	- 375	220 €		
(10% SC)	12.5	140.6	- 39.0	152 4	36.4	150.1	0.70	530.3	+32.2	
Control		1 000		1.10	1.00	100.1	- 33.4	350.1	+ 40.0	
Control	ı	730.7		239.7	r	240.3	1	250.0		

that the little weight of pupa could be due to lower total protein content in treated larvae as compared with control ones. Likewise, the highly pronounced reduction in mean pupal weight recorded for Pyriproxyfen in the present study are in agreement with findings of **Kelany** el al. (1991) indicating that the mean weight of *Musca domestica* pupae reared on medium treated with aqueous extract of neem seed kernel were about one half of those obtained from untreated medium.

7- Effect on pupal mortality:

Data in Tables (22-24) indicate the latent effects on the pupal morality when the 4th instar larvae were fed for 48 hr. on treated leaves collected at different time intervals after insecticides application.

a- Laboratory reference strain:

Data in Table (22) revealed slight latent pupal mortality ranged between 2-4 % in 1st testing interval (0-1 day) and between 4-7 % in the 2nd testing interval (7-8 day) whereas relatively low latent pupal mortality of 4-18% and 4-14 % was recorded at the 3rd (14-15 day)and 4th (21-22 day) testing intervals, respectively. However, the highest pupal mortality was recorded in Chromafenozide (16-18%) at the 3rd testing interval and in Azadirachtin (13-19 %) at the 4th testing interval.

b- Early-season strain (before spraying program):

Data in Table (23) revealed slight increase in latent pupal mortality relative to laboratory reference strain. The recorded latent pupal mortality ranged between 8-21% in 1st testing interval (0-1 day), 4-22 % in the 2nd testing interval (7-8 day) and 3-27 % in the 3rd testing interval (14-15 day) while it reached 2-18% in the 4th testing interval (21-22 day). However, the highest pupal mortality percentages were recorded by Ethoxazole treatments (13 and 21 %) in the 1st

Table (22). Pupal mortality of S. littoralis (Laboratory strain) for groups of larvae fed for 48 h. on treated cotton leaves collected at different time intervals after applying insecticides under field conditions.

		T	7	T	T	T	T		T	T	T
ved after	ed days.	4th (71 - 27th day)	(kpn 77 - 17) .	0 0	0 0	10		13	19	4 4	0
% Pupal mortality resulted from groups of larvae survived after	feeding on treated cotton leaves collected at the indicated days.	3rd (14 - 15th dav)	13	11	11	10	14	1 14	4		, ;
mortality resulted from	on treated cotton leaves	$2^{nd} (7 - 8^{th} day)$	5	5	9	4		9 4		7	F
% Pupal	feeding	$1^{st} (0 - 1^{st} day)$	3	2			4	3	•		6
Rate	(m)/	feddan	300	150	400	200	100	50	25	12.5	
9	Insecticides		Pyriproxyfen	(10% EC)	Chromafenozide	(5% EC)	Azadirachtin	(4.5% EC)	Ethoxazole	(10% SC)	Control

Table (23). Pupal mortality of S. littoralis early-season field strain (before spraying program) for groups of larvae fed for 48 h. on treated cotton leaves collected at different time intervals after applying insecticides under field conditions.

	Rate	% Pupal	mortality resulted fron	% Pupal mortality resulted from groups of larvae survived after	ved after
Insecticides	(ml)/	feeding	on treated cotton leaves	feeding on treated cotton leaves collected at the indicated days.	ed days.
	feddan	$1^{st} (0 - 1^{st} day)$	$2^{nd} (7 - 8^{th} day)$	3^{rd} (14 - 15 th day)	4th (21 - 22th day)
Pyriproxyfen	300	6	5	10	10
(10% EC)	150	13	4	3	11
Chromafenozide	400		8	13	4
(5% EC)	200	14	7	27	7
Azadirachtin	100	8	19	23	2
(4.5% EC)	50	13	22	23	18
Ethoxazole	25	13	10	3	15
(10% SC)	12.5	21	8	11	2
Control	•	6	11	12	13

testing interval, by Azadirachtin treatments (19 and 22 %) in the 2nd testing interval, by Chromafenozide treatments (13 and 27 %) in the 3rd testing interval.

Comparison between different treatments revealed that the highest pupal mortality during all testing intervals was recorded for Chromafenozide (13 and 27 %) and Azadirachtin (23 and 23 %) at the 3rd testing interval (14-15 day).

c- Late-season strain (after spraying program):

Data in Table (24) revealed slight percent of latent pupal mortality relative to both laboratory reference strain (Table 22) and the control of same strain (Table 24). The recorded low latent pupal mortality percentages averaged 1-8 %, 1-8 %, 1-5 % and 1-6 % in different treatments at 1st, 2nd, 3rd and 4th resting intervals versus 9,11,12 and 13 % for the control at same prementioned intervals. However, the least pupal mortality was recorded for Pyriproxyfen at 300 ml/fed. And Azadirachtin at 100 ml/fed., recording 1.5 and 1.75 % for both, respectively versus 11.25 % for control.

Such relatively low percentages of pupal mortality recorded her for the tested pesticides could be explained in view of the natural accumulated mortality through out larval and prepupal stages where the differences between treatments and the control were relatively slight.

El-Sayed and Sayed (1982) found that feeding 1st -5th instar larvae of S. littoralis on leaves treated with 0.2-0.5% suspension of Azadirachta indica caused 100% larval mortality by the end of larval stage and also caused high pupal mortality.

Recently **Zidan** *et al.* (1998) found that Ethoxazole (10 % SC) at 31.01 ppm reduced pupal weight. The latent effect of the compound caused 45.9 % pupal mortality.

Table (24). Pupal mortality of S. littoralis late-season field strain (after spraying program) for groups of larvae fed for 48 h. on treated cotton leaves collected at different time intervals after applying insecticides under field conditions.

	Rate	% Pupal	% Pupal mortality resulted from groups of larvae survived after	a groups of larvae survi	ved after
Treactioides	(lm)	feeding	feeding on treated cotton leaves collected at the indicated days.	collected at the indicat	ed days.
	foddan	1 st (0 - 1 st dav)	$2^{nd} (7 - 8^{th} day)$	3^{rd} (14 - 15 th day)	4^{th} (21 - 22 th day)
	300		2	1	2
Pyriproxyten			0	5	4
(10% EC)	150	∞		,	
Chromofonozide	400	2	5	1	ı
CIII UIII AI CIII CIII CIII CIII CIII CI	000	"	4	2	3
(3% EC)	7007		-0.2	•	C
Azadirachtin	100	2	2	1	4
()4 704 F)	50		. 5	2	9
(4.3% E.C.)	3	-	œ	co	ĸ
Ethoxazole	25	1			۲
(10% SC)	12.5	3	1	4	7 .
Control	•	6	11	12	13

8- Effect on adult emergence :

Data in Tables (25-27) illustrate the latent effects on adult emergence following feeding 4th instar larvae for 48h on cotton leaves treated with certain insecticides, followed by feeding for 3 days on untreated leaves.

a- Laboratory reference strain:

As shown in Table (25) it was obvious that the adult emergence was highly affected as demonstrated by the remarkably low percent of adult emergence which averaged 3-6, 6-12, 6-12 and 6-10 % in 1st (0-1 day) 2nd (7-8 day), 3rd (14-15 day) and 4th (21-22 day) testing intervals, respectively. However, it is of interest to note that no adults were obtained in Chromafenozide at different testing intervals, Azadirachtin at the 3rd and 4th testing intervals and in Ethoxazole at the 1st testing interval (0-1 day). In contrast, control treatment resulted in 80-87 % adult emergence.

Comparison between treatments in concern with adult emergence revealed that while control treatment resulted in the highest adult emergence (80-87 %), Chromafenozide treatments were the least (Zero %) and other treatments were considerably different than both prementioned ones.

b- Early-season field strain (before spraying program):

Data in Table (26) indicate in general that adult emergence percentages were slightly higher than in case of laboratory strain, recording averages of 3-9, 3-22, 14-19 and 3-14 % for the 1st (0-1 day), 2nd (7-8 day), 3rd (14-15 day) and 4th (21-22 day) testing intervals, respectively. However, Chromafenozide and Azadirachtin exhibited almostly results similar to those recorded for the laboratory strain where no adult emergence was recorded at the same testing intervals. Comparison between different treatments revealed that the highest adult emergence (80-87 %) was recorded for control treatment and came apart Ethoxazole (3-22 %) and Pyriproxyfen (4-18 %) while

Table (25). Adult emergence percentage of S. littoralis (Laboratory reference strain) for groups of larvae fed for 48h. on treated cotton leaves collected at different time intervals after applying insecticides under field conditions.

	Rate	% Adult em	ergence resulted from	% Adult emergence resulted from groups of larvae survived after 48h.	d after 48h.
Insecticides	(m)/	feeding	on treated cotton leave	feeding on treated cotton leaves collected at the indicated days.	ed days.
	feddan	$1^{57} (0 - 1^{57} day)$	$2^{nd} (7 - 8^{th} day)$	3^{rd} (14 - 15 th day)	4^{th} (21 - 22 th day)
Perincoxefen	300	4	12	7	9
(10% EC)		3	6	9	8
Chromafenozide	400	0	0	0	0
(5% EC)	200	0	0	0	0
Azadirachtin		9	9	0	0
(4.5% EC)	50	3	8	0	0
Fthoxazole	25	0	10	9	10
(10% SC)		0	12	12	9
Control		87	84	83	80

Table (26). Adult emergence percentage of S. littoralis early-season field strain (before spraying program) for groups of larvae fed for 48 h. on treated cotton leaves collected at different time intervals after applying insecticides under field conditions.

	Rate	% Adult en	dergence resulted from		
Insecticides	(ml)/	feeding	on treated cotton leave	feeding on treated cotton leaves collected at the indicated	d after 48h.
	feddan	$1^{57} (0 - 1^{57} dav)$	2nd (7 - 8th day)	3rd (14 15th)	ed days.
Pyriproxyfen	300		(Com (C) =	(Yeb CI - 41) C	4" (21 - 22" day)
() () () () () () () () () ()	_	,	CI	18	12
(10% EC)	130	4	18	14	7.
Chromafenozide	400	0	0	t, c	14
(5% EC)	200	n			0
		0	0	0	O
Azadirachtin	100	8	5		
(4.5% EC)	. 50	5			0
Ethoxazole	25	0	,	0	9
(100)	000		17	19	3
(10% SC)	17.5	8	22	18	-
Control		87	84	0.5	
				93	80

both of Chromafenozide (0.0-3.0 %) and Azadirachtin (3-8 %) being remarkably the least in this respect.

c- Late-season field strain (after spraying program):

Data in Tale (27) illustrate the adult emergence percentages of late-season field strain in response of feeding 4th instar larvae on cotton leaves treated with the tested insecticides. It was obvious that adult emergence percentages in treatments was remarkably low in compare to control treatment (80-87%) and averaged 2-5 %, 5-8 %, 4-10 % and 5-12 % for the 1st (0-1 day), 2nd (7-8 day), 3rd (14-15 day) and 4th (21-22 day) testing intervals, respectively. Comparison between treatments revealed that Chromafenozide treatments was the least and differences between all treatments were negligible but were severely lower than control treatment.

Our data agreed in general with findings of Loke et al. (1992) concerning the toxicity of neem oil Azadirachta indica against larvae of Plutella xylostella where it caused growth disruptive effects, delayed adult emergence and abnormal adults were observed with sublethal concentrations.

9- Effect on adult longevity:

The latent effect of feeding 4th larval instar larvae on cotton leaves treated with insecticides are shown in Tables (28-30) when adult longevity was concerned.

a- Laboratory reference strain:

Data in Table (28) revealed remarkable decrease than control in adult longevity by 35.3-46.7 % for Azadirachtin and by 17.7-25.3 % for Ethoxazole during different testing interval. In contrast adult longevity in Pyriproxyfen treatments increased by 5.8-86.7 % during the first three testing intervals. Comparison between different treatments revealed that variations between treatments and control

groups of larvae fed for 48 h. on treated cotton leaves collected at different time intervals after Table (27). Adult emergvence percentage of S. littoralis late-season field strain (after spraying program) for applying insecticides under field conditions.

	Rate	% Adult en	% Adult emergence resulted from groups of larvae survived after 48h.	groups of larvae survive	d after 48h.	
Insecticides	(m)/	feeding	feeding on treated cotton leaves collected at the indicated days.	collected at the indicat	ed days.	
	feddan	$1^{57} (0 - 1^{57} day)$	$2^{nd} (7 - 8^{th} day)$	3 rd (14 - 15 th day)	4th (21 - 22th day)	
Pyrinroxyfen	300	4	5	6	10	
EC)	150	0	9	8	6	
	400	2	0	4	5	
\Box	200	0	0	4	9	
Azadirachtin	100	4	5	9	11	
(4.5% EC)	50	4	7	10	12	
Ethoxazole	25	4	0	7	6	
% SC)	12.5	5	8	7	10	
Control	1	87	84	83	80	

Table (28). Adult longevity of S. littoralis (Laboratory strain) for groups of larvae fed for 48h. on treated cotton leaves collected at different time intervals after applying insecticides under field conditions.

			dult longev	Adult longevity (in days) resulted from groups of larvae survived after	resulted fr	om groups	f larvae su	rvived after	
	Doto	1	feeding 0	feeding on treated cotton leaves collected at the murant	tton leaves	collected at the man	Elle muncat	4th (21 - 22th day)	2 th day)
34	Kate	1st (0 - 1st day)	(st day)	$2^{nd} (7 - 8^m \text{ day})$	8" day)	3 (14 - 1	O.	Description	%
Insecticides	(mi)/	Duration	%	Duration	%	Duration	Срапое	Duranon	
	Icanan	Dui auon	Change		Change		158	5.1	- 43.3
		,	+ 53 3	10.3	+ 5.8	7.6	0.0		000
D. r. inroxvfen	300	11.5	0.00 +		+ 22 3	116	+33.3	5.2	- 44.2
Tyribrasher EC	150	14.0	+86.7	12.5	1.00.1				•
(10% EC)	22.			•					
Chromafenozide	400	ř.	•			•	ì		
Chromatenograe	000	,	٠	•					٠
(5% EC)	700		!		- 36 5	1	•		
	100	4.0	- 46.7	4.0	2.00		3		
Azadirachtın	201		413	5.5	- 35.3		-	(1	100
(4.5% EC)	20	4.4	2.11.		23.5	6.5	- 25.3	7.3	- 10.7
	35	•	1	6.0	- 43.3		401	7.4	17.8
Ethoxazole	77			7.0	- 17.7	7.0	C.61 -	-	
(10% SC)	12.5	1	•	2.7		8.7		0.6	1
	,	7.5	•	8.5	-				
Control									

in adult longevity were almostly of low magnitude, and the treatments could be arranged descendingly as follow: Pyriproxyfen, Control, Ethoxazole, Azadirachtin and Chromafenozide, respectively.

b- Early-season field strain (before spraying program):

Results tabulated in Table (29) revealed that the longevity of the resulted adults was mostly similar to control except in case of Pyriproxyfen low rate (150 ml/fed) and both rates of Chromafenozide where the adult longevity was remarkably longer in the former and shorter in the later, respectively. During the 1st testing interval (0-1) the lower rate of Chromafenozide (200 ml/fed) while the least reduction in adult longevity (-2.7 %) was recorded in Ethoxazole (12.5 ml/fed) whereas Pyriproxyfen treatments resulted in 14.7 and 80.0 % increase in adult longevity at 300 and 150 ml/fed., respectively. The latent effect fall down with time elapse where the change in adult longevity reached-16.7, zero, -27.8 and -5.6 % decrease in adult longevity for the prementioned treatments, respectively.

c- Late-season field strain (after spraying program):

Data in Table (30) revealed that the late-season field strain which tolerate several insecticides in the spraying program, exhibited also tolerance to the tested untraditional treatments tested in the present study which reflexed in negligible increase in adult longevity of +5.3, +4.0 and +4.4 % for Pyriproxyfen at 300 ml/fed., Azadirachtin at 100 and 50 ml/fed., respectively at the first testing interval (0-1 day) where the tested compounds were at its powerful concentration. No remarkable latent effect was recorded during the other three testing intervals where the adult longevity was almostly similar as control.

Our results agreed with findings of **Moawad** et al.(1996) that Pyriproxyfen at LC₅₀ level in leaf dipping experiments against larval

Adult longevity of S. littoralis early-season field strain (before spraying program) for groups of larvae fed for 48 h. on treated cotton leaves collected at different time intervals after applying insecticides under field conditions. Table (29).

- Le C	$1^{st} (0 - 1^{st} day)$	teering	on treated c	otton leaves	feeding on treated cotton leaves collected at the indicated days.	t the indica	ted days.	
		(x day)	$2^{nd} (7 - 8^{th} \text{ day})$	8th day)	3^{rd} (14 - 15 th day)	15 th day)	$4^{th} (21 - 22^{th} day)$	22th day)
	Jura flon	%	Duration	%	Duration	%	Duration	%
		Change		Change		Change		Change
Pyrinroxyten	8.6	+ 14.7	8.5	0	7.5	- 13.8	6.5	- 27.8
(10 % EC) 150	13.5	+ 80.0	11.5	+35.3	8.7	0	8.5	- 5.6
Chromafenozide 400		1	•		,	•	1	1
(5% EC) 200	4.1	- 45.3	•	•			•	•
Azadirachtin 100	5.4	- 28.0	7.6	- 10.6		ı		•
4.5% EC) 50	5.5	- 26.7	7.9	- 7.1	1.		9.0	0
Ethoxazole 25	7.2	- 4.0	8.0	-5.9	8.0	- 8.0	8.5	- 5.6
(10% SC) 12.5	7.3	- 2.7	8.5	0	8.5	- 2.3	9.0	0
Control	7.5		8.5		8.7	1	9.0	•

Adult longevity of S. littoralis late-season field strain (after spraying program) for groups of larvae fed for 48h. on treated cotton leaves collected at different time intervals after applying insecticides under field conditions. Table (30).

Rate cotton	cotton leaves collected at the indicated days.		from grou	ps of larvae	survived a	Adult longevity (in days) resulted from groups of larvae survived after feeding on treated	on treated
ides (ml)/ feddan Du en 300 % EC) 150 ozide 400 % EC) 200 in 100 % EC) 50	1st (0 = 1st day)	d at the indic	ated days.	e e e e e e e e e e e e e e e e e e e		D	
en 300 1% EC) 150 ozide 400 % EC) 200 in 100 % EC) 50	(an To)	$2^{nd} (7 - 8^{th} day)$	3th day)	3rd (14 - 15th day)	(5th day)	4th (21 - 22th dav)	22th dav)
en 300 % EC) 150 ozide 400 % EC) 200 in 100 % EC) 50	Duration %	Duration	%	Duration	%	Duration	%
en 300 1% EC) 150 ozide 400 % EC) 200 in 100 % EC) 50	Change		Change		Change		Change
% EC) 150 ozide 400 % EC) 200 in 100 % EC) 50 25 25	7.9 + 5.3	8.5	0	8.7	0	0.6	0
% EC) 200 in 100 % EC) 50	1	8.5	0	8.7	0	0.6	0
% EC) 200 in 100 % EC) 50 25	6.3 - 16.0	1		8.5	- 2.3	9.0	0
% EC) 50 25	1	i		8.6	-1.2	0.6	0
% EC) 50 25	7.8 + 4.0	8.5	0	8.7	0	9.0	0
25	7.8 + 4.0	8.5	0	8.7	0	0.6	0
	7.5	1	•	8.7	0	9.0	0
(10% SC) 12.5 7.5	7.5 0	8.5	0	8.7	0	9.0	0
Control - 7.5	7.5	8.5	ļ.	8.7		9.0	

stage caused prolongation in adult stage lately. Also are in agreement with data concerning feeding larval stage on sublethal concentrations of 1GR such as Diflubenzuron and Triflumuron (Attia and Ghattas, 1985) or/and Chlorfluazuron and Flufenoxuron (Hossain et al., 1996) where adult longevity was remarkably shortened.

10- Effect on adult fecundity:

Data in Tables (31-33) elucidate the latent indirect effects on fecundity of adults obtained when the 4th instar larvae of S. littoralis survived exposure and feeding on treated cotton leaves collected at different time intervals after application of insecticides under field conditions.

a- Laboratory reference strain:

Data recorded in Table (31) indicate the latent effects of the tested insecticide on adult fecundity when 4th instar larvae were fed and exposed to insecticide-treated cotton leaves collected at different time intervals after application. It was obvious that highly pronounced reduction in fecundity (100 %) was recorded at the 1st testing interval (0-1 day), in treatment of Pyriproxyfen at 300 ml/fed., whereas at the 2nd testing interval (7-8 day) 100% reduction in fecundity was recorded in both treatments of Ethoxazole (25 and 12.5 ml/fed.). However, at the 3rd (14-15 day) and 4th (21-22 day) testing intervals the maximum reduction in adult fecundity 91.7% and 80.0% was recorded for Ethoxazole at 25 ml/fed., respectively. However, it was of interest to note that due to no adults emerged at different testing intervals and treatments of Chromafenozide, the 3rd and 4th testing intervals of Azadirachtin treatments and the 1st testing interval of Ethoxazole, there was no deposited eggs entirely at these treatments.

Comparison based on number deposited eggs/ female revealed that the highest number (700 egg/female) was deposited in control treatment while the least deposited eggs/female was recorded in both

Adult fecundity of S. littoralis (Laboratory reference strain) for groups of larvae fed for 48h. on treated cotton leaves collected at different time intervals after applying insecticides under field conditions. Table (31).

	Rate	4	Average no.	of eggs/fem	ale resulted	Average no. of eggs/female resulted from group of larvae survived after feeding on treated cotton leaves collected at the indicated design	of larvae s	urvived after	1
Insecticides	(ml)/	$1^{57}(0 -$	$1^{st} (0 - 1^{st} day)$	2 nd (7 -	2"d (7 - 8th dav)	3rd (14 - 15th day)	15th day)	ou days.	. "
	feddan	Eggs no.	%	Eggs no.	%	Eggs no.	% ov	Fore 20 (4)	22 day)
			Change		Change	000	Change	E883 IIO.	%
Pyriproxyfen	300	0	- 100	39	- 94 4	100	000		Cuange
(10% EC)	150	25	06.4	10	1.1.7	100	- 03.3	1/3	- 65.0
		24	- 70.4	8/	- 88.9	200	- 66.7	350	- 30 0
Chromatenozide	400			1	ì				
(5% EC)	200	1		1					
Azadirachtin	100	000				1		1	
ACAUH ACHUIII	100	30	- 95.7	120	- 82.9	1			
(4.5% EC)	50	120	- 82.9	150	- 78.5		1		•
Ethoxazole	25			0	- 100	50	017		
(10% SC)	12.5	1		c	200	2	- 71.7	100	0.08 -
					- 100	100	- 83.3	150	- 70.0
Courrol		700		700		009		200	

treatments of Chromafenozide (zero), Azadirachtin at 100 ml/fed. (30 eggs) and Ethoxazole at 25 ml/fed. (50 eggs). Other treatments recorded mean number of eggs inbetween.

b- Early-season field strain (before spraying program):

As shown in Table (32), it was obvious that all tested insecticides performed similarly and resulted in reducing insect fecundity by averages of 52.9-83.6 %, 50.0-78.6 %, 50-75 % and 24.0-68.0% for adults resulted from groups of 1st, 2nd, 3rd and 4th testing intervals, respectively.

Comparison on the basis of mean eggs/female revealed that Chromafenozide resulted in remarkably the least number of deposited eggs/female. Considerable reduction in rate of eggs/female was recorded between other treatments including control where means eggs/female were lower than control treatment. The different insecticides could be arranged descendingly in this respect as follow: Pyriproxyfen, Ethoxazole and Azadirachtin, respectively.

c- Late-season field strain (after spraying program):

Data in Table (33) demonstrate that Chromafenozide at 400 ml/fed. and Ethoxazole at 25 ml/fed. resulted complete reduction (100 %) in deposited eggs at the 1st testing interval. The reduction in fecundity recorded averages of 78.6-100 %, 75-87.1 %, 60-93 % and 24-50 % for adults resulted from larval groups of 1st, 2nd, 3rd and 4th testing intervals. Comparison based on mean of eggs/females revealed remarkable reduction in fecundity between different treatment (40-380 eggs/female) and control treatment (500-700 eggs/female).

11- Effect on eggs viability:

Data in Tables (34-36) demonstrate the latent effect of different treatments on viability of eggs deposited by adults resulted after exposure and feeding of groups of 4th instar larvae to insecticide-treated cotton leaves collected at different testing intervals.

Table (32). Adult fecundity of S. littoralis early-season field strain (before spraying program) for groups of larvae fed for 48h. on treated cotton leaves collected at different time intervals after applying insecticides under field conditions.

Incontinidos	Rate	A	verage no.	of eggs/fem	ale resulted otton leaves	Average no. of eggs/female resulted from group of larvae survived after feeding on treated cotton leaves collected at the indicated days	of larvae s	urvived afte	J.
mscencines	(ml)/	$1^{st} (0 - 1^{st} day)$	1st day)	2nd (7 -	2"d (7 - 8th day)	3rd (14 - 15th dav)	15th dav)	4th (21 -	4th (21 - 22th day)
	feddan	Eggs no.	% Change	Eggs no.	% Change	Eggs no.	Change	Eggs no.	%
Pyriproxyfen	300	120	- 82.9	150	- 78.6	150	- 75.0	160	- 68 0
(10% EC)	150	135	- 80.7	250	- 64.3	270	- 550	310	38.0
Chromafenozide	400	ř		1					0.00
(5% EC)	200	115	- 83.6						
Azadirachtin	100	280	- 60.0	300	-57.1				
(4.5% EC)	50	330	- 52.9	350	- 50.0			380	0.80
Ethoxazole	25	185	- 73.6	200	- 71.4	230	-61.7	250	24.0
(10% SC)	12.5	255	- 63.6	270	- 61.4	300	- 50.0	320	36.0
Control	ı	700	ı	200	ī	009		500	

Adult fecundity of S. littoralis late-season field strain (after spraying program) for groups of larvae fed for 48h. on treated cotton leaves collected at different time intervals after applying insecticides under field conditions. Table (33).

		Average n	o. of eggs/fe	Average no. of eggs/female resulted from group of larvae survived after	ed from gro	up of larvae	survived a	fter	
	Rate	feeding on	treated cot	feeding on treated cotton leaves collected at the indicated days.	ollected at t	he indicated	days.		-
Insecticides	(m)/	$1^{37}(0 -$	$1^{37}(0-1^{37} day)$	$2^{nd} (7 - 8^{th} day)$	8''' day)	3^{rd} (14 - 15 th day)	15th day)	4" (21 - 22" day)	22''' day)
San Andrews	feddan	Fage no	%	Eggs no.	%	Eggs no.	%	Eggs no.	%
		- 555 m	Change	56	Change)	Change		Change
Designation	300	09	- 91.4	85	- 87.9	210	- 65.0	280	- 44.0
1 yriproxyren	150		,	06	- 87.1	240	- 60.0	310	- 38.0
Chromofonoride	400	0	- 100		•	40	- 93.3	250	- 50.0
CIII OIIII AI OIII OIII OIII OII OII OII	200					75	- 87.5	270	- 46.0
Andinohtin	100	100	- 85.7	130	- 81.4	150	- 75.0	320	- 36.0
AZAUH ACHUM	205	150	- 78.6	175	- 75.0	200	- 66.7	380	- 24.0
Ethoverole	25	0	- 100		1	100	- 83.3	300	40.0
(10% SC)	12.5	70	- 90.0	100	- 85.7	130	- 78.3	350	- 30.0
Control		700	a t	700	•	009		200	,

a- Laboratory reference strain:

Data in Table (34) revealed that the eggs deposited by adult females produced from different groups of susceptible strain larvae were susceptible to the latent effect of the tested insecticides. The inhibition in egg hatch reached 100 %, 100 %, 99-100 % and 98.6-100 %. In 1st (0-1 day), 2nd (7-8 day), 3rd (14-15 day) and 4th (21-27 day) testing intervals, respectively. However, in Chromafenozide no eggs were deposited where no adults resulted from the larval treatments in all testing intervals. Similarly, no deposited eggs was recorded in Ethoxazole's 1st and 2nd testing intervals and in the 3rd and 4th testing intervals of Azadirachtin treatments.

b- Early-season field strain (before spraying program):

As shown in Table (35) the latent effects of the tested compounds on eggs deposited laterly by adults produced from larval treatments of early-season field strain was relatively less susceptible than that for the laboratory reference strain. The recorded inhibition in eggs hatch reached 90.3-100 %, 86-98.6 %, 85-96 % and 75.7-95.2% at the 1st (0-1 day), 2nd (7-8 day), 3rd (14-15 day) and 4th (21-22 day), respectively. Comparison between different testing intervals revealed that the highest latent effect on eggs hatch inhibition was recorded in the 1st (0-1 day) testing interval and in general the maximum effect was recorded by Pyriproxyfen treatments (99.9-100 %) and was followed closely by Ethoxazole treatments (95.2-98.9 % and laterly came Azadirachtin treatments (90.3-95 %). However, it was worthy mentioning that no data was recorded in Chromafenozide treatments where no adult emerged from larval treatments.

c- Late-season field strain(after spraying program):

Data in Table (36) indicate in general that the late-season field strain performed slightly different than either laboratory or/and earlyseason field strains where it tolerate more the insecticidal treatments which in turn reflex on the latent effects. It was obvious that the latent

Table (34). Eggs viability deposited by females of S. littoralis (Laboratory strain) resulted from groups of larvae fed for 48h, on treated cotton leaves collected at different time intervals after applying insecticides under field conditions.

	Rate	% Sterility	% Sterility in eggs deposited by females resulted from groups of larvae	nales resulted from grou	tps of larvae
Insacticides	(m)/	survived a	survived after feeding on treated leaves collected at indicated days.	leaves collected at indic	ated days.
Santanas en la companya de la compan	feddan	$1^{st} (0 - 1^{st} day)$	$2^{nd} (7 - 8^{th} \operatorname{day})$	3^{rd} (14 - 15 th day)	4^{th} (21 - 22 th day)
Deminoscufon	300		100	100	100
ryriproxyten (10% FC)	150	100	100	100	100
Chromafenozide	400	•		•	,
(Fox FC)	200		nψ		1.
(3/0/20)	100	100	100	•	
Azadirachuli (4 5% FC)	205	100	100	1	•
(4.3.70 EV)	25			100	100
Ethoxazore (10% SC)	12.5			0.66	98.6
Control	-	0	0	0	0

Table (35). Eggs viability deposited by females of S. littoralis early-season field strain (before spraying program) resulted from groups of larvae fed for 48 h. on treated cotton leaves collected at different time intervals after applying insecticides under field conditions.

Trooping Town	Rate	% Sterility	in eggs deposited by fe	% Sterility in eggs deposited by females resulted from groups of larvae	ups of larvae
rusecucides	/(m)	survived	after feeding on treated	survived after feeding on treated leaves collected at indicated days	ated dave
	feddan	$1^{57} (0 - 1^{57} day)$	2"d (7 - 8th day)	3rd (11 15th 12)	oth cas anth
Pyriproxyfen	300	100	(6	(kgn cr - +r) c	4 (21 - 22 day)
The fact of the same of the sa		100	98.6	95.0	906
(10% EC)	150	6.66	956	200	0.00
Chromafenozide	400			30.7	85.4
		1			,
(5% EC)	200	100			
Azadirachtin	100	0.30			1
	TOO	93.0	90.0	1	-
(4.5% EC)	50	90.3	86.0		
Ethoxazole	25	0 80	000	# A1	13.7
		78.7	98.0	0.96	95.2
(10% SC)	12.5	95.2	0.06	850	000
Control		0	C		0.00
			0	0	

resulted from groups of larvae fed for 48h. on treated cotton leaves collected at different time Table (36). Eggs viability deposited by females of S. littoralis late-season field strain (after spraying program) intervals after applying insecticides under field conditions.

	,	0/ Ctomility	in ages denosited by fer	2/ Starility in same denosited by females resulted from groups of larvae	ns of larvae
	Kate (m)	survived	after feeding on treated	survived after feeding on treated leaves collected at indicated days.	ated days.
Insecticides	foddon f	1 st (0 - 1 st dav)	$2^{nd} (7 - 8^{th} dav)$	$3^{rd} (14 - 15^{th} \text{ day})$	4^{th} (21 - 22 th day)
n.	300	100	90.5	84.7	80.3
ryriproxylen	150		80.0	75.0	75.4
(10% EC)	400	ΑĒ		90.06	90.0
Curomatenozide	200		1	80.0	80.7
(3.70 EC)	100	80	75.3	70.0	70.0
Azadirachtin	20	71.3	65.7	0.09	60.5
Tethoverole	25		1	85.0	75.0
(10% SC)		06	85.0	70.7	70.0
Control	*	0	0	0	0

effect on eggs hatch inhibition reached 71.3-100 %, 65.7-90.5 %, 60-90 % and 60.5-90 % in the 1st (0-1 day), 2nd (7-8 day), 3rd (14-15 day) and 4th (21-22 day) testing intervals, respectively. No data was recorded only in the 1st and 2nd testing intervals of both treatments of Chromafenozide and the higher rate (25 ml/fed) of Ethoxazole which confirmed the relatively lower latent effect on the more tolerant late-season field strain. Data of eggs hatch inhibition in the 3rd and 4th testing intervals indicate that Chromafenozide (80-90 % and 80.7-90 %) was the first in this respect and was followed by the Pyriproxyfen (75-84.7 % and 75.4-80.3 %), Ethoxazole (70.7-85 % and 70-75 %) and Azadirachtin (60-70 % and 60.5-70 %), respectively.

In previous studies Schmidt et al. (1997) found that at concentrations of 10, 15 and 25 ppm Melia extract (Azadirachta indica) caused no adult emerged from pupae, fecundity and fertility were drastically reduced. Likewise, Sammour and Abdalla (1989) found that treatment of last instar larvae of Heliothis armigera with sublethal doses of Teflubenzuron caused considerable reduction in fecundity and fertility of the emerged adults with sterilizing effects amounted to ca. 91 %. In this respect. Gelbic and Matiola (1984) found the Juvenile hormone analogue (JHA) caused reduction in rate of eggs hatch and demonstrating that this performance could be due to numerous disorders in embryonic development. Recently, Aldebis et al. (1994) indicated that feeding on Fenoxycarb (JHA)-treated leaves resulted in alteration in the development of germinal cells ending in either malformed spermatozoa or eupyrene bundles.

4.3- The effect of non-traditional pesticides on feeding activity and nutritional indices:

4.3.1- Food consumption (C.W.):

On the basis of the overall mean weight of food consumed by early-season 4th instar larvae of *S. littoralis*, data in Table (37) indicate that the feeding ability of the larvae was significantly affected by insecticidal treatments, except for pyriproxyfen at the high rate, resulting in great and significant decrease in overall mean weight of food consumed. The overall weight of consumed food reached 144.3±23.1 mg by pyriproxyfen (150 ml/200 L), 107.5±15.8 and 110.5±19.2 mg by Chromafenozide (at 400 and 200 ml/200 L, respectively), 116.87±12.4 and 126.62±12.6 mg by Azadirachtin (at 100 and 50 ml/200 L, respectively) and 124.64±11.5 and 134.34±12.4 mg by Ethoxazole (at 25 and 12.5 ml/200 L, respectively) versus 154.16±1.62 mg for control larvae. However, the reduction in overall mean consumed food was in general positively correlated in most treatments with the concentration level, regardless of the tested insecticide.

Regarding the change in mean consumed food relative to the control larvae in details during different testing intervals, it was of interest to note that pyriproxyfen at 300 and 150 ml/200 L resulted in increase of +38.62 and +2.25 % in mean food consumed per larvae per day at the 1st (initial) testing interval whereas other treatments of Chromafenozide at 400 and 200 ml/200 L, Azadirachtin at 100 and 50 ml/200 L and Ethoxazole at 25 and 12.5 ml/200 L resulted in remarkable percent decrease of -41.34, -42.12; -36.58, -29.49; and -30.6, -24.5 % for the prementioned treatments, respectively. Also, it was obvious that the percent decrease in mean food consumed was gradually decline with time elapse where the least decrease recorded at the 4th testing intervals (21-22 day posttreatment) where the residues of sprayed insecticide was at its least level on treated leaves. However, the maximum reduction in mean food consumed (-24.49 and -26.32 %) was recorded by Pyriproxyfen 10 % EC at 300 and 150 ml/200 L whereas the least mean reduction (-11.49 and -3.37 %) was recorded

Table (37): Mean daily weight of consumed food (C.W.) for groups of early season S. littoralis field strain larvae fed for 48 h on treated leaves, collected at indicated day posttreatment, followed by 5 days feeding on untreated leaves.

	Rate/		Mean	daily we	ight of co	Mean daily weight of consumed food (mg/larva)	/gm) poo	larva)			
	, , , ,			at the	indicated	at the indicated testing intervals	ıtervals				-
Insecticides	feddan	1 <i>st</i> in	1st interval	2nd in	2nd interval	3rdin	3rdinterval	4th in	4th interval		
	(mi.)	0-1	0-1 day	7-8	7-8 day	14-1;	14-15 day	21-2	21-22 day	Mean ± S.D.	
		/gm	%	/gm	%	/gm	%	mø/	%		_
		larva	change	larva	change	larva	change	larva	change		-
Pyriproxyfen	300	216.93	+38.62	167.87	+9.74	166.05	+7.80	+7.80 115.65	24.40		-
(10 % EC)	150	160.01	+2.25	162.96		141 42	-8 10	117.05	26.30	166.63±41.36 a	-
Chromafenozide	400	91 79	-41 34	101 20	\top	1 00	0.10	117.03	-20.32	144.33±23.05 abc	-
(Da /0 4)		7	11.01	101.30	-22.72	107.99	-29.89	129.11	-15.71	107.57±15.83 d	
(3 % EC)	700	90.57	-42.12	105.62	-30.95	109.19	-29.11	136.63	-10 79	110 50±10 20 3	_
Azadirachtin	100	99.25	-36.58	119.05	-22 17	120 54	-217	130 63	1000	D 07'61±00'011	_
(4.5 % EC)	50	110 33	20 40	125 20			17.7	170.07	-10.03	116.87±12.47 cd	
Ethora		00.011	-47.47	02.621	-18.15	130.35	-15.37 140.61	140.61	-8.20	126.62±12.61 bcd	
Ethoxazole	72	108.60	-30.60	120.05	-21.52	125.33	-18.64	135.57	-11 49	124 64+11 50 1-4	_
(10 % SC)	12.5	118.95	-24.50	130.20	-14.88	140 81	-8 58	147 30	\neg	124.04±11.30 bcd	
Control		156.49		152 96		154 03		150 17		134.34±12.47 bcd	
I S D s s z .				2 2 1		101.00		173.17		154.16±1.62 ab	
.:0.0.0.										28.47	

Mean followed by the same letter are not significantly different (P = 0.05) according to Duncan's (1955) multiple range test. for Ethoxazole 10 % SC at 25 and 12.5 ml/200 L during the 4th testing interval (21-22 day posttreatment).

In respect to the late season *S. littoralis* strain, data presented in Table (38) revealed slight and insignificant decrease in overall mean weight of food consumed when different treatments were compared with control treatment.

Regarding the detailed change in mean consumed food relative to control during different testing intervals, it was obvious that the highest decrease at the 1st (initial) testing interval (-35.09 and -33.28 %) was recorded for Azadirachtin 45 % EC at 100 and 50 ml/200 L, respectively; while the least (-20.33 and -16.75 %) was recorded for Chromafenozide 5 % EC at 400 and 200 ml/200 L, respectively.

Also, it was obvious that the percent decrease in mean food consumed was gradually declined with time elapse to reach its least value at the 3rd testing interval at 14-15 day posttreatment, then a sudden increase in mean food consumed was exhibited at the 4th testing interval at 21-22 day posttreatment. However, the highest mean food consumed (+34.16 and +33.68 %) was recorded by Azadirachtin treatments (100 and 50 ml/200 L), while the least increase of +17.45 and +5.09 % was recorded for Pyriproxyfen 10 % EC at 300 and 150 ml/200 L, respectively.

Such performance of insignificant variations in response of larvae of late season strain between treatments and control larvae could be explained in view of the larvae being more tolerant to insecticidal treatments which reflect in their low response to insecticides particularly at the last (4th) testing interval where sublethal low residues of insecticides at such time on leaves exhibited stimulatory effect as manifested by increase in mean food consumed, reached its maximum (+34.16 and +33.68 %) by both treatments of Azadirachtin.

As for the laboratory strain (Table 39), its response to different treatments was almostly similar to the early-season strain in respect to mean food consumed.

Table (38): Mean daily weight of consumed food (C.W.) for groups of late season S. littoralis field strain larvae fed for 48 h on treated leaves, collected at indicated day posttreatment, followed by 5 days feeding on untreated leaves.

	Rate/		Mear	daily we	eight of co	Mean daily weight of consumed food (mg/larva) at the indicated testing intervals	food (mg/	larva)		
Insecticides	feddan (ml.)	1st in	1st interval	2nd in	2nd interval	3rdin	3rdinterval	4th in	4th interval	
		T-O	uay	2	/-8 day	14-1	14-15 day	21-2	21-22 day	Mean ≠ S.D.
		mg/	%	/gm	%	/gm	%	/au	%	
		larva	change	larva	change	larva	change	larva	change	
Pyriproxyfen	300	117.87	-24.18	118.32	-22.65	129 11	-16 18	170 01	117 AS	
(10 % EC)	150	116.26	-25.71	119.36	1	133 43	12 27	16007	C+./1-	130.32±29.57 a
Chromafenozide	400	124 67	2000	0, 10,		200.40	10.01-	16001	42.09	132.51±20.40 a
(F 1/ E)		174.01	-20.33	137.43	-10.15	145.98	-5.23	173.92	+13.55	145.5±20.87 a
(3 % EC)	200	130.28	-16.75	135.55	-11.38	146.65	-4.79	197 23	+78 77	150 42120 64
Azadirachtin	100	101.57	-35.09	127.37	-16 73	131 10	1	205 40	17.07	132.43±30.04 a
(4.5 % EC)	50	104 14	-33 28	136 23		11/00	110	447	+34.16	141.41±44.0 a
Ethoverole	20		07:70	120.32	-10.00	140.88	-4.64	204.76	+33.68	148.09±41.87 a
Trinovazone	C7	111.22	-28.93	137.37	-10.19	146.67	-4.78	175.21	+14.38	142 62+26 71 0
(10 % SC)	12.5	111.12	-28.93	142.17	-3.79	146.69	-4.77	181 47	+18.47	145 26 20 00
Control		156.49		152.96		154 03		152 17	10.1	143.30±28.80 a
L.S.D.0.05:						20:10		173.11		154.16±1.62 a
										43.11

Mean followed by the same letter are not significantly different (P = 0.05) according to Duncan's (1955) multiple

h on treated leaves, collected at indicated day posttreatment, followed by 5 days feeding on untreated Table (39): Mean daily weight of consumed food (C.W.) for groups of S. littoralis laboratory strain larvae fed for 48 leaves.

			Mean	daily wei	Mean daily weight of consumed food (mg/larva)	sumed fo	od (mg/l:	arva)		
	Rate/			at the i	at the indicated testing intervals	testing in	tervals			
	feddan	1st interval	erval	2nd interval	terval	3rdinterval	erval	4th interval	erval	Mean + S.D.
Insecticides	(m)	0-1 day	dav	7-8 day	day	14-15 day	day	21-22 day	day	MICAIL - D.D.
		/øm	%	/gm	%	/gm	%	/gm	%	
		larva	change	larva	change	larva	change	larva	change	
Designation	300	228 49	+46.01	181.97	+18.97	16.67	+8.21	+8.21 137.25	-10.39	178.60±38.09 a
ryrthroxyren	150	209.03		164.78	+7.73	145.50	-5.53	141.36	-7.71	165.28±31.19 a
Cl. mofonorido	400	81 78		89.38	-41.56	94.74	-38.49	105.23	-31.29	92.78±9.86 b
Chromatenoziue	200	85 14	-45 59	102.74	-32.83	102.74	-33.29	105.23	-31.29	98.80±9.20 b
(3 % EC)	100	98 94	-36.78	-36.78 113.23	_	-25.97 113.26	-26.47	127.22	-16.94	113.16±11.55 b
AZadirachum	9	105.69	-32.46	115.18	-	116.49	-24.37	136.56	-10.84	118.46±12.98 b
T-thorographo	25	104 16		119.50	-33.46	119.87	-22.11	132.55	-13.46	119.05±11.62 b
C10 % CC)	12.5	104.65		127.20	-16.84	127.20	-17.42	135.75	-11.33	123.54±13.63 b
Control	'	156.49		152.96	1	154.03	ı	153.17	ı	154.16±1.62 a
I S Door.										27.49

Mean followed by the same letter are not significantly different (P = 0.05) according to Duncan's (1955) multiple range test.

4.2.2- Consumption index (C.I.):

Data indicating consumption index (C.I.) values for the three tested strains (early-season, late-season and laboratory strains) are shown in Tables (40, 41 & 42). The data obviously indicate that C.I. was found to be associated with the amount of food consumed and though showed in general the same trend.

Data in Table (40) show that the overall mean of consumption index (C.I.) of the early-season strain larvae was significantly decreased below that of control in all treatments, except for those of Pyriproxyfen where C.I. recorded was insignificantly similar to control. Both treatments of Chromafenozide (400 and 200 ml/200 L) resulted in significantly the least C.I. values.

Regarding the detailed data at different testing intervals, a remarkable decline in C.I. values was recorded with time elapse where the longer the period after treatment the higher the C.I. value although it was still less than those recorded for control.

In respect to the late-season *S. littoralis* larvae, data shown in Table (41) revealed slight and insignificant decrease in overall mean C.I. for larvae in all treatments and control. However, regarding the change in detailed data during different testing intervals, it was obvious that the highest reduction in C.I. values (-48.48 and -45.45 %) at the 1st (initial) testing interval was recorded for Ethoxazole at 25 and 12.5 ml/200 L, respectively, while the least reduction in C.I. values (-21.21 and -21.21 %) was recorded for Chromafenozide at rate of 400 and 200 ml/200 L, respectively.

As for the overall mean of C.I. values for laboratory strain larvae, data in Table (42) revealed a trend of reduction similar to that recorded for larvae of early-season strain where Pyriproxyfen at the two tested rates exhibited almostly an overall mean similar and insignificant when compared with control. However, the other three insecticides Chromafenozide, Azadirachtin and Ethoxazole exhibited

Table (40): Consumption index (C.I.) for groups of early season S. littoralis field strain larvae fed for 48 h on treated leaves, collected at indicated day posttreatment, followed by 5 days feeding on untreated leaves.

			Mean (I. value	Mean C.I. values at the indicated testing intervals	dicated t	esting int	ervals		
Insecticides	Rate/	1 <i>st</i> int	1st interval	2nd interv	2nd interval	3rdinterval	erval	4th interval 21-22 day	erval	Mean ± S.D.
	(ml.)	C.I.	, %	C.I.	%	C.I.	%	C.I.	% change	
		t	change	0.37	+5 71	0.36	+2 86	0.30	-9.09	0.37±0.70 a
Pyriproxyfen	300	74.0	247.47	0.36	+2 86	0.31	-11.43	0.30	-9.03	0.35±0.50 a
(10 % EC)	150	0.10	CV CV	0.23	-40.0	0.21	-40.0	0.23	-3.30	0.24±0.02 d
Chromatenozide	200	0.17	-36 36	0.21	-40.0	0.23	-34.29	0.28	-15.15	0.23±0.03 d
(5 % EC)	100	0.25	-24 24	0.28	-20.0	0.29	-17.14	0.29	-12.12	0.28±0.02 bc
Azadirachtin	3 5	0.26	-21.21	0.29	-17.14	0.29	-17.14	0.30	-9.09	0.28±0.02 bc
(4.5 % EC)	25	0.20	-21.21	0.29	-17.14	0.29	-17.14	0.29	-12.12	0.28±0.01 bc
Ethoxazole	12.5	0.26	-21.21	0.29	-17.14	0.30	-14.29	0.31	-6.06	0.29±0.02 b
Control	C:21 -	0.33		0.35	1	0.35	ī	0.33	ı	0.34±0.01 a
100										0.07
L.S.D.0.05:										1. 1.1

Mean followed by the same letter are not significantly different (P = 0.05) according to Duncan's (1955) multiple range test.

Table (41): Consumption index (C.I.) for groups of late season S. littoralis field strain larvae fed for 48 h on treated leaves, collected at indicated day posttreatment, followed by 5 days feeding on untreated leaves.

			Mean	C.I. valu	es at the i	ndicated	Mean C.I. values at the indicated testing intervals	tormole			
Insecticides	Rate/ feddan	1 <i>st</i> in 0-1	1st interval 0-1 day	2nd ir 7-8	2nd interval 7-8 day	3rdin 14-1	3rdinterval	4th ir	4th interval	Mean ± S.D.	
	(ml.)	C.I.	% change	C.I.	% change	C.I.	%	CI	% .		
Pyriproxyfen	300	0.22	-33.33	0.22	-37 14	000	o a oo		change		-
(10 % EC)	150	0.22	-33 33	0.00	41.70	0.23	-34.29	0.34	+30.30	0.25±0.06 a	T
Chromafanozida	400	1000	00.00	0.43	-24.79	0.25	-28.57	0.36	+9.09	0.26±0.06 a	_
Can omatchozine	400	0.76	-21.21	0.26	-25.71	0.26	-25.71	0.39	+18 18	0 20+0 06 9	T
(5 % EC)	200	0.26	-21.21	0.26	-25.71	0.27	-22.86	0.30	110 10	0.23±0.00 a	_
Azadirachtin	100	0.21	-36.36	0.26	-25.71	0.27	2000	00.0	10.10	0.29±0.08 a	
(4.5 % EC)	50	0.10	17.17	200	10.71	77.0	-27.00	0.39	+18.18	0.28±0.08 a	
Fthowardle	200	71.0	74.74-	07.70	1/.cz-	0.26	-25.71	0.38	+15.15	0.27±0.07 a	
Trinovazone	57	0.17	-48.48	0.26	-25.71	0.29	-17.14	0.34	+30 30	0.76+0.07	_
(10 % SC)	12.5	0.18	-45.45	0.26	-25.71	0.28	-200	0.35	76.06	0.2020.07.8	
Control	,	0.33		0.35		0.35	0.01	0.00	40.00	0.2/±0.07 a	
L.S.D.0.05:						0.00	•	0.33		0.34±0.01 a	
Mer. 6 m										0.09	

Mean followed by the same letter are not significantly different (P = 0.05) according to Duncan's (1955) multiple

Table (42): Consumption index (C.I.) for groups of S. littoralis laboratory strain larvae fed for 48 h on treated leaves, collected at indicated day posttreatment, followed by 5 days feeding on untreated leaves.

			Mean C	.I. value	Mean C.I. values at the indicated testing intervals	dicated 1	esting int	ervals		
•	Rate/	1st int	1st interval	2nd in	2nd interval	3rdinterval	erval	4th interval	terval	Mean + S.D.
Insecticides	feddan	0-1 day	day	7-8	7-8 day	14-15 day	day	21-22 day	day	The state of the s
	(ml.)	į	%	1	%	7	%	1.7	%	
		C.I.	change	C.I.	change	C.I.	change	ניזי	change	
Pvriproxyfen	300	0.55	+66.67	0.38	+8.57	0.38	+8.57	0.29	-12.12	0.40±0.11 a
(10 % EC)	150	0.45	+36.36	0.36	+2.86	0.31	-11.92	0.28	-15.15	0.35±0.07 a
Chromafenozide	400	0.15	-54.55	0.20	-42.86	0.20	-42.86	0.22	-33.33	0.19±0.03 b
(5 % EC)	200	0.17	-48.49	0.20	-42.86	0.21	-40.0	0.24	-27.27	0.20±0.03 b
Azadirachtin	100	0.19	-42.42	0.21	-40.0	0.22	-37.14	0.25	-24.24	0.22±0.02 b
(4.5 % EC)	50	0.21	-36.36	0.21	-40.0	0.22	-37.14	0.28	-15.15	0.23±0.03 b
Ethoxazole	25	0.21	-36.36	0.21	-40.0	0.23	-34.29	0.28	-15.15	0.23±0.03 b
(10 % SC)	12.5	0.25	-24.24	0.25	-28.57	0.26	-25.71	0.30	-9.09	0.26±0.02 b
Control		0.33		0.35	ī	0.35		0.33	•	0.34±0.01 a
L.S.D.0 05:										0.07

Mean followed by the same letter are not significantly different (P = 0.05) according to Duncan's (1955) multiple range test.

overall mean C.I. values insignificant between each other, while significantly lower than Pyriproxyfen and control. Also, the initial effect (at 1st testing interval) demonstrate the positive effect of Pyriproxyfen, manifested as highly pronounced increase in C.I. values (+66.67 and +36.36 %) for both rates tested, compared with the remarkable decrease in C.I. value (-54.55 and -48.49, -42.42 and -36.36, 36.36 and 24.24 %) for the high and low rates of each of Chromafenozide, Azadirachtin and Ethoxazole, respectively.

The present results (Tables 37-42), generally show an inhibitory action of the tested compounds on the food consumption whether its determination was estimated as amount in mg (C.W.) or % (C.I.). Such an observations are in agreement with earlier reports which indicated that insect growth disruptors interfere with feeding (Mulder and Giswijt, 1973; Ascher and Nemny, 1976; Abid et al., 1978). Moreover, previous reports have also confirmed that diflubenzuron acts on the peritrophic membrane by affecting its chitin-protein structure, hindering its role in protecting secreting cells from any damage (Clarke et al., 1977).

Also, our findings agree with results recorded for several insect species such as S. littoralis fed on diflubenzuron-treated castor leaves (Sundaramurthy, 1977 and Radwan et al., 1986), S. littoralis fed on fenarimol or nuarimol (fungicides with anti-ecdysone activity)-treated cotton leaves (Farag, 1991). Also, Schmidt et al. (1997) found that a methanol extract of Melia azedarach fruits reduced food consumption. Furthermore Salama and Ahmed (1997) found that methanol extract, Melia azedarach exhibited decrease in respiration quotient (OR) and destroyed epithelial cells, the peritrophic membrane and basement membrane of the midgut of the tested larvae with 100 ppm. Outside Lepidoptera, Ismail (1995) recorded various reductions of relative consumption rate (RCR), consumed food amounts (C.W.) and faeces after topical application of fenoxycarb (a juvenoid) onto newly moulted last instar nymphs of S. gregaria. Ghoneim (1994), also

applied topically various doses of Pyriproxyfen onto newly moulted last instar nymphs of *S. gregaria*, a dose level of 10 μ g/nymph gave an inhibitory effect of this juvenoid and 100 and 150 μ g/nymph gave a reverse effect on food ingestion and consumption.

4.3.3- Growth Rate (G.R.):

The effect on growth rate (G.R.) after feeding 4th instar larvae of the three tested strains (early-season strain, late-season strain and laboratory strain) on cotton leaves at specific intervals after spraying of the four tested pesticides is demonstrated in Tables (43, 44 & 45).

Data in Table (43) representing the response of early-season larvae after feeding on treated cotton leaves collected at 0-1 day (initial) posttreatment revealed remarkable decrease in larval growth by percentage ranged between -51.66 % and -32.0 % for Chromafenozide, Azadirachtin and Ethoxazole whereas Pyriproxyfen at both rates tested resulted in highly pronounced increase in G.R. values of +122.6 % and +41.33 % for 300 and 150 ml/200 L, respectively.

Comparison on the basis of statistical analysis of the overall mean during the whole experimental periods indicate a remarkable reduction in overall mean growth rate (G.R.) in all treatments except both rates of Pyriproxyfen where a considerable and significant increase (0.425±0.1 and 0.33±0.09) relative to control (0.3±0.05) was recorded. Also, it was obvious that the lowest G.R. value (0.189±0.04) was recorded for larvae fed leaves treated with Chromafenozide at 400 ml/200 L, while the highest G.R. value (0.425±0.1) was recorded when the 4th instar larvae were fed on leaves treated with Pyriproxyfen at 300 ml/200 L which was also significantly higher than other treatments including the control (0.309±0.05).

As for the late-season *S. littoralis* strain (Table, 44), it was obvious that all tested treatments during the first three testing intervals exhibited G.R. values remarkably lower than the control. The reduction

Table (43): Growth rate (G.R.) for groups of early season S. littoralis field strain larvae fed for 48 h on treated leaves, collected at indicated day posttreatment, followed by 5 days feeding on untreated leaves.

		Mean	Mean growth rate values (G.R.) at the indicated testing intervals	rate valu	es (G.R.)	at the inc	licated te	sting inte	srvals	
Insecticides	Rate/	1 <i>st</i> in 0-1	1st interval	2nd ir	2nd interval	3rdin	3rdinterval	4th in	4th interval	Mean ± S.D.
	(ml.)		%		%		%	77	%	
	K (G.K.	change	G.R.	change	G.R.	change	G.R.	change	
Pyriproxyfen	300	0.668	+122.6	0.411	+30.47	0.406	+30.96	0.216	-30.99	0.425±0.1 a
(10 % EC)	150	0.424	+41.33	0.397	+26.03	0.282	-9.03	0.219	-30.03	0.330±0.09 ab
Chromafenozide	400	0.145	-51.66	0.164	-47.93	0.202	-34.83	0.247	-21.08	0.189±0.04 c
(5 % EC)	200	0.149	-50.33	0.189	-40.00	0.205	-33.87	0.265	-15.33	0.202±0.05 bc
Azadirachtin	100	0.158	-47.33	0.206	-34.60	0.223	-28.06	0.263	-15.97	0.212±0.04 bc
(4.5 % EC)	50	0.177	-41.00	0.240	-23.80	0.262	-15.48	0.287	-8.31	0.241±0.04 bc
Ethoxazole	25	0.185	-38.33	0.230	-26.98	0.250	-19.35.	0.273	-12.78	0.234±0.03 bc
(10 % SC)	12.5	0.204	-32.00	0.255	-19.04	0.277	-10.64	0.297	-5.11	0.258±0.04 bc
Control	5	0.300	,	0.315	ı	0.310	1	0.313	•	0.309±0.05 b
L.S.D.0.05:										90.0

Mean followed by the same letter are not significantly different (P = 0.05) according to Duncan's (1955) multiple range test.

Table (44): Growth rate (G.R.) for groups of late season S. littoralis field strain larvae fed for 48 h on treated leaves, collected at indicated day posttreatment, followed by 5 days feeding on untreated leaves.

		Mean	Mean growth rate values (G.R.) at the indicated testing intervals	rate value	es (G.R.)	at the inc	licated te	sting inte	rvals	
Insecticides	Rate/ feddan	1 <i>st</i> int 0-1	1st interval 0-1 day	2 <i>nd</i> in 7-8	2nd interval 7-8 day	3rdinterval	srdinterval	4th interval 21-22 day	terval 2 day	Mean ± S.D.
	(ml.)	G.R.	% change	G.R.	% change	G.R.	% change	G.R.	% change	
Pyriproxyfen	300	0.201	-33.0	0.223	-28.98	0.254	-18.06	0.412	+31.63	0.272±0.09 a
(10 % EC)	150	0.198	-34.0	0.224	-28.66	0.256	-17.42	0.401	+28.11	0.269±0.09 a
Chromafenozide	400	0.228	-24.0	0.274	-12.73	0.294	-5.16	0.451	+44.09	0.311±0.09 a
(5 % EC)	200	0.236	-21.33	0.271	-13.69	0.293	-5.48	0.513	+63.89	0.328±0.01 a
Azadirachtin	100	0.183	-39.0	0.235	-25.15	0.261	-15.8	0.523	+67.09	0.301±0.02 a
(4.5 % EC)	50	0.179	-40.33	0.265	-15.6	0.287	-7.42	0.511	+63.26	0.310±0.01 a
Ethoxazole	25	0.188	-37.33	0.239	-23.88	0.265	-1452	0.451	+44.09	0.382±0.01 a
(10 % SC)	12.5	0.177	-41.0	0.242	-22.92	0.296	-4.52	0.477	+52.39	0.298±0.01 a
Control	1	0.300	1	0.314	1	0.310	ī	0.313		0.309±0.06 a
L.S.D.0.05:										80.0

Mean followed by the same letter are not significantly different (P = 0.05) according to Duncan's (1955) multiple range test.

percentages in G.R. values were higher at the 1st testing interval (ranged between -41.0 and -21.33 %) and declined gradually by time elapse to reach the minimum level of reduction in G.R. values (ranged between -18.06 and -4.52 %) at the 3rd (14-18 day) testing interval. In contrast, the G.R. values at the 4th testing interval (21-22 day), where the pesticides residues (or metabolites) on leaves were at its lowest level, stimulate the growth rate and resulted in considerable increase in G.R. values ranged between +67.09 and +28.11 %. However, comparison on the basis of the overall means during the 4 testing intervals revealed insignificant differences between all treatments including the control.

Data in Table (45) demonstrate the response of laboratory-strain larvae of *S. littoralis* concerning the G.R. when larave were fed and subjected to insecticides-treated leaves, collected at specific testing intervals.

Generally, all treatments, except Pyriproxyfen, resulted in remarkable reduction in G.R. values which was at its highest level initially (during the 1st testing interval) then decreased gradually with progress in time after spraying. The percent reduction in G.R. values ranged between -40.67 and -65.67 % at the 1st (0-1 day) testing interval, -21.38 and -60.51 % at the 2nd (7-8 day) testing interval, -19.35 and -58.93 % at the 3rd (14-15 day) testing interval, while ranged between -9.9 and -40.25 % at the 4th (21-22 day) testing interval. On contrary, the G.R. values increased remarkably in most testing intervals of Pyriproxyfen treatments by percent ranged between +141.0 % and +30.0 %.

Comparison based on the overall mean during the whole experimental period revealed that G.R. values in different treatments could be arranged statistically in three categories, the first higher than control (GR = 0.471-0.39) which includes only Pyriproxyfen both treatments, the second include control only (GR = 0.309), while the third having GR values lower than control (GR = 0.235-0.138) and include treatments of the other three pesticides.

Table (45): Growth rate (G.R.) for groups of S. littoralis laboratory strain larvae fed for 48 h on treated leaves, collected at indicated day posttreatment, followed by 5 days feeding on untreated leaves.

		Mear	growth	rate valu	Mean growth rate values (G.R.) at the indicated testing intervals	at the inc	licated te	sting inte	rvals	
Insecticides	Rate/ feddan	1 <i>st</i> in: 0-1	1st interval 0-1 day	ai <i>pu</i> Z 7-8	2nd interval 7-8 day	3rdinterval	3rdinterval	4th interval 21-22 day	tth interval 21-22 day	Mean ± S.D.
	(ml.)	G.R.	% change	G.R.	% change	G.R.	% change	G.R.	% change	
Pyriproxyfen	300	0.723	+141.0	0.478	+52.23	0.403	+30.0	0.279	-9.90	0.471±0.18 a
(10 % EC)	150	0.576	+92.0	0.419	+33.44	0.292	-5.81	0.274	-12.46	0.390±0.13 ab
Chromafenozide	400	0.103	-65.67	0.124	-60.51	0.139	-55.16	0.187	-40.25	0.138±0.03 d
(5 % EC)	200	0.115	-61.67	0.169	-46.18	0.183	-58.93	0.215	-31.31	0.170±0.04 d
Azadirachtin	100	0.140	-53.33	0.170	-45.86	0.171	-44.84	0.203	-35.46	0.171±0.02 d
(4.5 % EC)	50	0.169	-43.67	0.172	-45.22	0.187	-39.68	0.268	-14.38	0.199±0.04 cd
Ethoxazole	25	0.168	-44.0	0.229	-27.07	0.230	-25.81.	0.257	-17.89	0.221±0.03 cd
(10 % SC)	12.5	0.178	-40.67	0.247	-21.38	0.250	-19.35	0.267	-14.70	0.235±0.03 cd
Control	1	0.300	ı	0.314	1	0.310		0.313		0.309±0.05 bc
L.S.D. _{0.05} :										0.04

Mean followed by the same letter are not significantly different (P = 0.05) according to Duncan's (1955) multiple range test.

It was rationally accepted that the larval growth had been hindered by the inhibitory action of these compounds on feeding and the metabolic abilities of *S. littoralis* larvae. In this respect, **Radwan** et al. (1986) found that larvae of *S. littoralis* fed on diflubenzuron and triflumuron-treated leaves have shown a proportional relationship between food consumed and values of both consumption index and growth rate throughout the study course and that both treatments showed considerable decrease in growth rate. Likewise, **Woodring** et al. (1978) and **Sundramurthy** (1977) indicated that the amount of growth reduction in *S. littoralis* was proportional in general to reduced food consumption. Similar results had been found by other authors after applying different IGRs against various insect species such as *Agrotis* ipsilon (Reese and Beck, 1976a, b & c), *Manduca sexta* (Dahlman, 1977), and *S. gregaria* (Ghoneim, 1994).

However, several authors explained the depressed G.R. or depleted R.W.G. by the decreasing amounts of food consumed. This interpretation may be also suggested in the present study because the data of Tables (37, 38 & 39) clearly show detrimentally reduced food consumed (C.W.).

4.3.4- Approximate Digestibility (A.D.):

Data in Tales (46, 47 & 48) reveals the approximate digestibility (A.D.) of food in larvae of *S. littoralis* representing the three different tested strains. As shown in Table (46), it was obvious that the overall mean of A.D. in control larvae of early-season strain was 87.65±1.48 %. Feeding the 4th instar larvae on leaves treated with different pesticides resulted in either decrease or increase in A.D. depending on the pesticide or/and the concentration used. Feeding the larvae on Chromafenozide-treated leaves increased significantly the percentage of digestibility, recording 92.3 % and 92.07 % at both tested rates of 400 and 200 ml/200 L, respectively. Other treatments except Pyriproxyfen (300 ml/200 L) came significantly next and no significant

Table (46): Approximate digestibility (A.D.) for groups of early season S. littoralis field strain larvae fed for 48 h on treated leaves, collected at indicated day posttreatment, followed by 5 days feeding on untreated leaves.

		Rate/	Appr 48	oximate by on tre	digestibil ated leav	Approximate digestibility (A.D. %) for 4th instar larvae fed for 48 h on treated leaves collected at the indicated intervals	%) for 4i	th instar indicated	larvae fec interval	l for S	
		feddan	1st interval	erval	2nd in	2nd interval	3rdinterval	erval	4th interva	terval	Moon + C D
	Insecticides	(ml.)	0-1 day	day	7-8 day	day	14-15 day	day	21-22 day	day	Mean + 5.D.
			A.D.	%	A.D.	%	A.D.	%	A.D.	%	
			%	change	%	change	%	change	%	change	
	Pvrinroxvfen	300	82.48	-4.56	84.37	-6.03	84.72	-3.13	88.61	+1.89	85.05±2.57 c
5	(10 % EC)	150	25.23	-1.38	87.44	-3.72	90.81	+3.84	90.84	+4.46	88.33±2.92 bc
1	Chromafenozide	400	97.15	+12.42	91.62	+2.05	90.48	+3.47	89.93	+3.42	92.30±3.31 a
, 	(5 % EC)	200	95.26	+10.23	92.85	+3.42	90.19	+3.13	89.97	+3.46	92.07±2.50 a
_	Azadirachtin	100	93.12	+7.76	90.43	+0.72	89.32	+2.64	89.33	+2.73	90.55±1.79 ab
4	(4.5 % EC)	50	91.36	+5.72	90.50	+0.80	89.75	+2.63	89.36	+2.76	90.24±0.88 ab
	Ethoxazole	25	90.36	-4.56	90.12	+0.38	89.29	+2.10	87.28	+0.37	89.26±1.40 ab
	(10 % SC)	12.5	91.45	+5.82	90.59	+0.80	89.89	+0.50	87.32	+0.41	89.31±2.02 ab
	Control		86.42	ť	89.78	1	87.45	•	96.98		87.65±1.48 bc.
-	L.S.D.0.05:										3.23

Mean followed by the same letter are not significantly different (P = 0.05) according to Duncan's (1955) multiple range test.

Table (47): Approximate digestibility (A.D.) for groups of late season S. littoralis field strain larvae fed for 48 h on treated leaves, collected at indicated day posttreatment, followed by 5 days feeding on untreated leaves.

	Rate/	App 4	roximate 8 h on tre	digestibi	Approximate digestibility (A.D. %) for 4th instar larvae fed for 48 h on treated leaves collected at the indicated integral.	%) for 4	th instar	larvae fe	d for		
-	feddan	1st in	1st interval	2nd in	2nd interval	3rdin	3rdinterval	Ath in	Ath intomol		
Insecticides	(ml.)	0-1	0-1 day	7-8	7-8 day	14-1	14-15 day	21-2	21-22 day	Mean ± S.D.	
		A.D.	%	A.D.	%	A.D.	%	A.D.	%		
		%	change	%	change	%	change	%	change		_
Pyriproxyfen	300	89.42	+3.47	89.91	+0.15	88.46	+1.16	81.37	-6 43	87 29+3 99 3	T
(10 % EC)	150	89.70	+3.79	90.90	+1.25	93.07	+6.43	84.98	-43 68	89 60+3 42 a	T
Chromafenozide	400	96.73	+11.93	92.14	+2.63	87.79	+0.39	86.37	0.68	90 76+4 68 3	T
(5 % EC)	200	91.43	+5.79	0.01	+0.26	87.14	+0.36	86.24	0.83	88 70+3 43 5	T
Azadirachtin	100	92.76	+7.34	91.95	+2.42	95 53	+9 24	86.55	0.02	01 70+3 76 3	_
(4.5 % EC)	50	91.60	+5.99	99.06	+0.98	90.41	+3.39	86.58	-0 44	89 81+2 21 9	_
Ethoxazole	25	90.92	+5.22	92.11	+2.6	87.61	+0.18	86.05	-1 14	89 17+7 87 9	_
(10 % SC)	12.5	91.31	+5.66	90.35	+0.64	87.88	+0.49	86.55	-0.47	89 07+7 19 3	_
Control		86.42	1	86.78	r	87.45		86.96		87 65+1 48 3	_
L.S.D.0.05:										4.57	_

Mean followed by the same letter are not significantly different (P = 0.05) according to Duncan's (1955) multiple range test.

S. littoralis laboratoty strain larvae fed for 48 h on treated leaves, collected at indicated day posttreatment, followed by 5 days feeding on untreated leaves. Table (48): Approximate digestibility (A.D.) for groups of

	Rate/	App.	Approximate digestibility (A.D. %) for 4th instar larvae fed for 48 h on treated leaves collected at the indicated intervals	digestibil ated leav	ity (A.D.	%) for 4.	th instar	larvae fec interval	d for Is		
9	feddan	1st int	1st interval	2nd in	2nd interval	3rdinterval	erval	4th interval	terval	Moon + C D	
Insecticides	(ml.)	0-1 day	day	7-8 day	day	14-15 day	5 day	21-22 day	day	Mean = 5.D.	
		A.D.	%	A.D.	%	A.D.	%	A.D.	%		_
		%	change	%	change	%	change	%	change		T
Pvriproxyfen	300	80.59	-6.75	82.58	-8.02	83.88	-4.08	87.85	+1.02	83.72±3.06 c	
(10 % EC)	150	83.96	-2.85	85.81	-4.42	89.29	+2.10	91.21	+4.89	87.57±3.28 bc	1
Chromafenozide	400	76.96	+12.12	92.99	+3.52	87.74	+0.33	87.71	+0.86	91.34±4.49 ab	
(5 % EC)	200	96.49	+11.65	93.05	+3.64	93.43	+6.84	91.89	+5.67	93.71±1.96 a	- 1
Azadirachtin	100	96.49	+11.58	93.01	+3.59	93.02	+6.37	86.38	+2.78	92.97±2.90 a	T
(4.5 % EC)	50	94.84	+9.74	91.95	+2.42	91.09	+4.16	90.17	+3.69	92.0±2.02 ab	T
Ethoxazole	25	93.78	+8.52	88.68	+0.11	99.68	+2.53	88.46	+1.72	90.43±2.28 ab	
(10 % SC)	12.5	98.20	+13.63	89.14	+0.71	89.21	+2.01	89.15	+2.51	91.42±4.52 ab	T
Control		86.42		86.78		87.45	•	96.98	•	87.65±1.48 bc	
L.S.D.o.o.s :										4.45	\neg
											1

Mean followed by the same letter are not significantly different (P = 0.05) according to Duncan's (1955) multiple range test.

differences were noticed between them and the control. Pyriproxyfen at rate of 300 ml/200 L recorded an approximate digestibility (85.05 %) insignificantly lower than control (87.65 %).

As for late-season strain larvae (Table, 47) detailed data indicate slight increase in A.D. values ranged between +3.47 % and +11.93; +0.15 % and +2.63 %; +0.18 % and +9.24 % at 1st (0-1 day), 2nd (7-8 day) and 3rd (14-15 day) testing intervals, respectively, whereas remarkable decrease in A.D. % values ranged between -0.44 % and -43.68 % was recorded at the 4th (21-22 day) testing interval. Generally, the variations between the overall mean A.D. values during the whole experimental period were insignificant and the A.D. values ranged between 87.29 % and 91.70 %.

Data in Table (48) demonstrate the estimated A.D. % values for larvae of the laboratory strain when fed and subjected to different levels of pesticides residues on leaves collected at different testing intervals.

Data clearly show no significant difference in overall mean A.D. % between Pyriproxyfen (83.72 % and 87.57 %) and the control (87.65 %), where both recorded the least A.D. achieved. However, remarkable and significant increase in overall mean A.D. % values ranged between 93.71 % and 90.43 % was recorded for Chromafenozide (200 ml/200 L) and Ethoxazole (25 ml/200 L), respectively.

Generally, the aforementioned data (Tables 46-48) revealed that the tested compounds except Pyriproxyfen induced the 4th larvae to exhibit increasing A.D. Such results confirms those obtained by **Radwan** et al. (1986) for S. littoralis due to the action of diflubenzuron and triflumuron throughout th larval period from the 4th instar to 6th instar. On the other hand, our data also agree with data of **El-Dessouki** and **Omar** (2000) where they found that the high concentration (37.5 ppm) of the JHM Pyriproxyfen when fed as treated

leaves to the 4th and 6th larval instar of Agrotis ipsilon resulted in a significant reduction in AD % when compared with Control.

4.3.5- Efficiency of Conversion of Ingested-food (E.C.I.):

The ability of insect to utilize food for growth is measured by the efficiency of conversion of ingested food (E.C.I.), together with the efficiency of conversion of digested food (E.C.D.) to body substances.

Data in Tables (49, 50 & 51) demonstrate the effect of feeding cotton leafworm 4th instar larvae of the three different strains for 48 h on insecticide- treated leaves, on the efficiency of conversion of ingested food (E.C.I.) to biomass. Data concerning early-season strain (Table, 49) show that all insecticides tested at the 1st testing interval (0-1 day) except Pyriproxyfen resulted in remarkable reduction in E.C.I. and recording reduction percentage ranged between -18.28 and -26.86. In contrast Pyriproxyfen recorded highly pronounced increase of +68.08 and +39.98 % in E.C.I. values at both rates tested.

Considering the bioresidual effect of th tested insecticides on E.C.I. values, it was obvious that the decrease in E.C.I. values decline with time elapse or/and the degradation of insecticide residues on treated leaves. For instance, it declined from -26.86 and -22.83; -22.78 and -21.07; -18.55 and -18.28 % at the 1st testing interval (0-1 day) to reach -9.61 and -8.29; -3.57 and -2.76; -2.08 and -2.04 % at the 4th testing interval (21-22 day) for Chromafenozide (400 and 200 ml/200 L), Azadirachtin (100 and 50 ml/200 L) and Ethoxazole (25 and 12.5 ml/200 L), respectively. On the other hand, the considerable percent increase in E.C.I. value recorded for Pyriproxyfen treatments at the 1st testing interval declined with time elapse and recorded slight percent decrease in E.C.I. values at the 4th testing interval.

Comparison on the basis of overall mean E.C.I. values within different testing intervals indicate that E.C.I. values in all treatments except Pyriproxyfen were slightly lower and were insignificant when

field strain larvae fed for 48 h on treated leaves, collected at indicated day posttreatment, followed by 5 Table (49): Efficiency of conversion of ingested food to body tissues (E.C.I.) for groups of early season S. littoralis days feeding on untreated leaves.

		Col	Conversion efficiency of ingested food to body tissues (E.C.I.)	efficiency	of ingest	ed food to	body tis	sues (E.C	(J.T.)	
	Kate/			at the	at the indicated tested intervals	tested in	tervals			
Insecticides	feddan	1st in	1st interval	2nd in	2nd interval	3rdin	3rdinterval	4th in	4th interval	1
	(ml.)	0-1	0-1 day	7-8	7-8 day	14-1	14-15 day	21-2	21-22 day	Mean ± S.D.
		E.C.I.	%	E.C.I.	%	E.C.I.	%	E.C.I.	%	
		%	change	%	change	%	change	%	change	
Pyriproxyfen	300	30.79	+60.45	24.49	+9.44	24.48	+19.82	19.66	-8.75	24.85±4.56 a
(10 % EC)	150	26.50	+38.09	24.35	+18.78	19.79	-2.25	19.45	-4.89	22.57±3.42 ab
Chromafenozide	400	15.78	-17.77	16.19	-21.02	18.72	-18.49	19.12	-6.50	17.45±1.71 c
(5 % EC)	200	16.45	-14.28	17.89	-12.73	18.81	-7.93	19.41	-5.09	18.14±1.29 c
Azadirachtin	100	15.97	-16.78	17.29	-15.66	18.49	-9.49	20.43	-0.09	18.04±1.89 c
(4.5 % EC)	50	16.02	-16.52	19.19	-6.39	19.82	-3.08	20.43	-0.09	18.86±1.96 bc
Ethoxazole	25	17.01	-11.36	19.18	-6.44	19.95	-7.55	20.10	-1.71	19.06±1.24 bc
(10 % SC)	12.5	17.14	-10.68	19.57	-4.54	19.70	-3.57	20.12	-0.78	19.13±1.35 bc
Control	í	19.19		20.50		20.43		20.45	ı	20.14±0.63 bc
L.S.D.0.05:										3.38

Mean followed by the same letter are not significantly different (P = 0.05) according to Duncan's (1955) multiple

strain larvae fed for 48 h on treated leaves, collected at indicated day posttreatment, followed by 5 days Table (50): Efficiency of conversion of ingested food to body tissues (E.C.I.) for groups of late season S. littoralis field feeding on untreated leaves.

	Rafe/	Cor	iversion e	fficiency at the i	of ingestendicated	iciency of ingested food to body at the indicated tested intervals	Conversion efficiency of ingested food to body tissues (E.C.I.) at the indicated tested intervals	sues (E.C	.I.)	
	feddan	1st interval	terval	2nd interval	terval	3rdinterval	erval	4th interva	terval	Mean ± S.D.
Insecticides	(m].)	0-1 day	day	7-8 day	day	14-15 day	day	21-22 day	day	-
		E.C.I.	%	E.C.I.	%	E.C.I.	%	E.C.I.	%	
		%	change	%	change	%	change	%	change	
Dyrinroxyfen	300	17.02	-11.31	18.87	-7.95	19.67	-3.72	22.89	+11.93	19.61±2.45 a
1 July 10 % FC)	150	17.01	-11.36	18.77	-8.44	19.17	-6.17	24.93	+21.91	19.97±3.44 a
Chromafenozide	400	18.31	-4.59	19.96	-2.63	20.14	-1.42	25.91	+26.69	21.08±3.32 a
(5 % FC)	200	18.13	-5.52	19.96	-2.63	20.01	-2.06	25.99	+27.91	21.02±3.42 a
A zodinochtin	100	17 99	-6.25	18.43	-10.09	19.86	-2.74	25.43	+24.35	20.43±3.49 a
Azaunachum	20	17.17	-10.53	19.41	-5.32	19.51	-4.50	24.93	+21.91	20.25±3.30 a
Fthoxazole	25	16.88	-12.04	17.41	-15.07	18.08	-11.50	25.72	+25.77	19.52±4.16 a
(10 % SC)	12.5	15.97	-16.78	16.46	-19.76	20.01	-2.06	26.31	+28.66	19.68±4.77 a
Control		19.19		20.50	ī	20.43		20.45	,	20.14±0.63 a
I S D s s z .										4.92
L.S.D.0.03 ·								1		

Mean followed by the same letter are not significantly different (P = 0.05) according to Duncan's (1955) multiple range test.

strain larvae fed for 48 h on treated leaves, collected at indicated day posttreatment, followed by 5 days Table (51): Efficiency of conversion of ingested food to body tissues (E.C.I.) for groups of S. littoralis laboratory feeding on untreated leaves.

	Rate/	CO	Conversion efficiency of ingested food to body tissues (E.C.I.)	efficiency	of ingest	ed food t	o body tis	sues (E.C	(T.T.)	
	12000			at the	at the indicated tested intervals	tested in	tervals			
Insecticides	feddan		1st interval	2nd ir	2nd interval	3rdin	3rdinterval	4th in	4th interval	
	(ml.)	0-1	0-1 day	7-8	7-8 day	14-1	14-15 day	21-2	21-22 day	Mean ± S.D.
	æ)	E.C.I.	%	E.C.I.	%	E.C.I.	%	E.C.I.	%	
		%	change	%	change	%	change	%	change	
Pyriproxyfen	300	31.66	+64.98	26.26	+28.09	24.17	+18.31	20.23	-1 08	25 58+4 25 2
(10 % EC)	150	27.56	+43.62	25.47	+24.24	26.07	-1.76	19 39	-5 18	27.50±1.20 a
Chromafenozide	400	12.64	-34.13	13.84	-34.49	14.65	-28 20	17.70	12.01	14.72.00.8
O3 % S)	200	12.40	07 00	16.40			77.07	71.17	10.01	14.73±2.20 d
(2000)	7007	13.40	-29.78	16.40	-20.0	17.86	-12.57	19.47	-4.79	16.80±2.45 bcd
Azadirachtin	100	14.18	-26.11	14.99	-26.88	15.09	-26.14	15.95	-22.01	15 05±0 72 cd
(4.5 % EC)	50	15.98	-16.73	16.83	-17.90	17.57	-13.99	19.60	-4 16	17 49±1 55 bod
Ethoxazole	25	16.09	-16.15	19.13	-6.68	19.21	-5.97	19.38	-5 23	18 45+1 76 bod
(10 % SC)	12.5	16.99	-5.73	19.47	-5.02	19.61	-4.01	19.63	-4 01	18 02±1 20 15
Control	ı	19.19	ï	20.50	ı	20.43		20.45	10:1	20 14±0 63 h
L.S.D.0.05:										3 57

Mean followed by the same letter are not significantly different (P = 0.05) according to Duncan's (1955) multiple

compared with control. However, Pyriproxyfen treatments only exhibited significantly higher E.C.I. values when compared with all other treatments including control.

Regarding the response of late-season strain larvae, data in Table (50) indicate no significant variations between different treatment and control based on overall mean E.C.I. values. However, detailed data at different testing intervals elucidate that the level of insecticides residues on leaves at the 4th testing interval (21-22 day) was low and not effective enough to affect the E.C.I. values specially against the tolerant larvae of the late-season strain. Furthermore, the level of residues at such testing interval (21-22 day) exhibited stimulatory effect and increase the E.C.I. values by percentages reached its maximum (+25.77 % and +28.66 %) for Ethoxazole at rate of 25 and 12.5 ml/200 L, while the minimum E.C.I. values of +11.93 and +21.91 % were recorded for Pyriproxyfen at rate of 300 and 150 ml/200 L, respectively.

As for the response of the laboratory strain larvae (Table, 51) it behaved almostly similar as those of the early season strain, where Pyriproxyfen at both rates recorded significantly the highest overall mean E.C.I. values (25.58 and 24.62 %), while Chromafenozide at 400 ml/200 L recorded significantly the least overall mean E.C.I. values (14.73 %).

4.3.6- Efficiency of Conversion of Digested-food (E.C.D.):

The second indicator of food utilization is the efficiency of conversion of digested food (E.C.D.) which is sometimes called "Net Growth Efficiency" (or "Metabolic Efficiency"). It estimates the percentage of assimilated food to biomass (Slansky and Scriber, 1985).

Data of E.C.D. shown in Table (52) indicate that at the 1st testing interval the initial effect of the tested compounds on early-season strain larvae was manifested by reduction percentages ranged between

field strain larvae fed for 48 h on treated leaves, collected at indicated day posttreatment, followed by 5 Table (52): Efficiency of conversion of digested food to body tissues (E.C.D.) for groups of early season S. littoralis days feeding on untreated leaves.

	Rate/	Ŝ	nversion	efficiency	of ingest	ed food t	Conversion efficiency of ingested food to body tissues (E.C.I.)	sues (E.C	(.I.)		
	Coddo			מו וווב	at the mulcated tested intervals	nestea in	itervais				
Insecticides	reddan	LST III	1st interval	2nd ii	2nd interval	3rdin	3rdinterval	4 <i>th</i> in	4th interval		
	(ml.)	0-1	0-1 day	7-8	7-8 day	14-1	14-15 day	21-2	21-22 day	Mean ± S.D.	
		E.C.D.	%	E.C.D.	%	E.C.D.	%	E.C.D.	%		
		%	change	%	change	%	change	%	change		
Pyriproxyfen	300	37.33	+68.08	29.02	+27.06	27.97	+21.65	21.06	-10.46	28.84±6.67.9	I
(10 % EC)	150	31.09	+39.98	28.17	+23.34	21.99	-4.43	21.41	-8.97	25 66+4 74 ah	T
Chromafenozide	400	16.24	-26.86	17.67	-22.64	20.69	-10.08	21.26	-9.61	18 69+2 40 c	T
(5 % EC)	200	17.14	-22.83	19.27	-15.63	20.86	-9.34	21.57	-8 29	19 71+1 9 60	T
Azadirachtin	100	17.15	-22.78	19.12	-16.28	20.70	-10.04	22.68	-3.57	19 91+7 35 6	T
(4.5 % EC)	50	17.53	-21.07	21.21	-7.13	22.42	-2.56	22.87	-2.76	21 01+2 43 bc	T
Ethoxazole	25	18.09	-18.55	21.21	-7.53	22.04	-4.22	23.03	-2.08	21 09+2 14 bc	1
(10 % SC)	12.5	18.15	-18.28	21.37	-6.44	22.42	-2.56	23.04	-2.04	21 24+2 18 bc	1
Control	1	22.21	1	22.84		23.01		23.52		22.89±0 54 bc	T
L.S.D.0.05:										4.78	_

Mean followed by the same letter are not significantly different (P = 0.05) according to Duncan's (1955) multiple

-18.28 % and -26.86 % for all tested pesticides except for Pyriproxyfen where a remarkable increase in E.C.D. relative to control reached +68.08 % and +39.98 % was recorded at both tested rates of 300 and 150 ml/200 L, respectively. However, such increase in E.C.D. value continued within the 2nd and 3rd testing intervals but was of gradually lower magnitude and then switch at the 4th testing interval to record low reduction percentage in E.C.D. of -10.46 and -8.97 % for both rates of Pyriproxyfen.

As for the late-season strain larvae (Table, 53), the E.C.D. was moderately affected for all insecticidal treatments and within most testing intervals recording percent reduction ranged between -10.72 % and -21.25 %; -2.93 % and -17.43 %; -0.3 % and -33.89 % for the 1st (0-1 day), the 2nd (7-8 day) and the 3rd (14-15 day) testing interval, respectively; while an increase percent ranged between +20.11 % and +29.21 % was recorded within the 4th testing interval (21-22 day).

Comparison based on overall mean E.C.D. revealed insignificant slight variations between E.C.D. values of different treatments including the control.

The response of the laboratory-strain larvae in relation to the obtained E.C.D. value was recorded in Table (54). The larvae performed similarly as the early-season strain larvae where reduction in E.C.D. values ranged between -41.29 % and -22.11 % for Chromafenozide, Azadirachtin and Ethoxazole versus an increase in E.C.D. value reached +76.86 % and +47.55 % for Pyriproxyfen at the 1st (0-1 day) testing interval.

Following the residual effect of pesticides residues after feeding the larvae on treated leaves collected at different time intervals after spray, it was obvious that the percent increase in E.C.D. values of Pyriproxyfen treatments was declined gradually with time elapse and switch to negligible decrease at the 4th (21-22 day) testing interval. Likewise, the highly decrease percent in E.C.D. value recorded at the

field strain larvae fed for 48 h on treated leaves, collected at indicated day posttreatment, followed by 5 Table (53): Efficiency of conversion of digested food to body tissues (E.C.D.) for groups of late season S. littoralis days feeding on untreated leaves.

	Rate/	రి	nversion	efficiency at the	iciency of ingested food to body at the indicated tested intervals	ed food t	Conversion efficiency of ingested food to body tissues (E.C.I.) at the indicated tested intervals	snes (E.	C.I.)	
Insecticides	feddan	1st in	1st interval	2nd it	2nd interval	3rdin	3rdinterval	4th in	4th interval	
	(ml.)	0-1	0-1 day	7-8	7-8 day	14-1;	14-15 day	21-2	21-22 day	Mean ± S.D.
		E.C.D.	%	E.C.D.	%	E.C.D.	%	E.C.D.	%	
		%	change	%	change	%	change	%	change	
Pyriproxyfen	300	19.03	-14.32	20.99	-8.09	22.23	-33.89	28.25	+20 11	22 62+3 70 0
(10 % EC)	150	18.97	-14.59	20.65	-9.59	20.59	-10.52	29.34	+24 75	22.32-3.12 a
Chromafenozide	400	18.93	-14.77	21.65	-5.21	22.94	-0.30	29.99	+27.51	21 63+1 90 a
(5 % EC)	200	19.83	-10.72	22.17	-2.93	22.95	-0.26	30.14	+28.15	23 77+4 45 0
Azadirachtin	100	19.39	-12.69	20.04	-12.26	20.79	-9.65	29.38	+24 92	22.77.11.13 a
(4.5 % EC)	50	18.75	-15.58	21.41	-6.26	21.58	-6.21	28.79	+22.41	22 63+4 31 9
Ethoxazole	25	18.57	-16.43	18.91	-17.21	20.64	-10.29	29.89	+27.08	22 00+5 34 a
(10 % SC)	12.5	17.49	-21.25	18.86	-17.43	22.77	-10.43	30.39	+29.21	22.38±5 79 a
Control	1	22.21	•	22.84	Ĭ	23.01	,	23.52		22.98±0 54 a
L.S.D.0.05:										6.19

Mean followed by the same letter are not significantly different (P = 0.05) according to Duncan's (1955) multiple range test.

strain larvae fed for 48 h on treated leaves, collected at indicated day posttreatment, followed by 5 days Table (54): Efficiency of conversion of digested food to body tissues (E.C.D.) for groups of S. littoralis laboratory feeding on untreated leaves.

	Rate/	Cor	iversion 6	efficiency at the i	Conversion efficiency of ingested food to body tissues (E.C.I.) at the indicated tested intervals	ed food to	body tis	sues (E.C	(T)	
1	feddan	1st in	1st interval	2nd in	2nd interval	3rdinterval	terval	4th interval	terval	Man - Ch
Insecticides	(ml.)	0-1 day	day	7-8	7-8 day	14-15	14-15 day	21-22 day	2 day	Mean = 5.D.
	10	E.C.D.	%	E.C.D.	%	E.C.D.	%	E.C.D.	%	
		%	change	%	change	%	change	%	change	
Pyriproxyfen	300	31.28	+76.86	36.68	+60.59	28.82	+25.25	23.12	-1.70	31.97±7.39 a
(10 % EC)	150	32.77	+47.55	31.14	+36.34	22.48	-2.30	21.26	-9.61	26.91±5.88 ab
Chromafenozide	400	13.04	-41.29	14.89	-34.81	16.69	-27.47	20.24	-13.73	16.23±3.09 d
(5 % EC)	200	13.97	-37.10	17.63	-22.81	19.12	-16.91	22.28	-5.27	18.25±3.45 cd
Azadirachtin	100	14.70	-33.81	16.12	-29.42	16.22	-29.51 17.84	17.84	-24.15	16.22±1.28 d
(4.5 % EC)	50	16.85	-24.13	16.24	-28.89	17.07	-25.82	21.74	-7.57	17.97±2.53 cd
Ethoxazole	25	17.16	-22.74	21.28	-6.83	21.42	-6.19	21.91	-6.85	20.44±2.20 cd
(10 % SC)	12.5	17.30	-22.11	21.81	-4.51	21.98	-4.48	22.03	-6.34	20.78±2.32 cd
Control	i	22.21	1	22.84	,	23.01	,	23.52	•	16.22±0.54 d
L.S.D. _{0.05} :										5.50

Mean followed by the same letter are not significantly different (P = 0.05) according to Duncan's (1955) multiple

1st (initial) time interval for the other three insecticides gradually decline with time elapse to reach the least percent decrease in E.C.D. values at the 4th testing interval.

Comparison based on overall mean E.C.D. values elucidate significantly the superiority of Pyriproxyfen both treatment and recording the highest E.C.D. values. Other treatments including the control came significantly next without any significant variations between each other.

The ability of insects to utilize food for growth is measured by the efficiency of conversion of ingested food (E.C.I.), together with efficiency of conversion of digested food (E.C.D.) to body substances. In the present study these two criteria (E.C.I. and E.C.D.) were decreased with most of pesticides tested (Chromafenozide, Azadirachtin and Ethoxazole). Similarly, previous studies indicate that different IGRs reduced E.C.I. and E.C.D. of various insect species such as diflubenzuron and triflumuron against *S. littoralis* (Radwan et al., 1986), fenarimol against *S. littoralis* (Farag, 1991) and fenoxycarb against *S. gregaria* (Ismail, 1995).

El-Shazly (1993) indicated that the E.C.I. will vary with the digestibility of food (A.D.) and the proportional amount of digestible portion of the food which is converted to body substance and metabolized for energy to maintain life. Furthermore, Waldbauer (1968) cited that the (E.C.I.) would raise and fall with the (A.D.). This statement extends to the results obtained in this study. Though the reduced (E.C.I.) and (E.C.D.) obtained may result from diversion of energy from production of biomass to overcoming the effect of treatment.

Several studies indicates that some IGRs action on E.C.I. and E.C.D. depended on the dose level, in addition to the nature of the chemical and insect species. In this respect, **Abo-Elghar** (1993) found

that abamectin at 10 and 25 ppm caused a significant decrease in E.C.I. and E.C.D., whereas all concentrations tested of the JHM fenoxycarb and buprofezin caused increase in the E.C.I. and E.C.D. over control. Furthermore, **Abo-Elghar (1994)** found that several herbicides such as fluazifop, bentazone, ametryne, flamprop and ioxynil reduced the E.C.I. and E.C.D. below those of untreated control in *S. littoralis*.